NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.

The Rules & Regulations of the City of New York
is current through June 30, 2009.

Introductory Notes:

Article 175 applies to all radiation equipment and radioactive material within the jurisdiction of the New York City Department of Health. Sections of Article 175 set forth under the heading, "Radiation Equipment," contain provisions for all radiation equipment and general and additional radiation control requirements applicable to specific radiation equipment. Sections of Article 175 set forth under the heading, "Radioactive Materials," contain provisions for all radioactive materials and general and additional radiation control requirements applicable to specific radioactive materials. Article 175 aims to protect the public generally, as well as workers in certain radiation installations, from the hazards inherent in the use of ionizing radiation. The Article is intended to serve as a framework for coordination of radiation control activities with the U.S. Atomic Energy Commission, the U.S. Food and Drug Administration, the N.Y. State Department of Labor, the N.Y. State Department of Health, the N.Y. State Atomic Energy Council, the N.Y. State Department of Environmental Conservation, and with other state and federal agencies, as well as with other N.Y. City agencies.

The present version of Article 175 follows the format of Part 16 of the New York Sanitary Code, relating to ionizing radiation, and is intended to be consistent with the standards included in Part 16 with certain standards deemed to be necessarily more restrictive. Article 175 is also intended to be compatible with safety standards of the U.S. Atomic Energy Commission in Title 10. Code of Federal Regulations; identical to standards established by the U.S. Food and Drug Administration in Title 21, Part 1020, of the Code of Federal Regulations; as well as consistent with corresponding standards for radiation control in the N.Y. State Industrial Code.

The present version of Article 175 continues many of the provisions which have been in such Articles since March 23, 1959, particularly the emphasis on the role of the radiation safety officer, safety standards identical with national recommendations, and information requirements for the location and use of all radiation sources. Article 175 now replaces the former registration on radiation equipment, such as medical x-ray units, with a permit system. Most of the former Notes in the previous Article 175 have been included within the text

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of the various sections. Reliance upon the recommendations in specific reports of the National Council on Radiation Protection and Measurements has been eliminated, with specific standards being included in appropriate sections of the Article. Such recommendations are to be used in complying with the requirements in Article 175, where appropriate. Guidelines to provide interpretations of several sections of Article 175 are to be issued by the Commissioner of Health after publication in The City Record.

Article 175 was first adopted in 1959, completely amended in 1962, and thereafter amended several times since. The present extensively revised version was adopted by the Board of Health on June 19, 1973 to provide greater uniformity with State and Federal requirements and was again repealed and reenacted on June 29, 1973 to incorporate necessary further amendments and revisions. Under the present Article 175, all radiation sources in the City must be obtained under either a permit or license or be specifically exempt. General licensing of small quantities of radioactive material is also provided for. However, the Department will be fully and currently informed of all radiation hazards within the City through its enforcement of the present Article 175.

This Article was amended by resolution adopted on June 19, 1975, to add a new table six (6) to Section Heading 175.117.

Article 175 was further amended by resolution adopted on September 18, 1975 which added a new section 175.70 to regulate equipment and procedures in the use of mammographic radiography for the detection of breast cancer and to prevent radiation hazards both to the person operating the applicable x-ray equipment and to the patient.

Article 175 was further amended by resolution adopted on December 6, 1976 to add new sections 175.118 through 175.124 which provide microwave radiation safety requirements designed to prevent the biologically deleterious effects of such radiation from improperly constructed, serviced or used microwave ovens. Such sections were intended to provide uniformity with applicable Federal performance standards governing microwave ovens contained in Section 1030.10 of Part 1030 of Subchapter J--"Radiological Health" of Chapter 1 of Title 21 of the Federal Code of Rules and Regulations (21 CFR Section 1030.10) promulgated pursuant to the applicable provisions of the Radiation Control for Health and Safety Act of 1968 (42 U.S.C. 263f, 263g, and 263n). Such sections were not intended to institute a regular program of inspection of microwave ovens used in homes.

The Section Headings for Section 175.09 and 175.10 were amended by resolution adopted on December 15, 1977 to conform them with the provisions contained in such sections as amended by resolution of the same date.

The Table of Section Headings was amended by resolution adopted on February 26, 1981 to conform them with the provisions contained in such sections as added or amended by resolution of the same date.

The Section Heading of Section 175.104 was amended by resolution adopted in

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October 23, 1986 to have the heading conform to changes made in the Section itself on the same date.

Section 175.13 was repealed by resolution adopted on June 26, 1990 to remove a maximum fee that is no longer imposed by New York State.

The repeal and reenactment of Article 175 was adopted by resolution by the Board of Health on June 7, 1994 to incorporate updated basic radiation protection standards as codified in federal regulations in all applicable provisions of the Article, to remove redundant language, inconsistencies and outdated provisions, to ensure a consistent numbering format, incorporate the use of metric and Système Internationale units, provide gender neutral references and to incorporate other provisions to maintain compatibility with federal regulations.

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ARTICLE 175. RADIATION CONTROL.
GENERAL PROVISIONS

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§ 175.01. Applicability and inapplicability, communications.

(a) Applicability. (1) Except as provided in § 175.01(b), this Code applies to any person who sells, transfers, assembles, receives, produces, possesses, or uses any radiation source in this City.

(b) Inapplicability. (1) This Code does not apply to any person with respect to any radiation source subject to regulation, as provided for by law, by the New York State Department of Labor. This exclusion does not apply to:

(A) the use of such sources in places where the general public may be exposed; or

(B) to persons with respect to radiation sources used at industrial or commercial establishments for the application of radiation to human beings.

(2) This Code does not apply to any common or contract carrier or any shipper operating within this City to the extent that such carrier or shipper is subject to regulation as provided for by law by the U.S. Department of Transportation or other agencies of the United States or agencies of the State or City of New York, except for compliance with provisions relating to transportation of radioactive materials set forth in § 175.105.

(c) Communications. (1) Except as otherwise provided for in this Code, or as authorized by the Department, all applications, notifications, reports or other communications filed pursuant to this Code shall be addressed to the Department at:

Bureau of Radiological Health 2 Lafayette Street, 11th Floor New York, New York 10007

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

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Subd. (c) amended City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

<General Materials (GM) - References, Annotations, or Tables>

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GENERAL PROVISIONS

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§ 175.02. Definitions.

(a) As used in this Code, the following definitions shall apply:

(1) "A1in. means the maximum activity of special form radioactive material permitted in a Type A package. "A2in. means the maximum activity of radioactive material, other than special form, LSA and SCO material, permitted in a Type A package. These values are either listed in Table A-1, Appendix A of § 175.105 of this Code or may be derived in accordance with the procedure prescribed in such Appendix A.

(2) "Absorbed dose" means the energy imparted by ionizing radiation per unit mass of irradiated material. The units of absorbed dose are the gray (Gy) and the rad.

(3) "Accelerator" means any machine capable of accelerating electrons, protons, deuterons, or other charged particles in a vacuum and of discharging the resultant particulate or other radiation into a medium at energies usually in excess of 1 MeV. For purposes of this definition, "particle accelerator" is an equivalent term.

(4) "Accelerator-produced material" means any material made radioactive by a particle accelerator.

(5) "Accessible surface" means the external surface of the enclosure or housing provided by the manufacturer.

(6) "Activity" means the rate of disintegration or transformation or decay of radioactive material. The units of activity are the becquerel (Bq) and the curie (Ci).

(7) "Added filtration" means any filtration which is in addition to the inherent filtration.

(8) "Adult" means an individual 18 or more years of age.

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(9) "Agreement State" means any State with which the U.S. Nuclear Regulatory Commission or the U.S. Atomic Energy Commission has entered into an effective agreement under subsection 274b. of the Atomic Energy Act of 1954, as amended (73 Stat. 689).

(10) "Airborne radioactive material" means any radioactive material dispersed in the air in the form of dusts, fumes, particulates, mists, vapors, or gases.

(11) "Airborne radioactivity area" means a room, enclosure, or area in which airborne radioactive materials exist in concentrations:

(i) in excess of the derived air concentrations (DACs) specified in Table 1, Appendix B of § 175.03 of this Code, or

(ii) to such a degree that an individual present in the area without respiratory protective equipment could exceed, during the hours an individual is present in a week, an intake of 0.6 percent of the annual limit on intake (ALI) or 12 DAC-hours.

(12) "Aluminum equivalent" means the thickness of aluminum (type 1100 alloy) affording the same attenuation, under specified conditions, as the material in question. The nominal chemical composition of type 1100 aluminum alloy is 99.00 percent minimum aluminum, 0.12 percent copper.

(13) "Annual limit on intake" (ALI) means the derived limit for the amount of radioactive material taken into the body of an adult worker by inhalation or ingestion in a year. ALI is the smaller value of intake of a given radionuclide in a year by the reference man that would result in a committed effective dose equivalent of 0.05 Sv (5 rem) or a committed dose equivalent of 0.5 Sv (50 rem) to any individual organ or tissue. ALI values for intake by ingestion and by inhalation of selected radionuclides are given in Table 1, Columns 1 and 2, of Appendix B of § 175.03 of this Code.

(14) "Area of use" means a portion of a physical structure, or a specified out-of-doors location, that has been set aside for the purpose of receiving, producing, using, or storing radioactive material.

(15) "As low as is reasonably achievable" (ALARA) means making every reasonable effort to maintain exposures to radiation as far below the dose limits in this Code as is practical, consistent with the purpose for which the licensed or registered activity is undertaken, taking into account the state of technology, the economics of improvements in relation to state of technology, the economics of improvements in relation to benefits to the public health and safety, and other societal and socioeconomic considerations, and in relation to utilization of nuclear energy and licensed or registered sources of radiation in the public interest.

(16) "Assembler" means any person engaged in the business of assembling, replacing, or installing one or more components into an x-ray system or subsystem. The term includes the owner of an x-ray system or such person's employee or agent who assembles components into an x-ray system that is subsequently used to provide professional or commercial services.

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(17) "Attenuation block" means a block or stack, having dimensions 20 centimeters by 20 centimeters by 3.8 centimeters (8 inches by 8 inches by 1.5 inches), of type 1100 aluminum alloy or other materials having equivalent attenuation.

(18) "Authorized user" means an individual who is identified as an authorized user on a Department, Agreement State, or U.S. Nuclear Regulatory Commission license that authorizes the use of radioactive material or who is named as an authorized user on a certified registration issued by the Department.

(19) "Automatic exposure control" means a device which automatically controls one or more technique factors in order to obtain a required quantity of radiation at a preselected location(s).

(20) "Background radiation" means radiation from cosmic sources; naturally occurring radioactive material, including radon (except as a decay product of source or special nuclear material); and global fallout as it exists in the environment from the testing of nuclear explosive devices or from past nuclear accidents such as Chernobyl that contribute to background radiation and are not under the control of the licensee. "Background radiation" does not include radiation from any regulated sources of radiation.

(21) "Barrier" [see "Protective Barrier"].

(22) "Beam axis" means a line from the source through the centers of the x-ray fields.

(23) "Beam-limiting device" means a device which provides a means to restrict the dimensions of the x-ray beam.

(24) "Beam monitoring system" means a system designed to detect and measure the radiation present in the useful beam.

(25) "Becquerel" (Bq) means the SI unit of activity. One becquerel is equal to 1 disintegration (d) or transformation (t) per second (d-s-1 or t-s-1).

(26) "Bioassay" means the determination of kinds, quantities or concentrations, and, in some cases, the locations of radioactive material in the human body, whether by direct measurement, in vivo counting, or by analysis and evaluation of materials excreted or removed from the human body. For purposes of this Code, "radiobioassay" is an equivalent term.

(27) "Brachytherapy" means a method of radiation therapy in which sealed sources are utilized to deliver a radiation dose at a distance of up to a few centimeters, by surface, intracavitary, or interstitial application. Brachytherapy includes radiation therapy using electronic remote after-loading devices.

(28) "Breast equivalent phantom" means a device which contains test objects of various specified dimensions as speck sets, masses and fibers representing low density areas and microcalcifications related to the imaging of breast lesions and

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which can be imaged by a mammographic x-ray system to visualize such test objects.

(29) "Byproduct material" means:

(i) Any radioactive material, except special nuclear material, yielded in or made radioactive by exposure to the radiation incident to the process of producing or utilizing special nuclear material; and

(ii) The tailings or wastes produced by the extraction or concentration of uranium or thorium from ore processed primarily for its source material content, including discrete surface wastes resulting from uranium or thorium solution extraction processes. Underground ore bodies depleted by these solution extraction operations do not constitute "byproduct material" within this definition.

(30) "Calendar quarter" means not less than 12 consecutive weeks nor more than 14 consecutive weeks. The first calendar quarter of each year shall begin in January and subsequent calendar quarters shall be so arranged such that no day is included in more than one calendar quarter and no day in any one year is omitted from inclusion within a calendar quarter. No licensee or registrant shall change the method observed by him/her of determining calendar quarters for purposes of this Code except at the beginning of a year.

(31) "Calibration" means the determination of:

(i) the response or reading of an instrument relative to a series of known radiation values over the range of the instrument, or

(ii) the strength of a source of radiation relative to a standard.

(32) "C-arm x-ray system" means an x-ray system in which the image receptor and x-ray tube housing assembly are connected by a common mechanical support system in order to maintain a desired spatial relationship. This system is designed to allow a change in the projection of the beam through the patient without a change in the position of the patient.

(33) "Carrier" means a person engaged in the transportation of passengers or property by land or water as a common, contract, or private carrier, or by civil aircraft.

(34) "Certified components" means components of x-ray systems which are subject to regulations promulgated under Public Law 90-602, the Radiation Control for Health and Safety Act of 1968.

(35) "Certified registration" means a registration for any therapeutic radiation machine issued by the Department upon review and approval of an application submitted pursuant to this Code.

(36) "Certified system" means any x-ray system which has one or more certified components.

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(37) "Certified Radiation Equipment Safety Officer" means an individual who holds an unexpired certificate as a radiation equipment safety officer issued by the New York State Department of Health.

(38) "CFR" means Code of Federal Regulations.

(39) "Chelating agent" means amine polycarboxylic acids, hydroxycarboxylic acids, gluconic acid, and polycarboxylic acids.

(40) "City" means the City of New York.

(41) "Class" means a classification scheme for inhaled material according to its rate of clearance from the pulmonary region of the lung. Materials are classified as D, W, or Y, which applies to a range of clearance half-times: for Class D, Days, of less than 10 days; for Class W, Weeks, from 10 to 100 days; and for Class Y, Years, of greater than 100 days. For purposes of this Code, "lung class" and "inhalation class" are equivalent terms.

(42) "Coefficient of Variation," or "C" means the ratio of the standard deviation to the mean value of a population of observations. It is estimated using the following equation:

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where

s = estimated standard deviation of the population.

X = mean value of observations in sample.

Xi = ith observation in sample.

n = number of observations in sample.

(43) "Collective dose" means the sum of the individual doses received in a given period of time by a specified population from exposure to a specified source of radiation.

(44) "Collimator " means a device by which a radiation beam is restricted in size.

(45) "Commissioner" means the Commissioner of Health and Mental Hygiene of the City of New York.

(46) "Committed dose equivalent" (HT,50) means the dose equivalent to organs or tissues of reference (T) that will be received from an intake of radioactive material by an individual during the 50-year period following the intake.

(47) "Committed effective dose equivalent" (HE,50) is the sum of the products of

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the weighting factors applicable to each of the body organs or tissues that are irradiated and the committed dose equivalent to each of these organs or tissues (HE,50 = <<SIGMA>> wT,HT,50).

(48) "Computed tomography" or "CT" means the production of a tomogram by the acquisition and computer processing of x-ray transmission data.

(49) "Cone" means a device used to indicate beam direction and to establish a minimum source-surface distance. It may or may not incorporate a collimator.

(50) "Contamination" means the presence in or on any animal, food, water supply, building or premises, body of water, municipal sewage disposal system, chattel or thing of a solid, liquid or gas emitting ionizing radiation which may constitute a danger to human beings.

(51) "Control panel" means that part of radiation equipment upon which is mounted the switches, knobs, pushbuttons, and other hardware necessary for manually setting the technique factors.

(52) "Conveyance" means: (i) "For transport by public highway or rail" any transport vehicle or large freight container;

(ii) "For transport by water" any vessel, or any hold, compartment, or defined deck area of a vessel including any transport vehicle on board the vessel; and

(iii) "For transport by aircraft" any aircraft.

(53) "Cooling curve" means the graphical relationship between heat units stored and cooling time.

(54) "Critical Group" means the group of individuals reasonably expected to receive the greatest exposure to residual radioactivity for any applicable set of circumstances.

(55) "Curie" means a unit of activity. One curie (Ci) is that quantity of radioactive material which decays at the rate of 3.7E+10 transformations per second (t-s-1).

(56) "Dead-man switch" means a switch so constructed that a circuit closing contact can be maintained only by continuous pressure on the switch by the operator.

(57) "Declared pregnant woman" means a woman who has voluntarily informed her employer, in writing, of her pregnancy and the estimated date of conception.

(58) "Decommission" means to remove a facility or site safely from service and reduce residual radioactivity to a level that permits--

(i) Release of the property for unrestricted use and termination of the license; or

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(ii) Release of the property under restricted conditions and the termination of the license.

(59) "Dedicated check source" means a radiation source that is used to assure the constant operation of a radiation detection or measurement device over several months or years.

(60) "Deep dose equivalent" (Hd), which applies to external whole body exposure, means the dose equivalent at a tissue depth of 1 centimeter (1000 mg/cm 2).

(61) "Department" means the New York City Department of Health and Mental Hygiene.

(62) "Depleted uranium" means the source material uranium in which the isotope uranium-235 is less than 0.711 weight percent of the total uranium present.

(63) "Derived air concentration" (DAC) means the concentration of a given radionuclide in air which, if breathed by the reference man for a working year of 2,000 hours under conditions of light work, results in an intake of one ALI. For purposes of this Code, the condition of light work is an inhalation rate of 1.2 cubic meters of air per hour for 2,000 hours in a year. DAC values are given in Table 1, Column 3, of Appendix B of § 175.03 of this Code.

(64) "Derived air concentration-hour" (DAC-hour) means the product of the concentration of radioactive material in air, expressed as a fraction or multiple of the derived air concentration for each radionuclide, and the time of exposure to that radionuclide, in hours. A licensee or registrant may take 2,000 DAC-hours to represent one ALI, equivalent to a committed effective dose equivalent of 0.05 Sv (5 rem).

(65) "Deterministic effect" [see "Nonstochastic effect"].

(66) "Diagnostic source assembly" means the tube housing assembly with a beam-limiting device attached.

(67) "Diagnostic type protective tube housing" means an x-ray tube housing so constructed that the leakage radiation at a distance of one meter from the tube housing does not exceed 2.58 E-5 C-kg-1 (100 milliroentgens) in one hour with a beam-limiting device attached and the tube operated at its leakage technique factors as specified by the manufacturer. Measurements may be averaged over an area of 100 cm 2 with no linear dimensions greater than 20 centimeters (8 inches).

(68) "Diagnostic x-ray system" means an x-ray system designed for irradiation of any part of the human body for the purpose of diagnosis or visualization.

(69) "Diaphragm" means a device or mechanism by which the radiation beam is restricted in size.

(70) "Distinguishable from background" means that the detectable concentration of

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a radionuclide is statistically different from the background concentration of that radionuclide in the vicinity of the site or, in the case of structures, in similar materials using adequate measurement technology, survey, and statistical techniques.

(71) "Dose" is a generic term that means absorbed dose, dose equivalent, effective dose equivalent, committed dose equivalent, committed effective dose equivalent, total organ dose equivalent, or total effective dose equivalent. For purposes of this Code, "radiation dose" is an equivalent term.

(72) "Dose equivalent (HT)" means the product of the absorbed dose in tissue, quality factor, and all other necessary modifying factors at the location of interest. The units of dose equivalent are the sievert (Sv) and rem.

(73) "Dose limits" means the permissible upper bounds of radiation doses established in accordance with this Code. For purposes of this Code, "limits" is an equivalent term.

(74) "Dose monitor unit" [See "Monitor unit".]

(75) "Dosimetry processor" means an individual or an organization that processes and evaluates individual monitoring devices in order to determine the radiation dose delivered to the monitoring devices.

(76) "Effective dose equivalent (HE)" means the sum of the products of the dose equivalent to each organ or tissue (HT) and the weighting factor (wT) applicable to each of the body organs or tissues that are irradiated (HE = << SIGMA>> wTHT).

(77) "Embryo/fetus" means the developing human organism from conception until the time of birth.

(78) "Entrance or access point" means any opening through which an individual or extremity of an individual could gain access to radiation areas or to licensed or registered radioactive materials. This includes entry or exit portals of sufficient size to permit human entry, irrespective of their intended use.

(79) "Entrance exposure rate" means the exposure per unit time at the point where the center of the useful beam enters the patient. For the purposes of this definition, "exposure" is defined in § 175.02(a)(80)(ii).

(80) "Equipment" means x-ray equipment.

(81) "Exclusive use" (also referred to in other regulations as "sole use" or "full load") means the sole use of a conveyance by a single consignor for which all initial, intermediate, and final loading and unloading are carried out in accordance with the direction of the consignor or consignee. The consignor and the carrier must ensure that any loading or unloading is performed by personnel having radiological training and resources appropriate for safe handling of the consignment. The consignor must issue specific instructions, in writing, for maintenance of exclusive use shipment controls, and include them with the shipping paper information

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provided to the carrier by the consignor.

(82) "Explosive material" means any chemical compound, mixture, or device which produces a substantial instantaneous release of gas and heat spontaneously or by contact with sparks or flame.

(83) "Exposure" means either:

(i) being exposed to ionizing radiation or to radioactive material; or

(ii) the quotient of dQ divided by dm where "dQ" is the absolute value of the total charge of the ions of one sign produced in air when all the electrons (negatrons and positrons) liberated by photons in a volume element of air having mass "dm" are completely stopped in air. The units of exposure are the coulomb per kilogram (C-kg-1) and the roentgen.

(84) "Exposure rate" means the exposure per unit of time.

(85) "External beam radiation therapy" means a method of radiation therapy utilized to deliver a radiation dose in which the source (sources) of radiation is (are) at a distance from the body. For the purposes of this Code "teletherapy" is an equivalent term.

(86) "External dose" means that portion of the dose equivalent received from any source of radiation outside the body.

(87) "Extremity" means hand, elbow, arm below the elbow, foot, knee, and leg below the knee.

(88) "Eye dose equivalent" means the external dose equivalent to the lens of the eye at a tissue depth of 0.3 centimeter (300 mg/cm 2).

(89) "Field emission equipment" means equipment which uses an x-ray tube in which electron emission from the cathode is due solely to the action of an electric field.

(90) "Filter" means material placed in the useful beam to absorb preferentially selected radiations.

(91) [Reserved]

(92) "Fissile material" means plutonium-239, plutonium-241, uranium-233, uranium-235 or any combination of these radionuclides. Fissile material means the fissile nuclides themselves, not material containing fissile nuclides. Unirradiated natural uranium and depleted uranium, and natural uranium or depleted uranium that has been irradiated in thermal reactors only are not included in this definition. Certain exclusions from fissile material controls are provided in 10 CFR 71.15

(93) "Fluoroscopic imaging assembly" means a subsystem in which x-ray photons produce a fluoroscopic image. It includes the image receptor(s) such as the image

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intensifier and spot-film device, electrical interlocks, if any, and structural material providing linkage between the image receptor and diagnostic source assembly.

(94) "Former U.S. Atomic Energy Commission (AEC) or U.S. Nuclear Regulatory Commission (NRC) licensed facilities" means nuclear reactors, nuclear fuel reprocessing plants, uranium enrichment plants, or critical mass experimental facilities where AEC or NRC licenses have been terminated.

(95) "Gantry" means that part of a radiation therapy system supporting and allowing movements of the radiation head about a center of rotation.

(96) "General purpose radiographic x-ray system" means any radiographic x-ray system which, by design, is not limited to radiographic examination of specific anatomical regions.

(97) "Generally applicable environmental radiation standards" means standards issued by the U.S. Environmental Protection Agency (EPA) under the authority of the Atomic Energy Act of 1954, as amended, that impose limits on radiation exposures or levels, or concentrations or quantities of radioactive material, in the general environment outside the boundaries of locations under the control of persons possessing or using radioactive material.

(98) "Gonad or gonadal shield" means a protective barrier for the ovaries or testes.

(99) "Gray" (Gy) means the SI unit of absorbed dose. One gray is equal to an absorbed dose of 1 joule per kilogram. One gray is equal to 100 rads.

(100) "Half-value layer (HVL)" means the thickness of specified material which, when introduced into the beam of a given path of radiation, reduces the exposure rate by one-half. In this definition, the contribution of all scattered radiation, other than any which might be present initially in the beam concerned, is deemed to be excluded.

(101) "Hazardous waste" means those wastes designated as hazardous by U.S. Environmental Protection Agency regulations in 40 CFR Part 261.

(102) "High radiation area" means an area, accessible to individuals, in which radiation levels could result in an individual receiving a dose equivalent in excess of 1 mSv (0.1 rem) in 1 hour at 30 centimeters (12 inches) from any source of radiation or from any surface that the radiation penetrates. For the purposes of this Code, rooms or areas in which diagnostic x-ray systems are used for healing arts purposes are not considered high radiation areas.

(103) "Human use" [see "Medical use"].

(104) "Image receptor" means any device, such as a fluorescent input phosphor or radiographic film, which transforms incident x-ray photons either into a visible image or into another form which can be made into a visible image by further transformation.

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(105) "Individual" means any human being.

(106) "Individual monitoring" means the assessment of:

(i) dose equivalent

(A) by the use of individual monitoring devices, or

(B) by the use of survey data; or

(ii) committed effective dose equivalent

(A) by bioassay, or

(B) by determination of the time-weighted air concentrations to which an individual has been exposed, that is, DAC-hours.

(107) "Individual monitoring devices" means devices designed to be worn by a single individual for the assessment of dose equivalent. For purposes of this Code, "personnel dosimeter" and "dosimeter" are equivalent terms. Examples of individual monitoring devices are film badges, thermoluminescent dosimeters (TLDs), pocket ionization chambers, and personal air sampling devices.

(108) "Inhalation class" [see "Class"].

(109) "Inherent filtration" means the filtration of the useful beam provided by the permanently installed components of the tube housing assembly.

(110) "Inspection" means an official examination or observation including, but not limited to, tests, surveys, and monitoring to determine compliance with rules, regulations, orders, requirements, and conditions of the Department.

(111) "Interlock" means a device arranged or connected such that the occurrence of an event or condition is required before a second event or condition can occur or continue to occur.

(112) "Internal dose" means that portion of the dose equivalent received from radioactive material taken into the body.

(113) "Kilo electron volt (keV)" means the energy equal to that acquired by a particle with one electron charge in passing through a potential difference of one thousand volts in a vacuum.

(114) "Kilovolt peak (kVp)" means the maximum value in kilovolts of the potential difference of a pulsating generator. When only one-half of the wave is used, the value refers to the useful half of the wave.

(115) "Lead equivalent" means the thickness of lead affording the same attenua-

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tion, under specified conditions, as the material in question.

(116) "Leakage radiation" means radiation emanating from the diagnostic source assembly except for:

(i) the useful beam, and

(ii) radiation produced when the exposure switch or timer is not activated.

(117) "Leakage technique factors" means the technique factors associated with the diagnostic or therapeutic source assembly which are used in measuring leakage radiation. They are defined as follows:

(i) for diagnostic source assemblies intended for capacitor energy storage equipment, the maximum-rated peak tube potential and the maximum-rated number of exposures in an hour for operation at the maximum-rated peak tube potential with the quantity of charge per exposure being 10 millicoulombs, i.e., 10 milliampere seconds, or the minimum obtainable from the unit, whichever is larger.

(ii) for diagnostic source assemblies intended for field emission equipment rated for pulsed operation, the maximum-rated peak tube potential and the maximum-rated number of x-ray pulses in an hour for operation at the maximum-rated peak tube potential.

(iii) for all other diagnostic or therapeutic source assemblies, the maximum-rated peak tube potential and the maximum-rated continuous tube current for the maximum-rated peak tube potential.

(118) "License" means a radioactive materials license issued by the Department for the transfer, receipt, production, possession or use of radioactive materials pursuant to this Code. There are two types of licenses: general and specific. A "general license" means a license to transfer, receive, possess, or use radioactive material in certain forms or quantities which is issued pursuant to the terms and conditions of this Code. General licenses are effective without the filing of an application with or the issuance of a license document by the Department. A "specific license" means a license evidenced by a license document issued by the Department to a licensee upon review and approval of an application submitted pursuant to this Code or a license similarly issued by the New York State Department of Health, the New York State Department of Labor, the U.S. Nuclear Regulatory Commission or any agreement state. Unless otherwise specified, the type of license referred to in this Code shall be a specific license.

(119) "Licensed material" means byproduct, source, or special nuclear material received, possessed, produced, used, transferred or disposed of under a general or specific license issued by the Department or any radioactive material which is subject to the licensure requirement of this Code.

(120) "Licensee" means any person who is licensed by the Department in accordance with this Code or any person who possesses radioactive material which is subject to the licensure requirements of this Code.

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(121) "Limits" [See "Dose limits"].

(122) "Light field" means the area illuminated by visible light, simulating the radiation field.

(123) "Linear accelerator" [See "Accelerator"]. For the purposes of this Code, "linac" is an equivalent term.

(124) "Line-voltage regulation" means the difference between the no-load line potentials expressed as a percent of the load line potential; that is,

Percent line-voltage regulation = 100 v1 = s1(Vn - V1)v2 = s1v3 = s2/2 (v1 + v3)_____,(v2 - v1)
(v1 + v3)
V1

where:

Vn = No-load line potential and

V1 = Line load potential.

(125) "Lost or missing licensed material" means licensed radioactive material whose location is unknown. This definition includes licensed material that has been shipped but has not reached its planned destination and whose location cannot be readily traced in the transportation system.

(126) "Low specific activity (LSA) material"' means radioactive material with limited specific activity which is nonfissile or is excepted under § 175.105(b)(2) that satisfies the descriptions and limits set forth below. Shielding materials surrounding the LSA material may not be considered in determining the estimated average specific activity of the package contents. LSA material must be in one of three groups:

(1) LSA-I.

(i) Uranium and thorium ores, concentrates of uranium and thorium ores, and other ores containing naturally occurring radioactive radionuclides that are not intended to be processed for the use of these radionuclides; or

(ii) Solid unirradiated natural uranium or depleted uranium or natural thorium or their solid or liquid compounds or mixtures; or

(iii) Radioactive material, other than fissile material, for which the A2 value is unlimited; or

(iv) Mill tailings, contaminated earth, concerete, rubble, other debris, and activated material in which the radioactive material is essentially uniformly distri-

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buted, and the average specific activity does not exceed 10-6 A2/g.

(2) LSA-II.

(i) Water with tritium concentration up to 0.8 TBq/liter (20.0 Ci/liter); or

(ii) Material in which the radioactive material is essentially uniformly distributed, and the average specific activity does not exceed 10-4 A2/g for solids and gases, and 10-5 A2/g for liquids.

(3) LSA-III. Solids (e.g., consolidated wastes, activated materials) excluding powders, that satisfy the requirements of § 71.77 in which:

(i) The radioactive material is essentially uniformly distributed throughout a solid or a collection of solid objects, or is essentially uniformly distributed in a solid compact binding agent (such as concrete, bitumen, ceramic, etc.);

(ii) The radioactive material is relatively insoluble, or it is intrinsically contained in a relatively insoluble material, so that, even under loss of packaging, the loss of radioactive material per package by leaching, when placed in water for 7 days, would not exceed 0.1 A2; and

(iii) The estimated average specific activity of the solid does not exceed 2X10-3 A2/g.

(127) "Lung class" [see "Class"].

(128) "Major processor" means a user processing, handling, or manufacturing radioactive material exceeding Type A quantities as unsealed sources or material, or exceeding 4 times Type B quantities as sealed sources, but does not include nuclear medicine programs, universities, industrial radiographers, or small industrial programs.

(129) "Management" means the chief executive officer or that individual's designee or designees.

(130) "Maximum line current" means the root-mean-square current in the supply line of an x-ray machine operating at its maximum rating.

(131) "Medical institution" means a facility as defined in Article 28 of the New York State Public Health Law.

(132) "Medical misadministration" means the administration of:

(i) a radiopharmaceutical, radiobiologic or radiation from a source other than the one ordered;

(ii) a radiopharmaceutical, radiobiologic or radiation to the wrong person;

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(iii) a radiopharmaceutical, radiobiologic or radiation by a route of administration, or to a part of the body, other than that in the order of the prescribing physician;

(iv) an activity of a diagnostic radiopharmaceutical or radiobiologic differing from the prescribed activity by more than 50 percent;

(v) an activity of a therapeutic radiopharmaceutical or radiobiologic differing from the prescribed activity by more than 10 percent;

(vi) a therapeutic radiation dose from any source other than a radiopharmaceutical, radiobiologic or brachytherapy source such that errors in computation, calibration, time of exposure, treatment geometry or equipment malfunction result in a calculated total treatment dose differing from the final prescribed total treatment dose ordered by more than 10 percent;

(vii) a therapeutic radiation dose from a brachytherapy source such that errors in computation, calibration, treatment time, source activity, source placement or equipment malfunction result in a calculated total treatment dose differing from the final total treatment dose ordered by more than 10 percent; or

(viii) a therapeutic radiation dose in any fraction of a fractionated treatment such that the administered dose in the individual treatment or fraction differs from the dose ordered for that individual treatment or fraction by more than 50 percent.

(133) "Medical use" means the intentional internal or external administration of radiation to humans in the practice of the healing arts in accordance with a license issued by a State or Territory of the United States, the District of Columbia, or the Commonwealth of Puerto Rico. For the purposes of this Code, "human use" is an equivalent term.

(134) "Mega electron volt (MeV)" means the energy equal to that acquired by a particle with one electron charge passing through a potential difference of one million volts in a vacuum.

(135) "Member of the public" means any individual, except when that individual is receiving an occupational dose.

(136) "Minor" means an individual less than 18 years of age.

(137) "Monitor unit" means a unit response from the beam monitoring system from which the absorbed dose can be calculated. For the purposes of this Code, "Dose monitor unit" is an equivalent term.

(138) "Monitoring" means the measurement of radiation, radioactive material concentrations, surface area activities or quantities of radioactive material and the use of the results of these measurements to evaluate potential exposures and doses. For purposes of this Code, "radiation monitoring" and "radiation protection monitoring" are equivalent terms.

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(139) "NARM" means any naturally occurring or accelerator-produced radioactive material. It does not include byproduct, source, or special nuclear material.

(140) "Natural radioactivity" means radioactivity of naturally occurring nuclides.

(141) "Nonstochastic effect" means a health effect, the severity of which varies with the dose and for which a threshold is believed to exist. Radiation-induced cataract formation is an example of a nonstochastic effect. For purposes of this Code, "deterministic effect" is an equivalent term.

(142) "Normal form radioactive material" means radioactive material which has not been demonstrated to qualify as special form radioactive material.

(143) "Nuclear Regulatory Commission" (NRC) means the U.S. Nuclear Regulatory Commission or its duly authorized representatives.

(144) "Occupational dose" means the dose received by an individual in the course of employment in which the individual's assigned duties involve exposure to radiation or to radioactive material from licensed and unlicensed sources of radiation, whether in the possession of the licensee, registrant, or other person. Occupational dose does not include doses received from background radiation, from any medical administration the individual has received, from exposure to individuals administered radioactive material and released under § 175.103(c)(9), from voluntary participation in medical research programs, or as a member of the public.

(145) "Operator" means any person conducting the business or activities carried on within a radiation installation or having by law the administrative control of a radiation source whether as owner, lessee, contractor, user or otherwise.

(146) "Output" means the exposure rate, dose rate, or a quantity related in a known manner to these rates of ionizing radiation from an external beam therapy unit for a specified set of exposure conditions.

(147) "Package" means the packaging together with its radioactive contents as presented for transport.

(i) "Fissile material package" or Type AF package, Type BF package, Type B(U)F package, or Type B(M)F package means a fissile material packaging together with its fissile material contents.

(2) Type A package means a Type A packaging together with its radioactive contents. A Type A package is defined and must comply with DOT regulations in 49 CFR Part 173.

(ii) "Type B package" means a Type B packaging together with its radioactive contents. On approval by the NRC, a Type B package design is designated by NRC as B(U) unless the package has a maximum normal operating pressure of more than 700 kPa (100 lb/in 2) gauge or a pressure relief device that would allow the release of ra-

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dioactive material to the environment under the tests specified in 10 CFR 71.73 (hypothetical accident conditions), in which case it will receive a designation B(M). B(U) refers to the need for unilateral approval of international shipments; B(M) refers to the need for multilateral approval of international shipments. There is no distinction made in how packages with these designations may be used in domestic transportation. To determine their distinction for international transportation, see USDOT regulations in 49 CFR Part 173. A Type B package approved before September 6, 1983, was designated only as Type B. Limitations on its use are specified in 10 CFR 71.13.

(148) "Packaging" means the assembly of components necessary to ensure compliance with the packaging requirements of 10 CFR Part 71. It may consist of one or more receptacles, absorbent materials, spacing structures, thermal insulation, radiation shielding, and devices for cooling or absorbing mechanical shocks. The vehicle, tie-down system, and auxiliary equipment may be designated as part of the packaging.

(149) "Particle accelerator" [See "Accelerator"].

(150) "Peak tube potential" means the maximum value of the potential difference across the x-ray tube during an exposure.

(151) "Person" means any individual, corporation, partnership, firm, association, trust, estate, public or private institution, group, agency, public authority or political subdivision of this State, any other State of the United States or political subdivision or agency thereof, and any legal successor, representative, agent or agency of the foregoing, but shall not include federal government agencies.

(152) "Personnel monitoring equipment" [See "Individual monitoring devices"].

(153) "Phantom" means an object behaving in essentially the same manner as tissue with respect to absorption or scattering of the ionizing radiation in question.

(154) "Planned special exposure" means an infrequent exposure to radiation, separate from and in addition to the annual occupational dose limits.

(155) "Position indicating device (PID)" means a device on dental x-ray equipment used to indicate the beam position and to establish a definite source-surface (skin) distance.

(156) "Positive collimating device" means a device which is permanently affixed to the x-ray tube housing and is intended to confine the emerging x-ray beam to the image receptor or area of clinical interest, whichever is smaller.

(157) "Primary protective barrier" [See "Protective barrier "].

(158) "Probabilistic effect" [See "Stochastic effect"].

(159) "Professional practice" means the practice of medicine, dentistry, podiatry, osteopathy, chiropractic or veterinary medicine.

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(160) "Professional practitioner" means any person licensed or otherwise authorized under the New York State Education Law to practice a professional practice.

(161) "Protective apron" means an apron made of radiation attenuating material(s), used to reduce radiation exposure.

(162) "Protective barrier" means a barrier of radiation absorbing material(s) used to reduce radiation exposure. The types of protective barriers are as follows:

(i) "Primary protective barrier" means the material, excluding filters, placed in the useful beam, for protection purposes, to reduce the radiation exposure.

(ii) "Secondary protective barrier" means a barrier sufficient to attenuate the stray radiation to the required degree.

(163) "Protective glove" means a glove made of radiation attenuating material(s) used to reduce radiation exposure.

(164) "Public dose" means the dose received by a member of the public from exposure to sources of radiation. It does not include occupational dose, dose received from background radiation, exposure to individuals administered radioactive material and released under § 175.103(c)(9), dose received as a patient from medical practices, or dose from voluntary participation in medical research programs.

(165) "Pyrophoric liquid" means any liquid that ignites spontaneously in dry or moist air at or below 54.4 degreeC (130 degreeF). A pyrophoric solid is any solid material, other than one classed as an explosive, which under normal conditions is liable to cause fires through friction, retained heat from manufacturing or processing, or which can be ignited readily and, when ignited, burns so vigorously and persistently as to create a serious transportation, handling, or disposal hazard. Included are spontaneously combustible and water-reactive materials.

(166) "Qualified expert" means an individual having the knowledge and training to measure ionizing radiation, to evaluate safety techniques, and to advise regarding radiation protection needs, e.g., individuals certified in the appropriate field by the American Board of Radiology or the American Board of Health Physics, or those having equivalent qualifications. With reference to the calibration of radiation therapy equipment, an individual having, in addition to the above qualifications, training and experience in the clinical applications of radiation physics to radiation therapy, e.g., individuals certified by the American Board of Medical Physics or in therapeutic Radiological Physics or X-Ray and Radium Physics by the American Board of Radiology.

(167) "Quality factor" (Q) means the modifying factor that is used to derive dose equivalent from absorbed dose.

(i) As used in this Code, the quality factors for converting absorbed dose to dose equivalent are shown in Table 1.

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Table 1 Quality Factors and Absorbed Dose Equivalents

Type of Radiation	Quality Factor (Q)	Absorbed Dose Equal to a Unit Dose Equivalent*
X, gamma, or beta radiation and high-speed electrons	1	1
Alpha particles, multiple-charged particles, fission fragments and heavy particles of unknown charge	20	0.05
Neutrons of unknown energy	10	0.1
High-energy protons	10	0.1

*Absorbed dose in gray equal to 1 Sv or the absorbed dose in rad equal to 1 rem.

(ii) If it is more convenient to measure the neutron fluence rate than to determine the neutron dose equivalent rate in sievert per hour or rem per hour, as provided in Table 1, 0.01 Sv (1 rem) of neutron radiation of unknown energies may, for purposes of this Code, be assumed to result from a total fluence of 25 million neutrons per square centimeter incident upon the body. If sufficient information exists to estimate the approximate energy distribution of the neutrons, the licensee or registrant may use the fluence rate per unit dose equivalent or the appropriate Q value from Table 2 to convert a measured tissue dose in gray or rad to dose equivalent in sievert or rem.

Table 2 Mean Quality Factors, Q, and Fluence per Unit Dose Equivalent for Monoenergetic Neutrons

Neutron Energy (MeV)	Quality Factora (Q)	Fluence per Unit Dose Equivalentb (neutrons cm-2 rem-1)	Fluence per Unit Dose Equivalentb (neutrons cm-2 Sv-1)
(thermal) 2.5E-8	2	980E+6	980E+8
1E-7	2	980E+6	980E+8
1E-6	2	810E+6	810E+8
1E-5	2	810E+6	810E+8
1E-4	2	840E+6	840E+8
1E-3	2	980E+6	980E+8
1E-2	2.5	1010E+6	1010E+8
1E-1	7.5	170E+6	170E+8
5E-1	11	39E+6	39E+8
1	11	27E+6	27E+8
2.5	9	29E+6	29E+8
5	8	23E+6	23E+8
7	7	24E+6	24E+8
10	6.5	24E+6	24E+8
14	7.5	17E+6	17E+8
20	8	16E+6	16E+8
40	7	14E+6	14E+8
60	5.5	16E+6	16E+8
1E+2	4	20E+6	20E+8

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2E+2	3.5	19E+6	19E+8
3E+2	3.5	16E+6	16E+8
4E+2	3.5	14E+6	14E+8

aValue of quality factor (Q) at the point where the dose equivalent is maximum in a 30-centimeter diameter cylinder tissue-equivalent phantom.

bMonoenergetic neutrons incident normally on a 30-centimeter diameter cylinder tissue-equivalent phantom.

(168) "Quarter" [See "Calendar quarter"].

(169) "Rad" means the special unit of absorbed dose. One rad is equal to an absorbed dose of 100 ergs per gram or 0.01 joule per kilogram (0.01 gray).

(170) "Radiation" means alpha particles, beta particles, gamma rays, x rays, neutrons, high-speed electrons, high-speed protons, and other particles capable of producing ions. For purposes of this Code, ionizing radiation is an equivalent term. Radiation, as used in this Code, does not include non-ionizing radiation, such as radiowaves or microwaves, visible, infrared, or ultraviolet light.

(171) "Radiation area" means any area, accessible to individuals, in which radiation levels could result in an individual receiving a dose equivalent in excess of 0.05 mSv (0.005 rem) in 1 hour at 30 centimeters (12 inches) from the source of radiation or from any surface that the radiation penetrates.

(172) "Radiation detector" means a device which in the presence of radiation provides a signal or other indication suitable for use in measuring one or more quantities of incident radiation.

(173) "Radiation dose" [See "Dose"].

(174) "Radiation equipment" means any equipment or device which can emit radiation by virtue of the application thereto of high voltage.

(175) "Radiation installation" means any place or facility, including vehicles such as a van or truck, where:

(i) radiation equipment, in operable condition or assembles and intended to be used, is located or used; or

(ii) radioactive material is transferred, received, produced, possessed or used.

Such installation shall include generally a hospital; medical, dental, chiropractic, osteopathic, podiatric, or veterinarian institution, clinic or office; van or truck providing services at non-permanent locations; educational institution; commercial, private or research laboratory performing diagnostic procedures or handling equipment or material for medical use; or any trucking, storage, messenger or delivery service establishment. Radiation installation shall include, whether or not it is specifically stated above, any place, facility or vehicle such as a van

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or truck where radiation is applied intentionally to a human. The limits of the radiation installation shall be as designated by the operator.

(176) "Radiation machine" means any device capable of producing radiation except those devices with radioactive material as the only source of radiation.

(177) "Radiation safety officer" means an individual who, under the authorization of the operator of a radiation installation, administers a radiation protection program in accordance with § 175.03 of this Code and who is qualified by training and experience in radiological health to evaluate the radiation hazards of such installation and administer such radiation protection program.

(178) "Radiation source" means any radioactive material or any radiation equipment.

(179) "Radiation therapy physicist" means the individual identified as the qualified radiation therapy physicist on a Department license or certified registration.

(180) "Radiation therapy simulation system" means a radiographic or fluoroscopic x-ray system intended for localizing the volume to be exposed during radiation therapy and confirming the position and size of the therapeutic irradiation field.

(181) "Radioactive material" means any solid, liquid, or gas which emits radiation spontaneously.

(182) "Radioactive material site" means a location, or contiguous and adjacent locations, under a single license in which radioactive materials are authorized to be received, produced, used, possessed (stored), or transferred and in which a specific use of said radioactive materials may be evaluated by a single set of Departmental inspection criteria concerning the procedures, equipment or shielding utilized by the licensee.

(183) "Radioactivity" means the transformation of unstable atomic nuclei by the emission of radiation.

(184) "Radiobioassay" [See "Bioassay"].

(185) "Rated line voltage" means the range of potentials, in volts, of the supply line specified by the manufacturer at which the x-ray machine is designed to operate.

(186) "Rated output current" means the maximum allowable load current of the x-ray high-voltage generator.

(187) "Rated output voltage" means the allowable peak potential, in volts, at the output terminals of the x-ray high-voltage generator.

(188) "Rating" means the operating limits specified by the manufacturer.

(189) "Recording" means producing a permanent form of an image resulting from x-

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ray photons (e.g., film, video tape).

(190) "Reference Man" means a hypothetical aggregation of human physical and physiological characteristics determined by international consensus. These characteristics may be used by researchers and public health workers to standardize results of experiments and to relate biological insult to a common base.

(191) "Registrant" means any person who is registered with the Department or who is legally obligated to register with the Department pursuant to this Code.

(192) "Registration" means registration with the Department in accordance with this Code.

(193) "Rem" means the special unit of any of the quantities expressed as dose equivalent. The dose equivalent in rem is equal to the absorbed dose in rad multiplied by the quality factor. One rem is equal to 0.01 sievert.

(194) "Research and development" means:

(i) theoretical analysis, exploration, or experimentation; or

(ii) the extension of investigative findings and theories of a scientific or technical nature into practical application for experimental and demonstration purposes, including the experimental production and testing of models, devices, equipment, materials, and processes.

Research and development does not include the internal or external administration of radiation or radioactive material to human beings.

(195) "Residual radioactivity" means radioactivity in structures, materials, soils, groundwater, and other media at a site resulting from activities under the licensee's control. This includes radioactivity from all licensed and unlicensed sources used by the licensee, but excludes background radiation. It also includes radioactive materials remaining at the site as a result of routine or accidental releases of radioactive material at the site and previous burials at the site, even if those burials were made in accordance with the provisions of 10 CFR Part 20.

(196) "Respiratory protective equipment" means an apparatus, such as a respirator, used to reduce an individual's intake of airborne radioactive materials.

(197) "Response time" means the time required for an instrument system to reach 90 percent of its final reading when the radiation-sensitive volume of the instrument system is exposed to a step change in radiation flux from zero sufficient to provide a steady state midscale reading.

(198) "Restricted area" means an area, access to which is limited by the licensee or registrant for the purpose of protecting individuals against undue risks from exposure to sources of radiation. A restricted area does not include any area used as residential quarters, but separate rooms in a residential building may be set apart as a restricted area.

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(199) "Roentgen" means the special unit of exposure. One roentgen (R) equals 2.58E-4 coulomb per kilogram of air (see "Exposure").

(200) "Sanitary sewerage" means a system of public sewers for carrying off waste water and refuse, but excluding sewage treatment facilities, septic tanks, and leach fields owned or operated by the licensee or registrant.

(201) "Scattered radiation" means radiation that, during passage through matter, has been deviated in direction. (The radiation also may have been modified by a decrease in energy.)

(202) "Sealed source" means radioactive material that is permanently bonded or fixed in a capsule or matrix designed to prevent release and dispersal of the radioactive material under the most severe conditions which are likely to be encountered in normal use and handling.

(203) "Secondary protective barrier" [See "Protective barrier"].

(204) "Shallow dose equivalent" (Hs), which applies to the external exposure of the skin of the whole body or the skin of an extremity, means the dose equivalent at a tissue depth of 0.007 centimeter (7 mg/cm 2).

(205) "Shutter" means a device attached to the tube housing assembly which can totally intercept the useful beam and which has a lead equivalency not less than that of the tube housing assembly.

(206) "SI" means the abbreviation for the International System of Units (Système Internationale).

(207) "Sievert" means the SI unit of any of the quantities expressed as dose equivalent. The dose equivalent in sievert is equal to the absorbed dose in gray multiplied by the quality factor. One sievert is equal to 100 rem.

(208) "Site boundary" means that line beyond which the land or property is not owned, leased, or otherwise controlled by the licensee or registrant.

(209) "Source" means, for the purposes of radiation equipment, the focal spot of the x-ray tube.

(210) "Source-image receptor distance (SID)" means the distance from the source to the center of the input surface of the image receptor.

(211) "Source material" means:

(i) Uranium or thorium, or any combination thereof, in any physical or chemical form; or

(ii) Ores that contain by weight one-twentieth of 1 percent (0.05 percent) or more

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of uranium, thorium or any combination of uranium and thorium. Source material does not include special nuclear material.

(212) "Source material milling" means any activity that results in the production of byproduct material as defined in § 175.02(a)(29)(ii).

(213) "Source of radiation" means any radioactive material or any device or equipment emitting, or capable of producing, radiation.

(214) "Source-skin distance or source-surface distance (SSD)" means the distance measured along the central ray from the center of the front surface of the source of the x-ray focal spot or sealed radioactive source to the surface of the irradiated object.

(215) "Special form radioactive material" means radioactive material which satisfies the following conditions:

(i) it is either a single solid piece or is contained in a sealed capsule that can be opened only by destroying the capsule;

(ii) the piece or capsule has at least one dimension not less than 5 mm (0.2 inch); and

(iii) it satisfies the requirements of 10 CFR 71.75. A special form encapsulation designed in accordance with the requirements of 10 CFR 71.4 in effect on June 30, 1983, (see 10 CFR Part 71, revised as of January 1, 1983), and constructed prior to July 1, 1985, and a special form encapsulation designed in accordance with the requirements of 10 CFR 71.4 in effect on March 31, 1996, (see 10 CFR Part 71, revised as of January 1, 1983), and constructed before April 1, 1998, may continue to be used. Any other special form encapsulation must meet the specifications of this definition.

(216) "Special nuclear material" means:

(i) Plutonium, uranium-233, uranium enriched in the isotope 233 or in the isotope 235, and any other material that the U.S. Nuclear Regulatory Commission, pursuant to the provisions of section 51 of the Atomic Energy Act of 1954, as amended, determines to be special nuclear material, but does not include source material; or

(ii) Any material artificially enriched by any of the foregoing, but does not include source material.

(217) "Special nuclear material in quantities not sufficient to form a critical mass" means uranium enriched in the isotope U-235 in quantities not exceeding 350 grams of contained U-235; uranium-233 in quantities not exceeding 200 grams; plutonium in quantities not exceeding 200 grams; or any combination of them in accordance with the following formula: For each kind of special nuclear material, determine the ratio between the quantity of that special nuclear material and the quantity specified above for the same kind of special nuclear material. The sum of such ratios for all of the kinds of special nuclear material in combination shall not

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exceed 1. For example, the following quantities in combination would not exceed the limitation and are within the formula:

$$\frac{175 \text{ (grams contained U-235)}}{350} + \frac{50 \text{ (grams U-233)}}{200} + \frac{50 \text{ (grams Pu)}}{200} = 1$$

(218) "Specific activity" of a radionuclide means the radioactivity of a radionuclide per unit mass of that nuclide. The specific activity of a material in which the radionuclide is essentially uniformly distributed is the radioactivity per unit mass of the material.

(219) "State" means the State of New York, unless the context of this Code clearly indicates that a different meaning is intended.

(220) "Stochastic effect" means a health effect that occurs randomly and for which the probability of the effect occurring, rather than its severity, is assumed to be a linear function of dose without threshold. Hereditary effects and cancer incidence are examples of stochastic effects. For purposes of this Code, "probabilistic effect" is an equivalent term.

(221) "Stray radiation" means the sum of leakage and scattered radiation.

(222) "Supervision" means:

(i) for radioactive materials licenses which do not authorize human use, the training of persons in the use of radioactive materials in other than medical procedures. Such training shall include at least thirty (30) hours of instruction in the principles and practices of radiation protection, radioactivity measurement, standardization and monitoring techniques and instruments, mathematics and calculations basic to the use and measurement of radioactivity, and biological effects of radiation; and

(ii) for radioactive materials licenses which do authorize human use, the training of a physician in the use of radioactive materials in the clinical treatment or diagnosis of disease. Such training shall provide that specified in § 175.102(j), as applicable.

(223) "Survey" means an evaluation of the radiological conditions and potential hazards incident to the production, use, transfer, release, disposal, or presence of sources of radiation. When appropriate, such evaluation includes, but is not limited to, tests, physical examinations, and measurements of levels of radiation or concentrations of radioactive material present.

(224) "Technique factors" means the conditions of operation of radiation equipment. They are specified as follows:

(i) For capacitor energy storage equipment, peak tube potential in kV and quantity of charge in mAs.

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(ii) For field emission equipment rated for pulsed operation, peak tube potential in kV and number of x-ray pulses.

(iii) For CT x-ray systems designed for pulsed operation, peak tube potential in kV, scan time in seconds, and either tube current in mA, x-ray pulse width in seconds, and the number of x-ray pulses per scan, or the product of tube current, x-ray pulse width, and the number of x-ray pulses in mAs;

(iv) For CT x-ray systems not designed for pulsed operation, peak tube potential in kV, and either tube current in mA and scan time in seconds, or the product of tube current and exposure time in mAs and the scan time when the scan time and exposure time are equivalent; and

(v) For all other equipment, peak tube potential in kV, and either tube current in mA and exposure time in seconds, or the product of tube current and exposure time in mAs.

(225) "Teletherapy" means a method of radiation therapy utilized to deliver a radiation dose in which the source (sources) of radiation is (are) at a distance from the body. For the purposes of this Code "external beam radiation therapy" is an equivalent term.

(226) "Test" means the process of verifying compliance with an applicable regulation.

(227) "Therapeutic-type protective tube housing" means:

(i) for x-ray therapy equipment not capable of operating at 500 kVp or above, the following definition applies: An x-ray tube housing so constructed that the leakage radiation averaged over any 100 cm 2 (15.5 inches 2) at a distance of 1 meter (3 feet) from the source does not exceed 2.58 E-4 C-kg-1 (1 roentgen) in an hour when the tube is operated at its maximum rated continuous current for the maximum rated tube potential.

(ii) For x-ray therapy equipment capable of operation at 500 kVp or above, the following definition applies: An x-ray tube housing so constructed that the leakage radiation averaged over an 100 cm 2 (15.5 inches 2) area at a distance of 1 meter (3 feet) from the source does not exceed 0.10 percent of the useful beam dose rate at 1 meter (3 feet) from the source for any of its operating conditions.

(228) "This Code" means Article 175 and all other parts of the New York City Health Code applicable to licensees and registrants or other persons subject to the provisions of Article 175.

(229) "Total effective dose equivalent" (TEDE) means the sum of the deep dose equivalent for external exposures and the committed effective dose equivalent for internal exposures.

(230) "Total organ dose equivalent" (TODE) means the sum of the deep dose equivalent and the committed dose equivalent to the organ receiving the highest dose as

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described in § 175.03(k)(7)(i)(F) of this Code.

(231) "Traceable to a national standard" means that a quantity or a measurement has been compared to a national standard directly or indirectly through one or more intermediate steps and that all comparisons have been documented.

(232) "Transport index (TI)" means the dimensionless number, rounded up to the next tenth, placed on the label of a package to designate the degree of control to be exercised by the carrier during transportation. The transport index is determined by multiplying the maximum radiation level in millisievert (mSv) per hour at 1 meter (3.3ft) from the external surface of the package by 100 (equivalent to the maximum radiation level in millirem per hour at 1 meter (3.3 ft)).

(233) "Tube housing assembly" means the tube housing with tube installed. It includes high-voltage and/or filament transformers and other appropriate elements when they are contained within the tube housing.

(234) "Tube rating chart" means the set of curves which specify the rated limits of operation of the tube in terms of the technique factors.

(235) "Type A package" means a packaging that, together with its radioactive contents limited to A1 or A2 as appropriate, meets the requirements of U.S. DOT 49 CFR 173.410 and 173.412 and is designed to retain the integrity of containment and shielding required by this part under normal conditions of transport as demonstrated by the tests set forth in 49 CFR 173.465 or 173.466, as appropriate.

(236) "Type A quantity" means a quantity of radioactive material, the aggregate radioactivity of which does not exceed A1 for special form radioactive material or A2 for normal form radioactive material. A1 and A2 are given in Appendix A of § 175.105 or may be determined by procedures described in such Appendix A.

(237) [Reserved]

(238) [Reserved]

(239) "Type B quantity" means a quantity of radioactive material greater than a Type A quantity.

(240) "U.S. Department of Energy" means the Department of Energy established by Public Law 95-91, August 4, 1977, 91 Stat. 565, 42 U.S.C. 7101 et seq., to the extent that the Department exercises functions formerly vested in the U.S. Atomic Energy Commission, its Chairman, members, officers and components and transferred to the U.S. Energy Research and Development Administration and to the Administrator thereof pursuant to sections 104(b), (c) and (d) of the Energy Reorganization Act of 1974 (Public Law 93-438, October 11, 1974, 88 Stat. 1233 at 1237, 42 U.S.C. 5814, effective January 19, 1975) and retransferred to the Secretary of Energy pursuant to section 301(a) of the Department of Energy Organization Act (Public Law 95-91, August 4, 1977, 91 Stat. 565 at 577-578, 42 U.S.C. 7151, effective October 1, 1977.)

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(241) "Unrefined and unprocessed ore" means ore in its natural form prior to any processing, such as grinding, roasting, beneficiating, or refining.

(242) "Unrestricted area" means an area, access to which is not controlled by the licensee or registrant for purposes of radiation protection.

(243) "Use" as used in radioactive materials licenses means to employ or apply radioactive materials for the licensed purpose. It shall include instruction of, and responsibility for, technical and support staff members. It does not include training others in the techniques of use of radioactive materials for the purpose of qualifying for licensure.

In licenses authorizing medical use of radioactive materials, "use" shall also include:

(i) ordering or directing the administration of radiation or radioactive materials to humans, including the method or route of administration;

(ii) actual use of, or direction of technologists or other paramedical personnel in the use of, radioactive material;

(iii) interpretation of results of diagnostic procedures; and

(iv) regular review of the progress of patients receiving therapy and modification of the originally prescribed dose as warranted by the patient's reaction to radiation therapy.

(244) "Useful beam" means the radiation emanating from the tube housing port or the radiation head and passing through the aperture of the beam limiting device when the exposure controls are in a mode to cause the system to produce radiation.

(245) "Variable-aperture beam-limiting device" means a beam-limiting device which has capacity for stepless adjustment of the x-ray field size at a given SID.

(246) "Very high radiation area" means an area, accessible to individuals, in which radiation levels could result in an individual receiving an absorbed dose in excess of 5 Gy (500 rad) in 1 hour at 1 meter (3 feet) from a source of radiation or from any surface that the radiation penetrates. At very high doses received at high dose rates, units of absorbed dose (gray and rad) are appropriate, rather than units of dose equivalent (sievert and rem).

(247) "Visible area" means that portion of the input surface of the image receptor over which incident x-ray photons produce a visible image.

(248) "Waste" means those low-level radioactive wastes that are acceptable for disposal in a land disposal facility. For the purposes of this definition, low-level waste has the same meaning as in the Low-Level Radioactive Waste Policy Act, P.L. 96-573, as amended by P.L. 99-240, effective January 15, 1986; that is, radioactive waste (a) not classified as high-level radioactive waste, spent nuclear fuel, or byproduct material as defined in Section lle.(2) of the Atomic Energy Act

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(uranium or thorium tailings and waste) and (b) classified as low-level radioactive waste consistent with existing law and in accordance with (a) by the U.S. Nuclear Regulatory Commission.

(249) "Waste handling licensees" mean persons licensed to receive and store radioactive wastes prior to disposal and/or persons licensed to dispose of radioactive waste.

(250) "Week" means 7 consecutive days starting on Sunday.

(251) "Weighting factor" wT for an organ or tissue (T) means the proportion of the risk of stochastic effects resulting from irradiation of that organ or tissue to the total risk of stochastic effects when the whole body is irradiated uniformly. For calculating the effective dose equivalent, the values of WT are:

Organ Dose Weighting Factors

Organ or Tissue	WT
Gonads	0.25
Breast	0.15
Red bone marrow	0.12
Lung	0.12
Thyroid	0.03
Bone surfaces	0.03
Remainder	0.30a
Whole Body	1.00b

a 0.30 results from 0.06 for each of 5 "remainder" organs, excluding the skin and the lens of the eye, that receive the highest doses.

b For the purpose of weighting the external whole body dose, for adding it to the internal dose, a single weighting factor, wT = 1.0, has been specified. The use of other weighting factors for external exposure will be approved on a case-by-case basis until such time as specific guidance is issued.

(252) "Whole body" means, for purposes of external exposure, head, trunk (including male gonads), arms above the elbow, or legs above the knee.

(253) "Worker" means an individual engaged in work under a license or registration issued by the Department and controlled by a licensee or registrant, but does not include the licensee or registrant.

(254) "Working level" (WL) means any combination of short-lived radon daughters in 1 liter of air that will result in the ultimate emission of 1.3E+5 MeV of potential alpha particle energy. The short-lived radon daughters are:

(i) for radon-222: polonium-218, lead-214, bismuth-214, and polonium-214; and

(ii) for radon-220: polonium-216, lead-212, bismuth-212, and polonium-212.

(255) "Working level month" (WLM) means an exposure to 1 working level for 170

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hours (2,000 working hours per year divided by 12 months per year is approximately equal to 170 hours per month).

(256) "X-ray control" means a device which controls input power to the x-ray high-voltage generator and/or the x-ray tube. It includes equipment such as timers, phototimers, automatic brightness stabilizers, and similar devices, which control the technique factors of an x-ray exposure.

(257) "X-ray equipment" means an x-ray system, subsystem, or component thereof. Types of x-ray equipment are as follows:

(i) "Mobile x-ray equipment" means x-ray equipment mounted on a permanent base with wheels and/or casters for moving while completely assembled.

(ii) "Portable x-ray equipment" means x-ray equipment designed to be hand carried.

(iii) "Stationary x-ray equipment" means x-ray equipment which is installed in a fixed location.

(258) "X-ray field" means that area of the intersection of the useful beam and any one of the set of planes parallel to and including the plane of the image receptor, whose perimeter is the locus of points at which the exposure rate is one-fourth of the maximum in the intersection.

(259) "X-ray high-voltage generator" means a device which transforms electrical energy from the potential supplied by the x-ray control to the tube operating potential. The device may also include means for transforming alternating current to direct current, filament transformers for the x-ray tube(s), high-voltage switches, electrical protective devices, and other appropriate elements.

(260) "X-ray system" means an assemblage of components for the controlled production of x-rays. It includes minimally an x-ray high-voltage generator, an x-ray control, a tube housing assembly, a beam-limiting device, and the necessary supporting structures. Additional components which function with the system are considered integral parts of the system.

(261) "X-ray subsystem" means any combination of two or more components of an x-ray system for which there are requirements specified in this Article.

(262) "X-ray tube" means any electron tube which is designed for the conversion of electrical energy into x-ray energy. For the purposes of permit fee requirements in Article 5 of this Code, an x-ray tube means any electrical device which produces x-rays of intensity exceeding 1.29 E-4 C-kg-1 (0.5 milliroentgen) per hour when measured 5 centimeters (2 inches) from any accessible surface thereof, and averaged over an area of 10 cm 2 (1.55 square inches).

(263) "Year" means the period of time beginning in January used to determine compliance with the provisions of this Code. The licensee or registrant may change the starting date of the year used to determine compliance by the licensee or registrant provided that the change is made at the beginning of the year and that no

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day is omitted or duplicated in consecutive years.

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Paragraphs (141) and (161) of subsection (a) of § 175.02 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 20.1003, which provides definitions for 10 CFR Part 20.

Subdivision (a) of § 175.02 was amended on September 26, 2006, to modify the definition of "shallow dose equivalent" to ensure compatibility with the revised NRC definition as published in the Federal Register, dated April 5, 2002, titled "Revision of the Skin Dose Limit", 67 FR 16298, that became effective April 5, 2002 (amending 10 CFR §§ 20.1003; 20.1201).

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (a) par (1) amended City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (20) amended City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) par (45) amended City Record Jan. 14, 2003 eff. Feb. 13, 2003. [See Vol. 9 Statements of Basis and Purpose No. 18-A]

Subd. (a) par (52) amended City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (54) added City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) pars (55)-(263) renumbered (former pars (54)-(260)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) par (58) added City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) par (61) renumbered (former par (59)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended (as par (59)) City Record Jan. 14, 2003 eff. Feb. 13, 2003. [See Vol. 9 Statements of Basis and Purpose No. 18-A]

Subd. (a) par (70) added City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9

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Statements of Basis and Purpose No. 28]

Subd. (a) par (81) renumbered (former par (78)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (78) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (91) renumbered (former par (88)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Repealed as par (88) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (92) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (92) renumbered (former par (89)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (89) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (119) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (119) renumbered (former par (116)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (116) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (126) open par amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (126) renumbered (former par (123)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (123) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (126) subpar (1) clause (i) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a). par (126) subpar (3) opening clause amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (126) subpar (3) clause (iii) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (135) renumbered and amended (former par (132)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) par (144) renumbered and amended (former par (141)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28].

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Amended as par (141) City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (a) par (147) open par amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (147) renumbered (former par (144)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (144) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (147) subpar (i) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (147) subpar (2) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (164) renumbered (former par (161)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (161) City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (a) par (195) added City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (a) par (204) renumbered (former par (200)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (200) City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (a) par (215) renumbered (former par (211)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (211) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (232) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (a) par (232) renumbered (former par (228)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (228) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (235) renumbered (former par (231)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Amended as par (231) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (237) renumbered (former par (233)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Repealed as par

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(233) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Subd. (a) par (238) renumbered (former par (234)) City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]. Repealed as par (234) City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

NOTES:

A number of definitions in § 175.02(a) in Article 175 of the NYC Health Code were added and/or amended on June 14, 2007 in order to assure compatibility with applicable federal U.S. Nuclear Regulatory Commission regulations on radioactive materials and licenses.

Paragraphs (141) and (161) of subsection (a) of § 175.02 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 20.1003, which provides definitions for 10 CFR Part 20.

Subdivision (a) of § 175.02 was amended on September 26, 2006, to modify the definition of "shallow dose equivalent" to ensure compatibility with the revised NRC definition as published in the Federal Register, dated April 5, 2002, titled "Revision of the Skin Dose Limit", 67 FR 16298, that became effective April 5, 2002 (amending 10 CFR §§ 20.1003; 20.1201).

Subdivisions (1), (52), (78), (89), (116), (123), (144), (228) and (231) of subsection (a) were amended and subdivisions (88), (233) and (234) of subsection (a) were repealed April 26, 1999 to conform to definitions in 10 CFR Part 71 to ensure compatibility with the NRC. In addition, the definition of "Licensed Material" was clarified to include radioactive material that requires a license even if one has not been obtained as well as radioactive material that is the subject of a general or specific license.

Subdivisions (45) and (59) of subsection (a) were amended by resolution adopted on December 12, 2002 following amendment of Chapter 22 of the New York City Charter by the electorate in the November 6, 2001 general election, which merged the Departments of Health and Mental Health, Mental Retardation and Alcoholism Services, and enactment of Local Law of 2002, which further amended the Charter and changed the name of the merged agency to Department of Health and Mental Hygiene, and the Commissioner to Commissioner of the Department of Health and Mental Hygiene.

Paragraphs (141) and (161) of subsection (a) of § 175.02 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 20.1003, which provides definitions for 10 CFR Part 20.

24 RCNY Hlth. Code § 175.02
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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
GENERAL PROVISIONS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.03. Standards for protection against radiation.

(a) General provisions. (1) Purpose. (i) This section establishes standards for protection against ionizing radiation resulting from activities conducted pursuant to licenses, certified registrations and/or registrations issued by the Department.

(ii) The requirements of this Code are designed to control the receipt, production, possession, use, transfer, and disposal of sources of radiation by any licensee or registrant so the total dose to an individual, including doses resulting from all sources of radiation other than background radiation, does not exceed the standards for protection against radiation prescribed in this section. However, nothing in this section shall be construed as limiting actions that may be necessary to protect health and safety.

(2) Scope. Except as specifically provided otherwise by this Code, this section applies to persons subject to licensure, certified registration or registration by the Department to receive, produce, possess, use, transfer, or dispose of sources of radiation. The limits in this section do not apply to doses due to background radiation, to exposure of patients to radiation for the purpose of medical diagnosis or therapy, or to voluntary participation in medical research programs.

(3) Implementation. (i) Any existing license, certified registration or registration condition that is more restrictive than the provisions in this section remains in force until there is an amendment or renewal of the license, certified registration or registration.

(ii) If a license, certified registration or registration condition exempts a licensee or registrant from a provision of this Code in effect on or before the effective date of this Code, it also exempts the licensee or registrant from the corresponding provision of this section.

(iii) If a license, certified registration or registration condition cites provisions of this Code in effect prior to the effective date of this Code, which do not correspond to any provisions of this section, the license, certified registration or registration condition remains in force until there is an amendment or renewal

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of the license, certified registration or registration that modifies or removes this condition.

(b) Radiation protection programs. (1) Radiation Protection Programs. Each person who operates or permits the operation of a radiation installation or who operates, transfers, receives, produces, possesses or uses, or permits the operation, transfer, receipt, production, possession or use of any radiation source shall:

(i) use, to the extent practicable, procedures and engineering controls based upon sound radiation protection principles to achieve occupational doses and public doses that are as low as reasonably achievable (ALARA) below the limits specified in this Code;

(ii) develop, document, and implement a radiation protection program commensurate with the scope and extent of the program and sufficient to ensure compliance with the provisions of this Code;

(iii) provide a radiation safety officer pursuant to § 175.03(b)(2) who shall be delegated authority to ensure the implementation of this radiation protection program. For licensed radioactive materials installations the radiation safety officer, or an authorized user designated to act as the radiation safety officer in the radiation safety officer's absence, shall be present on the premises at least 50 percent of the time that radioactive material is being handled or equipment containing radioactive material is being operated;

(iv) provide for a radiation safety committee to administer the radiation protection program in medical centers, hospitals and institutions of higher education. The committee shall include the facility operator or a person with the authority to act on behalf of the facility operator, and representation from departments within the facility where radiation sources are used. The committee shall oversee all uses of radiation-producing equipment and radioactive materials within the facility, shall review the activities of the radiation safety officer, and shall review the radiation safety program at least annually. The committee, or a subcommittee, shall oversee the administration of a quality assurance program as required by § 175.03(b)(1)(v);

(v) provide a quality assurance program for diagnostic and therapeutic uses of radiation-producing equipment and radioactive materials pursuant to § 175.07 and other applicable provisions of this Code;

(vi) ensure that all personnel involved in planning for or administering radiation doses to humans, or in the use of radiation-producing equipment or radioactive materials for other purposes, are supervised, are instructed as described in § 175.04(c) and are competent to safely use such radiation sources and services;

(vii) ensure that radiation equipment is used only for those procedures for which it is designed;

(viii) ensure that acceptance testing, by an individual competent to perform such testing, is performed on all medical and chiropractic diagnostic equipment and radiation therapy treatment and planning equipment before the first use of such equip-

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ment on humans; and

(ix) review the radiation protection program content and implementation at intervals not to exceed twelve months.

(2) Radiation safety officer. The radiation safety officer specified in § 175.03(b)(1)(iii) shall be:

(i) For human use radiation equipment installations:

(A) a physicist certified by the American Board of Health Physics, the American Board of Radiology or the American Board of Medical Physics in a branch of physics related to the type of use of radiation sources in the installation; or

(B) a person satisfying the radiation safety officer qualifications set forth in § 175.103(j) of this Code; or

(C) a professional practitioner as defined in § 175.02(a)(157), practicing within such person's professional practice as defined in § 175.02(a)(156).

(ii) For human use radiation equipment installations requiring a certified registration pursuant to § 175.64:

(A) a physicist certified by the American Board of Health Physics, the American Board of Radiology or the American Board of Medical Physics in a branch of physics related to the type of use of radiation sources in the installation; or

(B) a person satisfying the radiation safety officer qualifications set forth in § 175.103(j) of this Code; or

(C) an authorized user named on the facility's certified registration issued by the Department.

(iii) For non-human use radiation equipment installations:

(A) a veterinarian for veterinary installations; or

(B) a physicist certified by the American Board of Health Physics, the American Board of Radiology or the American Board of Medical Physics; or

(C) a person with equivalent training and experience as determined by the Department; or

(D) a researcher determined by the institution as qualified by training and experience for installations using only x-ray diffraction and fluorescence analysis equipment.

(iv) For non-human use radioactive materials installations:

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(A) a physicist certified by the American Board of Health Physics, the American Board of Radiology or the American Board of Medical Physics in a branch of physics related to the type and use of radioactive material in the installation; or

(B) a person with equivalent training and experience as determined by the Department; or

(C) an authorized user named on the radioactive materials license issued by the Department.

(c) Occupational dose limits. (1) Occupational dose limits for adults.

(i) Except for planned special exposures pursuant to § 175.03(c)(6), the licensee or registrant shall control the occupational dose to any individual adult from licensed or registered activities to ensure that such dose does not exceed:

(A) an annual limit, which is the lesser of:

(a) a total effective dose equivalent of 0.05 Sv (5 rem); or

(b) the sum of the deep dose equivalent and the committed dose equivalent to any individual organ or tissue, other than the lens of the eye, equal to 0.5 Sv (50 rem); and

(B) annual limits to the lens of the eye, to the skin of the whole body, and to the skin of the extremities of:

(a) an eye dose equivalent of 0.15 Sv (15 rem), and

(b) a shallow dose equivalent of 0.5 Sv (50 rem) to the skin of the whole body or to the skin of any extremity.

(ii) Doses received in excess of the annual limits, including doses received during accidents, emergencies, and planned special exposures, shall be subtracted from the limits for planned special exposures that the individual may receive during the current year and during the individual's lifetime.

(iii) The assigned deep dose equivalent must be for the part of the body receiving the highest exposure. The assigned shallow dose equivalent must be the dose averaged over the contiguous 10 square centimeters of skin receiving the highest exposure:

(A) the deep dose equivalent, eye dose equivalent and shallow dose equivalent may be assessed from surveys or other radiation measurements for the purpose of demonstrating compliance with the occupational dose limits, if the individual monitoring device was not in the region of highest potential exposure, or the results of individual monitoring are unavailable; or

(B) when a protective apron is worn during x-ray fluoroscopic procedures to be in compliance with § 175.62(i) of this Code and monitoring is conducted as specified

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in § 175.03(f)(2)(ii), the effective dose equivalent for external radiation may be determined for these individuals as follows:

(a) when only one individual monitoring device is used and it is located at the neck outside the protective apron, the reported deep dose equivalent value multiplied by 0.3 shall be the effective dose equivalent for external radiation; or

(b) when individual monitoring devices are worn, both under the protective apron at the waist and outside the protective apron at the neck, the effective dose equivalent for external radiation shall be assigned the value of the sum of the deep dose equivalent reported for the individual monitoring device located at the waist under the protective apron multiplied by 1.5 and the deep dose equivalent reported for the individual monitoring device located at the neck outside the protective apron multiplied by 0.04.

(iv) Derived air concentration (DAC) and annual limit on intake (ALI) values are presented in Table 1 of Appendix B of this section and may be used to determine the individual's dose and to demonstrate compliance with the occupational dose limits.

(v) Notwithstanding the annual dose limits, the licensee shall limit the soluble uranium intake by an individual to 10 milligrams in a week.

(vi) The licensee or registrant shall reduce the dose that an individual may be allowed to receive in the current year by the amount of occupational dose received while employed by any other person.

(2) Requirements for summation of external and internal doses. (i) If the licensee or registrant is required to monitor pursuant to both § 175.03(f)(2)(i) and (v), the licensee or registrant shall demonstrate compliance with the dose limits by summing external and internal doses. If the licensee or registrant is required to monitor only pursuant to § 175.03(f)(2)(i) or only pursuant to § 175.03(f)(2)(v), then summation is not required to demonstrate compliance with the dose limits. The licensee or registrant may demonstrate compliance with the requirements for summation of external and internal doses pursuant to § 175.03(c)(2)(ii), (iii) and (iv). The dose equivalents for the lens of the eye, the skin, and the extremities are not included in the summation, but are subject to separate limits.

(ii) Intake by inhalation. If the only intake of radionuclides is by inhalation, the total effective dose equivalent limit is not exceeded if the sum of the deep dose equivalent divided by the total effective dose equivalent limit, and one of the following, does not exceed unity:

(A) the sum of the fractions of the inhalation ALI for each radionuclide, or

(B) the total number of derived air concentration-hours (DAC-hours) for all radionuclides divided by 2,000, or

(C) the sum of the calculated committed effective dose equivalents to all significantly irradiated organs or tissues (T) calculated from bioassay data using appro-

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priate biological models and expressed as a fraction of the annual limit. For purposes of this requirement, an organ or tissue is deemed to be significantly irradiated if, for that organ or tissue, the product of the weighting factors, wT, and the committed dose equivalent, HT,50, per unit intake is greater than 10 percent of the maximum weighted value of H50, that is, wTHT,50, per unit intake for any organ or tissue.

(iii) Intake by oral ingestion. If the occupationally exposed individual also receives an intake of radionuclides by oral ingestion greater than 10 percent of the applicable oral ALI, the licensee or registrant shall account for this intake and include it in demonstrating compliance with the limits.

(iv) Intake through wounds or absorption through skin. The licensee or registrant shall evaluate and account for intakes through wounds or skin absorption. The intake through intact skin has been included in the calculation of DAC for hydrogen-3 and does not need to be evaluated or accounted for separately pursuant to § 175.03(c)(2)(iv).

(3) Determination of external dose from airborne radioactive material.

(i) Licensees, when determining the dose from airborne radioactive material, shall include the contribution to the deep dose equivalent, eye dose equivalent, and shallow dose equivalent from external exposure to the radioactive cloud. (See Appendix B of this section, footnotes 1 and 2.)

(ii) Airborne radioactivity measurements and DAC values shall not be used as the primary means to assess the deep dose equivalent when the airborne radioactive material includes radionuclides other than noble gases or if the cloud of airborne radioactive material is not relatively uniform. The determination of the deep dose equivalent to an individual shall be based upon measurements using instruments or individual monitoring devices.

(4) Determination of internal exposure. (i) For purposes of assessing dose used to determine compliance with occupational dose equivalent limits, the licensee shall, when required pursuant to § 175.03(f)(2) take suitable and timely measurements of:

(A) concentrations of radioactive materials in air in work areas; or

(B) quantities of radionuclides in the body; or

(C) quantities of radionuclides excreted from the body; or

(D) combinations of these measurements.

(ii) Unless respiratory protective equipment is used, as provided in § 175.03(h)(3), or the assessment of intake is based on bioassays, the licensee shall assume that an individual inhales radioactive material at the airborne concentration in which the individual is present.

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(iii) When specific information on the physical and biochemical properties of the radionuclides taken into the body or the behavior of the material in an individual is known, the licensee may:

(A) use that information to calculate the committed effective dose equivalent, and, if used, the licensee or registrant shall document that information in the individual's record; and

(B) upon prior approval of the Department, adjust the DAC or ALI values to reflect the actual physical and chemical characteristics of airborne radioactive material, for example, aerosol size distribution or density; and

(C) separately assess the contribution of fractional intakes of Class D, W, or Y compounds of a given radionuclide to the committed effective dose equivalent. (See Appendix B of this section.)

(iv) If the licensee chooses to assess intakes of Class Y material using the measurements given in § 175.03(c)(4)(i)(B) or (C), the licensee may delay the recording and reporting of the assessments for periods up to 7 months, unless otherwise required by § 175.03(l)(2) or (3). This delay permits the licensee to make additional measurements basic to the assessments.

(v) If the identity and concentration of each radionuclide in a mixture are known, the fraction of the DAC applicable to the mixture for use in calculating DAC-hours shall be either:

(A) the sum of the ratios of the concentration to the appropriate DAC value, that is, D, W, or Y, from Appendix B for each radionuclide in the mixture; or

(B) the ratio of the total concentration for all radionuclides in the mixture to the most restrictive DAC value for any radionuclide in the mixture.

(vi) If the identity of each radionuclide in a mixture is known, but the concentration of one or more of the radionuclides in the mixture is not known, the DAC for the mixture shall be the most restrictive DAC of any radionuclide in the mixture.

(vii) When a mixture of radionuclides in air exists, a licensee may disregard certain radionuclides in the mixture if:

(A) the licensee uses the total activity of the mixture in demonstrating compliance with the dose limits in § 175.03(c) and in complying with the monitoring requirements in § 175.03(f)(2)(v), and

(B) the concentration of any radionuclide disregarded is less than 10 percent of its DAC, and

(C) the sum of these percentages for all of the radionuclides disregarded in the mixture does not exceed 30 percent.

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(viii) When determining the committed effective dose equivalent, the following information may be considered:

(A) In order to calculate the committed effective dose equivalent, the licensee or registrant may assume that the inhalation of one ALI, or an exposure of 2,000 DAC-hours, results in a committed effective dose equivalent of 0.05 Sv (5 rem) for radionuclides that have their ALIs or DACs based on the committed effective dose equivalent.

(B) For an ALI and the associated DAC determined by the nonstochastic organ dose limit of 0.5 Sv (50 rem), the intake of radionuclides that would result in a committed effective dose equivalent of 0.05 Sv (5 rem), that is, the stochastic ALI, is listed in parentheses in Table 1 of Appendix B of this section. The licensee may, as a simplifying assumption, use the stochastic ALI to determine committed effective dose equivalent. However, if the licensee or registrant uses the stochastic ALI, the licensee or registrant shall also demonstrate that the limit in § 175.03(c)(1)(i)(A) is met.

(5) Determination of prior occupational dose. (i) For each individual who may enter the licensee's or registrant's restricted area and is likely to receive, in a year, an occupational dose requiring monitoring pursuant to § 175.03(f)(2), the licensee or registrant shall:

(A) determine the occupational radiation dose received during the current year; and

(B) request, in writing, the records of lifetime cumulative occupational radiation dose.

(ii) Prior to permitting an individual to participate in a planned special exposure, the licensee or registrant shall determine:

(A) the internal and external doses from all previous planned special exposures; and

(B) all doses in excess of the limits, including doses received during accidents and emergencies, received during the lifetime of the individual; and

(C) all lifetime cumulative occupational radiation doses.

(iii) In complying with the requirements of § 175.03(c)(5)(i), a licensee or registrant may:

(A) accept, as a record of the occupational dose that the individual received during the current year, a written signed statement from the individual, or from the individual's most recent employer for work involving radiation exposure, that discloses the nature and the amount of any occupational dose that the individual received during the current year; and

(B) accept, as the record of lifetime cumulative radiation dose, an up-to-date

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form RAD-4, "Cumulative Occupational Radiation Exposure History," or equivalent, signed by the individual and countersigned by an appropriate official of the most recent employer for work involving radiation exposure, or the individual's current employer, if the individual is not employed by the licensee or registrant; and

(C) obtain reports of the individual's dose equivalent from the most recent employer for work involving radiation exposure, or the individual's current employer, if the individual is not employed by the licensee or registrant, by telephone, telegram, facsimile, or letter. The licensee or registrant shall request a written verification of the dose data if the authenticity of the transmitted report cannot be established.

(iv) (A) The licensee or registrant shall record the exposure history, as required by § 175.03(d)(5)(i), on form RAD-4, "Cumulative Occupational Radiation Exposure History," or other clear and legible record, of all the information required on that form. The form or record shall show each period in which the individual received occupational exposure to radiation or radioactive material and shall be signed by the individual who received the exposure. For each period for which the licensee or registrant obtains reports, the licensee or registrant shall use the dose shown in the report in preparing form RAD-4 or equivalent. For any period in which the licensee or registrant does not obtain a report, the licensee or registrant shall place a notation on form RAD-4 or equivalent indicating the periods of time for which data are not available.

(B) Licensees or registrants are not required to reevaluate the separate external dose equivalents and internal committed dose equivalents or intakes of radionuclides assessed pursuant to this Code in effect before January 1, 1994. Further, occupational exposure histories obtained and recorded on form RAD-4 or equivalent before the effective date of this Code, would not have included effective dose equivalent, but may be used in the absence of specific information on the intake of radionuclides by the individual.

(v) If the licensee or registrant is unable to obtain a complete record of an individual's current and previously accumulated occupational dose, the licensee or registrant:

(A) in establishing administrative controls pursuant to § 175.03(c)(1) for the current year, shall assume that the allowable dose limit for the individual is reduced by 12.5 mSv (1.25 rem) for each quarter for which records were unavailable and the individual was engaged in activities that could have resulted in occupational radiation exposure; and

(B) shall not authorize the individual to receive any planned special exposures.

(vi) The licensee or registrant shall retain the records on form RAD-4, "Cumulative Occupational Radiation Exposure History," or equivalent until the Department authorizes their disposition. The licensee or registrant shall retain records used in preparing form RAD-4 or equivalent for 3 years after the record is made.

(6) Planned special exposures. A licensee or registrant may authorize an adult worker to receive doses in addition to, and accounted for separately from, the dos-

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es received under the limits specified in § 175.03(c)(1) provided that each of the following conditions is satisfied:

(i) The licensee or registrant authorizes a planned special exposure only in an exceptional situation when alternatives that might avoid the higher exposure are unavailable or impractical.

(ii) Before the exposure occurs, the licensee or registrant, and employer if the employer is not the licensee or registrant, specifically authorizes the planned special exposure in writing.

(iii) Before a planned special exposure, the licensee or registrant ensures that each individual involved is:

(A) informed of the purpose of the planned operation; and

(B) informed of the estimated doses and associated potential risks and specific radiation levels or other conditions that might be involved in performing the task; and

(C) instructed in the measures to be taken to keep the dose ALARA considering other risks that may be present.

(iv) Prior to permitting an individual to participate in a planned special exposure, the licensee or registrant shall ascertain prior doses as required by § 175.03(c)(5)(ii) during the lifetime of the individual for each individual involved.

(v) Subject to § 175.03(c)(1)(ii), the licensee or registrant shall not authorize a planned special exposure that would cause an individual to receive a dose from all planned special exposures and all doses in excess of the limits to exceed:

(A) the numerical values of any of the dose limits in § 175.03(c)(1)(i) in any year; and

(B) five times the annual dose limits in § 175.03(c)(1)(i) during the individual's lifetime.

(vi) The licensee or registrant maintains records of the conduct of a planned special exposure in accordance with § 175.03(k)(7) and submits a written report in accordance with § 175.03(l)(4).

(vii) The licensee or registrant records the best estimate of the dose resulting from the planned special exposure in the individual's record and informs the individual, in writing, of the dose within 30 days from the date of the planned special exposure. The dose from planned special exposures shall not be considered in controlling future occupational dose of the individual pursuant to § 175.03(c)(1)(i) but shall be included in evaluations required by § 175.03(c)(6)(iv) and (v).

(7) Occupational dose limits for minors. The annual occupational dose limits for

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minors are 10 percent of the annual occupational dose limits specified for adult workers in § 175.03(c)(1).

(8)

Dose to an embryo/fetus. (i) The licensee or registrant shall ensure that the dose to an embryo/fetus during the entire pregnancy, due to occupational exposure of a declared pregnant woman, does not exceed 5 mSv (0.5 rem). (See § 175.03(k)(8) for recordkeeping requirements.)

(ii) The licensee or registrant shall review exposure history and adjust working conditions so as to avoid a monthly exposure of more than 0.5 mSv (50 mrem) to a declared pregnant woman.

(iii) The dose to an embryo/fetus shall be taken as the sum of:

(A) the deep dose equivalent to the declared pregnant woman; and

(B) the dose to the embryo/fetus from radionuclides in the embryo/fetus and radionuclides in the declared pregnant woman.

(iv) If, by the time the woman declares pregnancy to the licensee or registrant, the dose to the embryo/fetus has exceeded 4.5 mSv (0.45 rem), the licensee or registrant shall be deemed to be in compliance with § 175.03(c)(8)(i) if the additional dose to the embryo/fetus does not exceed 0.5 mSv (0.05 rem) during the remainder of the pregnancy.

(d) Radiation dose limits for individual members of the public.

(1) Dose limits for individual members of the public. (i) Each licensee or registrant shall conduct operations so that:

(A) the dose in any unrestricted area from external sources does not exceed 0.02 mSv (0.002 rem) in any one hour; and

(B) the total effective dose equivalent to individual members of the public from the licensed or registered operation, exclusive of the dose contribution from the licensee's or registrant's disposal of radioactive material into sanitary sewerage in accordance with § 175.104 of this Code, does not exceed 1 mSv (0.1 rem) in a year.

(ii) Notwithstanding the provisions of § 175.03(d)(1)(i)(B), where structural modification to the physical plant or equipment is required to meet the 1 mSv (0.1 rem) per year limit at facilities with radiation equipment installed before the effective date of these requirements and neither the use nor the physical components or structure of the facility are changed after the effective date of these requirements, the total effective dose equivalent to a member of the general public shall not exceed 5 mSv (0.5 rem) in a year. However, any such change occurring after the effective date of these requirements shall require the licensee or registrant to comply with the provisions of § 175.03(d)(1)(i)(B).

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(iii) The Department may impose additional restrictions on radiation levels in any unrestricted area and on the total quantity of radionuclides that a licensee may release in effluents in order to assure that the limits set forth in this Code are not exceeded.

(2) Compliance with dose limits for individual members of the public.

(i) The licensee or registrant shall make or cause to be made surveys of radiation levels in unrestricted areas and radioactive materials in effluents released to unrestricted areas to demonstrate compliance with the dose limits for individual members of the public in § 175.03(d)(1).

(ii) A licensee or registrant shall show compliance with the annual dose limit in § 175.03(d)(1) by:

(A) demonstrating by measurement or calculation that the total effective dose equivalent to the individual likely to receive the highest dose from the licensed or registered operation does not exceed the annual dose limit; or

(B) demonstrating that:

(a) the annual average concentrations of radioactive material released in gaseous and liquid effluents at the boundary of the unrestricted area do not exceed the values specified in Table 2 of Appendix B of this section; and

(b) if an individual were continually present in an unrestricted area, the dose from external sources would not exceed 0.02 mSv (0.002 rem) in an hour and 0.5 mSv (0.05 rem) in a year.

(iii) Upon approval from the Department, the licensee or registrant may adjust the effluent concentration values in Table 2, Appendix B of this section, for members of the public, to take into account the actual physical and chemical characteristics of the effluents, such as, aerosol size distribution, solubility, density, radioactive decay equilibrium, and chemical form.

(e) Testing for leakage or contamination of sealed sources.

(1) Testing for leakage or contamination of sealed sources. (i) The licensee or registrant in possession of any sealed source shall assure that:

(A) each sealed source, except as specified in § 175.03(e)(1)(ii), is tested for leakage or contamination and the test results are received before the sealed source is put into use unless the licensee or registrant has a certificate from the transferor indicating that the sealed source was tested within 6 months before transfer to the licensee or registrant; and

(B) each sealed source that is not designed to emit alpha particles is tested for leakage or contamination at intervals not to exceed 6 months; and

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(C) each sealed source that is designed to emit alpha particles is tested for leakage or contamination at intervals not to exceed 3 months; and

(D) for each sealed source that is required to be tested for leakage or contamination, at any other time there is reason to suspect that the sealed source might have been damaged or might be leaking, the licensee or registrant shall assure that the sealed source is tested for leakage or contamination before further use; and

(E) tests for leakage for all sealed sources, except brachytherapy sources manufactured to contain radium, shall be capable of detecting the presence of 185 Bq (0.005 <<mu>>Ci) of radioactive material on a test sample. Test samples shall be taken from the sealed source or from the surfaces of the container in which the sealed source is stored or mounted on which one might expect contamination to accumulate. For a sealed source contained in a device, test samples are obtained when the source is in the "off" position; and

(F) the test for leakage for brachytherapy sources manufactured to contain radium shall be capable of detecting an absolute leakage rate of 37 Bq (0.001 <<mu>> Ci) of radon-222 in a 24 hour period when the collection efficiency for radon-222 and its daughters has been determined with respect to collection method, volume and time; and

(G) tests for contamination from radium daughters shall be taken on the interior surface of brachytherapy source storage containers and shall be capable of detecting the presence of 185 Bq (0.005 <<mu>>Ci) of a radium daughter which has a half-life greater than four (4) days; and

(H) the test for leakage for sealed sources containing iodine-125 shall be capable of detecting an absolute leakage rate of 185 Bq (0.005 <<mu>>Ci) in a 24 hour period.

(ii) A licensee or registrant need not perform tests for leakage or contamination on the following:

(A) sealed sources containing only radioactive material with a half-life of less than 30 days;

(B) sealed sources containing only radioactive material as a gas;

(C) sealed sources containing 3.7 MBq (100 <<mu>>Ci) or less of beta or photon-emitting material or 370 kBq (10 <<mu>>Ci) or less of alpha-emitting material;

(D) sealed sources containing only hydrogen-3;

(E) seeds of iridium-192 encased in nylon ribbon; and

(F) sealed sources, except teletherapy and brachytherapy sources, which are stored, not being used and identified as in storage. The licensee or registrant shall, however, test each such sealed source for leakage or contamination and receive the test results before any use or transfer unless it has been tested for

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leakage or contamination within 6 months before the date of use or transfer.

(iii) Tests for leakage or contamination from sealed sources shall be performed by persons specifically authorized by the Department to perform such services.

(iv) Test results shall be kept in units of becquerels (or microcuries) and maintained for inspection by the Department.

(v) The following shall be considered evidence that a sealed source is leaking:

(A) The presence of 185 Bq (0.005 <<mu>>Ci) or more of removable contamination on any test sample.

(B) Leakage of 37 Bq (0.001 <<mu>>Ci) of radon-222 per 24 hours for brachytherapy sources manufactured to contain radium.

(C) The presence of removable contamination resulting from the decay of 185 Bq (0.005 <<mu>>Ci) or more of radium.

(D) Leakage of 185 Bq (0.005 <<mu>>Ci) of iodine-125 per 24 hours for sealed sources containing iodine-125.

(vi) The licensee or registrant shall immediately withdraw a leaking sealed source from use and shall take action to prevent the spread of contamination. The leaking sealed source shall be repaired or disposed of in accordance with this Code.

(vii) Reports of test results for leaking or contaminated sealed sources shall be made pursuant to § 175.03(l)(7).

(f) Surveys and monitoring. (1) General. (i) Each licensee or registrant shall make, or cause to be made, surveys that:

(A) are necessary for the licensee or registrant to comply with this Code; and

(B) are necessary under the circumstances to evaluate:

(a) radiation levels; and

(b) concentrations or quantities of radioactive material; and

(c) the potential radiological hazards that could be present.

(ii) The licensee or registrant shall ensure that instruments and equipment used for quantitative radiation measurements, for example, dose rate and effluent monitoring, are calibrated at intervals not to exceed 12 months for the radiation measured.

(2) Personnel monitoring. (i) External radiation sources. Each person who pos-

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sesses any radiation source shall supply and require the proper use of appropriate, calibrated and operable individual monitoring devices by:

(A) adults likely to receive, in 1 year from sources external to the body, a dose in excess of 10 percent of the limits in § 175.03(c)(1)(i) and

(B) minors and declared pregnant women likely to receive, in 1 year from sources external to the body, a dose in excess of 10 percent of any of the applicable limits in § 175.03(c)(7) or § 175.03(c)(8); and

(C) individuals entering a high or very high radiation area.

(ii) A person supplying personnel monitoring devices to individuals pursuant to § 175.03(f)(2)(i) shall ensure that the individuals wear such devices as follows:

(A) An individual monitoring device used for monitoring the dose to the whole body shall be worn at the unshielded location of the whole body likely to receive the highest exposure.

(B) An individual monitoring device used for monitoring the dose to an embryo/fetus of a declared pregnant woman pursuant to § 175.03(c)(8) shall be located at the waist under any protective apron worn by the woman.

(C) An individual monitoring device used for monitoring the eye dose equivalent shall be located at the neck outside any protective apron worn by the individual, or at an unshielded location closer to the eye.

(D) An individual monitoring device used for monitoring the dose to the extremities shall be worn on the extremity likely to receive the highest exposure. The device shall be oriented to measure the highest dose to the extremity being monitored.

(iii) All personnel dosimeters, except for direct and indirect reading pocket ionization chambers and those dosimeters used to measure the dose to any extremity, that require processing to determine the radiation dose and that are used by licensees and registrants to comply with § 175.03(c)(1), with other applicable provisions of this Code, or with conditions specified in a license, certified registration or registration shall be processed and evaluated by a dosimetry processor:

(A) holding current personnel dosimetry accreditation from the National Voluntary Laboratory Accreditation Program (NVLAP) of the National Institute of Standards and Technology; and

(B) approved in this accreditation process for the type of radiation or radiations included in the NVLAP program that most closely approximates the type of radiation or radiations for which the individual wearing the dosimeter is monitored.

(iv) The licensee or registrant shall ensure that adequate precautions are taken to prevent a deceptive exposure of an individual monitoring device.

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(A) No licensee or registrant shall remove an exposure from an individual's exposure record without prior authorization from the Department.

(v) Each licensee or registrant shall monitor, to determine compliance with § 175.03(c)(4), the occupational intake of radioactive material by and assess the committed effective dose equivalent to:

(A) adults likely to receive, in 1 year, an intake in excess of 10 percent of the applicable ALI in Table 1, Columns 1 and 2, of Appendix B of this section; and

(B) minors and declared pregnant women likely to receive, in 1 year, a committed effective dose equivalent in excess of 0.5 mSv (0.05 rem).

(vi) The licensee or registrant shall submit the dosimeter for processing with due diligence and in no event in excess of the time period specified by the manufacturer of the dosimeter.

(g) Control of exposure from external sources in restricted areas. (1) Control of access to high radiation areas. (i) The licensee or registrant shall ensure that each entrance or access point to a high radiation area has one or more of the following features:

(A) a control device that, upon entry into the area, causes the level of radiation to be reduced below that level at which an individual might receive a deep dose equivalent of 1 mSv (0.1 rem) in 1 hour at 30 cm (12 in.) from the source of radiation or from any surface that the radiation penetrates; or

(B) a control device that energizes a conspicuous visible or audible alarm signal so that the individual entering the high radiation area and the supervisor of the activity are made aware of the entry; or

(C) entryways that are locked, except during periods when access to the areas is required, with positive control over each individual entry.

(ii) In place of the controls required by § 175.03(g)(1)(i) for a high radiation area, the licensee or registrant may substitute continuous direct or electronic surveillance that is capable of preventing unauthorized entry.

(iii) The licensee or registrant may apply to the Department for approval of alternative methods for controlling access to high radiation areas.

(iv) The licensee or registrant shall establish the controls required by § 175.03(g)(1)(i), (ii) or (iii) in a way that does not prevent individuals from leaving a high radiation area.

(v) The licensee or registrant is not required to control each entrance or access point to a room or other area that is a high radiation area solely because of the presence of radioactive materials prepared for transport and packaged and labeled in accordance with the regulations of the U.S. Department of Transportation provided that:

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(A) the packages do not remain in the area longer than 3 days; and

(B) the dose rate at 1 m (3 ft) from the external surface of any package does not exceed 0.1 mSv (0.01 rem) per hour.

(vi) The licensee or registrant is not required to control entrance or access to rooms or other areas in hospitals solely because of the presence of patients containing radioactive material, provided that there are personnel in attendance who are taking the necessary precautions to prevent the exposure of individuals to radiation or radioactive material in excess of the established limits in this section and to operate within the ALARA provisions of the licensee's or registrant's radiation protection program.

(vii) The registrant is not required to control entrance or access to rooms or other areas containing sources of radiation capable of producing a high radiation area as described in § 175.03(g)(1) if the registrant has met all the specific requirements for access and control specified in other applicable sections of this Code.

(2) Control of access to very high radiation areas. (i) In addition to the requirements in § 175.03(g)(1), the licensee or registrant shall institute measures to ensure that an individual is not able to gain unauthorized or inadvertent access to areas in which radiation levels could be encountered at 5 Gy (500 rad) or more in 1 hour at 1 m (3 ft) from a source of radiation or any surface through which the radiation penetrates. This requirement does not apply to rooms or areas in which diagnostic x-ray systems are the only source of radiation, or to non-self-shielded irradiators.

(ii) The registrant is not required to control entrance or access to rooms or other areas containing sources of radiation capable of producing a very high radiation area as described in § 175.03(g)(2)(i) if the registrant has met all the specific requirements for access and control specified in other applicable sections of this Code.

(3) Control of access to very high radiation areas--irradiators. (i) This subdivision applies to licensees or registrants with sources of radiation in non-self-shielded irradiators. Section 175.03(g)(3) does not apply to sources of radiation that are used in teletherapy, in industrial radiography, or in completely self-shielded irradiators in which the source of radiation is both stored and operated within the same shielding radiation barrier and, in the designed configuration of the irradiator, is always physically inaccessible to any individual and cannot create high levels of radiation in an area that is accessible to any individual.

(ii) Each area in which there may exist radiation levels in excess of 5 Gy (500 rad) in 1 hour at 1 m (3 ft) from a source of radiation that is used to irradiate materials shall meet the following requirements:

(A) Each entrance or access point shall be equipped with entry control devices which:

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(a) function automatically to prevent any individual from inadvertently entering a very high radiation area; and

(b) permit deliberate entry into the area only after a control device is actuated that causes the radiation level within the area, from the source of radiation, to be reduced below that at which it would be possible for an individual to receive a deep dose equivalent in excess of 1 mSv (0.1 rem) in 1 hour; and

(c) prevent operation of the source of radiation if it would produce radiation levels in the area that could result in a deep dose equivalent to an individual in excess of 1 mSv (0.1 rem) in 1 hour.

(B) Additional control devices shall be provided so that, upon failure of the entry control devices to function as required by § 175.03(g)(3)(ii)(A):

(a) the radiation level within the area, from the source of radiation, is reduced below that at which it would be possible for an individual to receive a deep dose equivalent in excess of 1 mSv (0.1 rem) in 1 hour; and

(b) conspicuous visible and audible alarm signals are generated to make an individual attempting to enter the area aware of the hazard and at least one other authorized individual, who is physically present, familiar with the activity, and prepared to render or summon assistance, aware of the failure of the entry control devices.

(C) The licensee or registrant shall provide control devices so that, upon failure or removal of physical radiation barriers other than the sealed source's shielded storage container:

(a) the radiation level from the source of radiation is reduced below that at which it would be possible for an individual to receive a deep dose equivalent in excess of 1 mSv (0.1 rem) in 1 hour; and

(b) conspicuous visible and audible alarm signals are generated to make potentially affected individuals aware of the hazard and the licensee or registrant or at least one other individual, who is familiar with the activity and prepared to render or summon assistance, aware of the failure or removal of the physical barrier.

(D) When the shield for stored sealed sources is a liquid, the licensee or registrant shall provide means to monitor the integrity of the shield and to signal, automatically, loss of adequate shielding.

(E) Physical radiation barriers that comprise permanent structural components, such as walls, that have no credible probability of failure or removal in ordinary circumstances need not meet the requirements of § 175.03(g)(3)(ii)(C) and (D).

(F) Each area shall be equipped with devices that will automatically generate conspicuous visible and audible alarm signals to alert personnel in the area before

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the source of radiation can be put into operation and in time for any individual in the area to operate a clearly identified control device, which must be installed in the area and which can prevent the source of radiation from being put into operation.

(G) Each area shall be controlled by use of such administrative procedures and such devices as are necessary to ensure that the area is cleared of personnel prior to each use of the source of radiation.

(H) Each area shall be checked by a radiation measurement to ensure that, prior to the first individual's entry into the area after any use of the source of radiation, the radiation level from the source of radiation in the area is below that at which it would be possible for an individual to receive a deep dose equivalent in excess of 1 mSv (0.1 rem) in 1 hour.

(I) The entry control devices required in § 175.03(g)(3)(ii)(A) shall be tested for proper functioning. (See § 175.03(k)(10) for recordkeeping requirements.)

(a) Testing shall be conducted prior to initial operation with the source of radiation on any day, unless operations were continued uninterrupted from the previous day; and

(b) testing shall be conducted prior to resumption of operation of the source of radiation after any unintentional interruption; and

(c) the licensee or registrant shall submit and adhere to a schedule for periodic tests of the entry control and warning systems.

(J) The licensee or registrant shall not conduct operations, other than those necessary to place the source of radiation in safe condition or to effect repairs on controls, unless control devices are functioning properly.

(K) Entry and exit portals that are used in transporting materials to and from the irradiation area, and that are not intended for use by individuals, shall be controlled by such devices and administrative procedures as are necessary to physically protect and warn against inadvertent entry by any individual through these portals. Exit portals for irradiated materials shall be equipped to detect and signal the presence of any loose radioactive material that is carried toward such an exit and automatically to prevent loose radioactive material from being carried out of the area.

(iii) Licensees, registrants, or applicants for licenses or registrations for sources of radiation within the purview of § 175.03(g)(3)(ii) which will be used in a variety of positions or in locations, such as open fields or forests, that make it impracticable to comply with certain requirements of § 175.03(g)(3)(ii), such as those for the automatic control of radiation levels, may apply to the Department for approval of alternative safety measures. Alternative safety measures shall provide personnel protection at least equivalent to those specified in § 175.03(g)(3)(ii). At least one of the alternative measures shall include an entry-preventing interlock control based on a measurement of the radiation that ensures the absence of high radiation levels before an individual can gain access to the

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area where such sources of radiation are used.

(iv) The entry control devices required by § 175.03(g)(3)(ii) and (iii) shall be established in such a way that no individual will be prevented from leaving the area.

(h) Respiratory protection and controls to restrict internal exposure in restricted areas. (1) Use of process or other engineering controls. The licensee or registrant shall use to the extent practicable, process or other engineering controls, such as containment, decontamination, or ventilation, to control the concentrations of radioactive material in air.

(2) Use of other controls. When it is not practicable to apply process or other engineering controls to control the concentrations of radioactive material in air to values below those that define an airborne radioactivity area, the licensee or registrant shall, consistent with maintaining the total effective dose equivalent ALARA, increase monitoring and limit intakes by one or more of the following means:

(i) control of access; or

(ii) limitation of exposure times; or

(iii) use of respiratory protection equipment; or

(iv) other controls.

If the licensee performs an ALARA analysis to determine whether or not respirators should be used, the licensee may consider safety factors other than radiological factors. The licensee should also consider the impact of respirator use on workers' industrial health and safety.

(3) Use of individual respiratory protection equipment. (i) If the licensee or registrant uses respiratory protection equipment to limit intakes pursuant to § 175.03(h)(2):

(A) except as provided in § 175.03(h)(3)(i)(B), the licensee or registrant shall use only respiratory protection equipment that is tested and certified or had certification extended by the National Institute for Occupational Safety and Health and the Mine Safety and Health Administration.

(B) If the licensee or registrant wishes to use equipment that has not been tested or certified by the National Institute for Occupational Safety and Health and the Mine Safety and Health Administration or has not had certification extended by the National Institute for Occupational Safety and Health and the Mine Safety and Health Administration, or for which there is no schedule for testing or certification, the licensee or registrant shall submit an application to the Department for authorized use of that equipment. The application must include a demonstration by licensee testing or a demonstration on the basis of reliable test information, that the material and performance characteristics of the equipment are capable of providing the proposed degree of protection under anticipated conditions of use.

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(C) The licensee or registrant shall implement and maintain a respiratory protection program that includes:

(a) air sampling sufficient to identify the potential hazard, permit proper equipment selection, and estimate doses; and

(b) surveys and bioassays, as necessary, to evaluate actual intakes; and

(c) testing of respirators for operability (user seal check for face sealing devices and functional check for others) immediately prior to each use; and

(d) written procedures regarding respirator selection; fit testing; inventory and control, storage, issuance; maintenance, repair and testing of respirators, including testing for operability immediately prior to each use; supervision and training of respirator users; monitoring, including air sampling and bioassays, and breathing air quality; and quality assurance and recordkeeping; and

(e) determination by a physician that the individual user is medically fit to use the respiratory protection equipment prior to initial fitting of a face sealing respirator; before the first field use of non-face sealing respirators and either every 12 months thereafter, or periodically at a frequency determined by a physician.

(f) Fit testing, with a fit factor 10 times the APF for negative pressure devices, and a fit factor greater than or equal to 500 for any positive pressure, continuous flow, and pressure demand devices, before the first field use of tight fitting, face sealing respirators and periodically thereafter at a frequency of at least once per year. Fit testing must be performed with the face-piece operating in the negative pressure mode.

(D) The licensee or registrant shall issue a written policy statement on respirator usage covering:

(a) the use of process or other engineering controls, instead of respirators; and

(b) the routine, nonroutine, and emergency use of respirators; and

(c) limitations on periods of respirator use and relief from respirator use.

(E) The licensee or registrant shall advise each respirator user that the user may leave the area at any time for relief from respirator use in the event of equipment malfunction, physical or psychological distress, procedural or communication failure, significant deterioration of operating conditions, or any other conditions that might require such relief.

(F) The licensee or registrant shall use respiratory protection equipment within the equipment manufacturer's expressed limitations for type and mode of use. When selecting respiratory devices, the licensee shall provide for low temperature work environments, and the concurrent use of other safety or radiological protection

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equipment or skin protection, when needed. The licensee shall use equipment in such a way as not to interfere with the proper operation of the respirator.

(ii) When estimating exposure of individuals to airborne radioactive materials, the licensee or registrant may make allowance for respiratory protection equipment used to limit intakes pursuant to § 175.03(h)(2), provided that the following conditions, in addition to those in § 175.03(h)(3)(i), are satisfied:

(A) the licensee or registrant selects respiratory protection equipment that provides a protection factor, specified in Appendix A of this section, greater than the multiple by which peak concentrations of airborne radioactive materials in the working area are expected to exceed the values specified in Table 1, Column 3 of Appendix B of this section. However, if the selection of respiratory protection equipment with a protection factor greater than this multiple of peak concentration is inconsistent with the goal specified in § 175.03(h)(2) of keeping the total effective dose equivalent ALARA, the licensee or registrant may select respiratory protection equipment with a lower protection factor provided that such a selection would result in a total effective dose equivalent that is ALARA. In estimating the dose to individuals from intake of airborne radioactive materials, the concentration of radioactive material in the air that is inhaled when respirators are worn is initially assumed to be the ambient concentration in air without respiratory protection, divided by the assigned protection factor. If the dose is later found to be greater than the initially estimated dose, the corrected value must be used. If the dose is later found to be less than the initially estimated dose, the corrected value may be used.

(B) The licensee or registrant shall obtain authorization from the Department before using assigned protection factors in excess of those specified in Appendix A of this section. The Department may authorize a licensee or registrant to use higher protection factors on receipt of an application that:

(a) describes the situation for which a need exists for higher protection factors, and

(b) demonstrates that the respiratory protection equipment provides these higher protection factors under the proposed conditions of use.

(iii) In an emergency, the licensee or registrant shall use as emergency equipment only respiratory protection equipment that has been specifically certified or had certification extended for emergency use by the National Institute for Occupational Safety and Health and the Mine Safety and Health Administration.

(iv) The licensee or registrant shall notify the Department in writing at least 30 days before the date that respiratory protection equipment is first used pursuant to either § 175.03(h)(3)(i) or (ii).

(4) Further restrictions on the use of respiratory protection equipment. The Department may impose restrictions, in addition to those in § 175.03(h)(2), § 175.03(h)(3) and in Appendix A to § 175.03 to:

(i) ensure that the respiratory protection program of the licensee is adequate to

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limit doses to individuals from intakes of airborne radioactive materials consistent with maintaining total effective dose equivalent ALARA; and

(ii) limit the extent to which a licensee may use respiratory protection equipment instead of process or other engineering controls.

(i) Storage and control of licensed or registered sources of radiation. (1) Security of stored sources of radiation. The licensee or registrant shall secure from unauthorized removal or access licensed or registered sources of radiation that are stored in unrestricted areas.

(2) Radioactive materials shall not be stored with either food or beverages.

(3) Control of sources of radiation not in storage.

(i) The licensee or registrant shall control and maintain constant surveillance of licensed radioactive material that is in an unrestricted area and that is not in storage or in a patient.

(ii) The registrant shall maintain control of radiation machines that are in an unrestricted area and that are not in storage.

(j) Precautionary procedures. (1) Caution Signs. (i) Standard radiation symbol. Unless otherwise authorized by the Department, the symbol prescribed herein shall use the colors magenta, or purple, or black on yellow background. The symbol prescribed is the three-bladed design as follows:

RADIATION SYMBOL

(A) Cross-hatched area is to be magenta, or purple, or black, and

(B) the background is to be yellow.

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(ii) Exception to color requirements for standard radiation symbol. Notwithstanding the requirements of § 175.03(j)(1)(i), licensees or registrants are authorized to label sources, source holders, or device components containing sources of radiation that are subjected to high temperatures, with conspicuously etched or stamped radiation caution symbols and without a color requirement.

(iii) Additional information on signs and labels. In addition to the contents of signs and labels prescribed in this section, the licensee or registrant shall provide, on or near the required signs and labels, additional information, as appropriate, to make individuals aware of potential radiation exposures and to minimize the exposures.

(2) Posting requirements. (i) Posting of radiation areas. The licensee or registrant shall post each radiation area with a conspicuous sign or signs bearing the radiation symbol and the words "CAUTION, RADIATION AREA."

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(ii) Posting of high radiation areas. The licensee or registrant shall post each high radiation area with a conspicuous sign or signs bearing the radiation symbol and the words "CAUTION, HIGH RADIATION AREA" or "DANGER, HIGH RADIATION AREA."

(iii) Posting of very high radiation areas. The licensee or registrant shall post each very high radiation area with a conspicuous sign or signs bearing the radiation symbol and words "GRAVE DANGER, VERY HIGH RADIATION AREA."

(iv) Posting of airborne radioactivity areas. The licensee or registrant shall post each airborne radioactivity area with a conspicuous sign or signs bearing the radiation symbol and the words "CAUTION, AIRBORNE RADIOACTIVITY AREA" or "DANGER, AIRBORNE RADIOACTIVITY AREA."

(v) Posting of areas or rooms in which licensed radioactive material is used or stored. The licensee shall post each area or room in which there is used or stored an amount of licensed material exceeding 10 times the quantity of such material specified in Appendix C of this section with a conspicuous sign or signs bearing the radiation symbol and the words "CAUTION, RADIOACTIVE MATERIAL(S)" or "DANGER, RADIOACTIVE MATERIAL(S)."

(vi) Such cautionary postings shall not be used except as required by this Code.

(3) Exceptions to posting requirements. (i) A licensee or registrant is not required to post caution signs in areas or rooms containing sources of radiation for periods of less than 8 hours, if each of the following conditions is met:

(A) the sources of radiation are constantly attended during these periods by an individual who takes the precautions necessary to prevent the exposure of individuals to sources of radiation in excess of the limits established in this section; and

(B) the area or room is subject to the licensee's or registrant's control.

(ii) Rooms or other areas in hospitals that are occupied by patients are not required to be posted with caution signs pursuant to § 175.03(j)(2) provided that the patient could be released from confinement pursuant to § 175.103(c)(9) of this Code.

(iii) A room or area is not required to be posted with a caution sign because of the presence of a sealed source provided the radiation level at 30 cm (12 in.) from the surface of the sealed source container or housing does not exceed 0.05 mSv (0.005 rem) per hour.

(iv) A room or area is not required to be posted with a caution sign because of the presence of radiation machines used solely for diagnosis in the healing arts.

(4) Labeling containers and radiation machines. (i) The licensee or registrant shall ensure that each container of licensed radioactive material bears a durable, clearly visible label bearing the radiation symbol and the words "CAUTION, RADIOAC-

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TIVE MATERIAL" or "DANGER, RADIOACTIVE MATERIAL." The label shall also provide information, such as the radionuclides present, an estimate of the quantity of radioactivity, the date for which the activity is estimated, radiation levels, kinds of materials, and mass enrichment, to permit individuals handling or using the containers, or working in the vicinity of the containers, to take precautions to avoid or minimize exposures.

(ii) Each licensee shall, prior to removal or disposal of empty uncontaminated containers to unrestricted areas, remove or deface the radioactive material label or otherwise clearly indicate that the container no longer contains radioactive materials.

(iii) Each registrant shall ensure that each radiation machine is labeled in a conspicuous manner which cautions individuals that radiation is produced when it is energized.

(iv) Such cautionary labels shall not be used except as required by this Code.

(5) Exemptions to labeling requirements. A licensee or registrant is not required to label:

(i) containers holding licensed radioactive material in quantities less than the quantities listed in Appendix C of this section; or

(ii) containers holding licensed radioactive material in concentrations less than those specified in Table 3 of Appendix B of this section; or

(iii) containers attended by an individual who takes the precautions necessary to prevent the exposure of individuals in excess of the limits established by this section; or

(iv) containers when they are in transport and packaged and labeled in accordance with the regulations of the U.S. Department of Transportation; or

(v) containers that are accessible only to individuals authorized to handle or use them, or to work in the vicinity of the containers, if the contents are identified to these individuals by a readily available written record. Examples of containers of this type are containers in locations such as water-filled canals, storage vaults, or hot cells. The record shall be retained as long as the containers are in use for the purpose indicated on the record; or

(vi) installed manufacturing or process equipment, such as piping and tanks.

(6) Procedures for receiving and opening packages. (i) Each licensee who expects to receive a package containing quantities of radioactive material in excess of a Type A quantity, as defined in § 175.02(a)(232) and Appendix A of § 175.105 of this Code, shall make arrangements to receive:

(A) the package when the carrier offers it for delivery; or

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(B) notification of the arrival of the package at the carrier's terminal and to take possession of the package expeditiously.

(ii) Each licensee shall:

(A) monitor the external surfaces of a package labeled with a U.S. Department of Transportation specified radioactive White I, Yellow II or Yellow III label for radioactive contamination unless the package contains only radioactive material in the form of gas or in special form as defined in § 175.02(a)(211) of this Code; and

(B) monitor the external surfaces of a package labeled with a U.S. Department of Transportation specified radioactive White I, Yellow II or Yellow III label for radiation levels unless the package contains quantities of radioactive material that are less than or equal to the Type A quantity, as defined in § 175.02(a)(232) and Appendix A of § 175.105 of this Code; and

(C) monitor all packages known to contain radioactive material for radioactive contamination and radiation levels if there is evidence of degradation of package integrity, such as packages that are crushed, wet, or damaged.

(iii) The licensee or registrant shall perform the monitoring required by § 175.03 (j)(6)(ii) as soon as practicable after receipt of the package, but not later than 3 hours after the package is received at the licensee's or registrant's facility if it is received during the licensee's or registrant's normal working hours, or not later than 3 hours from the beginning of the next working day if it is received after working hours.

(iv) The licensee or registrant shall immediately notify the final delivery carrier and, by telephone and telegram, mailgram, or facsimile, the Department when:

(A) removable radioactive surface contamination exceeds the limits specified in § 175.105 of the Code; or

(B) external radiation levels exceed the limits specified in § 175.105 of this Code.

(v) Each licensee or registrant shall:

(A) establish, maintain, and retain written procedures for safely opening packages in which radioactive material is received; and

(B) ensure that the procedures are followed and that due consideration is given to special instructions for the type of package being opened.

(vi) Licensees or registrants transferring special form sources in vehicles owned or operated by the licensee or registrant to and from a work site are exempt from the contamination monitoring requirements of § 175.03(j)(6)(ii), but are not exempt from the monitoring requirement in § 175.03(j)(6)(ii)(B) for measuring radiation levels that ensures that the source is still properly lodged in its shield.

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(k) Records. (1) General provisions. (i) Each licensee or registrant shall use SI units (becquerel, gray, sievert and coulomb per kilogram) or special units (curie, rad, rem and roentgen) including multiples and subdivisions, and shall clearly indicate the units of all quantities on records required by this Code.

(ii) The licensee or registrant shall make a clear distinction between the quantities entered on the records required by this section, such as, total effective dose equivalent, total organ dose equivalent, shallow dose equivalent, eye dose equivalent, deep dose equivalent, or committed effective dose equivalent.

(2) Records of radiation protection programs. (i) Each licensee or registrant shall maintain records of the radiation protection program, including:

(A) the provisions of the program; and

(B) audits and other reviews of program content and implementation.

(ii) The licensee or registrant shall retain the records required by § 175.03(k)(2)(i)(A) until the Department terminates each pertinent license, certified registration or registration requiring the record. The licensee or registrant shall retain the records required by § 175.03(k)(2)(i)(B) for 3 years after the record is made.

(3) Records of receipt, use and disposition of radioactive material. (i) Each licensee shall maintain records of the receipt, use and disposition of radioactive material in units of becquerels or microcuries and shall include from whom such materials were received and the ultimate disposition.

(ii) The licensee shall retain the records required by § 175.03(k)(3)(i) for 3 years after the record is made.

(4) Records of surveys. (i) Each licensee or registrant shall maintain records showing the results of surveys and calibrations required by § 175.03(f) and § 175.03(j)(6)(ii). The licensee or registrant shall retain these records for 3 years after the record is made.

(ii) The licensee or registrant shall retain each of the following records until the Department authorizes the disposition of these records:

(A) records of the results of surveys to determine the dose from external sources of radiation used, in the absence of or in combination with individual monitoring data, in the assessment of individual dose equivalents; and

(B) records of the results of measurements and calculations used to determine individual intakes of radioactive material and used in the assessment of internal dose; and

(C) records showing the results of air sampling, surveys, and bioassays required pursuant to § 175.03(h)(3)(i)(C)(a) and (b); and

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(D) records of the results of measurements and calculations used to evaluate the release of radioactive effluents to the environment.

(5) Records of tests for leakage or contamination of sealed sources. Records of tests for leakage or contamination of sealed sources required by § 175.03(e)(1) shall be kept in units of becquerel or microcurie and maintained for inspection by the Department for 5 years after the records are made.

(6) Records of prior occupational dose. (i) The licensee or registrant shall retain the records of prior occupational dose and exposure history as specified in § 175.03(c)(5) on form RAD-4, "Cumulative Occupational Radiation Exposure History" or equivalent, and the records used in preparing form RAD-4 until the Department authorizes their disposition.

(ii) Upon termination of the license or registration, the licensee or registrant shall permanently store records on form RAD-4, "Cumulative Occupational Radiation Exposure History," or equivalent, or shall make provision with the Department for transfer to the Department.

(7) Records of planned special exposures. (i) For each use of the provisions of § 175.03(c)(6) for planned special exposures, the licensee or registrant shall maintain records that describe:

(A) the exceptional circumstances requiring the use of a planned special exposure; and

(B) the name of the management official who authorized the planned special exposure and a copy of the signed authorization; and

(C) what actions were necessary; and

(D) why the actions were necessary; and

(E) what precautions were taken to assure that doses were main-

tained ALARA; and

(F) what individual and collective doses were expected to result; and

(G) the doses actually received in the planned special exposure.

(ii) The licensee or registrant shall retain the records until the Department authorizes their disposition.

(iii) Upon termination of the license or registration, the licensee or registrant shall permanently store records on form RAD-4, "Cumulative Occupational Radiation Exposure History," or until disposition is authorized by the Department.

(8) Records of individual monitoring results. (i) Recordkeeping Requirement. Each

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licensee or registrant shall maintain records of doses received by all individuals for whom monitoring was required pursuant to § 175.03(f)(2), and records of doses received during planned special exposures, accidents, and emergency conditions. Assessments of dose equivalent and records made using units in effect before the effective date of these requirements need not be changed. These records shall include, when applicable:

(A) the deep dose equivalent to the whole body, eye dose equivalent, shallow dose equivalent to the skin, and shallow dose equivalent to the extremities; and

(B) the estimated intake or body burden of radionuclides (see § 175.03(c)(2)); and

(C) the committed effective dose equivalent assigned to the intake or body burden of radionuclides; and

(D) the specific information used to calculate the committed effective dose equivalent pursuant to § 175.03(c)(4)(iii); and

(E) the total effective dose equivalent when required by § 175(c)(2); and

(F) the total of the deep dose equivalent and the committed dose to the organ receiving the highest total dose.

(ii) Recordkeeping frequency. The licensee or registrant shall make entries of the records specified in § 175.03(k)(8)(i) at intervals not to exceed 1 year.

(iii) Recordkeeping format. The licensee or registrant shall maintain the records specified in § 175.03(k)(8)(i) on form RAD-5, in accordance with the instructions for form RAD-5, or in clear and legible records containing all the information required by form RAD-5.

(iv) The licensee or registrant shall maintain the records of dose to an embryo/fetus with the records of dose to the declared pregnant woman. The declaration of pregnancy, including the estimated date of conception, shall also be kept on file, but may be maintained separately from the dose records.

(v) The licensee or registrant shall retain each required form or record until the Department authorizes disposition.

(vi) Upon termination of the license or registration, the licensee or registrant shall permanently store records on form RAD-4 or equivalent, or shall make provision with the Department for transfer to the Department.

(9) Records of dose to individual members of the public. (i) Each licensee or registrant shall maintain records sufficient to demonstrate compliance with the dose limit for individual members of the public as specified in § 175.03(d).

(ii) The licensee or registrant shall retain the records required by § 175.03(k)(9)(i) until the Department terminates each pertinent license, certified

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registration or registration requiring the record.

(10) Records of testing entry control devices for very high radiation areas.

(i) Each licensee or registrant shall maintain records of tests made pursuant to § 175.03(g)(3)(ii)(I) on entry control devices for very high radiation areas. These records must include the date, time, and results of each such test of function.

(ii) The licensee or registrant shall retain the records required by § 175.03(k)(10)(i) for 3 years after the record is made.

(11) Form of records. (i) Each record required by this section shall be legible throughout the specified retention period. The record shall be the original or a reproduced copy or a microform, provided that the copy or microform is authenticated by authorized personnel and that the microform is capable of producing a clear copy throughout the required retention period. The record may also be stored in electronic media with the capability for producing legible, accurate, and complete records during the required retention period. Records, such as letters, drawings, and specifications, shall include all pertinent information, such as stamps, initials, and signatures.

(ii) The licensee shall maintain adequate safeguards against tampering with and loss of records.

(iii) The discontinuance or curtailment of activities does not relieve any person who possesses any radiation source of responsibility for retaining all records required by this Code.

(l) Reports. (1) Reports of stolen, lost, or missing licensed or registered sources of radiation. (i) Telephone reports. Each licensee or registrant shall report to the Department by telephone as follows:

(A) immediately after its occurrence becomes known to the licensee or registrant, stolen, lost, or missing licensed or registered radioactive material in an aggregate quantity equal to or greater than 1,000 times the quantity specified in Appendix C of this section under such circumstances that it appears to the licensee or registrant that an exposure could result to individuals in unrestricted areas; or

(B) within 30 days after its occurrence becomes known to the licensee or registrant, lost, stolen, or missing licensed or registered radioactive material in an aggregate quantity greater than 10 times the quantity specified in Appendix C of this section that is still missing.

(C) immediately after its occurrence becomes known to the registrant, a stolen, lost, or missing radiation machine.

(ii) Written reports. Each licensee or registrant required to make a report pursuant to § 175.03(l)(1)(i) shall, within thirty (30) days after making the telephone report, make a written report to the Department setting forth the following information:

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(A) a description of the licensed or registered source of radiation involved, including, for radioactive material, the kind, quantity, and chemical and physical form; and, for radiation machines, the manufacturer, model and serial number, type and maximum energy of radiation emitted;

(B) a description of the circumstances under which the loss or theft occurred; and

(C) a statement of disposition, or probable disposition, of the licensed or registered source of radiation involved; and

(D) exposures of individuals to radiation, circumstances under which the exposures occurred, and the possible total effective dose equivalent to persons in unrestricted areas; and

(E) actions that have been taken, or will be taken, to recover the source of radiation; and

(F) procedures or measures that have been, or will be, adopted to ensure against a recurrence of the loss or theft of licensed or registered sources of radiation.

(iii) Subsequent to filing the written report, the licensee or registrant shall also report additional substantive information on the loss or theft within thirty (30) days after the licensee or registrant learns of such information.

(iv) The licensee or registrant shall prepare any report filed with the Department pursuant to § 175.03(l)(1)(ii) so that names of individuals who may have received exposure to radiation are stated in a separate and detachable portion of the report.

(2) Notification of incidents. (i) Immediate notification. Notwithstanding other requirements for notification, each licensee or registrant shall immediately report each event involving a source of radiation possessed by the licensee or registrant that may have caused or threatens to cause any of the following conditions:

(A) an individual to receive:

(a) a total effective dose equivalent of 0.25 Sv (25 rem) or more; or

(b) an eye dose equivalent of 0.75 Sv (75 rem) or more; or

(c) a shallow dose equivalent to the skin or extremities or a total organ dose equivalent of 2.5 Gy (250 rad) or more; or

(B) the release of radioactive material, inside or outside of a restricted area, so that, had an individual been present for 24 hours, the individual could have received an intake five (5) times the occupational ALI. This provision does not apply to locations where personnel are not normally stationed during routine operations, such as hot-cells or process enclosures.

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(ii) Twenty-four hour notification. Each licensee or registrant shall, within 24 hours of discovery of the event, report to the Department each event involving loss of control of a licensed or registered source of radiation possessed by the licensee or registrant that may have caused, or threatens to cause, any of the following conditions:

(A) an individual to receive, in a period of 24 hours:

(a) a total effective dose equivalent exceeding 0.05 Sv (5 rem); or

(b) an eye dose equivalent exceeding 0.15 Sv (15 rem); or

(c) a shallow dose equivalent to the skin or extremities or a total organ dose equivalent exceeding 0.5 Sv (50 rem); or

(B) the release of radioactive material, inside or outside of a restricted area, so that, had an individual been present for 24 hours, the individual could have received an intake in excess of one occupational ALI. This provision does not apply to locations where personnel are not normally stationed during routine operations, such as hot-cells or process enclosures.

(iii) The licensee or registrant shall prepare each report filed with the Department pursuant to § 175.03(l)(2) so that names of individuals who have received exposure to sources of radiation are stated in a separate and detachable portion of the report.

(iv) Licensees or registrants shall make the reports required by § 175.03(l)(2)(i) and (ii) to the Department by telephone, telegram, mailgram, or facsimile.

(v) The provisions of § 175.03(l)(2) do not apply to doses that result from planned special exposures, provided such doses are within the limits for planned special exposures and are reported pursuant to § 175.03(l)(4).

(3) Reports of exposures, radiation levels, and concentrations of radioactive material exceeding the limits. (i) Reportable events. In addition to the notification required by § 175.03(l)(2), each licensee or registrant shall submit a written report within thirty (30) days after learning of any of the following occurrences:

(A) incidents for which notification is required by § 175.03(l)(2); or

(B) doses in excess of any of the following:

(a) the occupational dose limits for adults in § 175.03(c)(1); or

(b) the occupational dose limits for a minor in § 175.03(c)(7); or

(c) the limits for an embryo/fetus of a declared pregnant woman in §

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175.03(c)(8); or

(d) the limits for an individual member of the public in § 175.03(d)(1); or

(e) any applicable limit in the license or registration; or

(C) levels of radiation or concentrations of radioactive material in:

(a) a restricted area in excess of applicable limits in the license or registration; or

(b) an unrestricted area in excess of 10 times the applicable limit set forth in this Code or in the license or registration, whether or not involving exposure of any individual in excess of the limits in § 175.03(d)(1).

(ii) Contents of reports. (A) Each report required by § 175.03(l)(3)(i) shall describe the extent of exposure of individuals to radiation and radioactive material, including, as appropriate:

(a) estimates of each individual's dose; and

(b) the levels of radiation and concentrations of radioactive material involved; and

(c) the cause of the elevated exposures, dose rates, or concentrations; and

(d) corrective steps taken or planned to ensure against a recurrence, including the schedule for achieving conformance with applicable limits, generally applicable environmental standards, and associated license or registration conditions.

(B) Each report filed pursuant to § 175.03(l)(3)(i) shall include for each individual exposed: the name, social security account number, and date of birth. With respect to the limit for the embryo/fetus in § 175.03(c)(8), the identifiers should be those of the declared pregnant woman. The report shall be prepared so that this information is stated in a separate and detachable portion of the report.

(iii) All licensees or registrants who make reports pursuant to § 175.03(l)(3)(i) shall submit the report in writing to the Department.

(4) Reports of planned special exposures. The licensee or registrant shall submit a written report to the Department within 30 days following any planned special exposure conducted in accordance with § 175.03(c)(6), informing the Department that a planned special exposure was conducted and indicating the date the planned special exposure occurred and the information required by § 175.03(k)(7).

(5) Reports of individual monitoring. (i) This section applies to each person licensed or registered by the Department to:

(A) possess or use at any time, for processing or manufacturing for distribution

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pursuant to this Code, radioactive material in quantities exceeding any one of the following quantities:

_____

Radionuclide	Activitya	Ci	GBq _____
Cesium-137	1	37	
Cobalt-60	1	37	
Gold-198	100	3,700	
Iodine-131	1	37	
Iridium-192	10	370	
Krypton-85	1,000	37,000	
Promethium-147	10	370	
Technetium-99m	1,000	37,000 _____	

a The Department may require as a license condition, or by rule, regulation, or order pursuant to § 175.03(n)(1), reports from licensees or registrants who are licensed or registered to use radionuclides not on this list, in quantities sufficient to cause comparable radiation levels.

(ii) Each licensee or registrant in a category listed in § 175.03(l)(5)(i) shall submit an annual report of the results of individual monitoring carried out by the licensee or registrant for each individual for whom monitoring was required by § 175.03(f)(2) during that year. The licensee or registrant may include additional data for individuals for whom monitoring was provided but not required. The licensee or registrant shall use form RAD-5 or equivalent or electronic media containing all the information required by form RAD-5.

(iii) The licensee or registrant shall file the report required by § 175.03(l)(5)(ii), covering the preceding year, on or before April 30 of each year. The licensee or registrant shall submit the report to the Department.

(6) Notifications and reports to individuals. (i) Requirements for notification and reports to individuals of exposure to radiation or radioactive material are specified in § 175.04 of this Code.

(ii) When a licensee or registrant is required pursuant to § 175.03(l)(3) to report to the Department any exposure of an individual to radiation or radioactive material, the licensee or registrant shall also notify the individual. Such notice shall be transmitted at a time not later than the transmittal to the Department, and shall comply with the provisions of § 175.04 of this Code.

(7) Reports of leaking or contaminated sealed sources. The licensee or registrant shall file a report within five (5) days with the Department if the test for leakage or contamination required pursuant to § 175.03(e)(1) indicates a sealed source is leaking or contaminated. The report shall include the equipment involved, the test results and the corrective action taken.

(8) Event reporting. (i) Immediate report. Each licensee or registrant shall notify the Department as soon as possible, but not later than four (4) hours, after the discovery of an event that prevents immediate preventive actions necessary to avoid

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exposures to radiation or radioactive material that could exceed regulatory limits or releases of licensed material that could exceed regulatory limits (events may include fires, explosions, toxic gas releases, etc.).

(ii) Twenty-four hour report. Each licensee or registrant shall notify the Department within twenty-four (24) hours after the discovery of any of the following events involving regulated sources of radiation:

(A) An unplanned contamination event that:

(a) requires access to the contaminated area, by workers or the public, to be restricted for more than 24 hours by imposing additional radiological controls or by prohibiting entry into the area;

(b) involves a quantity of material greater than five (5) times the lowest annual limit on intake specified in Appendix B of § 175.03 of this Code for the material; and

(c) has access to the area restricted for a reason other than to allow isotopes with a half-life of less than 24 hours to decay prior to decontamination.

(B) An event in which equipment is disabled or fails to function as designed when:

(a) the equipment is required by regulation or license condition to prevent releases exceeding regulatory limits, or to mitigate the consequences of an accident;

(b) the equipment is required to be available and operable when it is disabled or fails to function; and

(c) no redundant equipment is available and operable to perform the required safety function.

(C) An event that requires unplanned medical treatment at a medical facility of an individual with spreadable radioactive contamination on the individual's clothing or body.

(D) An unplanned fire or explosion damaging any regulated radiation source or any device, container or equipment containing licensed material when:

(a) the quantity of material involved is greater than five (5) times the lowest annual limit on intake specified in Appendix B of § 175.03 for the material; and

(b) the damage affects the integrity of the licensed material or its container.

(iii) Preparation and submission of reports. Reports made by licensees in response to requirements of § 175.03(l)(9)(i) and (ii) must be made as follows:

(A) Licensees shall make reports required by § 175.03(l)(9)(i) and (ii) by telephone to the Department. To the extent that the information is available at the

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time of notification, the information provided in these reports must include:

(a) the caller's name and call back telephone number;

(b) a description of the event, including date and time;

(c) the exact location of the event;

(d) the isotopes, quantities, and chemical and physical form of the licensed material involved; and

(e) any personnel radiation exposure data available.

(B) Written report. Each licensee or registrant who makes a report required by § 175.03(l)(9)(i) or (ii) shall submit a written follow up report to the Department within thirty (30) days of the initial report. The reports must include the following:

(a) a description of the event, including the probable cause and the manufacturer and model number, if applicable, of any equipment that failed or malfunctioned;

(b) the isotopes, quantities and chemical and physical form of the licensed material involved;

(c) corrective actions taken or planned and the results of any evaluations or assessments; and

(d) the extent of exposure of individuals to radiation or to radioactive materials without identification of individuals by name.

(m) Exemptions and variances. The Department may grant, upon either its own initiative or an application by an interested person, an exemption or variance from any requirement in this Code when the Department finds that such exemption or variance will not result in an undue danger to life or property from radiation hazards.

(n) Additional requirements. (1) Notwithstanding any exemptions set forth in this Code:

(i) The Department may, by rule, regulation or order, impose upon any person who sells, transfers, assembles, repairs, receives, produces, possesses, or uses any radiation source, such requirements, in addition to those set forth in this Code, as it deems appropriate or necessary to protect the public health and safety and to minimize danger to life or property from radiation hazards.

(ii) The Department may suspend, revoke, or amend any license, certified registration or registration issued pursuant to this Code when it finds that any person is not in compliance with this Code, or other laws, ordinances, rules, or regulations of the Department or this City.

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(iii) The Department may order the owner, person in charge or the radiation safety officer of an installation, or any other person owning or responsible for a radiation source, to take such additional precautions or procedures as it may determine are necessary to prevent contamination or the over-exposure of persons to ionizing radiation or to otherwise protect the public health and safety. Except where the public health requires immediate action, no such order shall be issued until the person to be ordered is notified by any effective means of communication and is given an opportunity to be heard by such personnel of the Department as the Commissioner may designate.

(2) Vacating premises. Each specific licensee or registrant shall notify the Department in writing of intent to vacate not less than 30 days before vacating or relinquishing possession or control of premises which may have been contaminated with radioactive material as a result of licensed or registered activities. When deemed necessary by the Department, the licensee shall decontaminate the premises to such levels as the Department may specify.

(3) The Department may by order require the removal through an authorized person, or the surrender to the Department, of any radiation source by any person who:

(i) does not hold, or continue to hold, a valid license, certified registration or registration issued by the Department;

(ii) is not able or equipped, or who fails to observe with regard to such radiation source those radiation protection standards as are established by the Department or who uses such radiation source in violation of law, this Code, order of the Department, or as set forth in a license, certified registration or registration issued therefor by the Department. Such person shall decontaminate any premises which may have been contaminated with radioactive material as a result of licensed or registered activities to such radiation levels as the Department may specify. The expenses incidental to such transfer, surrender, and/or decontamination shall be borne by such person responsible for the source.

(4) When necessary or desirable in order to aid in determining the extent of any individual's exposure to radiation subsequent to any radiation accident, contamination, theft or loss, the licensee or registrant shall comply with all orders of the Department directing such licensee or registrant to make available to such individual appropriate medical evaluation services or appropriate tests and to furnish to the Department a copy of the reports of such evaluation or test.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (b) par (2) amended City Record July 15, 1997 eff. Aug. 14, 1997. [See Vol. 9 Statements of Basis and Purpose No. 8]

Subd. (c) par (1) subpar (i) amended City Record Sept. 29, 2006 eff. Oct. 29, 2006.

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[See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (c) par (1) subpar (iii) amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (f) par (2) subpar (vi) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (h) amended in part City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (j) par (6) subpar (iv) amended City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Subdivisions (c) and (h) and Appendix A of § 175.03 were amended on September 26, 2006, to maintain compatibility with NRC requirements as published in the Federal Register, dated October 7, 1999, titled, "Respiratory Protection and Controls to Restrict Internal Exposures", 64 FR 54543; 64 FR 55524, dated October 13, 1999, that became effective February 2, 2000 (amending 10 CFR §§ 20.1003; 20.1701-.1704; adding 20.1705 and amending Appendix A to Part 20).

Section 175.03(b)(2)(i)(B) was relettered to (C) and new (B) was adopted; Section 170.03(b)(2)(ii)(B) was relettered to (C) and new (B) was adopted; Section 175.03(b)(2)(iii)(C) was relettered to (D) and new (C) was adopted and Section 175.03(b)(2)(iv)(B) was relettered to (C) and new (B) was adopted on June 30, 1997 to authorize individuals with the appropriate training and experience to become radiation safety officers even though they are not board certified. The amendment recognizes an option existing in Section 175.03(j) of the Health Code, 10 N.Y.C.R.R. Section 16.2(a)(99) and federal regulations codified at 10 C.F.R. Section 35.900.

In addition, Section 175.03(b)(2)(iv) was amended to clarify its applicability to non-human use radioactive materials.

Subparagraph (vi) of paragraph (2) of subsection (f) was added on March 21, 2001 to address the need for timely processing of personnel radiation monitoring devices.

Subparagraphs (A) and (B) of paragraph (iv) of subdivision (6) of subsection (j) were amended on April 26, 1999 to reflect the reenacted § 175.105.

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NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
GENERAL PROVISIONS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

APPENDIX A

Protection Factors for Respirators 1

Modes 3	Protection Factors 4	Particulates only	Particulates, gases & vapors 5	Tested & Certified Equipment National Institute for Occupational Safety & Health/Mine Health & Safety Administration tests for permeability
Facepiece, half-mask 7 11, Subpart K.	NP	10		30 CFR
Facepiece, full	NP	50		
Facepiece, half-mask full, or hood	PP	1000		
Facepiece, half-mask 11, Subpart J.	CF		1000	30 CFR
Facepiece, half-mask	D		5	
Facepiece, full	CF		2000	
Facepiece, full	D		5	
Facepiece, full	PD		2000	
Hood	CF		8	
Suit	CF		9	10
Facepiece, full 11, Subpart H.	D		50	30 CFR

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Facepiece, full	PD	10,000 1 1	
Facepiece, full	RD	50	
Facepiece, full	RP	5,000 1 2	
Any combination of air-purifying and atmosphere-supplying respirators		Protection factor for type and mode of operation as listed above.	30 CFR 11, Sec. 11.63(b)

Footnotes:

1. For use in the selection of respiratory protective equipment to be used only where the contaminants have been identified and the concentrations, or possible concentrations, are known.

2. The licensee shall ensure that no objects, materials or substances, such as facial hair, or any conditions that interfere with the face, face-piece seal or valve function, and that are under the control of the respirator wearer, are present between the skin of the wearer's face and the sealing surface of a tightfitting respirator face-piece. Hoods and suits are excepted.

3. The mode symbols are defined as follows:

3. CF = continuous flow

3. D = demand

3. NP = negative pressure, that is, negative phase during inhalation

3. PD = pressure demand, that is, always positive pressure

3. PP = positive pressure

3. RD = demand, recirculating or closed circuit

3. RP = pressure demand, recirculating or closed circuit

4. a. The protection factor is a measure of the degree of protection afforded by a respirator, defined as the ratio of the concentration of airborne radioactive material outside the respiratory protective equipment to that inside the equipment, usually inside the facepiece, under conditions of use. It is applied to the ambient airborne concentration to estimate the concentrations inhaled by the wearer according to the following formula:

Concentration inhaled = Ambient airborne concentration _____
Protection factor

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b. The protection factors apply:

(i) Only for individuals trained in using respirators and wearing properly fitted respirators that are used and maintained under supervision in a well-planned respiratory protective program.

(ii) For air-purifying respirators only when high efficiency particulate filters, above 99.97% removal efficiency by thermally generated 0.3 <<mu>>m dioctylphthalate (DOP) test or equivalent, are used in atmospheres not deficient in oxygen and not containing radioactive gas or vapor respiratory hazards.

(iii) No adjustment is to be made for the use of sorbents against radioactive material in the form of gases or vapors.

(iv) For atmosphere-supplying respirators only when supplied with adequate respirable air. Respirable air shall be provided of the quality and quantity required in accordance with the National Institute for Occupational Safety and Health and the Mine Safety and Health Administration certification described in 30 CFR 11. Oxygen and air shall not be used in the same apparatus.

5. Excluding radioactive contaminants that present an absorption or submersion hazard. For tritium oxide, approximately one-third of the intake occurs by absorption through the skin so that an overall protection factor of less than 2 is appropriate when atmosphere-supplying respirators are used to protect against tritium oxide. If the protection factor for respiratory protective equipment is 5, the effective protection factor for tritium is about 1.4; with protection factors of 10, the effective factor for tritium oxide is about 1.7; and with protection factors of 100 or more, the effective factor for tritium oxide is about 1.9. Air-purifying respirators are not suitable for protection against tritium oxide. See also footnote 9 concerning supplied-air suits.

6. Canisters and cartridges shall not be used beyond service-life limitations.

7. Under-chin type only. This type of respirator is not satisfactory for use where it might be possible, such as, if an accident or emergency were to occur, for the ambient airborne concentrations to reach instantaneous values greater than 10 times the pertinent values in Table 1, Column 3 of Appendix B of § 175.03. This type of respirator is not suitable for protection against plutonium or other high-toxicity materials. The mask is to be tested for fit prior to use, each time it is donned.

8. a. Equipment shall be operated in a manner that ensures that proper air flowrates are maintained. A protection factor of no more than 1000 may be utilized for tested-and-certified supplied-air hoods when a minimum air flow of 6 cubic feet per minute (0.17 m3/min) is maintained and calibrated air line pressure gauges or flow measuring devices are used. A protection factor of up to 2000 may be used for tested and certified hoods only when the air flow is maintained at the manufacturer's recommended maximum rate for the equipment, this rate is greater than 6 cubic feet per minute (0.17 m3/min) and calibrated air line pressure gauges or flow measuring devices are used.

b. The design of the supplied-air hood or helmet, with a minimum flow of 6 cubic

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feet per minute (0.17 m3/min) of air, may determine its overall efficiency and the protection it provides. For example, some hoods aspirate contaminated air into the breathing zone when the wearer works with hands-over-head. This aspiration may be overcome if a short cape-like extension to the hood is worn under a coat or overalls. Other limitations specified by the approval agency shall be considered before using a hood in certain types of atmospheres. See footnote 9.

9. Appropriate protection factors shall be determined, taking into account the design of the suit and its permeability to the contaminant under conditions of use. Standby rescue persons are required whenever one piece atmosphere-supplying suits, or any combination of supplied air respiratory protection device and personnel protective equipment are used from which an unaided individual would have difficulty extricating himself or herself. The standby persons must be equipped with communications devices and respiratory protection devices or other apparatus appropriate for the potential hazards. The standby rescue persons shall observe or otherwise maintain continuous communication with the workers (visual, voice, signal line, telephone, radio, or other suitable means), and be immediately available to assist them in case of a failure of the air supply or for any other reason that requires relief from distress. A sufficient number of standby rescue persons must be immediately available to assist all users of this type of equipment and to provide effective emergency rescue if needed.

10. No approval schedules are currently available for this equipment. Equipment is to be evaluated by testing or on the basis of reliable test information.

11. This type of respirator may provide greater protection and be used as an emergency device in unknown concentrations for protection against inhalation hazards. External radiation hazards and other limitations to permitted exposure, such as skin absorption, must be taken into account in such circumstances.

12. Quantitative fit testing shall be performed on each individual, and no more than 0.02% leakage is allowed with this type of apparatus. Perceptible outward leakage of gas from this or any positive pressure self-contained breathing apparatus is unacceptable because service life will be reduced substantially. Special training in the use of this type of apparatus shall be provided to the wearer.

Note 1:

Protection factors for respirators approved by the U.S. Bureau of Mines and the National Institute for Occupational Safety and Health, according to applicable approvals for respirators for type and mode of use to protect against airborne radionuclides, may be used to the extent that they do not exceed the protection factors listed in this table. The protection factors listed in this table may not be appropriate to circumstances where chemical or other respiratory hazards exist in addition to radioactive hazards. The selection and use of respirators for such circumstances should take into account applicable approvals of the U.S. Bureau of Mines and the National Institute for Occupational Safety and Health.

Note 2:

Radioactive contaminants, for which the concentration values in Table 1, Column 3

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of Appendix B of § 175.03 are based on internal dose due to inhalation, may present external exposure hazards at higher concentrations. Under these circumstances, limitations on occupancy may have to be governed by external dose limits.

Definitions:

"Airpurifying respirator" means a respirator with an air purifying filter, cartridge, or canister that removes specific air contaminants by passing ambient air through the air purifying element.

"Assigned protection factor" or "APF" means the expected workplace level of respiratory protection that would be provided by a properly functioning respirator or a class of respirators to properly fitted and trained users. Operationally, the inhaled concentration can be estimated by dividing the ambient airborne concentration by the APF.

"Atmosphere-supplying respirator" means a respirator that supplies the respirator user with breathing air from a source independent of the ambient atmosphere, and includes supplied air respirators (SARs) and self-contained breathing apparatus (SCBA) units.

"Demand respirator" means an atmosphere-supplying respirator that admits breathing air to the face-piece only when a negative pressure is created inside the face-piece by inhalation.

"Disposable respirator" means a respirator for which maintenance is not intended and that is designed to be discarded after excessive breathing resistance, sorbent exhaustion, physical damage, or end of service-life renders it unsuitable for use. Examples of this type of respirator are a disposable halfmask respirator or a disposable escape-only self contained breathing apparatus (SCBA).

"Filtering face-piece" or "dust mask" means a negative pressure particulate respirator with a filter as an integral part of the face-piece or with the entire face-piece composed of the filtering medium, not equipped with elastomeric sealing surfaces and adjustable straps.

"Fit factor" means a quantitative estimate of the fit of a particular respirator to a specific individual, and typically estimates the ratio of the concentration of a substance in ambient air to its concentration inside the respirator when worn.

"Fit test" means the use of a protocol to qualitatively or quantitatively evaluate the fit of a respirator on an individual.

"Helmet" means a rigid respiratory inlet covering that also provides head protection against impact and penetration.

"Hood" means a respiratory inlet covering that completely covers the head and neck and may also cover portions of the shoulders and torso.

"Loosefitting face-piece" means a respiratory inlet covering that is designed to

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form a partial seal with the face.

"Negative pressure respirator" or "tight fitting respirator" means a respirator in which the air pressure inside the face-piece is negative during inhalation with respect to the ambient air pressure outside the respirator.

"Positive pressure respirator" means a respirator in which the pressure inside the respiratory inlet covering exceeds the ambient air pressure outside the respirator.

"Powered air purifying respirator" or "PAPR" means an air-purifying respirator that uses a blower to force the ambient air through air purifying elements to the inlet covering.

"Pressure demand respirator" means a positive pressure atmosphere-supplying respirator that admits breathing air to the face-piece when the positive pressure is reduced inside the face-piece by inhalation.

"Qualitative fit test" or "QLFT" means a pass/fail fit test to assess the adequacy of respirator fit that relies on the individual's response to the test agent.

"Quantitative fit test" "QNFT" means an assessment of the adequacy of respirator fit by numerically measuring the amount of leakage into the respirator.

"Self-contained breathing apparatus" or "SCBA" means an atmosphere-supplying respirator for which the breathing air source is designed to be carried by the user.

"Supplied air respirator" or "SAR" or "airline respirator" means an atmosphere-supplying respirator for which the source of breathing air is not designed to be carried by the user.

"Tight-fitting face-piece" means a respiratory inlet covering that forms a complete seal with the face.

"User seal check" or "fit check" means an action conducted by the respirator user to determine if the respirator is properly sealed to the face. Examples include negative pressure check, positive pressure check, irritant smoke check, or isoamylacetate check.

HISTORICAL NOTE

Footnote 2 amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Footnote 9 amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Definitions added City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

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NOTES:

Subdivisions (c) and (h) and Appendix A of § 175.03 were amended on September 26, 2006, to maintain compatibility with NRC requirements as published in the Federal Register, dated October 7, 1999, titled, "Respiratory Protection and Controls to Restrict Internal Exposures", 64 FR 54543; 64 FR 55524, dated October 13, 1999, that became effective February 2, 2000 (amending 10 CFR §§ 20.1003; 20.1701-.1704; adding 20.1705 and amending Appendix A to Part 20).

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APPENDIX B. ANNUAL LIMITS ON INTAKE (ALI) AND DERIVED AIR CONCENTRATIONS (DAC) OF RADIONUCLIDES FOR OCCUPATIONAL EXPOSURE; EFFLUENT CONCENTRATIONS; CONCENTRATIONS FOR RELEASE TO SANITARY SEWERAGE

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Introduction

For each radionuclide, Table I indicates the chemical form which is to be used for selecting the appropriate ALI or DAC value. The ALIs and DACs for inhalation are given for an aerosol with an activity median aerodynamic diameter (AMAD) of 1 <<mu>>m, micron, and for three classes (D,W,Y) of radioactive material, which refer to their retention (approximately days, weeks or years) in the pulmonary region of the lung. This classification applies to a range of clearance half-times for D if less than 10 days, for W from 10 to 100 days, and for Y greater than 100 days. Table 2 provides concentration limits for airborne and liquid effluents released to the general environment. Table 3 provides concentration limits for discharges to sanitary sewerage.

Note: The values in Tables 1, 2, and 3 are presented in the computer "E" notation. In this notation a value of 6E-02 represents a value of 6 x 10-2 or 0.06, 6E+2 represents 6 x 102 or 600, and 6E+0 represents 6 x 100 or 6.

Table 1 "Occupational Values"

Note that the columns in Table I of this appendix captioned "Oral Ingestion ALI," "Inhalation ALI," and "DAC," are applicable to occupational exposure to radioactive material.

The ALIs in this appendix are the annual intakes of given radionuclide by "Reference Man" which would result in either (1) a committed effective dose equivalent of 0.05 Sv (5 rem), stochastic ALI, or (2) a committed dose equivalent of 0.5 Sv (50 rem) to an organ or tissue, non-stochastic ALI. The stochastic ALIs were derived to result in a risk, due to irradiation of organs and tissues, comparable to the risk associated with deep dose equivalent to the whole body of 0.05 Sv (5 rem). The derivation includes multiplying the committed dose equivalent to an organ or tissue by a weighting factor, wT. This weighting factor is the proportion of the risk of stochastic effects resulting from irradiation of the organ or tissue, T, to the total risk of stochastic

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effects when the whole body is irradiated uniformly. The values of wT are listed under the definition of weighting factor in § 175.02(a)(247). The non-stochastic ALIs were derived to avoid non-stochastic effects, such as prompt damage to tissue or reduction in organ function.

A value of wT = 0.06 is applicable to each of the five organs or tissues in the "remainder" category receiving the highest dose equivalents, and the dose equivalents of all other remaining tissues may be disregarded. The following portions of the gastrointestinal tract--stomach, small intestine, upper large intestine, and lower large intestine--are to be treated as four separate organs.

Note that the dose equivalents for an extremity, skin and lens of the eye are not considered in computing the committed effective dose equivalent, but are subject to limits that must be met separately.

When an ALI is defined by the stochastic dose limit, this value alone is given. When an ALI is determined by the non-stochastic dose limit to an organ, the organ or tissue to which the limit applies is shown, and the ALI for the stochastic limit is shown in parentheses. Abbreviated organ or tissue designations are used:

The use of the ALIs listed first, the more limiting of the stochastic and non-stochastic ALIs, will ensure that non-stochastic effects are avoided and that the risk of stochastic effects is limited to an acceptably low value. If, in a particular situation involving a radionuclide for which the non-stochastic ALI is limiting, use of that non-stochastic ALI is considered unduly conservative, the licensee may use the stochastic ALI to determine the committed effective dose equivalent. However, the licensee shall also ensure that the 0.5 Sv (50 rem) dose equivalent limit for any organ or tissue is not exceeded by the sum of the external deep dose equivalent plus the internal committed dose equivalent to that organ, not the effective dose. For the case where there is no external dose contribution, this would be demonstrated if the sum of the fractions of the non-stochastic ALIs (ALIns) that contribute to the committed dose equivalent to the organ receiving the highest dose does not exceed unity, that is, <<SIGMA>> (intake (in <<mu>>Ci) of each radionuclide/ALIns) # 1.0. If there is an external deep dose equivalent contribution of Hd, then this sum must be less than 1--(Hd/50), instead of 1.0.

Note that the dose equivalents for an extremity, skin, and lens of the eye are not considered in computing the committed effective dose equivalent, but are subject to limits that must be met separately.

The derived air concentration (DAC) values are derived limits intended to control chronic occupational exposures. The relationship between the DAC and the ALI is given by:

DAC = ALI (in <<mu>>Ci)/(2000 hr-working yr-1 x 60 min-hr-1 x 2 x 104 ml-min-1)

=[ALI/2.4 x 109]<<mu>>Ci/ml, where 2 x 104 Ml is the volume of air breathed per

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minute at work by Reference Man under working conditions of light work.

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The DAC values relate to one of two modes of exposure: either external submersion or the internal committed dose equivalents resulting from inhalation of radioactive materials. DACs based upon submersion are for immersion in a semi-infinite cloud of uniform concentration and apply to each radionuclide separately.

The ALI and DAC values include contributions to exposure by the single radionuclide named and any in-growth of daughter radionuclides produced in the body by decay of the parent. However, intakes that include both the parent and daughter radionuclides should be treated by the general method appropriate for mixtures.

The values of ALI and DAC do not apply directly when the individual both ingests and inhales a radionuclide, when the individual is exposed to a mixture of radionuclides by either inhalation or ingestion or both, or when the individual is exposed to both internal and external irradiation. See § 175.03(c)(2). When an individual is exposed to radioactive materials which fall under several of the translocation classifications of the same radionuclide, such as, Class D, Class W, or Class Y, the exposure may be evaluated as if it were a mixture of different radionuclides.

It should be noted that the classification of a compound as Class D, W, or Y is based on the chemical form of the compound and does not take into account the radiological half-life of different radionuclides. For this reason, values are given for Class D, W, and Y compounds, even for very short-lived radionuclides.

Table 2 "Effluent Concentrations"

The columns in Table 2 of this appendix captioned "Effluents," "Air" and "Water" are applicable to the assessment and control of dose to the public, particularly in the implementation of the provisions of § 175.03(d)(2). The concentration values given in Columns 1 and 2 of Table 2 are equivalent to the radionuclide concentrations which, if inhaled or ingested continuously over the course of a year, would produce a total effective dose equivalent of 0.5 mSv (0.05 rem).

Consideration of non-stochastic limits has not been included in deriving the air and water effluent concentration limits because non-stochastic effects are presumed not to occur at or below the dose levels established for individual members of the public. For radionuclides, where the non-stochastic limit was governing in deriving the occupational DAC, the stochastic ALI was used in deriving the corresponding airborne effluent limit in Table 2. For this reason, the DAC and airborne effluent limits are not always proportional as they were in the previous version of the New York City Health Code in Table 4 of § 175.117.

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The air concentration values listed in Table 2, Column 1 were derived by one of two methods. For those radionuclides for which the stochastic limit is governing, the occupational stochastic inhalation ALI was divided by 2.4 x 10 9, relating the inhalation ALI to the DAC, as explained above, and then divided by a factor of 300. The factor of 300 includes the following components: a factor of 50 to relate the 0.05 Sv (5 rem) annual occupational dose limit to the 0.1 rem limit for members of the public, a factor of 3 to adjust for the difference in exposure time and the inhalation rate for a worker and that for members of the public; and a factor of 2 to adjust the occupational values, derived for adults, so that they are applicable to other age groups.

For those radionuclides for which submersion, that is external dose, is limiting, the occupational DAC in Table 1, Column 3 was divided by 219. The factor of 219 is composed of a factor of 50, as described above, and a factor of 4.38 relating occupational exposure for 2,000 hours per year to full-time exposure (8,760 hours per year). Note that an additional factor of 2 for age considerations is not warranted in the submersion case.

The water concentrations were derived by taking the most restrictive occupational stochastic oral ingestion ALI and dividing by 7.3 x 10 7 . The factor of 7.3 x 10 7 (ml) includes the following components: the factors of 50 and 2 described above and a factor of 7.3 x 10 5 (ml) which is the annual water intake of Reference Man.

Note 2 of this appendix provides groupings of radionuclides which are applicable to unknown mixtures of radionuclides. These groupings, including occupational inhalation ALIs and DACs, air and water effluent concentrations and releases to sewer, require demonstrating that the most limiting radionuclides in successive classes are absent. The limit for the unknown mixture is defined when the presence of one of the listed radionuclides cannot be definitely excluded as being present either from knowledge of the radionuclide composition of the source or from actual measurements.

Table 3 "Releases to Sewers"

The monthly average concentrations for release to sanitary sewerage are applicable to the provisions in § 175.104(c). The concentration values were derived by taking the most restrictive occupational stochastic oral ingestion ALI and dividing by 7.3 x 10 6 (ml). The factor of 7.3 x 10 6 (ml) is composed of a factor of 7.3 x 10 5 (ml), the annual water intake by Reference Man, and a factor of 10, such that the concentrations, if the sewage released by the licensee were the only source of water ingested by a Reference Man during a year, would result in a committed effective dose equivalent of 5 mSv (0.5 rem).

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale

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following the tabular data displaying the character positions.]

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******** This is piece 1. -- It begins at character 1 of table line 1. ********
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LIST OF ELEMENTS

Name	Atomic Symbol	Number	Name	Symbol

LIST OF ELEMENTS

Name	Symbol	Number	Name	Symbol
Actinium	Ac	89	Mercury	Hg
Aluminum	Al	13	Molybdenum	Mo
Americium	Am	95	Neodymium	Nd
Antimony	Sb	51	Neptunium	Np
Argon	Ar	18	Nickel	Ni
Arsenic	As	33	Niobium	Nb
Astatine	At	85	Osmium	Os
Barium	Ba	56	Palladium	Pd
Berkelium	Bk	97	Phosphorus	P
Beryllium	Be	4	Platinum	Pt
Bismuth	Bi	83	Plutonium	Pu
Bromine	Br	35	Polonium	Po
Cadmium	Cd	48	Potassium	K
Calcium	Ca	20	Praseodymium	Pr
Californium	Cf	98	Promethium	Pm
Carbon	C	6	Protactinium	Pa
Cerium	Ce	58	Radium	Ra
Cesium	Cs	55	Radon	Rn
Chlorine	Cl	17	Rhenium	Re
Chromium	Cr	24	Rhodium	Rh
Cobalt	Co	27	Rubidium	Rb
Copper	Cu	29	Ruthenium	Ru
Curium	Cm	96	Samarium	Sm
Dysprosium	Dy	66	Scandium	Sc
Einsteinium	Es	99	Selenium	Se
Erbium	Er	68	Silicon	Si
Europium	Eu	63	Silver	Ag
Fermium	Fm	100	Sodium	Na
Fluorine	F	9	Strontium	Sr
Francium	Fr	87	Sulfur	S
Gadolinium	Gd	64	Tantalum	Ta
Gallium	Ga	31	Technetium	Tc
Germanium	Ge	32	Tellurium	Te
Gold	Au	79	Terbium	Tb
Hafnium	Hf	72	Thallium	Tl
Holmium	Ho	67	Thorium	Th
Hydrogen	H	1	Thulium	Tm
Indium	In	49	Tin	Sn
Iodine	I	53	Titanium	Ti
Iridium	Ir	77	Tungsten	W

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Iron	Fe	26	Uranium	U
Krypton	Kr	36	Vanadium	V
Lanthanum	La	57	Xenon	Xe
Lead	Pb	82	Ytterbium	Yb
Lutetium	Lu	71	Yttrium	Y
Magnesium	Mg	12	Zinc	Zn
Manganese	Mn	25	Zirconium	Zr
Mendelevium	Md	101		

Table 1
Occupational
Values

Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion Inhalation	Col. 2 ALI (mCi)
1	Hydrogen-3	Water, DAC includes skin absorption	8E+4	8E+4
		Gas (HT or T2) Submersion 1: Use above values as HT and T2 oxidize in air and in the body to HTO		
4	Beryllium-7	W, all compounds except those given for Y	4E+4	2E+4
		Y, oxides, halides, and nitrates	--	2E+4
4	Beryllium--10	W, see 7Be	1E+3	2E+2
			LLI wall (1E+3)	--
		Y, see 7Be	--	1E+1
6	Carbon-11 2	Monoxide	--	1E+6
		Dioxide	--	6E+5
		Compounds	4E+5	4E+5
6	Carbon-14	Monoxide	--	2E+6
		Dioxide	--	2E+5
		Compounds	2E+3	2E+3
9	Fluorine-18 2	D, fluorides of H,	5E+4	7E+4
		Li, Na, K, Rb, Cs, and Fr	St wall (5E+4)	--
		W, fluorides of Be, Mg, Ca, Sr, Ba, Ra, Al,	--	9E+4

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		Ga, In, Tl, As, Sb, Bi, Fe, Ru, Os, Co, Ni, Pd, Pt, Cu, Ag, Au, Zn, Cd, Hg, Sc, Y, Ti, Zr, V, Nb, Ta, Mn, Tc, and Re		
		Y, lanthanum fluoride	--	8E+4
11	Sodium-22	D, all compounds	4E+2	6E+2
11	Sodium-24	D, all compounds	4E+3	5E+3
12	Magnesium--28	D, all compounds except those given for W	7E+2	2E+3
		W, oxides, hydroxides, carbides, halides, and nitrates	--	1E+3
13	Aluminum-26	D, all compounds except those given for W	4E+2	6E+1
		W, oxides, hydroxides, carbides, halides, and nitrates	--	9E+1
14	Silicon-31	D, all compounds except those given for W and Y	9E+3	3E+4
		W, oxides, hydroxides, carbides, and nitrates	--	3E+4
		Y, aluminosili- cate glass	--	3E+4
14	Silicon-32	D, see 3 1Si	2E+3 LLI wall (3E+3)	2E+2 --
		W, see 3 1Si	--	1E+2
		Y, see 3 1Si	--	5E+0
15	Phosphorus--32	D, all compounds except phosphates given for W	6E+2	9E+2
		W, phosphates of Zn 2+, S 3+, Mg 2+, Fe 3+, Bi 3+, and lanthanides	--	4E+2
15	Phosphorus--33	D, see 3 2P	6E+3	8E+3
		W, see 3 2P	--	3E+3
16	Sulfur-35	Vapor	--	1E+4
		D, sulfides and sulfates except those given for W	1E+4 LLI wall (8E+3)	2E+4 --
		W, elemental sulfur, sulfides of Sr, Ba, Ge, Sn, Pb, As, Sb, Bi, Cu, Ag, Au, Zn, Cd, Hg, W, and Mo. Sulfates of Ca, Sr, Ba, Ra, As, Sb, and Bi	6E+3 --	 2E+3
17	Chlorine-36	D, chlorides of H, Li, Na, K, Rb, Cs, and Fr	2E+3	2E+3
		W, chlorides of	--	2E+2

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		lanthanides, Be, Mg, Ca, Sr, Ba, Ra, Al, Ga, In, Tl, Ge, Sn, Pb, As, Sb, Bi, Fe, Ru, Os, Co, Rh, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zn, Cd, Hg, Sc, Y, Ti, Zr, Hf, V, Nb, Ta, Cr, Mo, W, Mn, Tc, and Re		
17	Chlorine-38 2	D, see 3 6Cl	2E+4 St wall (3E+4)	4E+4 --
		W, see 3 6Cl	--	5E+4
17	Chlorine-39 2	D, see 3 6Cl	2E+4 St wall (4E+4)	5E+4 --
		W, see 3 6Cl	--	6E+4
18	Argon-37	Submersion 1	--	--
18	Argon-39	Submersion 1	--	--
18	Argon-41	Submersion 1	--	--
19	Potassium-- 40	D, all compounds	3E+2	4E+2
19	Potassium-- 42	D, all compounds	5E+3	5E+3
19	Potassium-- 43	D, all compounds	6E+3	9E+3
19	Potassium-- 44 2	D, all compounds	2E+4 St wall (4E+4)	7E+4 --
19	Potassium-- 45 2	D, all compounds	3E+4 St wall (5E+4)	1E+5 --
20	Calcium-41	W, all compounds	3E+3 Bone surf (4E+3)	4E+3 Bone surf (4E+3)
20	Calcium-45	W, all compounds	2E+3	8E+2
20	Calcium-47	W, all compounds	8E+2	9E+2
21	Scandium-43	Y, all compounds	7E+3	2E+4
21	Scandium-4- 4m	Y, all compounds	5E+2	7E+2
21	Scandium-44	Y, all compounds	4E+3	1E+4
21	Scandium-46	Y, all compounds	9E+2	2E+2
21	Scandium-47	Y, all compounds	2E+3 LLI wall (3E+3)	3E+3 --
21	Scandium-48	Y, all compounds	8E+2	1E+3
21	Scandium-49 2	Y, all compounds	2E+4	5E+4
22	Titanium-44	D, all compounds except those given for W and Y	3E+2	1E+1
		W, oxides, hydroxides, carbides, halides, and nitrates	--	3E+1
		Y, SrTi0	--	6E+0
22	Titanium-45	D, see 4 4Ti	9E+3	3E+4
		W, see 4 4Ti	--	4E+4

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		Y, see 4 4Ti	--	3E+4
23	Vanadium-47 2	D, all compounds	3E+4	8E+4
		except those given for W	St wall (3E+4)	--
		W, oxides, hydroxides, carbides, and halides	--	1E+5
23	Vanadium-48	D, see 4 7V	6E+2	1E+3
		W, see 4 7V	--	6E+2
23	Vanadium-49	D, see 4 7V	7E+4 LLI wall (9E+4)	3E+4 Bone surf (3E+4)
		W, see 4 7V	--	2E+4
24	Chromium-48	D, all compounds except those given for W and Y	6E+3	1E+4
		W, halides and nitrates	--	7E+3
		Y, oxides and hydroxides	--	7E+3
24	Chromium-49 2	D, see 4 8Cr	3E+4	8E+4
		W, see 4 8Cr	--	1E+5
		Y, see 4 8Cr	--	9E+4
24	Chromium-51	D, see 4 8Cr	4E+4	5E+4
		W, see 4 8Cr	--	2E+4
		Y, see 4 8Cr	--	2E+4
25	Manganese--51 2	D, all compounds except those given for W	2E+4	5E+4
		W, oxides, hydroxides, halides, and nitrates	--	6E+4
25	Manganese--52m 2	D, see 5 1Mn	3E+4 St wall (4E+4)	9E+4 --
		W, see 5 1Mn	--	1E+5
25	Manganese--52	D, see 5 1Mn	7E+2	1E+3
		W, see 5 1Mn	--	9E+2
25	Manganese--53	D, see 5 1Mn	5E+4 --	1E+4 Bone surf (2E+4)
		W, see 5 1Mn	--	1E+4
25	Manganese--54	D, see 5 1Mn	2E+3	9E+2
		W, see 5 1Mn	--	8E+2
25	Manganese--56	D, see 5 1Mn	5E+3	2E+4
		W, see 5 1Mn	--	2E+4
26	Iron-52	D, all compounds except those given for W	9E+2	3E+3
		W, oxides, hydroxides, and halides	--	2E+3
26	Iron-55	D, see 5 2Fe	9E+3	2E+3
		W, see 5 2Fe	--	4E+3
26	Iron-59	D, see 5 2Fe	8E+2	3E+2
		W, see 5 2Fe	--	5E+2

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26             Iron-60      D, see 5 2Fe              3E+1            6E+0
                            W, see 5 2Fe              --              2E+1
27             Cobalt-55    W, all compounds except   1E+3            3E+3
                              those given for Y
                            Y, oxides, hydroxides,    --              3E+3
                              halides, and nitrates
27             Cobalt-56    W, see 5 5Co              5E+2            3E+2
                            Y, see 5 5Co              4E+2            2E+2
27             Cobalt-57    W, see 5 5Co              8E+3            3E+3
                            Y, see 5 5Co              4E+3            7E+2
27             Cobalt-58m   W, see 5 5Co              6E+4            9E+4
                            Y, see 5 5Co              --              6E+4
27             Cobalt-58    W, see 5 5Co              2E+3            1E+3
                            Y, see 5 5Co              1E+3            7E+2
27             Cobalt-60m   W, see 5 5Co              1E+6 St wall    4E+6 --
                 2                                      (1E+6)
                            Y, see 5 5Co              --              3E+6
27             Cobalt-60    W, see 5 5Co              5E+2            2E+2
                            Y, see 5 5Co              2E+2            3E+1
27             Cobalt-61 2  W, see 5 5Co              2E+4            6E+4
                            Y, see 5 5Co              2E+4            6E+4
27             Cobalt-62m   W, see 5 5Co              4E+4 St wall    2E+5 --
                 2                                      (5E+4)
                            Y, see 5 5Co              --              2E+5
28             Nickel-56    D, all compounds except   1E+3            2E+3
                              those given for W
                            W, oxides, hydroxides,    --              1E+3
                              and carbides
                            Vapor                     --              1E+3
28             Nickel-57    D, see 5 6Ni              2E+3            5E+3
                            W, see 5 6Ni              --              3E+3
                            Vapor                     --              6E+3
28             Nickel-59    D, see 5 6Ni              2E+4            4E+3
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

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******* This is piece 2. -- It begins at character 80 of table line 1. ********
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  Atomic

  Number
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  80
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65
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Table 2 Effluent Concentrations

Col. 3 ALI (mCi)	Col. 1 DAC (mCi/ml)	Col. 2 Air (mCi/ml)	Water (mCi/ml)	Monthly Average Concentration (mCi/ml)

Table 2 Effluent Concentrations

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Col. 3	Col. 1	Col. 2		
ALI (mCi)	DAC (mCi/ml)	Air (mCi/ml)	Water (mCi/ml)	Monthly Average Concentration (mCi/ml)
2E-5	1E-7	1E-3	1E-2	
9E-6	3E-8	6E-4	6E-3	
8E-6	3E-8	--	--	
6E-8	2E-10	--	--	
--	--	2E-5	2E-4	
6E-9	2E-11	--	--	
5E-4	2E-6	--	--	
3E-4	9E-7	--	--	
2E-4	6E-7	6E-3	6E-2	
7E-4	2E-6	--	--	
9E-5	3E-7	--	--	
1E-6	3E-9	3E-5	3E-4	
3E-5	1E-7	--	--	
--	--	7E-4	7E-3	
4E-5	1E-7	--	--	
3E-5	1E-7	--	--	
3E-7	9E-10	6E-6	6E-5	
2E-6	7E-9	5E-5	5E-4	
7E-7	2E-9	9E-6	9E-5	
5E-7	2E-9	--	--	
3E-8	9E-11	6E-6	6E-5	
4E-8	1E-10	--	--	
1E-5	4E-8	1E-4	1E-3	

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1E-5	5E-8	--	--
1E-5	4E-8	--	--
1E-7	3E-10	--	--
--	--	4E-5	4E-4
5E-8	2E-10	--	--
2E-9	7E-12	--	--
4E-7	1E-9	9E-6	9E-5
2E-7	5E-10	--	--
4E-6	1E-8	8E-5	8E-4
1E-6	4E-9	--	--
6E-6	2E-8	--	--
7E-6	2E-8	--	--
--	--	1E-4	1E-3
9E-7	3E-9	--	--
1E-6	3E-9	2E-5	2E-4
1E-7	3E-10	--	--
2E-5 --	6E-8 --	-- 3E-4	-- 3E-3
2E-5	6E-8	--	--
2E-5 --	7E-8 --	-- 5E-4	-- 5E-3
2E-5	8E-8	--	--
1E+0	6E-3	--	--
2E-4	8E-7	--	--
3E-6	1E-8	--	--
2E-7	6E-10	4E-6	4E-5

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2E-6	7E-9	6E-5	6E-4
4E-6	1E-8	9E-5	9E-4
3E-5 --	9E-8 --	-- 5E-4	-- 5E-3
5E-5 --	2E-7 --	-- 7E-4	-- 7E-3
2E-6 --	-- 5E-9	-- 6E-5	-- 6E-4
4E-7	1E-9	2E-5	2E-4
4E-7	1E-9	1E-5	1E-4
9E-6	3E-8	1E-4	1E-3
3E-7	1E-9	7E-6	7E-5
5E-6	2E-8	5E-5	5E-4
1E-7	3E-10	1E-5	1E-4
1E-6 --	4E-9 --	-- 4E-5	-- 4E-4
6E-7	2E-9	1E-5	1E-4
2E-5	8E-8	3E-4	3E-3
5E-9	2E-11	4E-6	4E-5
1E-8	4E-11	--	--
2E-9	8E-12	--	--
1E-5	3E-8	1E-4	1E-3
1E-5	5E-8	--	--
1E-5	4E-8	--	--
3E-5	1E-7	--	--
--	--	4E-4	4E-3
4E-5	1E-7	--	--
5E-7	2E-9	9E-6	9E-5
3E-7	9E-10	--	--
1E-5 --	-- 5E-8	-- 1E-3	-- 1E-2
8E-6	2E-8	--	--
5E-6	2E-8	8E-5	8E-4
3E-6	1E-8	--	--
3E-6	1E-8	--	--
4E-5	1E-7	4E-4	4E-3

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4E-5	1E-7	--	--
4E-5	1E-7	--	--
2E-5	6E-8	5E-4	5E-3
1E-5	3E-8	--	--
8E-6	3E-8	--	--
2E-5	7E-8	3E-4	3E-3
3E-5	8E-8	--	--
4E-5 --	1E-7 --	-- 5E-4	-- 5E-3
4E-5	1E-7	--	--
5E-7	2E-9	1E-5	1E-4
4E-7	1E-9	--	--
5E-6 --	-- 3E-8	7E-4 --	7E-3 --
5E-6	2E-8	--	--
4E-7	1E-9	3E-5	3E-4
3E-7	1E-9	--	--
6E-6	2E-8	7E-5	7E-4
9E-6	3E-8	--	--
1E-6	4E-9	1E-5	1E-4
1E-6	3E-9	--	--
8E-7	3E-9	1E-4	1E-3
2E-6	6E-9	--	--
1E-7	5E-10	1E-5	1E-4
2E-7	7E-10	--	--
3E-9	9E-12	4E-7	4E-6
8E-9	3E-11	--	--
1E-6	4E-9	2E-5	2E-4
1E-6	4E-9	--	--
1E-7	4E-10	6E-6	6E-5
8E-8	3E-10	--	--
1E-6	4E-9	6E-5	6E-4
3E-7	9E-10	--	--
4E-5	1E-7	8E-4	8E-3
3E-5	9E-8	--	--
5E-7	2E-9	2E-5	2E-4
3E-7	1E-9	--	--
2E-3 --	6E-6 --	-- 2E-2	-- 2E-1
1E-3	4E-6	--	--
7E-8	2E-10	3E-6	3E-5
1E-8	5E-11	--	--
3E-5	9E-8	3E-4	3E-3

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```
  2E-5       8E-8     --                  --
  7E-5 --    2E-7 --  -- 7E-4             -- 7E-3

  6E-5       2E-7     --                  --
  8E-7       3E-9     2E-5                2E-4

  5E-7       2E-9     --                  --

  5E-7       2E-9     --                  --
  2E-6       7E-9     2E-5                2E-4
  1E-6       4E-9     --                  --
  3E-6       9E-9     --                  --
  2E-6       5E-9     3E-4                3E-3
80..+...90....+....0....+...10....+...20....+...30....+...40....+...50

*******************************************************************************
******* This is piece 3. -- It begins at character 150 of table line 1. *******
*******************************************************************************
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Table 3
Releases
to
Sewers

Table 3
Releases
to
Sewers

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150.....60.

TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

W, see 5 6Ni	--	7E+3	3E-6	1E-8	--	--
Vapor	--	2E+3	8E-7	3E-9	--	--

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28	Nickel-63	D, see 5 6Ni	9E+3	2E+3	7E-7	2E-9	1E-4	1E-3
		W, see 5 6Ni	--	3E+3	1E-6	4E-9	--	--
		Vapor	--	8E+2	3E-7	1E-9	--	--
28	Nickel-65	D, see 5 6Ni	8E+3	2E+4	1E-5	3E-8	1E-4	1E-3
		W, see 5 6Ni	--	3E+4	1E-5	4E-8	--	--
		Vapor	--	2E+4	7E-6	2E-8	--	--
28	Nickel-66	D, see 5 6Ni	4E+2 LLI wall (5E+-2)	2E+3 --	7E-7 --	2E-9 --	-- 6-E-6	-- 6-E--5
		W, see 5 6Ni	--	6E+2	3E-7	9E-10	--	--
		Vapor	--	3E+3	1E-6	4E-9	--	--
29	Copper-60 2	D, all compounds except those given for W and Y	3E+4 St wall (3E+4)	9E+4 --	4E-5 --	1E-7 --	-- 4E-4	-- 4E-3
		W, sulfides, halides, and nitrates	--	1E+5	5E-5	2E-7	--	--
		Y, oxides and hydroxides	--	1E+5	4E-5	1E-7	--	--
29	Copper-61	D, see 6 0Cu	1E+4	3E+4	1E-5	4E-8	2E-4	2E-3
		W, see 6 0Cu	--	4E+4	2E-5	6E-8	--	--
		Y, see 6 0Cu	--	4E+4	1E-5	5E-8	--	--
29	Copper-64	D, see 6 0Cu	1E+4	3E+4	1E-5	4E-8	2E-4	2E-3
		W, see 6 0Cu	--	2E+4	1E-5	3E-8	--	--
		Y, see 6 0Cu	--	2E+4	9E-6	3E-8	--	--
29	Copper-67	D, see 6 0Cu	5E+3	8E+3	3E-6	1E-8	6E-5	6E-4
		W, see 6 0Cu	--	5E+3	2E-6	7E-9	--	--
		Y, see 6 0Cu	--	5E+3	2E-6	6E-9	--	--
30	Zinc-62	Y, all compounds	1E+3	3E+3	1E-6	4E-9	2E-5	2E-4
30	Zinc-63 2	Y, all compounds	2E+4 St wall (3E+-4)	7E+4 --	3E-5 --	9E-8 --	-- 3-E-4	-- 3-E--3
30	Zinc-65	Y, all compounds	4E+2	3E+2	1E-7	4E-10	5E-6	5E-5
30	Zinc-69m	Y, all compounds	4E+3	7E+3	3E-6	1E-8	6E-5	6E-4
30	Zinc-69 2	Y, all compounds	6E+4	1E+5	6E-5	2E-7	8E-4	8E-3
30	Zinc-71m	Y, all compounds	6E+3	2E+4	7E-6	2E-8	8E-5	8E-4
30	Zinc-72	Y, all compounds	1E+3	1E+3	5E-7	2E-9	1E-5	1E-4
31	Gallium-65 2	D, all compounds except those given for W	5E+4 St wall (6E+4)	2E+5 --	7E-5 --	2E-7 --	-- 9E-4	-- 9E-3
		W, oxides, hydroxides, carbides, halides, and nitrates	--	2E+5	8E-5	3E-7	--	--
31	Gallium-66	D, see 6 5Ga	1E+3	4E+3	1E-6	5E-9	1E-5	1E-4
		W, see 6 5Ga	--	3E+3	1E-6	4E-9	--	--
31	Gallium-67	D, see 6 5Ga	7E+3	1E+4	6E-6	2E-8	1E-4	1E-3
		W, see 6 5Ga	--	1E+4	4E-6	1E-8	--	--

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31	Gallium-68	2	D, see 6 5Ga	2E+4	4E+4	2E-5	6E-8	2E-4	2E-3
			W, see 6 5Ga	--	5E+4	2E-5	7E-8	--	--
31	Gallium-70	2	D, see 6 5Ga	5E+4 St wall (7E+-4)	2E+5 --	7E-5 --	2E-7 --	-- 1-E-3	-- 1-E-2
			W, see 6 5Ga	--	2E+5	8E-5	3E-7	--	--
31	Gallium-72		D, see 6 5Ga	1E+3	4E+3	1E-6	5E-9	2E-5	2E-4
			W, see 6 5Ga	--	3E+3	1E-6	4E-9	--	--
31	Gallium-73		D, see 6 5Ga	5E+3	2E+4	6E-6	2E-8	7E-5	7E-4
			W, see 6 5Ga	--	2E+4	6E-6	2E-8	--	--
32	Germanium-66		D, all compounds except those given for W	2E+4	3E+4	1E-5	4E-8	3E-4	3E-3
			W, oxides, sulfides, and halides	--	2E+4	8E-6	3E-8	--	--
32	Germanium-67	2	D, see 6 6Ge	3E+4 St wall (4E+-4)	9E+4 --	4E-5 --	1E-7 --	-- 6-E-4	-- 6-E-3
			W, see 6 6Ge	--	1E+5	4E-5	1E-7	--	--
32	Germanium-68		D, see 6 6Ge	5E+3	4E+3	2E-6	5E-9	6E-5	6E-4
			W, see 6 6Ge	--	1E+2	4E-8	1E-10	--	--
32	Germanium-69		D, see 6 6Ge	1E+4	2E+4	6E-6	2E-8	2E-4	2E-3
			W, see 6 6Ge	--	8E+3	3E-6	1E-8	--	--
32	Germanium-71		D, see 6 6Ge	5E+5	4E+5	2E-4	6E-7	7E-3	7E-2
			W, see 6 6Ge	--	4E+4	2E-5	6E-8	--	--
32	Germanium-75	2	D, see 6 6Ge	4E+4 St wall (7E+-4)	8E+4 --	3E-5 --	1E-7 --	-- 9-E-4	-- 9-E-3
			W, see 6 6Ge	--	8E+4	4E-5	1E-7	--	--
32	Germanium-77		D, see 6 6Ge	9E+3	1E+4	4E-6	1E-8	1E-4	1E-3
			W, see 6 6Ge	--	6E+3	2E-6	8E-9	--	--
32	Germanium-78	2	D, see 6 6Ge	2E+4 St wall (2E+-4)	2E+4 --	9E-6 --	3E-8 --	-- 3-E-4	-- 3-E-3
			W, see 6 6Ge	--	2E+4	9E-6	3E-8	--	--
33	Arsenic-69	2	W, all compounds	3E+4 St wall (4E+-4)	1E+5 --	5E-5 --	2E-7 --	-- 6-E-4	-- 6-E-3
33	Arsenic-70	2	W, all compounds	1E+4	5E+4	2E-5	7E-8	2E-4	2E-3
33	Arsenic-71		W, all compounds	4E+3	5E+3	2E-6	6E-9	5E-5	5E-4
33	Arsenic-72		W, all compounds	9E+2	1E+3	6E-7	2E-9	1E-5	1E-4
33	Arsenic-73		W, all compounds	8E+3	2E+3	7E-7	2E-9	1E-4	1E-3
33	Arsenic-74		W, all compounds	1E+3	8E+2	3E-7	1E-9	2E-5	2E-4
33	Arsenic-76		W, all compounds	1E+3	1E+3	6E-7	2E-9	1E-5	1E-4
33	Arsenic-77		W, all compounds	4E+3 LLI wall	5E+3 --	2E-6 --	7E-9 --	-- 6-E-5	-- 6-E-

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			(5E+-3)					-4
33	Arsenic-78 2	W, all compounds	8E+3	2E+4	9E-6	3E-8	1E-4	1E-3
34	Selenium-70 2	D, all compounds except those given for W	2E+4	4E+4	2E-5	5E-8	1E-4	1E-3
		W, oxides, hydroxides, carbides, and elemental Se	1E+4	4E+4	2E-5	6E-8	--	--
34	Selenium-73m 2	D, see 7 0Se	6E+4	2E+5	6E-5	2E-7	4E-4	4E-3
		W, see 7 0Se	3E+4	1E+5	6E-5	2E-7	--	--
34	Selenium-73	D, see 7 0Se	3E+3	1E+4	5E-6	2E-8	4E-5	4E-4
		W, see 7 0Se	--	2E+4	7E-6	2E-8	--	--
34	Selenium-75	D, see 7 0Se	5E+2	7E+2	3E-7	1E-9	7E-6	7E-5
		W, see 7 0Se	--	6E+2	3E-7	8E-10	--	--
34	Selenium-79	D, see 7 0Se	6E+2	8E+2	3E-7	1E-9	8E-6	8E-5
		W, see 7 0Se	--	6E+2	2E-7	8E-10	--	--
34	Selenium-81m 2	D, see 7 0Se	4E+4	7E+4	3E-5	9E-8	3E-4	3E-3
		W, see 7 0Se	2E+4	7E+4	3E-5	1E-7	--	--
34	Selenium-81 2	D, see 7 0Se	6E+4 St wall (8E+-4)	2E+5 --	9E-5 --	3E-7 --	-- 1-E-3	-- 1-E-2
		W, see 7 0Se	--	2E+5	1E-4	3E-7	--	--
34	Selenium-83 2	D, see 7 0Se	4E+4	1E+5	5E-5	2E-7	4E-4	4E-3
		W, see 7 0Se	3E+4	1E+5	5E-5	2E-7	--	--
35	Bromine-74m 2	D, bromides of						
		H, Li, Na, K, Rb, Cs, and Fr	1E+4 St wall (2E+4)	4E+4 --	2E-5 --	5E-8 --	-- 3E-4	-- 3E-3
		W, bromides of lanthanides, Be, Mg, Ca, Sr, Ba, Ra, Al, Ga, In, Tl, Ge, Sn, Pb, As, Sb, Bi, Fe, Ru, Os, Co, Rh, Ir, Ni, Pd, Pt, Cu, Ag, Au, Zn, Cd, Hg, Sc, Y, Ti, Zr, Hf, V, Nb, Ta, Mn, Tc, and Re	--	4E+4	2E-5	6E-8	--	--
35	Bromine-74 2	D, see 7 4mBr	2E+4 St wall (4E+-4)	7E+4 --	3E-5 --	1E-7 --	-- 5-E-4	-- 5-E-3
		W, see 7 4mBr	--	8E+4	4E-5	1E-7	--	--

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35	Bromine-75 2	D, see 7 4mBr	3E+4 St wall (4E+-4)	5E+4 --	2E-5 --	7E-8 --	-- 5- E-4	-- 5- E-3
		W, see 7 4mBr	--	5E+4	2E-5	7E-8	--	--
35	Bromine-76	D, see 7 4mBr	4E+3	5E+3	2E-6	7E-9	5E-5	5E-4
		W, see 7 4mBr	--	4E+3	2E-6	6E-9	--	--
35	Bromine-77	D, see 7 4mBr	2E+4	2E+4	1E-5	3E-8	2E-4	2E-3
		W, see 7 4mBr	--	2E+4	8E-6	3E-8	--	--
35	Bromine-80m	D, see 7 4mBr	2E+4	2E+4	7E-6	2E-8	3E-4	3E-3
		W, see 7 4mBr	--	1E+4	6E-6	2E-8	--	--
35	Bromine-80 2	D, see 7 4mBr	5E+4 St wall (9E+-4)	2E+5 --	8E-5 --	3E-7 --	-- 1- E-3	-- 1- E-2
		W, see 7 4mBr	--	2E+5	9E-5	3E-7	--	--
35	Bromine-82	D, see 7 4mBr	3E+3	4E+3	2E-6	6E-9	4E-5	4E-4
		W, see 7 4mBr	--	4E+3	2E-6	5E-9	--	--
35	Bromine-83	D, see 7 4mBr	5E+4 St wall (7E+-4)	6E+4 --	3E-5 --	9E-8 --	-- 9- E-4	-- 9- E-3
		W, see 7 4mBr	--	6E+4	3E-5	9E-8	--	--
35	Bromine-84 2	D, see 7 4mBr	2E+4 St wall (3E+-4)	6E+4 --	2E-5 --	8E-8 --	-- 4- E-4	-- 4- E-3
		W, see 7 4mBr	--	6E+4	3E-5	9E-8	--	--
36	Krypton-74 2	Submersion 1	--	--	3E-6	1E-8	--	--
36	Krypton-76	Submersion 1	--	--	9E-6	4E-8	--	--
36	Krypton-77 2	Submersion 1	--	--	4E-6	2E-8	--	--
36	Krypton-79	Submersion 1	--	--	2E-5	7E-8	--	--
36	Krypton-81	Submersion 1	--	--	7E-4	3E-6	--	--
36	Krypton-83m 2	Submersion 1	--	--	1E-2	5E-5	--	--
36	Krypton-85m	Submersion 1	--	--	2E-5	1E-7	--	--
36	Krypton-85	Submersion 1	--	--	1E-4	7E-7	--	--
36	Krypton-87 2	Submersion 1	--	--	5E-6	2E-8	--	--
36	Krypton-88	Submersion 1	--	--	2E-6	9E-9	--	--
37	Rubidium-79 2	D, all compounds	4E+4 St wall (6E+-4)	1E+5 --	5E-5 --	2E-7 --	-- 8- E-4	-- 8- E-3
37	Rubidium-81m 2	D, all compounds	2E+5 St wall (3E+-5)	3E+5 --	1E-4 --	5E-7 --	-- 4- E-3	-- 4- E-2
37	Rubidium-81	D, all compounds	4E+4	5E+4	2E-5	7E-8	5E-4	5E-3
37	Rubidium-82m	D, all compounds	1E+4	2E+4	7E-6	2E-8	2E-4	2E-3
37	Rubidium-83	D, all compounds	6E+2	1E+3	4E-7	1E-9	9E-6	9E-5
37	Rubidium-84	D, all compounds	5E+2	8E+2	3E-7	1E-9	7E-6	7E-5
37	Rubidium-86	D, all compounds	5E+2	8E+2	3E-7	1E-9	7E-6	7E-5
37	Rubidium-87	D, all compounds	1E+3	2E+3	6E-7	2E-9	1E-5	1E-4

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37	Rubidium-88	D, all compounds	2E+4	St wall (3E+-4)	6E+4	--	3E-5	--	9E-8	-- -- 4- -- E-4 4-E--3
37	Rubidium-89	D, all compounds	4E+4	St wall (6E+-4)	1E+5	--	6E-5	--	2E-7	-- -- 9- -- E-4 9-E--3
38	Strontium-80	D, all soluble compounds except SrTiO	4E+3		1E+4		5E-6		2E-8	6E-5 6E-4
		Y, all insoluble compounds and SrTiO	--		1E+4		5E-6		2E-8	-- --
38	Strontium-81	D, see 8 0Sr	3E+4		8E+4		3E-5		1E-7	3E-4 3E-3
		Y, see 8 0Sr	2E+4		8E+4		3E-5		1E-7	-- --
38	Strontium-82	D, see 8 0Sr	3E+2	LLI wall (2E+-2)	4E+2	--	2E-7	--	6E-10 --	-- 3- -- E-6 3-E--5
		Y, see 8 0Sr	2E+2		9E+1		4E-8		1E-10	-- --
38	Strontium-83	D, see 8 0Sr	3E+3		7E+3		3E-6		1E-8	3E-5 3E-4
		Y, see 8 0Sr	2E+3		4E+3		1E-6		5E-9	-- --
38	Strontium-8-5m	D, see 8 0Sr	2E+5		6E+5		3E-4		9E-7	3E-3 3E-2
		Y, see 8 0Sr	--		8E+5		4E-4		1E-6	-- --
38	Strontium-85	D, see 8 0Sr	3E+3		3E+3		1E-6		4E-9	4E-5 4E-4
		Y, see 8 0Sr	--		2E+3		6E-7		2E-9	-- --
38	Strontium-8-7m	D, see 8 0Sr	5E+4		1E+5		5E-5		2E-7	6E-4 6E-3
		Y, see 8 0Sr	4E+4		2E+5		6E-5		2E-7	-- --
38	Strontium-89	D, see 8 0Sr	6E+2	LLI wall (6E+-2)	8E+2	--	4E-7	--	1E-9 --	-- 8- -- E-6 8-E--5
		Y, see 8 0Sr	5E+2		1E+2		6E-8		2E-10	-- --
38	Strontium-90	D, see 8 0Sr	3E+1	Bone surf (4E+-1)	2E+1	Bone surf (2E+-1)	8E-9	--	-- 3E-11	-- 5- -- E-7 5-E--6
		Y, see 8 0Sr	--		4E+0		2E-9		6E-12	-- --
38	Strontium-91	D, see 8 0Sr	2E+3		6E+3		2E-6		8E-9	2E-5 2E-4
		Y, see 8 0Sr	--		4E+3		1E-6		5E-9	-- --
38	Strontium-92	D, see 8 0Sr	3E+3		9E+3		4E-6		1E-8	4E-5 4E-4
		Y, see 8 0Sr	--		7E+3		3E-6		9E-9	-- --
39	Yttrium-86m	W, all compounds except those given for Y	2E+4		6E+4		2E-5		8E-8	3E-4 3E-3
		Y, oxides and	--		5E+4		2E-5		8E-8	-- --

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		hydroxides							
39	Yttrium-86	W, see 8 6mY	1E+3	3E+3	1E-6	5E-9	2E-5	2E-4	
		Y, see 8 6mY	--	3E+3	1E-6	5E-9	--	--	
39	Yttrium-87	W, see 8 6mY	2E+3	3E+3	1E-6	5E-9	3E-5	3E-4	
		Y, see 8 6mY	--	3E+3	1E-6	5E-9	--	--	
39	Yttrium-88	W, see 8 6mY	1E+3	3E+2	1E-7	3E-10	1E-5	1E-4	
		Y, see 8 6mY	--	2E+2	1E-7	3E-10	--	--	
39	Yttrium-90m	W, see 8 6mY	8E+3	1E+4	5E-6	2E-8	1E-4	1E-3	
		Y, see 8 6mY	--	1E+4	5E-6	2E-8	--	--	
39	Yttrium-90	W, see 8 6mY	4E+2	7E+2	3E-7	9E-10	7-	--	
			LLI wall (5E+-2)			--	E-6	7-E-5	
		Y, see 8 6mY	--	6E+2	3E-7	9E-10	--	--	
39	Yttrium-91m 2	W, see 8 6mY	1E+5	2E+5	1E-4	3E-7	2E-3	2E-2	
		Y, see 8 6mY	--	2E+5	7E-5	2E-7	--	--	
39	Yttrium-91	W, see 8 6mY	5E+2	2E+2	7E-8	2E-10	8-	--	
			LLI wall (6E+-2)			--	E-6	8-E-5	
		Y, see 8 6mY	--	1E+2	5E-8	2E-10	--	--	
39	Yttrium-92	W, see 8 6mY	3E+3	9E+3	4E-6	1E-8	4E-5	4E-4	
		Y, see 8 6mY	--	8E+3	3E-6	1E-8	--	--	
39	Yttrium-93	W, see 8 6mY	1E+3	3E+3	1E-6	4E-9	2E-5	2E-4	
		Y, see 8 6mY	--	2E+3	1E-6	3E-9	--	--	
39	Yttrium-94 2	W, see 8 6mY	2E+4 St wall (3E+-4)	8E+4	3E-5	1E-7	4-	--	
							E-4	4-E-3	
		Y, see 8 6mY	--	8E+4	3E-5	1E-7	--	--	
39	Yttrium-95 2	W, see 8 6mY	4E+4 St wall (5E+-4)	2E+5	6E-5	2E-7	7-	--	
							E-4	7-E-3	
		Y, see 8 6mY	--	1E+5	6E-5	2E-7	--	--	
40	Zirconium-86	D, all compounds except those given for W and Y	1E+3	4E+3	2E-6	6E-9	2E-5	2E-4	
		W, oxides, hydroxides, halides, and nitrates	--	3E+3	1E-6	4E-9	--	--	
		Y, carbide	--	2E+3	1E-6	3E-9	--	--	
40	Zirconium-88	D, see 8 6Zr	4E+3	2E+2	9E-8	3E-10	5E-5	5E-4	
		W, see 8 6Zr	--	5E+2	2E-7	7E-10	--	--	
		Y, see 8 6Zr	--	3E+2	1E-7	4E-10	--	--	
40	Zirconium-89	D, see 8 6Zr	2E+3	4E+3	1E-6	5E-9	2E-5	2E-4	
		W, see 8 6Zr	--	2E+3	1E-6	3E-9	--	--	
		Y, see 8 6Zr	--	2E+3	1E-6	3E-9	--	--	
40	Zirconium-93	D, see 8 6Zr	1E+3	6E+0	3E-9	--	4-	--	

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			Bone surf (3E+-3)	Bone surf (2E+-1)		2E-11	E-5	4-E--4
		W, see 8 6Zr		2E+1	1E-8	--	--	--
			--	Bone surf (6E+1)	--	9E-11	--	--
		Y, see 8 6Zr	--	6E+1	2E-8	--	--	--
			--	Bone surf (7E+1)	--	9E-11	--	--
40	Zirconium-95	D, see 8 6Zr	1E+3 --	1E+2	5E-8 --	--	2E-5	2E-4
				Bone surf (3E+-2)		4E-10	--	--
		W, see 8 6Zr	--	4E+2	2E-7	5E-10	--	--
		Y, see 8 6Zr	--	3E+2	1E-7	4E-10	--	--
40	Zirconium-97	D, see 8 6Zr	6E+2	2E+3	8E-7	3E-9	9E-6	9E-5
		W, see 8 6Zr	--	1E+3	6E-7	2E-9	--	--
		Y, see 8 6Zr	--	1E+3	5E-7	2E-9	--	--
41	Niobium-88 2	W, all compounds except those given for Y	5E+4	2E+5	9E-5	3E-7	--	--
			St wall (7E+4)	--	--	--	1E-3	1E-2
		Y, oxides and hydroxides	--	2E+5	9E-5	3E-7	--	--
41	Niobium-89 2 (66 min)	W, see 8 8Nb	1E+4	4E+4	2E-5	6E-8	1E-4	1E-3
		Y, see 8 8Nb	--	4E+4	2E-5	5E-8	--	--
41	Niobium-89 (122 min)	W, see 8 8Nb	5E+3	2E+4	8E-6	3E-8	7E-5	7E-4
		Y, see 8 8Nb	--	2E+4	6E-6	2E-8	--	--
41	Niobium-90	W, see 8 8Nb	1E+3	3E+3	1E-6	4E-9	1E-5	1E-4
		Y, see 8 8Nb	--	2E+3	1E-6	3E-9	--	--
41	Niobium-93m	W, see 8 8Nb	9E+3	2E+3	8E-7	3E-9	--	--
			LLI wall (1E+4)	--	--	--	2E-4	2E-3
		Y, see 8 8Nb	--	2E+2	7E-8	2E-10	--	--
41	Niobium-94	W, see 8 8Nb	9E+2	2E+2	8E-8	3E-10	1E-5	1E-4
		Y, see 8 8Nb	--	2E+1	6E-9	2E-11	--	--
41	Niobium-95m	W, see 8 8Nb	2E+3	3E+3 --	1E-6 --	4E-9 --	-- 3-	--
			LLI wall (2E+-3)				E-5	3-E--4
		Y, see 8 8Nb	--	2E+3	9E-7	3E-9	--	--
41	Niobium-95	W, see 8 8Nb	2E+3	1E+3	5E-7	2E-9	3E-5	3E-4
		Y, see 8 8Nb	--	1E+3	5E-7	2E-9	--	--
41	Niobium-96	W, see 8 8Nb	1E+3	3E+3	1E-6	4E-9	2E-5	2E-4
		Y, see 8 8Nb	--	2E+3	1E-6	3E-9	--	--
41	Niobium-97 2	W, see 8 8Nb	2E+4	8E+4	3E-5	1E-7	3E-4	3E-3

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		Y, see 8 8Nb	--	7E+4	3E-5	1E-7	--	--
41	Niobium-98 2	W, see 8 8Nb	1E+4	5E+4	2E-5	8E-8	2E-4	2E-3
		Y, see 8 8Nb	--	5E+4	2E-5	7E-8	--	--
42	Molybdenum-- 90	D, all compounds except those given for Y	4E+3	7E+3	3E-6	1E-8	3E-5	3E-4
		Y, oxides, hydroxides, and MoS	2E+3	5E+3	2E-6	6E-9	--	--
42	Molybdenum-- 93m	D, see 9 0Mo	9E+3	2E+4	7E-6	2E-8	6E-5	6E-4
		Y, see 9 0Mo	4E+3	1E+4	6E-6	2E-8	--	--
42	Molybdenum-- 93	D, see 9 0Mo	4E+3	5E+3	2E-6	8E-9	5E-5	5E-4
		Y, see 9 0Mo	2E+4	2E+2	8E-8	2E-10	--	--
42	Molybdenum-- 99	D, see 9 0Mo	2E+3 LLI wall (1E+-3)	3E+3 --	1E-6 --	4E-9 --	-- 2-E-5	-- 2-E--4
		Y, see 9 0Mo	1E+3	1E+3	6E-7	2E-9	--	--
42	Molybdenum-- 101 2	D, see 9 0Mo	4E+4 St wall (5E+-4)	1E+5 --	6E-5 --	2E-7 --	-- 7-E-4	-- 7-E--3
		Y, see 9 0Mo	--	1E+5	6E-5	2E-7	--	--
43	Technetium-- 93m 2	D, all compounds except those given for W	7E+4	2E+5	6E-5	2E-7	1E-3	1E-2
		W, oxides, hydroxides, halides, and nitrates	--	3E+5	1E-4	4E-7	--	--
43	Technetium-- 93	D, see 9 3mTc	3E+4	7E+4	3E-5	1E-7	4E-4	4E-3
		W, see 9 3mTc	--	1E+5	4E-5	1E-7	--	--
43	Technetium-- 94m 2	D, see 9 3mTc	2E+4	4E+4	2E-5	6E-8	3E-4	3E-3
		W, see 9 3mTc	--	6E+4	2E-5	8E-8	--	--
43	Technetium-- 94	D, see 9 3mTc	9E+3	2E+4	8E-6	3E-8	1E-4	1E-3
		W, see 9 3mTc	--	2E+4	1E-5	3E-8	--	--
43	Technetium-- 95m	D, see 9 3mTc	4E+3	5E+3	2E-6	8E-9	5E-5	5E-4
		W, see 9 3mTc	--	2E+3	8E-7	3E-9	--	--
43	Technetium-- 95	D, see 9 3mTc	1E+4	2E+4	9E-6	3E-8	1E-4	1E-3
		W, see 9 3mTc	--	2E+4	8E-6	3E-8	--	--
43	Technetium-- 96m 2	D, see 9 3mTc	2E+5	3E+5	1E-4	4E-7	2E-3	2E-2
		W, see 9 3mTc	--	2E+5	1E-4	3E-7	--	--

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[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

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******************************************************************************
******** This is piece 1. -- It begins at character 1 of table line 1. ********
******************************************************************************

43   Technetium-96       D, see 9 3mTc
                         W, see 9 3mTc
43   Technetium-97m      D, see 9 3mTc
                         W, see 9 3mTc
43   Technetium-97       D, see 9 3mTc
                         W, see 9 3mTc
43   Technetium-98       D, see 9 3mTc
                         W, see 9 3mTc
43   Technetium-99m      D, see 9 3mTc
                         W, see 9 3mTc
43   Technetium-99       D, see 9 3mTc

                         W, see 9 3mTc
43   Technetium-101 2    D, see 9 3mTc

                         W, see 9 3mTc
43   Technetium-104 2    D, see 9 3mTc

                         W, see 9 3mTc
44   Ruthenium-94 2      D, all compounds except those given for W and Y
                         W, halides
                         Y, oxides and hydroxides
44   Ruthenium-97        D, see 9 4Ru
                         W, see 9 4Ru
                         Y, see 9 4Ru
44   Ruthenium-103       D, see 9 4Ru
                         W, see 9 4Ru
                         Y, see 9 4Ru
44   Ruthenium-105       D, see 9 4Ru
                         W, see 9 4Ru
                         Y, see 9 4Ru
44   Ruthenium-106       D, see 9 4Ru

                         W, see 9 4Ru
                         Y, see 9 4Ru
45   Rhodium-99m         D, all compounds except those given for W and Y
                         W, halides
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		Y, oxides and hydroxides
45	Rhodium-99	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-100	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-101m	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-101	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-102m	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-102	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-103m 2	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-105	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-106m	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
45	Rhodium-107 2	D, see 9 9mRh
		W, see 9 9mRh
		Y, see 9 9mRh
46	Palladium-100	D, all compounds except those given for W and Y
		W, nitrates
		Y, oxides and hydroxides
46	Palladium-101	D, see 1 0 0Pd
		W, see 1 0 0Pd
		Y, see 1 0 0Pd
46	Palladium-103	D, see 1 0 0Pd
		W, see 1 0 0Pd
		Y, see 1 0 0Pd
46	Palladium-107	D, see 1 0 0Pd
		W, see 1 0 0Pd
		Y, see 1 0 0Pd
46	Palladium-109	D, see 1 0 0Pd
		W, see 1 0 0Pd
		Y, see 1 0 0Pd
47	Silver-102 2	D, all compounds
		except those

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		given for W
		and Y
		W, nitrates and sulfides
		Y, oxides and hydroxides
47	Silver-103 2	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-104m 2	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-104 2	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-105	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-106m	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-106 2	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-108m	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-110m	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-111	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-112	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
47	Silver-115 2	D, see 1 0 2Ag
		W, see 1 0 2Ag
		Y, see 1 0 2Ag
48	Cadmium-104 2	D, all compounds except those given for W and Y
		W, sulfides, halides, and nitrates
		Y, oxides and hydroxides
48	Cadmium-107	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-109	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-113m	D, see 1 0 4Cd

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		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-113	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-115m	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-117m	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
48	Cadmium-117	D, see 1 0 4Cd
		W, see 1 0 4Cd
		Y, see 1 0 4Cd
49	Indium-109	D, all compounds except those given for W
		W, oxides, hydroxides, halides, and nitrates
49	Indium-110 2 (69.1 min)	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-110 (4.9 h)	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-111	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-112	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-113m 2	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-114m	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-115m	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-115	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-116m 2	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-117m 2	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-117 2	D, see 1 0 9In
		W, see 1 0 9In
49	Indium-119m 2	D, see 1 0 9In
		W, see 1 0 9In
50	Tin-110	D, all compounds except those given for W
		W, sulfides, oxides, hydroxides, halides, nitrates, and stannic phosphate
50	Tin-111 2	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-113	D, see 1 1 0Sn

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		W, see 1 1 0Sn
50	Tin-117m	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-119m	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-121m	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-121	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-123m 2	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-123	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-125	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-126	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-127	D, see 1 1 0Sn
		W, see 1 1 0Sn
50	Tin-128 2	D, see 1 1 0Sn
		W, see 1 1 0Sn
51	Antimony-115 2	D, all compounds except those given for W
		W, oxides, hydroxides, halides, sulfides, sulfates, and nitrates
51	Antimony-116m 2	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-116 2	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-117	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-118m	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-119	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-120 2 (16 min)	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-120 (5.76 d)	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-122	D, see 1 1 5Sb
		W, see 1 1 5Sb
51	Antimony-124m 2	D, see 1 1 5Sb
		W, see 1 1 5Sb

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51   Antimony-124        D, see 1 1 5Sb
                         W, see 1 1 5Sb
51   Antimony-125        D, see 1 1 5Sb
                         W, see 1 1 5Sb
51   Antimony-126m 2     D, see 1 1 5Sb

                         W, see 1 1 5Sb
51   Antimony-126        D, see 1 1 5Sb
                         W, see 1 1 5Sb
51   Antimony-127        D, see 1 1 5Sb

                         W, see 1 1 5Sb
51   Antimony-128 2      D, see 1 1 5Sb
       (10.4 min)
                         W, see 1 1 5Sb
51   Antimony-128 (9.01  D, see 1 1 5Sb
       h)
                         W, see 1 1 5Sb
51   Antimony-129        D, see 1 1 5Sb
                         W, see 1 1 5Sb
51   Antimony-130 2      D, see 1 1 5Sb
                         W, see 1 1 5Sb
51   Antimony-131 2      D, see 1 1 5Sb

                         W, see 1 1 5Sb


52   Tellurium-116       D, all compounds except those given for W
                         W, oxides, hydroxides, and nitrates
52   Tellurium-121m      D, see 1 1 6Te

                         W, see 1 1 6Te
52   Tellurium-121       D, see 1 1 6Te
                         W, see 1 1 6Te
52   Tellurium-123m      D, see 1 1 6Te

                         W, see 1 1 6Te
52   Tellurium-123       D, see 1 1 6Te

                         W, see 1 1 6Te


52   Tellurium-125m      D, see 1 1 6Te

                         W, see 1 1 6Te
52   Tellurium-127m      D, see 1 1 6Te

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

*******************************************************************************
******* This is piece 2. -- It begins at character 80 of table line 1. ********
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2E+3	3E+3	1E-6	5E-9	3E-5
--	2E+3	9E-7	3E-9	--
5E+3 St wall --	7E+3 (7E+3)	3E-6 --	-- 1E-8	6E-5 --
--	1E+3	5E-7	2E-9	--
4E+4	5E+4	2E-5	7E-8	5E-4
--	6E+3	2E-6	8E-9	--
1E+3	2E+3	7E-7	2E-9	1E-5
--	3E+2	1E-7	4E-10	--
8E+4	2E+5	6E-5	2E-7	1E-3
--	2E+5	1E-4	3E-7	--
4E+3 --	5E+3 St wall (6E+3)	2E-6 --	-- 8E-9	6E-5 --
--	7E+2	3E-7	9E-10	--
9E+4 St wall (1E+5)	3E+5 --	1E-4 --	5E-7 --	-- 2E-3
--	4E+5	2E-4	5E-7	--
2E+4 St wall (3E+4)	7E+4 --	3E-5 --	1E-7 --	-- 4E-4
--	9E+4	4E-5	1E-7	--
2E+4	4E+4	2E-5	6E-8	2E-4
--	6E+4	3E-5	9E-8	--
--	6E+4	2E-5	8E-8	--
8E+3	2E+4	8E-6	3E-8	1E-4
--	1E+4	5E-6	2E-8	--
--	1E+4	5E-6	2E-8	--
2E+3	2E+3	7E-7	2E-9	3E-5
--	1E+3	4E-7	1E-9	--
--	6E+2	3E-7	9E-10	--
5E+3	1E+4	6E-6	2E-8	7E-5
--	1E+4	6E-6	2E-8	--
--	1E+4	5E-6	2E-8	--
2E+2 LLI wall (2E+2)	9E+1 --	4E-8 --	1E-10 --	-- 3E-6
--	5E+1	2E-8	8E-11	--
--	1E+1	5E-9	2E-11	--
2E+4	6E+4	2E-5	8E-8	2E-4
--	8E+4	3E-5	1E-7	--
--	7E+4	3E-5	9E-8	--
2E+3	3E+3	1E-6	4E-9	3E-5
--	2E+3	9E-7	3E-9	--
--	2E+3	8E-7	3E-9	--
2E+3	5E+3	2E-6	7E-9	2E-5
--	4E+3	2E-6	6E-9	--
--	4E+3	2E-6	5E-9	--
6E+3	1E+4	5E-6	2E-8	8E-5
--	8E+3	4E-6	1E-8	--
--	8E+3	3E-6	1E-8	--
2E+3	5E+2	2E-7	7E-10	3E-5
--	8E+2	3E-7	1E-9	--
--	2E+2	6E-8	2E-10	--
1E+3 LLI wall	5E+2 --	2E-7 --	7E-10 --	-- 2E-5

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(1E+3)				
--	4E+2	2E-7	5E-10	--
--	1E+2	5E-8	2E-10	--
6E+2	9E+1	4E-8	1E-10	8E-6
--	2E+2	7E-8	2E-10	--
--	6E+1	2E-8	8E-11	--
4E+5	1E+6	5E-4	2E-6	6E-3
--	1E+6	5E-4	2E-6	--
--	1E+6	5E-4	2E-6	--
4E+3 LLI wall	1E+4 --	5E-6 --	2E-8 --	-- 5E-5
(4E+3)				
--	6E+3	3E-6	9E-9	--
--	6E+3	2E-6	8E-9	--
8E+3	3E+4	1E-5	4E-8	1E-4
--	4E+4	2E-5	5E-8	--
--	4E+4	1E-5	5E-8	--
7E+4 St wall	2E+5 --	1E-4 --	3E-7 --	-- 1E-3
(9E+4)				
--	3E+5	1E-4	4E-7	--
--	3E+5	1E-4	3E-7	--
1E+3	1E+3	6E-7	2E-9	2E-5
--	1E+3	5E-7	2E-9	--
--	1E+3	6E-7	2E-9	--
1E+4	3E+4	1E-5	5E-8	2E-4
--	3E+4	1E-5	5E-8	--
--	3E+4	1E-5	4E-8	--
6E+3 LLI wall	6E+3 --	3E-6 --	9E-9 --	-- 1E-4
(7E+3)				
--	4E+3	2E-6	6E-9	--
--	4E+3	1E-6	5E-9	--
3E+4 LLI wall	2E+4 Kidneys	9E-6 --	-- 3E-8	-- 5E-4
(4E+4)	(2E+4)			
--	7E+3	3E-6	1E-8	--
--	4E+2	2E-7	6E-10	--
2E+3	6E+3	3E-6	9E-9	3E-5
--	5E+3	2E-6	8E-9	--
--	5E+3	2E-6	6E-9	--
5E+4	2E+5	8E-5	2E-7	--
St wall				
(6E+4)	--	--	--	9E-4
--	2E+5	9E-5	3E-7	--
--	2E+5	8E-5	3E-7	--
4E+4	1E+5	4E-5	1E-7	5E-4
--	1E+5	5E-5	2E-7	--
--	1E+5	5E-5	2E-7	--
3E+4	9E+4	4E-5	1E-7	4E-4
--	1E+5	5E-5	2E-7	--
--	1E+5	5E-5	2E-7	--
2E+4	7E+4	3E-5	1E-7	3E-4
--	1E+5	6E-5	2E-7	--
--	1E+5	6E-5	2E-7	--
3E+3	1E+3	4E-7	1E-9	4E-5

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--	2E+3	7E-7	2E-9	--
--	2E+3	7E-7	2E-9	--
8E+2	7E+2	3E-7	1E-9	1E-5
--	9E+2	4E-7	1E-9	--
--	9E+2	4E-7	1E-9	--
6E+4 St wall (6E+4)	2E+5 --	8E-5 --	3E-7 --	-- 9E-4
--	2E+5	9E-5	3E-7	--
--	2E+5	8E-5	3E-7	--
6E+2	2E+2	8E-8	3E-10	9E-6
--	3E+2	1E-7	4E-10	--
--	2E+1	1E-8	3E-11	--
5E+2	1E+2	5E-8	2E-10	6E-6
--	2E+2	8E-8	3E-10	--
--	9E+1	4E-8	1E-10	--
9E+2 LLI wall (1E+3)	2E+3 Liver (2E+3)	6E-7 --	-- 2E-9	-- 2E-5
--	9E+2	4E-7	1E-9	--
--	9E+2	4E-7	1E-9	--
3E+3	8E+3	3E-6	1E-8	4E-5
--	1E+4	4E-6	1E-8	--
--	9E+3	4E-6	1E-8	--
3E+4 St wall (3E+4)	9E+4 --	4E-5 --	1E-7 --	-- 4E-4
--	9E+4	4E-5	1E-7	--
--	8E+4	3E-5	1E-7	--
2E+4	7E+4	3E-5	9E-8	3E-4
--	1E+5	5E-5	2E-7	--
--	1E+5	5E-5	2E-7	--
2E+4	5E+4	2E-5	8E-8	3E-4
--	6E+4	2E-5	8E-8	--
--	5E+4	2E-5	7E-8	--
3E+2 Kidneys (4E+2)	4E+1 Kidneys (5E+1)	1E-8 --	-- 7E-11	-- 6E-6
-- --	1E+2 Kidneys (1E+2)	5E-8 --	-- 2E-10	-- --
--	1E+2	5E-8	2E-10	--
2E+1 Kidneys (4E+1)	2E+0 Kidneys (4E+0)	1E-9 --	-- 5E-12	-- 5E-7
-- --	8E+0 Kidneys (1E+1)	4E-9 --	-- 2E-11	-- --
--	1E+1	5E-9	2E-11	--
2E+1 Kidneys (3E+1)	2E+0 Kidneys (3E+0)	9E-10 --	-- 5E-12	-- 4E-7
-- --	8E+0 Kidneys (1E+1)	3E-9 --	-- 2E-11	-- --
--	1E+1	6E-9	2E-11	--
3E+2 --	5E+1 Kidneys (8E+1)	2E-8 --	-- 1E-10	4E-6 --
--	1E+2	5E-8	2E-10	--
--	1E+2	6E-8	2E-10	--
5E+3	1E+4	5E-6	2E-8	6E-5
--	2E+4	7E-6	2E-8	--

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--	1E+4	6E-6	2E-8	--
5E+3	1E+4	5E-6	2E-8	6E-5
--	2E+4	7E-6	2E-8	--
--	1E+4	6E-6	2E-8	--
2E+4	4E+4	2E-5	6E-8	3E-4
--	6E+4	3E-5	9E-8	--
2E+4	4E+4	2E-5	6E-8	2E-4
--	6E+4	2E-5	8E-8	--
5E+3	2E+4	7E-6	2E-8	7E-5
--	2E+4	8E-6	3E-8	--
4E+3	6E+3	3E-6	9E-9	6E-5
--	6E+3	3E-6	9E-9	--
2E+5	6E+5	3E-4	9E-7	2E-3
--	7E+5	3E-4	1E-6	--
5E+4	1E+5	6E-5	2E-7	7E-4
--	2E+5	8E-5	3E-7	--
3E+2 LLI wall (4E+2)	6E+1 --	3E-8 --	9E-11 --	-- 5E-6
--	1E+2	4E-8	1E-10	--
1E+4	4E+4	2E-5	6E-8	2E-4
--	5E+4	2E-5	7E-8	--
4E+1	1E+0	6E-10	2E-12	5E-7
--	5E+0	2E-9	8E-12	--
2E+4	8E+4	3E-5	1E-7	3E-4
--	1E+5	5E-5	2E-7	--
1E+4	3E+4	1E-5	5E-8	2E-4
--	4E+4	2E-5	6E-8	--
6E+4	2E+5	7E-5	2E-7	8E-4
--	2E+5	9E-5	3E-7	--
4E+4 St wall (5E+4)	1E+5 --	5E-5 --	2E-7 --	-- 7E-4
--	1E+5	6E-5	2E-7	--
4E+3	1E+4	5E-6	2E-8	5E-5
--	1E+4	5E-6	2E-8	--
7E+4	2E+5	9E-5	3E-7	1E-3
--	3E+5	1E-4	4E-7	--
2E+3 LLI wall (2E+3)	1E+3 --	5E-7 --	2E-9 --	-- 3E-5
--	5E+2	2E-7	8E-10	--
2E+3 LLI wall (2E+3)	1E+3 Bone surf (2E+3)	5E-7 --	-- 3E-9	-- 3E-5
--	1E+3	6E-7	2E-9	--
3E+3 LLI wall (4E+3)	2E+3 --	1E-6 --	3E-9 --	-- 6E-5
--	1E+3	4E-7	1E-9	--
3E+3 LLI wall (4E+3)	9E+2 --	4E-7 --	1E-9 --	-- 5E-5
--	5E+2	2E-7	8E-10	--
6E+3 LLI wall (6E+3)	2E+4 --	6E-6 --	2E-8 --	-- 8E-5
--	1E+4	5E-6	2E-8	--

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5E+4	1E+5	5E-5	2E-7	7E-4
--	1E+5	6E-5	2E-7	--
5E+2 LLI wall (6E+2)	6E+2 --	3E-7 --	9E-10 --	-- 9E-6
--	2E+2	7E-8	2E-10	--
4E+2 LLI wall (5E+2)	9E+2 --	4E-7 --	1E-9 --	-- 6E-6
--	4E+2	1E-7	5E-10	--
3E+2	6E+1	2E-8	8E-11	4E-6
--	7E+1	3E-8	9E-11	--
7E+3	2E+4	8E-6	3E-8	9E-5
--	2E+4	8E-6	3E-8	--
9E+3	3E+4	1E-5	4E-8	1E-4
--	4E+4	1E-5	5E-8	--
8E+4	2E+5	1E-4	3E-7	1E-3
--	3E+5	1E-4	4E-7	--
2E+4	7E+4	3E-5	1E-7	3E-4
--	1E+5	6E-5	2E-7	--
7E+4 St wall (9E+4)	3E+5 --	1E-4 --	4E-7 --	-- 1E-3
--	3E+5	1E-4	5E-7	--
7E+4	2E+5	9E-5	3E-7	9E-4
--	3E+5	1E-4	4E-7	--
6E+3	2E+4	8E-6	3E-8	7E-5
5E+3	2E+4	9E-6	3E-8	--
2E+4	5E+4	2E-5	6E-8	2E-4
2E+4	3E+4	1E-5	4E-8	--
1E+5 St wall (2E+5)	4E+5 --	2E-4 --	6E-7 --	-- 2E-3
--	5E+5	2E-4	7E-7	--
1E+3	2E+3	9E-7	3E-9	1E-5
9E+2	1E+3	5E-7	2E-9	--
8E+2 LLI wall (8E+2)	2E+3 --	1E-6 --	3E-9 --	-- 1E-5
7E+2	1E+3	4E-7	2E-9	--
3E+5	8E+5	4E-4	1E-6	3E-3
2E+5	6E+5	2E-4	8E-7	--
6E+2	9E+2	4E-7	1E-9	7E-6
5E+2	2E+2	1E-7	3E-10	--
2E+3	2E+3	1E-6	3E-9	3E-5
--	5E+2	2E-7	7E-10	--
5E+4 St wall (7E+4)	2E+5 --	8E-5 --	3E-7 --	-- 9E-4
--	2E+5	8E-5	3E-7	--
6E+2	1E+3	5E-7	2E-9	7E-6
5E+2	5E+2	2E-7	7E-10	--
8E+2 LLI wall (8E+2)	2E+3 --	9E-7 --	3E-9 --	-- 1E-5
7E+2	9E+2	4E-7	1E-9	--
8E+4 St wall (1E+5)	4E+5 --	2E-4 --	5E-7 --	-- 1E-3

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```
  --                4E+5              2E-4        6E-7        --
  1E+3              4E+3              2E-6        6E-9        2E-5

  --                3E+3              1E-6        5E-9        --
  3E+3              9E+3              4E-6        1E-8        4E-5
  --                9E+3              4E-6        1E-8        --
  2E+4              6E+4              3E-5        9E-8        3E-4
  --                8E+4              3E-5        1E-7        --
  1E+4 Thyroid      2E+4 Thyroid      1E-5 --     -- 6E-8     -- 2E-4
    (2E+4)            (4E+4)
  --                2E+4              1E-5        --          --
                    Thyroid
  --                (4E+4)            --          6E-8        --
  8E+3              2E+4              9E-6        3E-8        1E-4
  --                3E+4              1E-5        4E-8        --
  5E+2 Bone surf    2E+2 Bone surf    8E-8 --     -- 5E-10    -- 1E-5
    (7E+2)            (4E+2)
  --                4E+2              2E-7        6E-10       --
  3E+3              4E+3              2E-6        6E-9        4E-5
  --                3E+3              1E-6        4E-9        --
  6E+2 Bone surf    2E+2 Bone surf    9E-8 --     -- 8E-10    -- 1E-5
    (1E+3)            (5E+2)
  --                5E+2              2E-7        8E-10       --
  5E+2 Bone surf    2E+2 Bone surf    8E-8 --     -- 7E-10    -- 2E-5
    (1E+3)            (5E+2)
  --                4E+2              2E-7        --          --
                    Bone surf
  --                (1E+3)            --          2E-9        --
  1E+3 Bone surf    4E+2 Bone surf    2E-7 --     -- 1E-9     -- 2E-5
    (1E+3)            (1E+3)
  --                7E+2              3E-7        1E-9        --
  6E+2 --           3E+2 Bone surf    1E-7 --     -- 6E-10    9E-6 --
                      (4E+2)
80..+...90....+....0....+...10....+...20....+...30....+...40....+...50..

*******************************************************************************
******* This is piece 3. -- It begins at character 152 of table line 1. *******
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 3E-4
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 6E-4 --
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 5E-3
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-- 2E-2

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4E-4
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9E-5 --

152....60

TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

		W, see 1 1 6Te	--	3E+2	1E-7	4E-10	--	--
52	Tellurium-127	D, see 1 1 6Te	7E+3	2E+4	9E-6	3E-8	1E-4	1E-3
		W, see 1 1 6Te	--	2E+4	7E-6	2E-8	--	--
52	Tellurium-129m	D, see 1 1 6Te	5E+2	6E+2	3E-7	9E-10	7E-6	7E-5
		W, see 1 1 6Te	--	2E+2	1E-7	3E-10	--	--
52	Tellurium-129 2	D, see 1 1 6Te	3E+4	6E+4	3E-5	9E-8	4E-4	4E-3
		W, see 1 1 6Te	--	7E+4	3E-5	1E-7	--	--
52	Tellurium-131m	D, see 1 1 6Te	3E+2 T-	4E+2 T-	2E-7 --	-- 2E-9	-- 8-	--
			hyro-	hyro-			E-6	8-
			id (-	id (-				E-
			6E+2)	1E+3)				-5
		W, see 1 1 6Te	--	4E+2	2E-7	--	--	--
				Thyroid				
			--	(9E+2)	--	1E-9	--	--
52	Tellurium-131 2	D, see 1 1 6Te	3E+3 T-	5E+3 T-	2E-6 --	-- 2E-8	-- 8-	--
			hyro-	hyro-			E-5	8-
			id (-	id (-				E-
			6E+3)	1E+4)				-4
		W, see 1 1 6Te	--	5E+3	2E-6	--	--	--
				Thyroid				
			--	(1E+4)	--	2E-8	--	--
52	Tellurium-132	D, see 1 1 6Te	2E+2 T-	2E+2 T-	9E-8 --	-- 1E-9	-- 9-	--
			hyro-	hyro-			E-6	9-
			id (-	id (-				E-
			7E+2)	8E+2)				-5
		W, see 1 1 6Te	--	2E+2	9E-8	--	--	--
				Thyroid				
			--	(6E+2)	--	9E-10	--	--
52	Tellurium-133m 2	D, see 1 1 6Te	3E+3 T-	5E+3 T-	2E-6 --	-- 2E-8	-- 9-	--
			hyro-	hyro-			E-5	9-

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					id (-6E+3)	id (-1E+4)					E-4
		W, see 1 1 6Te		--	5E+3	2E-6	--		--	--	
				--	Thyroid (1E+4)	--	2E-8		--	--	
52	Tellurium-133 2	D, see 1 1 6Te	1E+4	T-	2E+4	T-	9E-6	-- -- 8E-8	-- 4-	--	
					hyro-id (-3E+4)	hyro-id (-6E+4)			E-4	4-E-3	
		W, see 1 1 6Te		--	2E+4	9E-6	--		--	--	
				--	Thyroid (6E+4)	--	8E-8		--	--	
52	Tellurium-134 2	D, see 1 1 6Te	2E+4	T-	2E+4	T-	1E-5	-- -- 7E-8	-- 3-	--	
					hyro-id (-2E+4)	hyro-id (-5E+4)			E-4	3-E-3	
		W, see 1 1 6Te		--	2E+4	1E-5	--		--	--	
				--	Thyroid (5E+4)	--	7E-8		--	--	
53	Iodine-120m 2	D, all compounds	1E+4	T-	2E+4	--	9E-6	-- 3E-8 --	-- 2-	--	
					hyro-id (-1E+4)				E-4	2-E-3	
53	Iodine-120 2	D, all compounds	4E+3	T-	9E+3	T-	4E-6	-- -- 2E-8	-- 1-	--	
					hyro-id (-8E+3)	hyro-id (-1E+4)			E-4	1-E-3	
53	Iodine-121	D, all compounds	1E+4	T-	2E+4	T-	8E-6	-- -- 7E-8	-- 4-	--	
					hyro-id (-3E+4)	hyro-id (-5E+4)			E-4	4-E-3	
53	Iodine-123	D, all compounds	3E+3	T-	6E+3	T-	3E-6	-- -- 2E-8	-- 1-	--	
					hyro-id (-1E+4)	hyro-id (-2E+4)			E-4	1-E-3	
53	Iodine-124	D, all compounds	5E+1	T-	8E+1	T-	3E-8	-- -- 4E-10	-- 2-	--	
					hyro-id (-2E+2)	hyro-id (-3E+2)			E-6	2-E-5	
53	Iodine-125	D, all compounds	4E+1	T-	6E+1	T-	3E-8	-- -- 3E-10	-- 2-	--	
					hyro-id (-1E+2)	hyro-id (-2E+2)			E-6	2-E-5	
53	Iodine-126	D, all compounds	2E+1	T-	4E+1	T-	1E-8	-- -- 2E-10	-- 1-	--	
					hyro-id (-7E+1)	hyro-id (-1E+2)			E-6	1-E-5	
53	Iodine-128 2	D, all compounds	4E+4	St	1E+5	--	5E-5	-- 2E-7 --	-- 8-	--	
					wall (6E+-4)				E-4	8-E-3	
53	Iodine-129	D, all compounds	5E+0	T-	9E+0	T-	4E-9	-- -- 4E-11	-- 2-	--	
					hyro-	hyro-			E-7	2-	

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			id (- 2E+1)	id (- 3E+1)					E- -6
53	Iodine-130	D, all compounds	4E+2 T- hyro- id (- 1E+3)	7E+2 T- hyro- id (- 2E+3)	3E-7	--	--	3E-9	-- 2- -- E-5 2- E- -4
53	Iodine-131	D, all compounds	3E+1 T- hyro- id (- 9E+1)	5E+1 T- hyro- id (- 2E+2)	2E-8	--	--	2E-10	-- 1- -- E-6 1- E- -5
53	Iodine-132m 2	D, all compounds	4E+3 T- hyro- id (- 1E+4)	8E+3 T- hyro- id (- 2E+4)	4E-6	--	--	3E-8	-- 1- -- E-4 1- E- -3
53	Iodine-132	D, all compounds	4E+3 T- hyro- id (- 9E+3)	8E+3 T- hyro- id (- 1E+4)	3E-6	--	--	2E-8	-- 1- -- E-4 1- E- -3
53	Iodine-133	D, all compounds	1E+2 T- hyro- id (- 5E+2)	3E+2 T- hyro- id (- 9E+2)	1E-7	--	--	1E-9	-- 7- -- E-6 7- E- -5
53	Iodine-134 2	D, all compounds	2E+4 T- hyro- id (- 3E+4)	5E+4 --	2E-5	--	6E-8	--	-- 4- -- E-4 4- E- -3
53	Iodine-135	D, all compounds	8E+2 T- hyro- id (- 3E+3)	2E+3 T- hyro- id (- 4E+3)	7E-7	--	--	6E-9	-- 3- -- E-5 3- E- -4
54	Xenon-120 2	Submersion 1	--	--	1E-5		4E-8		-- --
54	Xenon-121 2	Submersion 1	--	--	2E-6		1E-8		-- --
54	Xenon-122	Submersion 1	--	--	7E-5		3E-7		-- --
54	Xenon-123	Submersion 1	--	--	6E-6		3E-8		-- --
54	Xenon-125	Submersion 1	--	--	2E-5		7E-8		-- --
54	Xenon-127	Submersion 1	--	--	1E-5		6E-8		-- --
54	Xenon-129m	Submersion 1	--	--	2E-4		9E-7		-- --
54	Xenon-131m	Submersion 1	--	--	4E-4		2E-6		-- --
54	Xenon-133m	Submersion 1	--	--	1E-4		6E-7		-- --
54	Xenon-133	Submersion 1	--	--	1E-4		5E-7		-- --
54	Xenon-135m 2	Submersion 1	--	--	9E-6		4E-8		-- --
54	Xenon-135	Submersion 1	--	--	1E-5		7E-8		-- --
54	Xenon-138 2	Submersion 1	--	--	4E-6		2E-8		-- --
55	Cesium-125 2	D, all compounds	5E+4 St wall (9E+- 4)	1E+5 --	6E-5	--	2E-7	--	-- 1- -- E-3 1- E- -2
55	Cesium-127	D, all compounds	6E+4	9E+4	4E-5		1E-7		9E-4 9E-3
55	Cesium-129	D, all compounds	2E+4	3E+4	1E-5		5E-8		3E-4 3E-3
55	Cesium-130 2	D, all compounds	6E+4 St wall	2E+5 --	8E-5	--	3E-7	--	-- 1- -- E-3 1-

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			(1E+-5)					E--2
55	Cesium-131	D, all compounds	2E+4	3E+4	1E-5	4E-8	3E-4	3E-3
55	Cesium-132	D, all compounds	3E+3	4E+3	2E-6	6E-9	4E-5	4E-4
55	Cesium-134m	D, all compounds	1E+5 St wall (1E+-5)	1E+5 --	6E-5 --	2E-7 --	-- 2-E-3	-- 2-E--2
55	Cesium-134	D, all compounds	7E+1	1E+2	4E-8	2E-10	9E-7	9E-6
55	Cesium-135m 2	D, all compounds	1E+5	2E+5	8E-5	3E-7	1E-3	1E-2
55	Cesium-135	D, all compounds	7E+2	1E+3	5E-7	2E-9	1E-5	1E-4
55	Cesium-136	D, all compounds	4E+2	7E+2	3E-7	9E-10	6E-6	6E-5
55	Cesium-137	D, all compounds	1E+2	2E+2	6E-8	2E-10	1E-6	1E-5
55	Cesium-138 2	D, all compounds	2E+4 St wall (3E+-4)	6E+4 --	2E-5 --	8E-8 --	-- 4-E-4	-- 4-E--3
56	Barium-126 2	D, all compounds	6E+3	2E+4	6E-6	2E-8	8E-5	8E-4
56	Barium-128	D, all compounds	5E+2	2E+3	7E-7	2E-9	7E-6	7E-5
56	Barium-131m 2	D, all compounds	4E+5 St wall (5E+-5)	1E+6 --	6E-4 --	2E-6 --	-- 7-E-3	-- 7-E--2
56	Barium-131	D, all compounds	3E+3	8E+3	3E-6	1E-8	4E-5	4E-4
56	Barium-133m	D, all compounds	2E+3 LLI wall (3E+-3)	9E+3 --	4E-6 --	1E-8 --	-- 4-E-5	-- 4-E--4
56	Barium-133	D, all compounds	2E+3	7E+2	3E-7	9E-10	2E-5	2E-4
56	Barium-135m	D, all compounds	3E+3	1E+4	5E-6	2E-8	4E-5	4E-4
56	Barium-139 2	D, all compounds	1E+4	3E+4	1E-5	4E-8	2E-4	2E-3
56	Barium-140	D, all compounds	5E+2 LLI wall (6E+-2)	1E+3 --	6E-7 --	2E-9 --	-- 8-E-6	-- 8-E--5
56	Barium-141 2	D, all compounds	2E+4	7E+4	3E-5	1E-7	3E-4	3E-3
56	Barium-142 2	D, all	5E+4	1E+5	6E-5	2E-7	7E-4	7E-3

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		compounds						
57	Lanthanum-131 2	D, all compounds except those given for W	5E+4	1E+5	5E-5	2E-7	6E-4	6E-3
		W, oxides and hydroxides	--	2E+5	7E-5	2E-7	--	--
57	Lanthanum-132	D, see 1 3 1La	3E+3	1E+4	4E-6	1E-8	4E-5	4E-4
		W, see 1 3 1La	--	1E+4	5E-6	2E-8	--	--
57	Lanthanum-135	D, see 1 3 1La	4E+4	1E+5	4E-5	1E-7	5E-4	5E-3
		W, see 1 3 1La	--	9E+4	4E-5	1E-7	--	--
57	Lanthanum-137	D, see 1 3 1La	1E+4 --	6E+1	3E-8 --	-- 1E-10	2E-4	2E-3
				Liver (7E+-1)			--	--
		W, see 1 3 1La	--	3E+2	1E-7	--	--	--
			--	Liver (3E+2)	--	4E-10	--	--
57	Lanthanum-138	D, see 1 3 1La	9E+2	4E+0	1E-9	5E-12	1E-5	1E-4
		W, see 1 3 1La	--	1E+1	6E-9	2E-11	--	--
57	Lanthanum-140	D, see 1 3 1La	6E+2	1E+3	6E-7	2E-9	9E-6	9E-5
		W, see 1 3 1La	--	1E+3	5E-7	2E-9	--	--
57	Lanthanum-141	D, see 1 3 1La	4E+3	9E+3	4E-6	1E-8	5E-5	5E-4
		W, see 1 3 1La	--	1E+4	5E-6	2E-8	--	--
57	Lanthanum-142 2	D, see 1 3 1La	8E+3	2E+4	9E-6	3E-8	1E-4	1E-3
		W, see 1 3 1La	--	3E+4	1E-5	5E-8	--	--
57	Lanthanum-143 2	D, see 1 3 1La	4E+4 St wall (4E+-4)	1E+5 --	4E-5 --	1E-7 --	-- 5-E-4	-- 5-E-3
		W, see 1 3 1La	--	9E+4	4E-5	1E-7	--	--
58	Cerium-134	W, all compounds except those given for Y	5E+2	7E+2	3E-7	1E-9	--	--
			LLI wall (6E+2)	--	--	--	8E-6	8E-5
		Y, oxides, hydroxides, and fluorides	--	7E+2	3E-7	9E-10	--	--
58	Cerium-135	W, see 1 3 4Ce	2E+3	4E+3	2E-6	5E-9	2E-5	2E-4
		Y, see 1 3 4Ce	--	4E+3	1E-6	5E-9	--	--
58	Cerium-137m	W, see 1 3 4Ce	2E+3 LLI wall (2E+-3)	4E+3 --	2E-6 --	6E-9 --	-- 3-E-5	-- 3-E-4
		Y, see 1 3 4Ce	--	4E+3	2E-6	5E-9	--	--
58	Cerium-137	W, see 1 3 4Ce	5E+4	1E+5	6E-5	2E-7	7E-4	7E-3
		Y, see 1 3 4Ce	--	1E+5	5E-5	2E-7	--	--
58	Cerium-139	W, see 1 3 4Ce	5E+3	8E+2	3E-7	1E-9	7E-5	7E-4
		Y, see 1 3 4Ce	--	7E+2	3E-7	9E-10	--	--
58	Cerium-141	W, see 1 3 4Ce	2E+3	7E+2 --	3E-7 --	1E-9 --	-- 3-	--

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			LLI wall (2E+-3)					E-5	3-E--4
		Y, see 1 3 4Ce	--	6E+2	2E-7	8E-10		--	--
58	Cerium-143	W, see 1 3 4Ce	1E+3	2E+3	-- -- 8E-7	3E-9 --		-- 2-	--
			LLI wall (1E+-3)					E-5	2-E--4
		Y, see 1 3 4Ce	--	2E+3	7E-7	2E-9		--	--
58	Cerium-144	W, see 1 3 4Ce	2E+2	3E+1	-- 1E-8 --	4E-11 --		-- 3-	--
			LLI wall (3E+-2)					E-6	3-E--5
		Y, see 1 3 4Ce	--	1E+1	6E-9	2E-11		--	--
59	Praseodymium-136 2	W, all compounds except those given for Y	5E+4	2E+5	1E-4	3E-7		--	--
			St wall (7E+4)	--	--	--		1E-3	1E-2
		Y, oxides, hydroxides, carbides, and fluorides	--	2E+5	9E-5	3E-7		--	--
59	Praseodymium-137 2	W, see 1 3 6Pr	4E+4	2E+5	6E-5	2E-7		5E-4	5E-3
		Y, see 1 3 6Pr	--	1E+5	6E-5	2E-7		--	--
59	Praseodymium-13-8m	W, see 1 3 6Pr	1E+4	5E+4	2E-5	8E-8		1E-4	1E-3
		Y, see 1 3 6Pr	--	4E+4	2E-5	6E-8		--	--
59	Praseodymium-139	W, see 1 3 6Pr	4E+4	1E+5	5E-5	2E-7		6E-4	6E-3
		Y, see 1 3 6Pr	--	1E+5	5E-5	2E-7		--	--
59	Praseodymium-14-2m 2	W, see 1 3 6Pr	8E+4	2E+5	7E-5	2E-7		1E-3	1E-2
		Y, see 1 3 6Pr	--	1E+5	6E-5	2E-7		--	--
59	Praseodymium-142	W, see 1 3 6Pr	1E+3	2E+3	9E-7	3E-9		1E-5	1E-4
		Y, see 1 3 6Pr	--	2E+3	8E-7	3E-9		--	--
59	Praseodymium-143	W, see 1 3 6Pr	9E+2	8E+2	-- 3E-7 --	1E-9 --		-- 2-	--
			LLI wall (1E+-3)					E-5	2-E--4
		Y, see 1 3 6Pr	--	7E+2	3E-7	9E-10		--	--
59	Praseodymium-144 2	W, see 1 3 6Pr	3E+4	St 1E+5	-- 5E-5 --	2E-7 --		-- 6-	--
			wall (4E+-4)					E-4	6-E--3
		Y, see 1 3 6Pr	--	1E+5	5E-5	2E-7		--	--
59	Praseodymium-145	W, see 1 3 6Pr	3E+3	9E+3	4E-6	1E-8		4E-5	4E-4
		Y, see 1 3 6Pr	--	8E+3	3E-6	1E-8		--	--
59	Praseodymium-147	W, see 1 3 6Pr	5E+4	St 2E+5	-- 8E-5 --	3E-7 --		-- 1-	--

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	2		wall (8E+-4)				E-3	1-E--2
		Y, see 1 3 6Pr	--	2E+5	8E-5	3E-7	--	--
60	Neodymium-136 2	W, all compounds except those given for Y	1E+4	6E+4	2E-5	8E-8	2E-4	2E-3
		Y, oxides, hydroxides, carbides, and fluorides	--	5E+4	2E-5	8E-8	--	--
60	Neodymium-138	W, see 1 3 6Nd	2E+3	6E+3	3E-6	9E-9	3E-5	3E-4
		Y, see 1 3 6Nd	--	5E+3	2E-6	7E-9	--	--
60	Neodymium-139m	W, see 1 3 6Nd	5E+3	2E+4	7E-6	2E-8	7E-5	7E-4
		Y, see 1 3 6Nd	--	1E+4	6E-6	2E-8	--	--
60	Neodymium-139 2	W, see 1 3 6Nd	9E+4	3E+5	1E-4	5E-7	1E-3	1E-2
		Y, see 1 3 6Nd	--	3E+5	1E-4	4E-7	--	--
60	Neodymium-141	W, see 1 3 6Nd	2E+5	7E+5	3E-4	1E-6	2E-3	2E-2
		Y, see 1 3 6Nd	--	6E+5	3E-4	9E-7	--	--
60	Neodymium-147	W, see 1 3 6Nd	1E+3	9E+2 --	4E-7 --	1E-9 --	-- 2-	--
			LLI wall (1E+-3)				E-5	2-E--4
		Y, see 1 3 6Nd	--	8E+2	4E-7	1E-9	--	--
60	Neodymium-149 2	W, see 1 3 6Nd	1E+4	3E+4	1E-5	4E-8	1E-4	1E-3
		Y, see 1 3 6Nd	--	2E+4	1E-5	3E-8	--	--
60	Neodymium-151 2	W, see 1 3 6Nd	7E+4	2E+5	8E-5	3E-7	9E-4	9E-3
		Y, see 1 3 6Nd	--	2E+5	8E-5	3E-7	--	--
61	Promethium-141 2	W, all compounds except those given for Y	5E+4	2E+5	8E-5	3E-7	--	--
			St wall (6E+4)	--	--	--	8E-4	8E-3
		Y, oxides, hydroxides, carbides, and fluorides	--	2E+5	7E-5	2E-7	--	--
61	Promethium-143	W, see 1 4 1Pm	5E+3	6E+2	2E-7	8E-10	7E-5	7E-4
		Y, see 1 4 1Pm	--	7E+2	3E-7	1E-9	--	--
61	Promethium-144	W, see 1 4 1Pm	1E+3	1E+2	5E-8	2E-10	2E-5	2E-4
		Y, see 1 4 1Pm	--	1E+2	5E-8	2E-10	--	--
61	Promethium-145	W, see 1 4 1Pm	1E+4	2E+2	7E-8	--	1E-4	1E-3
			--	Bone surf (2E+2)	--	3E-10	--	--
		Y, see 1 4 1Pm	--	2E+2	8E-8	3E-10	--	--
61	Promethium-146	W, see 1 4 1Pm	2E+3	5E+1	2E-8	7E-11	2E-5	2E-4
		Y, see 1 4 1Pm	--	4E+1	2E-8	6E-11	--	--
61	Promethium-147	W, see 1 4 1Pm	4E+3	1E+2	5E-8 --	-- 3E-10	-- 7-	--
			LLI	Bone			E-5	7-

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				wall (5E+-3)	surf (2E+-2)				E-4
		Y, see 1 4 1Pm	--		1E+2	6E-8	2E-10	--	--
61	Promethium-148m	W, see 1 4 1Pm	7E+2		3E+2	1E-7	4E-10	1E-5	1E-4
		Y, see 1 4 1Pm	--		3E+2	1E-7	5E-10	--	--
61	Promethium-148	W, see 1 4 1Pm	4E+2		5E+2 --	2E-7 --	8E-10 --	--	7- --
				LLI wall (5E+-2)				E-6	7-E-5
		Y, see 1 4 1Pm	--		5E+2	2E-7	7E-10	--	--
61	Promethium-149	W, see 1 4 1Pm	1E+3		2E+3 --	8E-7 --	3E-9 --	--	2- --
				LLI wall (1E+-3)				E-5	2-E-4
		Y, see 1 4 1Pm	--		2E+3	8E-7	2E-9	--	--
61	Promethium-150	W, see 1 4 1Pm	5E+3		2E+4	8E-6	3E-8	7E-5	7E-4
		Y, see 1 4 1Pm	--		2E+4	7E-6	2E-8	--	--
61	Promethium-151	W, see 1 4 1Pm	2E+3		4E+3	1E-6	5E-9	2E-5	2E-4
		Y, see 1 4 1Pm	--		3E+3	1E-6	4E-9	--	--
62	Samarium-141m 2	W, all compounds	3E+4		1E+5	4E-5	1E-7	4E-4	4E-3
62	Samarium-141 2	W, all compounds	5E+4	St wall (6E+-4)	2E+5 --	8E-5 --	2E-7 --	--	8- --
								E-4	8-E-3
62	Samarium-142 2	W, all compounds	8E+3		3E+4	1E-5	4E-8	1E-4	1E-3
62	Samarium-145	W, all compounds	6E+3		5E+2	2E-7	7E-10	8E-5	8E-4
62	Samarium-146	W, all compounds	1E+1		4E+2	1E-11	-- 9E-14	--	3- --
				Bone surf (3E+-1)	Bone surf (6E--2)	--		E-7	3-E-6
62	Samarium-147	W, all compounds	2E+1		4E+2	2E-11	-- 1E-13	--	4- --
				Bone surf (3E+-1)	Bone surf (7E--2)	--		E-7	4-E-6
62	Samarium-151	W, all compounds	1E+4		1E+2	4E-8 --	-- 2E-10	--	2- --
				LLI wall (1E+-4)	Bone surf (2E+-2)			E-4	2-E-3
62	Samarium-153	W, all compounds	2E+3		3E+3 --	1E-6 --	4E-9 --	--	3- --
				LLI wall (2E+-3)				E-5	3-E-4
62	Samarium-155 2	W, all	6E+4	St	2E+5 --	9E-5 --	3E-7 --	--	1- --

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		compounds	wall (8E+-4)				E-3	1-E--2
62	Samarium-156	W, all compounds	5E+3	9E+3	4E-6	1E-8	7E-5	7E-4
63	Europium-145	W, all compounds	2E+3	2E+3	8E-7	3E-9	2E-5	2E-4
63	Europium-146	W, all compounds	1E+3	1E+3	5E-7	2E-9	1E-5	1E-4
63	Europium-147	W, all compounds	3E+3	2E+3	7E-7	2E-9	4E-5	4E-4
63	Europium-148	W, all compounds	1E+3	4E+2	1E-7	5E-10	1E-5	1E-4
63	Europium-149	W, all compounds	1E+4	3E+3	1E-6	4E-9	2E-4	2E-3
63	Europium-150 (12.62h)	W, all compounds	3E+3	8E+3	4E-6	1E-8	4E-5	4E-4
63	Europium-150 (34.2 y)	W, all compounds	8E+2	2E+1	8E-9	3E-11	1E-5	1E-4
63	Europium-152m	W, all compounds	3E+3	6E+3	3E-6	9E-9	4E-5	4E-4
63	Europium-152	W, all compounds	8E+2	2E+1	1E-8	3E-11	1E-5	1E-4
63	Europium-154	W, all compounds	5E+2	2E+1	8E-9	3E-11	7E-6	7E-5
63	Europium-155	W, all compounds	4E+3 --	9E+1 Bone surf (1E+-2)	4E-8 --	-- 2E-10	5E-5 --	5E-4 --
63	Europium-156	W, all compounds	6E+2	5E+2	2E-7	6E-10	8E-6	8E-5
63	Europium-157	W, all compounds	2E+3	5E+3	2E-6	7E-9	3E-5	3E-4
63	Europium-158 2	W, all compounds	2E+4	6E+4	2E-5	8E-8	3E-4	3E-3
64	Gadolinium-145 2	D, all compounds	5E+4	2E+5	6E-5	2E-7	--	--
		except those given for W	St wall (5E+4)	--	--	--	6E-4	6E-3
		W, oxides, hydroxides, and fluorides	--	2E+5	7E-5	2E-7	--	--
64	Gadolinium-146	D, see 1 4 5Gd	1E+3	1E+2	5E-8	2E-10	2E-5	2E-4
		W, see 1 4 5Gd	--	3E+2	1E-7	4E-10	--	--
64	Gadolinium-147	D, see 1 4 5Gd	2E+3	4E+3	2E-6	6E-9	3E-5	3E-4
		W, see 1 4 5Gd	--	4E+3	1E-6	5E-9	--	--
64	Gadolinium-148	D, see 1 4 5Gd	1E+1 Bone surf (2E+-1)	8E+3 Bone surf (2E+-2)	3E-12 --	-- 2E-14	-- 3-E-7	-- 3-E--6

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		W, see 1 4 5Gd	--	3E-2 Bone surf	1E-11	--	--	--
			--	(6E-2)	--	8E-14	--	--
64	Gadolinium-149	D, see 1 4 5Gd	3E+3	2E+3	9E-7	3E-9	4E-5	4E-4
		W, see 1 4 5Gd	--	2E+3	1E-6	3E-9	--	--
64	Gadolinium-151	D, see 1 4 5Gd	6E+3 --	4E+2 Bone surf (6E+-2)	2E-7 --	-- 9E-10	9E-5 --	9E-4 --
		W, see 1 4 5Gd	--	1E+3	5E-7	2E-9	--	--
64	Gadolinium-152	D, see 1 4 5Gd	2E+1 Bone surf (3E+-1)	1E-2 Bone surf (2E--2)	4E-12 --	-- 3E-14	-- 4-E-7	-- 4-E-6
		W, see 1 4 5Gd	--	4E-2 Bone surf	2E-11	--	--	--
			--	(8E-2)	--	1E-13	--	--
64	Gadolinium-153	D, see 1 4 5Gd	5E+3 --	1E+2 Bone surf (2E+-2)	6E-8 --	-- 3E-10	6E-5 --	6E-4 --
		W, see 1 4 5Gd	--	6E+2	2E-7	8E-10	--	--
64	Gadolinium-159	D, see 1 4 5Gd	3E+3	8E+3	3E-6	1E-8	4E-5	4E-4
		W, see 1 4 5Gd	--	6E+3	2E-6	8E-9	--	--
65	Terbium-147 2	W, all compounds	9E+3	3E+4	1E-5	5E-8	1E-4	1E-3
65	Terbium-149	W, all compounds	5E+3	7E+2	3E-7	1E-9	7E-5	7E-4
65	Terbium-150	W, all compounds	5E+3	2E+4	9E-6	3E-8	7E-5	7E-4
65	Terbium-151	W, all compounds	4E+3	9E+3	4E-6	1E-8	5E-5	5E-4
65	Terbium-153	W, all compounds	5E+3	7E+3	3E-6	1E-8	7E-5	7E-4
65	Terbium-154	W, all compounds	2E+3	4E+3	2E-6	6E-9	2E-5	2E-4
65	Terbium-155	W, all compounds	6E+3	8E+3	3E-6	1E-8	8E-5	8E-4
65	Terbium-156m (5.0 h)	W, all compounds	2E+4	3E+4	1E-5	4E-8	2E-4	2E-3
65	Terbium-156m (24.4 h)	W, all compounds	7E+3	8E+3	3E-6	1E-8	1E-4	1E-3
65	Terbium-156	W, all compounds	1E+3	1E+3	6E-7	2E-9	1E-5	1E-4
65	Terbium-157	W, all compounds	5E+4 LLI wall (5E+-	3E+2 Bone surf (6E+-	1E-7 --	-- (6E+2)	-- 7-E-4	-- 7-E-3

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			4)	2)				
65	Terbium-158	W, all compounds	1E+3	2E+1	8E-9	3E-11	2E-5	2E-4
65	Terbium-160	W, all compounds	8E+2	2E+2	9E-8	3E-10	1E-5	1E-4
65	Terbium-161	W, all compounds	2E+3 LLI wall (2E+-3)	2E+3 --	7E-7 --	2E-9 --	-- 3- E-5	-- 3- E- -4
66	Dysprosium-155	W, all compounds	9E+3	3E+4	1E-5	4E-8	1E-4	1E-3
66	Dysprosium-157	W, all compounds	2E+4	6E+4	3E-5	9E-8	3E-4	3E-3
66	Dysprosium-159	W, all compounds	1E+4	2E+3	1E-6	3E-9	2E-4	2E-3

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[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

66	Dysprosium-165	W, all compounds
66	Dysprosium-166	W, all compounds
67	Holmium-155 2	W, all compounds
67	Holmium-157 2	W, all compounds
67	Holmium-159 2	W, all compounds
67	Holmium-161	W, all compounds
67	Holmium-162m 2	W, all compounds
67	Holmium-162 2	W, all compounds
67	Holmium-164m 2	W, all compounds
67	Holmium-164 2	W, all compounds
67	Holmium-166m	W, all compounds
67	Holmium-166	W, all compounds
67	Holmium-167	W, all compounds
68	Erbium-161	W, all compounds
68	Erbium-165	W, all compounds
68	Erbium-169	W, all compounds

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68	Erbium-171	W, all compounds
68	Erbium-172	W, all compounds
69	Thulium-162 2	W, all compounds
69	Thulium-166	W, all compounds
69	Thulium-167	W, all compounds
69	Thulium-170	W, all compounds
69	Thulium-171	W, all compounds
69	Thulium-172	W, all compounds
69	Thulium-173	W, all compounds
69	Thulium-175 2	W, all compounds
70	Ytterbium-162 2	W, all compounds except those given for Y
		Y, oxides, hydroxides, and fluorides
70	Ytterbium-166	W, see 1 6 2Yb
		Y, see 1 6 2Yb
70	Ytterbium-167 2	W, see 1 6 2Yb
		Y, see 1 6 2Yb
70	Ytterbium-169	W, see 1 6 2Yb
		Y, see 1 6 2Yb
70	Ytterbium-175	W, see 1 6 2Yb
		Y, see 1 6 2Yb
70	Ytterbium-177 2	W, see 1 6 2Yb
		Y, see 1 6 2Yb
70	Ytterbium-178 2	W, see 1 6 2Yb
		Y, see 1 6 2Yb
71	Lutetium-169	W, all compounds except those given for Y
		Y, oxides, hydroxides, and fluorides
71	Lutetium-170	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-171	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-172	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-173	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-174m	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-174	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-176m	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-176	W, see 1 6 9Lu

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		Y, see 1 6 9Lu
71	Lutetium-177m	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-177	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-178m 2	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-178 2	W, see 1 6 9Lu
		Y, see 1 6 9Lu
71	Lutetium-179	W, see 1 6 9Lu
		Y, see 1 6 9Lu
72	Hafnium-170	D, all compounds except those given for W
		W, oxides, hydroxides, carbides, and nitrates
72	Hafnium-172	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-173	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-175	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-177m 2	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-178m	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-179m	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-180m	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-181	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-182m 2	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-182	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-183 2	D, see 1 7 0Hf
		W, see 1 7 0Hf
72	Hafnium-184	D, see 1 7 0Hf
		W, see 1 7 0Hf
73	Tantalum-172 2	W, all compounds except those given for Y
		Y, elemental Ta, oxides, hydroxides, halides,

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		carbides, nitrates, and nitrides
73	Tantalum-173	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-174 2	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-175	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-176	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-177	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-178	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-179	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-180m	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-180	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-182m 2	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-182	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-183	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-184	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-185 2	W, see 1 7 2Ta
		Y, see 1 7 2Ta
73	Tantalum-186 2	W, see 1 7 2Ta
		Y, see 1 7 2Ta
74	Tungsten-176	D, all compounds
74	Tungsten-177	D, all compounds
74	Tungsten-178	D, all compounds
74	Tungsten-179 2	D, all compounds
74	Tungsten-181	D, all compounds
74	Tungsten-185	D, all compounds
74	Tungsten-187	D, all compounds
74	Tungsten-188	D, all compounds
75	Rhenium-177 2	D, all compounds except those given for W
		W, oxides, hydroxides, and nitrates
75	Rhenium-178 2	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-181	D, see 1 7 7Re
		W, see 1 7 7Re

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75	Rhenium-182 (12.7 h)	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-182 (64.0 h)	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-184m	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-184	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-186m	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-186	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-187	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-188m 2	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-188	D, see 1 7 7Re
		W, see 1 7 7Re
75	Rhenium-189	D, see 1 7 7Re
		W, see 1 7 7Re
76	Osmium-180 2	D, all compounds except those given for W and Y
		W, halides and nitrates
		Y, oxides and hydroxides
76	Osmium-181 2	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-182	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-185	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-189m	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-191m	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-191	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-193	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
76	Osmium-194	D, see 1 8 0Os
		W, see 1 8 0Os
		Y, see 1 8 0Os
77	Iridium-182 2	D, all compounds
		except those

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		given for W
		and Y
		W, halides, nitrates, and metallic iridium
		Y, oxides and hydroxides
77	Iridium-184	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-185	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-186	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-187	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-188	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-189	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-190m 2	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-190	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-192m	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-192	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-194m	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-194	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-195m	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
77	Iridium-195	D, see 1 8 2Ir
		W, see 1 8 2Ir
		Y, see 1 8 2Ir
78	Platinum-186	D, all compounds
78	Platinum-188	D, all compounds
78	Platinum-189	D, all compounds
78	Platinum-191	D, all compounds
78	Platinum-193m	D, all compounds

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78   Platinum-193           D, all compounds

78   Platinum-195m          D, all compounds

78   Platinum-197m 2        D, all compounds
78   Platinum-197           D, all compounds
78   Platinum-199 2         D, all compounds
78   Platinum-200           D, all compounds
79   Gold-193               D, all compounds except those given for W and Y
                            W, halides and nitrates
                            Y, oxides and hydroxides
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

*******************************************************************************
******* This is piece 2. -- It begins at character 80 of table line 1. ********
*******************************************************************************

  1E+4              5E+4              2E-5        6E-8        2E-4
  6E+2 LLI wall     7E+2 --           3E-7 --     1E-9 --     -- 1E-5
    (8E+2)
  4E+4              2E+5              6E-5        2E-7        6E-4
  3E+5              1E+6              6E-4        2E-6        4E-3
  2E+5              1E+6              4E-4        1E-6        3E-3
  1E+5              4E+5              2E-4        6E-7        1E-3
  5E+4              3E+5              1E-4        4E-7        7E-4
  5E+5 St wall      2E+6 --           1E-3 --     3E-6 --     -- 1E-2
    (8E+5)
  1E+5              3E+5              1E-4        4E-7        1E-3
  2E+5 St wall      6E+5 --           3E-4 --     9E-7 --     -- 3E-3
    (2E+5)
  6E+2              7E+0              3E-9        9E-12       9E-6
  9E+2 LLI wall     2E+3 --           7E-7 --     2E-9 --     -- 1E-5
    (9E+2)
  2E+4              6E+4              2E-5        8E-8        2E-4
  2E+4              6E+4              3E-5        9E-8        2E-4
  6E+4              2E+5              8E-5        3E-7        9E-4
  3E+3 LLI wall     3E+3 --           1E-6 --     4E-9 --     -- 5E-5
    (4E+3)
  4E+3              1E+4              4E-6        1E-8        5E-5
  1E+3 LLI wall     1E+3 --           6E-7 --     2E-9 --     -- 2E-5
    (E+3)
  7E+4 St wall      3E+5 --           1E-4 --     4E-7 --     -- 1E-3
    (7E+4)
  4E+3              1E+4              6E-6        2E-8        6E-5
  2E+3 LLI wall     2E+3 --           8E-7 --     3E-9 --     -- 3E-5
    (2E+3)
  8E+2 LLI wall     2E+2 --           9E-8 --     3E-10 --    -- 1E-5
    (1E+3)
  1E+4 LLI wall     3E+2 Bone surf    1E-7 --     -- 8E-10    -- 2E-4
    (1E+4)            (6E+2)
  7E+2 LLI wall     1E+3 --           5E-7 --     2E-9 --     -- 1E-5
    (8E+2)
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4E+3	1E+4	5E-6	2E-8	6E-5
7E+4 St wall (9E+4)	3E+5 --	1E-4 --	4E-7 --	-- 1E-3
7E+4	3E+5	1E-4	4E-7	1E-3
--	3E+5	1E-4	4E-7	--
1E+3	2E+3	8E-7	3E-9	2E-5
--	2E+3	8E-7	3E-9	--
3E+5	8E+5	3E-4	1E-6	4E-3
--	7E+5	3E-4	1E-6	--
2E+3	8E+2	4E-7	1E-9	2E-5
--	7E+2	3E-7	1E-9	--
3E+3 LLI wall (3E+3)	4E+3 --	1E-6 --	5E-9 --	-- 4E-5
--	3E+3	1E-6	5E-9	--
2E+4	5E+4	2E-5	7E-8	2E-4
--	5E+4	2E-5	6E-8	--
1E+4	4E+4	2E-5	6E-8	2E-4
--	4E+4	2E-5	5E-8	--
3E+3	4E+3	2E-6	6E-9	3E-5
--	4E+3	2E-6	6E-9	--
1E+3	2E+3	9E-7	3E-9	2E-5
--	2E+3	8E-7	3E-9	--
2E+3	2E+3	8E-7	3E-9	3E-5
--	2E+3	8E-7	3E-9	--
1E+3	1E+3	5E-7	2E-9	1E-5
--	1E+3	5E-7	2E-9	--
5E+3 --	3E+2 Bone surf (5E+2)	1E-7 --	-- 6E-10	7E-5 --
--	3E+2	1E-7	4E-10	--
2E+3 LLI wall (3E+3)	2E+2 Bone surf (3E+2)	1E-7 --	-- 5E-10	-- 4E-5
--	2E+2	9E-8	3E-10	--
5E+3 --	1E+2 Bone surf (2E+2)	5E-8 --	-- 3E-10	7E-5 --
--	2E+2	6E-8	2E-10	--
8E+3	3E+4	1E-5	3E-8	1E-4
--	2E+4	9E-6	3E-8	--
7E+2 --	5E+0 Bone surf (1E+1)	2E-9 --	-- 2E-11	1E-5 --
--	8E+0	3E-9	1E-11	--
7E+2 --	1E+2 Bone surf (1E+2)	5E-8 --	-- 2E-10	1E-5 --
--	8E+1	3E-8	1E-10	--
2E+3 LLI wall (3E+3)	2E+3 --	9E-7 --	3E-9 --	-- 4E-5
--	2E+3	9E-7	3E-9	--
5E+4 St wall (6E+4)	2E+5 --	8E-5 --	3E-7 --	-- 8E-4
--	2E+5	7E-5	2E-7	--
4E+4 St wall (4E+4)	1E+5 --	5E-5 --	2E-7 --	-- 6E-4
--	1E+5	5E-5	2E-7	--
6E+3	2E+4	8E-6	3E-8	9E-5

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--	2E+4	6E-6	3E-8	--
3E+3	6E+3	2E-6	8E-9	4E-5
--	5E+3	2E-6	6E-9	--
1E+3 --	9E+0 Bone surf (2E+1)	4E-9 --	-- 3E-11	2E-5 --
-- --	4E+1 Bone surf (6E+1)	2E-8 --	-- 8E-11	-- --
5E+3	1E+4	5E-6	2E-8	7E-5
--	1E+4	5E-6	2E-8	--
3E+3 --	9E+2 Bone surf (1E+3)	4E-7 --	-- 1E-9	4E-5 --
--	1E+3	5E-7	2E-9	--
2E+4	6E+4	2E-5	8E-8	3E-4
--	9E+4	4E-5	1E-7	--
3E+2 --	1E+0 Bone surf (2E+0)	5E-10 --	-- 3E-12	3E-6 --
-- --	5E+0 Bone surf (9E+0)	2E-9 --	-- 1E-11	-- --
1E+3 --	3E+2 Bone surf (6E+2)	1E-7 --	-- 8E-10	1E-5 --
--	6E+2	3E-7	8E-10	--
7E+3	2E+4	9E-6	3E-8	1E-4
--	3E+4	1E-5	4E-8	--
1E+3 --	2E+2 Bone surf (4E+2)	7E-8 --	-- 6E-10	2E-5 --
--	4E+2	2E-7	6E-10	--
4E+4	9E+4	4E-5	1E-7	5E-4
--	1E+5	6E-5	2E-7	--
2E+2 Bone surf (4E+2)	8E-1 Bone surf (2E+0)	3E-10 --	-- 2E-12	-- 5E-6
-- --	3E+0 Bone surf (7E+0)	1E-9 --	-- 1E-11	-- --
2E+4	5E+4	2E-5	6E-8	3E-4
--	6E+4	2E-5	8E-8	--
2E+3	8E+3	3E-6	1E-8	3E-5
--	6E+3	3E-6	9E-9	--
4E+4	1E+5	5E-5	2E-7	5E-4
--	1E+5	4E-5	1E-7	--
7E+3	2E+4	8E-6	3E-8	9E-5
--	2E+4	7E-6	2E-8	--
3E+4	1E+5	4E-5	1E-7	4E-4
--	9E+4	4E-5	1E-7	--
6E+3	2E+4	7E-6	2E-8	8E-5
--	1E+4	6E-6	2E-8	--
4E+3	1E+4	5E-6	2E-8	5E-5
--	1E+4	5E-6	2E-8	--
1E+4	2E+4	8E-6	3E-8	2E-4
--	2E+4	7E-6	2E-8	--
2E+4	9E+4	4E-5	1E-7	2E-4
--	7E+4	3E-5	1E-7	--
2E+4	5E+3	2E-6	8E-9	3E-4
--	9E+2	4E-7	1E-9	--

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2E+4	7E+4	3E-5	9E-8	3E-4
--	6E+4	2E-5	8E-8	--
1E+3	4E+2	2E-7	6E-10	2E-5
--	2E+1	1E-8	3E-11	--
2E+5 St wall (2E+5)	5E+5 --	2E-4 --	8E-7 --	-- 3E-3
--	4E+5	2E-4	6E-7	--
8E+2	3E+2	1E-7	5E-10	1E-5
--	1E+2	6E-8	2E-10	--
9E+2 LLI wall (1E+3)	1E+3 --	5E-7 --	2E-9 --	-- 2E-5
--	1E+3	4E-7	1E-9	--
2E+3	5E+3	2E-6	8E-9	3E-5
--	5E+3	2E-6	7E-9	--
3E+4	7E+4	3E-5	1E-7	4E-4
--	6E+4	3E-5	9E-8	--
5E+4 St wall (7E+4)	2E+5 --	1E-4 --	3E-7 --	-- 1E-3
--	2E+5	9E-5	3E-7	--
1E+4	5E+4	2E-5	7E-8	1E-4
2E+4	9E+4	4E-5	1E-7	3E-4
5E+3	2E+4	8E-6	3E-8	7E-5
5E+5	2E+6	7E-4	2E-6	7E-3
2E+4	3E+4	1E-5	5E-8	2E-4
2E+3 LLI wall (3E+3)	7E+3 --	3E-6 --	9E-9 --	-- 4E-5
2E+3	9E+3	4E-6	1E-8	3E-5
4E+2 LLI wall (5E+2)	1E+3 --	5E-7 --	2E-9 --	-- 7E-6
9E+4	3E+5	1E-4	4E-7	--
St wall (1E+5)	--	--	--	2E-3
--	4E+5	1E-4	5E-7	--
7E+4 St wall (1E+5)	3E+5 --	1E-4 --	4E-7 --	-- 1E-3
--	3E+5	1E-4	4E-7	--
5E+3	9E+3	4E-6	1E-8	7E-5
--	9E+3	4E-6	1E-8	--
7E+3	1E+4	5E-6	2E-8	9E-5
--	2E+4	6E-6	2E-8	--
1E+3	2E+3	1E-6	3E-9	2E-5
--	2E+3	9E-7	3E-9	--
2E+3	3E+3	1E-6	4E-9	3E-5
--	4E+2	2E-7	6E-10	--
2E+3	4E+3	1E-6	5E-9	3E-5
--	1E+3	6E-7	2E-9	--
1E+3 St wall (2E+3)	2E+3 St wall (2E+3)	7E-7 --	-- 3E-9	-- 2E-5
--	2E+2	6E-8	2E-10	--
2E+3	3E+3	1E-6	4E-9	3E-5
--	2E+3	7E-7	2E-9	--
6E+5 St wall --	8E+5 (9E+5)	4E-4 --	-- 1E-6	8E-3 --
--	1E+5	4E-5	1E-7	--

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8E+4	1E+5	6E-5	2E-7	1E-3
--	1E+5	6E-5	2E-7	--
2E+3	3E+3	1E-6	4E-9	2E-5
--	3E+3	1E-6	4E-9	--
3E+3	5E+3	2E-6	7E-9	4E-5
--	4E+3	2E-6	6E-9	--
1E+5	4E+5	2E-4	5E-7	1E-3
--	5E+5	2E-4	7E-7	--
--	5E+5	2E-4	6E-7	--
1E+4	4E+4	2E-5	6E-8	2E-4
--	5E+4	2E-5	6E-8	--
--	4E+4	2E-5	6E-8	--
2E+3	6E+3	2E-6	8E-9	3E-5
--	4E+3	2E-6	6E-9	--
--	4E+3	2E-6	6E-9	--
2E+3	5E+2	2E-7	7E-10	3E-5
--	8E+2	3E-7	1E-9	--
--	8E+2	3E-7	1E-9	--
8E+4	2E+5	1E-4	3E-7	1E-3
--	2E+5	9E-5	3E-7	--
--	2E+5	7E-5	2E-7	--
1E+4	3E+4	1E-5	4E-8	2E-4
--	2E+4	8E-6	3E-8	--
--	2E+4	7E-6	2E-8	--
2E+3 LLI wall (3E+3)	2E+3 --	9E-7 --	3E-9 --	-- 3E-5
--	2E+3	7E-7	2E-9	--
--	1E+3	6E-7	2E-9	--
2E+3 LLI wall (2E+3)	5E+3 --	2E-6 --	6E-9 --	-- 2E-5
--	3E+3	1E-6	4E-9	--
--	3E+3	1E-6	4E-9	--
4E+2 LLI wall (6E+2)	4E+1 --	2E-8 --	6E-11 --	-- 8E-6
--	6E+1	2E-8	8E-11	--
--	8E+0	3E-9	1E-11	--
4E+4	1E+5	6E-5	2E-7	--
St wall (4E+4)	--	--	--	6E-4
--	2E+5	6E-5	2E-7	--
--	1E+5	5E-5	2E-7	--
8E+3	2E+4	1E-5	3E-8	1E-4
--	3E+4	1E-5	5E-8	--
--	3E+4	1E-5	4E-8	--
5E+3	1E+4	5E-6	2E-8	7E-5
--	1E+4	5E-6	2E-8	--
--	1E+4	4E-6	1E-8	--
2E+3	8E+3	3E-6	1E-8	3E-5
--	6E+3	3E-6	9E-9	--
--	6E+3	2E-6	8E-9	--
1E+4	3E+4	1E-5	5E-8	1E-4
--	3E+4	1E-5	4E-8	--

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  --                3E+4              1E-5        4E-8        --
  2E+3              5E+3              2E-6        6E-9        3E-5
  --                4E+3              1E-6        5E-9        --
  --                3E+3              1E-6        5E-9        --
  5E+3 LLI wall     5E+3 --           2E-6 --     7E-9 --     -- 7E-5
    (5E+3)
  --                4E+3              2E-6        5E-9        --
  --                4E+3              1E-6        5E-9        --
  2E+5              2E+5              8E-5        3E-7        2E-3
  --                2E+5              9E-5        3E-7        --
  --                2E+5              8E-5        3E-7        --
  1E+3              9E+2              4E-7        1E-9        1E-5
  --                1E+3              4E-7        1E-9        --
  --                9E+2              4E-7        1E-9        --
  3E+3              9E+1              4E-8        1E-10       4E-5
  --                2E+2              9E-8        3E-10       --
  --                2E+1              6E-9        2E-11       --
  9E+2              3E+2              1E-7        4E-10       1E-5
  --                4E+2              2E-7        6E-10       --
  --                2E+2              9E-8        3E-10       --
  6E+2              9E+1              4E-8        1E-10       9E-6
  --                2E+2              7E-8        2E-10       --
  --                1E+2              4E-8        1E-10       --
  1E+3              3E+3              1E-6        4E-9        1E-5
  --                2E+3              9E-7        3E-9        --
  --                2E+3              8E-7        3E-9        --
  8E+3              2E+4              1E-5        3E-8        1E-4
  --                3E+4              1E-5        4E-8        --
  --                2E+4              9E-6        3E-8        --
  1E+4              4E+4              2E-5        6E-8        2E-4
  --                5E+4              2E-5        7E-8        --
  --                4E+4              2E-5        6E-8        --
  1E+4              4E+4              2E-5        5E-8        2E-4
  2E+3              2E+3              7E-7        2E-9        2E-5
  1E+4              3E+4              1E-5        4E-8        1E-4
  4E+3              8E+3              4E-6        1E-8        5E-5
  3E+3 LLI wall     6E+3 --           3E-6 --     8E-9 --     -- 4E-5
    (3E+4)
  4E+4 LLI wall     2E+4 --           1E-5 --     3E-8 --     -- 6E-4
    (5E+4)
  2E+3 LLI wall     4E+3 --           2E-6 --     6E-9 --     -- 3E-5
    (2E+3)
  2E+4              4E+4              2E-5        6E-8        2E-4
  3E+3              1E+4              4E-6        1E-8        4E-5
  5E+4              1E+5              6E-5        2E-7        7E-4
  1E+3              3E+3              1E-6        5E-9        2E-5
  9E+3              3E+4              1E-5        4E-8        1E-4
  --                2E+4              9E-6        3E-8        --
  --                2E+4              8E-6        3E-8        --
80..+...90....+....0....+...10....+...20....+...30....+...40....+...50..

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******* This is piece 3. -- It begins at character 152 of table line 1. *******
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2E-3
-- 1E-4

6E-3
4E-2
3E-2
1E-2
7E-3
-- 1E-1

1E-2
-- 3E-2

9E-5
-- 1E-4

2E-3
2E-3
9E-3
-- 5E-4

5E-4
-- 2E-4

-- 1E-2

6E-4
-- 3E-4

-- 1E-4

-- 2E-3

-- 1E-4

6E-4
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1E-2
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2E-4
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2E-4
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3E-5 --

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4E-3
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152....60

TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

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79	Gold-194	D, see 1 9 3Au	3E+3
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-195	D, see 1 9 3Au	5E+3
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-198m	D, see 1 9 3Au	1E+3
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-198	D, see 1 9 3Au	1E+3
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-199	D, see 1 9 3Au	3E+3 LLI wall (3E+3)
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-200m	D, see 1 9 3Au	1E+3
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-200 2	D, see 1 9 3Au	3E+4
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
79	Gold-201 2	D, see 1 9 3Au	7E+4 St wall (9E+4)
		W, see 1 9 3Au	--
		Y, see 1 9 3Au	--
80	Mercury-193m	Vapor	--
		Organic D	4E+3
		D, sulfates	3E+3
		W, oxides, hydroxides, halides, nitrates, and sulfides	--
80	Mercury-193	Vapor	--
		Organic D	2E+4
		D, see 1 9 3mHg	2E+4
		W, see 1 9 3mHg	--
80	Mercury-194	Vapor	--
		Organic D	2E+1
		D, see 1 9 3mHg	8E+2
		W, see 1 9 3mHg	--
80	Mercury-195m	Vapor	--
		Organic D	3E+3
		D, see 1 9 3mHg	2E+3
		W, see 1 9 3mHg	--
80	Mercury-195	Vapor	--
		Organic D	2E+4
		D, see 1 9 3mHg	1E+4
		W, see 1 9 3mHg	--
80	Mercury-197m	Vapor	--
		Organic D	4E+3
		D, see 1 9 3mHg	3E+3
		W, see 1 9 3mHg	--
80	Mercury-197	Vapor	--
		Organic D	7E+3

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		D, see 1 9 3mHg	6E+3
		W, see 1 9 3mHg	--
80	Mercury-199m 2	Vapor	--
		Organic D	6E+4 St wall (1E+5)
		D, see 1 9 3mHg	6E+4
		W, see 1 9 3mHg	--
80	Mercury-203	Vapor	--
		Organic D	5E+2
		D, see 1 9 3mHg	2E+3
		W, see 1 9 3mHg	--
81	Thallium-194m 2	D, all compounds	5E+4 St wall (7E+4)
81	Thallium-194 2	D, all compounds	3E+5 St wall (3E+5)
81	Thallium-195 2	D, all compounds	6E+4
81	Thallium-197	D, all compounds	7E+4
81	Thallium-198m 2	D, all compounds	3E+4
81	Thallium-198	D, all compounds	2E+4
81	Thallium-199	D, all compounds	6E+4
81	Thallium-200	D, all compounds	8E+3
81	Thallium-201	D, all compounds	2E+4
81	Thallium-202	D, all compounds	4E+3
81	Thallium-204	D, all compounds	2E+3
82	Lead-195m 2	D, all compounds	6E+4
82	Lead-198	D, all compounds	3E+4
82	Lead-199 2	D, all compounds	2E+4
82	Lead-200	D, all compounds	3E+3
82	Lead-201	D, all compounds	7E+3
82	Lead-202m	D, all compounds	9E+3
82	Lead-202	D, all compounds	1E+2
82	Lead-203	D, all compounds	5E+3
82	Lead-205	D, all compounds	4E+3
82	Lead-209	D, all compounds	2E+4
82	Lead-210	D, all compounds	6E1 Bone surf (1E+0)
82	Lead-211 2	D, all compounds	1E+4
82	Lead-212	D, all compounds	8E+1 Bone surf (1E+2)
82	Lead-214 2	D, all compounds	9E+3
83	Bismuth-200 2	D, nitrates	3E+4
		W, all other compounds	--
83	Bismuth-201 2	D, see 2 0 0Bi	1E+4
		W, see 2 0 0Bi	--
83	Bismuth-202 2	D, see 2 0 0Bi	1E+4
		W, see 2 0 0Bi	--
83	Bismuth-203	D, see 2 0 0Bi	2E+3
		W, see 2 0 0Bi	--
83	Bismuth-205	D, see 2 0 0Bi	1E+3
		W, see 2 0 0Bi	--
83	Bismuth-206	D, see 2 0 0Bi	6E+2
		W, see 2 0 0Bi	--
83	Bismuth-207	D, see 2 0 0Bi	1E+3

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		W, see 2 0 0Bi	--
83	Bismuth-210m	D, see 2 0 0Bi	4E+1 Kidneys (6E+1)
		W, see 2 0 0Bi	--
83	Bismuth-210	D, see 2 0 0Bi	8E+2 --
		W, see 2 0 0Bi	--
83	Bismuth-212 2	D, see 2 0 0Bi	5E+3
		W, see 2 0 0Bi	--
83	Bismuth-213 2	D, see 2 0 0Bi	7E+3
		W, see 2 0 0Bi	--
83	Bismuth-214 2	D, see 2 0 0Bi	2E+4 St wall (2E+4)
		W, see 2 0 0Bi	--
84	Polonium-203 2	D, all compounds except those given for W	3E+4
		W, oxides, hydroxides, and nitrates	--
84	Polonium-205 2	D, see 2 0 3Po	2E+4
		W, see 2 0 3Po	--
84	Polonium-207	D, see 2 0 3Po	8E+3
		W, see 2 0 3Po	--
84	Polonium-210	D, see 2 0 3Po	3E+0
		W, see 2 0 3Po	--
85	Astatine-207 2	D, halides	6E+3
		W	--
85	Astatine-211	D, halides	1E+2
		W	--
86	Radon-220	With daughters removed	--
		With daughters present	--
86	Radon-222	With daughters removed	--
		With daughters present	--
87	Francium-222 2	D, all compounds	2E+3
87	Francium-223 2	D, all compounds	6E+2
88	Radium-223	W, all compounds	5E+0 Bone surf (9E+0)
88	Radium-224	W, all compounds	8E+0 Bone surf (2E+1)
88	Radium-225	W, all compounds	8E+0 Bone surf (2E+1)
88	Radium-226	W, all compounds	2E+0 Bone surf (5E+0)
88	Radium-227 2	W, all compounds	2E+4 Bone surf (2E+4)
88	Radium-228	W, all compounds	2E+0 Bone surf (4E+0)
89	Actinium-224	D, all compounds except those given for W and Y	2E+3 LLI wall (2E+3)
		W, halides and nitrates	--

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		Y, oxides and hydroxides	--
89	Actinium-225	D, see 2 2 4Ac	5E+1 LLI wall (5E+1)
		W, see 2 2 4Ac	--
		Y, see 2 2 4Ac	--
89	Actinium-226	D, see 2 2 4Ac	1E+2 LLI wall (1E+2)
		W, see 2 2 4Ac	--
		Y, see 2 2 4Ac	--
89	Actinium-227	D, see 2 2 4Ac	2E-1 Bone surf (4E-1)
		W, see 2 2 4Ac	-- --
		Y, see 2 2 4Ac	--
89	Actinium-228	D, see 2 2 4Ac	2E+3 --
		W, see 2 2 4Ac	-- --
		Y, see 2 2 4Ac	--
90	Thorium-226 2	W, all compounds except those given for Y	5E+3 St wall (5E+3)
		Y, oxides and hydroxides	--
90	Thorium-227	W, see 2 2 6Th	1E+2
		Y, see 2 2 6Th	--
90	Thorium-228	W, see 2 2 6Th	6E+0 Bone surf (1E+1)
		Y, see 2 2 6Th	--
90	Thorium-229	W, see 2 2 6Th	6E-1 Bone surf (1E+0)
		Y, see 2 2 6Th	-- --
90	Thorium-230	W, see 2 2 6Th	4E+0 Bone surf (9E+0)
		Y, see 2 2 6Th	-- --
90	Thorium-231	W, see 2 2 6Th	4E+3
		Y, see 2 2 6Th	--
90	Thorium-232	W, see 2 2 6Th	7E-1 Bone surf (2E+0)
		Y, see 2 2 6Th	-- --
90	Thorium-234	W, see 2 2 6Th	3E+2 LLI wall (4E+2)
		Y, see 2 2 6Th	--
91	Protactinium-2-27 2	W, all compounds except those given for Y	4E+3
		Y, oxides and hydroxides	--
91	Protactinium-2-28	W, see 2 2 7Pa	1E+3 --
		Y, see 2 2 7Pa	--
91	Protactinium-2-30	W, see 2 2 7Pa	6E+2 Bone surf (9E+2)
		Y, see 2 2 7Pa	--
91	Protactinium-2-31	W, see 2 2 7Pa	2E-1 Bone surf (5E-1)
		Y, see 2 2 7Pa	-- --
91	Protactinium-2-32	W, see 2 2 7Pa	1E+3 --

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		Y, see 2 2 7Pa	-- --
91	Protactinium-2-33	W, see 2 2 7Pa	1E+3 LLI wall (2E+3)
		Y, see 2 2 7Pa	--
91	Protactinium-2-34	W, see 2 2 7Pa	2E+3
		Y, see 2 2 7Pa	--
92	Uranium-230	D, UF, UOF, UO(NO)	4E+0 Bone surf (6E+0)
		W, UO, UF, UC1	--
		Y, UO, UO	--
92	Uranium-231	D, see 2 3 0U	5E+3 LLI wall (4E+3)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-232	D, see 2 3 0U	2E+0 Bone surf (4E+0)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-233	D, see 2 3 0U	1E+1 Bone surf (2E+1)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-234 3	D, see 2 3 0U	1E+1 Bone surf (2E+1)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-235 3	D, see 2 3 0U	1E+1 Bone surf (2E+1)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-236	D, see 2 3 0U	1E+1 Bone surf (2E+1)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-237	D, see 2 3 0U	2E+3 LLI wall (2E+3)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-238 3	D, see 2 3 0U	1E+1 Bone surf (2E+1)
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-239 2	D, see 2 3 0U	7E+4
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-240	D, see 2 3 0U	1E+3
		W, see 2 3 0U	--
		Y, see 2 3 0U	--
92	Uranium-natural 3	D, see 2 3 0U	1E+1 --
		W, see 2 3 0U	--
		Y, see 2 3 0U	--

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93   Neptunium-232 2  W, all compounds                        1E+5 --
93   Neptunium-233 2  W, all compounds                        8E+5
93   Neptunium-234    W, all compounds                        2E+3
93   Neptunium-235    W, all compounds                        2E+4 LLI wall
                                                               (2E+4)
93   Neptunium-236    W, all compounds                        3E+0 Bone surf
       (1.15E+5 y)                                             (6E+0)
93   Neptunium-236    W, all compounds                        3E+3 Bone surf
       (22.5 h)                                                (4E+3)
93   Neptunium-237    W, all compounds                        5E-1 Bone surf
                                                               (1E+0)
93   Neptunium-238    W, all compounds                        1E+3 --
93   Neptunium-239    W, all compounds                        2E+3 LLI wall
                                                               (2E+3)
93   Neptunium-240 2  W, all compounds                        2E+4
94   Plutonium-234    W, all compounds except PuO             8E+3
                      Y, see PuO                              --
94   Plutonium-235 2  W, see 2 3 4Pu                          9E+5
                      Y, see 2 3 4Pu                          --
94   Plutonium-236    W, see 2 3 4Pu                          2E+0 Bone surf
                                                               (4E+0)
                      Y, see 2 3 4Pu                          --
94   Plutonium-237    W, see 2 3 4Pu                          1E+4
                      Y, see 2 3 4Pu                          --
94   Plutonium-238    W, see 2 3 4Pu                          9E-1 Bone surf
                                                               (2E+0)
                      Y, see 2 3 4Pu                          --
94   Plutonium-239    W, see 2 3 4Pu                          8E-1 Bone surf
                                                               (1E+0)
                      Y, see 2 3 4Pu                          -- --
94   Plutonium-240    W, see 2 3 4Pu                          8E-1 Bone surf
                                                               (1E+0)
                      Y, see 2 3 4Pu                          -- --
94   Plutonium-241    W, see 2 3 4Pu                          4E+1 Bone surf
                                                               (7E+1)
                      Y, see 2 3 4Pu                          -- --
94   Plutonium-242    W, see 2 3 4Pu                          8E-1 Bone surf
                                                               (1E+0)
                      Y, see 2 3 4Pu                          -- --
94   Plutonium-243    W, see 2 3 4Pu                          2E+4
                      Y, see 2 3 4Pu                          --
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

*******************************************************************************
******* This is piece 2. -- It begins at character 80 of table line 1. ********
*******************************************************************************

  8E+3                    3E-6              1E-8              4E-5
  5E+3                    2E-6              8E-9              --
  5E+3                    2E-6              7E-9              --
  1E+4                    5E-6              2E-8              7E-5
  1E+3                    6E-7              2E-9              --
```

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4E+2	2E-7	6E-10	--
3E+3	1E-6	4E-9	1E-5
1E+3	5E-7	2E-9	--
1E+3	5E-7	2E-9	--
4E+3	2E-6	5E-9	2E-5
2E+3	8E-7	3E-9	--
2E+3	7E-7	2E-9	--
9E+3 --	4E-6 --	1E-8 --	-- 4E-5
4E+3	2E-6	6E-9	--
4E+3	2E-6	5E-9	--
4E+3	1E-6	5E-9	2E-5
3E+3	1E-6	4E-9	--
2E+4	1E-6	3E-9	--
6E+4	3E-5	9E-8	4E-4
8E+4	3E-5	1E-7	--
7E+4	3E-5	1E-7	--
2E+5 --	9E-5 --	3E-7 --	-- 1E-3
2E+5	1E-4	3E-7	--
2E+5	9E-5	3E-7	--
8E+3	4E-6	1E-8	--
1E+4	5E-6	2E-8	6E-5
9E+3	4E-6	1E-8	4E-5
8E+3	3E-6	1E-8	--
3E+4	1E-5	4E-8	--
6E+4	3E-5	9E-8	3E-4
4E+4	2E-5	6E-8	2E-4
4E+4	2E-5	6E-8	--
3E+1	1E-8	4E-11	--
3E+1	1E-8	4E-11	2E-7
4E+1	2E-8	6E-11	1E-5
1E+2	5E-8	2E-10	--
4E+3	2E-6	6E-9	--
6E+3	3E-6	8E-9	4E-5
5E+3	2E-6	7E-9	3E-5
4E+3	2E-6	5E-9	--
3E+4	1E-5	4E-8	--
5E+4	2E-5	6E-8	2E-4
4E+4	1E-5	5E-8	2E-4
3E+4	1E-5	5E-8	--
5E+3	2E-6	7E-9	--
9E+3	4E-6	1E-8	5E-5
7E+3	3E-6	1E-8	4E-5
5E+3	2E-6	7E-9	--
8E+3	4E-6	1E-8	--
1E+4	6E-6	2E-8	9E-5
1E+4	5E-6	2E-8	8E-5
9E+3	4E-6	1E-8	--
8E+4	3E-5	1E-7	--
2E+5 --	7E-5 --	2E-7 --	-- 1E-3

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1E+5	6E-5	2E-7	8E-4
2E+5	7E-5	2E-7	--
8E+2	4E-7	1E-9	--
8E+2	3E-7	1E-9	7E-6
1E+3	5E-7	2E-9	3E-5
1E+3	5E-7	2E-9	--
2E+5 --	6E-5 --	2E-7 --	-- 1E-3
6E+5 --	2E-4 --	8E-7 --	-- 4E-3
1E+5	5E-5	2E-7	9E-4
1E+5	5E-5	2E-7	1E-3
5E+4	2E-5	8E-8	4E-4
3E+4	1E-5	5E-8	3E-4
8E+4	4E-5	1E-7	9E-4
1E+4	5E-6	2E-8	1E-4
2E+4	9E-6	3E-8	2E-4
5E+3	2E-6	7E-9	5E-5
2E+3	9E-7	3E-9	2E-5
2E+5	8E-5	3E-7	8E-4
6E+4	3E-5	9E-8	4E-4
7E+4	3E-5	1E-7	3E-4
6E+3	3E-6	9E-9	4E-5
2E+4	8E-6	3E-8	1E-4
3E+4	1E-5	4E-8	1E-4
5E+1	2E-8	7E-11	2E-6
9E+3	4E-6	1E-8	7E-5
1E+3	6E-7	2E-9	5E-5
6E+4	2E-5	8E-8	3E-4
2E1 Bone surf (4E-1)	1E-10 --	-- 6E-13	-- 1E-8
6E+2	3E-7	9E-10	2E-4
3E+1 --	1E-8 --	5E-11 --	-- 2E-6
8E+2	3E-7	1E-9	1E-4
8E+4	4E-5	1E-7	4E-4
1E+5	4E-5	1E-7	--
3E+4	1E-5	4E-8	2E-4
4E+4	2E-5	5E-8	--
4E+4	2E-5	6E-8	2E-4
8E+4	3E-5	1E-7	--
7E+3	3E-6	9E-9	3E-5
6E+3	3E-6	9E-9	--
3E+3	1E-6	3E-9	2E-5
1E+3	5E-7	2E-9	--
1E+3	6E-7	2E-9	9E-6
9E+2	4E-7	1E-9	--
2E+3	7E-7	2E-9	1E-5
4E+2	1E-7	5E-10	--
5E+0 Kidneys (6E+0)	2E-9 --	-- 9E-12	-- 8E-7
7E-1	3E-10	9E-13	--
2E+2 Kidneys (4E+2)	1E-7 --	-- 5E-10	1E-5 --

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3E+1	1E-8	4E-11	--
2E+2	1E-7	3E-10	7E-5
3E+2	1E-7	4E-10	--
3E+2	1E-7	4E-10	1E-4
4E+2	1E-7	5E-10	--
8E+2 --	3E-7 --	1E-9 --	-- 3E-4
9E-2	4E-7	1E-9	--
6E+4	3E-5	9E-8	3E-4
9E+4	4E-5	1E-7	--
4E+4	2E-5	5E-8	3E-4
7E+4	3E-5	1E-7	--
3E+4	1E-5	3E-8	1E-4
3E+4	1E-5	4E-8	--
6E-1	3E-10	9E-13	4E-8
6E-1	3E-10	9E-13	--
3E+3	1E-6	4E-9	8E-5
2E+3	9E-7	3E-9	--
8E+1	3E-8	1E-10	2E-6
5E+1	2E-8	8E-11	--
2E+4	7E-6	2E-8	--
2E+1 (or 12 working level months)	9E-9	3E-11 (or 1.0 working level)	--
1E+4	4E-6	1E-8	--
1E+2 (or 4 working level months)	3E-8 (or 0.33 working level)	1E-10	--
5E+2	2E-7	6E-10	3E-5
8E+2	3E-7	1E-9	8E-6
7E+1 --	3E-10 --	9E-13 --	-- 1E-7
2E+0 --	7E-10 --	2E-12 --	-- 2E-7
7E-1 --	3E-10 --	9E-13 --	-- 2E-7
6E-1 --	3E-10 --	9E-13 --	-- 6E-8
1E+4 Bone surf (2E+4)	6E-6 --	-- 3E-8	-- 3E-4
1E+0 --	5E-10 --	2E-12 --	-- 6E-8
3E+1	1E-8	--	--
Bone surf (4E+1)	--	5E-11	3E-5
5E+1	2E-8	7E-11	--
5E+1	2E-8	6E-11	--
3E-1 Bone surf (5E-1)	1E-10 --	-- 7E-13	-- 7E-7
6E-1	3E-10	9E-13	--
6E-1	3E-10	9E-13	--

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3E+0 Bone surf (4E+0)	1E-9 --	-- 5E-12	-- 2E-6
5E+0	2E-9	7E-12	--
5E+0	2E-9	6E-12	--
4E-4 Bone surf (8E-4)	2E-13 --	-- 1E-15	-- 5E-9
2E-3 Bone surf (3E-3)	7E-13 --	-- 4E-15	-- --
4E-3	2E-12	6E-15	--
9E+0 Bone surf (2E+1)	4E-9 --	-- 2E-11	3E-5 --
4E+1 Bone surf (6E+1)	2E-8 --	-- 8E-11	-- --
4E+1	2E-8	6E-11	--
2E+2	6E-8	2E-10	--
--	--	--	7E-5
1E+2	6E-8	2E-10	--
3E-1	1E-10	5E-13	2E-6
3E-1	1E-10	5E-13	--
1E-2 Bone surf (2E-2)	4E-12 --	-- 3E-14	-- 2E-7
2E-2	7E-12	2E-14	--
9E-4 Bone surf (2E-3)	4E-13 --	-- 3E-15	-- 2E-8
2E-3 Bone surf (3E-3)	1E-12 --	-- 4E-15	-- --
6E-3 Bone surf (2E-2)	3E-12 --	-- 2E-14	-- 1E-7
2E-2 Bone surf (2E-2)	6E-12 --	-- 3E-14	-- --
6E+3	3E-6	9E-9	5E-5
6E+3	3E-6	9E-9	--
1E-3 Bone surf (3E-3)	5E-13 --	-- 4E-15	-- 3E-8
3E-3 Bone surf (4E-3)	1E-12 --	-- 6E-15	-- --
2E+2 --	8E-8 --	3E-10 --	-- 5E-6
2E+2	6E-8	2E-10	--
1E+2	5E-8	2E-10	5E-5
1E+2	4E-8	1E-10	--
1E+1 Bone surf (2E+1)	5E-9 --	-- 3E-11	2E-5 --
1E+1	5E-9	2E-11	--
5E+0 --	2E-9 --	7E-12 --	-- 1E-5
4E+0	1E-9	5E-12	--
2E-3 Bone surf (4E-3)	6E-13 --	-- 6E-15	-- 6E-9
4E-3 Bone surf (6E-3)	2E-12 --	-- 8E-15	-- --
2E+1 Bone surf (6E+1)	9E-9 --	-- 8E-11	2E-5 --
6E+1 Bone surf (7E+1)	2E-8 --	-- 1E-10	-- --
7E+2 --	3E-7 --	1E-9 --	-- 2E-5
6E+2	2E-7	8E-10	--
8E+3	3E-6	1E-8	3E-5

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7E+3	3E-6	9E-9	--
4E-1 Bone surf (6E-1)	2E-10 --	-- 8E-13	-- 8E-8
4E-1	1E-10	5E-13	--
3E-1	1E-10	4E-13	--
8E+3 --	3E-6 --	1E-8 --	-- 6E-5
6E+3	2E-6	8E-9	--
5E+3	2E-6	6E-9	--
2E-1 Bone surf (4E-1)	9E-11 --	-- 6E-13	-- 6E-8
4E-1	2E-10	5E-13	--
8E-3	3E-12	1E-14	--
1E+0 Bone surf (2E+0)	5E-10 --	-- 3E-12	-- 3E-7
7E-1	3E-10	1E-12	--
4E-2	2E-11	5E-14	--
1E+0 Bone surf (2E+0)	5E-10 --	-- 3E-12	-- 3E-7
7E-1	3E-10	1E-12	--
4E-2	2E-11	5E-14	--
1E+0 Bone surf (2E+0)	6E-10 --	-- 3E-12	-- 3E-7
8E-1	3E-10	1E-12	--
4E-2	2E-11	6E-14	--
1E+0 Bone surf (2E+0)	5E-10 --	-- 3E-12	-- 3E-7
8E-1	3E-10	1E-12	--
4E-2	2E-11	6E-14	--
3E+3 --	1E-6 --	4E-9 --	-- 3E-5
2E+3	7E-7	2E-9	--
2E+3	6E-7	2E-9	--
1E+0 Bone surf (2E+0)	6E-10 --	-- 3E-12	-- 3E-7
8E-1	3E-10	1E-12	--
4E-2	2E-11	6E-14	--
2E+5	8E-5	3E-7	9E-4
2E+5	7E-5	2E-7	--
2E+5	6E-5	2E-7	--
4E+3	2E-6	5E-9	2E-5
3E+3	1E-6	4E-9	--
2E+3	1E-6	3E-9	--
1E+0 Bone surf (2E+1)	5E-10 Bone surf (2E+0)	-- 3E-12	-- 3E-7
8E-1	3E-10	9E-13	--
5E-2	2E-11	9E-14	--
2E+3 Bone surf (5E+2)	7E-7 --	-- 6E-9	2E-3 --
3E+6	1E-3	4E-6	1E-2
3E+3	1E-6	4E-9	3E-5
8E+2 Bone surf (1E+3)	3E-7 --	-- 2E-9	-- 3E-4

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```
  2E-2 Bone surf (5E-2)   9E-12 --          -- 8E-14          -- 9E-8

  3E+1 Bone surf (7E+1)   1E-8 --           -- 1E-10          -- 5E-5

  4E-3 Bone surf (1E-2)   2E-12 --          -- 1E-14          -- 2E-8

  6E+1 Bone surf (2E+2)   3E-8 --           -- 2E-10          2E-5 --
  2E+3 --                 9E-7 --           3E-9 --           -- 2E-5

  8E+4                    3E-5              1E-7              3E-4
  2E+2                    9E-8              3E-10             1E-4
  2E+2                    8E-8              3E-10             --
  3E+6                    1E-3              4E-6              1E-2
  3E+6                    1E-3              3E-6              --
  2E-2 Bone surf (4E-2)   8E-12 --          -- 5E-14          -- 6E-8

  4E-2                    2E-11             6E-14             --
  3E+3                    1E-6              5E-9              2E-4
  3E+3                    1E-6              4E-9              --
  7E-3 Bone surf (1E-2)   3E-12 --          -- 2E-14          -- 2E-8

  2E-2                    8E-12             2E-14             --
  6E-3 Bone surf (1E-2)   3E-12 --          -- 2E-14          -- 2E-8

  2E-2 Bone surf (2E-2)   7E-12 --          -- 2E-14          -- --
  6E-3 Bone surf (1E-2)   3E-12 --          -- 2E-14          -- 2E-8

  2E-2 Bone surf (2E-2)   7E-12 --          -- 2E-14          -- --
  3E-1 Bone surf (6E-1)   1E-10 --          -- 8E-13          -- 1E-6

  8E-1 Bone surf (1E+0)   3E-10 --          -- 1E-12          -- --
  7E-3 Bone surf (1E-2)   3E-12 --          -- 2E-14          -- 2E-8

  2E-2 Bone surf (2E-2)   7E-12 --          -- 2E-14          -- --
  4E+4                    2E-5              5E-8              2E-4
  4E+4                    2E-5              5E-8              --
80..+...90....+....0....+...10....+...20....+...30....+...40....+...50..

*******************************************************************************
******* This is piece 3. -- It begins at character 152 of table line 1. *******
*******************************************************************************

 4E-4
 --
 --
 7E-4
 --
 --
 1E-4
 --
 --
 2E-4
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152....60
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******************************************************************************
******** This is piece 1. -- It begins at character 1 of table line 1. ********
******************************************************************************

94    Plutonium-244                W, see 2 3 4Pu            8E-1 Bone surf
                                                                 (2E+0)
                                   Y, see 2 3 4Pu            -- --
94    Plutonium-245                W, see 2 3 4Pu            2E+3
```

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		Y, see 2 3 4Pu	--
94	Plutonium-246	W, see 2 3 4Pu	4E+2 LLI wall (4E+2)
		Y, see 2 3 4Pu	--
95	Americium-237 2	W, all compounds	8E+4
95	Americium-238 2	W, all compounds	4E+4 --
95	Americium-239	W, all compounds	5E+3
95	Americium-240	W, all compounds	2E+3
95	Americium-241	W, all compounds	8E-1 Bone surf (1E+0)
95	Americium-242m	W, all compounds	8E-1 Bone surf (1E+0)
95	Americium-242	W, all compounds	4E+3 --
95	Americium-243	W, all compounds	8E-1 Bone surf (1E+0)
95	Americium-244m 2	W, all compounds	6E+4 St wall (8E+4)
95	Americium-244	W, all compounds	3E+3 --
95	Americium-245	W, all compounds	3E+4
95	Americium-246m 2	W, all compounds	5E+4 St wall (6E+4)
95	Americium-246 2	W, all compounds	3E+4
96	Curium-238	W, all compounds	2E+4
96	Curium-240	W, all compounds	6E+1 Bone surf (8E+1)
96	Curium-241	W, all compounds	1E+3 --
96	Curium-242	W, all compounds	3E+1 Bone surf (5E+1)
96	Curium-243	W, all compounds	1E+0 Bone surf (2E+0)
96	Curium-244	W, all compounds	1E+0 Bone surf (3E+0)
96	Curium-245	W, all compounds	7E-1 Bone surf (1E+0)
96	Curium-246	W, all compounds	7E-1 Bone surf (1E+0)
96	Curium-247	W, all compounds	8E-1 Bone surf (1E+0)
96	Curium-248	W, all compounds	2E-1 Bone surf (4E-1)
96	Curium-249 2	W, all compounds	5E+4 --
96	Curium-250	W, all compounds	4E-2 Bone surf (6E-2)
97	Berkelium-245	W, all compounds	2E+3
97	Berkelium-246	W, all compounds	3E+3
97	Berkelium-247	W, all compounds	5E-1 Bone surf (1E+0)
97	Berkelium-249	W, all compounds	2E+2 Bone surf (5E+2)
97	Berkelium-250	W, all compounds	9E+3 --
98	Californium-244 2	W, all compounds except those given for Y	3E+4 St wall (3E+4)

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		Y, oxides and hydroxides	--
98	Californium-246	W, see 2 4 4Cf	4E+2
		Y, see 2 4 4Cf	--
98	Californium-248	W, see 2 4 4Cf	8E+0 Bone surf (2E+1)
		Y, see 2 4 4Cf	--
98	Californium-249	W, see 2 4 4Cf	5E-1 Bone surf (1E+0)
		Y, see 2 4 4Cf	-- --
98	Californium-250	W, see 2 4 4Cf	1E+0 Bone surf (2E+0)
		Y, see 2 4 4Cf	--
98	Californium-251	W, see 2 4 4Cf	5E-1 Bone surf (1E+0)
		Y, see 2 4 4Cf	-- --
98	Californium-252	W, see 2 4 4Cf	2E+0 Bone surf (5E+0)
		Y, see 2 4 4Cf	--
98	Californium-253	W, see 2 4 4Cf	2E+2 Bone surf (4E+2)
		Y, see 2 4 4Cf	--
98	Californium-254	W, see 2 4 4Cf	2E+0
		Y, see 2 4 4Cf	--
99	Einsteinium-250	W, all compounds	4E+4 --
99	Einsteinium-251	W, all compounds	7E+3 --
99	Einsteinium-253	W, all compounds	2E+2
99	Einsteinium-254m	W, all compounds	3E+2 LLI wall (3E+2)
99	Einsteinium-254	W, all compounds	8E+0 Bone surf (2E+1)
100	Fermium-252	W, all compounds	5E+2
100	Fermium-253	W, all compounds	1E+3
100	Fermium-254	W, all compounds	3E+3
100	Fermium-255	W, all compounds	5E+2
100	Fermium-257	W, all compounds	2E+1 Bone surf (4E+1)
101	Mendelevium-257	W, all compounds	7E+3 --
101	Mendelevium-258	W, all compounds	3E+1 Bone surf (5E+1)

-- Any single radionuclide not listed above with decay mode other than alpha emission or spontaneous fission and with radioactive half-life less than 2 hours Submersion 1

-- Any single radionuclide not listed above with decay mode other than alpha emission or spontaneous fission and with radioactive half-life

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```
        greater than 2 hours
--    Any single radionuclide not
        listed above that decays
        by alpha emission or
        spontaneous fission, or
        any mixture for which
        either the identity or
        the concentration of any
        radionuclide in the
        mixture is not known
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

*******************************************************************************
******* This is piece 2. -- It begins at character 80 of table line 1. ********
*******************************************************************************

  7E-3 Bone surf (1E-2)   3E-12 --    -- 2E-14    -- 2E-8     -- 2E-7

  2E-2 Bone surf (2E-2)   7E-12 --    -- 2E-14    -- --       -- --
  5E+3                    2E-6        6E-9        3E-5        3E-4
  4E+3                    2E-6        6E-9        --          --
  3E+2 --                 1E-7 --     4E-10 --    -- 6E-6     -- 6E-5

  3E+2                    1E-7        4E-10       --          --
  3E+5                    1E-4        4E-7        1E-3        1E-2
  3E+3 Bone surf (6E+3)   1E-6 --     -- 9E-9     5E-4 --     5E-3 --
  1E+4                    5E-6        2E-8        7E-5        7E-4
  3E+3                    1E-6        4E-9        3E-5        3E-4
  6E-3 Bone surf (1E-2)   3E-12 --    -- 2E-14    -- 2E-8     -- 2E-7

  6E-3 Bone surf (1E-2)   3E-12 --    -- 2E-14    -- 2E-8     -- 2E-7

  8E+1 Bone surf (9E+1)   4E-8 --     -- 1E-10    5E-5 --     5E-4 --
  6E-3 Bone surf (1E-2)   3E-12 --    -- 2E-14    -- 2E-8     -- 2E-7

  4E+3 Bone surf (7E+3)   2E-6 --     -- 1E-8     -- 1E-3     -- 1E-2

  2E+2 Bone surf (3E+2)   8E-8 --     -- 4E-10    4E-5 --     4E-4 --
  8E+4                    3E-5        1E-7        4E-4        4E-3
  2E+5 --                 8E-5 --     3E-7 --     -- 8E-4     -- 8E-3

  1E+5                    4E-5        1E-7        4E-4        4E-3
  1E+3                    5E-7        2E-9        2E-4        2E-3
  6E-1 Bone surf (6E-1)   2E-10 --    -- 9E-13    -- 1E-6     -- 1E-5

  3E+1 Bone surf (4E+1)   1E-8 --     -- 5E-11    2E-5 --     2E-4 --
  3E-1 Bone surf (3E-1)   1E-10 --    -- 4E-13    -- 7E-7     -- 7E-6

  9E-3 Bone surf (2E-2)   4E-12 --    -- 2E-14    -- 3E-8     -- 3E-7

  1E-2 Bone surf (2E-2)   5E-12 --    -- 3E-14    -- 3E-8     -- 3E-7
```

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6E-3 Bone surf (1E-2)	3E-12 --	-- 2E-14	-- 2E-8	-- 2E-7
6E-3 Bone surf (1E-2)	3E-12 --	-- 2E-14	-- 2E-8	-- 2E-7
6E-3 Bone surf (1E-2)	3E-12 --	-- 2E-14	-- 2E-8	-- 2E-7
2E-3 Bone surf (3E-3)	7E-13 --	-- 4E-15	-- 5E-9	-- 5E-8
2E+4 Bone surf (3E+4)	7E-6 --	-- 4E-8	7E-4 --	7E-3 --
3E-4 Bone surf (5E-4)	1E-13 --	-- 8E-16	-- 9E-10	-- 9E-9
1E+3	5E-7	2E-9	3E-5	3E-4
3E+3	1E-6	4E-9	4E-5	4E-4
4E-3 Bone surf (9E-3)	2E-12 --	-- 1E-14	-- 2E-8	-- 2E-7
2E+0 Bone surf (4E+0)	7E-10 --	-- 5E-12	-- 6E-6	-- 6E-5
3E+2 Bone surf (7E+2)	1E-7 --	-- 1E-9	1E-4 --	1E-3 --
6E+2	2E-7	8E-10	--	--
--	--	--	4E-4	4E-3
6E+2	2E-7	8E-10	--	--
9E+0	4E-9	1E-11	5E-6	5E-5
9E+0	4E-9	1E-11	--	--
6E-2 Bone surf (1E-1)	3E-11 --	-- 2E-13	-- 2E-7	-- 2E-6
1E-1	4E-11	1E-13	--	--
4E-3 Bone surf (9E-3)	2E-12 --	-- 1E-14	-- 2E-8	-- 2E-7
1E-2 Bone surf (1E-2)	4E-12 --	-- 2E-14	-- --	-- --
9E-3 Bone surf (2E-2)	4E-12 --	-- 3E-14	-- 3E-8	-- 3E-7
3E-2	1E-11	4E-14	--	--
4E-3 Bone surf (9E-3)	2E-12 --	-- 1E-14	-- 2E-8	-- 2E-7
1E-2 Bone surf (1E-2)	4E-12 --	-- 2E-14	-- --	-- --
2E-2 Bone surf (4E-2)	8E-12 --	-- 5E-14	-- 7E-8	-- 7E-7
3E-2	1E-11	5E-14	--	--
2E+0 --	8E-10 --	3E-12 --	-- 5E-6	-- 5E-5
2E+0	7E-10	2E-12	--	--
2E-2	9E-12	3E-14	3E-8	3E-7
2E-2	7E-12	2E-14	--	--
5E+2 Bone surf (1E+3)	2E-7 --	-- 2E-9	6E-4 --	6E-3 --
9E+2 Bone surf (1E+3)	4E-7 --	-- 2E-9	1E-4 --	1E-3 --
1E+0	6E-10	2E-12	2E-6	2E-5
1E+1 --	4E-9 --	1E-11 --	-- 4E-6	-- 4E-5
7E-2 Bone surf (1E-1)	3E-11 --	-- 2E-13	-- 2E-7	-- 2E-6
1E+1	5E-9	2E-11	6E-6	6E-5
1E+1	4E-9	1E-11	1E-5	1E-4

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9E+1	4E-8	1E-10	4E-5	4E-4	
2E+1	9E-9	3E-11	7E-6	7E-5	
2E-1 Bone surf (2E-1)	7E-11 --	-- 3E-13	-- 5E-7	-- 5E-6	
8E+1 Bone surf (9E+1)	4E-8 --	-- 1E-10	1E-4 --	1E-3 --	
2E-1 Bone surf (3E-1)	1E-10 --	-- 5E-13	-- 6E-7	-- 6E-6	
--	2E+2	1E-7	1E-9	--	--
--	2E-1	1E-10	1E-12	1E-8	1E-7
--	4E-4	2E-13	1E-15	2E-9	2E-8

80..+...90....+....0....+...10....+...20....+...30....+...40....+...50....+.

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Notes:

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

Table 1 Occupational Values

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_____
Atomic No. Radionuclide
Oral Ingestion Inhalation ____ ALI (mCi)

If it is known that Ac-227-D and Cm-250-W are not present
If, in addition, it is known that Ac-227-W, Y, Th-229-W, Y, Th-230-W, Th-232-W, Y, Pa-231-W, Y, Np-237-W, Pu-239-W, Pu-240-W, Pu-242-W, Am-241-W, Am-242m-W, Am-243-W, Cm-245-W, Cm-246-W, Cm-247-W, Cm-248-W, Bk-247-W, Cf-249-W, and Cf-251-W are not present
If, in addition, it is known that Sm-146-W, Sm-147-W, Gd-148-D, W, Gd-152-D, W, Th-228-W, Y, Th-230-Y, U-232-Y, U-233-Y, U-234-Y, U-235-Y, U-236-Y, U-238-Y, Np-236-W, Pu-236-W, Y, Pu-238-W, Y, Pu-239-Y, Pu-240-Y, Pu-242-Y, Pu-244-W, Y, Cm-243-W, Cm-244-W, Cf-248-W, Cf-249-Y, Cf-250-W, Y, Cf-251-Y, Cf-252-W, Y, and Cf-254-W, Y are not present
If, in addition, it is known that Pb-210-D, Bi-210m-W, Po-210-D, W, Ra-223-W, Ra-225-W, Ra-226-W, Ac-225-D, W, Y, Th-227-W, Y, U-230-D, W, Y, U-232-D, W, Pu-241-W, Cm-240-W, Cm-242-W, Cf-248-Y, Es-254-W, Fm-257-W, and Md-258-W are not present
If, in addition, it is known that Si-32-Y, Ti-44-Y, Fe-60-D, Sr-90-Y, Zr-93-D, Cd-113m-D, Cd-113-D, In-115-D, W, La-138-D, Lu-176-W, Hf-178m-D, W, Hf-182-D, W, Bi-210m-D, Ra-224-W, Ra-228-W, Ac-226-D, W, Y, Pa-230-W, Y, U-233-D, W, U-234-D, W, U-235-D, W, U-236-D, W, U-238-D, W, Pu-241-Y, Bk-249-W, Cf-253-W, Y, and Es-253-W are not present
If it is known that Ac-227-D, W, Y, Th-229-W, Y, Th-232-W, Y, Pa-231-W, Y, Cm-248-W, and, Cm-250-W are not present
If, in addition, it is known that Sm-146-W, Gd-148-D, W, Gd-152-D, Th-228-W, Y, Th-230-W, Y, U-232-Y, U-233-Y, U-234-Y, U-235-Y, U-236-Y, U-238-Y, U-Nat-Y, Np-236-W, Np-237-W, Pu-236-W, Y, Pu-238-W, Y, Pu-239-W, Y, Pu-240-W, Y, Pu-242-W, Y, Pu-244-W, Y, Am-241-W, Am-242m-W, Am-243-W, Cm-243-W, Cm-244-W, Cm-245-W, Cm-246-W, Cm-247-W, Bk-247-W, Cf-249-W, Y, Cf-250-W, Y, Cf-251-W, Y, Cf-252-W, Y, and Cf-254-W, Y are not present
If, in addition, it is known that Sm-147-W, Gd-152-W, Pb-210-D, Bi-210m-W, Po-210-D, W, Ra-223-W, Ra-225-W, Ra-226-W, Ac-225-D, W, Y, Th-227-W, Y, U-230-D, W, Y, U-232-D, W, U-Nat-W, Pu-241-W, Cm-240-W, Cm-242-W, Cf-248-W, Y, Es-254-W, Fm-257-W, and Md-258-W are not present
If, in addition, it is known that Fe-60, Sr-90, Cd-113m, Cd-113, In-115, I-129, Cs-134, Sm-145, Sm-147, Gd-148, Gd-152, Hg-194 (organic), Bi-210m, Ra-223, Ra-224, Ra-225, Ac-225, Th-228,

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  Th-230, U-233, U-234, U-235, U-236, U-238, U-Nat, Cm-242,
  Cf-248, Es-254, Fm-257, and Md-258 are not present
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

*******************************************************************************
******* This is piece 2. -- It begins at character 80 of table line 1. ********
*******************************************************************************
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		Table 2 Effluent Concentrations			Table 3 Releases to Sewers	
	Col. 1	Col. 2	Col. 3	Col. 1	Col. 2	
Class ALI (mCi)	DAC (-mCi-/ml)	Air (mCi/ml)	Water (mC-i/m-l)	Monthly Average Concen-tration (mCi/m-l)		
		--	7E-4	3E-13	--	--
		--	7E-3	3E-12	--	--
		--	7E-2	3E-11	--	--
		--	7E-1	3E-10	--	--
		--	7E+0	3E-9	--	--
		--	--	--	1E-14	--
		--	--	--	1E-13	--

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                    --                  --      --         1E-12       --


                    --                  --      --         --          1E-6


80..+...90....+....0....+...10....+...20....+...30....+...40....+...50....+.

*******************************************************************************
******* This is piece 3. -- It begins at character 156 of table line 1. *******
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1E-5

156..

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3. If a mixture of radionuclides consists of uranium and its daughters in ore dust (10 mm AMAD particle distribution assumed) prior to chemical separation of the uranium from the ore, the following values may be used for the DAC of the mixture: 6E-11 mCi of gross alpha activity from uranium-238, uranium-234, thorium-230, and radium-226 per milliliter of air; 3E-11 mCi of natural uranium per milliliter of air; or 45 micrograms of natural uranium per cubic meter of air. 4. If the identity and concentration of each radionuclide in a mixture are known, the limiting values should be derived as follows: determine, for each radionuclide in the mixture, the ratio between the concentration present in the mixture and the concentration otherwise established in Appendix B for the specific radionuclide when not in a mixture. The sum of such ratios for all of the radionuclides in the mixture may not exceed "1"' (i.e., "unity"').

Example: If radionuclides "A,"' "B,"' and "C"' are present in concentrations CA, CB, and CC, and if the applicable DACs are DACA, DACB, and DACC, respectively, then the concentrations shall be limited so that the following relationship exists:

CA + CB + CC # 1
DACA DACB DACC

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APPENDIX C QUANTITIES* OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

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Radionuclide Quantity (mCi)** Radionuclide Quantity (mCi)**

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Radionuclide Quantity (mCi)** Radionuclide Quantity (mCi)**

(cont'd)

QUANTITIES* OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

Radionuclide	Quantity (mCi)**	Radionuclide	Quantity (mCi)**
Hydrogen-3	1,000	Chromium-48	1,000
Beryllium-7	1,000	Chromium-49	1,000
Beryllium-10	1	Chromium-51	1,000
Carbon-11	1,000	Manganese-51	1,000
Carbon-14	1,000	Manganese-52m	1,000
Fluorine-18	1,000	Manganese-52	100
Sodium-22	10	Manganese-53	1,000
Sodium-24	100	Manganese-54	100
Magnesium-28	100	Manganese-56	1,000
Aluminum-26	10	Iron-52	100
Silicon-31	1,000	Iron-55	100
Silicon-32	1	Iron-59	10
Phosphorus-32	10	Iron-60	1
Phosphorus-33	100	Cobalt-55	100
Sulfur-35	100	Cobalt-56	10
Chlorine-36	10	Cobalt-57	100
Chlorine-38	1,000	Cobalt-58m	1,000
Chlorine-39	1,000	Cobalt-58	100
Argon-39	1,000	Cobalt-60m	1,000
Argon-41	1,000	Cobalt-60	1
Potassium-40	100	Cobalt-61	1,000
Potassium-42	1,000	Cobalt-62m	1,000
Potassium-43	1,000	Nickel-56	100
Potassium-44	1,000	Nickel-57	100
Potassium-45	1,000	Nickel-59	100
Calcium-41	100	Nickel-63	100
Calcium-45	100	Nickel-65	1,000
Calcium-47	100	Nickel-66	10
Scandium-43	1,000	Copper-60	1,000
Scandium-44m	100	Copper-61	1,000
Scandium-44	100	Copper-64	1,000
Scandium-46	10	Copper-67	1,000
Scandium-47	100	Zinc-62	100
Scandium-48	100	Zinc-63	1,000
Scandium-49	1,000	Zinc-65	10
Titanium-44	1	Zinc-69m	100
Titanium-45	1,000	Zinc-69	1,000
Vanadium-47	1,000	Zinc-71m	1,000
Vanadium-48	100	Zinc-72	100
Vanadium-49	1,000	Gallium-65	1,000

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Gallium-66	100	Krypton-81	1,000
Gallium-67	1,000	Krypton-83m	1,000
Gallium-68	1,000	Krypton-85m	1,000
Gallium-70	1,000	Krypton-85	1,000
Gallium-72	100	Krypton-87	1,000
Gallium-73	1,000	Krypton-88	1,000
Germanium-66	1,000	Rubidium-79	1,000
Germanium-67	1,000	Rubidium-81m	1,000
Germanium-68	10	Rubidium-81	1,000
Germanium-69	1,000	Rubidium-82m	1,000
Germanium-71	1,000	Rubidium-83	100
Germanium-75	1,000	Rubidium-84	100
Germanium-77	1,000	Rubidium-86	100
Germanium-78	1,000	Rubidium-87	100
Arsenic-69	1,000	Rubidium-88	1,000
Arsenic-70	1,000	Rubidium-89	1,000
Arsenic-71	100	Strontium-80	100
Arsenic-72	100	Strontium-81	1,000
Arsenic-73	100	Strontium-83	100
Arsenic-74	100	Strontium-85m	1,000
Arsenic-76	100	Strontium-85	100
Arsenic-77	100	Strontium-87m	1,000
Arsenic-78	1,000	Strontium-89	10
Selenium-70	1,000	Strontium-90	0.1
Selenium-73m	1,000	Strontium-91	100
Selenium-73	100	Strontium-92	100
Selenium-75	100	Yttrium-86m	1,000
Selenium-79	100	Yttrium-86	100
Selenium-81m	1,000	Yttrium-87	100
Selenium-81	1,000	Yttrium-88	10
Selenium-83	1,000	Yttrium-90m	1,000
Bromine-74m	1,000	Yttrium-90	10
Bromine-74	1,000	Yttrium-91m	1,000
Bromine-75	1,000	Yttrium-91	10
Bromine-76	100	Yttrium-92	100
Bromine-77	1,000	Yttrium-93	100
Bromine-80m	1,000	Yttrium-94	1,000
Bromine-80	1,000	Yttrium-95	1,000
Bromine-82	100	Zirconium-86	100
Bromine-83	1,000	Zirconium-88	10
Bromine-84	1,000	Zirconium-89	100
Krypton-74	1,000	Zirconium-93	1
Krypton-76	1,000	Zirconium-95	10
Krypton-77	1,000	Zirconium-97	100
Krypton-79	1,000		
Niobium-88	1,000	Palladium-101	1,000
Niobium-89m		Palladium-103	100
(66 min)	1,000	Palladium-107	10
Niobium-89		Palladium-109	100
(122 min)	1,000	Silver-102	1,000
Niobium-90	100	Silver-103	1,000
Niobium-93m	10	Silver-104m	1,000
Niobium-94	1	Silver-104	1,000

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Niobium-95m	100	Silver-105	100
Niobium-95	100	Silver-106m	100
Niobium-96	100	Silver-106	1,000
Niobium-97	1,000	Silver-108m	1
Niobium-98	1,000	Silver-110m	10
Molybdenum-90	100	Silver-111	100
Molybdenum-93m	100	Silver-112	100
Molybdenum-93	10	Silver-115	1,000
Molybdenum-99	100	Cadmium-104	1,000
Molybdenum-101	1,000	Cadmium-107	1,000
Technetium-93m	1,000	Cadmium-109	1
Technetium-93	1,000	Cadmium-113m	0.1
Technetium-94m	1,000	Cadmium-113	100
Technetium-94	1,000	Cadmium-115m	10
Technetium-96m	1,000	Cadmium-115	100
Technetium-96	100	Cadmium-117m	1,000
Technetium-97m	100	Cadmium-117	1,000
Technetium-97	1,000	Indium-109	1,000
Technetium-98	10	Indium-110m	
Technetium-99m	1,000	(69.1m)	1,000
Technetium-99	100	Indium-110	
Technetium-101	1,000	(4.9h)	1,000
Technetium-104	1,000	Indium-111	100
Ruthenium-94	1,000	Indium-112	1,000
Ruthenium-97	1,000	Indium-113m	1,000
Ruthenium-103	100	Indium-114m	10
Ruthenium-105	1,000	Indium-115m	1,000
Ruthenium-106	1	Indium-115	100
Rhodium-99m	1,000	Indium-116m	1,000
Rhodium-99	100	Indium-117m	1,000
Rhodium-100	100	Indium-117	1,000
Rhodium-101m	1,000	Indium-119m	1,000
Rhodium-101	10	Tin-110	100
Rhodium-102m	10	Tin-111	1,000
Rhodium-102	10	Tin-113	100
Rhodium-103m	1,000	Tin-117m	100
Rhodium-105	100	Tin-119m	100
Rhodium-106m	1,000	Tin-121m	100
Rhodium-107	1,000	Tin-121	1,000
Palladium-100	100		
Tin-123m	1,000	Tellurium-133	1,000
Tin-123	10	Tellurium-134	1,000
Tin-125	10	Iodine-120m	1,000
Tin-126	10	Iodine-120	100
Tin-127	1,000	Iodine-121	1,000
Tin-128	1,000	Iodine-123	100
Antimony-115	1,000	Iodine-124	10
Antimony-116m	1,000	Iodine-125	1
Antimony-116	1,000	Iodine-126	1
Antimony-117	1,000	Iodine-128	1,000
Antimony-118m	1,000	Iodine-129	1
Antimony-119	1,000	Iodine-130	10
Antimony-120		Iodine-131	1

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(16m)	1,000	Iodine-132m	100
Antimony-120		Iodine-132	100
(5.76d)	100	Iodine-133	10
Antimony-122	100	Iodine-134	1,000
Antimony-124m	1,000	Iodine-135	100
Antimony-124	10	Xenon-120	1,000
Antimony-125	100	Xenon-121	1,000
Antimony-126m	1,000	Xenon-122	1,000
Antimony-126	100	Xenon-123	1,000
Antimony-127	100	Xenon-125	1,000
Antimony-128		Xenon-127	1,000
(10.4m)	1,000	Xenon-129m	1,000
Antimony-128		Xenon-131m	1,000
(9.01h)	100	Xenon-133m	1,000
Antimony-129	100	Xenon-133	1,000
Antimony-130	1,000	Xenon-135m	1,000
Antimony-131	1,000	Xenon-135	1,000
Tellurium-116	1,000	Xenon-138	1,000
Tellurium-121m	10	Cesium-125	1,000
Tellurium-121	100	Cesium-127	1,000
Tellurium-123m	10	Cesium-129	1,000
Tellurium-123	100	Cesium-130	1,000
Tellurium-125m	10	Cesium-131	1,000
Tellurium-127m	10	Cesium-132	100
Tellurium-127	1,000	Cesium-134m	1,000
Tellurium-129m	10	Cesium-134	10
Tellurium-129	1,000	Cesium-135m	1,000
Tellurium-131m	10	Cesium-135	100
Tellurium-131	100	Cesium-136	10
Tellurium-132	10	Cesium-137	10
Tellurium-133m	100	Cesium-138	1,000
Barium-126	1,000	Promethium-141	1,000
Barium-128	100	Promethium-143	100
Barium-131m	1,000	Promethium-144	10
Barium-131	100	Promethium-145	10
Barium-133m	100	Promethium-146	1
Barium-133	100	Promethium-147	10
Barium-135m	100	Promethium-148m	10
Barium-139	1,000	Promethium-148	10
Barium-140	100	Promethium-149	100
Barium-141	1,000	Promethium-150	1,000
Barium-142	1,000	Promethium-151	100
Lanthanum-131	1,000	Samarium-141m	1,000
Lanthanum-132	100	Samarium-141	1,000
Lanthanum-135	1,000	Samarium-142	1,000
Lanthanum-137	10	Samarium-145	100
Lanthanum-138	100	Samarium-146	1
Lanthanum-140	100	Samarium-147	100
Lanthanum-141	100	Samarium-151	10
Lanthanum-142	1,000	Samarium-153	100
Lanthanum-143	1,000	Samarium-155	1,000
Cerium-134	100	Samarium-156	1,000
Cerium-135	100	Europium-145	100

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Cerium-137m	100
Cerium-137	1,000
Cerium-139	100
Cerium-141	100
Cerium-143	100
Cerium-144	1
Praseodymium-136	1,000
Praseodymium-137	1,000
Praseodymium-138m	1,000
Praseodymium-139	1,000
Praseodymium-142m	1,000
Praseodymium-142	100
Praseodymium-143	100
Praseodymium-144	1,000
Praseodymium-145	100
Praseodymium-147	1,000
Neodymium-136	1,000
Neodymium-138	100
Neodymium-139m	1,000
Neodymium-139	1,000
Neodymium-141	1,000
Neodymium-147	100
Neodymium-149	1,000
Neodymium-151	1,000
Terbium-147	1,000
Terbium-149	100
Terbium-150	1,000
Terbium-151	100
Terbium-153	1,000
Terbium-154	100
Terbium-155	1,000
Terbium-156m (5.0h)	1,000
Terbium-156m (24.4 h)	1,000
Terbium-156	100
Terbium-157	10
Terbium-158	1
Terbium-160	10
Terbium-161	100
Dysprosium-155	1,000
Dysprosium-157	1,000
Dysprosium-159	100
Dysprosium-165	1,000
Dysprosium-166	100
Holmium-155	1,000
Holmium-157	1,000
Holmium-159	1,000
Holmium-161	1,000
Holmium-162m	1,000
Holmium-162	1,000
Holmium-164m	1,000
Europium-146	100
Europium-147	100
Europium-148	10
Europium-149	100
Europium-150 (12.62h)	100
Europium-150 (34.2y)	1
Europium-152m	100
Europium-152	1
Europium-154	1
Europium-155	10
Europium-156	100
Europium-157	100
Europium-158	1,000
Gadolinium-145	1,000
Gadolinium-146	10
Gadolinium-147	100
Gadolinium-148	0.001
Gadolinium-149	100
Gadolinium-151	10
Gadolinium-152	100
Gadolinium-153	10
Gadolinium-159	100
Ytterbium-162	1,000
Ytterbium-166	100
Ytterbium-167	1,000
Ytterbium-169	100
Ytterbium-175	100
Ytterbium-177	1,000
Ytterbium-178	1,000
Lutetium-169	100
Lutetium-170	100
Lutetium-171	100
Lutetium-172	100
Lutetium-173	10
Lutetium-174m	10
Lutetium-174	10
Lutetium-176m	1,000
Lutetium-176	100
Lutetium-177m	10
Lutetium-177	100
Lutetium-178m	1,000
Lutetium-178	1,000
Lutetium-179	1,000
Hafnium-170	100
Hafnium-172	1
Hafnium-173	1,000
Hafnium-175	100
Hafnium-177m	1,000
Hafnium-178m	0.1
Hafnium-179m	10

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Holmium-164	1,000	Hafnium-180m	1,000
Holmium-166m	1	Hafnium-181	10
Holmium-166	100	Hafnium-182m	1,000
Holmium-167	1,000	Hafnium-182	0.1
Erbium-161	1,000	Hafnium-183	1,000
Erbium-165	1,000	Hafnium-184	100
Erbium-169	100	Tantalum-172	1,000
Erbium-171	100	Tantalum-173	1,000
Erbium-172	100	Tantalum-174	1,000
Thulium-162	1,000	Tantalum-175	1,000
Thulium-166	100	Tantalum-176	100
Thulium-167	100	Tantalum-177	1,000
Thulium-170	10	Tantalum-178	1,000
Thulium-171	10	Tantalum-179	100
Thulium-172	100	Tantalum-180m	1,000
Thulium-173	100	Tantalum-180	100
Thulium-175	1,000	Tantalum-182m	1,000
Tantalum-182	10	Iridium-188	100
Tantalum-183	100	Iridium-189	100
Tantalum-184	100	Iridium-190m	1,000
Tantalum-185	1,000	Iridium-190	100
Tantalum-186	1,000	Iridium-192m	
Tungsten-176	1,000	(1.4m)	10
Tungsten-177	1,000	Iridium-192	
Tungsten-178	1,000	(73.8d)	1
Tungsten-179	1,000	Iridium-194m	10
Tungsten-181	1,000	Iridium-194	100
Tungsten-185	100	Iridium-195m	1,000
Tungsten-187	100	Iridium-195	1,000
Tungsten-188	10	Platinum-186	1,000
Rhenium-177	1,000	Platinum-188	100
Rhenium-178	1,000	Platinum-189	1,000
Rhenium-181	1,000	Platinum-191	100
Rhenium-182		Platinum-193m	100
(12.7h)	1,000	Platinum-193	1,000
Rhenium-182		Platinum-195m	100
(64.0h)	100	Platinum-197m	1,000
Rhenium-184m	10	Platinum-197	100
Rhenium-184	100	Platinum-199	1,000
Rhenium-186m	10	Platinum-200	100
Rhenium-186	100	Gold-193	1,000
Rhenium-187	1,000	Gold-194	100
Rhenium-188m	1,000	Gold-195	10
Rhenium-188	100	Gold-198m	100
Rhenium-189	100	Gold-198	100
Osmium-180	1,000	Gold-199	100
Osmium-181	1,000	Gold-200m	100
Osmium-182	100	Gold-200	1,000
Osmium-185	100	Gold-201	1,000

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Osmium-189m	1,000	Mercury-193m	100
Osmium-191m	1,000	Mercury-193	1,000
Osmium-191	100	Mercury-194	1
Osmium-193	100	Mercury-195m	100
Osmium-194	1	Mercury-195	1,000
Iridium-182	1,000	Mercury-197m	100
Iridium-184	1,000	Mercury-197	1,000
Iridium-185	1,000	Mercury-199m	1,000
Iridium-186	100	Mercury-203	100
Iridium-187	1,000		
Thallium-194m	1,000	Francium-223	100
Thallium-194	1,000	Radium-223	0.1
Thallium-195	1,000	Radium-224	0.1
Thallium-197	1,000	Radium-225	0.1
Thallium-198m	1,000	Radium-226	0.1
Thallium-198	1,000	Radium-227	1,000
Thallium-199	1,000	Radium-228	0.1
Thallium-201	1,000	Actinium-224	1
Thallium-200	1,000	Actinium-225	0.01
Thallium-202	100	Actinium-226	0.1
Thallium-204	100	Actinium-227	0.001
Lead-195m	1,000	Actinium-228	1
Lead-198	1,000	Thorium-226	10
Lead-199	1,000	Thorium-227	0.01
Lead-200	100	Thorium-228	0.001
Lead-201	1,000	Thorium-229	0.001
Lead-202m	1,000	Thorium-230	0.001
Lead-202	10	Thorium-231	100
Lead-203	1,000	Thorium-232	100
Lead-205	100	Thorium-234	10
Lead-209	1,000	Thorium-natural	100
Lead-210	0.01	Protactinium-227	10
Lead-211	100	Protactinium-228	1
Lead-212	1	Protactinium-230	0.1
Lead-214	100	Protactinium-231	0.001
Bismuth-200	1,000	Protactinium-232	1
Bismuth-201	1,000	Protactinium-233	100
Bismuth-202	1,000	Protactinium-234	100
Bismuth-203	100	Uranium-230	0.01
Bismuth-205	100	Uranium-231	100
Bismuth-206	100	Uranium-232	0.001
Bismuth-207	10	Uranium-233	0.001
Bismuth-210m	0.1	Uranium-234	0.001
Bismuth-210	1	Uranium-235	0.001
Bismuth-212	10	Uranium-236	0.001
Bismuth-213	10	Uranium-237	100
Bismuth-214	100	Uranium-238	100
Polonium-203	1,000	Uranium-239	1,000
Polonium-205	1,000	Uranium-240	100
Polonium-207	1,000	Uranium-natural	100
Polonium-210	0.1	Neptunium-232	100
Astatine-207	100	Neptunium-233	1,000
Astatine-211	10	Neptunium-234	100

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Radon-220	1	Neptunium-235	100
Radon-222	1	Neptunium-236	
Francium-222	100	(1.15E+5)	0.001
Neptunium-236		Curium-242	0.01
(22.5h)	1	Curium-243	0.001
Neptunium-237	0.001	Curium-244	0.001
Neptunium-238	10	Curium-245	0.001
Neptunium-239	100	Curium-246	0.001
Neptunium-240	1,000	Curium-247	0.001
Plutonium-234	10	Curium-248	0.001
Plutonium-235	1,000	Curium-249	1,000
Plutonium-236	0.001	Berkelium-245	100
Plutonium-237	100	Berkelium-246	100
Plutonium-238	0.001	Berkelium-247	0.001
Plutonium-239	0.001	Berkelium-249	0.1
Plutonium-240	0.001	Berkelium-250	10
Plutonium-241	0.01	Californium-244	100
Plutonium-242	0.001	Californium-246	1
Plutonium-243	1,000	Californium-248	0.01
Plutonium-244	0.001	Californium-249	0.001
Plutonium-245	100	Californium-250	0.001
Americium-237	1,000	Californium-251	0.001
Americium-238	100	Californium-252	0.001
Americium-239	1,000	Californium-253	0.1
Americium-240	100	Californium-254	0.001
Americium-241	0.001	Einsteinium-250	100
Americium-242m	0.001	Einsteinium-251	100
Americium-242	10	Einsteinium-253	0.1
Americium-243	0.001	Einsteinium-254m	1
Americium-244m	100	Einsteinium-254	0.01
Americium-244	10	Fermium-252	1
Americium-245	1,000	Fermium-253	1
Americium-246m	1,000	Fermium-254	10
Americium-246	1,000	Fermium-255	1
Curium-238	100	Fermium-257	0.01
Curium-240	0.1	Mendelevium-257	10
Curium-241	1	Mendelevium-258	0.01
Any alpha-emitting radionuclide not listed above or mistures of alpha emitters of unknown composition	0.001	Any radionuclide other than alpha-emitting radionuclides not listed above, or mixtures of beta emitters of unknown composition	0.01

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NOTE: For purposes of § 175.03(j)(2)(v), § 175.03(j)(5)(i) and § 175.03(l)(1)(i) where there is involved a combination of radionuclides in known amounts, the limit for the combination shall be derived as follows: determine, for each radionuclide in the combination, the ratio between the quantity present in the combination and the limit otherwise established for the specific radionuclide when not in combination. The sum of such ratios for all

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radionuclides in the combination may not exceed "1in.--that is, unity.

FOOTNOTES:

* The quantities listed above were derived by taking 1/10th of the most restrictive ALI listed in Table I, Columns 1 and 2, of Appendix B to § 175.03, rounding to the nearest factor of 10, and constraining the values listed between 37 Bq and 37 MBq (0.001 and 1,000 <<mu>>Ci). Values of 3.7 MBq (100 <<mu>>Ci)
have been assigned for radionuclides having a radioactive half-life in excess of E+9 years, except rhenium, 37 MBq (1,000 <<mu>>Ci), to take into account their low specific activity.

** To convert <<mu>>Ci to kBq, multiply the <<mu>>Ci value by 37.

APPENDIX D
RADIOACTIVE SURFACE
CONTAMINATION LIMITS

Application	Alpha (dpm/100cm 2)		Beta/Gamma 1	
	Total	Removable	Total (mR/hr)	Removable (dpm/100cm 2)
Controlled area				
Basic guide	25,000 Max. 5,000 Av.	500	1.0	5,000
Clean area	1,000	100	0.5	1,000
Non-controlled area				
Skin, personal clothing	500	ND 2	0.1	ND 2
Release of material or facilities	2,500 (Max.) 500 (Av.)	100	0.2	1,000

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Controlled area

Basic guide 25,000 Max. 500 1.0 5,000

5,000 Av.

Clean area 1,000 100 0.5 1,000

Non-controlled area

Skin, personal clothing 500 ND 2 0.1 ND 2

Release of material or 2,500 (Max.) 100 0.2 1,000

facilities 500 (Av.)

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1 Measured at 1 cm from the surface

2 ND-Non-detectable

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NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
GENERAL PROVISIONS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.04. Notices, instructions and reports to workers; inspections.

(a) Purpose and scope. (1) This section establishes requirements for notices, instructions and reports by licensees and registrants to individuals engaged in activities under a license or registration and options available to such individuals in connection with Department inspections of licensees or registrants to ascertain compliance with the provisions of all applicable regulations and conditions stated on the license or registration regarding radiological working conditions. The regulations in this section apply to all persons who receive, possess, produce, use, own or transfer sources of radiation registered with or licensed by the Department or are otherwise subject to this Code.

(b) Posting of notices to workers. (1) Each licensee and/or registrant shall post current copies of the following documents:

(i) this Code;

(ii) the radioactive materials license and the conditions or documents incorporated by reference into the license and any amendments thereto;

(iii) the certified registration and the conditions or documents incorporated by reference into the certified registration and any amendments thereto;

(iv) the certificate of registration;

(v) the operating procedures applicable to the work under the license, registration and/or certified registration;

(vi) any notice of violation involving radiological working conditions, any proposed imposition of civil penalty or order issued pursuant to this Code and any response from the licensee and/or registrant.

(2) If posting of a document specified in § 175.04(b)(1)(i), (ii), (iii) or (v) is not practicable, the licensee and/or registrant may post a notice which describes the document and states where it may be examined.

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(3) A current copy of the "Notice to Employees" prescribed by the Department shall be posted by each licensee and/or registrant wherever individuals work in or frequent any portion of a restricted area.

(4) Documents, notices, or forms posted pursuant to this Code shall appear in a sufficient number of places to permit individuals engaged in work under the license, certified registration and/or registration to observe them on the way to or from any particular work location to which the document applies, shall be conspicuous, and shall be replaced if defaced or altered.

(5) Department documents posted pursuant to § 175.04(b)(1)(vi) shall be posted within two (2) working days after receipt of the documents from the Department; the licensee's and/or registrant's response, if any, shall be posted within two (2) working days after dispatch from the licensee and/or registrant. Such documents shall remain posted for a minimum of five (5) working days or until action correcting the violation has been completed, whichever is later.

(c) Instructions to workers. (1) All individuals working in or frequenting any portion of a restricted area:

(i) shall be kept informed of the storage, transfer, or use of radioactive material, of radiation producing equipment or of radiation in such portions of the restricted area;

(ii) shall be instructed in the operating procedures applicable to work under the license, registration or certified registration and in the health protection problems associated with exposure to such radioactive material or radiation, in precautions or procedures to minimize exposure, in the purposes and functions of protective devices employed, and shall be required to demonstrate familiarity with such precautions, procedures and devices;

(iii) shall be instructed in, and instructed to observe, to the extent within the worker's control, the applicable provisions of this Code, licenses, certified registrations and registrations for the protection of personnel from exposures to radiation or radioactive material occurring in such areas;

(iv) shall be instructed of their responsibility to report promptly to the licensee and/or registrant any condition which may lead to or cause a violation of this Code, licenses and registrations or unnecessary exposure to radiation or radioactive material;

(v) shall be instructed in the appropriate response to warnings made in the event of any unusual occurrence or malfunction that may involve exposure to radiation or radioactive material; and

(vi) shall be advised as to the radiation exposure reports which workers must be given or may request pursuant to § 175.04(d).

(2) The extent of these instructions shall be commensurate with potential radio-

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logical health protection problems in the restricted area.

(3) Instruction shall be given before an individual begins work in a restricted area and at least annually thereafter.

(4) Records documenting individual worker instruction shall be maintained for inspection by the Department for a period of three (3) years.

(d) Notification and reports to workers. (1) Radiation exposure data for an individual and the results of any measurements, analyses, and calculations of radioactive material deposited or retained in the body of an individual shall be reported to the individual as specified herein. The information reported shall include data and results obtained pursuant to this Code or license or certified registration conditions as shown in records maintained by the licensee and/or registrant pursuant to § 175.03(k)(8) of this Code. Each notification and report shall:

(i) be in writing;

(ii) include appropriate identifying data such as the name of the licensee and/or registrant, the name of the individual and the individual's social security number;

(iii) include the individual's exposure information; and

(iv) contain the following statement: "This report is furnished to you under the provisions of § 175.04 of the New York City Health Code. You should preserve this report for further reference."

(2) Each licensee and/or registrant shall advise each worker annually of the worker's exposure to radiation or radioactive material as shown in records maintained by the licensee and/or registrant pursuant to § 175.03(k)(8).

(3) Each licensee and/or registrant shall furnish a report of the worker's exposure to sources of radiation at the request of the worker formerly engaged in activities controlled by the licensee or registrant. The report shall include the dose record for each year the worker was required to be monitored pursuant to § 175.03(f)(2) of this Code, or the equivalent provisions of previous versions. Such report shall be furnished within thirty (30) days from the date of the request, or within thirty (30) days after the dose of the individual has been determined by the licensee or registrant, whichever is later. The report shall cover the period of time that the worker's activities involved exposure to sources of radiation and shall include the dates and locations of licensed or registered (including certified registrations) activities.

(4) When a licensee and/or registrant is required pursuant to § 175.03(l)(3) to report to the Department any exposure of an individual to radiation or radioactive material, the licensee and/or registrant shall also provide the individual a report on the exposure data included therein. Such reports shall be transmitted at a time not later than the transmittal to the Department.

(5) At the request of a worker who is terminating employment with the licensee

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and/or registrant in work involving exposure to radiation or radioactive material, during the current year, each licensee and/or registrant shall provide at termination to each such worker, or to the worker's designee, a written report regarding the radiation dose received by that worker from operations of the licensee or registrant during the current year or fraction thereof. If the most recent individual monitoring results are not available at that time, a written estimate of the dose shall be provided together with a clear indication that this is an estimate.

(e) Presence of representatives of licensees and/or registrants and workers during inspections. (1) Each licensee and/or registrant shall afford the Department at all reasonable times opportunity to perform an inspection of materials, machines, activities, facilities, premises and records pursuant to this Code.

(2) During an inspection, Department inspectors may consult privately with workers as specified in § 175.04(f). The licensee and/or registrant, or that person's representative, may accompany the Department inspectors during other phases of an inspection.

(3) If, at the time of inspection, an individual has been authorized by the workers to represent them during Department inspections, the licensee and/or registrant shall notify the inspectors of such authorization and shall give the workers' representatives an opportunity to accompany the inspectors during the inspection of physical working conditions.

(4) Each workers' representative shall be routinely engaged in work under control of the licensee and/or registrant and shall have received instructions as specified in § 175.04(c).

(5) Different representatives of licensees or registrants and workers may accompany the inspectors during different phases of an inspection if there is no resulting interference with the conduct of the inspection. However, only one workers' representative at a time may accompany the inspectors.

(6) With the approval of the licensee and/or registrant and the workers' representative, an individual who is not routinely engaged in work under control of the licensee and/or registrant, for example a consultant to the licensee and/or registrant or to the workers' representative, shall be afforded the opportunity to accompany inspectors during the inspection of physical working conditions.

(7) Notwithstanding the other provisions of § 175.04(e), Department inspectors are authorized to refuse to permit accompaniment by any individual who deliberately interferes with a fair and orderly inspection. With regard to any area containing proprietary information, the workers' representative for that area shall be an individual previously authorized by the licensee and/or registrant to enter that area.

(f) Consultation with workers during inspections. (1) Department inspectors may consult privately with workers concerning matters of occupational radiation protection and other matters related to applicable provisions of this Code, licenses, certified registrations and registrations to the extent the inspectors deem necessary for the conduct of an effective and thorough inspection.

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(2) During the course of an inspection any worker may bring privately to the attention of the inspectors, either orally or in writing, any past or present condition which the worker has reason to believe may have contributed to or caused any violations of this Code, license condition, certified registration condition or registration, or any unnecessary exposure of any individual to radiation from sources of radiation under the licensee's and/or registrant's control. Any such notices in writing shall comply with the requirements of § 175.04(g)(1).

(3) The provisions of § 175.04(f)(2) shall not be interpreted as authorization to disregard instructions pursuant to § 175.04(c).

(g) Requests by workers for inspections. (1) Any worker or representative of workers who believes that a violation of this Code, registration, certified registration or license conditions exists or has occurred in work under a license and/or certificate of registration with regard to radiological working conditions in which the worker is engaged, may request an inspection by giving notice of the alleged violation to the Bureau of Radiological Health. Any such notice shall be in writing, shall set forth the specific grounds for the notice, and shall be signed by the worker or representative of the workers. A copy shall be provided to the licensee and/or registrant by the Bureau of Radiological Health no later than at the time of inspection except that, upon the request of the worker giving such notice, such worker's name and the name of individuals referred to therein shall not appear in such copy or any record published, released, or made available by the Department, except for good cause shown.

(2) If, upon receipt of such notice, the Bureau of Radiological Health determines that the complaint meets the requirements set forth in § 175.04(g)(1), and that there are reasonable grounds to believe that the alleged violation exists or has occurred, an inspection shall be made as soon as practicable, to determine if such alleged violation exists or has occurred. Inspections performed pursuant to § 175.04(g) need not be limited to matters referred to in the complaint.

(3) No licensee, registrant, or contractor or subcontractor of a licensee or registrant shall discharge or in any manner discriminate against any worker because such worker has filed any complaint or instituted or caused to be instituted any proceeding under this Code or has testified or is about to testify in any such proceeding or because of the exercise by such worker on behalf of such worker or others of any option afforded by this Code.

(h) Inspections not warranted; informal review. (1) If the Bureau of Radiological Health determines, with respect to a complaint under § 175.04(g), that an inspection is not warranted because there are no reasonable grounds to believe that a violation exists or has occurred, the Bureau of Radiological Health shall notify the complainant in writing of such determination. The complainant may obtain review of such determination by submitting a written statement of position with the Deputy Commissioner for Environmental Health Services, who will provide the licensee and/or registrant with a copy of such statement by certified mail, excluding, at the request of the complainant, the name of the complainant. The licensee or registrant may submit an opposing written statement of position with the Deputy Commissioner of Environmental Health Services who will provide the complainant with a

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copy of such statement by certified mail.

(2) Upon the request of the complainant, the Deputy Commissioner for Environmental Health Services may hold an informal conference in which the complainant and the licensee and/or registrant may orally present their views. An informal conference may also be held at the request of the licensee and/or registrant, but disclosure of the identity of the complaint will be made only following receipt of written authorization from the complainant. After considering all written or oral views presented, the Deputy Commissioner for Environmental Health Services shall affirm, modify, or reverse the determination of the Bureau of Radiological Health and furnish the complainant and the licensee and/or registrant a written notification of the decision and the reason therefor.

(3) If the Bureau of Radiological Health determines that an inspection is not warranted because the requirements of § 175.04(g)(1) have not been met, the complainant shall be notified in writing of such determination. Such determination shall be without prejudice to the filing of a new complaint meeting the requirements of § 175.04(g)(1).

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.05. Fees.

(a) Inspection fees. Notwithstanding any other provision of this Code, the Department is authorized to charge the following inspection fees pursuant to Section 225 of the Public Health Law and the regulations promulgated thereunder:

(1) For radiation equipment facilities required to have quality assurance programs pursuant to § 175.07 of this Code, the following inspection fees apply:

(i) Annually inspected facilities:

(A) Large hospital (more than 40 tubes) base fee: $1960.00

(B) Medium hospital (21-40 tubes) base fee: $1585.00

(C) Small hospital (1-20 tubes) base fee: $1290.00

(D) Large (more than 2500 examinations per year, excluding mammography) non-hospital base fee: $670.00

(E) Small (less than 2500 examinations per year, excluding mammography) non-hospital base fee: $375.00

(ii) Biennially inspected facilities:

(A) Large (more than 2500 examinations per year) facility base fee: $670.00

(B) Small (less than 2500 examinations per year) facility base fee: $375.00

(iii) For each tube inspected at annually or biennially inspected facilities, the following inspection fees apply in addition to the base fee:

(A) Radiographic: $120.00

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(B) Fluoroscopic: $175.00

(C) Mammographic: $295.00

(D) Dental: $60.00

(E) All other: $60.00

(iv) For radiation equipment facilities not required to have quality assurance programs pursuant to § 175.07 of this Code, the following inspection fees apply:

(A) First tube: $170.00

(B) Each additional tube: $60.00

(2) For linear accelerator facilities, the following fee applies:

(i) Base fee: $715.00

(3) For facilities licensed to possess and use radioactive materials, the following inspection fees apply:

(i) Specific licenses authorizing teletherapy

(A) Base fee: $320.00

(ii) Specific licenses of limited scope authorizing medical use (except for teletherapy)

(A) Base fee: $610.00

(B) Per site fee: $140.00

(iii) Specific licenses of limited scope authorizing non-human use

(A) Base fee: $385.00

(B) Per site fee: $160.00

(iv) Specific licenses of broad scope authorizing medical use (except for teletherapy)

(A) Base fee: $3515.00

(B) Per site fee: $140.00

(v) Specific license of broad scope authorizing research and development (non-

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human use)

(A) Base fee: $2450.00

(B) Per site fee: $160.00

(b) Due date for inspection fees. (1) Payment for inspection fees is due and payable thirty (30) days from the billing date.

(2) Failure to pay any inspection fee may result in the suspension or revocation of a registration, certified registration or radioactive materials license.

HISTORICAL NOTE

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§ 175.06. Professional practitioners and related provisions.

(a) Nothing in these regulations shall limit any human use of radiation in diagnostic and therapeutic procedures pursuant to sections below, provided that with respect to human use of radioactive materials, such use is in accordance with a specific license issued pursuant to these regulations, or an exemption therefrom, or under a license issued by the New York State Department of Health or the U.S. Nuclear Regulatory Com- mission.

(b) Each professional practitioner who treats or diagnoses any alleged or proven case of radiation illness or radiation injury to any individual, except that which can be expected in the normal course of radiation therapy, shall report to the Commissioner in writing within seven (7) days such treatment or diagnosis, the fact thereof and the full name, address, social security number, and age of such individual.

(c) No person other than a professional practitioner shall direct or order the application of radiation to a human being; nor shall any person other than a professional practitioner, or a person working under the direction, order, or direct supervision of a professional practitioner apply radiation to a human being. Such direction, order to apply, application of, or administration of radiation shall be in the course of the practitioner's professional practice and shall comply with the following:

(1) The provisions of the license or other authorization of the professional practitioner under the Education Law of the State of New York, or any successor law or regulation, and all regulations pertinent thereto, including, but not limited to provisions as to those parts of the human body and those persons which the professional practitioner may diagnose, analyze or treat or to which he may direct or order the application of, or apply, radiation, and provisions as to the type of radiation which the professional practitioner may use and the purpose for which the professional practitioner may use it; and

(2) The applicable provisions of Part 89 of Title 10 of the Codes, Rules and Regulations of the State of New York and Article 35 of the Public Health Law of the

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State of New York, or any successor law or regulation, relating to the practice of radiologic technology including licensure requirements and the limitations under which radiologic technologists and other persons, other than professional practitioners, may apply x-rays to human beings and all regulations pertinent thereto.

(3) A professional practitioner shall be responsible for the supervision of any radiation employee who administers radiation to human beings to assure that each exposure is given consistent with expected medical benefit and in accordance with any standards or requirements relating to the practice for which he/she is licensed.

(d) A radiologic technologist shall be responsible for complying with the requirements of such technologist's license and the limitations established by the New York State Department of Health, Bureau of Radiologic Technology. Each activity carried out as a radiologic technologist shall be such as to assure the maximum medical benefit with the minimum radiation exposure to patients and employees.

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§ 175.07. Quality assurance programs and misadministration records and reports.

(a) Purpose and scope. This section establishes requirements for the use of machine-produced radiation, radioactive materials or the radiation therefrom for diagnostic and therapeutic uses in the healing arts. These requirements and provisions provide for the protection of the public health and safety and are in addition to, and not in substitution for, others in this Code. The requirements of this section apply to all applicants, licensees and registrants subject to this Code.

(b) Diagnostic facilities. A quality assurance program is a system of plans, actions, reviews, reports and records, the purpose of which is to ensure that diagnostic facilities achieve consistent high quality imaging and other diagnostic results, while maintaining radiation output and personnel doses within limits prescribed by the Department.

(1) Each radiation installation performing diagnostic x-ray and/or radioactive materials procedures, except dental, podiatric or veterinary facilities, shall implement a quality assurance program including at a minimum:

(i) the adoption of a manual containing written policies and procedures for radiation protection and describing the facility's quality assurance program. Policies and procedures must be consistent with the types of equipment and services provided including, but not limited to, identification of patients, use of gonadal or scoliosis shielding, personnel monitoring, protection of pregnant workers and patients, and holding of patients. The quality assurance manual must describe the various processing, generator and systems quality control tests appropriate for the types of equipment and services provided in sufficient detail to ensure that they will be performed properly;

(ii) the performance of quality control tests and the correction of deficiencies as specified in the quality assurance manual;

(iii) the maintenance of equipment records for each diagnostic imaging system, containing test results, records of equipment repairs and other pertinent information;

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(iv) the provisions of a formalized in-service training program for employees including, but not limited to, quality assurance and radiation safety procedures;

(v) the determination of radiation output at the point of skin entry for common x-ray examinations;

(vi) the measurement of the amount of activity of each dose of a radiopharmaceutical/radiobiologic administered to each patient;

(vii) the calculated absorbed dose for diagnostic procedures involving radioactive materials;

(viii) the provision of the information described in § 175.07(b)(1)(v), (vi) and (vii) to any patient upon request;

(ix) the performance of an ongoing program of analysis of repeated, rejected or misadministered diagnostic studies which is designed to identify and correct problems and to optimize quality; and

(x) the performance of an ongoing film processing quality assurance (sensitometry) program, including the use of sensitometry tools (densitometer, sensitometer and thermometer), the determination and plotting of daily sensitometry data, and corrective action when sensitometry values exceed tolerance limits.

(2) Each licensee or registrant shall maintain written records documenting processing quality assurance and audit activities for review by the Department. Such records shall be maintained by the licensee or registrant until after the next scheduled inspection is completed by the Department.

(c) External beam and brachytherapy. A quality assurance program for external beam therapy and brachytherapy is a system of plans, actions, reviews, reports and records, the purpose of which is to ensure a consistent and safe fulfillment of the dose prescription to the target volume, with minimal dose to normal tissue.

(1) Each licensee or registrant who uses external beam therapy and/or brachytherapy in humans shall implement a quality assurance program which includes at a minimum:

(i) the adoption of a quality assurance manual containing written policies and procedures designed to ensure effective supervision, safety, proper performance of equipment, effective communication and quality control. These must include policies and procedures to ensure that:

(A) each patient's evaluation and intended treatment is documented in the patient's record;

(B) a written, signed and dated order for medical use of radiation or radioactive material is made for each patient in accordance with § 175.06 of this Code;

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(C) each patient is positively identified;

(D) all orders and other treatment records are clear and legible;

(E) staff will be instructed to obtain clarification before treating a patient if any element of the order or other record is confusing, ambiguous or suspected of being erroneous;

(F) each patient's response to treatment is assessed by an authorized user physician, or a physician under the supervision of an authorized user physician, for external beam therapy and/or brachytherapy, as appropriate, and that unusual responses are evaluated as possible indications of treatment errors;

(G) complete treatment records containing data recorded at the time of each treatment are maintained;

(H) the treatment charts of patients undergoing fractionated treatment are checked for completeness and accuracy at weekly intervals;

(I) final plans of treatment and related calculations are checked for accuracy before 25 percent of the prescribed dose for external beam therapy or 50 percent of the prescribed dose for brachytherapy is administered. If a treatment plan and related calculations were originally prepared by a radiation therapy physicist possessing the qualifications specified in § 175.64(c)(2) or § 175.103(j)(10), it may be checked by the same person using a different calculational method. Treatment plans and related calculations prepared by all other personnel must be checked by a second person using procedures specified in the treatment planning procedures manual required pursuant to § 175.07(c)(2), and who has received training in the use of such manual;

(J) there is quality control for all physical components of radiation therapy such as: equipment function and safety (including treatment planning equipment), treatment planning procedures and computer codes, treatment application procedures, dosimetry and personnel radiation safety;

(K) that the quality control tests to be performed are documented, including:

(a) detailed procedures for performing each test;

(b) the frequency of each test;

(c) acceptable results for each test;

(d) corrective actions to be taken; and

(e) recordkeeping and reporting procedures for test results.

(2) Each licensee or registrant shall ensure that a radiation therapy physicist possessing the qualifications specified in § 175.64(c)(2) or § 175.103(j)(10), prepares a procedures manual describing how radiation therapy treatment planning is to

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be performed at the licensee's or registrant's facility. The treatment planning manual may be part of the quality assurance manual required by § 175.07(c)(1) and shall include the calculation methods and formulas to be used at the facility, including the methods for performing the checks of treatment plans and related calculations as required by § 175.07(c)(1). The treatment planning manual shall be reviewed annually by a radiation therapy physicist and shall be included in training given pursuant to § 175.04(c) of this Code to facility staff who will participate in treatment planning.

(3) Each licensee or registrant shall ensure that all equipment used in planning and administering radiation therapy is properly functioning and is designed and used for the intended purpose and is maintained in accordance with the manufacturer's instructions and the quality assurance program described in the licensee's or registrant's quality assurance manual. Such equipment shall be calibrated prior to use on patients, at least annually thereafter and following any change, repair or replacement of any component which may alter the radiation output.

(4) Each licensee or registrant shall implement procedures for auditing the effectiveness of the radiation therapy quality assurance program as specified below. Audit procedures must specify either that:

(i) external audits will be conducted at intervals not to exceed twelve (12) months by radiation therapy physicists possessing the qualifications specified in § 175.64(c)(2) or § 175.103(j)(10) and by physicians who are active in the practice of the type of radiation therapy conducted by the licensee or registrant. These shall be individuals who are not involved in the therapy program being audited; and

(A) the individuals who conduct the audit will prepare and deliver to the licensee or registrant a written report which contains an assessment of the effectiveness of the quality assurance program and makes recommendations for any needed modifications or improvements; and

(B) the licensee or registrant shall promptly review the audit findings, address the need for modifications or improvements and document actions taken. If recommendations are not acted on, the reasons for this also shall be documented; or

(ii) internal audits will be conducted at intervals not to exceed twelve (12) months by program staff who will prepare and deliver to the licensee or registrant a written report as specified in § 175.07(c)(4)(i)(A), and external audits will be conducted at intervals not to exceed five (5) years by an organized review program supervised by the American College of Radiology, or a program found to be equivalent by the Department based on the scope of the audit and the experience of the sponsoring organization in performing such audits; and

(A) the licensee or registrant shall promptly review the audit findings, address the need for modifications or improvements and document actions taken. If recommendations are not acted on, the reasons for this also shall be documented.

(5) Each licensee or registrant shall maintain written records documenting quality assurance and audit activities for review by the Department.

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(d) Therapy with radiopharmaceuticals and/or radiobiologics. A quality assurance program for radiopharmaceutical/radiobiologic therapy is a system of plans, actions, reviews, reports and records, the purpose of which is to ensure a consistent and safe fulfillment of the dose prescription.

(1) Each licensee who uses radiopharmaceuticals and/or radiobiologics for therapy in humans shall implement a quality assurance program which includes at a minimum:

(i) the adoption of a manual containing written policies and procedures designed to assure effective supervision, safety, proper performance of equipment, effective communication and quality control. These must include procedures to assure that:

(A) each patient's evaluation and intended treatment is documented in the patient's record;

(B) a written, signed and dated order for medical use of radioactive material is made in accordance with § 175.06 of this Code;

(C) each patient is positively identified;

(D) all orders and other treatment records are clear and legible;

(E) staff will be instructed to obtain clarification before treating a patient if any element of the order or other record is confusing, ambiguous or suspected of being erroneous;

(F) each patient's response to treatment is assessed by an authorized user physician, or a physician under the supervision of an authorized user physician, for radiopharmaceutical/radiobiologic therapy and that unusual responses are evaluated as possible indications of treatment errors; and

(G) complete treatment records containing data recorded at the time of each treatment are maintained.

(2) Each licensee shall ensure that all equipment used in planning and administering radiopharmaceutical/radiobiologic therapy is designed and used for the intended purpose and is properly functioning, is properly calibrated and is maintained in accordance with the manufacturer's instructions and the quality assurance program described in the licensee's or registrant's quality assurance manual.

(3) Each licensee shall audit the radiopharmaceutical/radiobiologic quality assurance program at intervals not to exceed twelve (12) months to assess the effectiveness of the program, document the audit and any modifications or improvements found to be needed and institute corrective actions and improvements as indicated by the audit findings.

(e) Records and reports of misadministrations. (1) Diagnostic misadministrations.

(i) Records of misadministrations as defined in § 175.02(a)(129) of this Code which involve diagnostic procedures and the corrective actions taken pursuant to §

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175.07(b)(1)(ix) shall be retained for three (3) years; and

(ii) if such a misadministration results in a dose to the patient exceeding 50 millisieverts (5 rem) to the whole body or 500 millisieverts (50 rem) to any individual organ, or involves the administration of iodine-125 or iodine-131 in the form of iodide in a quantity greater than 1 megabecquerel (30 microcuries), the licensee or registrant shall notify the Department in writing within fifteen (15) days and make and retain a record pursuant to § 175.07(e)(3).

(2) Therapy misadministrations. (i) When a misadministration described in § 175.02(a) (129)(v), (vi) or (vii), in which the percentage of error is equal to or less than 20 percent is discovered, the licensee or registrant shall immediately investigate the cause and take corrective action; and

(A) the licensee or registrant shall make and retain a record of all therapy misadministrations described in § 175.07(e)(2). The record shall contain all the information required by § 175.07(e)(3) and shall be retained for six (6) years.

(ii) When a therapy misadministration described in § 175.02(a)(129)(i), (ii), (iii) or (viii) is discovered, or when a misadministration described in § 175.02(a)(129)(v), (vi) or (vii) is discovered in which the percentage of error is greater than 20 percent, the licensee or registrant shall notify the Department by telephone within 24 hours. The licensee or registrant shall also notify the referring physician of the affected patient and the patient of any therapy misadministration described herein, with the exception of the misadministration defined in § 175.02(a)(129)(viii). When it is not medically advisable to give such information to the patient, the information shall be made available to the patient's responsible relative or guardian on the patient's behalf. These notifications must be made within 24 hours after the misadministration is discovered. If the referring physician, patient or the patient's responsible relative or guardian cannot be reached within 24 hours, the licensee or registrant shall notify them as soon as practicable. It is not required that the patient be notified without first consulting the referring physician; however, medical care for the patient shall not be delayed because of this.

(iii) Within seven (7) days after an initial therapy misadministration report, the licensee or registrant shall send a written report to the Department. The written report must contain the name of the licensee or registrant, the information required by § 175.07(e)(3) and whether the licensee or registrant notified the patient or the patient's responsible relative or guardian. This reporting requirement may be satisfied by submitting to the Department a copy of the incident report filed with the New York State Department of Health pursuant to 10 NYCRR Part 405 provided, however, that such report contains all information required by this Code.

(3) Each licensee or registrant shall maintain a record of each reportable diagnostic misadministration and each therapy misadministration for six (6) years. The record shall contain the names of all individuals involved in the event (including the treating physician, allied health personnel, the patient and the patient's referring physician), the patient's social security number or identification number if one has been assigned, a description of the event, the effect on the patient (including sequelae, prognosis and follow-up actions) and actions taken to prevent

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recurrence.

(4) Within seven (7) days after an initial therapy misadministration report made pursuant to § 175.07(e)(2)(ii), the licensee or registrant shall provide the patient a written report, with a copy to the patient's referring physician. The report shall contain a brief description of the event, the effect on the patient including any change in the patient's health status which resulted from or could result from the misadministration, and recommendations for the appropriate course of treatment or follow-up. If it is not medically advisable to give such information to the patient, the report shall be made available to the patient's responsible relative or guardian on the patient's behalf. Such action shall be documented in the patient's treatment record.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (b) par (1) subpars (viii), (ix) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (b) par (1) subpar (x) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (e) par (1) subpar (ii) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Subparagraphs (viii) and (ix) of paragraph (1) of subsection (b) were amended and subparagraph (x) of paragraph (1) and paragraph (2) of subsection (b) were adopted on March 21, 2001 to provide additional quality assurance requirements for the processing of radiographic film after the x-ray film has been produced and to insure the proper maintenance of records.

Subparagraph (ii) of paragraph (1) of subsection (e) was amended on March 21, 2001 to delete the reference to iodine 123 to make the regulation consistent with state and federal regulations which have been amended owing to the lesser potency of iodine 123.

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§ 175.51. Registration and inspection of installations with radiation equipment; other permitted activities.

(a) Applicability. The provisions of this section apply to all radiation installations using any radiation equipment, and to those persons who sell, assemble and install such equipment as applicable. However, those facilities possessing equipment subject to the requirements of § 175.64(b) of this Code are required to obtain a certified registration for that equipment. If such facility has additional radiation equipment not subject to the requirements of § 175.64(b) of this Code, such equipment shall be registered pursuant to this section.

(b) Registration required. (1) Prior to establishing, maintaining or operating any radiation installation at which is located any radiation equipment in operable condition, or prior to installing such equipment which is intended to be used, the owner or operator of such installation shall have obtained a current certificate of registration or, for a therapeutic radiation machine subject to the requirements of § 175.64(b) of this Code, a certified registration from the Department.

(2) For professional practitioners in private practice, registrations shall not be issued to anyone other than natural persons who shall be responsible for the use and operation of the equipment and shall be liable for violations of the conditions of the registration or the provisions of this Code.

(c) Application for a certificate of registration as described in § 175.51(b)(1) shall be made to the Department on a written form and in a manner prescribed by the Department.

(d) Facilities at which either the operator or location will be changed shall apply for a new registration at least thirty (30) days prior to such change.

(1) Facilities without a current certificate of registration shall apply as follows:

No registrant shall apply x-rays to treat or diagnose any patient's medical condition at a facility that does not possess a current, non-expired Certificate of

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Registration from the Department. This shall not prohibit the installation of radiation-producing equipment by a registrant at a facility solely for testing purposes by medical physicists.

(i) All new facilities possessing radiation-producing equipment, excluding dental, podiatric, and veterinary facilities, shall apply for a certificate of registration at least thirty (30) days prior to the .expected start date' for clinical operation of the facility.

(ii) The applicant for registration shall submit the following information:

(A) a completed application form; and

(B) a medical physicist report detailing the results of initial quality control tests conducted on all radiation-producing equipment in the facility. In this context, the initial quality control tests shall be the sum of all quality control tests mandated to be conducted for the facility type at daily, weekly, monthly, semiannual, annual and biennial frequencies. In addition, a radiation protection survey shall be conducted for each room housing a radiographic unit. In this context, a medical physicist shall be an individual possessing a current, non-expired CRESO certification in New York State, or a license to practice the specialty of diagnostic medical physics in New York State.

(iii) The Department has the right to refuse to grant a facility's registration until such time as the facility's physicist report contains all quality control mandated tests as submitted by an individual licensed in New York State or possessing a CRESO certification in New York State.

(iv) Upon completion of the review process for the submitted quality control tests by the facility, if reasons exist to refuse authorization to register the facility's radiation-producing equipment for clinical usage, the facility shall be notified of the reasons for such a decision by the Department in writing.

(2) All new dental, podiatric, and veterinary facilities without a current certificate of registration shall apply for a new registration at least thirty (30) days before establishing the installation and/or installing the x-ray equipment. All new dental, podiatric, and veterinary facilities shall be prohibited from commencing diagnostic clinical examinations until such time that the facility has complied with items (i) and (ii) below:

(i) New facilities shall file a completed application form with the Department; and

(ii) Prior to any clinical usage of radiation-producing equipment, all such new facilities shall be inspected by the Department and shall correct all deficiencies noted at the time of such inspection.

(iii) The Department shall have the right to refuse to issue a certificate of registration to any facility that refuses to allow the Department to conduct an inspection of all of the facility's x-ray equipment and/or refuses to correct any vi-

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olations of the Health Code noted during the inspection provided for in subparagraph (ii).

(e) Renewal registrations. Facilities with current, valid certificates of registration shall apply for renewal at least thirty (30) days prior to the expiration of such certificate of registration unless such certificate of registration is revoked or unless the installation is discontinued on or before the expiration of the certificate of registration.

(f) Fees for each registration shall be paid pursuant to § 5.09(j). For certified registrations, fees shall be paid pursuant to § 175.05.

(g) Suspension and revocation of registrations. (1) A registration may be denied, suspended or revoked pursuant to § 5.17 of this Code or if the Commissioner finds that:

(i) the information submitted in the application is incorrect or incomplete; or

(ii) the installation is, has been or will be established, maintained, or operated in violation of this Code or any other applicable law, rule, regulation, or order; or

(iii) the certificate of registration has not been issued correctly; or

(iv) the fees for registration have not been paid as required.

(h) A certificate of registration shall be issued for a limited period of time extending from the date of issuance to the date of expiration as specified on the certificate of registration. The length of such period of time shall not exceed two years except that the Department may issue a certificate of registration for a longer period of time in order to stagger expiration dates for administrative purposes and may charge a proportionate increase in fees therefor.

(i) Expiration of registrations. (1) The registration issued for a radiation installation to the operator thereof shall expire and may be required to be surrendered to the Department upon:

(i) the expiration date specified on the certificate of registration; or

(ii) revocation by the Commissioner; or

(iii) a change of the person to whom the certificate of registration is issued; or

(iv) a change in address of the radiation installation if it is not a mobile unit; or

(v) a change in the name of the installation; or

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(vi) the discontinuance of the installation.

(j) A certificate of registration shall not be transferable or assignable.

(k) A certificate of registration issued for a radiation installation shall be posted in accordance with the provisions of § 5.15 of this Code.

(l) The registration shall not imply endorsement or approval by the Department and shall not be used to advertise or promote business.

(m) The operator of a radiation installation shall keep correct registration by reporting to the Department within ten (10) days any change affecting such information.

(n) Inspections. (1) Any radiation installation subject to the registration requirements of this section or the certified registration requirements of § 175.64 of this Code shall be inspected periodically to assure compliance with the provisions of this Code and the maintenance of radiation exposures as far below the limits set forth in this Code as practicable.

(2) Except as otherwise provided in § 175.51(n)(3), such inspections shall be made at a frequency specified herein, with the first inspection to be made at the time of the beginning of operation.

(A) Hospitals, clinics, radiologists and any other type of facility as specified by the Department shall be inspected annually.

(B) Dental, podiatric, veterinary installations shall be inspected triennially.

(C) All other facilities shall be inspected biennially.

(D) Reinspections or other appropriate follow-up activities shall be made at intervals of sixty (60) days or less to ensure correction of any violation found during an initial inspection and remaining uncorrected at the conclusion thereof.

(E) The inspections shall be made by the Department or, as the Department shall direct for dental and podiatric installations, by Certified Radiation Equipment Safety Officers approved by the Department, to use the New York City Health Code, including Article 175 thereof, for compliance purposes.

(i) Certified Radiation Equipment Safety Officers shall furnish an inspection report, in a form prescribed by the Department, signed by the person who made the inspection to the operator of the installation and a copy thereof to the Department in accordance with the instructions of the prescribed form.

(ii) Certified Radiation Equipment Safety Officers shall not charge or propose to charge a fee for an inspection in excess of a fair and reasonable amount as determined by the New York State Department of Health.

(3) The Department may establish inspection frequencies of any installation dif-

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ferent from those specified in § 175.51(n)(2).

(o) Other activities requiring a permit. (1) No person shall engage in the business of selling new or used radiation equipment to, or assembling, installing, or repairing such equipment for, professional practitioners or any other holder of a registration to operate a radiation installation in the City without a permit therefor issued by the Department. Application for such permit shall be made to the Department on a written form in a manner prescribed by the Department and fees paid pursuant to § 5.07 of this Code.

(2) Each person who is engaged in the business of selling new or used radiation equipment to, or assembling, installing, or repairing such equipment for professional practitioners or any other person holding a registration to operate a radiation installation in the City shall comply with the reporting requirements in this Code.

(3) Any person who sells, assembles, installs or repairs new or used radiation equipment shall file a report with the Department pursuant to 21 CFR 1020.30(d), or any successor law or regulation, within fifteen (15) days of the completion of the activity.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (d) amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (n) par (2) subpar (E) added City Record July 30, 2004 eff. Aug. 29, 2004. [See Vol. 9 Statements of Basis and Purpose No. 21]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Subdivision (d) of § 175.51 was amended on September 26, 2006, in order to implement a NYS Comptroller's Audit Report (2001-N-9) recommendation that all new x-ray facilities be inspected by the Department prior to initiating clinical x-ray exams.

Paragraph (2) of subdivision (n) of § 175.51 was added on July 22, 2004 to authorize the Department to also utilize Certified Radiation Equipment Safety Officers for the inspections of dental and podiatric installations to assure timely inspections for such installations.

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§ 175.52. Exemptions of radiation equipment.

(a) The provisions of these regulations shall not apply to:

(1) radiation equipment constructed so that it cannot emit radiation at a level greater than 1.3 E-7 C/kg (0.5 milliroentgen) per hour, measured 5 cm (2 in.) from any accessible surface thereof, and averaged over an area of 10 cm 2 (1.55 in 2) provided, however, that such exemption shall not apply to the testing or servicing of such equipment during its production; or

(2) radiation equipment during its storage, shipment, retail sale or other similar use (but not including installation) during which such equipment is not connected to a voltage source and does not emit radiation, provided however, that such equipment is not exempt from the labeling requirement of § 175.56(d).

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.53. Prohibited uses and activities.

(a) Prohibited uses.

(1) Hand-held fluoroscopic screens shall not be used.

(2) Shoe-fitting fluoroscopic devices shall not be used.

(3) Intraoral fluoroscopy in dental examinations shall not be used.

(4) Photofluorographic equipment shall not be used.

(5) Equipment employing bare overhead or uninsulated conductors shall not be used.

(6) Non-image intensified fluoroscopes shall not be used.

(b) Prohibited activities. (1) Individuals shall not operate x-ray equipment such that the useful beam is applied to human beings unless such individual is a professional practitioner or is otherwise authorized to operate x-ray equipment pursuant to New York State law.

(2) The sale, lease, transfer or loan of x-ray or fluoroscopic equipment or the supplies appertaining thereto, except to persons engaged in an occupation where such use is permitted or to institutions where such use is permitted, is prohibited. However, this restriction shall not apply to persons intending to use x-ray or fluoroscopic equipment and supplies solely for the application of radiation to other than human beings, nor to the acquisition of such equipment or supplies by wholesalers, distributors or retailers in the regular course of their trade or business.

(3) No person shall sell, lease, transfer, lend or install any radiation-producing equipment, or the supplies used in connection with such equipment, unless such supplies or equipment when properly placed into operation or properly used

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will meet the requirements of this Code. A person who undertakes to repair such equipment shall repair the same properly so that when it is placed in operation or properly used after the repair the equipment will meet the requirements of this Code for radiation protection and safety generally.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.54. Surveys, shielding requirements and operator protection for diagnostic radiation machines.

(a) When appropriate, the Department may require the applicant for a certificate of registration to utilize the services of a qualified expert in shielding design to determine the shielding requirements prior to installation of radiation equipment. The shielding requirements shall be submitted to the Department for review.

(b) When appropriate, the Department may require that the continuity and adequacy of any protective barriers be verified by a protection survey performed by a qualified expert or a Certified Radiation Equipment Safety Officer. The findings shall be submitted to the Department for review, and shall be maintained by the owner or operator as part of the installation's radiation records and reports. (If the Department finds that the radiation surveys and reports of any qualified expert or Certified Radiation Equipment Safety Officer employed in this City are inadequate to assess radiation exposures, the Department may require the registrants and licensees to have such surveys and reports performed by other qualified experts or Certified Radiation Equipment Safety Officers and may notify the appropriate certifying agency of such action.

(c) Operator protection. (1) Fixed radiographic installations (except dental, mobile, portable, podiatric or mammographic systems or spot-film devices as defined in 21 CFR Section 1020.30(b)).

(i) The operator of radiographic equipment shall initiate x-ray exposures from a control console that satisfies the following requirements:

(A) the control console shall be located within a structure so constructed that

(a) radiation has to be scattered at least twice before entering the structure; or

(b) the structure shall be provided with a protective door that is interlocked to prevent an x-ray exposure unless the door is closed and which door shall have sufficient shielding to ensure compliance with the requirements of § 175.03; and

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(c) any walls of the structure shall be permanently fixed barriers of at least 2.13 m (7 ft) in height and shall provide sufficient shielding to ensure compliance with the requirements of § 175.03; and

(B) the operator shall be provided with a viewing system to observe the patient during any exposure and which has been so placed that the operator can view any entry into the radiographic room; and

(C) the operator shall be provided with a means of communication with the patient from the operator's position at the control console; and

(D) the operator shall be allotted unobstructed floor space at the control console; and

(E) for control consoles located in structures without an interlocked door, the x-ray exposure control shall be fixed in the structure and placed at least 1.02 m (40 in.) from any open edge of the structure where radiation may enter.

(2) Mobile, portable, podiatric and dental radiographic installations, excluding mammographic systems. (i) Mobile, portable, podiatric and dental x-ray equipment shall be provided with the means to allow an operator to stand at least 2 m (6 ft) from the patient or behind a protective barrier and not in the path of the primary x-ray beam whenever an x-ray exposure is initiated.

(ii) Mobile and portable x-ray systems, excluding dental and podiatric systems, that are used continuously for greater than one week in the same location shall be deemed a fixed radiographic installation and shall meet the operator protection standards of either § 175.54(c)(1)(i)(A) or (B).

(iii) Each operator of a mobile or portable radiographic x-ray unit, excluding dental and podiatric units, shall be provided with personnel monitoring as provided in § 175.03 and shall wear a protective apron of at least 0.25 mm lead equivalent.

(3) Mammographic installations. (i) The operator of the mammographic equipment shall initiate x-ray exposures from the control console of the mammographic equipment such that:

(A) protective shielding is provided for the operator that meets the following criteria:

(a) the shielding shall be provided to a height of 2m (6 ft) from the floor, with the lower edge not more than 7.5 cm (3 in.) from the floor. Such shielding shall be permanently attached to the mammographic x-ray unit in such a manner that an air gap does not exist between the shield and the mammographic unit or shall be constructed as a permanent operator shield such that the operator is able to stand completely within the shielded area during the exposure; and

(b) the exposure control shall be permanently fixed on the mammographic control console; and

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(c) the operator shall be able to communicate with and view the patient from the operator's protected position during the exposure.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.55. Compliance with federal standards and precedence thereof.

(a) Whenever any requirement of this Code relating to radiation equipment conflicts with the Federal performance standard (21 CFR 1020.30, or any successor law or regulation) for diagnostic x-ray systems and/or their major components, the Federal performance standard shall take precedence over and supersede any conflicting requirements of this Code pertaining to radiation equipment.

(b) The operator shall retain, and shall present to the Department for examination when requested, all information provided by the manufacturer to the purchaser in accordance with the requirements of the applicable Federal standard and shall transfer this information to any subsequent owner of the equipment.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.56. General requirements for radiation equipment.

(a) In addition to the requirements set forth in this section, all radiation equipment shall meet all applicable specific provisions of the sections of this Code set forth under the heading "Radiation Equipment" and all possession and use thereof shall comply with the requirements set forth in the sections of this Code set forth under the heading "General Provisions" and with any other applicable requirement deemed necessary by the Department.

(b) Radiation equipment which is not intended to be used must be made inoperable to the satisfaction of the Department by dismantling or sealing with an official Department seal or other suitable method, and shall not be unsealed or restored to operable condition without prior authorization by the Department.

(c) All radiation equipment, except equipment used solely in research and development, installed in a radiation installation shall, where applicable, be listed by the Underwriters' Laboratories, Inc., or shall comply with the National Electrical Code of the National Fire Protection Association or an equivalent safety standard.

(d) Labels. (1) The control panel for radiation equipment used for diagnostic or therapeutic radiology shall be clearly labeled with the words "CAUTION--THIS EQUIPMENT PRODUCES X-RAYS WHEN ENERGIZED" or other cautionary wording as specified in applicable performance standards established by the United States Department of Health and Human Services, Food and Drug Administration.

(2) Such cautionary labels shall not be used except as required by this Code.

(e) Equipment accessories. Film processing materials and techniques shall be those recommended by the x-ray film manufacturer or those otherwise tested to ensure maximum information content of the developed x-ray film and quality control methods shall be employed to ensure optimum results as specified in § 175.07 of this Code. X-ray film shall not be used past the manufacturer's expiration date.

(f) Beam quality--half-value layer. (1) The half-value layer (HVL) of the useful beam for a given x-ray tube potential shall not be less than the appropriate value

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shown in Table 1 below under "Specified dental systems" for any dental system designed for use with intraoral image receptors and manufactured after December 1, 1980, and under "Other x-ray systems" for all other x-ray systems subject to this Code. If it is necessary to determine such half-value layer at an x-ray tube potential which is not listed in Table 1, linear interpolation or extrapolation may be made.

(2) Positive means shall be provided to insure that at least the minimum filtration needed to achieve the beam quality requirements specified in § 175.56(f)(1) is in the useful beam during each exposure.

Table 1

X-ray tube voltage (kilovolt peak)	Designed operating range	Measured operating potential	Minimum HVL (mm of Al)	Specified dental systems	Other x-ray systems

Table 1 (cont'd)

X-ray tube voltage (kilovolt peak)	Designed operating range	Measured operating potential	Minimum HVL (mm of Al)	Specified dental systems	Other x-ray systems
Below 50	30		0.3		
	40		0.4		
	49		0.5		
50 to 70	50	1.5	1.2		
	60	1.5	1.3		
	70	1.5	1.5		
Above 70	71	2.1	2.1		
	80	2.3	2.3		
	90	2.5	2.5		
	100	2.7	2.7		
	110	3.0	3.0		
	120	3.2	3.2		
	130	3.5	3.5		
	140	3.8	3.8		
	150	4.1	4.1		

(g) Radiographic protective tube housings and diaphragm protection.

(1) All radiographic protective tube housings shall be of the diagnostic type.

(2) All diaphragms, collimators, cones, shutters or other devices used to define the useful beam shall provide the same degree of attenuation as that required of the tube housing.

(h) The tube head shall remain stationary when placed in any exposure position and during the exposure.

(i) Additional requirements. In addition to other applicable requirements of this Code, diagnostic x-ray equipment manufactured after August 1, 1974, shall meet the

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following requirements:

(1) Aluminum equivalent of material between patient and image receptor. The aluminum equivalent of each of the items listed in Table 2 below, which are used between the patient and image receptor, shall not exceed the indicated limits. Compliance shall be determined by x-ray measurements made at a potential of 100 kVp and with an x-ray beam which has a half-value layer of 2.7 mm of aluminum. This requirement is applicable to front panel(s) of cassette holders and film changers provided by the manufacturer for purposes of patient support and/or to prevent foreign object intrusions. It does not apply to such items as a screen and its associated mechanical support panel or grids.

Table 2
Item Aluminum equivalent (in mm)

Table 2 (cont'd)

Item	Aluminum equivalent (in mm)
Front panel(s) of cassette holder (total of all)	1.0
Front panel(s) of film changer (total of all)	1.0
Stationary tabletop	1.0
Moveable tabletop (including stationary subtop)	1.5
Cradle	2.0

(2) On battery-powered generators, visual means shall be provided on the control panel to indicate whether the battery is in a state of charge adequate for proper operation.

(j) The x-ray equipment, and all related items, including all individual components, whether related to the setting of exposure technique factors, the taking of the x-ray or post-exposure processing of the x-ray film, shall be maintained in proper working order. This includes, but is not limited to, dials, buttons, indicators, switches, controls and meters.

(k) All x-ray producing equipment shall be provided with a light, or other visual indication, which functions whenever x-rays are produced. This indication shall be visible at the operator's protected position.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (e) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (j), (k) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

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NOTES:

Subsection (e) was amended on March 21, 2001 to prohibit the use of outdated x-ray film. In addition, the reference to paragraph (I) was deleted as unnecessary.

Subsections (j) and (k) were added on March 21, 2001 to include additional requirements for x-ray producing equipment.

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§ 175.57. Diagnostic radiography (other than veterinary).

(a) Applicability. The provisions of this section apply to equipment for the recording of images.

(b) Control and indication of technique factors. (1) Visual indication. The technique factors to be used during an exposure shall be indicated before the exposure begins, except when automatic exposure controls are used, in which case the technique factors which are set prior to the exposure shall be indicated. On equipment having fixed technique factors, this requirement may be met by permanent markings. Indication of technique factors shall be visible from the operator's position except in the case of spot films made by the fluoroscopist.

(2) Timers. Means shall be provided to terminate the exposure at a preset time interval, a preset product of current and time, a present number of pulses, or a preset radiation exposure to the image receptor.

(i) Except during serial radiography, the operator shall be able to terminate the exposure at any time during an exposure of greater than one-half section. Termination of exposure shall cause automatic resetting of the timer to its initial setting or zero. It shall not be possible to make an exposure when the timer is set to a zero or off position if either position is provided.

(ii) During serial radiography, the operator shall be able to terminate the x-ray exposure(s) at any time, but means may be provided to permit completion of any single exposure of the series in progress.

(3) Automatic exposure controls. When an automatic exposure control is provided:

(i) indication shall be made on the control panel when this mode of operation is selected;

(ii) when the x-ray tube potential is equal to or greater than 51 kVp, the minimum exposure time for field emission equipment rated for pulsed operation shall be equal to or less than a time interval equivalent to two pulses and the minimum ex-

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posure time for all other equipment shall be equal to or less than 1/60 second or a time interval required to deliver 5 mAs, whichever is greater;

(iii) either the product of peak x-ray tube potential, current and exposure time shall be limited to not more than 60 kWs per exposure or the product of x-ray tube current and exposure time shall be limited to not more than 600 mAs per exposure except when the x-ray tube potential is less than 51 kVp in which case the product of x-ray tube current and exposure time shall be limited to not more than 2,000 mAs per exposure; and

(iv) a visible signal shall indicate when an exposure has been terminated at the limits described in § 175.57(a)(3)(iii) and manual resetting shall be required before further automatically timed exposures can be made.

(c) Accuracy. Deviation of technique factors from indicated values shall not exceed the limits given in the information provided by the manufacturer in accordance with applicable Federal standards (21 CFR Part 1020, or any successor law or regulation).

(d) Reproducibility. The following requirements shall apply when the equipment is operated on an adequate power supply as specified by the manufacturer in accordance with the requirements of applicable Federal standards (21 CFR Part 1020, or any successor law or regulation):

(1) Coefficient of variation. For any specific combination of selected technique factors, the estimated coefficient of variation of radiation exposures shall be no greater than 0.05.

(e) Linearity. The following requirements apply when the equipment is operated on a power supply as specified by the manufacturer in accordance with the requirements of applicable Federal standards (21 CFR Part 1020 or any successor law or regulation) for any fixed x-ray tube potential within the range of 40 percent to 100 percent of the maximum rated.

(1) Equipment having independent selection of x-ray tube current (mA). The average ratios of exposure to the indicated milliampere-seconds product, C-kg-1/mAs (mR/mAs), obtained at any two consecutive tube current settings shall not differ by more than 0.10 times their sum. This is:

' X1 - X2 ' # 0.10 (X1 + X2) where X1 and X2 are the average C-kg-1/mAs (or mR/mAs) values obtained at each of two consecutive tube current settings or at two settings differing by no more than a factor of 2 when the tube current selection is continuous.

(2) Equipment having selection of x-ray tube current-exposure time product (mAs). For equipment manufactured after May 3, 1994, the average ratios of exposure to the indicated milliampere-seconds product, C-kg-1/mAs (mR/mAs), obtained at any two consecutive mAs selector settings shall not differ by more than 0.10 times their sum. This is: ' X1 - X2 ' # 0.10 (X1 + X2) where X1 and X2 are the average C-kg-1/mAs (or mR/mAs) values obtained at each of two consecutive mAs selector settings or at two settings differing by no more than a factor of 2 where the mAs selector

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provides continuous selection.

(f) Field limitation and alignment for mobile and stationary general purpose x-ray systems. Except where spot-film devices are used, mobile and stationary general purpose radiographic x-ray systems shall meet the following requirements:

(1) Variable x-ray field limitation. A means for stepless adjustment of the size of the x-ray field shall be provided. Each dimension of the minimum field size at an SID of 100 cm shall be equal to or less than 5 cm.

(2) Visual definition:

(i) Means for visually defining the perimeter of the x-ray field shall be provided. The total misalignment of the edges of the visually defined field with the respective edges of the x-ray field along either the length or width of the visually defined field shall not exceed 2 percent of the distance from the source of the center of the visually defined field when the surface upon which it appears is perpendicular to the axis of the x-ray beam.

(ii) When a light localizer is used to define the x-ray field, it shall provide an average illuminance of not less than 160 lux (15 footcandles) at 100 cm or at the maximum SID, whichever is less. The average illuminance shall be based upon measurements made in the approximate center of each quadrant of the light field. Radiation therapy simulation systems are exempt from this requirement.

(iii) The edge of the light field at 100 cm or at the maximum SID, whichever is less, shall have a contrast ratio, corrected for ambient lighting, of not less than 4 in the case of beam-limiting devices designed for use on stationary equipment, and a contrast ratio of not less than 3 in case of beam-limiting devices designed for use on mobile equipment. The contrast ratio is defined as I1/I2 where I1 is the illuminance 3 mm from the edge of the light field toward the center of the field and I2 is the illuminance 3 mm from the edge of the light field away from the center of the field.

(g) Field indication and alignment on stationary general purpose x-ray equipment. Except when spot-film devices are used, stationary general purpose x-ray systems shall meet the following requirements in addition to those prescribed in § 175.57(e):

(1) Means shall be provided to indicate when the axis of the x-ray beam is perpendicular to the plane of the image receptor, to align the center of the x-ray field with respect to the center of the image receptor to within 2 percent of the SID, and to indicate the SID to within 2 percent;

(2) The beam-limiting device shall numerically indicate the field size in the plane of the image receptor to which it is adjusted;

(3) Indication of field size dimensions and SIDs shall be specified in centimeters and/or inches, and shall be such that aperture adjustments result in x-ray field dimensions in the plane of the image receptor which correspond to those indi-

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cated by the beam-limiting device to within 2 percent of the SID when the beam axis is indicated to be perpendicular to the plane of the image receptor; and

(h) Field limitation on radiographic x-ray equipment other than general purpose radiographic systems. (1) X-ray systems designed for one image receptor size. Radiographic equipment designed for only one image receptor size at a fixed SID shall be provided with means to limit the field at the plane of the image receptor to dimensions no greater than those of the image receptor, and to align the center of the x-ray field with the center of the image receptor to within 2 percent of the SID or shall be provided with means to both size and align the x-ray field such that the x-ray field at the plane of the image receptor does not extend beyond any edge of the image receptor.

(2) Other x-ray systems. Radiographic systems not specifically covered in § 175.57(f), (g), (h)(1), and (j) and systems covered in § 175.58(b), which are also designed for use with extraoral image receptors and when used with an extraoral image receptor, shall be provided with means to limit the x-ray field in the plane of the image receptor so that such field does not exceed each dimension of the image receptor by more than 2 percent of the SID, when the axis of the x-ray beam is perpendicular to the plane of the image receptor. In addition, means shall be provided to align the center of the x-ray field such that the x-ray field at the plane of the image receptor does not extend beyond any edge of the image receptor. These requirements may be met with:

(i) a system which performs in accordance with § 175.57(f) and (g); or, when alignment means are also provided, may be met with either:

(ii) an assortment of removable fixed-aperture, beam-limiting devices sufficient to meet the requirement for each combination of image receptor size and SID for which the unit is designed. Each such device shall have clear and permanent markings to indicate the image receptor size and SID for which it is designed; or

(iii) a beam-limiting device having multiple fixed apertures sufficient to meet the requirement for each combination of image receptor size and SID for which the unit is designed. Permanent, clearly legible markings shall indicate the image receptor size and SID for which each aperture is designed and shall indicate which aperture is in position for use.

(i) Positive beam limitation (PBL). These requirements apply to radiographic systems which contain PBL.

(1) Field size. When a PBL system is provided, it shall prevent x-ray production when:

(i) either the length or width of the x-ray field in the plane of the image receptor differs from the corresponding image receptor dimension by more than 3 percent of the SID; or

(ii) the sum of the length and width differences as stated in § 175.57(i)(1)(i), without regard to sign exceeds 4 percent of the SID.

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(iii) The beam limiting device is at an SID for which PBL is not designed for sizing.

(2) Conditions for PBL. When provided, the PBL system shall function as described in § 175.57(i)(1) whenever all the following conditions are met:

(i) the image receptor is inserted into a permanently mounted cassette holder;

(ii) the image receptor length and width are less than 50 cm;

(iii) the x-ray beam axis is within +- 3 degrees of vertical and the SID is 90 cm to 130 cm inclusive, or the x-ray beam is within +- 3 degrees of horizontal and the SID is 90 cm to 205 cm inclusive;

(iv) the x-ray beam axis is perpendicular to the plane of the image receptor to within +- 3 degrees; and

(v) neither tomographic nor stereoscopic radiography is being performed.

(3) Operator initiated undersizing. The PBL system shall be capable of operation such that, at the discretion of the operator, the size of the field may be made smaller than the size of the image receptor through stepless adjustment of the field size. Each dimension of the minimum field size at an SID of 100 cm shall be equal to, or less than, 5 cm. Return to PBL function as described in § 175.57(i)(1) shall occur automatically upon any change of image receptor size or SID.

(4) Override of PBL. A capability may be provided for overriding PBL in case of system failure and for servicing the system. This override may be for all SIDs and image receptor sizes. A key shall be required for any override capability that is accessible to the operator. It shall not be possible to remove the key while PBL is overridden. Each such key switch or key shall be clearly and durably labeled as follows:

For X-Ray Field Limitation System Failure The override capability is considered accessible to the operator if it is referenced in the operator's manual or in other material intended for the operator or if its location is such that the operator would consider it part of the operational controls.

(j) Field limitation and alignment for spot-film devices. The following requirements shall apply to spot-film devices, except when the spot-film device is provided for use with a radiation therapy simulation system:

(1) Means shall be provided between the source and the patient for adjustment of the x-ray field size in the plane of the image receptor to the size of that portion of the image receptor which has been selected on the spot-film selector. Such adjustment shall be accomplished automatically when the x-ray field size in the plane of the image receptor is greater than the selected portion of the image receptor. If the x-ray field size is less than the size of the selected portion of the image receptor, the field size shall not open automatically to the size of the selected

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portion of the image receptor unless the operator has selected that mode of operation.

(2) Neither the length nor the width of the x-ray field in the plane of the image receptor shall differ from the corresponding dimensions of the selected portion of the image receptor by more than 3 percent of the SID when adjusted for full coverage of the selected portion of the image receptor. The sum, without regard to sign, of the length and width differences shall not exceed 4 percent of the SID. On spot-film devices manufactured after February 25, 1978, if the angle between the plane of the image receptor and beam axis is variable, means shall be provided to indicate when the axis of the x-ray beam is perpendicular to the plane of the image receptor, and compliance shall be determined with the beam axis indicated to be perpendicular to the plane of the image receptor.

(3) The center of the x-ray field in the plane of the image receptor shall be aligned with the center of the selected portion of the image receptor to within 2 percent of the SID.

(4) Means shall be provided to reduce the x-ray field size in the plane of the image receptor to a size smaller than the selection portion of the image receptor such that:

(i) for spot-film devices used on fixed-SID fluoroscopic systems which are not required to, and do not, provide stepless adjustment of the x-ray field, the minimum field size, at the greatest SID, does not exceed 125 cm 2; or

(ii) for spot-film devices used on fluoroscopic systems that have a variable SID and/or stepless adjustment of the field size, the minimum field size, at the greatest SID, shall be containable in a square of 5 cm by 5 cm.

(5) A capability may be provided for overriding the automatic x-ray field size adjustment in case of a system failure. If it is so provided, a signal visible at the fluoroscopist's position shall indicate whenever the automatic x-ray field size adjustment override is engaged. Each such system failure override switch shall be clearly labeled as follows:

For X-Ray Field Limitation System Failure.

(k) Source-skin distance. Mobile and portable x-ray systems other than dental shall be provided with means to limit the source-skin distance to not less than 30 cm.

(l) Beam-on indicators. The x-ray control shall provide visual indication whenever x-rays are produced. In addition, a signal audible to the operator shall indicate that the exposure has terminated.

(m) Multiple tubes. Where two or more radiographic tubes are controlled by one exposure switch, the tube or tubes which have been selected shall be clearly indicated before initiation of the exposure. This indication shall be both on the x-ray control and at or near the tube housing assembly which has been selected.

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(n) Radiation from capacitor energy storage equipment. Radiation emitted from the x-ray tube shall not exceed:

(1) 8.6 E -9 C-kg-1 (0.03 mR) in one minute at 5 cm from any accessible surface of the diagnostic source assembly, with the beam-limiting device fully open, the system fully charged, and the exposure switch, timer, or any discharge mechanism not activated. Compliance shall be determined by measurements averaged over an area of 100 cm 2, with no linear dimension greater than 20 cm; and

(2) 2.58 E -5 C-kg-1 (100 mR) in one (1) hour at 100 cm from the x-ray source, with the beam-limiting device fully open, when the system is discharged through the x-ray tube either manually or automatically by use of a discharge switch or deactivation of the input power. Compliance shall be determined by measurement of the maximum exposure per discharge multiplied by the total number of discharges in one (1) hour (duty cycle). The measurements shall be averaged over an area of 100 cm 2 with no linear dimension greater than 20 cm.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (a) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

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NOTES:

Subsection (a) was amended on March 21, 2001 to remove language which is confusing and which does not appear in state or federal regulations.

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§ 175.58. Dental radiography.

(a) The requirements of this section apply specifically to x-ray equipment and associated facilities used for dental radiography and are in addition to, and not in substitution for, other requirements of this Code.

(b) Non-certified dental units shall meet the equipment and use conditions outlined in § 175.58(b)(1) through (8), with exceptions as noted. Certified equipment shall meet the conditions in § 175.58(b)(1) through (9).

(1) Source-to-skin distance. X-ray systems designed for use with an intraoral image receptor shall be provided with means to limit source-to-skin distance to not less than 17.8 cm (7 in.) or 10.2 cm (4 in.) for equipment operating at 50 kVp.

(2) Field limitation. (i) Radiographic systems designed for use with an intraoral image receptor shall be provided with means to limit the x-ray field such that the x-ray field at the minimum SSD shall be containable in a circle having a diameter of not more than 7 cm (2.75 in.), except dental x-ray units manufactured before August 1, 1974 in which the x-ray beam shall not exceed 7.6 cm (3 in.) at the minimum SSD.

(ii) The operator shall position the end of the position indicating device (PID) within 1 cm (0.4 in.) of the skin of the patient, if such device is routinely used in conducting dental radiography.

(3) Radiation exposure control. (i) Exposure initiation.

(A) A device shall be provided to terminate the radiation exposure after a preset time interval or exposure. The exposure switch shall be of the dead-man type, and where protective barriers are required shall be so arranged that it cannot be operated outside the shielded area.

(B) An x-ray control shall be incorporated into each x-ray system such that an exposure can be terminated by the operator at any time, except for exposures of 0.5 sec or less.

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(C) Termination of an exposure shall cause automatic resetting of the timer to its initial setting or to "zero."

(D) A timer setting beyond the necessary exposure time shall not be used.

(ii) Exposure indication. The x-ray control shall provide visual indication observable at or from the operator's protected position whenever x-rays are produced. A signal audible to the operator shall indicate that the exposure has terminated on dental x-ray units manufactured after August 1, 1974.

(4) Timer reproducibility. With a timer setting of 0.5 sec or less, the difference between the maximum exposure time (Tmax) and the minimum exposure time (Tmin) shall be less than or equal to 10 percent of the average exposure time (T), when four timing tests are performed:

(Tmax - Tmin) # 0.10 T.

(5) Accuracy. Deviation of technique factors from indicated values shall not exceed the limits specified for that system by its manufacturer. In the absence of manufacturer's specifications, the deviation shall not exceed 10% of the indicated value.

(6) Kilovolts peak limitations. Dental x-ray machines with a nominal fixed kVp of less than 50 kVp shall not be used to make diagnostic dental radiographs of humans.

(7) Conditions for operation of equipment. (i) Patient film holding devices shall be used when techniques permit. The x-ray system shall always be operated in such a manner that the useful beam at the patient's skin does not exceed the requirements of § 175.58(b)(2).

(ii) The tube housing and the PID shall not be hand-held during an exposure.

(iii) Dental fluoroscopy shall not be used.

(iv) Time-temperature techniques or automatic processing shall be used to develop pre-operative diagnostic dental x-ray films. Processing techniques shall be consistent with those recommended by the x-ray film manufacturer. Sight developing of dental radiographs is prohibited except for films taken during operative procedures.

(v) Dental x-ray exposure technique factors and dental processing conditions shall yield entrance skin exposure (ESE) values for the bitewing x-ray projection that are identical to the range of ESE values for dental "D" and "E" speed film designations as published in HHS document #FDA-85-8245 (August 1985) or superseding documents.

(vi) Dental intraoral x-ray radiography shall be conducted with dental film classified American National Standards Institute (ANSI) speed group "D" or faster.

(vii) The tube head shall remain stationary when placed in the clinical exposure

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position.

(viii) All x-ray units manufactured before August 1, 1974 shall be equipped with electronic means (timers) for exposure control not later than January 1, 1997.

(ix) Persons not required for the dental x-ray procedure shall not be present in the dental x-ray room.

(8) For extraoral dental radiography, the x-ray film used as the recording medium during the x-ray examination shall show substantial evidence of cut-off (beam delineation).

(9) Additional requirements applicable only to certified systems. Certified dental x-ray units shall meet the following additional requirements:

(i) Reproducibility. When the equipment is operated on an adequate power supply as specified by the manufacturer, the estimated coefficient of variation of radiation exposures shall be no greater than 0.05 for any specific combination of selected technique factors.

(ii) Linearity. When the equipment allows a choice of x-ray tube current settings and is operated on a power supply as specified by the manufacturer in accordance with the requirements of applicable federal standards, for any fixed x-ray tube potential within the range of 40-100 percent of the maximum rating, the average ratios of exposure to the indicated mAs product obtained at any two consecutive tube current settings shall not differ by more than 0.10 times their sum:

(X1 - X2) # 0.10 (X1 + X2) where X1 and X2 are the average mR/mAs values obtained at each of two consecutive tube current settings.

(iii) Accuracy. Deviation of technique factors from indicated values shall not exceed the limits specified for that system by its manufacturer.

(iv) Beam quality. All certified dental x-ray systems manufactured on or after December 1, 1980 shall have a minimum half-value layer not less than 1.5 mm aluminum equivalent. Systems operating above 70 kVp are subject to the filtration requirements of § 175.56(f).

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.59. Podiatric radiography.

(a) Equipment. (1) Collimating devices capable of restricting the useful beam to the area of clinical interest shall be used.

(2) The x-ray films used as the recording medium during the x-ray examination shall show substantial evidence of cut-off (beam delineation).

(3) A device shall be provided which terminates the exposure after a preset time interval or exposure. The exposure switch shall be of the dead-man type and where protective barriers are required shall be so arranged that it cannot be operated outside the shielded area.

(4) Each installation shall be arranged so that the operator can stand at least two (2) meters (6 feet) from the patient, the x-ray tube and the useful beam during exposure. A protective barrier shall be provided when the operator cannot stand at least 2 meters (6 feet) away from the patient, the x-ray tube and the useful beam during exposures.

(b) Conditions for operation of equipment. (1) No person shall hold film during the exposure.

(2) Only persons required for the radiographic procedure shall be in the radiographic room during exposure.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.60. Fixed radiography (excluding dental, veterinary and podiatric radiography).

(a) Equipment. (1) Collimating devices capable of restricting the useful beam to the area of clinical interest shall be used.

(2) The x-ray films used as recording medium during the x-ray examination shall show substantial evidence of cut-off (beam delineation). In addition, general purpose equipment, exclusive of spot-film devices, shall be equipped with adjustable collimators with a means of visually defining the entire field. The total misalignment of the visually defined field with the x-ray field, along either the length or width dimensions, shall not exceed 2 percent of the SID. Such devices shall be calibrated in terms of the size of the projected useful beam at specified source-film distances with the same degree of accuracy.

(3) The technique factors to be used during an exposure shall be indicated before the exposure begins, except when automatic exposure controls are used, in which case the technique factors which are set prior to the exposure shall be indicated. On equipment having fixed technique factors, this requirement may be met by permanent markings. Indication of technique factors shall be visible from the operator's position except in the case of spot films made by the fluoroscopist.

(4) A device shall be provided which terminates the exposure after a preset time interval or exposure.

(5) A dead-man type of exposure switch shall be used and so arranged such that it cannot be operated outside a shielded area. Exposure switches for spot-film devices used in conjunction with fluoroscopic equipment are excepted from this shielding requirement.

(b) Conditions for operation of equipment. (1) No person shall be regularly employed to hold patients or films during exposures nor shall such duty be performed by an individual occupationally exposed to radiation in the course of that individual's other duties. When it is necessary to restrain the patient, mechanical supporting or restraining devices should be used. If patients or films must be held by

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an individual, that individual shall be protected with appropriate shielding devices such as protective gloves and a protective apron of at least 0.25 mm lead equivalent. No part of the holding individual's body shall be in the useful beam. The exposure of any individual used for holding patients shall be monitored. Pregnant women and individuals under 18 years of age shall not hold patients under any conditions.

(2) Only persons required for the radiographic procedure shall be in the radiographic room during the exposure and, except for the patient, all such persons shall be equipped with appropriate shielding devices such as protective gloves and a protective apron of at least 0.25 mm lead equivalent.

(3) For patients who have not passed the reproductive age, gonadal shielding of not less than 0.5 mm lead equivalent shall be used during radiographic procedures in which the gonads are in the useful beam, except for cases in which this would interfere with the diagnostic procedure.

HISTORICAL NOTE

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§ 175.61. Portable, bedside or mobile equipment (excluding dental, veterinary and podiatric radiography).

(a) Equipment. (1) Collimating devices capable of restricting the useful beam to the area of clinical interest shall be used.

(2) The x-ray films used as the recording medium during the x-ray examination shall show substantial evidence of cut-off (beam delineation). General purpose equipment, exclusive of portable equipment, shall be equipped with adjustable collimators with a means of visually defining the entire field. The total misalignment of the visually defined field with the x-ray field, along either the length or width dimensions, shall not exceed 2 percent of the SID. Such devices shall be calibrated in terms of the size of the projected useful beam at specified source-film distances with the same degree of accuracy.

(3) A device shall be provided which terminates the exposure after a preset time interval or exposure.

(4) All mobile, portable or bedside equipment shall be provided with cones or metal frames so that the minimum source-to-skin distance is at least 31 cm (12 in.).

(b) Conditions for operation of equipment. (1) No person shall be regularly employed to hold patients or films during exposures, nor shall such duty be performed by an individual occupationally exposed to radiation in the course of that individual's other duties. When it is necessary to restrain the patient, mechanical supporting or restraining devices should be used. If patient or films must be held by an individual, that individual shall be protected with appropriate shielding devices such as protective gloves and a protective apron of at least 0.25 mm lead equivalent. No part of the holding individual's body shall be in the useful beam. The exposure of any individual used for holding patients shall be monitored. Pregnant women and individuals under 18 years of age shall not hold patients under any conditions.

(2) For patients who have not passed the reproductive age, gonadal shielding of

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not less than 0.5 mm lead equivalent shall be used during radiographic procedures in which the gonads are in the useful beam, except for cases in which this would interfere with the diagnostic procedure.

(3) Personnel monitoring shall be required for all persons operating mobile or portable x-ray equipment.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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§ 175.62. Fluoroscopy.

Fluoroscopic systems and associated components shall meet the following requirements:

(a) Primary protective barrier. (1) Limitation of useful beam. The entire cross-section of the useful beam shall be intercepted by the primary protective barrier of the fluoroscopic image assembly irrespective of position. The fluoroscopic tube shall not produce x-rays unless the barrier is in position to intercept the entire useful beam. The exposure rate due to transmission through the barrier with the attenuation block in the useful beam combined with radiation from the image intensifier, if provided, shall not exceed 5.16 E-7 C-kg-1 (2 milliroentgens) per hour at 10 cm (4 in.) from any accessible surface of the fluoroscopic imaging assembly beyond the plane of the image receptor for each roentgen per minute of entrance exposure rate.

(b) Field limitation. (1) Image-intensified fluoroscopy. For image-intensified fluoroscopic equipment the total misalignment of the edges of the x-ray field with the respective edges of the visible area of the image receptor along any dimension of the visually defined field in the plane of the image receptor shall not exceed 3 percent of the SID. The sum, without regard to sign, of the misalignment along any two orthogonal dimensions intersecting at the center of the visible area of the image receptor shall not exceed 4 percent of the SID. For rectangular x-ray fields used with circular image receptors, the error in alignment shall be determined along the length and width dimensions of the x-ray field which pass through the center of the visible area of the image receptor. Means shall be provided to permit further limitation of the field. The minimum field size, at the greatest SID, shall be equal to or less than 5 by 5 cm (2 by 2 in.).

(c) Activation of tube. X-ray production in the fluoroscopic mode shall be controlled by a device which requires continuous pressure by the operator for the entire time of any exposure. When recording serial fluoroscopic images, the operator shall be able to terminate the x-ray exposure(s) at any time, but means may be provided to permit completion of any single exposure of the series in process.

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(d) Entrance exposure rate limits.

(1) The fluoroscopic exposure rate when measured under the following conditions shall not exceed 5 Roentgens per minute:

(i) the controls are set to the dose rate mode used for the fluoroscopic procedure most commonly performed on that fluoroscopic unit; and

(ii) the image intensifier is set to the largest field of view; and

(iii) the image intensifier is at 12 inches (30 cm) above the tabletop or the over table fluoro tube is at a source to image distance normally used for an average patient; and

(iv) a patient phantom composed of 1 and 1/2 inch (3.8 cm) thickness of Type 1100 aluminum and 0.02 inch (0.5 mm) thickness of copper or an equivalent device is completely intercepting the useful beam; and

(v) the measurement is made at the measurement location specified in 21 CFR § 1020.32(d)(3).

(vi) If the exposure rate cannot be measured, the exposure integrated for one minute under the same conditions as subparagraph (i) or paragraph (7) shall not exceed 5 Roentgens.

(2) The maximum exposure rate measured in air shall not exceed 10 Roentgens per minute, when measured in the manner as specified in 21 CFR § 1020.32(d)(3), except as follows:

(i) Equipment manufactured before May 19, 1995 and certified in accordance with 21 CFR Part 1020 and having an optional high level control is limited to a maximum output of 5 Roentgens per minute unless the high level control is activated and an audible signal to that effect is provided. When the high level is activated, the maximum exposure rate measured in air at a point where the center of the useful beam enters the patient shall not exceed 20 Roentgens per minute.

(ii) Certified equipment manufactured after May 19, 1995 with automatic exposure rate and having an optional high level control is limited to a maximum output of 10 Roentgens per minute unless the high level control is activated and an audible signal to that effect is provided. When the high level control is activated, the maximum exposure rate measured in air at a point where the center of the useful beam enters the patient shall not exceed 20 Roentgens per minute.

(iii) Certified equipment manufactured after May 19, 1995 without automatic exposure rate is limited to 5 Roentgens per minute unless the high level control is activated and an audible signal to that effect is provided. When the high level control is activated, the maximum exposure rate measured in air at a point where the center of the useful beam enters the patient shall not exceed 20 Roentgens per minute.

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(3) With the system configured for the most frequently performed fluoroscopic procedure, exposure rates shall be measured with each of the following attenuators in the beam:

(i) 0.75 inches (19 mm) of aluminum (pediatric patient--25 kg.),

(ii) 1.50 inches (38 mm) of aluminum (small adult patient--50 kg.),

(iii) 1.50 inches (38 mm) of aluminum and 0.02 inches (0.5 mm) of copper (average adult patient--75 kg.),

(iv) 1.50 inches (38 mm) of aluminum and 0.08 inches (2.0 mm) of copper (large adult patient--100 kg.),

(v) 1.50 inches (38 mm) of aluminum and 0.08 inches (2.0 mm) of copper and 0.12 inches (3.0 mm) of lead (for maximum fluoroscopic exposure rate only).

The fluoroscopic exposure rates for the most frequently performed procedure shall be posted so that they are conspicuous to the operator.

(e) Indication of potential and current. During fluoroscopy and cinefluorography, x-ray tube potential and current shall be continuously indicated. Deviation of x-ray tube potential and current from the indicated values shall not exceed the maximum deviation as stated by the manufacturer in accordance with applicable Federal standards (21 CFR Part 1020 or successor regulations).

(f) Source-skin distance. Means shall be provided to limit the source-skin distance to not less than 38 cm (15 in.) on stationary fluoroscopes and to not less than 30 cm (12 in.) on mobile fluoroscopes. In addition, for image-intensified fluoroscopes intended for specific surgical applications that would be prohibited at the source-skin distances specified in § 175.62(f), provision may be made for operation at shorter source skin distances but in no case less than 20 cm (8 in.). When provided, the manufacturer must set forth precautions with respect to the optional means of spacing in addition to other information as required in accordance with applicable Federal standards.

(g) Fluoroscopic timer. Fluoroscopy equipment shall not be operated for medical use unless a means is provided to preset the cumulative on-time of the fluoroscopic tube. The maximum cumulative time of the timing device shall not exceed 5 minutes without resetting. A signal audible to the fluoroscopist shall indicate the completion of any present cumulative on-time. Such signal shall continue to sound while x-rays are produced until the timing device is reset.

(h) Contrast tests. (1) The spatial resolution of the fluoroscopic system shall be measured using a test tool composed of a line pair (lp) plate with discreet line pair groups and a maximum lead foil thickness of 0.1 mm or an equivalent device. The test tool shall be placed on a 0.75 inch (19 mm) thickness of type 1100 aluminum, large enough to completely intercept the useful beam, with the test tool 12 inches (30 cm) from the entrance surface of the image receptor assembly. If the system has variable source-to-image distance (SID), the measurement SID shall not

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exceed 40 inches (100 cm). The image receptor of the fluoroscopic system shall be operated in the six inches (15 cm) field of view (FOV) to conduct this test. If six inches (15 cm) FOV is not available, the system shall be operated in the smallest FOV that exceeds the six inches (15 cm) FOV. The minimum spatial resolution at the center of the beam for all FOVs shall be determined by the following equation:

2 lp/mm X (6 inches (15cm)/size of FOV used) = minimum number of lp/mm.

(2) The low contrast performance of the fluoroscopic system shall be capable of resolving a minimum hole size of 3 mm using a test tool composed of a 1.0 mm aluminum sheet with two sets of four holes of dimension 1.0, 3.0, 5.0 and 7.0 mm and a phantom composed of a 1 and 1/2 inch (3.8 cm) thickness of Type 1100 aluminum large enough to completely intercept the useful beam or an equivalent device. The test tool shall be 12 inches (30 cm) from the entrance surface of the image receptor assembly. The image receptor of the fluoroscopic system shall be operated in the six inches (15 cm) FOV to conduct this test. If six inches (15 cm) FOV is not available, the system shall be operated in the smallest FOV that exceeds the six inches (15 cm) FOV.

(i) Conditions for operation of equipment. (1) The operator of any fluoroscopic installation shall determine and record the outputs made pursuant to § 175.62(d) where the center of the useful beam enters the patient during routine fluoroscopy and cinefluorography. The rate shall be determined at least annually, each time a major component is serviced or replaced, when the outputs exceed the limits specified in § 175.62 or when there is a reason to believe that the operating factors have changed significantly. Notwithstanding the requirements of § 175.54, such measurements and measurement of stray radiation to operators and observers, are required when the equipment is first placed in operation.

(2) Protective garments of at least 0.25 mm lead equivalent shall be available and worn by the fluoroscopist during every examination.

(3) Unless measurements indicate that they are not needed, protective gloves and aprons of at least 0.25 mm lead equivalent each shall be worn by any person within the fluoroscopy room.

(4) Only persons needed in the fluoroscopy room shall be present during the exposure.

(j) Exemptions. (1) Fluoroscopic radiation therapy simulation systems are exempt from the requirements of § 175.62(a), (d), and (g) provided that:

(i) the systems are designed and used in a manner such that no individual other than the patient is in the x-ray room when the system is producing x-rays; and

(ii) systems which do not meet the requirements of § 175.62(g) are provided with a means of indicating the cumulative time that an individual patient has been exposed to x-rays. Procedures shall require that the timer be reset between cases.

HISTORICAL NOTE

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Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Open par amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (d) amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

Subd. (d) par (3) added City Record Apr. 3, 2001 eff. May 3, 2001. This par (3) was bracketed out of rule in City Record Sept. 29, 2006 amendment.

Subd. (h) amended City Record Sept. 29, 2006 eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Subdivisions (d) and (h) of § 175.62 were amended on September 26, 2006, to conform with the requirements of the New York State Sanitary Code § 16.58 for fluoroscopic equipment, and FDA performance standards in 21 CFR 1020.32.

The first sentence of § 175.62 was amended and paragraph (3) of subsection (d) was added on March 21, 2001 to include additional requirements for entrance exposure rate limits for fluoroscopic equipment in compliance with state and federal regulations.

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
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NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIATION EQUIPMENT

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.63. Mammography.

(a) Applicability. The provisions of this section apply to all facilities which produce, process or interpret mammograms for screening and/or diagnostic purposes and are in addition to, and not in substitution for, other requirements of this Code.

(b) Requirement for certification. (1) Except for facilities holding provisional certificates as described in § 175.63(b)(2), effective October 1, 1994 each mammography facility shall have received a certificate indicating approval by the U.S. Food and Drug Administration (FDA) to provide screening and diagnostic mammography services pursuant to 21 CFR Section 900.11, or any successor law or regulation.

(2) A provisional certificate issued pursuant to 21 CFR Section 900.11, or any successor law or regulation, will be accepted in lieu of the certificate required by § 175.63(b)(1) for a period of not longer than six (6) months from the date of issuance plus one ninety (90) day extension.

(c) Equipment. (1) Radiographic equipment designed for conventional radiographic procedures that has been modified or equipped with special attachments for mammography shall not be used for mammography.

(2) Radiographic equipment used for mammography shall:

(i) be certified by the U.S. Food and Drug Administration pursuant to 21 CFR 1010.2, or any successor law or regulation, as meeting the applicable requirements of 21 CFR 1020.30 and 21 CFR 1020.31 in effect on the date of manufacture; and

(ii) be specifically designed for mammography; and

(iii) incorporate a breast compression device; and

(iv) have the provision for operating with a removable grid, except for xeromammography systems.

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(3) Beam quality for mammographic systems with a molybdenum target--molybdenum filter combination.

(i) When used with screen-film image receptors, and the contribution to filtration made by the compression device is included, the useful beam shall have a half-value layer (HVL) between the values of measured kVp/100 and measured kVp/100 + 0.1 mm aluminum.

(ii) For xeromammography, the HVL of the useful beam with the compression device in place shall be at least 1.0 and not greater than 1.6 mm aluminum, measured at 49 kVp with a tungsten target tube.

(d) Dose. (1) The average glandular dose delivered during a single cranio-caudal view of an approved phantom simulating a 4.5 cm thick, compressed breast consisting of 50 percent adipose and 50 percent glandular tissue, shall not exceed:

(i) 1 mGy (100 millirads) per exposure for screen-film mammography procedures without a grid;

(ii) 3 mGy (300 millirads) per exposure for screen-film mammography procedures with a grid; or

(iii) 4 mGy (400 millirads) per exposure for xeromammography procedures.

The dose shall be determined at least annually using the technique factors and conditions that are used to produce the phantom images submitted for accreditation.

(e) Personnel. The following requirements apply to personnel involved in any aspect of mammography, including the production, processing and interpretation of mammograms and related quality assurance activities.

(1) Interpreting physicians shall meet the following requirements:

(i) be licensed to practice medicine in the State of New York; and

(ii) have had the following training:

(A) be certified in an accepted speciality area by one of the bodies approved by FDA to certify interpreting physicians; or

(B) have had at least two (2) months of documented full-time training in the interpretation of mammograms, including instruction in radiation physics, radiation effects and radiation protection; and

(C) have forty (40) hours of documented continuing medical education in mammography. Time spent in residency specifically devoted to mammography is acceptable, if documented in writing by a fully qualified interpreting physician; and

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(iii) have had the following initial experience:

(A) have read and interpreted the mammograms from the examinations of at least 240 patients in the six (6) months preceding application; or

(B) read and interpret mammograms as specified in § 175.63(e)(1)(iii)(A) under the direct supervision of a fully qualified interpreting physician; and

(iv) have the following continuing experience:

(A) continue to read and interpret mammograms from the examination of an average of at least 40 patients per month over 24 months; and

(B) continue to participate in education programs, either by teaching or completing an average of at least five (5) continuing medical education credits in mammography per year.

(2) Radiologic technologists shall meet the following requirements:

(i) have a New York State license to perform radiographic procedures; and

(ii) have satisfied the requirements set forth in 21 C.F.R. § 900.12(a)(2), or its successor regulation.

(3) Medical physicists shall meet the following requirements:

(i) have approval by the Department to conduct evaluations of mammography equipment and procedures required under the Public Health Service Act; or

(ii) be certified in an accepted speciality area by one of the bodies approved by FDA to certify medical physicists; or

(iii) for those medical physicists associated with facilities applying for accreditation before October 27, 1997, meet the following criteria:

(A) have a masters, or higher, degree in physics, radiological physics, applied physics, biophysics, health physics, medical physics, engineering, radiation science, or in public health with a bachelor's degree in the physical sciences; and

(B) have one (1) year of training in medical physics specific to diagnostic radiological physics; and

(C) have two (2) years of experience in conducting performance evaluation of mammography equipment; and

(iv) participate in continuing education programs related to mammography, either by teaching or completing an average of at least five (5) continuing education units per year.

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(f) Quality assurance. (1) Each registrant performing mammography examinations shall establish and maintain a quality assurance program to assure the adequate performance of the radiographic equipment and other equipment and materials used in conjunction with such equipment to ensure the reliability and clarity of its mammograms. The program shall also require periodic monitoring of the dose delivered by the facility's examination procedures to ensure that it does not exceed the limit specified in § 175.63(d) and is appropriate for the image receptor used.

(2) For screen-film systems, the mammography quality assurance program required by § 175.63(f)(1) shall be substantially the same as that described in the 1992 edition of "Mammography Quality Control: Radiologist's Manual, Radiologic Technologist's Manual and Medical Physicist's Manual," prepared by the American College of Radiology, Committee on Quality Assurance in Mammography, or in any superseding document.

(3) For systems with alternate image receptor modalities, the mammography quality assurance program required by § 175.63(f)(1) shall be substantially the same as the quality assurance program recommended by the image receptor manufacturer, which, if followed, will allow a facility to maintain high image quality.

(4) The mammography quality assurance program required by § 175.63(f)(1) shall provide for the maintenance of log books documenting compliance with the requirements of § 175.63(f)(1) through (3) and recording corrective actions taken.

(5) Prior to performing patient mammography, using a breast equivalent phantom specified in § 175.63(f)(7)(i), (ii) or (iii), the registrant shall optimize the mammographic system and determine image resolution, which shall be the reference image resolution for the mammographic system.

(6) The registrant shall use the breast equivalent phantom used to satisfy the requirement of § 175.63(f)(5) to test image resolution of the mammographic system at monthly intervals; mobile and portable equipment shall be so tested each time the unit is moved or at monthly intervals, whichever is less.

(7) No patient mammogram shall be performed unless the mammographic system is capable of imaging, using phantom objects as follows:

(i) the phantom image shall achieve at least the minimum score established by an accreditation body approved by the FDA in accordance with 21 C.F.R. § 900.3(d) or 900.4(a)(8), or successor regulations; or

(ii) the equivalent test object resolution on another phantom approved by the Department.

(8) Diminished phantom test object resolution and facility follow-up.

(i) If, when tested pursuant to § 175.63(f)(6), the mammographic system detects two (or more) fewer test objects than the reference resolution image made pursuant to § 175.63(f)(5), the cause of resolution loss shall be determined and corrected and the mammographic system re-optimized pursuant to § 175.63(f)(5).

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(ii) If the imaging system resolves less than seven (7) test objects in the phantom, in addition to the requirements of § 175.63(f)(8)(i), there shall be:

(A) a review of monthly phantom images to determine at which point the image resolution fell below the minimum specified in § 175.63(f)(7); and

(B) a review, by a physician(s) not from the facility who is approved by the Department and meets the requirements of § 175.63(e)(1), to determine the diagnostic quality of the mammographic images.

(iii) The review required by § 175.63(f)(8)(ii)(B) shall include:

(A) images from the range of studies performed by the facility which such physician(s) ascertains to be sufficient to determine that the clinical images are of diagnostic quality; and

(B) images from the time interval from when such review is required to the date when the system met the requirements of § 175.63(f)(7).

(iv) If film images reviewed by the physician(s) pursuant to § 175.63(f)(8)(iii)(B) are identified as not being of diagnostic quality, the facility shall, within five (5) business days, notify:

(A) the referring physician or other authorized referring practitioner, or the patient, if not referred by a practitioner, of the need for follow-up; and

(B) the Department of the results of the investigation and follow-up contacts.

(v) A record of the reviews and findings made pursuant to § 175.63(f)(8)(ii)(B) shall be maintained by the registrant at the facility for review by the Department.

(vi) A record of the results of investigations and actions taken to correct any deficiency pursuant to this section shall be maintained for review by the Department for three (3) years.

(9) Each facility shall establish and maintain a clinical image quality control program, including at a minimum:

(i) monitoring of mammograms repeated due to poor image quality; and

(ii) maintenance of records, analysis of results and a description of any remedial action taken on the basis of such monitoring.

(10) Each facility shall establish a system for reviewing outcome date from all mammography performed, including follow-up on the disposition of positive mammograms and correlation of surgical biopsy results with mammogram reports.

(11) As part of its overall quality assurance program, each facility shall have a

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physicist with the qualifications specified in § 175.63(e)(3) establish, monitor and direct the procedures required by § 175.63(f) and perform a survey of the facility to assure that it meets the quality control and equipment standards as specified in § 175.63(c)(2). Such surveys shall be performed at least annually and reports of such surveys shall be prepared and submitted to the body which accredited the facility. Each such report shall be retained by the facility for inspection by the Department.

(g) Medical records. (1) Each facility shall maintain mammograms and associated records in a permanent medical record of the patient as follows:

(i) for a period of not less than five (5) years or not less than ten (10) years, if no additional mammograms of the patient are performed at the facility, or longer as mandated by any applicable law or regulation; or

(ii) until requested by the patient to permanently transfer the records to a medical institution, to a physician of the patient or to the patient, and the records are so transferred.

(2) Each facility shall prepare a written report of the results of any mammography examination. Such report shall be completed as soon as reasonably possible and shall:

(i) be signed by the interpreting physician; and

(ii) be provided to the patient's physician(s) (if any); or

(A) if the patient's physician is not available or if the patient does not have a physician, the report shall be sent directly to the patient; and

(B) if such report is sent to the patient, it shall include a summary written in language easily understood by a lay person; and

(iii) be maintained in the patient's record pursuant to § 175.63(g)(1).

(h) Revocation of accreditation and accrediting body approval. (1) If a facility's accreditation is revoked by an accrediting body (as defined in 21 CFR Section 900.2), the facility's certificate (as defined in 21 CFR Section 900.2) shall remain in effect until such time as determined by the FDA or other certifying body on a case-by-case basis after an investigation into the reasons for the revocation. If the FDA or other certifying body determines that the revocation was justified by violations of applicable quality standards, the FDA or other certifying body will suspend or revoke the facility's certificate and/or require the submission and implementation of a corrective action plan, whichever action will protect the public health in the least burdensome way.

(2) If the approval of an accrediting body is revoked by FDA, the certificates of the facilities accredited by such body shall remain in effect for a period of one (1) year after the date of such revocation subject to FDA's determination that the facility is continuing to perform mammography of acceptable quality. The facility

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must obtain accreditation from an approved accrediting body within one (1) year of the date of revocation.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (e) par (2) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Subd. (f) par (7) amended City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

Subparagraph (ii) of paragraph (2) of subsection (e) was amended and subparagraphs (iii) and (iv) were repealed on March 21, 2001 to amend the requirements for radiologic technologists in accordance with federal regulations.

Subparagraph (i) was amended; subparagraph (ii) was repealed; and subparagraph (iii) was relettered to be known as subparagraph (ii) of paragraph (7) of subsection (f) on March 21, 2001 to reflect federal regulations now in place for testing the minimum imaging quality of equipment prior to performing patient mammograms.

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RADIATION EQUIPMENT

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§ 175.64. Therapeutic radiation machines.

(a) Purpose and scope. (1) This section establishes requirements for the use of radiation machines used for therapeutic purposes in the healing arts to protect public health and safety and for which the registrant is responsible. The requirements of this section are in addition to, and not in substitution for, other applicable provisions of this Code. Except as specifically provided otherwise in this section, the requirements of this section are applicable to all therapeutic radiation machines used in the City of New York for the treatment of humans, regardless of the date of installation.

(b) Certified registration for therapeutic radiation machines. (1) Certified registration required.

(i) Section 175.64(b) applies only to therapeutic radiation machines subject to the requirements of § 175.64(g) of this Code.

(ii) All new facilities with therapeutic radiation machines subject to the requirements of § 175.64(g) of this Code, and all such existing facilities not holding a certified registration on August 1, 1994, shall obtain a certified registration from the Department in accordance with the provisions of this section.

(2) Certified registration application. (i) If the application is for use sited in a medical institution, only the institution's management may apply; for use not sited in a medical institution, any professional practitioner may apply.

(ii) An application for a certified registration shall be filed in duplicate (original plus one copy) on, and shall contain completely and accurately all information called for by, a written form prescribed by the Department.

(iii) When a change affecting a radiation source or installation subject to the certified registration requirements of this section is considered by a registrant, including but not limited to changes ordered pursuant to this Code, so that the information on file with the Department, either in the initial certified registration application, or subsequent request for amendments, or in the initial certified reg-

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istration or amendment previously granted, will no longer be accurate, the registrant shall so inform the Department in writing.

(A) For ministerial changes, such notification shall be within 10 days of effecting such change.

(B) For all other changes, an amendment must be requested and received pursuant to § 175.64(b)(2)(xiv).

(C) Failure to notify the Department of a change of ownership or address of a radiation installation may result in revocation of the installation's certified registration under § 175.64(b)(2)(xii).

(iv) At any time subsequent to the filing of an application for a certified registration and before the termination of a certified registration issued in response thereto, the Department may require the applicant to submit supplementary statements containing additional information to enable the Department to determine whether such application should be approved or denied, or whether a previously issued certified registration should be amended, suspended or revoked.

(v) Each application or supplementary statement shall be signed by either the applicant personally or a person duly authorized by the applicant to sign for and on the applicant's behalf.

(vi) The Department will approve an application for, and issue in response thereto, a certified registration if the Department determines that the following requirements have been met:

(A) the applicant's proposed use, equipment, facilities and procedures will protect health and safety and will minimize danger to life and property from radiation hazards; and

(B) the applicant's instrumentation is appropriate for detecting and measuring the type(s) of radiation produced (either directly or indirectly) by the radiation source requested in the application; and

(C) the applicant, or the applicant's personnel, if the applicant is not an individual, is qualified by training and experience to use such radiation source for the purposes covered by the application so as to protect public health and safety and to minimize danger to life and property from radiation hazards; and

(D) the applicant submits sufficient information to support a determination that the requirements of § 175.64(b)(2)(vi)(A), (B) and (C) are met.

(vii) Certified registrations issued by the Department shall be in the form of a written authorization permitting possession and use of radiation therapy machines capable of operating at 500 kV (photons) and/or 500 keV (electrons) or higher. Such possession and use provided for in the foregoing shall be subject to the requirements of:

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(A) all applicable provisions of this Code; and

(B) all conditions as stated on the certified registration issued by the Department.

(viii) Except as otherwise provided in this Code, each certified registration shall expire on the expiration date stated on the certified registration. If any registrant duly files with the Department an application in proper form for renewal of such certified registration, or for a new and superseding certified registration, not less than 30 days prior to the stated expiration date, such certified registration shall not be deemed to have expired until the Department has finally determined such application.

(ix) The Department may terminate any certified registration upon the written request of the registrant.

(x) The Department may at any time set forth in any certified registration or incorporate by reference therein, additional conditions, restrictions or requirements applicable to the registrant's transfer, receipt, possession or use of any radiation source covered by such certified registration in order to protect the public health and safety and to minimize danger to life and property from radiation hazards.

(xi) Any certified registration may be amended or revoked by the Department by reason of amendment of this Code, or any other applicable law or regulation.

(xii) Any certified registration may be suspended or revoked by the Department for:

(A) any material misstatement in the application therefor or in any supplementary statement thereto;

(B) any condition revealed by such application, supplementary statement, report, record, inspection or other means, which would warrant the Department's refusal to grant a certified registration on an original application; or

(C) any violation or failure to observe any condition of such certified registration, this Code, or any other applicable rule, regulation or law now or hereafter in effect.

(xiii) Any application by a registrant for the renewal of a certified registration, including amendments, shall be considered as an application for a certified registration and shall be filed on, and shall contain completely and accurately all information called for by, a written form or other manner prescribed by the Department. In considering any such application for renewal, the Department will apply the requirements set forth in § 175.64(b)(2)(vi).

(xiv) A registrant shall apply for, and shall have received approval for, a certified registration amendment before:

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(A) permitting anyone to work as an authorized user under the certified registration;

(B) permanently changing the radiation safety officer or radiotherapy physicist;

(C) making any change in the treatment room shielding;

(D) making any change in the location of the therapeutic radiation machine within the treatment room;

(E) using the therapeutic radiation machine in a manner that could result in increased radiation levels in areas outside the treatment room;

(F) relocating the therapeutic radiation machine;

(G) allowing an individual not listed on the registrant's certified registration to perform the duties of the radiation therapy physicist, except during the temporary absence of the radiation therapy physicist when a person who is otherwise qualified to perform such duties may perform such duties. Such temporary absence shall not exceed sixty (60) days; or

(H) before changing non-ministerial statements, representations and/or procedures incorporated by reference into the certified registration.

Any application by a registrant for an amendment of a certified registration shall be filed in writing with the Department and shall set forth in detail the reasons for such requested amendment. In considering any such application for amendment, the Department will apply the requirements set forth in § 175.64(b)(2)(vi).

(c) Training requirements. (1) The registrant of any therapeutic radiation machine shall require the authorized user to be a physician who:

(i) is certified in:

(A) Radiology, therapeutic radiology or radiation oncology by the American Board of Radiology; or

(B) Radiation oncology by the American Osteopathic Board of Radiology; or

(C) Radiology, with specialization in radiotherapy, as a British "Fellow of the Faculty of Radiology" or "Fellow of the Royal College of Radiology"; or

(D) Therapeutic radiology by the Canadian Royal College of Physicians and Surgeons; or

(ii) is in the active practice of therapeutic radiology, and has completed 200 hours of instruction in basic radiation techniques applicable to the use of an external beam radiation therapy unit, 500 hours of supervised work experience, and a

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minimum of three (3) years of supervised clinical experience.

(A) To satisfy the requirement for instruction, the classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radiation; and

(d) radiation biology.

(B) To satisfy the requirement for supervised work experience, training shall be under the supervision of an authorized user at an institution and shall include:

(a) review of the full calibration measurements and periodic quality assurance checks;

(b) preparing treatment plans and calculating treatment times;

(c) using administrative controls to prevent misadministrations;

(d) implementing emergency procedures to be followed in the event of the abnormal operation of an external beam radiation therapy unit or its console; and

(e) checking and using survey meters.

(C) To satisfy the requirement for a period of supervised clinical experience, training shall include one (1) year in a formal training program approved by the Residency Review Committee for Radiology of the Accreditation Council for Graduate Medical Education or the Committee on Postdoctoral Training of the American Osteopathic Association and an additional two (2) years of clinical experience in therapeutic radiology under the supervision of an authorized user at a medical institution. The supervised clinical experience shall include:

(a) examining individuals and reviewing their case histories to determine their suitability for external beam radiotherapy treatment, and any limitations or contraindications;

(b) selecting the proper dose and how it is to be administered;

(c) calculating the external beam radiotherapy doses and collaborating with the authorized user in the review of patients' progress and consideration of the need to modify originally prescribed doses as warranted by patients' reaction to radiation; and

(d) post-administration followup and review of case histories.

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(2) The registrant of any therapeutic radiation machine shall require the radiation therapy physicist to:

(i) be certified by the American Board of Radiology in:

(A) Therapeutic radiological physics; or

(B) Roentgen ray and gamma ray physics; or

(C) X-ray and radium physics; or

(D) Radiological physics; or

(ii) be certified by the American Board of Medical Physics in Radiation Oncology Physics; or

(iii) be certified by the Canadian College of Medical Physics; or

(iv) hold a master's or doctor's degree in physics, biophysics, radiological physics, or health physics, and have completed one (1) year of full time training in therapeutic radiological physics and an additional year of full time work experience under the supervision of a radiotherapy physicist at a medical institution. To meet this requirement, the individual shall have performed the tasks listed in § 175.64(e)(1) and § 175.64(g)(8) and (9) of this Code under the supervision of a radiation therapy physicist during the year of work experience.

(v) Notwithstanding the provision of § 175.64(c)(2)(iv), as of January 1, 2000, the radiation therapy physicist shall be an individual who is certified as described in § 175.64(c)(2)(i), (ii) or (iii).

(3) Qualifications of operators. (i) Any person, other than a qualified physician, who operates a therapeutic radiation machine for medical use shall be a licensed and registered radiation therapist, or a student currently enrolled in an approved program of study in radiation therapy technology and under the direct supervision of a qualified physician or licensed radiation therapist. Such direction or order to apply, or application of, radiation shall be in compliance with all applicable provisions of Title 10 of the New York Code of Rules and Regulations, Part 89, Subchapter L and Article 35 of the Public Health Law of the State of New York, or any successor laws or regulations.

(ii) The names and training of all personnel who currently operate any therapeutic radiation machine(s) shall be kept on file at the facility. Information on former operators shall be retained for a period of three (3) years beyond the last date such a person was authorized to operate a therapeutic radiation machine at the facility.

(4) The training and experience specified in § 175.64(c)(1), (2) or (3) shall have been obtained within the 5 years preceding the date of application or the individual shall demonstrate continuing applicable experience since the required training and experience was completed.

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(d) General administrative requirements. (1) The registrant shall be responsible for directing the operation of any therapeutic radiation machine and shall ensure that the requirements of this section are met in the operation of the therapeutic radiation machine(s).

(2) A therapeutic radiation machine which does not meet the requirements of this Code shall not be used for irradiation of patients.

(3) A physician shall not act as an authorized user for any therapeutic radiation machine which is subject to the provisions of § 175.64(g) until such time as said physician's training has been reviewed and approved by the Department.

(4) Written safety procedures and rules. (i) Written safety procedures and rules, including any restrictions required for the safe operation of a particular therapeutic radiation machine, shall be developed by a radiation therapy physicist. At a minimum, these shall include:

(A) the procedure to be followed to ensure that, except for contact therapy machines, only the patient is in the treatment room before turning the primary beam of radiation on to begin a treatment or after a door interlock interruption; and

(B) the procedure to be followed if the operator is unable to turn the primary beam of radiation off with controls outside the treatment room or in the event that potential harm to the patient or personnel is imminent; and

(C) the names and telephone numbers of the authorized users, radiation safety officer and radiation therapy physicist to be contacted immediately if the therapeutic radiation machine or console operates abnormally.

(ii) Such procedures and rules shall be provided to each individual who operates a therapeutic radiation machine and shall be available at the control console of the therapeutic radiation machine.

(iii) The operator shall be able to demonstrate familiarity with these rules.

(iv) The registrant shall provide instruction in the written safety procedures and rules required by § 175.64(d)(4)(i) to all individuals who operate a therapeutic radiation machine and shall provide appropriate refresher training to such individuals at intervals not to exceed one (1) year. Such instruction shall include "dry runs" of emergency procedures.

(v) The registrant shall retain for three (3) years a record of individuals receiving instruction required by § 175.64(d)(4)(iv), a description of the instruction, the date of instruction, and the name of the individual who gave the instruction.

(5) All individuals associated with the operation of a therapeutic radiation machine shall be instructed in, and shall comply with, the provisions of the registrant's quality assurance program as required by § 175.07 of this Code.

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(6) Individuals shall not be exposed to the useful beam except for medical therapy purposes and unless such exposure has been ordered in writing by an authorized user who meets the training requirements of § 175.64(c). This provision specifically prohibits deliberate exposure of an individual for training, demonstration or other non-healing arts purposes.

(7) For radiation therapy machines installed after August 1, 1994, the registrant shall maintain the following information for each therapeutic radiation machine for inspection by the Department:

(i) report of acceptance testing;

(ii) records of calibrations and periodic spot checks for the therapeutic radiation machine, as well as the name(s) of the person(s) who performed such activities;

(iii) records of major maintenance and modifications performed on the therapeutic radiation machine after August 1, 1994, as well as the name(s) of the person(s) who performed such services; and

(iv) copies of all correspondence with this Department regarding that therapeutic radiation machine.

(8) For each radiation therapy machine, the registrant shall retain records of all surveys for inspection by the Department.

(9) Record retention. (i) Unless specified otherwise, all records required by this section shall be retained until disposal is authorized by the Department. All required records shall be retained in an active file from at least the time of generation until the next Departmental inspection. Any required record generated prior to the last Departmental inspection may be microfilmed or otherwise archived providing a complete copy of such record can be retrieved until such time as the Department authorizes final disposal.

(e) General technical requirements. (1) Protection surveys.

(i) The registrant shall ensure that radiation protection surveys of all new facilities, and existing facilities not previously surveyed, are performed with an operable radiation measurement survey instrument calibrated in accordance with § 175.64(h). The radiation protection survey shall be performed by a radiation therapy physicist or by a physicist certified by the American Board of Health Physics or by the American Board of Radiology, the American Board of Medical Physics or the Canadian College of Medical Physics in the appropriate specialty and shall verify, with the therapeutic radiation machine in a "BEAM-ON" condition with the largest clinically available treatment field and with a scattering phantom in the useful beam of radiation, that:

(A) radiation levels in restricted areas are not likely to cause personnel exposures in excess of the limits specified in § 175.03(c); and

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(B) radiation levels in unrestricted areas do not exceed the limits specified in § 175.03(d).

(ii) In addition to the requirements of § 175.64(e)(1)(i), a radiation protection survey also shall be performed prior to any subsequent medical use:

(A) after making any change in the treatment room shielding;

(B) after making any change in the location of the therapeutic radiation machine within the treatment room;

(C) after relocating the therapeutic radiation machine;

(D) before using the therapeutic radiation machine in a manner that could result in increased radiation levels in areas outside the external beam radiation therapy treatment room; or

(E) whenever there is reason to believe that radiation levels in unrestricted areas may have increased.

(iii) The survey record shall indicate all instances where the facility, in the opinion of the physicist performing the survey, is in violation of applicable regulations. The survey record also shall include the date of the measurements, the reason the survey is required, the manufacturer's name, model and serial number or other unambiguous identification of the therapeutic radiation machine and of the instrument(s) used to measure radiation levels, a plan of the areas surrounding (including accessible areas above and below) the treatment room which were surveyed, the measured dose rate at several points in each area expressed in <<mu>>Sv (millirems) per hour, the calculated maximum radiation exposure in a period of one (1) week for each restricted and unrestricted area, and the signature of the individual conducting the survey.

(iv) If the results of the surveys required by § 175.64(e)(1)(i) or (ii) indicate any radiation levels in excess of the respective limits specified in § 175.64(e)(1)(i), the registrant shall lock the control in the "OFF" position and not use the unit:

(A) except as may be necessary to repair, replace or test the therapeutic radiation machine, the therapeutic radiation machine shielding or the treatment room shielding; or

(B) until the registrant has applied for, and received, a specific exemption from the Department.

(2) If the survey required by § 175.64(e)(1) indicates that an individual in an unrestricted area may be exposed to levels of radiation greater than those permitted by § 175.03(d) of this Code, prior to the first medical use of the radiation therapy machine the registrant shall:

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(i) equip the unit with beam direction interlocks or install additional radiation shielding to ensure compliance with § 175.03(d) of this Code;

(ii) perform the survey required by § 175.64(e)(1) again; and

(iii) include in the report required by § 175.64(e)(3) the results of the initial survey, a description of the modification made to comply with § 175.64(e)(2)(i), and the results of the second survey; or

(iv) request and receive an amendment to the certified registration that authorizes radiation levels in unrestricted areas greater than those permitted by § 175.03(d) of this Code.

(3) Reports of external beam radiation therapy surveys and measurements. (i) The registrant for any therapeutic radiation machine subject to the requirements of § 175.64(f) or § 175.64(g) of this Code shall furnish a copy of the records required in § 175.64(e)(1) and (2) to the Department within thirty (30) days following completion of the action that initiated the record requirement.

(4) A registrant shall control access to a therapeutic radiation machine treatment room by a door at each entrance.

(5) A registrant shall equip each entrance to a therapeutic radiation machine room with a beam condition indicator light.

(6) Dosimetry equipment. (i) The registrant shall have a calibrated dosimetry system available for use. The system shall have been calibrated for cobalt-60 by the National Institute of Standards and Technology (NIST, formerly the National Bureau of Standards) or by an American Association of Physicists in Medicine (AAPM) Accredited Dosimetry Calibration Laboratory (ADCL). The calibration shall have been performed within the previous two (2) years and after any servicing that may have affected system calibration.

(ii) The registrant shall have available for use a dosimetry system for quality assurance check measurements. To meet this requirement, the system shall be compared with a system calibrated pursuant to § 175.64(e)(6)(i) and shall have been performed within the previous twelve (12) months and after each servicing that may have affected system calibration. Alternatively, the spot check system may be the same system used to meet the requirement in § 175.64(e)(6)(i).

(iii) The registrant shall maintain a record of each dosimetry system calibration, intercomparison, and comparison for the duration of the registration or certified registration. For each calibration, intercomparison, or comparison, the record shall include the date, the model numbers and serial numbers of the instruments that were calibrated, intercompared, or compared as required by § 175.64(e)(6)(i) and (ii), the correction factors that were determined, the names of the individuals who performed the calibration, intercomparison, or comparison, and evidence that the intercomparison was performed by, or under the direct supervision of, a radiation therapy physicist.

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(f) Therapeutic radiation machines incapable of operating at 500 kV or above. (1) Leakage radiation.

(i) When the x-ray tube is operated at its maximum rated tube current for the maximum kV, the leakage air kerma rate shall not exceed:

(A) for 5-50 kV systems: 1 mGy (100 mrad) in any one hour at any position 5 cm from the tube housing assembly; or

(B) for 50 and kV systems: 1 cGy (1 rad) in any one hour in any direction at 1 m from the source. This air kerma measurement may be averaged over areas not larger than 100 cm 2. In addition, the air kerma rate at a distance of 5 cm from the surface of the tube housing shall not exceed 30 cGy (30 rad) per hour.

(2) Permanent beam limiting devices. (i) Permanent diaphragms or cones used for limiting the useful beam shall provide at least the same degree of attenuation as required for the tube housing assembly.

(3) Adjustable or removable beam limiting devices. (i) All adjustable or removable beam limiting devices, diaphragms, cones or blocks shall not transmit more than 5 percent of the useful beam for the most penetrating beam used.

(ii) When adjustable beam limiting devices are used, the position and shape of the radiation field shall be indicated by a light beam.

(4) Filter system. (i) The filter system shall be so designed that:

(A) filters cannot be accidentally displaced at any possible tube orientation;

(B) for equipment installed after August 1, 1994, an interlock system prevents irradiation if the proper filter is not in place; and

(C) the air kerma rate outside the useful beam measured 1 m from the filter slot shall not exceed 1 cGy (1 rad) in any hour under any operating conditions.

(ii) Each filter shall be marked as to its material of construction and its thickness.

(5) Tube immobilization. (i) The x-ray tube shall be mounted so that it cannot accidentally turn or slide with respect to the housing aperture; and

(ii) the tube housing assembly shall be capable of being immobilized for stationary portal treatments.

(6) Source marking. (i) The tube housing assembly shall be marked so that it is possible to determine the location of the source to within 5 mm, and such marking shall be readily accessible for use during calibration procedures.

(7) Beam block. (i) Contact therapy tube housing assemblies shall have a remova-

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ble shield of material, equivalent in attenuation to 0.5 mm of lead at 100 kV, which can be positioned over the entire useful beam exit port during periods when the beam is not in use.

(8) Timer. (i) A suitable irradiation control device shall be provided to terminate the irradiation after a pre-set time interval.

(ii) A timer which has a display shall be provided at the treatment control panel. The timer shall have a pre-set time selector.

(iii) The timer shall be a cumulative timer which activates with an indication of "BEAM-ON" and retains its reading after irradiation is interrupted or terminated. After irradiation is terminated and before irradiation can be reinitiated, it shall be necessary to reset the timer.

(iv) The timer shall terminate irradiation when a pre-selected time has elapsed, if any dose monitoring system present has not previously terminated irradiation.

(v) The timer shall permit accurate pre-setting and determination of exposure times as short as 1 sec.

(vi) The timer shall not permit an exposure if set at zero.

(vii) The timer shall not activate until the shutter is opened when irradiation is controlled by a shutter mechanism unless calibration includes a timer error correction to compensate for mechanical lag.

(viii) The timer shall be accurate to within 1 percent of the selected value or 1 second, whichever is greater.

(9) Control panel functions. (i) The control panel, in addition to the displays required by other provisions in § 175.64(f), shall have:

(A) an indication of whether electrical power is available at the control panel and if activation of the x-ray tube is possible;

(B) an indication of whether x-rays are being produced;

(C) a means for indicating x-ray tube potential and current;

(D) a means for terminating an exposure at any time;

(E) a locking device which will prevent unauthorized use of the therapeutic radiation machine; and

(F) for therapeutic radiation machines installed after August 1, 1994, a positive display of the specific filter(s) in the beam.

(10) Multiple tubes. (i) When a control panel may energize more than one x-ray

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tube:

(A) it shall be possible to activate only one x-ray tube at any time;

(B) there shall be an indication at the control panel identifying which x-ray tube is energized; and

(C) there shall be an indication at the tube housing assembly when that tube is energized.

(11) Target-to-skin distance (TSD). (i) There shall be a means of determining the central axis TSD to within 1 cm and of reproducing this measurement to within 2 mm thereafter.

(12) Shutters. (i) Unless it is possible to bring the x-ray output to the prescribed exposure parameters within 5 sec after the x-ray "ON" switch is energized, the beam shall be attenuated by a shutter having a lead equivalency not less than that of the tube housing assembly. In addition, after the unit is at operating parameters, the shutter shall be controlled electrically by the operator from the control panel. An indication of shutter position shall appear at the control panel.

(13) Low filtration x-ray tubes. (i) Each therapeutic radiation machine equipped with a beryllium or other low-filtration window shall be clearly labeled as such upon the tube housing assembly and shall be provided with a permanent warning device on the control panel that is activated when no addition filtration is present to indicate that the dose rate is very high.

(14) Facility design requirements for therapeutic radiation machines capable of operating in the range of greater than 50 kV to 500 kV.

(i) In addition to shielding adequate to meet the requirements of § 175.03(d), the treatment room shall provide:

(A) a system for continuous two-way aural communication between the patient and the operator at the control panel; and

(B) a system to permit continuous observation of the patient during irradiation by the operator from the control panel.

(ii) The therapeutic radiation machine shall not be used for patient irradiation unless both aural communication with and continuous observation of the patient are possible.

(15) Additional requirements. (i) Treatment rooms which contain a therapeutic radiation machine capable of operating above 150 kV shall meet the following additional requirements:

(A) all protective barriers shall be fixed except for entrance doors or beam interceptors;

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(B) the control panel shall be located outside the treatment room, or in a totally enclosed and shielded booth which has a ceiling, inside the room;

(C) interlocks shall be provided such that all entrance doors, including doors to any interior booths, shall be closed before treatment can be initiated or continued; and

(D) if the radiation beam is interrupted by any door opening, it shall not be possible to restore the machine to operation without closing the door and reinitiating irradiation by manual action at the control panel.

(16) Full calibration measurements. (i) Full calibration of a therapeutic radiation machine subject to the requirements of § 175.64(f) of this Code shall be performed by, or under the direct supervision of, a radiation therapy physicist:

(A) before the first medical use following installation or reinstallation of the therapeutic radiation machine;

(B) annually; and

(C) before medical use under the following conditions:

(a) whenever quality assurance check measurements indicate that the radiation output differs by more than 5 percent from the value obtained at the last full calibrations; or

(b) following any component replacement, major repair or modification of components that could affect the characteristics of the radiation beam.

(D) Notwithstanding the requirements of § 175.64(f)(16)(i)(C):

(a) full calibration of therapeutic radiation machines with multi-energy capabilities is required only for those modes and/or energies that are not within their acceptable ranges; and

(b) if the repair, replacement or modification does not affect all energies, full calibration shall be performed at the affected energy that is in most frequent clinical use at the facility and the remaining energies may be validated by quality assurance check procedures using the criteria in § 175.64(f)(16)(i)(C)(a).

(ii) Whenever a quality assurance check required by § 175.64(f)(17) indicates a significant change in the operating characteristics of a system, as specified in the radiation therapy physicist's quality assurance manual, the system shall be recalibrated as specified herein.

(iii) To satisfy the requirement of § 175.64(f)(16)(i), full calibration shall include all measurements recommended for annual calibration by the National Council on Radiation Measurement and Protection (NCRP) Report 69, "Dosimetry of X-Ray and Gamma Ray Beams for Radiation Therapy in the Energy Range 10 keV to 50 MeV" (1981), or any successor protocol, and be performed with a dosimetry system calibrated pur-

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suant to § 175.64(e)(6)(i).

(iv) The registrant shall maintain a record of each calibration for the duration of the registration. The record shall include the date of the calibration, the manufacturer's name, model and serial number or other unambiguous identification of both the therapeutic radiation machine and the x-ray tube, the model numbers and serial numbers or other unambiguous identification of the instruments used to calibrate the therapeutic radiation machine, and the signature of the radiation therapy physicist responsible for performing the calibration.

(17) Periodic quality assurance checks. (i) Periodic quality assurance checks shall be performed on therapeutic radiation machines subject to the requirements of § 175.64(f) of this Code which are capable of operation at greater than 50 kV.

(ii) To satisfy the requirement of § 175.64(f)(17)(i), quality assurance checks shall meet the following requirements:

(A) the registrant shall perform quality assurance checks in accordance with written procedures established by the radiation therapy physicist;

(B) the quality assurance check procedures shall specify that the quality assurance check shall be performed during the calibration specified in § 175.64(f)(16)(i); and

(C) the quality assurance check procedures shall specify the frequency at which tests or measurements are to be performed and the acceptable tolerance for each parameter measured in the check, when compared to the value for that parameter determined in the full calibration specified in § 175.64(f)(16)(i).

(iii) The cause for a parameter exceeding a tolerance set by the radiation therapy physicist shall be investigated and corrected before the system is used for patient irradiation.

(iv) The registrant shall use the dosimetry system specified in § 175.64(e)(6)(ii) to make the periodic quality assurance check required in § 175.64(f)(17)(i).

(v) The registrant shall have the radiation therapy physicist review and sign the results of each radiation output quality assurance check at intervals not to exceed one month.

(vi) Therapeutic radiation machines subject to the requirements of § 175.64(f) of this Code shall have safety quality assurance checks of each external beam radiation therapy facility performed monthly. This requirement does not apply to facilities which have not been used for more than one month, except that such safety quality assurance checks shall be performed before the first clinical treatment when the facility is returned to use.

(vii) To satisfy the requirements of § 175.64(f)(17)(vii), safety quality assurance checks shall ensure proper operation of:

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(A) electrical interlocks at each external beam radiation therapy room entrance, including those at any interior control booths;

(B) proper operation of the "BEAM-ON" and termination switches;

(C) beam condition indicator lights on the access door(s), control console and in the radiation therapy room;

(D) viewing and aural communication systems; and

(E) electrically operated treatment room doors from inside and outside the treatment room.

(viii) The registrant shall promptly repair any system identified in § 175.64(f)(17)(vii) that is not operating properly.

(ix) The registrant shall maintain a record of each quality assurance check required by § 175.64(f)(17)(i) and § 175.64(f)(17)(vi) for three (3) years. The record shall include the date of the quality assurance check, the manufacturer's name, model and serial number or other unambiguous identification of the therapeutic radiation machine and of the instrument(s) used to measure the radiation output of the therapeutic radiation machine, and the signature of the individual who performed the periodic quality assurance check.

(18) Operating procedures. (i) The therapeutic radiation machine shall not be used for irradiation of patients unless the requirements of § 175.64(f)(16) and § 175.64(f)(17) have been met.

(ii) Therapeutic radiation machines shall not be left unattended unless it is secured pursuant to § 175.64(f)(9)(i)(E).

(iii) When a patient must be held in position for radiation therapy, mechanical supporting or restraining devices shall be used.

(iv) The tube housing assembly shall not be held by an individual during operation unless the assembly is designed to require such holding and the peak tube potential of the system does not exceed 50 kV. In such cases, the holder shall wear protective gloves and apron of not less than 0.5 mm lead equivalency at 100 kV.

(v) A copy of the current operating and emergency procedures shall be maintained at the therapeutic radiation machine control console.

(vi) No individual other than the patient shall be in the treatment room during exposures from therapeutic radiation machines operating above 150 kV. At energies less than or equal to 150 kV, an individual, other than the patient, in the treatment room shall be protected by a barrier sufficient to meet the requirements of § 175.03 of this Code.

(19) Possession of survey instrument(s). (i) Each facility location authorized to

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use a therapeutic radiation machine in accordance with § 175.64(f) of this Code shall possess appropriately calibrated portable monitoring equipment. At a minimum, such equipment shall include a portable radiation measurement survey instrument capable of measuring dose rates over the range of 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The survey instruments shall be operable and calibrated in accordance with § 175.64(h).

(g) Therapeutic radiation machines: photon therapy systems capable of operating at 500 kV and above and/or electron therapy systems capable of operating at 500 keV and above. (1) All therapeutic radiation machines installed after August 1, 1994 shall have an active (not withdrawn or terminated), approved Premarket Approval Application (PMA) issued by the U.S. Food and Drug Administration pursuant to 21 CFR Part 814, or any successor law or regulation.

(2) Leakage radiation outside the maximum useful beam.

(i) The absorbed dose rate due to leakage radiation (excluding neutrons) at any point outside the maximum sized useful beam, but within a circular plane of radius two (2) meters which is perpendicular to and centered on the central axis of the useful beam at the nominal treatment distance (i.e. patient plane), shall not exceed a maximum of 0.2 percent and an average of 0.1 percent of the absorbed dose rate on the center axis at the treatment distance. Measurements shall be averaged over an area not exceeding 100 cm 2 at a minimum of 8 points uniformly distributed in the plane.

(ii) The neutron absorbed dose rate outside the useful beam shall be kept as low as practicable. Measurements of the portion of the leakage radiation dose contributed by neutrons shall be averaged over an area not exceeding 800 cm 2.

(iii) For each therapeutic radiation machine, the registrant shall determine, or obtain from the manufacturer, the leakage radiation existing at the positions specified in § 175.64(g)(1)(i) and (ii) for the specified operating conditions.

(iv) Records on leakage radiation measurements shall be maintained at the installation for inspection by the Department.

(3) Filters and wedges. (i) Each filter and/or wedge which is removable from the system shall be clearly marked with an identification number. For removable wedge filters, the nominal wedge angle shall appear on the wedge or wedge tray (if permanently mounted to the tray). If the wedge or wedge tray is damaged, the wedge transmission factor shall be redetermined.

(4) Termination switches. (i) It shall be possible to terminate irradiation and equipment movement or go from an interruption condition to termination condition at any time from the operator's position at the treatment control panel.

(5) Facility design requirements for therapeutic radiation machines capable of operating above 500 kV or 500 keV. (i) In addition to shielding adequate to meet the requirements of § 175.03 of this Code, the following design requirements are made:

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(A) All protective barriers shall be fixed, except for access doors to the treatment room or movable beam interceptors.

(B) In addition to other requirements specified in this section, the control panel shall:

(a) be located outside the treatment room;

(b) provide an indication of whether electrical power is available at the control panel and if activation of the radiation source is possible;

(c) provide an indication of whether radiation is being produced; and

(d) include a locking access control device which will prevent unauthorized use of the therapeutic radiation machine.

(C) There shall be a system for continuous two-way aural communication between the patient and the operator at the control panel.

(D) There shall be a system to permit continuous observation of the patient following positioning and during irradiation by the operator from the control panel.

(E) Treatment room entrances shall be provided with warning lights in a readily observable position near the outside of all access doors which will indicate when the useful beam is "ON" and when it is "OFF."

(F) Interlocks shall be provided such that all entrance doors must be closed before treatment can be initiated or continued; if the radiation beam is interrupted by any door opening, it shall not be possible to restore the machine to operation without closing the door and reinitiating irradiation by manual action at the control panel.

(G) If the shielding material in any protective barrier requires the presence of a beam interceptor to ensure compliance with § 175.03(d) of this Code, interlocks shall be provided to prevent the production of radiation, unless the beam interceptor is in place, whenever the useful beam is directed at the designated barrier(s).

(H) In addition to the termination switch required by § 175.64(g)(4), at least one (1) emergency power cutoff switch or button (e.g. "scram button") shall be located in the radiation therapy treatment room and shall terminate all equipment electrical power including radiation and mechanical motion; all emergency power cutoff switches shall include a manual reset so that the therapeutic radiation machine cannot be restarted from the unit's control panel without resetting the emergency cutoff switch.

(I) All safety interlocks shall be designed so that any defect or component failure in the safety interlock system prevents or terminates operation of the therapeutic radiation machine.

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(J) Surveys for residual radioactivity shall be conducted on all therapeutic radiation machines capable of generating photon and electron energies above 10 MV or MeV prior to machining, removing or working on such machine's components which may have become activated due to photo-neutron production.

(6) Radiation therapy physicist. (i) The radiation therapy physicist named on the registrant's certified registration shall be responsible for:

(A) all full calibrations required by § 175.64(g)(8) and protection surveys required by § 175.64(e)(1);

(B) supervision and review of dosimetry as required by § 175.07(c) of this Code;

(C) beam data acquisition and transfer for computerized dosimetry, and supervision of its use;

(D) quality assurance required by § 175.07 of this Code and quality assurance check review required by § 175.64(g)(9)(v);

(E) consultation with the authorized user(s) in treatment planning, as needed; and

(F) performance of calculations or other assessments regarding misadministrations.

(ii) If the radiation therapy physicist named on the registrant's certified registration is not a full-time employee of the registrant, the operating procedures required by § 175.64(g)(7) shall specifically address how the radiation therapy physicist is to be contacted for problems or emergencies, as well as the specific actions to be taken until the radiation therapy physicist can be contacted.

(7) Operating procedures. (i) No individual, other than the patient, shall be in the treatment room during treatment.

(ii) The therapeutic radiation machine shall not be used for patient irradiation unless both aural and visual communication systems are operational and continuous observation of the patient is maintained.

(iii) No individual shall be in the treatment room during irradiation for testing or calibration purposes.

(iv) Therapeutic radiation machines shall not be used for medical use unless the requirements of § 175.64(e)(1), § 175.64(g)(8) and § 175.64(g)(9) have been met.

(v) Therapeutic radiation machines, when not in operation, shall be secured to prevent unauthorized use.

(vi) If a patient must be held in position during treatment, mechanical supporting or restraining devices shall be used.

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(vii) A copy of the current operating and emergency procedures shall be maintained at the therapeutic radiation machine control console.

(viii) Emergency procedures shall be posted in the treatment room.

(8) Full calibration measurements. (i) Full calibration of a therapeutic radiation machine subject to the requirements of § 175.64(g) of this Code shall be performed by, or under the direct supervision of, the radiation therapy physicist named on the registrant's certified registration:

(A) before first medical use following installation or reinstallation of the therapeutic radiation machine;

(B) annually; and

(C) before medical use under the following conditions:

(a) whenever quality assurance check measurements indicate that the output differs by more than five (5) percent from the output obtained at the last full calibration; and

(b) following any component replacement, major repair, or modification of components that could affect the characteristics of the radiation beam; and

(D) Notwithstanding the requirements of § 175.64(g)(8)(i)(C):

(a) full calibration of therapeutic radiation machines with multi-energy and/or multi-mode capabilities is required only for those modes and/or energies that are not within their acceptable ranges; and

(b) if the repair, replacement or modification does not affect all modes and/or energies, full calibration shall be performed on the affected mode/energy that is in most frequent clinical use for that machine at the facility and the remaining energies/modes may be validated with quality assurance check procedures pursuant to the criteria in § 175.64(g)(8)(i)(C)(a).

(ii) To satisfy the requirement of § 175.64(g)(8)(i), full calibration shall include all measurements required for annual calibration by "Comprehensive QA for Radiation Oncology: Report of AAPM Radiation Therapy Committee Task Group 40," Medical Physics 21(4):581-, 1994, or any successor protocol. Additionally, the facility shall conduct the output calibration for the therapeutic radiation machine's photon and electron beam modes according to the protocol as outlined in AAPM Report Task Group Report 21 ("A Protocol for the Determination of Absorbed Dose from High-Energy Photon and Electron Beams" Task Group 21 (TG21), Radiation Therapy Committee of American Association of Physicists in Medicine, published in Medical Physics, Volume 10(8), Nov/Dec 1983); or, AAPM Report Task Group Report 51(TG51) ("Protocol for Clinical Reference Dosimetry of High-Energy Photon and Electron Beams", Medical Physics, Volume 29(9), September 1999) or any successor to these documents.

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(iii) The registrant shall use the dosimetry system described in § 175.64(e)(6)(i) to measure the output at each discrete photon and electron energy (which the therapeutic radiation machine can produce) for one (1) set of exposure conditions. The remaining radiation measurements required in § 175.64(g)(8)(ii) may then be made using a dosimetry system that indicates relative dose rates;

(iv) The registrant shall maintain a record of each calibration for the duration of the certified registration. The record shall include the date of the calibration, the manufacturer's name, model and serial number or other unambiguous identification of the therapeutic radiation machine along with the instrument's certificate of calibration, all measured beam output data collected during the calibration, the derivation for all the correction factors (as delineated in AAPM Reports TG 21 or TG 51 or any successor publication) applied to the .measured beam output data' in the calculation of the therapeutic radiation machine's beam output dose rate (the latter shall be conducted for each photon and electron beam clinically utilized at the facility), and the signature of the radiation therapy physicist named on the certified registration.

(9) Periodic quality assurance checks. (i) Periodic quality assurance checks shall be performed on each therapeutic radiation machine subject to the requirements of § 175.64(g) of this Code.

(ii) To satisfy the requirement of § 175.64(g)(9)(i), quality assurance checks shall include determination of all parameters for periodic quality assurance checks at the intervals contained in "Comprehensive QA for Radiation Oncology: Report of AAPM Radiation Therapy Committee Task Group 40," Medical Physics 21(4):581-, 1994, or any successor protocol.

(iii) The registrant shall use a dosimetry system described in § 175.64(e)(6)(ii) to make the periodic quality assurance checks required by § 175.64(g)(9)(ii).

(iv) The registrant shall perform periodic quality assurance checks required by § 175.64(g)(8)(i) in accordance with procedures established by the radiation therapy physicist named on the registrant's certified registration.

(v) The registrant shall review the results of each periodic radiation output check according to the following procedures:

(A) the authorized user and radiation therapy physicist shall be notified immediately if any parameter is not within its acceptable range as determined pursuant to § 175.64(g)(9)(iv). The therapeutic radiation machine shall not be made available for subsequent medical use until the radiation therapy physicist has determined that all parameters are within their acceptable ranges;

(B) if all quality assurance check parameters appear to be within their acceptable ranges, the quality assurance check shall be reviewed and signed by either the authorized user or the radiation therapy physicist within ten (10) days; and

(C) the radiation therapy physicist shall review and sign the results of each radiation output quality assurance check at intervals not to exceed one (1) month.

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(vi) Therapeutic radiation machines subject to the requirements of § 175.64(g) of this Code shall have safety quality assurance checks of each external beam radiotherapy facility weekly.

(vii) To satisfy the requirement of § 175.64(g)(9)(vi), safety quality assurance checks shall ensure proper operation of:

(A) electrical interlocks at each therapeutic radiation machine treatment room entrance;

(B) proper operation of "BEAM-ON", interrupt and termination switches;

(C) beam condition indicator lights on the access doors, control console, and in the radiation therapy treatment room;

(D) viewing and aural communication systems;

(E) electrically operated treatment room door(s) from inside and outside the treatment room;

(F) at least one (1) emergency power cutoff switch. If more than one (1) emergency power cutoff switch is installed and not all switches are tested at once, each switch shall be tested on a rotating basis.

(viii) The registrant shall lock the control console in the "off" position if any door interlock malfunctions. No registrant shall use the therapeutic radiation machine until the interlock system is repaired unless specifically authorized by the Department.

(ix) The registrant shall promptly repair any system identified in § 175.64(g)(9)(vii) that is not operating properly.

(x) The registrant shall maintain a record of each quality assurance check required by § 175.64(g)(9)(i) and (vii) for three (3) years. The record shall include the date of the quality assurance check, the manufacturer's name, model and serial number or other unambiguous identification of the therapeutic radiation machine of the instrument(s) used to measure the output of the therapeutic radiation machine, and the signature of the individual who performed the periodic quality assurance check.

(10) Reports of calibrations. (i) The registrant shall furnish a copy of the initial full calibration report required by § 175.64(g)(8)(i)(A) to the Bureau of Radiological Health within thirty (30) days following completion of the calibration.

(11) Possession of survey instrument(s). (i) Each facility location authorized to use a therapeutic radiation machine in accordance with § 175.64(g) of this Code shall possess appropriately calibrated portable monitoring equipment. At a minimum, such equipment shall include a portable radiation measurement survey instrument capable of measuring dose rates over the range of 10 <<mu>>Sv (1 mrem) per hour to 10

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mSv (1000 mrem) per hour. The survey instruments shall be operable and calibrated in accordance with § 175.64(h).

(h) Calibration and check of survey instruments. (1) The registrant shall ensure that the survey instruments used to show compliance with the requirements of this section and other applicable parts of this Code have been calibrated before first use, at intervals not to exceed twelve (12) months and following repair.

(2) To satisfy the requirements of § 175.64(h)(1), the registrant shall:

(i) calibrate all required scale readings up to 10 mSv (100 mrem) per hour with an appropriate radiation source;

(ii) calibrate at least two (2) points on each scale to be calibrated. These points should be at approximately 1/3 and 2/3 of the full scale.

(3) To satisfy the requirements of § 175.64(h)(2), the registrant shall:

(i) consider a point as calibrated if the indicated dose rate differs from the calculated dose rate by not more than ten (10) percent; or

(ii) consider a point as calibrated if the indicated dose rate differs from the calculated dose rate by not more than twenty (20) percent if a correction factor or graph is conspicuously attached to the instrument.

(4) The registrant shall retain a record of each calibration required in § 175.64(h)(1) for three (3) years and which shall include:

(i) a description of the calibration procedure;

(ii) the manufacturer, model and serial number of the instrument;

(iii) a description of the source used and the certified dose rates from the source (as evidenced by NIST traceability);

(iv) the rates indicated by the instrument being calibrated, the correction factors determined from the calibration data; and

(v) the signature of individual who performed the calibration and the date of calibration.

(5) Records of calibrations which contain information required by § 175.64(h)(4) shall be maintained by the registrant at the facility.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

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Subd. (g) par (8) subpars (ii), (iv) amended City Record Sept. 29, 2006, eff. Oct. 29, 2006. [See Vol. 9 Statements of Basis and Purpose No. 25]

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NOTES:

Paragraph (8) of subdivision (g) of § 175.64 was amended on September 26, 2006, in order to specify the use of x-ray therapy protocols developed by the American Association of Physicists in Medicine to be followed by each licensed facility in conducting the annual radiation output measurement for each therapy unit, and to delineate the minimal content that should be contained in each annual calibration report by the facility for review by Department inspectors.

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TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIATION EQUIPMENT

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.65. Veterinary radiography and fluoroscopy.

(a) Fixed radiographic installations. (1) Equipment.

(i) Collimating devices capable of restricting the useful beam to the area of clinical interest shall be used.

(ii) The x-ray films used as the recording medium during the x-ray examination shall show substantial evidence of cut-off (beam delineation).

(iii) A device shall be provided which terminates the exposure after a preset time interval or exposure. The exposure switch shall be of the dead-man type and shall be so arranged that it cannot be operated outside a shielded area.

(2) Conditions for operation of equipment. (i) Only persons required for the x-ray procedure shall be in the x-ray room during exposures.

(ii) When an animal patient must be held in position during exposures, mechanical supporting or restraining devices shall be used.

(iii) Animal patients or films shall be held by an individual only under extreme conditions when clinically necessary. Such individuals shall wear protective gloves having at least 0.5 mm lead equivalent, a protective apron of at least 0.25 mm lead equivalent, and shall keep all parts of his/her body out of the useful beam.

(iv) The exposure of any individual used for holding animals shall be monitored.

(v) Pregnant women and individuals under 18 years of age shall not hold animal patients or films under any conditions.

(b) Portable or mobile radiographic installations. (1) Equipment.

(i) Collimating devices capable of restricting the useful beam to the area of clinical interest shall be used.

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(ii) The x-ray film used as the recording medium during the x-ray examination shall show evidence of cut-off (beam delineation).

(iii) A device shall be provided which terminates the exposure after a preset time interval or exposure.

(iv) A dead-man type of exposure switch shall be provided with a cord of sufficient length so that the operator can stand at least two (2) m (6 ft) from the animal patient, the x-ray tube and out of the useful beam.

(2) Conditions for operation of equipment. (i) No person shall be regularly employed to support or hold animals or film during x-ray exposures.

(ii) When an animal patient must be held in position during exposures, mechanical supporting or restraining devices shall be used.

(iii) Animal patients or films shall be held by an individual only under extreme conditions when clinically necessary. Such individuals shall wear protective gloves having at least 0.5 mm lead equivalent, a protective apron of at least 0.25 mm lead equivalent, and shall keep all parts of his/her body out of the useful beam.

(iv) The exposure of any individual used for holding animals shall be monitored.

(v) Pregnant women and individuals under 18 years of age shall not hold animal patients or films under any conditions.

(c) Fluoroscopic installations. (1) Equipment.

(i) Equipment shall be so constructed that the entire cross-section of the useful beam is always intercepted by a primary protective barrier (usually a lead glass screen or image intensifier assembly) regardless of the panel-screen distance. For conventional fluoroscopes, this requirement may be assumed to have been met if, when the collimating system is opened to its fullest extent, an unilluminated margin is left on all edges of the fluorescent screen regardless of the position of the screen during use. Equipment with an image intensifier shall be so constructed that the useful beam cannot exceed the limits of the input phosphor.

(ii) The exposure shall automatically terminate when the barrier is removed from the useful beam.

(iii) With the fluoroscope operating at the highest potential employed and with the fluorescent screen 36 cm (14 in.) from the panel of the tabletop and in the useful beam without a patient, the exposure rate 5 cm (2 in.) beyond the viewing surface of the screen shall not exceed 7.74 E-6 C-kg-1-hr-1 (30 mR-hr-1) for each 2.58 E-4 C-kg-1- min-1 (R-min-1) at the tabletop.

(iv) The fluoroscopic exposure switch shall be of the dead-man type.

(v) Provision shall be made to intercept the scattered x-rays from the undersur-

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face of the tabletop and other structures under the table.

(vi) The source-panel or source-tabletop distance shall in no case be less than 30 cm (12 in.) and is recommended to be not less than 38 cm (15 in.).

(2) Mobile fluoroscopic equipment is subject to the following additional requirements. (i) In the absence of a tabletop, a cone or spacer frame shall limit the source-to-skin distance to not less than 30 cm (12 in.).

(ii) Image intensification shall always be provided.

(iii) It shall not be possible to operate a machine unless the useful beam is intercepted by the image intensifier.

(3) Conditions for operation of equipment. (i) Protective garments of at least 0.25 mm lead equivalent shall be available and shall be worn by the fluoroscopist during every examination.

(ii) Unless physical measurements indicate that they are not needed, protective garments of at least 0.25 mm lead equivalent each shall be worn by the physician, nurse, technician and all other persons within the fluoroscopy room.

(iii) Only persons needed in the fluoroscopic room shall be present during the exposure.

(iv) The fluoroscopic room shall be free of extraneous light that interferes with the examination.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

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ARTICLE 175. RADIATION CONTROL.
RADIATION EQUIPMENT

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.66. Miscellaneous and special types of radiation equipment.

(a) Types or uses of radiation producing equipment not specifically listed or covered by these regulations, and not specifically exempted, shall be manufactured, operated or used such that the radiation level measured 5 cm (2 inches) from any accessible surface, and averaged over an area of 10 cm 2 (1.55 in. 2), shall not exceed 1.3 E-7 C/kg (0.5 milliroentgen) per hour.

HISTORICAL NOTE

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TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.101. General requirements for radioactive materials licenses.

(a) License required. (1) Except for the removal of source material from its place of deposit in nature or as otherwise provided in this Code, no person shall transfer, receive, produce, possess or use any radioactive material except pursuant to a license issued by the Department.

(2) Fees for each license shall be paid pursuant to § 5.07 of this Code.

(3) The requirements of this section are in addition to, and not in substitution for, other requirements of this Code. In any conflict between the requirements of this section and a specific requirement in another part of this Code, the specific requirement governs.

(b) Exempt source material. (1) Any person is exempt from the provisions of this code to the extent that such person receives, possesses, uses or transfers unrefined and unprocessed ore containing source material; provided that, except as authorized in a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, such person shall not refine or process such ore.

(2) Any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses or transfers source material in any chemical mixture, compound, solution or alloy in which the source material is by weight less than 1/20 of 1 percent (0.05%) of the mixture, compound, solution or alloy.

(3) Any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses or transfers:

(i) any quantities of thorium contained in:

(A) incandescent gas mantles;

(B) vacuum tubes;

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(C) welding rods;

(D) electric lamps for illuminating purposes provided that each lamp does not contain more than 50 milligrams of thorium;

(E) germicidal lamps, sunlamps, and lamps for outdoor or industrial lighting provided that each lamp does not contain more than 2 grams of thorium;

(F) rare earth metals and compounds, mixtures, and products containing not more than 0.25 percent, by weight of thorium, uranium, or any combination of these; or

(G) personnel neutron dosimeters, provided that each dosimeter does not contain more than 50 milligrams of thorium;

(ii) any source material contained in the following products:

(A) glazed ceramic tableware, provided that the glaze contains not more than 20 percent by weight source material;

(B) glassware, containing not more than 10 percent by weight source material, but not including commercially manufactured glass brick, pane glass, ceramic tile or other glass, glass enamel or ceramic used in construction;

(C) glass enamel or glass enamel frit containing not more than 10 percent by weight of source material imported or ordered for importation into the United States, or initially distributed by manufacturers in the United States, before July 25, 1983; or

(D) piezoelectric ceramic, containing not more than 2 percent by weight source material;

(iii) photographic film, negatives, and prints containing uranium or thorium;

(iv) any finished or partly fabricated product of, or containing, tungstenthorium or magnesium-thorium alloys, provided that the thorium content of the alloy does not exceed 4 percent by weight and that this exemption shall not be deemed to authorize the chemical, physical, or metallurgical treatment or processing of any such product or part;

(v) uranium contained in counterweights installed in aircraft, rockets, projectiles, and missiles, or stored or handled in connection with installation or removal of such counterweights, provided that:

(A) each counterweight is manufactured in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission, authorizing distribution by the licensee pursuant to 10 CFR Part 40;

(B) each counterweight has been impressed with the following legend clearly legi-

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ble through any plating or other covering: "DEPLETED URANIUM";

(C) each counterweight is durably and legibly labeled or marked with the identification of the manufacturer and the statement: "UNAUTHORIZED ALTERATIONS PROHIBITED"; and

(D) this exemption shall not be deemed to authorize the chemical, physical, or metallurgical treatment or processing of any such counterweights other than repair or restoration of any plating or other covering;

(vi) natural or depleted uranium metal used as shielding constituting part of any shipping container, provided that:

(A) the shipping container is conspicuously and legibly impressed with the legend "Caution--Radioactive Shielding--Uranium"; and

(B) the uranium metal is encased in mild steel or equally fire resistant metal or minimum wall thickness of one-eighth inch (3.2 mm);

(vii) thorium contained in finished optical lenses, provided that each lens does not contain more than 30 percent by weight of thorium, and that this exemption shall not be deemed to authorize either:

(A) the shaping, grinding, or polishing of such lens or manufacturing processes other than the assembly of such lens into optical systems and devices without any alteration of the lens; or

(B) the receipt, possession, use, or transfer of thorium contained in contact lenses, or in spectacles, or in eyepieces in binoculars or other optical instruments;

(viii) uranium contained in detector heads for use in fire detection units, provided that each detector head contains not more than 0.005 microcurie of uranium;

(ix) thorium contained in any finished aircraft engine part containing nickel-thoria alloy, provided that:

(A) the thorium is dispersed in the nickel-thoria alloy in the form of finely divided thoria (thorium dioxide); and

(B) the thorium content in the nickel-thoria alloy does not exceed 4 percent by weight.

(x) The exemptions contained in § 175.101(b)(2) and (3)(i) through (ix) shall not authorize the manufacturer of any of the products described.

(c) Exempt radioactive material other than source material. (1) Exempt concentrations.

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(i) Except as provided in § 175.101(c)(1)(ii), any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, owns or acquires products containing radioactive material introduced in concentrations not in excess of those listed in Appendix A of this section.

(ii) No person may introduce radioactive material into a product or material knowing or having reason to believe that it will be transferred to persons exempt under § 175.101(c)(1)(i) or equivalent regulations of the U.S. Nuclear Regulatory Commission or an agreement state, except in accordance with a specific license issued pursuant to this Code or a general license provided for in this Code.

(2) Exempt quantities. (i) Except as provided in § 175.101(c)(2)(ii) and (iii), any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, owns or acquires radioactive material in individual quantities, each of which does not exceed the applicable quantity set forth in Appendix B of this section.

(ii) Section 175.101(c)(2)(i) does not authorize the production, packaging or repackaging of radioactive material for purposes of commercial distribution, or the incorporation of radioactive material into products intended for commercial distribution.

(iii) No person shall, for purposes of commercial distribution, transfer radioactive material in the individual quantities set forth in Appendix B of this section, knowing or having reason to believe that such quantities of radioactive material will be transferred to persons exempt under § 175.101(c)(2)(i) or equivalent regulations of the U.S. Nuclear Regulatory Commission or an agreement state, except in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission pursuant to § 32.18 of 10 CFR Part 32, or by the Department, which license states that the radioactive material may be transferred by the licensee to persons exempt under § 175.101(c)(2)(i) or the equivalent regulations of the U.S. Nuclear Regulatory Commission or any agreement state.

(3) Exempt items. Except for persons who apply radioactive material to, or persons who incorporate radioactive material into the following products, or persons who initially transfer such products for sale or distribution, any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, owns or acquires the following products:

(i) Timepieces or timepiece hands or dials containing radium which were manufactured under a specific license issued by the Department or an agreement state and which meet the following or equivalent conditions:

(A) The timepiece or timepiece hands or dials contain no more than the following specified quantities of radium:

(a) 5.55 kBq (0.15 <<mu>>Ci) per watch;

(b) 1.11 kBq (0.03 <<mu>>Ci) per watch hand;

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(c) 3.33 kBq (0.09 <<mu>>Ci) per watch dial;

(d) 7.4 kBq (0.2 <<mu>>Ci) per clock;

(e) 1.48 kBq (0.04 <<mu>>Ci) per clock hand; or

(f) 4.44 kBq (0.12 <<mu>>Ci) per clock dial.

(B) The timepiece is not a pocket watch.

(C) The timepiece is marked or coded to identify the date of manufacture and that it contains radium.

(D) The timepiece emits sufficient luminosity, omitting photoactivation, that its dial can be read in the dark during its entire design lifetime.

(ii) Timepieces or hands or dials containing not more than the following specified quantities of radioactive material and not exceeding the following specified radiation dose rates:

(A) 925 MBq (25 mCi) of hydrogen-3 per timepiece.

(B) 185 MBq (5 mCi) of hydrogen-3 per hand.

(C) 555 MBq (15 mCi) of hydrogen-3 per dial (bezels when used shall be considered as part of the dial).

(D) 3.7 MBq (100 <<mu>>Ci) of promethium-147 per watch or 7.4 MBq (200 <<mu>>Ci) of promethium-147 per any other timepiece.

(E) 0.74 MBq (20 <<mu>>Ci) of promethium-147 per watch hand or 1.48 MBq (40 <<mu>>Ci) of promethium-147 per other timepiece hand.

(F) 2.22 MBq (60 <<mu>>Ci) of promethium-147 per watch dial or 4.44 MBq (120 <<mu>>Ci) of promethium-147 per other timepiece dial (bezels when used shall be considered as part of the dial).

(G) The radiation dose rate from hands and dials containing promethium-147 will not exceed, when measured through 50 milligrams per square centimeter of absorber:

(a) for wristwatches, 1 <<mu>>Gy (0.1 millirad) per hour at 10 centimeters from any surface;

(b) for pocket watches, 1 <<mu>>Gy (0.1 millirad) per hour at 1 centimeter from any surface;

(c) for any other timepiece, 2 <<mu>>Gy (0.2 millirad) per hour at 10 centimeters from any surface.

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(H) 37 kBq (1 <<mu>>Ci) of radium-226 per timepiece in timepieces acquired prior to September 1, 1984.

(iii) Lock illuminators containing not more than 555 MBq (15 millicuries) of hydrogen-3 or not more than 74 MBq (2 millicuries) of promethium-147 installed in automobile locks. The radiation dose rate from each lock illuminator containing promethium-147 will not exceed 10 <<mu>>Gy (1 millirad) per hour at one centimeter from any surface when measured through 50 milligrams per square centimeter of absorber.

(iv) Precision balances containing not more than 37 MBq (1 millicurie) of hydrogen-3 per balance or not more than 18.5 MBq (0.5 millicurie) of hydrogen-3 per balance part.

(v) Automobile shift quadrants containing not more than 925 MBq (25 millicuries) of hydrogen-3.

(vi) Marine compasses containing not more than 27.8 GBq (750 millicuries) of hydrogen-3 gas and other marine navigational instruments containing not more than 9.25 GBq (250 millicuries) of hydrogen-3 gas.

(vii) Thermostat dials and pointers containing not more than 925 MBq (25 millicuries) of hydrogen-3 per thermostat.

(ix) Electron tubes, provided, that each tube does not contain more than one of the following specified qualities of radioactive material:

(A) 5.55 GBq (150 millicuries) of hydrogen-3 per microwave receiver detector tube or 370 MBq (10 millicuries) of hydrogen-3 per any other electron tube;

(B) 37 kBq (1 <<mu>>Ci) of cobalt-60;

(C) 185 kBq (5 <<mu>>Ci) of nickel-63;

(D) 1.11 MBq (30 <<mu>>Ci) of krypton-85;

(E) 185 kBq (5 <<mu>>Ci) of cesium-137;

(F) 1.11 MBq (30 <<mu>>Ci) of promethium-147;

and, provided further, that the radiation dose rate due to radioactive material contained in each electron tube does not exceed 10 <<mu>>Gy (1 millirad) per hour at one centimeter from any surface when measured through 7 milligrams per square centimeter of absorber.

(x) Ionizing radiation measuring instruments containing, for purposes of internal calibration or standardization, one or more sources of radioactive material, provided that:

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(A) each source contains no more than one exempt quantity set forth in Appendix B of this section, and

(B) each instrument contains no more than 10 exempt quantities. For purposes of this requirement, an instrument's source(s) may contain either one or different types of radioactive materials and an individual exempt quantity may be composed of fractional parts of one or more of the exempt quantities in Appendix B of this section, provided that the sum of such fractions shall not exceed unity; and

(C) for the purposes of § 175.101(c)(3)(x), 1.85 kBq (0.05 <<mu>>Ci) of americium-241 shall be considered one exempt quantity.

(xi) Spark gap irradiators containing not more than 37 kBq (1 <<mu>>Ci) of cobalt-60 per spark gap irradiator for use in electrically ignited fuel oil burners having a firing rate of at least 3 gallons (11.4 liter) per hour.

(xii) The exemptions contained in § 175.101(c)(3)(i) through (xi) shall not authorize the application or incorporation of radioactive materials into the listed devices.

(4) Any person, except those who manufacture, process, or produce self-luminous products containing hydrogen-3, krypton-85, or promethium-147, is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, owns, or acquires hydrogen-3, krypton-85 or promethium-147 in self-luminous products manufactured, processed, produced, imported, or transferred in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission pursuant to 10 CFR § 32.22, which license authorizes the transfer of the product to persons who are exempt from regulatory requirements. This exemption does not apply to hydrogen-3, krypton-85, or promethium-147 used in products for frivolous purposes or in toys or adornments.

(5) Any person is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, or owns articles containing less than 3.7 kBq (0.1 <<mu>>Ci) of radium-226 which were acquired prior to October 30, 1986.

(6) Any person, except those who manufacture, process, or produce gas and aerosol detectors containing radioactive material, is exempt from the provisions of this Code to the extent that such person receives, possesses, uses, transfers, owns or acquires radioactive material in gas and aerosol detectors designed to protect life or property from fires and airborne hazards provided that detectors containing radioactive material shall have been manufactured, imported, or transferred in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission pursuant to 10 CFR § 32.26, or by an agreement state pursuant to equivalent regulations.

(i) Gas and aerosol detectors previously manufactured and distributed to general licensees in accordance with a specific license issued by an agreement state shall be considered exempt, provided that the device is labeled in accordance with the specific license authorizing distribution of the generally licensed device, and provided further that they meet requirements equivalent to 10 CFR § 32.26.

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(d) Licensees and contractors of the U.S. Department of Energy and U.S. Nuclear Regulatory Commission. (1) The owner or the person in charge of any radiation installation licensed by the U.S. Department of Energy and/or the U.S. Nuclear Regulatory Commission is exempt from the requirements of this Code provided that such owner or person in charge shall:

(i) afford the Department access to all records which such person is required to maintain pursuant to the U.S. Department of Energy or U.S. Nuclear Regulatory Commission license or contract issued to such person;

(ii) afford the Department opportunity to sample effluents, and to conduct such surveys of levels of radiation and radioactive contamination, as will not substantially interfere with or interrupt for any substantial period of time any activity licensed by or contracted for by the U.S. Department of Energy or U.S. Nuclear Regulatory Commission; and

(iii) afford inspectors or officers of the Department access to any installation in which such radioactive materials are present to accomplish the foregoing review of records, sampling of effluents, and conduct of surveys.

(2) Any U.S. Department of Energy or U.S. Nuclear Regulatory Commission contractor or subcontractor of the following categories operating within the City is exempt from the requirements of this Code to the extent that such contractor, or subcontractor under such contractor, transfers, receives, possesses, uses or acquires sources of radiation:

(i) prime contractors performing work for the U.S. Department of Energy at United States government-owned or controlled sites including the transportation of sources of radiation to or from such sites and the performance of contract services during temporary interruptions of such transportation;

(ii) prime contractors of the U.S. Department of Energy performing research in, or development, manufacture, storage, testing, or transportation of, atomic weapons or components thereof;

(iii) prime contractors of the U.S. Department of Energy using or operating nuclear reactors or other nuclear devices in a United States-owned vehicle or vessel; and

(iv) any other prime contractor or subcontractor of the U.S. Department of Energy or of the U.S. Nuclear Regulatory Commission when the City and the U.S. Nuclear Regulatory Commission jointly determine:

(A) that the exemption of the prime contractor or subcontractor is authorized by law; and

(B) that under the terms of the contract or subcontract, there is adequate assurance that the work thereunder can be accomplished without undue risk to the public health and safety.

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(e) General requirements for issuing specific licenses. (1) The Department will approve an application for, and issue in response thereto, a specific license to receive, produce, possess, use and transfer any radioactive material, if the Department determines that the following requirements have been met:

(i) the applicant's proposed use, equipment, facilities and procedures will protect health and safety and will minimize danger to life and property from radiation hazards; and

(ii) the applicant's radiation detection and measuring instrumentation is appropriate for the radioactive materials and uses thereof requested in the application; and

(iii) the applicant, or the applicant's personnel, if the applicant is not an individual, is qualified by training and experience to use such radioactive material for the purposes covered by the application so as to protect public health and safety and to minimize danger to life and property from radiation hazards; and

(iv) the applicant submits sufficient information to support a determination that the requirements of § 175.101(e)(1)(i) through (iii) are satisfied.

(f) Applications for specific licenses. (1) A license application shall be made in writing on forms prescribed by the Department and shall contain completely and accurately the information required thereon. Such application shall be filed in duplicate (original plus one copy) and may incorporate, by clear specific reference, information contained in any previous application, supplementary statement, notification or report filed with the Department.

(2) Each application or supplementary statement shall be signed by either the applicant personally or a person duly authorized by the applicant to sign for and on the applicant's behalf.

(3) For those applicants or licensees who are required to establish and maintain a radiation safety committee pursuant to this Code, each application or supplementary statement shall be transmitted with a letter signed by the chairman of the radiation safety committee indicating the committee's approval of the requested licensing action.

(4) At any time subsequent to the filing of an application for a license, including amendments, and before the termination of a license issued in response thereto, the Department may require the applicant to submit one or more supplementary statements containing additional information to enable the Department to determine whether such application should be approved or denied, or whether a previously issued license should be amended, suspended or revoked.

(5) A single application may apply for a license covering more than one radioactive material.

(6) Specific licenses issued by the Department shall be in the form of a written authorization permitting possession of certain specific radioactive materials in

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not more than certain specific quantities, and certain specific uses of these radioactive materials. Such possession and use of radioactive materials provided for in the foregoing shall be subject to the requirements of:

(i) all applicable provisions of this Code; and

(ii) all conditions as stated on the license issued by the Department.

(g) Amendments. (1) When a change affecting the licensed operation or facility is considered by a licensee, including but not limited to changes ordered pursuant to this Code, so that the information on file with the Department, either in the initial license application or subsequent requests for amendments, or in the initial license or amendment previously granted, will no longer be accurate, the licensee shall request and receive an amendment for such change prior to causing such change.

(2) Any application by a licensee for a license amendment to conform with the provisions of § 175.101(g)(1) shall be filed in writing with the Department and shall set forth in detail the reasons for such requested amendment. In considering any such application for amendment, the Department shall apply the requirements set forth in § 175.101(e).

(3) A corrective amendment of any license may be issued by the Department at any time upon its initiative.

(4) Any license may be amended or revoked by the Department by reason of the amendment of this Code, or any other applicable law.

(h) Expiration, renewal and termination of licenses. (1) Except as otherwise provided in this Code, each license shall expire at the end of the day on the expiration date stated in the license. If, not less than 30 days prior to such expiration date, a licensee duly files with the Department an application in proper form for license renewal, or for a new and superseding license, the existing license shall not be deemed to have expired until the Department has finally determined such application.

(2) Any application by a licensee for the renewal of such license, including amendments, shall be considered as an application for a license and shall be filed on, and shall contain completely and accurately all information called for by, a written form or other manner prescribed by the Department. In considering any such application for renewal, the Department shall apply the requirements set forth in § 175.101(e).

(3) Each licensee shall notify the Department in writing and request termination of the license when the licensee decides to terminate all activities authorized under the license. This notification and request for termination shall include the reports and information specified in § 175.101(h)(4)(v) and a plan for completion of decommissioning.

(4) If a licensee does not submit an application for renewal pursuant to §

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175.101(h)(1), the licensee shall on or before the expiration date stated in the license:

(i) terminate use of radioactive material;

(ii) dispose of all radioactive material in accordance with all applicable regulations in effect at the time of disposal;

(iii) submit a written certification of the disposition of all radioactive materials authorized by the license on forms prescribed by the Department;

(iv) remove radioactive contamination to the extent practicable; and

(v) conduct a radiation survey of the premises where the licensed activities were carried out and submit a report of the results of this survey to the Department. Such survey shall be subject to confirmation by the Department and shall include:

(A) levels of radiation in units (or multiples) of Gy-hr-1 (millirads-hr-1) at one (1) cm for beta-gamma radiation or at one (1) m for gamma radiation;

(B) levels of removable and fixed contamination, including alpha, in units of disintegrations (transformations) per min (becquerels) per 100 cm 2 for surfaces;

(C) becquerels-ml-1 (<<mu>>Ci-ml-1) for water;

(D) becquerels-g-1 (pCi-ml-1) for solids such as soil or concrete; and

(E) a description of the survey or other measuring instrument(s) used, including manufacturer(s) and model number(s) and date of most recent calibration.

(vi) If the information submitted pursuant to § 175.101(h)(4)(v) does not adequately demonstrate that the premises are suitable for unrestricted use, the Department shall inform the licensee of the appropriate further actions required for the termination of the license, including, but not limited to, decontamination of the licensed premises to such levels and within such time frames as the Department may prescribe.

(vii) Each specific license shall continue in effect, beyond the expiration date if necessary, with respect to possession of residual radioactive materials present as contamination until the Department issues an amendment terminating the license. During this time the licensee shall:

(A) limit activities involving radioactive material to those related to decommissioning; and

(B) continue to control entry to restricted areas until the Department determines they are suitable for release for unrestricted use and the Department issues an amendment terminating the license.

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(viii) The Department will approve a request for termination of a specific license, and issue an amendment terminating such license, when the Department determines that:

(A) radioactive material has been properly disposed; and

(B) premises have been decontaminated to such levels that the total effective dose equivalent (TEDE) from residual radioactivity distinguishable from background radiation, to an average member of the public will not exceed 25 mrem (0.25 mSv) per year;

(C) a radiation survey has been performed which describes all radiation levels and levels of fixed and removable contamination; and

(D) the licensee submits sufficient information to support a determination that the requirements of § 175.101(h)(4)(viii)(A) through (C) have been met.

(i) Amendment, suspension or revocation of licenses. (1) Specific and general licenses shall be subject to amendment, suspension or revocation by reason of amendment of the New York State Public Health Law, enactment or amendment of any other applicable law, amendment of the New York State Sanitary Code, amendment of this Code, or amendment or promulgation of any other applicable rule, regulation or order.

(2) In addition to the provisions of § 5.17 of this Code, the Department may amend, revoke or suspend any license, in whole or in part, for:

(i) Any material misstatement in the application therefor or in any supplementary statement thereto.

(ii) Any condition revealed by such application, supplementary statement, report, record, inspection or other means, which would warrant the Department's refusal to grant a license on an original application.

(iii) Any violation or failure to observe any condition of such license, this Code, or any other applicable rule, regulation or law now or hereafter in effect.

(iv) Failure to notify the Department of a change in ownership or address of a radiation installation.

(j) Emergency response plan. (1) Each application for a license to possess radioactive materials in unsealed form, on foils or plated sources, or sealed in glass and in excess of the quantities specified in Appendix C of this section shall include either:

(i) an evaluation showing that the maximum dose to a person offsite due to a release of radioactive materials would not exceed 10 mSv (1 rem) effective dose equivalent or 50 mSv (5 rem) to the thyroid; or

(ii) an emergency plan for responding to a release of radioactive material.

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(2) One or more of the following factors may be used to support an evaluation submitted pursuant to § 175.101(j)(1)(i):

(i) The radioactive material is physically separated so that only a portion could be involved in an accident.

(ii) All or part of the radioactive material is not subject to release during an accident because of the way it is stored or packaged.

(iii) The release fraction in the respirable size range would be lower than the release fraction shown in Appendix C of this section due to the chemical or physical form of the material.

(iv) The solubility of the radioactive material would reduce the dose received.

(v) Facility design or engineered safety features in the facility would cause the release fraction to be lower than that shown in Appendix C of this section.

(vi) Operating restrictions or procedures would prevent a release fraction as large as that shown in Appendix C of this section.

(vii) Other factors appropriate for the specific facility.

(3) An emergency plan for responding to a release of radioactive material submitted pursuant to § 175.101(j)(1)(ii) shall include the following information:

(i) A brief description of the licensee's facility and area near the site.

(ii) An identification of each type of radioactive materials accident for which protective actions may be needed.

(iii) A classification system for classifying accidents as alerts or site area emergencies.

(iv) Identification of the means of detecting each type of accident in a timely manner.

(v) A brief description of the means and equipment for mitigating the consequences of each type of accident, including those provided to protect workers onsite, and a description of the program for maintaining the equipment.

(vi) A brief description of the methods and equipment to assess releases of radioactive materials.

(vii) A brief description of the responsibilities of licensee personnel should an accident occur, including identification of personnel responsible for promptly notifying offsite response organizations and the Department.

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(viii) A brief description of the responsibilities for developing, maintaining and updating the plan.

(ix) A commitment to, and brief description of, the means to promptly notify offsite response organizations and to request offsite assistance, including medical assistance for the treatment of contaminated injured onsite workers when appropriate. A control point must be established. The notification and coordination must be planned so that unavailability of some personnel, parts of the facility, and some equipment will not prevent the notification and coordination. The licensee shall also commit to notify the Department immediately after notification of the appropriate offsite response organizations and not later than one (1) hour after the licensee declares an emergency.

(x) A brief description of the types of information on facility status, radioactive releases, and recommended protective actions, if necessary, to be given to offsite response organizations and the Department.

(xi) A brief description of the frequency, performance objectives and plans for the training that the licensee will provide workers on how to respond to an emergency including any special instructions and orientation tours the licensee would offer to fire, police, medical and other emergency personnel. The training shall familiarize personnel with site-specific emergency procedures. The training shall thoroughly prepare site personnel for their responsibilities in the event of accident scenarios postulated as most probable for the specific site, including the use of team training for such scenarios.

(xii) A brief description of the means of restoring the facility to a safe condition after an accident.

(xiii) Provisions for conducting quarterly communications checks with offsite response organizations and biennial onsite exercises to test response to simulated emergencies. Quarterly communications checks with offsite response organizations must include the check and update of all necessary telephone numbers. The licensee shall invite offsite response organizations to participate in the biennial exercises. Participation of offsite response organizations is recommended, but not required. Exercises must use accident scenarios postulated as most probable for the specific site and the scenarios shall not be known to most exercise participants. The licensee shall critique each exercise using individuals not having direct implementation responsibility for the plan. Critiques of exercises must evaluate the appropriateness of the plan, emergency procedures, facilities, equipment, training of personnel and overall effectiveness of the response. Plan deficiencies identified in the critiques shall be corrected.

(xiv) A certification that the applicant has met its responsibilities under the Emergency Planning and Community Right-to-Know Act of 1986 codified at 42 USCA 11001 et seq., if applicable to the applicant's activities at the proposed place of use of the radioactive materials.

(4) The licensee shall allow the offsite response organizations expected to respond in case of an accident sixty (60) days to comment on the licensee's emergency response plan before submitting it to the Department. The licensee shall provide

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any comments received within the sixty (60) days to the Department with the plan.

(k) Conditions of specific licenses. (1) Each of the following is hereby made a condition of each specific license:

(i) The licensee thereunder shall comply with all applicable provisions of the New York State Public Health Law, all applicable provisions of this Code, all other laws now or hereafter in effect, and with all applicable rules, regulations, codes and orders now or hereafter in effect of the Department and of all appropriate regulatory agencies.

(ii) Neither such license, nor any right, title or interest in, of or to such license, shall be disposed of by assignment, transfer or otherwise, either voluntarily or involuntarily, either directly or indirectly, unless the Department shall, after securing complete and accurate pertinent information, have approved in writing of such disposal.

(iii) The licensee shall confine the possession and use of radioactive material to such location or locations and for such purpose or purposes as the license may authorize; provided, however, that except as otherwise provided in such license or this Code, such license shall be deemed to authorize the licensee to transfer the material covered by such license to any other person authorized to receive it by the Department, the U.S. Nuclear Regulatory Commission or an agreement state.

(iv) No person, in any advertisement, expressly or by implication, shall refer to the fact that an installation is licensed by the Department, and no person shall state or imply that an installation or its activities have been approved by the Board of Health, the Department or the Commissioner.

(v) The licensee shall notify the Department, in writing, within thirty (30) days if an authorized user, radiation safety officer or radiation therapy physicist permanently discontinues performance of duties under the license.

(vi) Each licensee shall notify the Department, in writing, immediately following the filing of a voluntary or involuntary petition for bankruptcy under any chapter of Title 11 (Bankruptcy) of the United States Code by or against:

(A) the licensee;

(B) an entity (as that term is defined in 11 U.S.C. § 101(14)) controlling the licensee or listing the license or licensee as property of the estate; or

(C) an affiliate (as that term is defined in 11 U.S.C. § 101(2)) of the licensee.

This notification must indicate:

(A) the bankruptcy court in which the petition for bankruptcy was filed; and

(B) the date of the filing of the petition.

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(vii) Any license covering the use of special nuclear material, in the course of which licensed use additional special nuclear material is produced, shall be deemed to cover any such special nuclear material so produced, provided however, that the total quantity of special nuclear material possessed by the licensee is not sufficient to form a critical mass.

(viii) A licensee, employee of a licensee, contractor, subcontractor, or employee of a contractor or subcontractor shall not:

(A) engage in a deliberate misconduct that causes or would have caused if not detected, a licensee or applicant to be in violation of any provision of this Code or license condition issued by the Department; or

(B) deliberately submit to the Department information that the person submitting the information knows to be incomplete or inaccurate in some respect to the Department.

(2) The Department may at any time set forth in any license or incorporate by reference therein, additional conditions, restrictions or requirements applicable to the licensee's receipt, production, possession, use or transfer of radioactive material covered by such license in order to protect the public health and safety and to minimize danger to life and property from radiation hazards.

(3) All licensees subject to the criteria to implement Increased Controls pursuant to the U.S. Nuclear Regulatory Commission (NRC) Order EA 05-090, 70 FR 72128, dated December 1, 2005, shall have as part of their Increased Control Program, a Fingerprinting and Criminal History Records Check procedure established for all individuals whom the licensee wishes to allow unescorted access to radioactive material quantities of concern. Such Fingerprinting and Criminal History Records Check procedures shall adhere to the requirements in NRC Order EA-07-305, 72 FR 70901, or any successor order, law or regulation. The requirements of this provision shall apply to all affected licensees upon its effective date.

(l) Transfer of radioactive materials. (1) No licensee shall transfer radioactive material except as authorized pursuant to this subsection.

(2) Except as provided otherwise by the license, and subject to the provisions of § 175.101(l)(3) and (4), a licensee may transfer radioactive material:

(i) to the Department, after receiving prior written approval of the Department;

(ii) to the U.S. Nuclear Regulatory Commission;

(iii) to any person exempt from the provisions of this Code to the extent permitted under such exemption;

(iv) to any person authorized to receive radioactive material under the terms of a general license or its equivalent, or a specific license or equivalent licensing document, issued by the Department, the U.S. Nuclear Regulatory Commission, or an agreement state, or to any person otherwise authorized to receive radioactive ma-

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terial by the federal government or any agency thereof, the department, or an agreement state; or

(v) as otherwise authorized by the Department in writing.

(3) Before transferring radioactive material to a specific licensee of the Department, the U.S. Nuclear Regulatory Commission or an agreement state, or to a general licensee who is required to register with the Department, the U.S. Nuclear Regulatory Commission or an agreement state prior to the receipt of the radioactive materials, the licensee transferring the radioactive material shall verify that the transferee's license or registration certificate authorizes the receipt of the type, form and quantity of radioactive material to be transferred.

(4) The following methods for the verification required by § 175.101(l)(3) are acceptable:

(i) the transferor may have in his/her possession, and read, a current copy of the transferee's specific license or registration certificate; or

(ii) the transferor may have in his/her possession a written certification by the transferee that he/she is authorized by license or registration certificate to receive the type, form and quantity of radioactive material to be transferred, specifying the license or registration certificate number, issuing agency and expiration date; or

(iii) for emergency shipments the transferor may accept oral certification by the transferee that he is authorized by license or registration certificate to receive the type, form, and quantity of radioactive material to be transferred, specifying the license or registration certificate number, issuing agency and expiration date, provided that the oral certification is confirmed in writing within 10 days; or

(iv) the transferor may obtain other sources of information compiled by a reporting service from official records of the Department, the U.S. Nuclear Regulatory Commission or an agreement state as to the identity of licensees and the scope and expiration dates of licenses and registrations; or

(v) when none of the methods of verification described in § 175.101(l)(4)(i) through (iv) are readily available, or when a transferor desires to verify that information received by one of such methods is correct or up-to-date, the transferor may obtain and record confirmation from the Department, the U.S. Nuclear Regulatory Commission, or an agreement state that the transferee is licensed to receive the radioactive material.

(5) Preparation for shipment and transport of radioactive material shall be in accordance with the provisions of § 175.105 of this Code.

(m) Reciprocity. (1) The holder of a license issued by the New York State Department of Labor, the New York State Department of Health, the U.S. Nuclear Regulatory Commission or any agreement state, may bring, possess or use radioactive material covered by such license within the Department's jurisdiction for a period not in

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excess of 180 days in any twelve consecutive months without obtaining a license from the Department, provided that:

(i) such license does not limit the holder's possession or use of such material to a specific installation or installations;

(ii) such holder, prior to bringing such material into the city, files with the Department a notice indicating the period, type and location of proposed possession and use within the Department's jurisdiction, and a copy of the license;

(iii) such holder supplies such additional information as the Department may reasonably request;

(iv) such holder, during the period of this possession and use of such material within the city, complies with all applicable sections of this Code except § 175.101(a)(1); and

(v) such holder, during such period, complies with all the terms and conditions of his license, except such terms or conditions which may be inconsistent with this Code.

(2) The holder of a license issued by the New York State Department of Labor, the New York State Department of Health, the New York City Department of Health and Mental Hygiene or an Agreement State must obtain reciprocity approval from the U.S. Nuclear Regulatory Commission to conduct licensed activity in areas of exclusive federal jurisdiction within New York City. At least three days before engaging in each activity for the first time in a calendar year, the licensee will provide the U.S. Nuclear Regulatory Commission with advanced notice of its proposed activity in areas under exclusive federal jurisdiction within New York City.

(n) Financial assurance and recordkeeping for decommissioning.

(1)(a) Each applicant for a specific license authorizing the possession and use of unsealed radioactive material of half-life greater than 120 days and in quantities exceeding 10 5 times the applicable quantities set forth in Appendix B to this section shall submit a decommissioning funding plan as described in § 175.101(n)(5). The decommissioning funding plan must also be submitted when a combination of isotopes is involved if R divided by 10 5 is greater than one (1) (unity rule), where R is defined here as the sum of the ratios of the quantity of each isotope to the applicable value in Appendix B to this section.

(1)(b) Each holder of, or applicant for, any specific license authorizing the possession and use of sealed sources or plated foils of half-life greater than 120 days and in quantities exceeding 10 times the applicable quantities set forth in Appendix B to this section shall submit a decommissioning funding plan as described in § 175.101(n)(5). The decommissioning funding plan must also be submitted when a combination of isotopes is involved if R, as defined in § 175.101(n)(1)(a), divided by 10 1 2 is greater than one (1) (unity rule). The decommissioning funding plan must be submitted to the Department within 2 years of the effective date of this provision.

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(1)(c) Each applicant for a specific license authorizing the possession and use of more than 100 mCi of source material in a readily dispersible form shall submit a decommissioning funding plan as described in § 175.101(n)(5).

(1)(d) Each applicant for a specific license authorizing possession and use of quantities of source material greater than 10 mCi but less than or equal to 100 mCi in a readily dispersible form shall either:

(i) Submit a decommissioning funding plan as described in § 175.101(n)(5); or

(ii) Submit a certification that financial assurance for decommissioning has been provided in the amount of $225,000 within eighteen months of the effective date of this provision using one of the methods described in § 175.101(n)(6). For an applicant, this certification may state that the appropriate assurance will be obtained after the application has been approved and the license issued but before the receipt of licensed material. If the applicant defers execution of the financial instrument until after the license has been issued, a signed original of the financial instrument obtained to satisfy the requirements of paragraph § 175.101(n)(6) of this section must be submitted to the Department prior to receipt of licensed material. If the applicant does not defer execution of the financial instrument, the applicant shall submit to the Department as part of the certification, a signed original of the financial instrument obtained to satisfy the requirements of paragraph § 175.101(n)(6) of this section.

(1)(e) Each applicant for a specific license authorizing the possession and use of unsealed special nuclear material in quantities exceeding 10 5 times the applicable quantities set forth in Appendix B to this section shall submit a decommissioning funding plan as described in § 175.101(n)(5). A decommissioning funding plan must also be submitted when a combination of isotopes is involved if R, as defined in § 175.101(n)(1)(a), divided by 10 5 is greater than one (1) (unity rule).

(2) Each applicant for a specific license authorizing possession and use of radioactive material of half-life greater than 120 days and in quantities specified in § 175.101(n)(4) shall either:

(i) submit a decommissioning funding plan as described in § 175.101(n)(4); or

(ii) submit a certification that financial assurance for decommissioning has been provided in the amount prescribed by § 175.101(n)(4) using one of the methods described in § 175.101(n)(6). For an applicant, this certification may state that the appropriate assurance will be obtained after the application has been approved and the license issued but before the receipt of licensed material. If the applicant defers execution of the financial instrument until after the license has been issued, a signed original of the financial instrument obtained to satisfy the requirements of paragraph § 175.101(n)(6) of this section must be submitted to the Department prior to receipt of licensed material. If the applicant does not defer execution of the financial instrument, the applicant shall submit to the Department as part of the certification, a signed original of the financial instrument obtained to satisfy the requirements of paragraph § 175.101(n)(6) of this section.

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(3) (i) Each holder of a specific license issued on or after July 27, 1990, which is of a type described in § 175.101(n)(1) or (2), shall provide financial assurance for decommissioning in accordance with the criteria set forth in this section.

(ii) Each holder of a specific license issued before July 27, 1990, and of a type described in § 175.101(n)(1) or (2), shall submit a decommissioning funding plan as described in § 175.101(n)(5) or a certification of financial assurance for decommissioning in an amount at least equal to $1,125,000 in accordance with the criteria set forth in this section. If the licensee submits the certification of financial assurance rather than a decommissioning funding plan, the licensee shall include a decommissioning funding plan in any application for license renewal.

(iii) Each holder of a specific license issued before July 27, 1990, and of a type described in § 175.101(n)(2), shall submit a certification of financial assurance for decommissioning or a decommissioning funding plan in accordance with the criteria set forth in this section.

(4) Table of required amounts of financial assurance for decommissioning by quantity of material. Licensees required to submit the $1,125,000 amount must do so within 1 year of the effective date of this provision. Licensees required to submit the $113,000 or $225,000 amount must do so within 18 months of the effective date of this provision. Licensees having possession limits exceeding the upper bounds of this table must base financial assurance on a decommissioning funding plan.

(i) Greater than 10 4 but less than or equal to 10 5 times the applicable quantities of Appendix B to this section in unsealed form (for a combination of isotopes, if R, as defined in § 175.101(n)(1)(a), divided by 10 4 is greater than 1, but R divided by 10 5 is less than or equal to 1)--$1,125,000.

(ii) Greater than 10 3 but less than or equal to 10 4 times the applicable quantities of Appendix B to this section in unsealed form (for a combination of isotopes, if R, as defined in § 175.101(n)(1)(a), divided by 10 3 is greater than 1, but R divided by 10 4 is less than or equal to 1)--$225,000.

(iii) Greater than 10 1 0 but less than or equal to 10 1 2 times the applicable quantities of Appendix B to this section in sealed sources or plated foils (for a combination of isotopes, if R, as defined in § 175.101(n)(1)(a), divided by 10 1 0 is greater than 1, but R divided by 10 1 2 is less than or equal to 1)--$113.000.

(5) Each decommissioning funding plan must contain a cost estimate for decommissioning and a description of the method of assuring funds for decommissioning from § 175.101(n)(6), including means of adjusting cost estimates and associated funding levels periodically over the life of the facility. Cost estimates must be adjusted at intervals not to exceed 3 years. The decommissioning funding plan must also contain a certification by the licensee that financial assurance for decommissioning has been provided in the amount of the cost estimate for decommissioning and a signed original of the financial instrument obtained to satisfy the requirements of § 175.101(n)(6).

(6) Financial assurance for decommissioning must be provided by one or more of the following methods:

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(i) Prepayment. Prepayment is the deposit prior to the start of operation into an account segregated from licensee assets and outside the licensee's administrative control of cash or liquid assets such that the amount of funds would be sufficient to pay decommissioning costs. Prepayment may be in the form of a trust, escrow account, government fund, certificate of deposit, or deposit of government securities.

(ii) A surety method, insurance, or other guarantee method. These methods guarantee that decommissioning costs will be paid. A surety method may be in the form of a surety bond, letter of credit, or line of credit. A parent company guarantee of funds for decommissioning costs based on a financial test may be used if the guarantee and test are as contained in Appendix D. A parent company guarantee may not be used in combination with other financial methods to satisfy the requirements of this section. A guarantee of funds by the applicant or licensee for decommissioning costs based on a financial test may be used if the guarantee and test are as contained in Appendix E. A guarantee by the applicant or licensee may not be used in combination with any other financial methods to satisfy the requirements of this section or in any situation where the applicant or licensee has a parent company holding majority control of the voting stock of the company. Any surety method or insurance used to provide financial assurance for decommissioning must contain the following conditions:

(A) the surety method or insurance must be open-ended or, if written for a specified term, such as five years, must be renewed automatically unless 90 days or more prior to the renewal date, the issuer notifies the Department, the beneficiary, and the licensee of its intention not to renew. The surety method or insurance must also provide that the full face amount be paid to the beneficiary automatically prior to the expiration without proof of forfeiture if the licensee fails to provide a replacement acceptable to the Department within 30 days after receipt of notification of cancellation.

(B) The surety method or insurance must be payable to a trust established for decommissioning costs. The trustee and trust must be acceptable to the Department. An acceptable trustee includes an appropriate state or federal government agency or an entity which has the authority to act as a trustee and whose trust operations are regulated and examined by a federal or state agency.

(C) The surety method or insurance must remain in effect until the Department has terminated the license.

(iii) An external sinking fund in which deposits are made at least annually, coupled with a surety method or insurance, the value of which may decrease by the amount being accumulated in the sinking fund. An external sinking fund is a fund established and maintained by setting aside funds periodically in an account segregated from licensee assets and outside the licensee's administrative control in which the total amount of funds would be sufficient to pay decommissioning costs at the time termination of operation is expected. An external sinking fund may be in the form of a trust, escrow account, government fund, certificate of deposit, or depost of government securities. The surety or insurance provisions must be as stated in § 175.101(n)(6)(ii).

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(iv) In the case of federal, state or local government licensees, a statement of intent containing a cost estimate for decommissioning or an amount based on the Table in § 175.101(n)(4), and indicating that funds for decommissioning will be obtained when necessary.

(7) Each licensed person shall keep records of information important to the safe and effective decommissioning of the facility in an identified location until the license is terminated by the Department and the site is released for unrestricted use. If records of relevant information are kept for other purposes, reference to these records and their location may be used. Information the Department considers important to decommissioning consists of:

(i) Records of spills or other unusual occurrences involving the spread of contamination in and around the facility, equipment or site. These records may be limited to instances when contamination remains after any cleanup procedures or when there is reasonable likelihood that contaminants may have spread to inaccessible areas as in the case of possible seepage into porous materials such as concrete. These records must include any known information on identification of involved nuclides, quantities, forms and concentrations.

(ii) As-built drawings and modifications of structures and equipment in restricted areas where radioactive materials are used and/or stored, and of locations of possible inaccessible contamination such as buried pipes which may be subject to contamination. If required drawings are referenced, each relevant document need not be indexed individually. If drawings are not available, the licensee shall substitute appropriate records of available information concerning these areas and locations.

(iii) Except for areas containing only sealed sources (provided the sources have not leaked or no contamination remains after any leak) or radioactive materials having only half-lives of less than 65 days, a list contained in a single document and updated every two (2) years, of the following:

(A) All areas designated and formerly designated restricted areas as defined in § 175.02(a)(194);

(B) all areas outside of restricted areas that require documentation under § 175.101(n)(7)(i);

(C) all areas where current and/or previous wastes have been buried; and

(D) all areas outside of restricted areas which contain material such that, if the license expired, the licensee would be required to decontaminate the area to unrestricted release levels or to apply for approval for disposal under § 175.104(b).

(iv) Records of the cost estimate performed for the decommissioning funding plan or of the amount certified for decommissioning, and records of the funding method used for assuring funds if either a funding plan or certification is used.

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HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Subd. (h) par (4) subpar (vi) amended City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (h) par (4) subpar (viii) clause (B) amended City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (k) par (1) subpar (viii) added City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (k) par (3) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (m) par (2) amended City Record June 20, 2007 eff. July 20, 2007. [See Vol. 9 Statements of Basis and Purpose No. 28]

Subd. (m) par (2) added City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (n) par (1) amended City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (n) par (1)(a) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (1)(b) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (1)(c) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (1)(d) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (1)(e) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (2) open par amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (2) subpar (ii) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

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Subd. (n) par (3) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (4) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statements of Basis and Purpose No. 30]

Subd. (n) par (5) amended City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Subd. (n) par (7) open par amended City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

On September 17, 2008, the Board of Health amended Health Code § 175.101(k) pursuant to NRC Order EA-07-305 to incorporate a fingerprinting and criminal history records check requirement on licensees, who possess certain radioactive materials in quantities of concern. This requirement applies to those employees of affected licensees who are allowed unescorted access to certain radioactive materials in quantities of concern. Also, the Board amended several provisions of § 175.101(n) of the Health Code, including the addition of new § 175.101(n)(1) (c)-(e), in order to assure compatibility with applicable NRC regulations concerning financial assurance and decommissioning of a licensed facility. The NRC amended its regulations (e.g., 10 CFR §§ 30.35, 40.36, and 70.25) for financial assurance for certain materials licensees to bring the amount of financial assurance required more in line with current decommissioning costs. The objective of this action is to ensure that licensees maintain adequate financial assurance so that timely decommissioning can be carried out following shutdown of a licensed facility.

§ 175.101(m) in Article 175 of the NYC Health Code was amended on June 14, 2007 to provide reciprocity approval by the U.S. Nuclear Regulatory Commission (NRC). NRC required that the Department amend the NYC Health Code to include a 3-day advance notice requirement to NRC for prior approval to any initial activity by any Agreement State licensee/non-NRC licensee in areas under exclusive federal jurisdiction within NYC, as per 10 CFR § 150.20.

Subparagraph (vi) of paragraph (4) of subsection (h) of § 175.101 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 30.36 in assuring that licensees who wish to terminate licensed activities must perform decommissioning within timeframes prescribed by the Department.

Subparagraph (viii)(B) of paragraph (4) of subsection (h) of § 175.101 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 20.1403, which provides specific dose limits to members of the public from sites released for unrestricted use.

Subparagraph (viii) of paragraph (1) of subsection (k) of § 175.101 of the New York City Health Code was added on March 10, 2005 to reflect federal regulation 10

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§ 30.10 which enables the Department to conduct enforcement actions against individuals who provide inaccurate information to the Department.

Paragraph (2) of subsection (m) of § 175.101 of the New York City Health Code was added on March 10, 2005 to reflect 10 CFR § 150.20 which clarifies that Agreement State licensees can seek reciprocal recognition of their license from the NRC when working within areas of exclusive Federal jurisdiction in New York City.

Paragraph (1)(b) of subsection (n) of § 175.101 of the New York City Health Code was added on March 10, 2005 to reflect 10 CFR § 30.35 which requires licensees that are authorized to possess sealed sources or plated foils of half-life greater than 120 days and in quantities exceeding 10 to the 12th times the applicable quantities set forth in Appendix B of the New York City Health Code to submit a decommissioning funding plan.

Paragraph (5) of subsection (n) of § 175.101 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 30.35, which addresses adjusting cost estimates for decommissioning at intervals not to exceed 3 years.

Paragraph (7) of subsection (n) of § 175.101 of the New York City Health Code was amended on March 10, 2005 to reflect 10 CFR § 30.35.

APPENDIX A
EXEMPT CONCENTRATIONS

Element (atomic number)	Isotope	Column I Gas concentration <<mu>>Ci/ml 1	Column II Liquid and solid concentration <<mu>>Ci/ml 2

APPENDIX A (cont'd)
EXEMPT CONCENTRATIONS

Element (atomic number)	Isotope	Column I Gas concentration <<mu>>Ci/ml 1	Column II Liquid and solid concentration <<mu>>Ci/ml 2
Antimony (51)	Sb 122		3E-4
	Sb 124		2E-4
	Sb 125		1E-3
Argon (18)	A 37	1E-3	
	A 41	4E-7	
Arsenic (33)	As 73		5E-3
	As 74		5E-4
	As 76		2E-4
	As 77		8E-4
Barium (56)	Ba 131		2E-3
	Ba 140		3E-4
Beryllium (4)	Be 7		2E-2
Bismuth (83)	Bi 206		4E-4
Bromine (35)	Br 82	4E-7	3E-3
Cadmium (48)	Cd 109		2E-3

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	Cd 115m		3E-4
	Cd 115		3E-4
Calcium (20)	Ca 45		9E-5
	Ca 47		5E-4
Carbon (6)	C 14	1E-6	8E-3
Cerium (58)	Ce 141		9E-4
	Ce 143		4E-4
	Ce 144		1E-4
Cesium (55)	Cs 131		2E-2
	Cs 134m		6E-2
	Cs 134		9E-5
Chlorine (17)	Cl 38	9E-7	4E-3
Chromium (24)	Cr 51		2E-2
Cobalt (27)	Co 57		5E-3
	Co 58		1E-3
	Co 60		5E-4
Copper (29)	Cu 64		3E-3
Dysprosium (66)	Dy 165		4E-3
	Dy 166		4E-4
Erbium (68)	Er 169		9E-4
	Er 171		1E-3
Europium (63)	Eu 152		6E-4
	(T/2=9.2 Hrs)		
	Eu 155		2E-3
Fluorine (9)	F 18	2E-6	8E-3
Gadolinium (64)	Gd 153		2E-3
	Gd 159		8E-4
Gallium (31)	Ga 72		4E-4
Germanium (32)	Ge 71		2E-2
Gold (79)	Au 196		2E-3
	Au 198		5E-4
	Au 199		2E-3
Hafnium (72)	Hf 181		7E-4
Hydrogen (1)	H 3	5E-6	3E-2
Indium (49)	In 113m		1E-2
	In 114m		2E-4
Iodine (53)	I 126	3E-9	2E-5
	I 131	3E-9	2E-5
	I 132	8E-8	6E-4
Iodine (53)	I 133	1E-8	7E-5
	I 134	2E-7	1E-3
Iridium (77)	Ir 190		2E-3
	Ir 192		4E-4
	Ir 194		3E-4
Iron (26)	Fe 55		8E-3
	Fe 59		6E-4
Krypton (36)	Kr 85m	1E-6	
	Kr 85	3E-6	
Lanthanum (57)	La 140		2E-4
Lead (82)	Pb 203		4E-3
Lutetium (71)	Lu 177		1E-3
Manganese (25)	Mn 52		3E-4
	Mn 54		1E-3

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	Mn 56		1E-3
Mercury (80)	Hg 197m		2E-3
	Hg 197		3E-3
	Hg 203		2E-4
Molybdenum (42)	Mo 99		2E-3
Neodymium (60)	Nd 147		6E-4
	Nd 149		3E-3
Nickel (28)	Ni 65		1E-3
Niobium (Columbium) (41)	Nb 95 Nb 97		1E-3 9E-3
Osmium (76)	Os 185		7E-4
	Os 191m		3E-2
	Os 191		2E-3
	Os 193		6E-4
Palladium (46)	Pd 103		3E-3
	Pd 109		9E-4
Phosphorus (15)	P 32		2E-4
Platinum (78)	Pt 191		1E-3
	Pt 193m		1E-2
	Pt 197m		1E-2
	Pt 197		1E-3
Polonium (84)	Po 210	2E-10	7E-6
Potassium (19)	K 42		3E-3
Praseodymium (59)	Pr 142		3E-4
	Pr 143		5E-4
Promethium (61)	Pm 147		2E-3
	Pm 149		4E-4
Radium (88)	Ra 226	1E-11	1E-7
	Ra 228	2E-11	3E-7
Rhenium (75)	Re 183		6E-3
	Re 186		9E-4
	Re 188		6E-4
Rhodium (45)	Rh 103m		1E-1
	Rh 105		1E-3
Rubidium (37)	Rb 86		7E-4
Ruthenium (44)	Ru 97		4E-3
	Ru 103		8E-4
	Ru 105		1E-3
	Ru 106		1E-4
Samarium (62)	Sm 153		8E-4
Scandium (21)	Sc 46		4E-4
	Sc 47		9E-4
	Sc 48		3E-4
Selenium	Se 75		3E-3
Silicon (14)	Si 31		9E-3
Silver (47)	Ag 105		1E-3
	Ag 110m		3E-4
	Ag 111		4E-4
Sodium (11)	Na 24		2E-3
Strontium (38)	Sr 85		1E-3
	Sr 89		1E-4

APPENDIX A

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EXEMPT CONCENTRATIONS

Element (atomic number) Isotope Column I Gas Column II Liquid and

concentration solid

Ci/ml 1 concentration

Ci/ml 2

APPENDIX A (cont'd)

EXEMPT CONCENTRATIONS

Element (atomic number) Isotope Column I Gas Column II Liquid and

concentration solid

Ci/ml 1 concentration

Ci/ml 2

Antimony (51) Sb 122 3E-4

Sb 124 2E-4

Sb 125 1E-3

Argon (18) A 37 1E-3

A 41 4E-7

Arsenic (33) As 73 5E-3

As 74 5E-4

As 76 2E-4

As 77 8E-4

Barium (56) Ba 131 2E-3

Ba 140 3E-4

Beryllium (4) Be 7 2E-2

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Bismuth (83)	Bi 206	4E-4	
Bromine (35)	Br 82	4E-7	3E-3
Cadmium (48)	Cd 109	2E-3	
Cd 115m	3E-4		
Cd 115	3E-4		
Calcium (20)	Ca 45	9E-5	
Ca 47	5E-4		
Carbon (6)	C 14	1E-6	8E-3
Cerium (58)	Ce 141	9E-4	
Ce 143	4E-4		
Ce 144	1E-4		
Cesium (55)	Cs 131	2E-2	
Cs 134m	6E-2		
Cs 134	9E-5		
Chlorine (17)	Cl 38	9E-7	4E-3
Chromium (24)	Cr 51	2E-2	
Cobalt (27)	Co 57	5E-3	
Co 58	1E-3		
Co 60	5E-4		
Copper (29)	Cu 64	3E-3	
Dysprosium (66)	Dy 165	4E-3	
Dy 166	4E-4		
Erbium (68)	Er 169	9E-4	
Er 171	1E-3		

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Europium (63) Eu 152 6E-4

(T/2=9.2 Hrs)

Eu 155 2E-3

Fluorine (9) F 18 2E-6 8E-3

Gadolinium (64) Gd 153 2E-3

Gd 159 8E-4

Gallium (31) Ga 72 4E-4

Germanium (32) Ge 71 2E-2

Gold (79) Au 196 2E-3

Au 198 5E-4

Au 199 2E-3

Hafnium (72) Hf 181 7E-4

Hydrogen (1) H 3 5E-6 3E-2

Indium (49) In 113m 1E-2

In 114m 2E-4

Iodine (53) I 126 3E-9 2E-5

I 131 3E-9 2E-5

I 132 8E-8 6E-4

Iodine (53) I 133 1E-8 7E-5

I 134 2E-7 1E-3

Iridium (77) Ir 190 2E-3

Ir 192 4E-4

Ir 194 3E-4

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Iron (26) Fe 55 8E-3

Fe 59 6E-4

Krypton (36) Kr 85m 1E-6

Kr 85 3E-6

Lanthanum (57) La 140 2E-4

Lead (82) Pb 203 4E-3

Lutetium (71) Lu 177 1E-3

Manganese (25) Mn 52 3E-4

Mn 54 1E-3

Mn 56 1E-3

Mercury (80) Hg 197m 2E-3

Hg 197 3E-3

Hg 203 2E-4

Molybdenum (42) Mo 99 2E-3

Neodymium (60) Nd 147 6E-4

Nd 149 3E-3

Nickel (28) Ni 65 1E-3

Niobium (Columbium) (41) Nb 95 Nb 97 1E-3 9E-3

Osmium (76) Os 185 7E-4

Os 191m 3E-2

Os 191 2E-3

Os 193 6E-4

Palladium (46) Pd 103 3E-3

Pd 109 9E-4

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Phosphorus (15) P 32 2E-4

Platinum (78) Pt 191 1E-3

Pt 193m 1E-2

Pt 197m 1E-2

Pt 197 1E-3

Polonium (84) Po 210 2E-10 7E-6

Potassium (19) K 42 3E-3

Praseodymium (59) Pr 142 3E-4

Pr 143 5E-4

Promethium (61) Pm 147 2E-3

Pm 149 4E-4

Radium (88) Ra 226 1E-11 1E-7

Ra 228 2E-11 3E-7

Rhenium (75) Re 183 6E-3

Re 186 9E-4

Re 188 6E-4

Rhodium (45) Rh 103m 1E-1

Rh 105 1E-3

Rubidium (37) Rb 86 7E-4

Ruthenium (44) Ru 97 4E-3

Ru 103 8E-4

Ru 105 1E-3

Ru 106 1E-4

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Samarium (62)	Sm 153	8E-4		
Scandium (21)	Sc 46	4E-4		
	Sc 47	9E-4		
	Sc 48	3E-4		
Selenium	Se 75	3E-3		
Silicon (14)	Si 31	9E-3		
Silver (47)	Ag 105	1E-3		
	Ag 110m	3E-4		
	Ag 111	4E-4		
Sodium (11)	Na 24	2E-3		
Strontium (38)	Sr 85	1E-3		
	Sr 89	1E-4		
	Sr 91		7E-4	
	Sr 92		7E-4	
Sulfur (16)	S 35	9E-8	6E-4	
Tantalum (73)	Ta 182		4E-4	
Technetium (43)	Tc 96m		1E-1	
	Tc 96		1E-3	
Tellurium (52)	Te 125m		2E-3	
	Te 127m		6E-4	
	Te 127		3E-3	
	Te 129m		3E-4	
	Te 131m		6E-4	
	Te 132		3E-4	
Terbium (65)	Tb 160		4E-4	
Thallium (81)	Tl 200		4E-3	
	Tl 201		3E-3	
	Tl 202		1E-3	
	Tl 204		1E-3	
Thulium (69)	Tm 170		5E-4	
	Tm 171		5E-3	
Tin (50)	Sn 113		9E-4	
	Sn 125		2E-4	
Tungsten (Wolfram) (74)	W 181 W 187		4E-3	7E-4
Vanadium (23)	V 48		3E-4	
Xenon (54)	Xe 131m	4E-6		
	Xe 133	3E-6		
	Xe 135	1E-6		

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Ytterbium (70)	Yb 175		1E-3
Yttrium (39)	Y 90		2E-4
	Y 91m		3E-2
	Y 91		3E-4
	Y 92		6E-4
	Y 93		3E-4
Zinc (30)	Zn 65		1E-3
	Zn 69m		7E-4
	Zn 69		2E-2
Zirconium (40)	Zr 95		6E-4
	Zr 97		2E-4
Beta and/or gramma emitting radioactive material not listed above with half-life less than 3 years.		1E-10	1E-6
	Sr 91	7E-4	
	Sr 92	7E-4	
Sulfur (16)	S 35	9E-8	6E-4
Tantalum (73)	Ta 182		4E-4
Technetium (43)	Tc 96m		1E-1
	Tc 96	1E-3	
Tellurium (52)	Te 125m		2E-3
	Te 127m	6E-4	
	Te 127	3E-3	
	Te 129m	3E-4	
	Te 131m	6E-4	
	Te 132	3E-4	
Terbium (65)	Tb 160		4E-4
Thallium (81)	Tl 200		4E-3
	Tl 201	3E-3	
	Tl 202	1E-3	
	Tl 204	1E-3	

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Thulium (69) Tm 170 5E-4

Tm 171 5E-3

Tin (50) Sn 113 9E-4

Sn 125 2E-4

Tungsten (Wolfram) (74) W 181 W 187 4E-3 7E-4

Vanadium (23) V 48 3E-4

Xenon (54) Xe 131m 4E-6

Xe 133 3E-6

Xe 135 1E-6

Ytterbium (70) Yb 175 1E-3

Yttrium (39) Y 90 2E-4

Y 91m 3E-2

Y 91 3E-4

Y 92 6E-4

Y 93 3E-4

Zinc (30) Zn 65 1E-3

Zn 69m 7E-4

Zn 69 2E-2

Zirconium (40) Zr 95 6E-4

Zr 97 2E-4

Beta and/or gramma emitting radioactive 1E-10 1E-6

material not listed above with half-life

less than 3 years.

FOOTNOTES:

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1 Values are given only for those materials normally used as gases.

2 <<mu>>Ci-g-1 for solids.

NOTE 1: Many radioisotopes disintegrate into isotopes which are also radioactive. In expressing the concentrations in Appendix A, the activity stated is that of the parent isotope and takes into account the daughters.

NOTE 2: For purposes of § 175.101(c)(1) where there is involved a combination of isotopes, the limit for the combination should be derived as follows:

Determine for each isotope in the product the ratio between the concentration present in the product and the exempt concentration present in the product and the exempt concentration established in Schedule A for the specific isotope when not in combination. The sum of such ratios may not exceed "1in. (i.e., unity).

Example:

Concentration of Isotope A in Product 1 / Exempt concentration of Isotope A + Concentration of Isotope B in Product 1 / Exempt concentration of Isotope B # 1

Concentration of Isotope A in Product 1 / Exempt concentration of Isotope A Concentration of Isotope B in Product 1 / Exempt concentration of Isotope B # 1

HISTORICAL NOTE

Appendix A in original publication July 1, 1991.

Polonium (84) added City Record Apr. 3, 2001 eff. May 3, 2001.

Radium (88) added City Record Apr. 3, 2001 eff. May 3, 2001.

NOTES:

Appendix A was amended on March 21, 2001 to be consistent with the New York State Sanitary Code, Appendix 16-A, Table 2.

APPENDIX B
EXEMPT QUANTITIES

APPENDIX B

EXEMPT QUANTITIES

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Radioactive material Microcuries 1

Radioactive material Microcuries 1

APPENDIX B

APPENDIX B

24 RCNY Hlth. Code § 175.101
END OF DOCUMENT

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

(cont'd)

EXEMPT QUANTITIES

TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

Radioactive material	Microcuries 1
Americium 241	.01
Antimony 122 (Sb 122)	100
Antimony 124 (Sb 124)	10
Antimony 125 (Sb 125)	10
Arsenic 73 (As 73)	100
Arsenic 74 (As 74)	10
Arsenic 76 (As 76)	10
Arsenic 77 (As 77)	100
Barium 131 (Ba 131)	10
Barium 133 (Ba 133)	10
Barium 140 (Ba 140)	10
Beryllium 7 (Be 7)	100
Bismuth 210 (Bi 210)	1
Bromine 82 (Br 82)	10
Cadmium 109 (Cd 109)	10
Cadmium 115m (Cd 115m)	10
Cadmium 115 (Cd 115)	100
Calcium 45 (Ca 45)	10
Calcium 47 (Ca 47)	10
Carbon 14 (C14)	100
Cerium 141 (Ce 141)	100
Cerium 143 (Ce 143)	100
Cerium 144 (Ce 144)	1
Cesium 131 (Cs 131)	1,000
Cesium 134m (Cs 134m)	100
Cesium 134 (Cs 134)	1
Cesium 135 (Cs 135)	10

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Cesium 136 (Cs 136)	10
Cesium 137 (Cs 137)	10
Chlorine 36 (Cl 36)	10
Chlorine 38 (Cl 38)	10
Chromium 51 (Cr 51)	1,000
Cobalt 58m (Co 58m)	10
Cobalt 58 (Co 58)	10
Cobalt 60 (Co 60)	1
Copper 64 (Cu 64)	100
Dysprosium 165 (Dy 165)	10
Dysprosium 166 (Dy 166)	100
Erbium 169 (Er 169)	100
Erbium 171 (Er 171)	100
Europium 152 9.2h (Eu 152 9.2h)	100
Europium 152 13 yr (Eu 152 13yr)	1
Europium 154 (Eu 154)	1
Europium 155 (Eu 155)	10
Fluorine 18 (F 18)	1,000
Gadolinium 153 (Gd 153)	10
Gadolinium 159 (Gd 159)	100
Gallium 72 (Ga 72)	10
Germanium 71 (Ga 71)	100
Gold 198 (Au 198)	100
Gold 199 (Au 199)	100
Hafnium 181 (Hf 181)	10
Holmium 166 (Ho 166)	100
Hydrogen 3 (H3)	1,000
Indium 113m (In 113m)	100
Indium 114m (In 114m)	10
Indium 114 (In 114)	1
Indium 115m (In 115m)	100
Indium 115 (In 115)	10
Iodine 125 (I 125)	1
Iodine 126 (I 126)	1
Iodine 129 (I 129)	0.1
Iodine 131 (I 131)	1
Iodine 132 (I 132)	10
Iodine 133 (I 133)	1
Iodine 134 (I 134)	10
Iodine 135 (I 135)	10
Iridium 192 (Ir 192)	10
Iridium 194 (Ir 194)	100
Iron 55 (Fe 55)	100
Iron 59 (Fe 59)	10
Krypton 85 (Kr 85)	100
Krypton 87 (Kr 87)	10
Lanthanum 140 (La 140)	10
Lutetium 177 (Lu 177)	100
Manganese 52 (Mn 52)	10
Manganese 54 (Mn 54)	10
Manganese 56 (Mn 56)	10
Mercury 197m (Hg 197m)	100
Mercury 197 (Hg 197)	100

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Mercury 203 (Hg 203)	10
Molybdenum 99 (Mo 99)	100
Neodymium 147 (Nd 147)	100
Neodymium 149 (Nd 149)	100
Nickel 59 (Ni 59)	100
Nickel 63 (Ni 63)	10
Nickel 65 (ni 65)	100
Niobium 93m (Nb 93m)	10
Niobium 95 (Nb 95)	10
Niobium 97 (Nb 97)	10
Osmium 185 (Os 185)	10
Osmium 191m (Os 191m)	100
Osmium 191 (Os 191)	100
Osmium 193 (Os 193)	100
Palladium 103 (Pd 103)	100
Palladium 109 (Pd 109)	100
Phosphorous 32 (P 32)	10
Platinum 191 (Pt 191)	100
Platinum 193m (Pt 193m)	100
Platinum 193 (Pt 193)	100
Platinum 197m (Pt 197m)	100
Platinum 197 (Pt 197)	100
Plutonium 239	.01
Polonium 210 (Po 210)	0.1
Potassium 42 (K 42)	10
Praseodymium 142 (Pr 142)	100
Praseodymium 143 (Pr 143)	100
Promethium 147 (Pm 147)	10
Promethium 149 (Pm 149)	10
Radium 226 (Ra 226)	0.1
Rhenium 186 (Re 186)	100
Rhenium 188 (Re 188)	100
Rhodium 103m (Rh 103m)	100
Rhodium 105 (Rh 105)	100
Rubidium 86 (Rb 86)	10
Rubidium 87 (Rb 87)	10
Ruthenium 97 (Ru 97)	100
Ruthenium 103 (Ru 103)	10
Ruthenium 105 (Ru 105)	10
Ruthenium 106 (Ru 106)	1
Samarium 151 (Sm 151)	10
Samarium 153 (Sm 153)	100
Scandium 46 (Sc 46)	10
Scandium 47 (Sc 47)	100
Scandium 48 (Sc 48)	10

TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

Selenium 75 (Sc 75)	10
Silicon 31 (Si 31)	100
Silver 105 (Ag 105)	10
Silver 110m (Ag 110m)	1

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Silver 111 (Ag 111)	100
Sodium 22 (Na 22)	10
Sodium 24 (Na 24)	10
Strontium 85 (Sr 85)	10
Strontium 89 (Sr 89)	1
Strontium 90 (Sr 90)	0.1
Strontium 91 (Sr 91)	10
Strontium 92 (Sr 92)	10
Sulfur 35 (S 35)	100
Tantalum 182 (Ta 182)	10
Technetium 96 (Tc 96)	10
Technetium 97m (Tc 97m)	100
Technetium 97 (Tc 97)	100
Technetium 99m (Tc 99m)	100
Technetium 99 (Tc 99)	10
Tellurium 125m (Te 125m)	10
Tellurium 127m (Te 127m)	10
Tellurium 127 (Tel 127)	100
Tellurium 129m (Te 129m)	10
Tellurium 129 (Te 129)	100
Tellurium 131m (Te 131m)	10
Tellurium 132 (Te 132)	10
Terbium 160 (Tb 160)	10
Thallium 200 (Tl 200)	100
Thallium 201 (Tl 201)	100
Thallium 202 (Tl 202)	100
Thallium 204 (Tl 204)	10
Thulium 170 (Tm 170)	10
Thulium 171 (Tm 171)	10
Tin 113 (Sn 113)	10
Tin 125 (Sn 125)	10
Tungsten 181 (W 181)	10
Tungsten 185 (W 185)	10
Tungsten 187 (W 187)	100
Vanadium 48 (V 48)	10
Xenon 131m (Xe 131m)	1,000
Xenon 133 (Xe 133)	100
Xenon 135 (Xe 135)	100
Ytterbium 175 (Yb 175)	100
Yttrium 90 (Y 90)	10
Yttrium 91 (Y 91)	10
Yttrium 92 (Y 92)	100
Yttrium 93 (Y 93)	100
Zinc 65 (Zn 65)	10
Zinc 69m (Zn 69m)	100
Zinc 69 (Zn 69)	1,000
Zirconium 93 (Zr 93)	10
Zirconium 95 (Zr 95)	10
Zirconium 97 (Zr 97)	10
Any radioactive material not listed above other than alpha emitting radioactive material	0.1
Any alpha radionuclide not listed above or mixtures of alpha emitters of unknown composition	0.1

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TABULAR OR GRAPHIC MATERIAL SET FORTH AT THIS POINT IS NOT DISPLAYABLE

1 To convert microcuries (<<mu>>Ci) to kilobecquerels (kBq), multiply microcuries
by 37.

HISTORICAL NOTE

Appendix B in original publication July 1, 1991.

Americium 241 added City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Arsenic 73 (As 74) repealed City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Arsenic 74 (As 74) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Beryllium 7 (Be 7) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Carbon 14 (C14) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Indium 114 (In 114) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Plutonium 239 added City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

Radium 226 (Ra 226) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Sodium 22 (Na 22) added City Record Apr. 3, 2001 eff. May 3, 2001. [See Vol. 9 Statements of Basis and Purpose No. 16]

Any alpha radionuclide . . . added (after Zirconium 97) City Record Mar. 16, 2005 eff. Apr. 15, 2005. [See Vol. 9 Statements of Basis and Purpose No. 23]

NOTES:

Appendix B was amended on Mar. 10, 2005 to be consistent with 10 CFR Part 30 Appendix B.

Appendix B was amended on March 21, 2001 to be consistent with the New York State Sanitary Code, Appendix 16-A, Table 3.

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APPENDIX C
QUANTITIES OF RADIOACTIVE MATERIAL REQUIRING
CONSIDERATION OF THE NEED FOR AN EMERGENCY PLAN
FOR RESPONDING TO A RELEASE

Radioactive material 1	Release fraction	Quantity (curies) 2

APPENDIX C (cont'd)
QUANTITIES OF RADIOACTIVE MATERIAL REQUIRING
CONSIDERATION OF THE NEED FOR AN EMERGENCY PLAN
FOR RESPONDING TO A RELEASE

Radioactive material 1	Release fraction	Quantity (curies) 2
Actinium-228	0.001	4,000
Americium-241	.001	2
Americium-242	.001	2
Americium-243	.001	2
Antimony-124	.01	4,000
Antimony-126	.01	6,000
Barium-133	.01	10,000
Barium-140	.01	30,000
Bismuth-207	.01	5,000
Bismuth-210	.01	600
Cadmium-109	.01	1,000
Cadmium-113	.01	80
Calcium-45	.01	20,000
Californium-252	.001	9 (20 mg)
Carbon-14	.01	50,000
	Non CO	
Cerium-141	.01	10,000
Cerium-144	.01	300
Cesium-134	.01	2,000
Cesium-137	.01	3,000
Chlorine-36	.5	100
Chromium-51	.01	300,000
Cobalt-60	.001	5,000
Copper-64	.01	200,000
Curium-242	.001	60
Curium-243	.001	3
Curium-244	.001	4
Curium-245	.001	2
Europium-152	.01	500
Europium-154	.01	400
Europium-155	.01	3,000
Germanium-68	.01	2,000
Gadolinium-153	.01	5,000
Gold-198	.01	30,000
Hafnium-172	.01	400
Hafnium-181	.01	7,000
Holmium-166m	.01	10C
Hydrogen-3	.5	20,000
Iodine-125	.5	10
Iodine-131	.5	10
Indium-114m	.01	1,000

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Iridium-192	.001	40,000
Iron-55	.01	40,000
Iron-59	.01	7,000
Krypton-85	1.0	6,000,000
Lead-210	.01	8
Manganese-58	.01	60,000
Mercury-203	.01	10,000
Molybdenum-99	.01	30,000
Neptunium-237	.001	2
Nickel-63	.01	20,000
Niobium-94	.01	300
Phosphorus-32	.5	100
Phosphorus-33	.5	1,000
Polonium-210	.01	10
Potassium-42	.01	9,000
Promethium-145	.01	4,000
Promethium-147	.01	4,000
Ruthenium-106	.01	200
Samarium-151	.01	4,000
Scandium-46	.01	3,000
Selenium-75	.01	10,000
Silver-110m	.01	1,000
Sodium-22	.01	9,000
Sodium-24	.01	10,000
Strontium-89	.01	3,000
Strontium-90	.01	90
Sulfur-35	.5	900
Technetium-99	.01	10,000
Technetium-99m	.01	400,000
Tellurium-127m	.01	5,000
Tellurium-129m	.01	5,000
Terbium-160	.01	4,000
Thulium-170	.01	4,000
Tin-113	.01	10,000
Tin-123	.01	3,000
Tin-126	.01	1,000
Titanium-44	.01	100
Vanadium-48	.01	7,000
Xenon-133	1.0	900,000
Yttrium-91	.01	2,000
Zinc-65	.01	5,000
Zirconium-93	.01	400
Zirconium-95	.01	5,000
Any other beta-gamma emitter	.01	10,000
Mixed fission products	.01	1,000
Mixed corrosion products	.01	10,000
Contaminated equipment beta-gamma	.001	10,000
Irradiated material, any form other than solid noncombustible	.01	1,000
Irradiated material, solid noncombustible	.001	10,000
Mixed radioactive waste, beta-gamma	.01	1,000
Packaged mixed waste, beta-gamma 3	.001	10,000
Any other alpha emitter	.001	2

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Contaminated equipment, alpha	.0001	20
Packaged waste, alpha 3	.0001	20
Combination of radioactive materials listed above 1		

APPENDIX C

QUANTITIES OF RADIOACTIVE MATERIAL REQUIRING

CONSIDERATION OF THE NEED FOR AN EMERGENCY PLAN

FOR RESPONDING TO A RELEASE

Radioactive material 1 Release Quantity

fraction (curies) 2

APPENDIX C (cont'd)

QUANTITIES OF RADIOACTIVE MATERIAL REQUIRING

CONSIDERATION OF THE NEED FOR AN EMERGENCY PLAN

FOR RESPONDING TO A RELEASE

Radioactive material 1 Release Quantity

fraction (curies) 2

Actinium-228	0.001	4,000
Americium-241	.001	2
Americium-242	.001	2
Americium-243	.001	2
Antimony-124	.01	4,000
Antimony-126	.01	6,000
Barium-133	.01	10,000
Barium-140	.01	30,000
Bismuth-207	.01	5,000
Bismuth-210	.01	600

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Cadmium-109	.01	1,000
Cadmium-113	.01	80
Calcium-45	.01	20,000
Californium-252	.001	9 (20 mg)
Carbon-14	.01	50,000
Non CO		
Cerium-141	.01	10,000
Cerium-144	.01	300
Cesium-134	.01	2,000
Cesium-137	.01	3,000
Chlorine-36	.5	100
Chromium-51	.01	300,000
Cobalt-60	.001	5,000
Copper-64	.01	200,000
Curium-242	.001	60
Curium-243	.001	3
Curium-244	.001	4
Curium-245	.001	2
Europium-152	.01	500
Europium-154	.01	400
Europium-155	.01	3,000
Germanium-68	.01	2,000
Gadolinium-153	.01	5,000

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Gold-198	.01	30,000
Hafnium-172	.01	400
Hafnium-181	.01	7,000
Holmium-166m	.01	10C
Hydrogen-3	.5	20,000
Iodine-125	.5	10
Iodine-131	.5	10
Indium-114m	.01	1,000
Iridium-192	.001	40,000
Iron-55	.01	40,000
Iron-59	.01	7,000
Krypton-85	1.0	6,000,000
Lead-210	.01	8
Manganese-58	.01	60,000
Mercury-203	.01	10,000
Molybdenum-99	.01	30,000
Neptunium-237	.001	2
Nickel-63	.01	20,000
Niobium-94	.01	300
Phosphorus-32	.5	100
Phosphorus-33	.5	1,000
Polonium-210	.01	10
Potassium-42	.01	9,000
Promethium-145	.01	4,000

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Promethium-147	.01	4,000
Ruthenium-106	.01	200
Samarium-151	.01	4,000
Scandium-46	.01	3,000
Selenium-75	.01	10,000
Silver-110m	.01	1,000
Sodium-22	.01	9,000
Sodium-24	.01	10,000
Strontium-89	.01	3,000
Strontium-90	.01	90
Sulfur-35	.5	900
Technetium-99	.01	10,000
Technetium-99m	.01	400,000
Tellurium-127m	.01	5,000
Tellurium-129m	.01	5,000
Terbium-160	.01	4,000
Thulium-170	.01	4,000
Tin-113	.01	10,000
Tin-123	.01	3,000
Tin-126	.01	1,000
Titanium-44	.01	100
Vanadium-48	.01	7,000
Xenon-133	1.0	900,000

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Yttrium-91	.01	2,000
Zinc-65	.01	5,000
Zirconium-93	.01	400
Zirconium-95	.01	5,000
Any other beta-gamma emitter	.01	10,000
Mixed fission products	.01	1,000
Mixed corrosion products	.01	10,000
Contaminated equipment beta-gamma	.001	10,000
Irradiated material, any form other than solid noncombustible	.01	1,000
Irradiated material, solid noncombustible	.001	10,000
Mixed radioactive waste, beta-gamma	.01	1,000
Packaged mixed waste, beta-gamma 3	.001	10,000
Any other alpha emitter	.001	2
Contaminated equipment, alpha	.0001	20
Packaged waste, alpha 3	.0001	20
Combination of radioactive materials listed above 1		

FOOTNOTES:

1 For combinations of radioactive materials, consideration of the need for an emergency plan is required if the sum of the ratios of the quantity of each radioactive material authorized to the quantity listed for that material in Appendix C exceeds one.

2 To convert curies (Ci) to gigabecquerels (GBq), multiply curies by 37.

3 Waste packaged in Type B containers does not require an emergency plan.

HISTORICAL NOTE

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Appendix C in original publication July 1, 1991.

APPENDIX D CRITERIA RELATING TO USE OF FINANCIAL TESTS AND PARENT COMPANY GUARANTEES FOR PROVIDING REASONABLE ASSURANCE OF FUNDS FOR DECOMMISSIONING I. Introduction. An applicant or licensee may provide reasonable assurance of the availability of funds for decommissioning based on obtaining a parent company guarantee that funds will be available for decommissioning costs and on a demonstration that the parent company passes a financial test. This appendix established criteria for passing the financial test and for obtaining the parent company guarantee.II. Financial Test. A. To pass the financial test, the parent company must meet the criteria of either paragraph A.1 or A.2 of this section:1. The parent company must have:i. Two of the following three ratios: A ratio of total liabilities to net worth less than 2.0; a ratio of the sum of net income plus depreciation, depletion, and amortization to total liabilities greater than 0.1; and a ratio of current assets to current liabilities greater than 1.5; andii. Net working capital and tangible net worth each at least six times the current decommissioning cost estimates (or prescribed amount if a certification is used); andiii. tangible net worth of at least $10 million; andiv. assets located in the United States amounting to at least 90 percent of total assets or at least six times the current decommissioning cost estimates (or prescribed amount if a certification is used).2. The parent company must have:i. A current rating for its most recent bond issuance of AAA, AA, A or BBB as issued by Standard and Poor's or Aaa, Aa, A or Baa as issued by Moody's; andii. tangible net worth of at least $10 million; andiii. tangible net worth at least six times the current decommissioning cost estimate (or prescribed amount if a certification is used); andiv. assets located in the United States amounting to at least 90 percent of total assets or at least six times the current decommissioning cost estimates (or prescribed amount if a certification is used).B. The parent company's independent certified public accountant must have compacted the data used by the parent company in the financial test, which is derived from the independently audited, year end financial statements for the latest fiscal year, with the amounts in such financial statement. In connection with that procedure, the licensee shall inform the Department within 90 days of any matters coming to the auditor's attention which cause the auditor to believe that the data specified in the financial test should be adjusted and that the company no longer passes the test.C. 1. After the initial financial test, the parent company must repeat the passage of the test within 90 days after the close of each succeeding fiscal year.2. If the parent company no longer meets the requirements of paragraph A of this section, the license, the licensee must send notice to the Department of intent to establish alternate financial assurance as specified in this Code. The notice must be sent by certified mail within 90 days after the end of the fiscal year for which the year end financial data show that the parent company no longer meets the financial test requirements. The licensee must provide alternate financial assurance within 120 days after the end of such fiscal year.III. Parent Company Guarantee. The terms of a parent company guarantee which an applicant or licensee obtains must provide that:A. The parent company guarantee will remain in force unless the guarantor sends notice of cancellation by certified mail to the licensee and the Department. Cancellation may not occur, however, during the 120 days beginning on the date of receipt of the notice of cancellation by both the licensee and the Department, as evidenced by the return receipts.B. If the licensee fails to provide alternate financial assurance as specified by this Code within 90 days after receipt by the licensee and Department of a notice of cancellation of the parent company guarantee

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from the guarantor, the guarantor will provide such alternative financial assurance in the name of the licensee.C. The parent company guarantee and financial test provisions must remain in effect until the Department has terminated the license.D. If a trust is established for decommissioning costs, the trustee and trust must be acceptable to the Department. An acceptable trustee includes an appropriate state or federal government agency or an entity which has the authority to act as a trustee and whose trust operations are regulated and examined by a federal or state agency.

HISTORICAL NOTE

Appendix D in original publication July 1, 1991.

APPENDIX E CRITERIA RELATING TO USE OF FINANCIAL TESTS AND SELF GUARANTEES FOR PROVIDING REASONABLE ASSURANCE OF FUNDS FOR DECOMMISSIONING I. Introduction. An applicant or licensee may provide reasonable assurance of the availability of funds for decommissioning based on furnishing its own guarantee that funds will be available for decommissioning costs and on a demonstration that the company passes the financial test of section II of this appendix. The terms of the self-guarantee are in section III of this appendix. This appendix establishes criteria for passing the financial test for the self-guarantee and establishes the terms for a self-guarantee.II. Financial Test. A. To pass the financial test, a company must meet all of the following criteria:1. tangible net worth at least 10 times the total current decommissioning cost estimate (or the current amount required if certification is used) for all decommissioning activities for which the company is responsible as self-guaranteeing licensee and as parent-guarantor.2. assets located in the United States amounting to at least 90 percent of total assets or at least 10 times the total current decommissioning cost estimate (or the current amount required if certification is used) for all decommissioning activities for which the company is responsible as self-guaranteeing licensee and as parent-guarantor.3. A current rating for its most recent bond issuance of AAA, AA or A as issued by Standard and Poor's or Aaa, Aa or A as issued by Moody's.B. To pass the financial test, a company must meet all of the following additional requirements:1. the company must have at least one class of equity securities registered under the Securities Exchange Act of 1934.2. the company's independent certified public accountant must have compared the data used by the company in the financial test which is derived from the independently audited, year end financial statements for the latest fiscal year, with the amounts in such financial statement. In connection with that procedure, the licensee shall inform the Department within 90 days of any matters coming to the attention of the auditor that cause the auditor to believe that the data specified in the financial test should be adjusted and that the company no longer passes the test.3. After the initial financial test, the company must repeat passage of the test within 90 days after the close of each succeeding fiscal year.C. If the licensee no longer meets the requirements of Section II.A. of this appendix, the licensee must send immediate notice to the Department of its intent to establish alternate financial assurance as specified in this Code within 120 days of such notice.III. Company Self-Guarantee. The terms of a self-guarantee which an applicant or licensee furnishes must provide that:A. The guarantee will remain in force unless the licensee sends notice of cancellation by certified mail to the Department. Cancellation may not occur, however, during the 120 days beginning on the date of receipt of the notice of cancellation by the Department, as evidenced by the return receipt.B. The licensee shall provide alternative financial assurance as specified in this Code within 90 days following receipt by the Department of a notice of can-

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cellation of the guarantee.C. The guarantee and financial test provisions must remain in effect until the Department has terminated the license or until another financial assurance method acceptable to the Department has been put into effect by the licensee.D. The licensee will promptly forward to the Department and the licensee's independent auditor all reports covering the latest fiscal year filed by the licensee with the Securities and Exchange Commission pursuant to the requirements of Section 13 of the Securities and Exchange Act of 1934.E. If, at any time, the licensee's most recent bond issuance ceases to be rated in any category of "A" or above by either Standard and Poor's or Moody's, the licensee will provide notice in writing of such fact to the Department within 20 days after publication of the change by the rating service. If the licensee's most recent bond issuance ceases to be rated in any category of "A" or above by both Standard and Poor's and Moody's, the licensee no longer meets the requirements of Section II.A. of this appendix.F. The applicant or licensee must provide to the Department a written guarantee (a written commitment by a corporate officer) which states that the licensee will fund and carry out the required decommissioning activities or, upon issuance of an order by the Department, the licensee will set up and fund a trust in the amount of the current cost estimates for decommissioning.

HISTORICAL NOTE

Appendix E in original publication July 1, 1991.

RCNY Art. 175, Refs & Annos
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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.102. Requirements for specific types of radioactive materials licenses.

(a) Types of license. (1) For the purposes of § 5.07(a) and § 175.05 of this Code, the following special designations of radioactive materials licenses shall apply:

(i) A specific license of limited scope for teletherapy means a license that authorizes receipt, acquisition, ownership, possession, use and transfer of specified quantities and types of radioactive material for use in teletherapy programs.

(ii) A specific license of limited scope for medical use means a license that authorizes receipt, acquisition, ownership, possession, use and transfer of specified quantities and types of radioactive material for use in or on humans in a medical program, but does not include teletherapy.

(iii) A specific license of limited scope for other use means a license that authorizes receipt, production, acquisition, ownership, possession, use and transfer of specified quantities and types of radioactive material for uses other than in or on humans.

(iv) A specific license of broad scope for medical use means a license that authorizes receipt, acquisition, ownership, possession, use and transfer of any chemical or physical form of the radioactive materials specified in the license, for use in or on humans in a medical program, but does not include teletherapy.

(v) A specific license of broad scope for research and development means a license that authorizes receipt, acquisition, ownership, possession, use and transfer of any chemical or physical form of radioactive materials specified in the license, in quantities not exceeding those specified in the license, for uses other than in or on humans.

(b) The requirements specified in this section are in addition to, and not in substitution for, others in this Code. In particular, the provisions of § 175.101 apply to all license applications and all specific radioactive materials licenses.

(c) Specific licenses for human use of radioactive materials in institutions.

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(1) An application by an institution for a specific license for medical use of radioactive material will be approved if:

(i) the applicant satisfies the requirements specified in § 175.101 and § 175.103 of this Code; and

(ii) the applicant possesses adequate facilities for the clinical care of patients; and

(iii) the physician(s) designated on the application as the individual authorized user(s) has training and experience as specified in § 175.103(j) in the proposed use, the handling and administration of radionuclides and the clinical management of radioactive patients. The physician shall furnish evidence of such experience with his/her application. A statement from his/her preceptor at the institution where he/she acquired such training and experience, indicating their amount and nature, may be submitted as evidence of such experience.

(iv) If the application is for a license to use unspecified quantities or multiple types of radioactive materials with atomic numbers 3 through 83 the applicant shall have had previous experience operating under a specific institutional license and have been engaged in the use of radioisotopes in medical research, as well as routine diagnosis and therapy.

(v) The license application is signed by the chairman of the radiation safety committee and an authorized representative of the institution.

(d) Specific licenses to individual physicians for human use of radioactive materials. (1) An application by an individual physician for a specific license for human use of radioactive material will be approved if:

(i) the applicant satisfies the requirements specified in § 175.101 and § 175.103 of this Code; and

(ii) the applicant has training and experience as specified in § 175.103(j) in the proposed use, the handling and administration of radionuclides, and the clinical management of radioactive patients. The physician shall furnish evidence of such training and experience with his/her application. A statement from his/her preceptor at the institution where he/she acquired such training and experience, indicating their amount and nature, may be submitted as evidence of such experience.

(e) Specific licenses of broad scope. (1) A specific license of broad scope shall be issued only to medical institutions or institutions of higher education; such licenses shall not be issued to individuals.

(2) An application for a specific license of broad scope will be approved if:

(i) the applicant satisfies the general requirements specified in § 175.101(e) of this Code; and

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(ii) the applicant has engaged in a reasonable number of activities involving the use of radioactive material; and

(iii) the applicant has established administrative controls and provisions relating to organization and management, procedures, recordkeeping, material control, and accounting and management review that are necessary to assure safe operations, including:

(A) the establishment of a radiation safety committee pursuant to § 175.03 of this Code; and

(B) the appointment of a full-time radiation safety officer pursuant to § 175.03 of this Code; and

(C) the establishment of appropriate administrative procedures to assure:

(a) control of procurement and use of radioactive material; and

(b) completion of safety evaluations of proposed uses of radioactive material which take into consideration such matters as the adequacy of facilities and equipment, training and experience of the user, and the operating or handling procedures; and

(c) review, approval, and recording by the radiation safety committee of safety evaluations of proposed uses prepared in accordance with § 175.102(e)(2)(iii)(C)(b) prior to the use of radioactive materials.

(3) The following are conditions of all specific licenses of broad scope:

(i) Unless specifically authorized pursuant to other provisions of this Code, broad scope licensees shall not:

(A) conduct tracer studies in the environment involving direct release of radioactive material;

(B) receive, acquire, own, possess, use, transfer, or import devices containing 3.7 E+6 GBq (100,000 Ci) or more of radioactive material in sealed sources used for irradiation of materials;

(C) add, or cause the addition of, radioactive material to any food, beverage, cosmetic, drug or other product designed for ingestion by, or application to, a human being except as authorized in the license.

(f) Specific licenses for non-human use. (1) An application for a specific license authorizing non-human use of radioactive materials will be approved if:

(i) the applicant satisfies the general requirements specified in § 175.101(e) of this Code; and

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(ii) the applicant, or the applicant's personnel, has training and experience commensurate with the proposed amounts, types and uses of radioactive materials which shall include at a minimum:

(A) a college degree at the bachelor level in a physical, biological, environmental or engineering science; and

(B) at least forty (40) hours of training and experience in the safe handling of radioactive materials appropriate to the type and forms of such materials to be used, which shall include:

(a) characteristics of ionizing radiation;

(b) units of radiation dose and quantities;

(c) radiation detection instrumentation; and

(d) biological hazards of exposure to radiation.

(g) General licenses. (1) A general license is hereby issued to own radioactive material without regard to quantity. Notwithstanding any other provisions of this Code, this general license does not authorize the manufacture, production, transfer, receipt, possession or use of radioactive material.

(2) Source material. (i) A general license is hereby issued authorizing use and transfer of not more than 6.8 kilograms (15 pounds) of source material at any one time by persons in the following categories:

(A) pharmacists using the source material solely for the compounding of medicinals;

(B) physicians using the source material for medicinal purposes;

(C) persons receiving possession of source material from pharmacists and physicians in the form of medicinals or drugs;

(D) commercial and industrial firms, and research, educational, and medical institutions for research, development, educational or commercial purposes;

and provided, that no such person shall, pursuant to this general license, receive more than 68 kilograms (150 pounds) of source material in any one (1) calendar year.

(ii) Persons who transfer, receive, possess or use source material pursuant to the general license issued in § 175.102(g)(2)(i) are exempt from the provisions of § 175.03, § 175.04, § 175.06, § 175.104 and § 175.105 of this Code to the extent that such transfer, receipt, possession or use is within the terms of such general license, provided however, that this exemption shall not be deemed to apply to any such person who is also in possession of source material under a specific license issued pursuant to this Code.

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(iii) A general license is hereby issued authorizing the receipt of title to source material without regard to quantity. This general license does not authorize any person to transfer, receive, possess or use source material.

(3) Certain devices and equipment. (i) A general license is hereby issued to transfer, receive, possess or use radioactive material incorporated in the following devices or equipment which have been manufactured, tested and labeled in accordance with a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, authorizing distribution under this general license or its equivalent.

(A) Static elimination devices. Devices designed for use as static eliminators which contain, as a sealed source or sources, radioactive material consisting of a total of not more than 18.5 MBq (500 mCi) of polonium-210 per device.

(B) Ion generating tubes. Devices designed for ionization of air which contain, as a sealed source or sources, radioactive material consisting of a total of not more than 18.5 MBq (500 <<mu>>CI) of polonium-210 per device or a total of not more than 1.85 GBq (50 mCi) per device.

(4) Certain measuring, gauging or controlling devices. (i) A general license is hereby issued to receive, possess or use radioactive material when contained in devices used at a fixed location and designed and manufactured for the purpose of detecting, measuring, gauging or controlling thickness, density, level, interface location, radiation leakage, or qualitative or quantitative chemical composition, or for producing light or an ionized atmosphere, when such devices are manufactured or imported in accordance with the specifications contained in a specific license issued to the supplier by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, and authorizing distribution under this general license or its equivalent, provided that:

(A) such devices are labeled in accordance with the provisions of the specific license which authorizes the distribution of the devices; 1 [FN5]

[FN5]. 1 Regulations under the Federal Food, Drug, and Cosmetic Act authorizing the use of radioactive control devices in food production require certain additional labeling thereon which is found in § 121.3001 of 21 CFR Part 121.

(B) such devices bear a label containing the following or a substantially similar statement which contain the information called for in the following statement:

The transfer, receipt, possession or use of this device, Model _____ , Serial [FN6] number _____ , are subject to a general license or the equivalent and the regulations of the U.S. Nuclear Regulatory Commission or of a State with which the Nuclear Regulatory Commission has entered into an agreement for the exercise of regulatory authority. Removal of this label is prohibited.

[FN6]. 2 The model, serial number, and name of supplier may be omitted from this label provided they are elsewhere specified in labeling affixed to the device.

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_____
(Name of supplier 2)

(C) such devices are installed on the premises of the general licensee by a person authorized to install such devices under a specific license issued to the installer by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, if a label affixed to the device at the time of receipt states that installation by a specific licensee is required. This requirement does not apply while devices are held in storage in the original shipping container pending installation by a specific licensee.

(ii) Persons who receive, possess or use a device pursuant to the general license of § 175.102(g)(3)(i):

(A) shall, within ten (10) days after the receipt of the device, notify the Department of the type of device and the name and address of the supplier;

(B) shall not transfer, abandon, or dispose of the device except by transfer to a person duly authorized to receive such device by a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, and shall furnish to the Department, within thirty (30) days after any such transfer, a report containing the name of the manufacturer of the device, the type of device, the manufacturer's serial number of the device, and the name and address of the person receiving the device;

(C) shall assure that all labels affixed to the device at the time of receipt and bearing the statement, "Removal of this label is prohibited" are maintained thereon and shall comply with all instructions contained in such labels;

(D) shall have the device tested for leakage of radioactive material and proper operation of the on-off mechanism and indicator, if any, at the time of installation of the device or replacement of the radioactive material on the premises of the general licensee and thereafter at no longer than six (6) month intervals or at such longer intervals not to exceed three (3) years as are specified in the label required by § 175.102(g)(3)(i)(A), provided, that devices containing only krypton-85 need not be tested for leakage, and devices containing only hydrogen-3 need not be tested for any purpose;

(E) shall have all the tests required by § 175.102(g)(3)(ii)(D) and all other services involving the radioactive material, its shielding and containment, performed by the supplier or other person duly authorized by a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, to manufacture, install or service such devices;

(F) shall, within thirty (30) days after the occurrence of a failure or of damage to the shielding of the radioactive material or the on-off mechanism or indicator or upon the detection of 0.185 kBq (0.005 <<mu>>Ci) or more of removable radioactive material, furnish to the Department a report containing the name of the manufacturer of the device and a brief description of the event and the remedial action

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taken; and shall maintain records of all tests performed on the devices as required herein, including the dates and results of the tests and the names of the persons conducting the tests;

(G) shall, upon the occurrence of a failure of or damage to, or any indication of a possible failure of or damage to, the shielding or containment of the radioactive material or the on-off mechanism or indicator, immediately suspend operation of the device until it has been repaired by a person holding a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state, to manufacture, install or service such devices, or disposed of by transfer to a person holding a specific license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state to receive the radioactive material contained in the device;

(H) shall be exempt from the provisions of § 175.03, § 175.04, § 175.06, § 175.104 and § 175.105 of this Code, except that such persons shall comply with the provisions of § 175.03(l)(1).

(iii) Any holder of a license issued by the Department, the U.S. Nuclear Regulatory Commission or an agreement state which authorizes the holder to manufacture, install or service a device subject to the general license in § 175.102(g)(3)(i), and which is not limited as to specific installation or installations, may install or service such devices without obtaining a license from the Department provided that:

(A) such person shall file a report with the Department within thirty (30) days after the end of each calendar quarter in which any device is transferred to or installed within the Department's jurisdiction. Each such report shall identify the name and address of each person receiving a device, the type of device transferred, and the quantity and type of radioactive material contained in the device;

(B) the device is manufactured, labeled, installed and serviced in accordance with terms and conditions of the license issued to such person;

(C) such person shall assure that any labels required to be affixed to the device bear a statement that "Removal of this label is prohibited"; and

(D) the person to whom such device is transferred, or on whose premises such device is installed or serviced, has a copy of the general license requirements or equivalent requirements specified in § 175.102(g)(3)(ii).

(iv) The Department may withdraw, limit or qualify its acceptance of any specific license issued by another agency, or any product distributed pursuant to such licensing document, upon determining that such action is necessary to prevent undue hazard to public health and safety or property.

(4) Luminous safety devices for aircraft. (i) A general license is hereby issued to receive, possess or use hydrogen-3 or promethium-147 contained in luminous safety devices for use in aircraft, provided:

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(A) each device contains not more than 370 GBq (10 Ci) of hydrogen-3 or 11.1 GBq (300 mCi) of promethium-147; and

(B) each device has been manufactured, assembled or imported in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission, or each device has been manufactured, assembled or imported in accordance with the specifications contained in a specific license issued by the Department or an agreement state pursuant to licensing requirements equivalent to those in § 32.53 of 10 CFR Part 32.

(ii) Persons who receive, possess or use luminous safety devices pursuant to the general license in § 175.102(g)(4)(i) are exempt from the provisions of § 175.03, § 175.04, § 175.06, § 175.104 and § 175.105 of this Code, except that such persons shall comply with the provisions of § 175.03(l)(1).

(iii) This general license does not authorize the manufacture, assembly or repair of luminous safety devices containing hydrogen-3 or promethium-147.

(iv) This general license does not authorize the receipt, possession or use of promethium-147 contained in instrument dials.

(5) Calibration and reference sources. (i) A general license is hereby issued to those persons listed below to transfer, receive, possess or use, in accordance with the provisions of § 175.102(g)(5)(iii) and (iv), americium-241 in the form of calibration or reference sources:

(A) any person who holds a specific license issued by the Department, the New York State Department of Health or the New York State Department of Labor which authorizes the transfer, receipt, possession or use of radioactive material; and

(B) any person who holds a specific license issued by the U.S. Nuclear Regulatory Commission which authorizes the transfer, receipt, possession or use of special nuclear material.

(ii) A general license is hereby issued to transfer, receive, possess or use plutonium in the form of calibration or reference sources in accordance with the provisions of § 175.102(g)(5)(iv) and (v) to any person who holds a specific license issued by the Department, the New York State Department of Health or the New York State Department of Labor which authorizes the transfer, receipt, possession or use of radioactive material.

(iii) The general licenses in § 175.102(g)(5)(i) and (ii) apply only to calibration or reference sources which have been manufactured in accordance with the specifications contained in a specific license issued to the manufacturer or importer of the sources by the U.S. Nuclear Regulatory Commission pursuant to § 32.57 of 10 CFR Part 32 or § 70.39 of 10 CFR Part 70 or which have been manufactured or imported in accordance with the specifications contained in a specific license issued by the Department or an agreement state pursuant to licensing requirements equivalent to those contained in § 32.57 of 10 CFR Part 32 or § 70.39 of 10 CFR Part 70.

(iv) Persons who transfer, receive, possess or use one or more calibration or

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reference sources pursuant to these general licenses:

(A) shall not possess at any one time, at any one location of storage or use, more than 185 kBq (5 <<mu>>Ci) of americium-241 and 185 kBq (5 <<mu>>Ci) of plutonium in such sources;

(B) shall not transfer, receive, possess or use such source unless the source, or the storage container, bears a label which includes the following statement or a substantially similar statement which contains the information called for in the following statement:

The transfer, receipt, possession or use of this device, Model_____ , Serial number _____ , are subject to a general license and the regulations of the U.S. Nuclear Regulatory Commission or of a State with which the Nuclear Regulatory Commission has entered into an agreement for the exercise of regulatory authority. Do not remove this label. CAUTION--RADIOACTIVE MATERIAL-- THIS SOURCE CONTAINS (AMERICIUM-241) (PLUTONIUM) 3 [FN7]. DO NOT TOUCH RADIOACTIVE PORTION OF THIS SOURCE.

[FN7]. 3 Showing only the name of the appropriate material.

_____

(Name of manufacturer or importer);

(C) shall not transfer, abandon or dispose of such source except by transfer to a person authorized by a license from the Department, the U.S. Nuclear Regulatory Commission or an agreement state to receive the source;

(D) shall store such source, except when the source is being used, in a closed container adequately designed and constructed to contain americium-241 or plutonium which might otherwise escape during storage; and

(E) shall not use such source for any purpose other than the calibration of radiation detectors or the standardization of other sources.

(v) These general licenses do not authorize the manufacture of calibration or reference sources containing americium-241 or plutonium.

(vi) A general license is hereby issued to transfer, receive, possess or use sealed radioactive materials sources in individual quantities each of which does not exceed the applicable quantity set forth in Appendix B of § 175.101 of this Code, to any person who holds a specific license issued by the Department.

(vii) Persons who transfer, receive, possess or use sources pursuant to the general license in § 175.102(g)(5)(vi):

(A) shall not transfer, abandon or dispose of such sources except by transfer to a person duly authorized to receive such sources by the Department, the U.S. Nuclear Regulatory Commission or an agreement state; and

(B) shall store such sources, except when being used, in a secure location.

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(6) Ice detection devices. (i) A general license is hereby issued to transfer, receive, possess or use strontium-90 contained in ice detection devices, provided each device contains not more than 1.85 MBq (50 <<mu>>Ci) of strontium-90 and each device has been manufactured or imported in accordance with a specific license issued by the U.S. Nuclear Regulatory Commission pursuant to § 32.61 of 10 CFR Part 32 or by the Department or an agreement state pursuant to licensing requirements equivalent to those contained in § 32.61 of 10 CFR Part 32.

(ii) Persons who transfer, receive, possess or use strontium-90 contained in ice detection devices pursuant to this general license:

(A) shall, upon occurrence of visually observable damage, such as a bend or crack or discoloration from overheating, to the device, discontinue use of the device until it has been inspected, tested for leakage and repaired by a person holding a specific license from the U.S. Nuclear Regulatory Commission or an agreement state to manufacture or service such devices; or shall dispose of the device pursuant to § 175.104 of this Code;

(B) shall assure that all labels affixed to the device at the time of receipt, and which bear a statement which prohibits removal of the labels, are maintained thereon; and

(C) are exempt from the provisions of § 175.03, § 175.04, § 175.06 and § 175.105 of this Code, except that such persons shall comply with the provisions of § 175.03(l)(1).

(iii) This general license does not authorize the manufacture, assembly, disassembly or repair of strontium-90 sources in ice detection devices.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

<General Materials (GM) - References, Annotations, or Tables>

24 RCNY Hlth. Code § 175.102
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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.103. Medical use of radioactive materials.

(a) General information. (1) Purpose and scope. This section establishes requirements and provisions for the use of radioactive material in the healing arts and for issuance of licenses authorizing the medical use of this material. These requirements and provisions provide for the protection of the public health and safety. The requirements and provisions of this section are in addition to, and not in substitution for, others in this Code. The requirements and provisions of this Code apply to applicants and licensees subject to this section unless specifically exempted.

(2) License required. (i) No person shall manufacture, produce, acquire, receive, possess, use, or transfer radioactive material for medical use except in accordance with a specific license issued pursuant to this section or as otherwise provided in this section.

(ii) Unless prohibited by license condition an individual may receive, possess, use, or transfer radioactive material in accordance with this Code under the supervision of an authorized user as provided in § 175.103(b)(5).

(3) Application for license, amendment or renewal. (i) If the application is for medical use sited in a medical institution, only the institution's management may apply. If the application is for medical use not sited in a medical institution, any professional practitioner may apply.

(ii) An application for a license for medical use of radioactive material, or renewal thereof, must be made by filing an original and one copy on Form BRC-M100, "Application for Radioactive Materials License". A request for license amendment may be submitted as an original and one copy in letter format.

(4) License amendments. A licensee shall apply for and shall have received a license amendment before:

(i) receiving or using radioactive material for a method or type of medical use not permitted by the license;

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(ii) permitting anyone, except a visiting authorized user as specified by license condition, to work as an authorized user under the license;

(iii) changing the radiation safety officer or radiation therapy physicist;

(iv) receiving radioactive material in excess of the amount, or radionuclide or form different from that authorized by the license;

(v) adding to or changing the area of use or mailing address identified on the license; and

(vi) changing statements, representations, and procedures which are incorporated by reference into the license.

(5) Notifications. A licensee shall notify the Department in writing within 30 days when an authorized user, radiation safety officer, or radiation therapy physicist, permanently discontinues performance of duties under the license. This requirement is not intended to relieve the licensee of the requirements of § 175.103(a)(4).

(b) General administrative requirements. (1) ALARA Program.

(i) Each licensee shall develop and implement a written program to maintain radiation doses and releases of radioactive material in effluents to unrestricted areas as low as reasonably achievable in accordance with § 175.03(b) of this Code.

(ii) To satisfy the requirement of § 175.103(b)(1)(i):

(A) for licensees that are medical institutions, the management, radiation safety officer and all authorized users shall participate in the establishment, implementation, and operation of the program as required by this Code or required by the radiation safety committee; or

(B) for licensees that are not medical institutions, management and all authorized users shall participate in the program as requested by the radiation safety officer.

(iii) The ALARA program shall include notice to workers of the program's existence and workers' responsibility to help keep dose equivalents ALARA.

(iv) The ALARA program shall include an annual review by the radiation safety committee for licensees that are medical institutions, or management, all authorized users and the radiation safety officer for licensees that are not medical institutions, of summaries of the types and amounts of radioactive material used, occupational dose reports, and continuing education and training for all personnel who work with or in the vicinity of radioactive material.

(v) The purpose of the review required by § 175.103(b)(1)(iv) is to ensure that individuals make every reasonable effort to maintain occupational doses, doses to

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the general public, and releases of radioactive material to unrestricted areas as low as reasonably achievable, taking into account the state of technology, and the cost of improvements in relation to benefits.

(vi) The licensee shall retain a current written description of the ALARA program for the duration of the license. The written description must include:

(A) a commitment by management to keep occupational doses as low as reasonably achievable;

(B) a requirement that the radiation safety officer brief management once each year on the radiation safety program;

(C) personnel exposure investigational levels that, when exceeded, will initiate an investigation by the radiation safety officer of the cause of the exposure; and

(D) personnel exposure investigational levels that, when exceeded, will initiate a prompt investigation by the radiation safety officer of the cause of the exposure and a consideration of actions that might be taken to reduce the probability of recurrence.

(2) Radiation safety officer. (i) A licensee shall appoint a radiation safety officer responsible for implementing the radiation safety program. The licensee, through the radiation safety officer, shall ensure that radiation safety activities are being performed in accordance with approved procedures and regulatory requirements in the daily operation of the licensee's radioactive material program.

(ii) The radiation safety officer shall:

(A) investigate overexposures, misadministrations, accidents, spills, losses, thefts, unauthorized receipts, uses, transfers, and disposals, and other deviations from approved radiation safety practice and implement corrective actions as necessary;

(B) establish, implement and maintain written policy and procedures for:

(a) authorizing the purchase of radioactive material;

(b) receiving and opening packages of radioactive material;

(c) storing radioactive material;

(d) keeping an inventory record of radioactive material;

(e) using radioactive material safely;

(f) taking emergency action if control of radioactive material is lost;

(g) performing periodic radiation surveys;

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(h) performing checks of survey instruments and other safety equipment;

(i) disposing of radioactive material;

(j) training personnel who work in or frequent areas where radioactive material is used or stored; and

(k) keeping copies of this Code, all records and reports required by this Code, each licensing request and license and amendments, and the written policies and procedures required by this Code;

(C) brief management at least once each year on the radioactive materials program; and

(D) for medical use sited at a medical institution, assist the radiation safety committee in the performance of its duties; or

(E) for medical use not sited at a medical institution, approve or disapprove radiation safety program changes with the advice and consent of management prior to submittal to the Department for licensing action.

(3) Radiation safety committee. Each medical institution licensee shall establish a radiation safety committee to oversee the use of radioactive material.

(i) The committee shall meet the following administrative requirements:

(A) Membership shall consist of at least three individuals and shall include an authorized user of each type of use permitted by the license, the radiation safety officer, a representative of the nursing service, and a representative of management who is neither an authorized user nor the radiation safety officer. Other members may be included as the licensee deems appropriate.

(B) The committee shall meet at least quarterly.

(C) To establish a quorum and to conduct business, at least one-half of the committee's membership shall be present, including the radiation safety officer and the management's representative.

(D) The minutes of each radiation safety committee meeting shall include:

(a) the date of the meeting;

(b) members present;

(c) members absent;

(d) summary of deliberations and discussions;

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(e) recommended actions and the numerical results of all ballots; and

(f) document any reviews required in § 175.103(b)(1)(iv) and (b)(3)(ii).

(E) The committee shall provide each member with a copy of the meeting minutes, and retain one copy for the duration of the license.

(ii) To oversee the use of licensed material, the committee shall:

(A) be responsible for monitoring the institutional program to maintain individual and collective doses as low as reasonably achievable;

(B) review, on the basis of safety and with regard to the training and experience standards of this Code, and approve or disapprove any individual who is to be listed as an authorized user, the radiation safety officer, or radiation therapy physicist before submitting a license application or request for amendment or renewal;

(C) review on the basis of safety and approve or disapprove each proposed method of use of radioactive material;

(D) review on the basis of safety, and approve or disapprove with the advice and consent of the radiation safety officer and the management representative, minor changes in radiation safety procedures that are not potentially important to safety and are permitted under § 175.103(b)(3)(iii);

(E) review on the basis of safety, and approve or disapprove with the advice and consent of the radiation safety officer and the management representative, procedures and radiation safety program changes prior to submittal to the Bureau of Radiological Health for licensing action;

(F) review quarterly, with the assistance of the radiation safety officer, occupational radiation exposure records of all personnel working with radioactive material;

(G) review quarterly, with the assistance of the radiation safety officer, all incidents involving radioactive material with respect to cause and subsequent actions taken;

(H) review annually, with the assistance of the radiation safety officer, the radioactive materials program; and

(I) establish a table of investigational levels for occupational dose that, when exceeded, will initiate investigations and considerations of action by the radiation safety officer.

(iii) A licensee may make minor changes in radiation safety procedures that are not potentially important to safety (e.g. editing of procedures for clarity, updating names or telephone numbers, replacement of equipment or assignment of service contracts), except for changes in § 175.103(a)(4) or § 175.103(i)(3). A licensee is

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responsible for assuring that any change made is in compliance with the requirements of this Code and the license.

(iv) A licensee shall retain a record of each change made pursuant to § 175.103 (b)(3)(iii) until the license has been renewed or terminated. The record must include the effective date of the change, a copy of the old and new radiation safety procedures, the reason for the change, a summary of radiation safety matters that were considered before making the change, the signature of the radiation safety officer, and the signatures of the affected authorized user and of management or, in a medical institution, the radiation safety committee's chairman and the management representative.

(4) Statement of authorities and responsibilities. (i) A licensee shall provide the radiation safety officer, and at a medical institution the radiation safety committee, sufficient authority and organizational freedom to:

(A) identify radiation safety problems;

(B) initiate, recommend, or provide corrective actions; and

(C) verify implementation of corrective actions.

(ii) A licensee shall establish in writing the authorities, duties, responsibilities, and radiation safety activities of the radiation safety officer, and at a medical institution the radiation safety committee, and retain the current edition of these statements for the duration of the license.

(5) Supervision. (i) A licensee who permits the receipt, possession, use, or transfer of radioactive material by an individual under the supervision of an authorized user as allowed by § 175.103(a)(2) shall:

(A) instruct the supervised individual in the principles and practices of radiation protection, radioactivity measurement standardization and monitoring techniques and instruments, mathematics and calculations basic to the use and measurement of radioactivity, and biological effects of radiation which are appropriate to that individual's use of radioactive material;

(B) if the supervised individual is a professional practitioner, provide further instruction by the supervising authorized user which shall include that training stipulated in § 175.103(j) which is appropriate for the uses in which the supervised individual is being trained. The licensee shall maintain records of such training for three years.

(C) require the supervising authorized user to periodically review the supervised individual's use of radioactive material, review records kept to reflect this use, and provide reinstruction as needed; and

(D) require that only those individuals specifically trained, and designated by the authorized user, shall be permitted to administer radionuclides or radiation to patients.

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(ii) A licensee shall require the supervised individual receiving, possessing, using or transferring radioactive material under § 175.103(a)(2) to:

(A) follow the instructions of the supervising authorized user;

(B) follow the procedures established by the radiation safety officer; and

(C) comply with this Code and the license conditions with respect to the use of radioactive material.

(iii) Personnel, other than physicians or registered professional nurses, at institutional licensees involved in the performance of diagnostic procedures utilizing radioactive material which includes performing parenteral administration of radioactive material by intravenous, intramuscular or subcutaneous methods shall:

(A) have satisfactorily completed an educational program in nuclear medicine technology accredited by the Committee on Allied Health Education and Accreditation or the accrediting agency of the state in which the program was completed, provided such state accreditation requires education and training in the above methods of parenteral administration; or

(B) possess certification as a nuclear medicine technologist by the American Registry of Radiologic Technologists or certification by the Nuclear Medicine Technology Board; and

(iv) prior to permitting parenteral administration by a nuclear medicine technologist, the medical board of a hospital, or the radiation safety committee of an institution who have no medical board, shall adopt with governing authority approval:

(A) procedures to assure that the nuclear medicine technologist possesses the education and training or certification set forth in § 175.103(b)(5)(iii) and is proficient in the competent performance of parenteral administration; and

(B) requirements for authorized user physician which at a minimum shall require supervision by such a physician on the premises when parenteral administration of radioactive material for diagnostic testing is performed by a qualified nuclear medicine technologist.

(v) A licensee that supervises an individual is responsible for the acts and omissions of the supervised individual.

(6) Suppliers. (i) A licensee may use for medical use only:

(A) radioactive material manufactured, labeled, packaged, and distributed in accordance with a license issued for such activities by an Agreement State or the U.S. Nuclear Regulatory Commission; and

(B) reagent kits that have been manufactured, labeled, packaged, and distributed in accordance with an approval issued by the U.S. Department of Health and Human

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Services, Food and Drug Administration ("FDA"); and

(C) teletherapy sources manufactured and distributed in accordance with a license issued pursuant to 10 CFR Part 30 or the equivalent regulations of an Agreement State.

(c) General technical requirements. (1) Possession, use, calibration, and check of dose calibrators.

(i) A medical use licensee authorized to administer radioactive materials shall possess a dose calibrator and use it to measure the amount of activity administered to each patient.

(ii) A licensee shall:

(A) check each dose calibrator for constancy with a dedicated check source at the beginning of each day of use. To satisfy this requirement, the check must be done on a frequently used setting with a sealed source of not less than 370 kBq (10 <<mu>>Ci) of radium-226 or 1.85 MBq (50 <<mu>>Ci) of any other photon-emitting radionuclide with a half-life greater than 90 days; and

(B) test each dose calibrator for accuracy upon installation and at intervals not to exceed 12 months thereafter by assaying at least 2 sealed sources containing different radionuclides, the activity of which the manufacturer has determined by traceability to a national standard to be within 5 percent of the stated activity, with minimum activity of 370 kBq (10 <<mu>>Ci) for radium-226 and 1.85 MBq (50 <<mu>>Ci) for any other photon-emitting radionuclide, and at least one of which has a principal photon energy between 100 keV and 500 keV; and

(C) test each dose calibrator for linearity upon installation and at intervals not to exceed three months thereafter over the range of use between 370 kBq (10 <<mu>>Ci) and the highest dosage that will be administered; and

(D) test each dose calibrator for geometry dependence upon installation over the range of volumes and volume configurations for which it will be used. The licensee shall keep a record of this test for the duration of the use of the dose calibrator.

(iii) Notwithstanding the provisions of § 175.103(c)(1)(ii), a licensee that must use a dose calibrator to measure the activity of beta-emitting radioactive materials to be administered to a patient shall perform additional checks specified in § 175.103(c)(1)(ii)(A) and (B) using the same radionuclide to be administered and having an activity of at least 50 percent, but not more than 200 percent, of the prescribed activity or by equivalent procedures approved by the Department. Records shall be kept pursuant to § 175.103(c)(1)(vi).

(iv) A licensee shall mathematically correct dosage readings for any geometry or linearity error that exceeds +- 10 percent if the dosage is greater than 370 kBq (10 <<mu>>Ci) and shall repair or replace the dose calibrator if the accuracy or constancy error exceeds +- 10 percent.

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(v) A licensee shall also perform checks and tests required by § 175.103(c)(1)(ii) following adjustment or repair of the dose calibrator.

(vi) A licensee shall retain a record of each check and test required by § 175.103(c)(1)(ii), (iii), and (v) for 3 years. Such records shall include:

(A) for § 175.103(c)(1)(ii)(A), the models and serial numbers of the dose calibrator and check source, the identity and calibrated activity of the radionuclide contained in the check source, the date of the check, the activity measured, the instrument settings, and the name of the individual who performed the check;

(B) for § 175.103(c)(1)(ii)(B), the model and serial number of the dose calibrator, the model and serial number of each source used and the identity of the radionuclide contained in the source and its activity, proof of traceability to a national standard, the date of the test, the results of the test, the instrument settings, and the signature of the radiation safety officer;

(C) for § 175.103(c)(1)(ii)(C), the model and serial number of the dose calibrator, the calculated activities, the measured activities, the date of the test, and the signature of the radiation safety officer; and

(D) for § 175.103(c)(1)(ii)(D), the model and serial number of the dose calibrator, the configuration and calibrated activity of the source measured, the activity of the source, the activity measured and the instrument setting for each volume measured, the date of the test, and the signature of the radiation safety officer.

(2) Calibration and check of survey instruments. (i) A licensee shall ensure that the survey instruments used to show compliance with this Code have been calibrated before first use, annually, and following repair.

(ii) To satisfy the requirements of § 175.103(c)(2)(i) the licensee shall:

(A) calibrate all scales with readings up to 10 mSv (1000 mrem) per hour with a radiation source, the intensity of which is determined to within 10 percent accuracy;

(B) for each scale that shall be calibrated, calibrate two readings at approximately one-third and two-thirds of the full scale reading; and

(C) conspicuously note on the instrument the apparent exposure rate from a dedicated check source as determined at the time of calibration, and the date of calibration.

(iii) To satisfy the requirements of § 175.103(c)(2)(ii) the licensee shall:

(A) consider a point as calibrated if the indicated exposure rate differs from the calculated exposure rate by not more than 10 percent; and

(B) consider a point as calibrated if the indicated exposure rate differs from

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the calculated exposure rate by not more than 20 percent and if a correction chart or graph is conspicuously attached to the instrument.

(iv) A licensee shall check each survey instrument for proper operation with the dedicated check source before each use. The licensee shall not be not required to keep records of these checks.

(v) The licensee shall retain a record of each calibration required in § 175.103(c)(2)(i) for 3 years. The record shall include:

(A) a description of the calibration procedure; and

(B) a description of the source used and the certified exposure rates from the source, and the rates indicated by the instrument being calibrated, the correction factors deduced from the calibration data, the signature of the individual who performed the calibration and the date of calibration.

(vi) To meet the requirements of § 175.103(c)(2)(i), (ii) and (iii), the licensee shall perform such calibrations as authorized by specific license condition or shall obtain the services of persons licensed by the U.S. Nuclear Regulatory Commission or an agreement state to perform calibrations of survey instruments. Records of calibrations which contain information required by § 175.103(c)(2)(v), shall be maintained by the licensee.

(3) Assay of radiopharmaceutical dosages. A licensee shall:

(i) assay, before medical use, the activity of each radiopharmaceutical dosage that contains more than 370 kBq (10 <<mu>>Ci) of any radionuclide;

(ii) assay, before medical use, the activity of each radiopharmaceutical dosage with a prescribed activity of 370 kBq (10 <<mu>>Ci) or less of a photon-emitting radionuclide to verify that the dosage does not exceed 370 kBq (10 <<mu>> Ci); and

(iii) Retain a record of the assays required by § 175.103(c)(3)(i) and (ii) for 3 years. To satisfy this requirement, the record shall contain the:

(A) generic name, trade name, or abbreviation of the radiopharmaceutical, its lot number, and expiration dates and the radionuclide;

(B) patient's name, and identification number if one has been assigned;

(C) prescribed dosage and activity of the dosage at the time of assay, or a notation that the total activity is less than 370 kBq (10 <<mu>>Ci);

(D) date and time of the assay; and

(E) initials of the individual who performed the assay.

(4) Authorization for calibration and reference sources. Any person authorized by

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§ 175.103(a)(2) for medical use of radioactive material may receive, possess, and use the following radioactive material for check, calibration and reference use:

(i) Sealed sources manufactured and distributed by persons specifically licensed pursuant to regulations of the U.S. Nuclear Regulatory Commission or equivalent regulations of an agreement state and that do not exceed 555 MBq (15 mCi) each;

(ii) Any radioactive material listed in § 175.103(d)(1) or (e)(1) with a half-life of 100 days or less in individual amounts not to exceed 555 MBq (15 mCi);

(iii) Any radioactive material listed in § 175.103(d)(1) or (e)(1) with a half life greater than 100 days in individual amounts not to exceed 7.4 MBq (200 <<mu>> Ci) each; and

(iv) Technetium-99m in individual amounts not to exceed 1.85 GBq (50 mCi).

(5) Requirements for possession of sealed sources and brachytherapy sources. (i) A licensee in possession of any sealed source or brachytherapy source shall follow the radiation safety and handling instructions approved by the U.S. Nuclear Regulatory Commission or any agreement state and supplied by the manufacturer on the label attached to the source, device or permanent container thereof, or in the leaflet or brochure that accompanies the source or device, and shall maintain the instructions for the duration of source use in a legible form convenient to users.

(ii) A licensee in possession of a sealed source or brachytherapy source shall conduct a physical inventory of all such sources at intervals not to exceed 3 months. The licensee shall retain each inventory record for 5 years. The inventory records must contain the model number of each source, and serial number if one has been assigned, the identity of each source radionuclide and its estimated activity, the location of each source, date of the inventory and the signature of the radiation safety officer.

(iii) A licensee in possession of a sealed source or brachytherapy source shall survey with a radiation survey instrument at intervals not to exceed 3 months all areas where such sources are stored. This shall not apply to teletherapy sources in teletherapy units or sealed sources in diagnostic devices.

(iv) A licensee shall retain a record of each survey required in § 175.103(c)(5)(iii) for 3 years. The record must include the date of the survey, a sketch of each area that was surveyed, the measured dose rate at several points in each area expressed in microsieverts (mrem) per hour, the model number and serial number of the survey instrument used to make the survey, and the signature of the radiation safety officer.

(6) Syringe shields and labels. (i) A licensee shall keep syringes that contain radioactive material to be administered in a radiation shield.

(ii) A licensee shall require each individual who prepares or administers radiopharmaceuticals/radiobiologics to use a syringe radiation shield unless the use of the shield is contraindicated for that patient.

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(iii) A licensee shall conspicuously label each syringe, or syringe radiation shield that contains a syringe with radioactive material, with the radiopharmaceutical/radiobiologic name or its abbreviation, the type of diagnostic study or therapy procedure to be performed, or the patient's name.

(7) Vial shields and labels. (i) A licensee shall require each individual preparing or handling a vial that contains radioactive material to keep the vial in a vial radiation shield.

(ii) A licensee shall conspicuously label each vial radiation shield that contains a vial of radioactive material with the radiopharmaceutical/radiobiologic name or its abbreviation.

(8) Surveys for contamination and ambient radiation dose rate. (i) A licensee shall survey with a radiation detection survey instrument at the end of each day of use all areas where radioactive materials are routinely prepared for use or administered.

(ii) A licensee shall survey with a radiation detection survey instrument at least once each week all areas where unsealed radioactive materials or radioactive wastes are stored.

(iii) A licensee shall conduct the surveys required by § 175.103(c)(8)(i) and (ii) so as to be able to detect and measure dose rates as low as 1 <<mu>>Sv (0.1 mrem) per hour.

(iv) A licensee shall establish dose rate action levels for the surveys required by § 175.103(c)(8)(i) and (ii) and shall require that the individual performing the survey immediately notify the radiation safety officer if a dose rate exceeds an action level.

(v) A licensee shall perform wipe tests for removable contamination once each week on all areas where radioactive materials are routinely prepared for use or administered and where unsealed sources of radioactive materials are stored.

(vi) A licensee shall perform the wipe tests required by § 175.103(c)(8)(v) so as to be able to detect contamination on each wipe sample of 35 Bq (2000 disintegrations or transformations per minute).

(vii) A licensee shall establish removable contamination action levels for the surveys required by § 175.103(c)(8)(v) and shall require that the individual performing the survey immediately notify the radiation safety officer if contamination exceeds action levels.

(viii) A licensee shall retain a record of each survey or wipe test required by § 175.103(c)(8)(i), (ii) and (v) of this section for 3 years. The record must include the date of the survey, a sketch of each area surveyed, action levels established for each area, the measured dose rate at several points in each area expressed in <<mu>>Sv (mrem) per hour or the removable contamination in each area expressed in

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becquerels (disintegrations or transformations per minute) per 100 square centimeters, the serial number and the model number of the instrument used to make the survey or analyze the samples, and the initials of the individual who performed the survey.

(9) Release of patients containing radioactive materials. (i) Notwithstanding the provisions of § 175.03(d)(1), a licensee shall not authorize release from confinement for medical care any patient administered radioactive material as either a radiopharmaceutical/radiobiologic or as a permanent source implant until the activity in the patient is such that the total effective dose equivalent for the individual (other than the patient) likely to receive the greatest dose is 5 mSv (500 mrem) or less.

(ii) When the total effective dose equivalent to any individual that could result from the release of a patient is likely to exceed 1 mSv (100 mrem), the licensee shall provide the patient, or the patient's competent representative, written information on risks of radiation and methods for reducing the exposure of individuals, and shall keep records of such patient releases for a period of five (5) years.

(iii) [Reserved]

(iv) Radioactive cadavers. (A) If any patient containing radioactive material administered/implanted for therapeutic purposes dies, it shall be the responsibility of the physician who pronounces such patient as dead to notify immediately the physician in charge of the case or such physician's designated representative.

(B) No person shall commence any autopsy on any cadaver that contains more than 185 MBq (5 mCi) of radioactive material administered/implanted for therapeutic purposes without first having consulted with, and having been advised by, the radiation safety officer of the hospital or the physician responsible for the administration/implantation of the radioactive material. If neither is available, a designated representative may serve.

(C) A radioactivity report on every cadaver containing more than 185 MBq (5 mCi) of radioactive material administered/implanted for therapeutic purposes shall be completed by the radiation safety officer or the physician responsible for the administration of the radioactive material or their designated representative. The report must include the name, address and radioactive materials license number of the hospital; the name of the deceased; the name, address and telephone number of the next of kin; the name, address and telephone number of the funeral home to which the deceased will be sent; the radionuclide involved; the approximate activity on the date of the report and the physical form; the location(s) of the radioactive materials within the body and the external dose rate at the body surface closest to the source; the precautions to be observed during autopsy or handling of the body by the funeral director; and the name of the person who prepared the form. This report shall accompany the body, whether autopsied or not, when it is surrendered to the funeral director. The Department shall be notified in person, by telephone, by mailgram or by facsimile within 24 hours of the death and a copy of the radioactivity report shall be sent to the Department within fifteen (15) days of the date of death.

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(10) Storage of volatiles and gases. (i) A licensee shall store volatile radiopharmaceuticals and radioactive gases in the shippers' radiation shield and container.

(ii) After drawing the first dosage, a licensee shall store and use a multidose container in a properly functioning fume hood.

(11) Decay-in-storage. (i) A licensee may hold radioactive material with a physical half-life of 65 days or less for decay-in-storage before disposal in ordinary trash and is exempt from the requirements of § 175.104 of this Code if the licensee:

(A) holds radioactive material for decay a minimum of 10 half-lives;

(B) monitors radioactive material at the container surface before disposal as ordinary trash and determines that its radioactivity cannot be distinguished from natural background radiation levels with a radiation detection survey instrument set on its most sensitive scale and with no interposed shielding;

(C) removes or obliterates all radiation labels; and

(D) separates and monitors each generator column individually with all radiation shielding removed to ensure that its contents have decayed to natural background radiation levels before disposal.

(ii) For radioactive material disposed in accordance with § 175.103 (c)(11)(i), the licensee shall retain a record of each disposal for 3 years. The record must include the date of the disposal, the date on which the radioactive material was placed in storage, the model and serial number of the survey instrument used, the background dose rate, the radiation dose rate measured at the surface of each waste container, and the name of the individual who performed the disposal.

(d) Uptake, dilution and excretion. (1) Use of radiopharmaceuticals for uptake, dilution or excretion studies.

(i) A licensee may use any radioactive material authorized in a license in any radiopharmaceutical or radiobiologic for diagnostic studies involving the measurement of uptake, dilution or excretion.

(ii) A licensee shall prepare and use authorized radiopharmaceuticals or radiobiologics in accordance with the manufacturer's instructions for radiation safety.

(2) Possession of a survey instrument. A licensee authorized to use radioactive material for uptake, dilution, and excretion studies shall have in its possession a portable radiation detection survey instrument capable of detecting dose rates over the range 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour. The instrument shall be operable and calibrated in accordance with § 175.103(c)(2).

(e) Imaging and localization. (1) Use of radiopharmaceuticals, generators, and

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reagent kits for imaging and localization studies.

(i) A licensee may use any radioactive material authorized in a license in any diagnostic radiopharmaceutical, radiobiologic or reagent kit, except for aerosol or gaseous forms, for preparation and diagnostic use of a radiopharmaceutical or radiobiologic for studies involving imaging and localization.

(ii) A licensee may use generators upon approval of the Department.

(iii) Provided the conditions of § 175.103(e)(3) are met, a licensee shall use radioactive aerosols or gases only if specific application is made to and approved by the Department.

(iv) A licensee shall prepare and use authorized radiopharmaceuticals or radiobiologics in accordance with the manufacturer's instructions for radiation safety.

(v) A licensee shall elute generators in compliance with § 175.103(e)(2) and prepare radiopharmaceuticals from kits in accordance with the manufacturer's instructions for radiation safety.

(2) Permissible molybdenum-99 concentration. (i) A licensee shall not administer to humans a radiopharmaceutical containing more than 5.55 kBq (0.15 <<mu>>Ci) of molybdenum-99 per 37 MBq (1.0 mCi) of technetium-99m.

(ii) A licensee preparing technetium-99m radiopharmaceuticals from molybdenum-99/technetium-99m generators shall measure the molybdenum-99 concentration in each eluate or extract.

(iii) A licensee who must measure molybdenum concentration shall retain a record of each measurement for 3 years. The record shall include, for each elution or extraction of technetium-99m, the measured activity of the technetium-99m expressed in MBq (mCi), the measured activity of the molybdenum expressed in kBq (<<mu>>Ci), the ratio of the measures expressed as kBq (<<mu>>Ci) of molybdenum per MBq (mCi) of technetium-99m, the date of the test, and the initials of the individual who performed the test.

(iv) A licensee shall report immediately to the Bureau of Radiological Health each occurrence of molybdenum-99 concentration exceeding the limits specified in § 175.103(e)(2)(1).

(3) Control of aerosols and gases. (i) A licensee who administers radioactive aerosols or gases shall do so with a system that will keep airborne concentrations within the limits prescribed by § 175.03 of this Code.

(ii) The system shall provide for collection and decay or disposal of the aerosol or gas in a shielded container.

(iii) Before receiving, producing, using, or storing a radioactive gas, the licensee shall calculate the amount of time needed after a release to reduce the concentration in the area of use to the ALI listed in Table 1 of Appendix A of §

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175.03 of this Code. The calculation shall be based on the highest activity of gas handled in a single container and the measured available air exhaust rate.

(iv) A licensee shall post the time calculated in § 175.103(e)(3)(iii) at the area of use, as well as safety measures to be instituted in case of a spill at the area of use.

(v) A licensee shall check the operation of collection systems monthly and measure the ventilation rates in areas of use at intervals not to exceed 6 months. Records of these checks and measurements shall be maintained for 3 years.

(vi) A copy of the calculations, including assumptions, measurements and calculations made, required in § 175.103(e)(3)(iii) shall be recorded and retained for the duration of the license.

(4) Possession of survey instruments. A licensee authorized to use radioactive material for imaging and localization studies shall have in its possession a portable, radiation detection survey instrument capable of detecting dose rates over the range of 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour, and a portable radiation measurement survey instrument capable of measuring dose rates over the range 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The instruments shall be operable and calibrated in accordance with § 175.103(c)(2).

(f) Radiopharmaceuticals/radiobiologics for therapy. (1) Use of radiopharmaceuticals/radiobiologics for therapy.

(i) A licensee may use any radioactive material authorized in a license in any radiopharmaceutical or radiobiologic for a therapeutic use.

(ii) The licensee shall prepare and use authorized radiopharmaceuticals or radiobiologics in accordance with the manufacturer's instructions for radiation safety.

(2) Safety instruction. (i) A licensee shall provide oral and written radiation safety instruction for all personnel caring for patients undergoing radiopharmaceutical therapy. Refresher training shall be provided at intervals not to exceed 1 year.

(ii) To satisfy § 175.103(f)(2)(i), the instruction shall describe the licensee's procedures for:

(A) patient control;

(B) visitor control;

(C) contamination control;

(D) waste control; and

(E) notification of the radiation safety officer or authorized user in case of the patient's death or medical emergency.

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(iii) A licensee shall keep for 3 years a list of individuals receiving instruction required by § 175.103(f)(2)(i), a description of the instruction, the date of instruction, and the name of the individual who gave the instruction.

(3) Safety precautions. (i) For each patient receiving radiopharmaceutical therapy and hospitalized for compliance with § 175.103(c)(9), a licensee shall:

(A) provide a private room with a private sanitary facility;

(B) post the patient's door with a "Caution: Radioactive Material" sign and note on the door or on the patient's chart where and how long visitors may stay in the patient's room;

(C) authorize visits by individuals under age 18 only on a patient-by-patient basis with the approval of the authorized user after consultation with the radiation safety officer;

(D) promptly after administration of the dosage, measure the dose rates in contiguous restricted and unrestricted areas with a radiation measurement survey instrument to demonstrate compliance with the requirements of § 175.03 of this Code and retain for 3 years a record of each survey that includes the time and date of the survey, a plan of the area or list of points surveyed, the measured dose rate at several points expressed in <<mu>>Sv (mrem) per hour, the instrument used to make the survey, and the initials of the individual who made the survey;

(E) either monitor material and items removed from the patient's room to determine that any contamination cannot be distinguished from the natural background radiation level with a radiation detection survey instrument set on its most sensitive scale and with no interposed shielding, or handle these materials and items as radioactive waste;

(F) provide the patient with radiation safety guidance that will help to keep radiation dose to household members and the public as low as reasonably achievable before authorizing release of the patient;

(G) survey the patient's room and private sanitary facility for removable contamination with a radiation detection survey instrument before assigning another patient to the room. The room must not be reassigned until removable contamination is less than 5 Bq (l200 disintegrations per minute) per 100 square centimeters; and

(H) measure the thyroid burden of each individual who helped prepare or administer a dosage of iodine-131 within 3 days after administering the dosage, and retain for the period required by § 175.03(k) a record of each thyroid burden measurement, date of measurement, the name of the individual whose thyroid burden was measured, and the initials of the individual who made the measurements.

(ii) A licensee shall notify the radiation safety officer or the authorized user immediately if the patient dies or has a medical emergency.

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(4) Possession of survey instruments. A licensee authorized to use radioactive material for radiopharmaceutical therapy shall have in its possession a portable radiation detection survey instrument capable of detecting dose rates over the range 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour, and a portable radiation measurement survey instrument capable of measuring dose rates over the range 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The instruments shall be operable and calibrated in accordance with § 175.103(c)(2).

(g) Sealed sources for diagnosis. (1) Use of sealed sources for diagnosis. A licensee shall use the following sealed sources in accordance with the manufacturer's radiation safety, handling and maintenance instructions:

(i) Iodine-125 as a sealed source in a device for bone mineral analysis;

(ii) Americium-241 as a sealed source in a device for bone mineral analysis;

(iii) Gadolinium-153 as a sealed source in a device for bone mineral analysis; and

(iv) Iodine-125 as a sealed source in a portable device for imaging.

(2) Availability of survey instrument. A licensee authorized to use radioactive material as a sealed source for diagnostic purposes shall have available for use a portable radiation detection survey instrument capable of detecting dose rates over the range 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour or a portable radiation measurement survey instrument capable of measuring dose rates over the range 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The instrument shall be operable and calibrated in accordance with § 175.103(c)(2).

(h) Sealed sources for brachytherapy. (1) Use of sealed sources for brachytherapy. A licensee shall use the following sources in accordance with the manufacturer's radiation safety, handling and maintenance instructions:

(i) Cesium-137 as a sealed source in needles and applicator cells for topical, interstitial, and intracavitary treatment of cancer;

(ii) Cobalt-60 as a sealed source in needles and applicator cells for topical, interstitial, and intracavitary treatment of cancer;

(iii) Gold-198 as a sealed source in seeds for interstitial treatment of cancer;

(iv) Iodine-125 as a sealed source in seeds for interstitial treatment of cancer as a permanent implant;

(v) Iridium-192 as seeds encased in nylon ribbon for interstitial treatment of cancer;

(vi) Palladium-103 as seeds for the interstitial treatment of cancer;

(vii) Strontium-90 as a sealed source in an applicator for treatment of superfi-

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cial eye conditions; and

(viii) any other brachytherapy source authorized in the license.

(2) Safety instruction. (i) The licensee shall provide oral and written radiation safety instruction to all personnel caring for a patient receiving implant therapy. Refresher training shall be provided at intervals not to exceed 1 year.

(ii) To satisfy § 175.103(h)(2)(i), the instruction shall describe:

(A) size and appearance of the brachytherapy sources;

(B) safe handling and shielding instructions in case of a dislodged source;

(C) procedures for patient control;

(D) procedures for visitor control; and

(E) procedures for notification of the radiation safety officer or authorized user if the patient dies or has a medical emergency.

(iii) A licensee shall retain for 3 years a record of individuals receiving instruction required by § 175.103(h)(2)(i), a description of the instruction, the date of instruction, and the name of the individual who gave the instruction.

(3) Safety precautions. (i) For each patient receiving implant therapy a licensee shall:

(A) not place the patient in the same room with a patient who is not receiving radiation therapy unless the licensee can demonstrate compliance with the requirements of § 175.03 of this Code at a distance of one meter from the implant;

(B) post the patient's door with a "Caution: Radioactive Materials" sign and note on the door or in the patient's chart where and how long visitors may stay in the patient's room;

(C) authorize visits by individuals under age 18 only on a case-by-case basis with the approval of the authorized user after consultation with the radiation safety officer;

(D) promptly after implanting the sources, survey the dose rates in contiguous restricted and unrestricted areas with a radiation measurement survey instrument to demonstrate compliance with § 175.03 of this Code and retain for 3 years a record of each survey that includes the time and date of the survey, a sketch of the area or list of points surveyed, the measured dose rate at several points expressed in <<mu>>Sv (mrem) per hour, the instrument used to make the survey, and the initials of the individual who made the survey; and

(E) provide the patient with radiation safety guidance that will help to keep the

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radiation dose to household members and the public as low as reasonably achievable before releasing the patient if the patient was administered a permanent implant.

(ii) A licensee shall notify the radiation safety officer or authorized user immediately if the patient dies or has a medical emergency.

(4) Brachytherapy sources inventory. (i) A licensee shall promptly return brachytherapy sources to their place of storage after removal from the patient.

(ii) Each time brachytherapy sources are returned to an area of storage from an area of use, the licensee shall immediately count or otherwise verify the number returned to ensure that all sources taken from the storage area have been returned.

(iii) A licensee shall make a record of brachytherapy source use which includes:

(A) the names of the individuals permitted to handle the sources;

(B) the number and activity of sources removed from storage, the room number of use and patient's name, the time and date they were removed from storage, the number and activity of the sources in storage after the removal, and the initials of the individual who removed the sources from storage; and

(C) the number and activity of sources returned to storage, the room number of use and patient's name, the time and date they were returned to storage, the number and activity of sources in storage after the return, and the initials of the individual who returned the sources to storage.

(iv) Immediately after implanting sources in a patient and immediately after removal of sources from a patient, including those sources inserted and removed by remote after loading devices, the licensee shall make a radiation survey of the patient and the area of use to confirm that no sources have been misplaced, left in the patient, or otherwise not returned to the appropriate place of storage. The licensee shall make a record of each survey.

(v) A licensee shall retain the records required in § 175.103(h)(4)(iii) and (iv) for 3 years.

(5) Release of patients treated with temporary implants. (i) Immediately after removing the last temporary implant source from a patient, the licensee shall make a radiation survey of the patient with a radiation detection survey instrument to confirm that all sources have been removed. The licensee shall not release from confinement for medical care a patient treated by temporary implant until all sources have been removed.

(ii) A licensee shall retain a record of patient surveys which demonstrate compliance with § 175.103(h)(5)(i) for 3 years. Each record must include the date of the survey, the name of the patient, the dose rate from the patient expressed as <<mu>>Sv (mrem) per hour and measured within 1 meter from the patient, and the initials of the individual who made the survey.

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(6) Possession of survey instruments. A licensee authorized to use radioactive material for implant therapy shall have in its possession a portable radiation detection survey instrument capable of detecting rates over the range 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour, and a portable radiation measurement survey instrument capable of measuring dose rates over the range 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The instruments shall be operable and calibrated in accordance with § 175.103(c)(2).

(i) Teletherapy. (1) Use of a sealed source in a teletherapy unit. A licensee shall use cobalt-60 as a sealed source in a teletherapy unit for medical use in accordance with the manufacturer's radiation safety and operating instructions.

(2) Maintenance, repair and inspection requirements. (i) Routine maintenance of the teletherapy unit shall be performed according to the manufacturer's recommended schedule, including periodic replacement of parts. Replacement parts shall be those specified by the manufacturer of the teletherapy unit.

(ii) A licensee shall have each teletherapy unit fully inspected and serviced during teletherapy source replacement or at intervals not to exceed 5 years, whichever comes first.

(iii) Repairs found to be needed shall be made promptly.

(iv) Only a person specifically licensed by the U.S. Nuclear Regulatory Commission or an agreement state to perform teletherapy unit inspections and servicing shall perform such functions, including, but not limited to installation, relocation, or removal of a teletherapy sealed source or a teletherapy unit that contains a sealed source or maintenance, adjustment, or repair of the source drawer, the shutter or other mechanism of a teletherapy unit that could expose the source, reduce the shielding around the source, result in increased radiation levels or compromise the safety of the unit.

(v) A licensee shall keep record of the inspections and servicing for the duration of the license. The record shall contain the name of the person performing the inspection or service, such person's license number, the date of inspection, the manufacturer's name and model number and serial number for both the teletherapy unit and source, a list of components inspected, the findings of the inspection, a list of components serviced and the type of service, a list of components replaced, and the signature of the licensed individual performing the service or inspection.

(3) Amendments. In addition to the requirements specified in § 175.103(a)(4), a licensee shall apply for and shall have received a license amendment before:

(i) making any change in the treatment room shielding;

(ii) making any change in the location of the teletherapy unit within the treatment room;

(iii) using the teletherapy unit in a manner that could result in increased radiation levels in areas outside the teletherapy treatment room;

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(iv) relocating the teletherapy unit; or

(v) allowing an individual not listed on the licensee's license to perform the duties of the radiation therapy physicist.

(4) Safety instruction. (i) A licensee shall post written instructions at the teletherapy unit console. To satisfy this requirement, these instructions shall inform the operator of:

(A) the procedure to be followed to ensure that only the patient is in the treatment room before turning the primary beam of radiation on to begin a treatment or after a door interlock interruption; and

(B) the procedure to be followed if:

(a) the operator is unable to turn the primary beam of radiation off with controls outside the treatment room or any other abnormal operation occurs; and

(b) the names and telephone numbers of the authorized users and radiation safety officer to be immediately contacted if the teletherapy unit or console operates abnormally.

(ii) A licensee shall provide instruction in the topics identified in § 175.103(i)(4)(i) of this section to all individuals who operate a teletherapy unit and shall provide appropriate refresher training to individuals at intervals not to exceed 1 year. Such instruction shall include "dry runs" of emergency procedures.

(iii) A licensee shall retain for 3 years a record of individuals receiving instruction required by § 175.103(2)(4)(ii) of this section, a description of the instruction, the date of instruction, and the name of the individual who gave the instruction.

(5) Doors, interlocks, and warning systems. (i) A licensee shall control access to the teletherapy room by a door at each entrance.

(ii) A licensee shall equip each entrance to the teletherapy room with an electrical interlock system that shall:

(A) prevent the operator from turning the primary beam of radiation on unless each treatment room entrance door is closed;

(B) turn the primary beam of radiation off immediately when an entrance door is opened; and

(C) prevent the primary beam of radiation from being turned on following an interlock interruption until all treatment room entrance doors are closed and the beam on-off control is reset at the console.

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(iii) A licensee shall equip each entrance to the teletherapy room with a beam condition indicator light.

(6) Possession of survey instrument. A licensee authorized to use radioactive material in a teletherapy unit shall have in its possession either a portable radiation detection survey instrument capable of detecting dose rates over the range 1.0 <<mu>>Sv (0.1 mrem) per hour to 1000 <<mu>>Sv (100 mrem) per hour, or a portable radiation measurement survey instrument capable of measuring dose rates over the range 10 <<mu>>Sv (1 mrem) per hour to 10 mSv (1000 mrem) per hour. The instruments shall be operable and calibrated in accordance with § 175.103(c)(2).

(7) Radiation monitoring device. (i) A licensee shall have in each teletherapy room a permanent radiation monitor capable of continuously monitoring beam status.

(ii) Each radiation monitor shall be capable of providing visible notice of a teletherapy unit malfunction that results in an exposed or partially exposed source. The visible indicator of high radiation levels must be observable by an individual entering the teletherapy room.

(iii) Each radiation monitor shall be equipped with a backup power supply separate from the power supply to the teletherapy unit. This backup power supply may be a battery system.

(iv) A radiation monitor must be checked with a dedicated check source for proper operation each day before the teletherapy unit is used for treatment of patients.

(v) A licensee shall maintain a record of the check required by § 175.103(i)(7)(iv) for 3 years. The record shall include the date of the check, notation that the monitor indicates when the source is exposed, and the initials of the individual who performed the check.

(vi) If a radiation monitor is inoperable, the licensee shall require any individual entering the teletherapy room to use a survey instrument or audible alarm personal dosimeter to monitor for any malfunction of the source exposure mechanism that may result in an exposed or partially exposed source. The instrument or dosimeter shall be checked with a dedicated check source for proper operation at the beginning of each day of use. The licensee shall keep a record as described in § 175.103(i)(7)(v).

(vii) A licensee shall promptly repair or replace the radiation monitor if it is inoperable.

(8) Viewing and communications system. (i) A licensee shall construct or equip each teletherapy room to permit continuous observation of the patient by a licensed radiation therapy technician or physician from the teletherapy unit console during irradiation.

(ii) A licensee shall construct or equip each teletherapy room to provide a means of communication with the patient from the teletherapy unit console during irradiation.

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(9) Dosimetry equipment. (i) A licensee shall have a calibrated dosimetry system available for use. The system shall have been calibrated for cobalt-60 by the National Institute of Standards and Technology (formerly the National Bureau of Standards) or by an American Association of Physicists in Medicine (AAPM) Accredited Dosimetry Calibration Laboratory (ADCL). The calibration shall have been performed within the previous two (2) years and after any servicing that may have affected system calibration.

(ii) The licensee shall have available for use a dosimetry system for spot check measurements. To meet this requirement, the system may be compared with a system that has been calibrated in accordance with § 175.103(i)(9)(i). This comparison shall have been performed within the previous twelve (12) months (6 months if an ionization chamber) and after each servicing that may have affected system calibration. Alternatively, the spot check system may be the same system used to meet the requirement in § 175.103(i)(9)(i).

(iii) The licensee shall retain a record of each calibration, intercomparison, and comparison for the duration of the license. For each calibration, intercomparison, or comparison, the record shall include the date, the model numbers and serial numbers of the instruments that were calibrated, intercompared, or compared as required by § 175.103(i)(9)(i) and (ii), the correction factors that were deduced, the names of the individuals who performed the calibration, intercomparison, or comparison, and evidence that the intercomparison meeting was sanctioned by a calibration laboratory or radiologic physics center accredited by AAPM.

(10) Full calibration measurements. (i) A licensee authorized to use a teletherapy unit for medical use shall perform full calibration measurements on each teletherapy unit:

(A) before the first medical use of the unit; and

(B) before medical use under the following conditions:

(a) whenever spot check measurements indicate that the output differs by more than 5 percent from the output obtained at the last full calibration corrected mathematically for radioactive decay;

(b) following replacement of the source or following reinstallation of the teletherapy unit in a new location;

(c) following any repair of the teletherapy unit that includes removal of the source or major repair of the components associated with the source exposure assembly; and

(C) at intervals not exceeding 1 year.

(ii) To satisfy the requirement of § 175.103(i)(10)(i), full calibration measurements shall include determination of:

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(A) the output within +- 3 percent for the entire range of field sizes and for the distance or entire range of distances used for medical use;

(B) the coincidence of the radiation field and the field indicated by the light beam localizing device;

(C) the uniformity of the radiation field and its dependence on the orientation of the useful beam;

(D) timer constancy and linearity over the range of uses;

(E) on-off error; and

(F) the accuracy of all distance measuring and localization devices in medical use.

(iii) A licensee shall use the dosimetry system described in § 175.103(i)(9) to measure the output for one set of exposure conditions. The remaining radiation measurements required in § 175.103(i)(10)(ii)(A) of this section may then be made using a dosimetry system that indicates relative dose rates.

(iv) A licensee shall make full calibration measurements required by § 175.103(i)(10)(i) in accordance with either the procedures recommended by the Scientific Committee on Radiation Dosimetry of the AAPM that are described in Physics in Medicine and Biology Vol. 16, No. 3, 1971, pp. 379-396, or by Task Group 21 of the Radiation Therapy Committee of the American Association of Physicists in Medicine that are described in Medical Physics Vol. 10, No. 6, 1983, pp. 741-771, and Vol. 11, No. 2, 1984, p. 213, or any superseding protocol.

(v) A licensee shall correct mathematically the outputs determined in § 175.103(i)(10)(ii)(A) for physical decay for intervals not exceeding one month for cobalt-60.

(vi) Full calibration measurements required by § 175.103(i)(10)(i) and physical decay corrections required by § 175.103(i)(10)(v) shall be performed by the radiation therapy physicist named on the license.

(vii) A licensee shall retain a record of each calibration for the duration of the license. The record shall include the date of the calibration, the manufacturer's name, model number, and serial number or other unambiguous identification of both the teletherapy unit and the source and of the instruments used to calibrate the teletherapy unit, tables that describe the output of the unit over the range of field sizes and for the range of distances used in radiation therapy, a determination of the coincidence of the radiation field and the field indicated by the light beam localizing device, the measured timer accuracy for a typical treatment time, the calculated on-off error, the estimated accuracy of each distance measuring or localization device, and the signature of the radiation therapy physicist.

(11) Periodic spot checks. (i) A licensee authorized to use teletherapy units for medical use shall perform output spot checks on each teletherapy unit once each ca-

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lendar month.

(ii) To satisfy the requirement of § 175.103(i)(11)(i), measurements shall include determination of:

(A) timer constancy and timer linearity over the range of use;

(B) on-off error;

(C) the coincidence of the radiation field and the field indicated by the light beam localizing device;

(D) the accuracy of all distance measuring and localization devices used for medical use;

(E) the output for one typical set of operating conditions; and

(F) the difference between the measurement made in § 175.103 (i)(11)(ii)(E) and the anticipated output, expressed as a percentage of the anticipated value obtained at last full calibration corrected mathematically for physical decay.

(iii) A licensee shall use the dosimetry system described in § 175.103(i)(9) to make the measurement required in § 175.103(i)(11)(ii)(E).

(iv) A licensee shall perform measurements required by § 175.103(i)(11)(i) in accordance with procedures established by the radiation therapy physicist. That individual does not need to actually perform the output spot check measurements.

(v) A licensee shall have the radiation therapy physicist review the results of each output spot check within 15 days. The radiation therapy physicist shall promptly notify the licensee in writing of the results of each output spot check. The licensee shall keep a copy of each written notification for 3 years.

(vi) A licensee authorized to use a teletherapy unit for medical use shall perform safety spot checks of each teletherapy facility once each calendar month.

(vii) To satisfy the requirement of § 175.103(i)(11)(vi), checks shall ensure proper operation of:

(A) electrical interlocks at each teletherapy room entrance;

(B) electrical or mechanical stops installed for the purpose of limiting use of the primary beam of radiation, restriction of source housing angulation or elevation, carriage or stand travel and operation of the beam on-off mechanism;

(C) beam condition indicator lights on the teletherapy unit, on the control console, and in the facility;

(D) viewing systems;

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(E) treatment room doors from inside and outside the treatment room; and

(F) electrically assisted treatment room doors with the teletherapy unit electrical power turned "off".

(viii) A licensee shall lock the control console in the "off" position if any door interlock malfunctions. No licensee shall use the unit until the interlock system is repaired unless specifically authorized by the Department.

(ix) A licensee shall promptly repair any system identified in § 175.103(i)(11)(vii) that is not operating properly. The teletherapy unit shall not be used until all repairs are completed.

(x) A licensee shall retain a record of each spot check required by § 175.103(i)(11)(i) and (vi) for 3 years. The record shall include the date of the spot check, the manufacturer's name, model number, and serial number for both the teletherapy unit and source, the manufacturer's name, model number and serial number of the instrument used to measure the output of the teletherapy unit, the measured timer accuracy, the calculated on-off error, a determination of the coincidence of the radiation field and the field indicated by the light beam localizing device, the measured timer accuracy for a typical treatment time, the calculated on-off error, the estimated accuracy of each distance measuring or localization device, the difference between the anticipated output and the measured output, notations indicating the operability of each entrance door electrical interlock, each electrical or mechanical stop, each beam condition indicator light, the viewing system and doors, and the signature of the individual who performed the periodic spot check.

(12) Radiation surveys for teletherapy facilities. (i) Before medical use, after each installation of a teletherapy source, and after making any change for which an amendment is required by § 175.103(i)(3) the licensee shall perform radiation surveys with an operable radiation measurement survey instrument calibrated in accordance with § 175.103(c)(2) to verify that:

(A) the maximum and average radiation levels at 1 meter from the teletherapy source with the source in the off position and the collimators set for a normal treatment field do not exceed 100 <<mu>>Sv (10 mrem) per hour and 20 <<mu>>Sv (2 mrem) per hour, respectively; and

(B) with the teletherapy source in the on position with the largest clinically available treatment field and with a scattering phantom in the primary beam of radiation, that:

(a) radiation levels in restricted areas are not likely to cause personnel exposures in excess of the limits specified in § 175.03; and

(b) radiation levels in unrestricted areas do not exceed the limits specified in § 175.03.

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(ii) If the results of the surveys required in § 175.103(i)(12)(i) indicate any radiation levels in excess of the respective limit specified in § 175.103(i)(12)(i)(A) or (B), the licensee shall lock the control in the "off" position and not use the unit:

(A) except as may be necessary to repair, replace, or test the teletherapy unit, the teletherapy unit shielding or the treatment room shielding; or

(B) until the licensee has received a specific exemption from the Department.

(iii) A licensee shall retain a record of the radiation measurements made following installation of a source for the duration of the license. The record shall include the date of the measurements, the reason the survey is required, the manufacturer's name, model number and serial number of the teletherapy unit, the source, and the instrument used to measure radiation levels, each dose rate measured around the teletherapy source while in the off position and the average of all measurements, a plan of the areas surrounding the treatment room that were surveyed, the measured dose rate at several points in each area expressed in <<mu>>Sv (mrem) per hour, the calculated maximum level of radiation over a period of 1 week for each restricted and unrestricted area, and the signature of the radiation safety officer.

(13) Safety checks for teletherapy facilities. (i) A licensee shall promptly check all systems listed in § 175.103(i)(11) for proper function after each installation of a teletherapy source and after making any change for which an amendment is required by § 175.103(i)(3).

(ii) If the results of the checks required in § 175.103(i)(12)(i) indicate the malfunction of any system specified in § 175.103(i)(11), the licensee shall lock the control console in the "off" position and not use the unit except as may be necessary to repair, replace, or check the malfunctioning system.

(iii) A licensee shall retain for 3 years a record of the facility checks following installation of a source. The record shall include notations indicating the operability of each entrance door interlock, each electrical or mechanical stop, each beam condition indicator light, the viewing system, and doors, and the signature of the radiation safety officer.

(14) Modification of a teletherapy unit or room before beginning a treatment program. (i) If the survey required by § 175.103(i)(l2) indicates that an individual in an unrestricted area may be exposed to levels of radiation greater than those permitted by § 175.03, before beginning the treatment program the licensee shall:

(A) either equip the unit with stops or add additional radiation shielding to ensure compliance with § 175.03;

(B) perform the survey required by § 175.103(i)(12) again; and

(C) include in the report required by § 175.103(i)(15) the results of the initial survey, a description of the modification made to comply with § 175.103(i)(14)(i)

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and the results of the second survey; or

(D) request and receive a license amendment that authorizes radiation levels in unrestricted areas greater than those permitted by § 175.03.

(15) Reports of teletherapy surveys, checks, tests, and measurements. A licensee shall furnish a copy of the records required in § 175.103(i)(12), (13) and (14) and the output from the teletherapy source expressed as Sv (rem) per hour at one meter from the source and determined during the full calibration required in § 175.103(i)(10) to the Bureau of Radiological Health within 30 days following completion of the action that initiated the record requirement.

(j) Training and experience requirements. (1) Radiation safety officer. Except as provided in § 175.103(j)(2), an individual fulfilling the responsibilities of the radiation safety officer as provided in § 175.103(b)(2) shall:

(i) be certified by the:

(A) American Board of Health Physics in Comprehensive Health Physics; or

(B) American Board of Radiology; or

(C) American Board of Nuclear Medicine; or

(D) American Board of Science in Nuclear Medicine; or

(E) American Board of Medical Physics; or

(F) Board of Pharmaceutical Specialties in Nuclear Pharmacy; or

(ii) have had 200 hours of classroom and laboratory training as follows:

(A) radiation physics and instrumentation;

(B) radiation protection;

(C) mathematics pertaining to the use and measurement of radioactivity;

(D) radiation biology;

(E) radiopharmaceutical chemistry; and

(F) 1 year of full time experience in radiation safety at a medical institution under the supervision of the individual identified as the radiation safety officer on a Department, agreement state, or U.S. Nuclear Regulatory Commission license that authorizes the medical use of radioactive material; or

(iii) be an authorized user for those radioactive material uses that come within the radiation safety officer's responsibilities.

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(2) Training for experienced radiation safety officer. An individual identified as a radiation safety officer on a Department, agreement state or U.S. Nuclear Regulatory Commission license on August 15, 1990 who oversees only the use of radioactive material for which the licensee was authorized on that date need not comply with the training requirements of § 175.103(j)(1).

(3) Training for uptake, dilution, or excretion studies. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user of radioactive material listed in § 175.103(d)(1)(i) to be a physician who:

(i) is certified in:

(A) Nuclear medicine by the American Board of Nuclear Medicine; or

(B) Diagnostic radiology by the American Board of Radiology; or

(C) Diagnostic radiology or radiology by the American Osteopathic Board of Radiology; or

(ii) has completed 40 hours of instruction in basic radioisotope handling techniques applicable to the use of prepared radiopharmaceuticals, and 20 hours of supervised clinical experience:

(A) to satisfy the basic instruction requirement, 40 hours of classroom and laboratory instruction shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity;

(d) radiation biology; and

(e) radiopharmaceutical chemistry.

(B) to satisfy the requirement for 20 hours of supervised clinical experience, training must be under the supervision of an authorized user at a medical institution and shall include:

(a) examining patients and reviewing their case histories to determine their suitability for radioisotope diagnosis, limitations, or contraindications;

(b) selecting the suitable radiopharmaceuticals and calculating and measuring the dosages;

(c) administering dosages to patients and using syringe radiation shields;

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(d) collaborating with the authorized user in the interpretation of radioisotope test results; and

(e) patient followup; or

(C) has successfully completed a 6 month training program in nuclear medicine as part of a training program that has been approved by the Accreditation Council for Graduate Medical Education and that included classroom and laboratory training, work experience, and supervised clinical experience in all the topics identified in § 175.103(j)(3)(ii).

(4) Training for imaging and localization studies. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user of a radiopharmaceutical, radiobiologic, generator, or reagent kit specified in § 175.103(e)(1) to be a physician who:

(i) is certified in:

(A) Nuclear medicine by the American Board of Nuclear Medicine; or

(B) Diagnostic radiology by the American Board of Radiology; or

(C) Diagnostic radiology or radiology by the American Osteopathic Board of Radiology; or

(ii) has completed 200 hours of instruction in basic radioisotope handling techniques applicable to the use of prepared radiopharmaceuticals, generators, and reagent kits, 500 hours of supervised work experience and 500 hours of supervised clinical experience:

(A) to satisfy the basic instruction requirement, 200 hours of classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity;

(d) radiopharmaceutical chemistry; and

(e) radiation biology.

(B) to satisfy the requirement for 500 hours of supervised work experience, training shall be under the supervision of an authorized user at a medical institution and shall include:

(a) ordering, receiving, and unpacking radioactive materials safely and performing the related radiation surveys;

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(b) calibrating dose calibrators and diagnostic instruments and performing checks for proper operation of survey meters;

(c) calculating and safely preparing patient dosages;

(d) using administrative controls to prevent the misadministration of radioactive material;

(e) using emergency procedures to contain spilled radioactive material safely and using proper decontamination procedures; and

(f) eluting technetium-99m from generator systems, assaying and testing the eluate for molybdenum-99 and alumina contamination, and processing the eluate with reagent kits to prepare technetium-99m labeled radiopharmaceuticals or radiobiologics.

(C) to satisfy the requirement for 500 hours of supervised clinical experience, training shall be under the supervision of an authorized user at a medical institution and shall include:

(a) examining patients and reviewing their case histories to determine their suitability for radioisotope diagnosis, limitations, or contraindications;

(b) selecting the suitable radiopharmaceuticals or radiobiologics and calculating and measuring the dosages;

(c) administering dosages to patients and using syringe radiation shields;

(d) collaborating with the authorized user in the interpretation of radioisotope test results; and

(e) patient followup; or

(iii) has successfully completed a 6 month training program in nuclear medicine that has been approved by the Accreditation Council for Graduate Medical Education and that included classroom and laboratory training, work experience, and supervised clinical experience in all the topics identified in § 175.103(j)(d)(ii).

(5) Training for therapeutic use of radiopharmaceuticals or radiobiologics. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user of a radioactive material listed in § 175.103(f)(1)(i) for therapy to be a physician who:

(i) is certified by the:

(A) American Board of Nuclear Medicine; or

(B) American Board of Radiology in radiology, therapeutic radiology, or radiation

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oncology; or

(ii) has completed 80 hours of instruction in basic radioisotope handling techniques applicable to the use of therapeutic radiopharmaceuticals, and has had supervised clinical experience:

(A) to satisfy the requirement for instruction, 80 hours of classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity; and

(d) radiation biology;

(B) to satisfy the requirement for supervised clinical experience, training shall be under the supervision of an authorized user at a medical institution and shall include:

(a) use of iodine-131 for diagnosis of thyroid function and the treatment of hyperthyroidism or cardiac dysfunction in ten individuals;

(b) use of strontium-89 as strontium chloride for the treatment of bone pain in patients with painful bone metastases in three individuals; and

(c) use of soluble phosphorus-32 for the treatment of ascites polycythemia vera, leukemia, or bone metastases in three individuals; and

(d) use of iodine-131 for treatment of thyroid carcinoma in three individuals; and

(e) use of colloidal chromic phosphorus-32 or of colloidal gold-198 for intracavitary treatment of malignant effusions in three individuals.

(6) Training for therapeutic use of brachytherapy sources. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user using a brachytherapy source specified in § 175.103(h)(1) for therapy to be a physician who:

(i) is certified in:

(A) Radiology, therapeutic radiology, or radiation oncology by the American Board of Radiology; or

(B) Radiation oncology by the American Osteopathic Board of Radiology; or

(C) Radiology, with a specialization in radiotherapy, as a British "Fellow of the Faculty of Radiology" or "Fellow of the Royal College of Radiology"; or

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(D) Therapeutic radiology by the Canadian Royal College of Physicians and Surgeons; or

(ii) is in the active practice of therapeutic radiology, has completed 200 hours of instruction in basic radioisotope handling techniques applicable to the therapeutic use of brachytherapy sources and 500 hours of supervised work experience and a minimum of 3 years of supervised clinical experience:

(A) to satisfy the requirement for instruction, 200 hours of classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity; and

(d) radiation biology.

(B) to satisfy the requirement for 500 hours of supervised work experience, training shall be under the supervision of an authorized user at an institution and shall include:

(a) ordering, receiving, and unpacking radioactive materials safely and performing the related radiation surveys;

(b) checking survey meters for proper operation;

(c) preparing, implanting, and removing sealed sources;

(d) maintaining running inventories of material on hand;

(e) using administrative controls to prevent the misadministration of radioactive material; and

(f) using emergency procedures to control radioactive material.

(C) to satisfy the requirement for a period of supervised clinical experience, training shall include 1 year in a formal training program approved by the Residency Review Committee for Radiology of the Accreditation Council for Graduate Medical Education or the Committee on Postdoctoral Training of the American Osteopathic Association, and an additional 2 years of clinical experience in therapeutic radiology under the supervision of an authorized user at a medical institution. The supervised clinical experience shall include:

(a) examining individuals and reviewing their case histories to determine their suitability for brachytherapy treatment, and any limitations or contraindications;

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(b) selecting the proper brachytherapy sources and dose and method of administration;

(c) calculating the dose; and

(d) post-administration followup and review of case histories in collaboration with the authorized user.

(7) Training for ophthalmic use of strontium-90. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user using only strontium-90 for ophthalmic radiotherapy to be a physician who:

(i) Is certified in radiology, therapeutic radiology, or radiation oncology by the American Board of Radiology; or

(ii) Is in the active practice of therapeutic radiology or ophthalmology, and has completed 24 hours of instruction in basic radioisotope handling techniques applicable to the use of strontium-90 for ophthalmic radiotherapy, and a period of supervised clinical training in ophthalmic radiotherapy:

(A) to satisfy the requirement for instruction, the classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity; and

(d) radiation biology.

(B) to satisfy the requirement for a period of supervised clinical training in ophthalmic radiotherapy, training must be under the supervision of an authorized user at a medical institution and must include the use of strontium-90 for the ophthalmic treatment of five individuals that includes:

(a) examination of each individual to be treated;

(b) calculation of the dose to be administered;

(c) administration of the dose; and

(d) followup and review of each individual's case history.

(8) Training for use of sealed sources for diagnosis. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user using a sealed source in a device specified in § 175.103(g)(1) to be a physician, dentist, or podiatrist who:

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(i) is certified in:

(A) radiology, diagnostic radiology, therapeutic radiology, or radiation oncology by the American Board of Radiology; or

(B) nuclear medicine by the American Board of Nuclear Medicine; or

(C) diagnostic radiology or radiology by the American Osteopathic Board of Radiology; or

(ii) Has completed 8 hours of instruction in basic radioisotope handling techniques specifically applicable to the use of the device. To satisfy the requirement for instruction, the training shall include:

(A) radiation physics, mathematics pertaining to the use and measurement of radioactivity, and instrumentation;

(B) radiation biology; and

(C) radiation protection and training in the use of the device for the purposes authorized by the license.

(9) Training for teletherapy. Except as provided in § 175.103(j)(11), the licensee shall require the authorized user of a sealed source specified in § 175.103(i)(1) in a teletherapy unit to be a physician who:

(i) is certified in:

(A) radiology, therapeutic radiology or radiation oncology by the American Board of Radiology; or

(B) radiation oncology by the American Osteopathic Board of Radiology; or

(C) radiology, with specialization in radiotherapy, as a British "Fellow of the Faculty of Radiology" or "Fellow of the Royal College of Radiology"; or

(D) therapeutic radiology by the Canadian Royal College of Physicians and Surgeons; or

(ii) is in the active practice of therapeutic radiology, and has completed 200 hours of instruction in basic radioisotope techniques applicable to the use of a sealed source in a teletherapy unit, 500 hours of supervised work experience, and a minimum of 3 years of supervised clinical experience:

(A) to satisfy the requirement for instruction, the classroom and laboratory training shall include:

(a) radiation physics and instrumentation;

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(b) radiation protection;

(c) mathematics pertaining to the use and measurement of radioactivity; and

(d) radiation biology.

(B) to satisfy the requirement for supervised work experience, training shall be under the supervision of an authorized user at an institution and shall include:

(a) review of the full calibration measurements and periodic spot checks;

(b) preparing treatment plans and calculating treatment times;

(c) using administrative controls to prevent misadministrations;

(d) implementing emergency procedures to be followed in the event of the abnormal operation of a teletherapy unit or console; and

(e) checking and using survey meters.

(C) to satisfy the requirement for a period of supervised clinical experience, training shall include 1 year in a formal training program approved by the Residency Review Committee for Radiology of the Accreditation Council for Graduate Medical Education or the Committee on Postdoctoral Training of the American Osteopathic Association and an additional 2 years of clinical experience in therapeutic radiology under the supervision of an authorized user at a medical institution. The supervised clinical experience shall include:

(a) examining individuals and reviewing their case histories to determine their suitability for teletherapy treatment, and any limitations or contraindications;

(b) selecting the proper dose and how it is to be administered;

(c) calculating the teletherapy doses and collaborating with the authorized user in the review of patients' progress and consideration of the need to modify originally prescribed doses as warranted by patients' reaction to radiation; and

(d) post-administration followup and review of case histories.

(10) Training for radiation therapy physicist. The licensee shall require the radiation therapy physicist to:

(i) be certified by the American Board of Radiology or the American Board of Medical Physics in a branch of medical physics which deals with the therapeutic application of roentgen rays, gamma rays, electrons or other charged particles beams, neutrons and radiation from sealed radionuclide sources and the equipment associated with the production and use of these radiations; or

(ii) hold a master's or doctor's degree in physics, biophysics, radiological

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physics, or health physics, and have completed 1 year of full time training in therapeutic radiological physics and an additional year of full time work experience under the supervision of a radiation therapy physicist at a medical institution. To meet this requirement, the individual shall have performed the tasks listed in § 175.103(c)(5) and (i)(10), (11) and (12) under the supervision of a radiation therapy physicist during the year of work experience.

However, as of January 1, 2000 only individuals certified as in § 175.103(j)(10)(i) shall be named as radiation therapy physicists.

(11) Training for experienced authorized users. Practitioners of the healing arts identified as authorized users for the human use of radioactive material on a Department, agreement state, or U.S. Nuclear Regulatory Commission license issued before August 15, 1990 who perform only those methods of use for which they were authorized on that date need not comply with the training requirements of § 175.103(j)(1) through (9).

(12) Recentness of training. The training and experience specified in § 175.103(j)(1) through (10) must have been obtained within the 5 years preceding the date of application or the individual shall have had continuing applicable experience since the required training and experience was completed.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

<General Materials (GM) - References, Annotations, or Tables>

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.104. Waste disposal.

(a) General requirements. (1) A licensee shall dispose of licensed material only:

(i) by transfer to an authorized recipient as provided in § 175.101 or § 175.104(f) of this Code, or to the U.S. Department of Energy; or

(ii) by decay in storage; or

(iii) by release in effluents within the limits in § 175.03(d); or

(iv) as authorized pursuant to § 175.104(b), (c), (d) or (e).

(2) A person shall be specifically licensed to receive waste containing licensed material from other persons for:

(i) treatment prior to disposal; or

(ii) treatment or disposal by incineration; or

(iii) decay in storage; or

(iv) disposal at a land disposal facility licensed pursuant to 10 CFR Part 61 or the equivalent regulations of an agreement state; or

(v) storage until transferred to a storage or disposal facility authorized to receive the waste.

(3) A licensee or applicant for a license shall obtain any permits required by the New York State Department of Environmental Conservation pursuant to 6 NYCRR Part 380, or any successor law or regulation.

(4) A licensee or applicant for a license shall develop, document and implement a discharge minimization program required by the New York State Department of Envi-

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ronmental Conservation pursuant to 6 NYCRR Section 380-7, or any successor law or regulation.

(b) Method for obtaining approval of proposed disposal procedures. (1) A licensee or applicant for a license may apply to the Department for approval of proposed disposal procedures, not otherwise authorized in this Code, but which will conform to state and federal regulations, to dispose of licensed material generated in the licensee's operations. Each application shall include:

(i) a description of the waste containing licensed material to be disposed of, including the physical and chemical properties that have an impact on risk evaluation, and the proposed manner and conditions of waste disposal; and

(ii) an analysis and evaluation of pertinent information on the nature of the environment; and

(iii) the nature and location of other potentially affected facilities; and

(iv) analyses and procedures to ensure that doses are maintained ALARA and within the dose limits in § 175.03.

(c) Disposal by release into sanitary sewerage. (1) A licensee may discharge licensed material into sanitary sewerage if each of the following conditions is satisfied:

(i) the material is readily soluble in water or is biological material that is readily dispersible in water; and

(ii) the quantity of licensed radioactive material that the licensee releases into the sewer in 1 month divided by the average monthly volume of water released into the sewer by the licensee does not exceed the concentration listed in Table 3 of Appendix B of § 175.03; and

(iii) if more than one radionuclide is released, the following conditions must also be satisfied:

(A) the licensee shall determine the fraction of the limit in Table 3 of Appendix B of § 175.03 represented by discharges into sanitary sewerage by dividing the actual monthly average concentration of each radionuclide released by the licensee into the sewer by the concentration of that radionuclide listed in Table 3 of Appendix B of § 175.03; and

(B) the sum of the fractions for each radionuclide required by § 175.104(c)(1)(iii)(A) does not exceed unity; and

(iv) the total quantity of licensed radioactive material that the licensee releases into the sanitary sewerage in a year does not exceed 37 GBq (1 Ci) of all radioactive materials combined.

(2) Excreta from individuals undergoing medical diagnosis or therapy with ra-

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dioactive material are not subject to the limitations contained in § 175.104(c)(1).

(d) Treatment or disposal by incineration or burial. (1) No person shall treat or dispose of licensed radioactive material by incineration except as specifically approved by the Department pursuant to § 175.104(b).

(2) No person shall bury any licensed radioactive materials within this City.

(e) Disposal of specific wastes. (1) A licensee may ship for disposal outside of this City the following licensed material as if it were not radioactive, provided however, that the receiving jurisdiction regulates such materials as if they were not radioactive:

(i) 1.85 kBq (0.05 <<mu>>Ci), or less, of hydrogen-3 or carbon-14 per gram of medium used for liquid scintillation counting; and

(ii) 1.85 kBq (0.05 <<mu>>Ci), or less, of hydrogen-3 or carbon-14 per gram of animal tissue, averaged over the weight of the entire animal.

(2) A licensee shall not dispose of tissue pursuant to § 175.104(e)(1)(ii) in a manner that would permit its use either as food for humans or as animal feed.

(3) The licensee shall maintain records in accordance with § 175.03(k)(l0).

(f) Transfer for disposal and manifests. (1) The requirements of § 175.104(f) and Appendix A of § 175.104 are designed to control transfers of low-level radioactive waste intended for disposal at a licensed low-level radioactive waste disposal facility, establish a manifest tracking system, and supplement existing requirements concerning transfers and recordkeeping for those wastes.

(2) Each shipment of radioactive waste designated for disposal at a licensed low-level radioactive waste disposal facility shall be accompanied by a shipment manifest as specified in Section I of Appendix A of § 175.104.

(3) Each shipment manifest shall include a certification by the waste generator as specified in Section II of Appendix A of § 175.104.

(4) Each person involved in the transfer of waste for disposal or in the disposal of waste, including the waste generator, waste collector, waste processor, and disposal facility operator, shall comply with the requirements specified in Section III of Appendix A of § 175.104.

(5) The licensee or applicant for a license shall comply with the requirements of the New York State Department of Environmental Conservation as codified in 6 NYCRR Part 381, or any successor law or regulation.

(g) Compliance with environmental and health protection regulations.

(1) Nothing in this section relieves the licensee from complying with other applicable Federal, State and local regulations governing any other toxic or hazard-

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ous properties of materials that may be disposed of pursuant to this section.

(h) Records of waste disposal. (1) The licensee shall maintain records of the disposal of licensed materials made under § 175.104(b), (c), (d), (e) and 10 CFR Part 61 or the equivalent regulations of an agreement state.

(2) The licensee shall retain the records required by § 175.104(h)(1) until the Department authorizes disposition.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

APPENDIX A REQUIREMENTS FOR TRANSFER OF LOW-LEVEL RADIOACTIVE WASTE FOR DISPOSAL AT LAND DISPOSAL FACILITIES AND MANIFESTS I. Manifest. The shipment manifest shall contain the name, address, and telephone number of the person generating the waste. The manifest shall also include the name, address, and telephone number or the name and U.S. Environmental Protection Agency hazardous waste identification number of the person transporting the waste to the land disposal facility. The manifest shall also indicate: a physical description of the waste, the volume, radionuclide identity and quantity, the total radioactivity, and the principal chemical form. The solidification agent shall be specified. Waste containing more than 0.1% chelating agents by weight shall be identified and the weight percentage of the chelating agent estimated. Wastes classified as Class A, Class B, or Class C in Section I of Appendix B shall be clearly identified as such in the manifest. The total quantity of the radionuclides hydrogen-3, carbon-14, technetium-99, and iodine-129 shall be shown. The manifest required by this paragraph may be shipping papers used to meet U.S. Department of Transportation or U.S. Environmental Protection Agency regulations or requirements of the receiver, provided all the required information is included. Copies of manifests required by this section may be legible carbon copies or legible photocopies.II. Certification. The waste generator shall include in the shipment manifest a certification that the transported materials are properly classified, described, packaged, marked, and labeled and are in proper condition for transportation according to the applicable regulations of the U.S. Department of Transportation, the New York State Department of Environmental Protection and this Department. An authorized representative of the waste generator shall sign and date the manifest.III. Control and Tracking. (a) Any radioactive waste generator who transfers radioactive waste to a land disposal facility or a licensed waste collector shall comply with the requirements in III. (a)(1) through (8). Any radioactive waste generator who transfers waste to a licensed waste processor who treats or repackages waste shall comply with the requirements of III. (a)(4) through (8). A licensee shall:(1) Prepare all wastes so that the waste is classified according to Section I of Appendix B and meets the waste characteristics requirements in Section II of Appendix B;(2) Label each package of waste to identify whether it is Class A waste, Class B waste, or Class C waste, in accordance with Section I of Appendix B;(3) Conduct a quality control program to ensure compliance with Section I and II of Appendix B; the program shall include management evaluation of audits;(4) Prepare shipping manifests to meet the requirements of Section I and II;(5) Forward a copy of the manifest to the intended recipient, at the time of shipment, or deliver

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to a collector at the time the waste is collected, obtaining acknowledgment of receipt in the form of a signed copy of the manifest or equivalent documentation from the collector; (6) Include one copy of the manifest with the shipment;(7) Retain a copy of the manifest and documentation of acknowledgment of receipt as the record of transfer of licensed material as required by § 175.101 of this Code; and(8) For any shipments or any portion of a shipment for which acknowledgment of receipt has not been received within the times set forth in this section, conduct an investigation in accordance with Section III. (e).(b) Any waste collector licensee who handles only prepackaged waste shall:(1) Acknowledge receipt of the waste from the generator within 1 week of receipt by returning a signed copy of the manifest or equivalent documentation;(2) Prepare a new manifest to reflect consolidated shipments; the new manifest shall serve as a listing or index for the detailed generator manifests. Copies of the generator manifests shall be a part of the new manifest. The waste collector may prepare a new manifest without attaching the generator manifests, provided the new manifest contains for each package the information specified in Section I of this appendix. The collector licensee shall certify that nothing has been done to the waste that would invalidate the generator's certification;(3) Forward a copy of the new manifest to the land disposal facility operator at the time of shipment;(4) Include the new manifest with the shipment to the disposal site; (5) Retain a copy of the manifest and documentation of acknowledgment of receipt as the record of transfer of licensed material as required by § 175.101 of this Code, and retain information from generator manifest until disposition is authorized by the Department; and(6) For any shipments or any portion of a shipment for which acknowledgment of receipt is not received within the times set forth in this section, conduct an investigation in accordance with Section III. (e).(c) Any licensed waste processor who treats or repackages wastes shall:(1) Acknowledge receipt of the waste from the generator within 1 week of receipt by returning a signed copy of the manifest or equivalent documentation; (2) Prepare a new manifest that meets the requirements of Sections I and II of this appendix. Preparation of the new manifest reflects that the processor is responsible for the waste;(3) Prepare all wastes so that the waste is classified according to Section I of Appendix B and meets the waste characteristics requirements in Section II of Appendix B;(4) Label each package of waste to identify whether it is Class A waste, Class B waste, or Class C waste, in accordance with Sections I and III of Appendix B;(5) Conduct a quality control program to ensure compliance with Sections I and II of Appendix B. The program shall include management evaluation of audits;(6) Forward a copy of the new manifest to the disposal site operator or waste collector at the time of shipment, or deliver to a collector at the time the waste is collected, obtaining acknowledgment of receipt in the form of a signed copy of the manifest or equivalent documentation by the collector;(7) Include the new manifest with the shipment;(8) Retain copies of original manifests and new manifests and documentation of acknowledgment of receipt as the record of transfer of licensed material required by § 175.101 of this Code; and(9) For any shipment or portion of a shipment for which acknowledgment is not received within the times set forth in this section, conduct an investigation in accordance with Section III. (e).(d) The land disposal facility operator shall: (1) Acknowledge receipt of the waste within 1 week of receipt by returning a signed copy of the manifest or equivalent documentation to the shipper. The shipper to be notified is the licensee who last possessed the waste and transferred the waste to the operator. The returned copy of the manifest or equivalent documentation shall indicate any discrepancies between materials listed on the manifest and materials received;(2) Maintain copies of all completed manifests or equivalent documentation until the Agency authorizes their disposition; and(3) Notify the shipper, that is, the generator, the collector, or processor, and the

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Agency when any shipment or portion of a shipment has not arrived within 60 days after the advance manifest was received.(e) Any shipment or portion of a shipment for which acknowledgment is not received within the times set forth in this section shall:(1) Be investigated by the shipper if the shipper has not received notification or receipt within 20 days after transfer.(i) Such investigation shall include tracing the shipment and filing a report with the Department. Each licensee who conducts a trace investigation shall file a written report with the Department within 2 weeks of completion of the investigation.

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NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

APPENDIX B CLASSIFICATION AND CHARACTERISTICS OF LOW-LEVEL RADIOACTIVE WASTE

I. Classification of Radioactive Waste for Land Disposal. (a) Considerations. Determination of the classification of radioactive waste involves two considerations. First, consideration must be given to the concentration of long-lived radionuclides (and their shorter-lived precursors) whose potential hazard will persist long after such precautions as institutional controls, improved waste form, and deeper disposal have ceased to be effective. These precautions delay the time when long-lived radionuclides could cause exposures. In addition, the magnitude of the potential dose is limited by the concentration and availability of the radionuclide at the time of exposure. Second, consideration must be given to the concentration of shorter-lived radionuclides for which requirements on institutional controls, waste form, and disposal methods are effective.

(b) Classes of waste. (1) Class A waste is waste that is usually segregated from other waste classes at the disposal site. The physical form and characteristics of Class A waste must meet the minimum requirements set forth in Section II. (a). If Class A waste also meets the stability requirements set forth in Section II. (b), it is not necessary to segregate the waste for disposal.

(2) Class B waste is waste that must meet more rigorous requirements on waste form to ensure stability after disposal. The physical form and characteristics of Class B waste must meet both the minimum and stability requirements set forth in Section II.

(3) Class C waste is waste that not only must meet more rigorous requirements on waste form to ensure stability but also requires additional measures at the disposal facility to protect against inadvertent intrusion. The physical form and characteristics of Class C waste must meet both the minimum and stability requirements set forth in Section II.

(c) Classification determined by long-lived radionuclides. If the radioactive waste contains only radionuclides listed in Table I, classification shall be determined as follows:

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(1) If the concentration does not exceed 0.1 times the value in Table I, the waste is Class A.

(2) If the concentration exceeds 0.1 times the value in Table I, but does not exceed the value in Table I, the waste is Class C.

(3) If the concentration exceeds the value in Table I, the waste is not generally acceptable for land disposal.

(4) For wastes containing mixtures of radionuclides listed in Table I, the total concentration shall be determined by the sum of fractions rule described in Section I. (g).

Table I

Radionuclide Concentration curie/cubic metera nanocurie/gramb

Table I (cont'd)

Radionuclide	Concentration	curie/cubic metera	nanocurie/gramb
C-14	8		
C-14 in activated metal	80		
Ni-59 in activated metal	220		
Nb-94 in activated metal	0.2		
Tc-99	3		
I-129	0.08		
Alpha emitting transuranic radionuclides with half-life greater than five years		100	
Pu-241		3,500	
Cm-242		20,000	
Ra-226		100	

a To convert the Ci-m-3 values to gigabecquerels (GBq) per cubic meter, multiply the Ci-m-3 by 37.

b To convert the nCi/g values to becquerel (Bq) per gram, multiply the nCi/g value by 37.

(d) Classification determined by short-lived radionuclides. If the waste does not contain any of the radionuclides listed in Table I, classification shall be determined based on the concentrations shown in Table II. However, as specified in Section I. (f), if radioactive waste does not contain any nuclides listed in either Table I or II, it is Class A.

(1) If the concentration does not exceed the value in Column 1, the waste is Class A.

(2) If the concentration exceeds the value in Column 1 but does not exceed the

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value in Column 2, the waste is Class B.

(3) If the concentration exceeds the value in Column 2 but does not exceed the value in Column 3, the waste is Class C.

(4) If the concentration exceeds the value in Column 3, the waste is not generally acceptable for near-surface disposal.

(5) For wastes containing mixtures of the radionuclides listed in Table II, the total concentration shall be determined by the sum of fractions rule described in Section I. (g).

Table II

Radionuclide	Concentration, curie/cubic meter*	Column 1	Column 2	Column 3
Total of all radionuclides with less than 5-year half-life	700	*	*	
H-3	40	*	*	
Co-60	700	*	*	
Ni-63	3.5	70	700	
Ni-63 in activated metal	35	700	7000	
Sr-90	0.04	150	7000	
Cs-137	1	44	4600	

* To convert the Ci/m 3 value to gigabecquerel (GBq) per cubic meter, multiply the Ci/m 3 value by 37. There are no limits established for these radionuclides in Class B or C wastes. Practical considerations such as the effects of external radiation and internal heat generation on transportation, handling, and disposal will limit the concentrations for these wastes. These wastes shall be Class B unless the concentrations of other radionuclides in Table II determine the waste to be Class C independent of these radionuclides.

(e) Classification determined by both long- and short-lived radionuclides. If the radioactive waste contains a mixture of radionuclides, some of which are listed in Table I and some of which are listed in Table II, classification shall be determined as follows:

(1) If the concentration of a radionuclide listed in Table I is less than 0.1 times the value listed in Table I, the class shall be that determined by the concentration of radionuclides listed in Table II.

(2) If the concentration of a radionuclide listed in Table I exceeds 0.1 times the value listed in Table I, but does not exceed the value in Table 1, the waste shall be Class C, provided the concentration of radionuclides listed in Table II does not exceed the value shown in Column 3 of Table II.

(f) Classification of wastes with radionuclides other than those listed in Tables I and II. If the waste does not contain any radionuclides listed in either Table I or II, it is Class A.

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(g) The sum of the fractions rule for mixtures of radionuclides. For determining classification for waste that contains a mixture of radionuclides, it is necessary to determine the sum of fractions by dividing each radionuclide's concentration by the appropriate limit and adding the resulting values. The appropriate limits must all be taken from the same column of the same table. The sum of the fractions for the column must be less than 1.0 if the waste class is to be determined by that column. Example: A waste contains Sr-90 in a concentration of 1.85 TBq/m 3 (50 Ci/m 3) and Cs-137 in a concentration of 814 GBq/m 3 (22 Ci/m 3). Since the concentrations both exceed the values in Column 1, Table II, they must be compared to Column 2 values. For Sr-90 fraction, 50/150 = 0.33., for Cs-137 fraction, 22/44 = 0.5; the sum of the fractions = 0.83. Since the sum is less than 1.0, the waste is Class B.

(h) Determination of concentrations in wastes. The concentration of a radionuclide may be determined by indirect methods such as use of scaling factors which relate the inferred concentration of one radionuclide to another that is measured, or radionuclide material accountability, if there is reasonable assurance that the indirect methods can be correlated with actual measurements. The concentration of a radionuclide may be averaged over the volume of the waste, or weight of the waste if the units are expressed as becquerel (nanocurie) per gram.

II. Radioactive Waste Characteristics. (a) The following are minimum requirements for all classes of waste and are intended to facilitate handling and provide protection of health and safety of personnel at the disposal site.

(1) Wastes shall be packaged in conformance with the conditions of the license issued to the site operator to which the waste will be shipped. Where the conditions of the site license are more restrictive than the provisions of Part D, the site license conditions shall govern.

(2) Wastes shall not be packaged for disposal in cardboard or fiberboard boxes.

(3) Liquid waste shall be packaged in sufficient absorbent material to absorb twice the volume of the liquid.

(4) Solid waste containing liquid shall contain as little free-standing and noncorrosive liquid as is reasonably achievable, but in no case shall the liquid exceed 1% of the volume.

(5) Waste shall not be readily capable of detonation or of explosive decomposition or reaction at normal pressures and temperatures, or of explosive reaction with water.

(6) Waste shall not contain, or be capable of generating, quantities of toxic gases, vapors, or fumes harmful to persons transporting, handling, or disposing of the waste. This does not apply to radioactive gaseous waste packaged in accordance with Section II. (a)(8).

(7) Waste must not be pyrophoric. Pyrophoric materials contained in wastes shall be treated, prepared, and packaged to be nonflammable. [FN8]

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[FN8]. See § 175.02(a)(162) of this Code for a definition of "pyrophoric."

(8) Wastes in a gaseous form shall be packaged at an absolute pressure that does not exceed 1.5 atmospheres at 20 degreeC. Total activity shall not exceed 3.7 TBq (100 Ci) per container.

(9) Wastes containing hazardous, biological, pathogenic, or infectious material shall be treated to reduce to the maximum extent practicable the potential hazard from the non-radiological materials.

(b) The following requirements are intended to provide stability of the waste. Stability is intended to ensure that the waste does not degrade and affect overall stability of the site through slumping, collapse, or other failure of the disposal unit and thereby lead to water infiltration. Stability is also a factor in limiting exposure to an inadvertent intruder, since it provides a recognizable and nondispersible waste.

(1) Waste shall have structural stability. A structurally stable waste form will generally maintain its physical dimensions and its form, under the expected disposal conditions such as weight of overburden and compaction equipment, the presence of moisture, and microbial activity, and internal factors such as radiation effects and chemical changes. Structural stability can be provided by the waste form itself, processing the waste to a stable form, or placing the waste in a disposal container or structure that provides stability after disposal.

(2) Notwithstanding the provisions in Sections II. (a)(3) and (4), liquid wastes, or wastes containing liquid, shall be converted into a form that contains as little free-standing and non-corrosive liquid as is reasonably achievable, but in no case shall the liquid exceed 1% of the volume of the waste when the waste is in a disposal container designed to ensure stability, or 0.5% of the volume of the waste for waste processed to a stable form.

(3) Void spaces within the waste and between the waste and its package shall be reduced to the extent practicable.

III. Labeling. Each package of waste shall be clearly labeled to identify whether it is Class A, Class B, or Class C waste, in accordance with Section I.

<General Materials (GM) - References, Annotations, or Tables>

24 RCNY Hlth. Code § 175.104
END OF DOCUMENT

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.105. Transportation and Packaging of Radioactive Materials.

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Appendices

Appendix A--Determination of A1 and A2

Table A-1--A1 and A2 Values for Radionuclides

Table A-2--General Values for A1 and A2

(a) General Provisions. (1) Purpose and Scope.

(i) This section establishes requirements for packaging, preparation for shipment, and transportation of licensed material. The packaging and transport of licensed material are also subject to other sections of this Code (e.g., §§ 175.03, 175.101) and to the regulations of other agencies (e.g., the U.S. Nuclear Regulatory Commission (NRC), the U.S. Department of Transportation (USDOT) and the U.S. Postal Service*) [FN13] having jurisdiction over means of transport and other applicable state and local laws and regulations. The requirements of this section are in addition to, and not in substitution for, other requirements.

[FN13]. Postal Service Manual (Domestic Mail Manual), § 124.3, which is incorporated by reference at 39 CFR 111.1.

(ii) This section applies to any licensee authorized by specific or general license issued by the Department to receive, possess, use, or transfer licensed material, if the licensee delivers that material to a carrier for transport, transports the material outside the site of usage as specified in the Department license, or transports that material on public highways. No provision of this section authorizes possession of licensed material.

(iii) The requirements of this section apply to any person who has a license or who is required to obtain a license pursuant to this Code, if the person delivers radioactive material to a common or contract carrier for transport or transports the material outside the confines of the person's facility or other authorized place of use.

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(2) Records. Each record required by this section must be legible throughout the retention period specified by this Code. The record may be the original or a reproduced copy or a microform provided that the copy or microform is authenticated by authorized personnel and that the microform is capable of producing a clear copy throughout the required retention period. The record may also be stored in electronic media with the capability for producing legible, accurate, and complete records during the required retention period. Records such as letters, drawings, specifications, must include all pertinent information such as stamps, initials, and signatures. The licensee shall maintain adequate safeguards against tampering with and loss of records.

(3) Completeness and accuracy of information.

(i) Information provided to the Department by an applicant for a license, or by a licensee, or information required by applicable laws or regulations, or licensed conditions to be maintained by the applicant or the licensee must be complete and accurate in all material respects.

(ii) Each applicant or licensee shall notify the Department of information identified by the applicant or licensee as having, for the regulated activity, a significant implication for public health and safety or common defense and security. An applicant or licensee violates this requirement if the applicant or licensee fails to notify the Department of information that the applicant or licensee has identified as having a significant implication for public health and safety or common defense and security. Notification must be provided to the Department within two working days of identifying the information. This requirement is not applicable to information that is already required to be provided to the Department by other reporting or updating requirements.

(4) Requirement for license.

(i) Except as authorized in a general license or a specific license issued by the Department, or as exempted in this section, no person may--

(A) Deliver licensed material to a carrier for transport; or

(B) Transport licensed material.

(ii) Exemptions from the requirement for license in § 175.105(a)(4) are specified in § 175.105(b)(2). General licenses for which no NRC package approval is required are issued in §§ 175.105(c)(3) and 175.105(c)(4). The general license in § 175.105(c)(1) requires that an NRC certificate of compliance or other package approval be issued for the package to be used under the general license. The transport of licensed material or delivery of licensed material to a carrier for transport is subject to the operating controls and procedures requirements of § 175.105(d), to the quality assurance requirements of § 175.105(e), and to the general provisions of § 175.105(a), including USDOT regulations referenced in § 175.105(a)(6).

(5) Definitions. The following terms are defined herein for the purpose of this section. These definitions are in addition to those in § 175.02. To ensure compati-

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bility with international transportation standards, all limits in this section are given in terms of dual units: The International System of Units (SI) followed or preceded by U.S. standard or customary units. The U.S. customary units are not exact equivalents, but are rounded to a convenient value, providing a functionally equivalent unit. For the purpose of this section, either unit may be used.

(i) "Certificate holder" means a person who has been issued a certificate of compliance or other package approval by the U.S. Nuclear Regulatory Commission.

(ii)

"Certificate of Compliance (CoC)" means the certificate issued by the NRC which approves the design of a package for the transportation of radioactive material.

(iii) "Close reflection by water" means immediate contact by water of sufficient thickness for maximum reflection of neutrons.

(iv) "Containment system" means the assembly of components of the packaging intended to retain the radioactive material during transport.

(v) "Criticality Safety Index (CSI)"' means the dimensionless number (rounded up to the next tenth) assigned to and placed on the label of a fissile material package, to designate the degree of control of accumulation of packages containing fissile material during transportation. Determination of the criticality safety index is described in 10 CFR §§ 71.22, 71.23, and 71.59.

(vi) "Deuterium" means, for the purposes of 10 CFR §§ 71.15 and 71.22, deuterium and any deuterium compounds, including heavy water, in which the ratio of deuterium atoms to hydrogen atoms exceeds 1:5000.

(vii) "Graphite" means, for the purposes of 10 CFR §§ 71.15 and 71.22, graphite with a boron equivalent content less than 5 parts per million and density greater than 1.5 grams per cubic centimeter.

(viii) "Low toxicity alpha emitters" means natural uranium, depleted uranium, natural thorium; uranium-235, uranium-238, thorium-232, thorium-228 or thorium-230 when contained in ores or physical or chemical concentrates or tailings; or alpha emitters with a half-life of less than 10 days.

(ix) "Maximum normal operating pressure" means the maximum gauge pressure that would develop in the containment system in a period of 1 year under the heat condition specified in 10 CFR 71.71(c)(1), in the absence of venting, external cooling by an ancillary system, or operational controls during transport.

(x) "Natural thorium" means thorium with the naturally occurring distribution of thorium isotopes (essentially 100 weight percent thorium-232).

(xi) "Optimum interspersed hydrogenous moderation" means the presence of hydrogenous material between packages to such an extent that the maximum nuclear reactivity results.

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(xii) "Spent nuclear fuel"' means fuel that has been withdrawn from a nuclear reactor following irradiation, has undergone at least 1 year's decay since being used as a source of energy in a power reactor, and has not been chemically separated into its constituent elements by reprocessing. Spent fuel includes the special nuclear material, byproduct material, source material, and other radioactive materials associated with fuel assemblies.

(xiii) "Surface Contaminated Object (SCO)" means a solid object that is not itself classed as radioactive material, but which has radioactive material distributed on any of its surfaces. SCO must be in one of two groups with surface activity not exceeding the following limits:

(A) SCO-1: A solid object on which:

(a) The non-fixed contamination on the accessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 4 Bq/cm2 (10-4 microcurie/cm2) for beta and gamma and low toxicity alpha emitters, or 0.4 Bq/cm2 (10-5 microcurie/cm2) for all other alpha emitters;

(b) The fixed contamination on the accessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 4X104 Bq/cm2 (1.0 microcurie/cm2) for beta and gamma and low toxicity alpha emitters, or 4X103 Bq/cm2 (0.1 microcurie/cm2) for all other alpha emitters; and

(c) The non-fixed contamination plus the fixed contamination on the inaccessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 4X104 Bq/cm2 (1 microcurie/cm2) for beta and gamma and low toxicity alpha emitters, or 4X103 Bq/cm2 (0.1 microcurie/cm2) for all other alpha emitters.

(B) SCO-II: A solid object on which the limits for SCO-I are exceeded and on which:

(a) The non-fixed contamination on the accessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 400 Bq/cm2 (10-2 microcurie/cm2) for beta and gamma and low toxicity alpha emitters or 40 Bq/cm2 (10-3 microcurie/cm2) for all other alpha emitters;

(b) The fixed contamination on the accessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 8X105 Bq/cm2 (20 microcuries/cm2) for beta and gamma and low toxicity alpha emitters, or 8X104 Bq/cm2 (2 microcuries/cm2) for all other alpha emitters; and

(c) The non-fixed contamination plus the fixed contamination on the inaccessible surface averaged over 300 cm2 (or the area of the surface if less than 300 cm2) does not exceed 8X105 Bq/cm2 (20 microcuries/cm2) for beta and gamma and low toxicity alpha emitters, or 8X104 Bq/cm2 (2 microcuries/cm2) for all other alpha emitters.

(xiv) "Unirradiated uranium" means uranium containing not more than 2 x 103 Bq of

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plutonium per gram of uranium-235, not more than 9 x 106 Bq of fission products per gram of uranium-235, and not more than 5 x 10-3 g of uranium-236 per gram of uranium-235.

(xv) Uranium--natural, depleted, enriched

(A) "Natural uranium" means uranium with the naturally occurring distribution of uranium isotopes (approximately 0.711 weight percent uranium-235, and the remainder by weight essentially uranium-238).

(B) "Depleted uranium" means uranium containing less uranium-235 than the naturally occurring distribution of uranium isotopes.

(C) "Enriched uranium" means uranium containing more uranium-235 than the naturally occurring distribution of uranium isotopes.

(6) Transportation of licensed material.

(i)

Each licensee who transports licensed material outside the site of usage, as specified in the license or where transport is on public highways, or who delivers licensed material to a carrier for transport, shall comply with the applicable requirements of the USDOT regulations in 49 CFR Parts 107, 171 through 180 and 390 through 397 appropriate to the mode of transport.

(A) The licensee shall particularly note USDOT regulations in the following areas:

(a) Packaging--49 CFR Part 173; Subparts A and B and I.

(b) Marking and labeling--49 CFR Part 172: Subpart D; and Sections 172.400 through 172.407 and Sections 172.436 through 172.441 of Subpart E.

(c) Placarding--49 CFR Part 172: Subpart F, especially Sections 172.500 through 172.519, 172.556, and Appendices B and C.

(d) Accident reporting--49 CFR Part 171: Sections 171.15 and 171.16.

(e) Shipping papers and emergency information--49 CFR Part 172: Subparts C and G.

(f) Hazardous material employee training--49 CFR Part 172: Subpart H.

(g) Security plans--49 CFR Part 172: subpart I.

(h) Hazardous material shipper/carrier registration--49 CFR Part 107: Subpart G.

(B) The licensee shall also note USDOT regulations pertaining to the following modes of transportation:

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(a) Rail--49 CFR Part 174: Subparts A through D and K.

(b) Air--49 CFR Part 175.

(c) Vessel--49 CFR Part 176: Subparts A through F and M.

(d) Public Highway--49 CFR Part 177 and Parts 390 through 397.

(ii) If USDOT regulations are not applicable to a shipment of licensed material, the licensee shall conform to the standards and requirements of the USDOT specified in § 175.105(a)(6)(i) to the same extent as if the shipment or transportation were subject to USDOT regulations. A request for modification, waiver, or exemption from those requirements, and any notification referred to in those requirements, must be filed with, or made to, the Director, Office of Nuclear Material Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001.

(b) Exemption. (1) Exemption of physicians. Any physician licensed by a State to dispense drugs in the practice of medicine is exempt from § 175.105(a)(6) with respect to transport by the physician of licensed material for use in the practice of medicine. However, any physician operating under this exemption must be licensed under applicable sections of this Code, 10 CFR Part 35 or the equivalent Agreement State regulations. Such transport must not be by public modes of transportation including, but not limited to, buses, subways, trams, taxicabs, car services, trains, ferries, or other means which would be returned immediately to public use after transporting licensed material.

(2) Exemption for low-level materials.

(i)A licensee is exempt from all requirements of this section with respect to shipment or carriage of the following low-level materials:

(A) Natural material and ores containing naturally occurring radionuclides that are not intended to be processed for use of these radionuclides, provided the activity concentration of the material does not exceed 10 times the values specified in Appendix A, Table A-2 of this section.

(B) Materials for which the activity concentration is not greater than the activity concentration values specified in Appendix A, Table A-2 of this section , or for which the consignment activity is not greater than the limit for an exempt consignment found in Appendix A, Table A-2 of this section.

(3) Exemption from classification as fissile material.

(i) Fissile material meeting the requirements of at least one of the paragraphs of this section are exempt from classification as fissile material and from the fissile material package standards of 10 CFR §§ 71.55 and 71.59, but are subject to all other requirements of this part, except as noted.

(ii) Individual package containing 2 grams or less of fissile material

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(iii) Individual or bulk packaging containing 15 grams or less of fissile material provided the package has at least 200 grams of solid nonfissile material for every gram of fissile material. Lead, beryllium, graphite and hydrogenous material enriched in deuterium may be present in the package but must not be included in determining the required mass for solid nonfissile material.

(iv) Low concentrations of solid fissile material commingled with solid nonfissile material provided that:

(A) There is at least 2000 grams of solid nonfissile material for every gram of fissile material, and

(B) There is no more than 180 grams of fissile material distributed within 360 kg of contiguous nonfissile material.

(C) Lead, beryllium, graphite, and hydrogenous material may be present in the package but must not be included in determining the required mass of solid nonfissile material.

(v) Uranium enriched in uranium-235 to a maximum of 1 percent by weight, and with a total plutonium and uranium-233 content of up to 1 percent of the mass of the uranium235, provided that the mass of any beryllium, graphite, and hydrogenous material enriched in deuterium constitutes less than 5 percent of the uranium mass.

(vi) Liquid solutions of uranyl nitrate enriched in uranium-235 to a maximum of 2 percent by mass, with a total plutonium and uranium-233 content not exceeding 0.002 percent of the mass of uranium, and with a minimum nitrogen to uranium atomic ratio (N/U) of 2. The material must be contained in at least a DOT Type A package.

(vii) Packages containing, individually, a total plutonium mass of not more than 1000 grams, of which not more than 20 percent by mass may consist of plutonium-239, plutonium-241, or any combination of these radionuclides.

(c) General Licenses. (1) General license: NRC-approved package.

(i) A general license is hereby issued to any licensee of the Department to transport, or to deliver to a carrier for transport, licensed material in a package for which a license, certificate of compliance, or other approval has been issued by the NRC.

(ii) This general license applies only to a licensee who--

(A) Has a quality assurance program approved by the Department as satisfying the provisions of § 175.105(e) of this Code.

(B) Has a copy of the certificate of compliance, or other approval of the package, and has the drawings and other documents referenced in the approval relating to the use and maintenance of the packaging and to the actions to be taken before shipment;

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(C) Complies with the terms and conditions of the license, certificate, or other approval, as applicable, and applicable provisions of the operating controls and procedures requirements of § 175.105(d), the quality assurance requirements of § 175.105(e), and the general provisions of § 175.105(a); and

(D) Submits in writing to the Department, before the licensee's first use of the package, the licensee's name and license number and the package identification number specified in the package approval.

(iii) This general license applies only when the package approval authorizes use of the package under this general license.

(iv) For a Type B or fissile material package, the design of which was approved by NRC before April 1, 1996, the general license is subject to the additional restrictions contained in 10 CFR 71.13.

(2) Previously approved package.

(i) A Type B package previously approved by NRC but not designated as B(U) or B(M) in the identification number of the NRC Certificate of Compliance, may be used under the general license of § 175.105(c)(1) with the following additional conditions:

(A) Fabrication of the packaging was satisfactorily completed by August 31, 1986, as demonstrated by application of its model number in accordance with § 175.105(d)(2)(iii);

(B) A package used for a shipment to a location outside the United States is subject to multilateral approval, as defined in USDOT regulations at 49 CFR 173.403; and

(C) A serial number that uniquely identifies each packaging which conforms to the approved design is assigned to, and legibly and durably marked on, the outside of each packaging.

(ii) A Type B(U) package, a Type B(M) package, a low specific activity (LSA) material package or a fissile material package, previously approved by the NRC but without the designation "-85in. in the identification number of the NRC Certificate of Compliance, may be used under the general license of § 175.105(c)(1) with the following additional conditions:

(A) Fabrication of the package is satisfactorily completed by April 1, 1999 as demonstrated by application of its model number in accordance with § 175.105(d)(2)(iii) of this Code;

(B) A package used for a shipment to a location outside the United States is subject to multilateral approval as defined in USDOT regulations at 49 CFR 173.403; and

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(C) A serial number which uniquely identifies each packaging which conforms to the approved design is assigned to and legibly and durably marked on the outside of each packaging.

(3) General license: U.S. Department of Transportation specification container.

(i) A general license is issued to any licensee of the Department to transport, or to deliver to a carrier for transport, licensed material in a specification container for fissile material or for a Type B quantity of radioactive material as specified in USDOT regulations at 49 CFR Parts 173 and 178.

(ii) This general license applies only to a licensee who has a quality assurance program approved by the Department as satisfying the provisions of § 175.105(e) of this Code.

(iii) This general license applies only to a licensee who--

(A) Has a copy of the specification; and

(B) Complies with the terms and conditions of the specification and the applicable provisions of the operating and procedures requirements in § 175.105(d), the quality assurance requirements in § 175.105(e) and the general provisions contained in § 175.105(a).

(iv) This general license is subject to the limitation that the specification container may not be used for a shipment to a location outside the United States, except by multilateral approval, as defined in USDOT regulations at 49 CFR 173.403.

(v) The requirements of § 175.105(c)(3) shall expire October 1, 2008.

(4) General License: Use of foreign approved package.

(i) A general license is issued to any licensee of the Department to transport, or to deliver to a carrier for transport, licensed material in a package the design of which has been approved in a foreign national competent authority certificate that has been revalidated by USDOT as meeting the applicable requirements of 49 CFR 171.12.

(ii) Except as otherwise provided herein, the general license applies only to a licensee who has a quality assurance program approved by the Department as satisfying the applicable provisions of § 175.105(e) of this Code.

(iii) This general license applies only to shipments made to or from locations outside the United States.

(iv) This general license applies only to a licensee who--

(A) Has a copy of the applicable certificate, the revalidation, and the drawings and other documents referenced in the certificate, relating to the use and maintenance of the packaging and to the actions to be taken before shipment; and

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(B) Complies with the terms and conditions of the certificate and revalidation, and with the applicable provisions of the operating and procedures requirements in § 175.105(d), the quality assurance requirements in § 175.105(e) and the general provisions in § 175.105(a). With respect to the quality assurance provisions of § 175.105(e) of this Code, the licensee is exempt from design, construction, and fabrication considerations.

(5) General License: Fissile Material.

(i) A general license is issued to any licensee of the Department to transport fissile material, or to deliver fissile material to a carrier for transport, if the material is shipped in accordance with this section. The fissile material need not be contained in a package which meets the standards of-subparts E and F of 10 CFR 71.22; however the material must be contained in a Type A package. The Type A package must also meet the DOT requirements of 49 CFR 173.417(a).

(ii) The general license applies only to a licensee who has a quality assurance program approved by the Department as satisfying the provisions of § 175.105(e) of this part

(iii) The general license applies only when a package's contents:

(A) Contain less than a Type A quantity of fissile material; and

(B) Contains less than 500 total grams of beryllium, graphite, or hydrogenous material enriched in deuterium.

(iv) The general license applies only to packages containing fissile material that are labeled with a CSI which:

(A) Has been determined in accordance with section (5) of this section

(B) Has a value less than or equal to 10; and

(C) For a shipment of multiple packages containing fissile material, the sum of the CSIs must be less than or equal to 50 (for shipment on a nonexclusive use conveyance) and less than or equal to 100 (for shipment on an exclusive use conveyance).

(v) (A) The value for the CSI must be greater than or equal to the number calculated by the following equation:

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(B) The calculated CSI must be rounded up to the first decimal place;

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(C) The values of X, Y, and Z used in the CSI equation must be taken from Tables-71.1 or 71.2, as appropriate;

(D) If Table 71-2 is used to obtain the value of X, then the values of the terms in the equation for uranium-233 and plutonium must be assumed to be zero; and,

(E) Table 71-1 values for X, Y, and Z must be used to determine the CSI if:

(a) Uranium-233 is present in the package;

(b) The mass of plutonium exceeds 1 percent of the mass of uranium-235;

(c) The uranium is of unknown uranium-235 enrichment or greater than 24 weight percent enrichment; or

(d) Substances having a moderating effectiveness (i.e., an average hydrogen density greater than H2O) (e.g., certain hydrocarbon oils or plastics) are present in any form, except as polyethylene used for packing or wrapping.

(6) General license: Plutonium/Beryllium special form material.

(i) A general license is issued to any licensee of the Department to transport fissile material in the form of plutonium-beryllium (Pu-Be) special form sealed sources, or to deliver Pu-Be sealed sources to a carrier for transport, if the material is shipped in accordance with this section. This material need not be contained in a package which meets the standards of subparts E and F of 10 CFR Part 71; however, the material must be contained in a Type A package. The Type A package must also meet the USDOT requirements of 49 CFR § 173.417(a).

(ii) The general license applies only to a licensee who has a quality assurance program approved by the Department as satisfying § 175.105(e)(1) of this section.

(iii) The general license applies only when a package's contents:

(A) Contain less than a Type A quantity of material; and

(B) Contain less than 1000 g of plutonium, provided that: plutonium-239, plutonium-241, or any combination of these radionuclides, constitute less than 240 g of the total quantity of plutonium in the package.

(iv) The general license applies only to packages labeled with a CSI which:

(A) Has been determined in accordance with part (v) of this section;

(B) Has a value less than or equal to 100;

(C) For a shipment of multiple packages containing Pu-Be sealed sources, the sum of the CSI must be less than or equal to 50 (for shipment on a nonexclusive use conveyance) and less than or equal to 100 (for shipment on an exclusive use con-

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veyance).

(v) (A) The value for the CSI must be greater than or equal to the number calculated by the following equation:

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(B) The calculated CSI must be rounded up to the first decimal place.

(d) Operating Controls and Procedures. (1) Applicability of operating controls and procedures. A licensee subject to this section, who, under a general or specific license, transports licensed material or delivers licensed material to a carrier for transport, shall comply with the requirements of this subsection, with the quality assurance requirements of § 175.105(e), and with the general provisions of § 175.105(a) of this Code.

(2) Preliminary determinations. Before the first use of any packaging for the shipment of licensed material--

(i) The licensee shall ascertain that there are no cracks, pinholes, uncontrolled voids, or other defects that could significantly reduce the effectiveness of the packaging:

(ii) Where the maximum normal operating pressure will exceed 35 kPa (5 lbf/in2) gauge, the licensee shall test the containment system at an internal pressure at least 50 percent higher than the maximum normal operating pressure, to verify the capability of that system to maintain its structural integrity at that pressure; and

(iii) The licensee shall conspicuously and durably mark the packaging with its model number, serial number, gross weight, and a package identification number assigned by the NRC. Before applying the model number, the licensee shall determine that the packaging has been fabricated in accordance with the design approved by the NRC.

(3) Routine determinations. Before each shipment of licensed material, the licensee shall ensure that the package with its contents satisfies the applicable requirements of this section and of the license. The licensee shall determine that--

(i) The package is proper for the contents to be shipped;

(ii) The package is in unimpaired physical condition except for superficial defects such as marks or dents;

(iii) Each closure device of the packaging, including any required gasket, is properly installed and secured and free of defects;

(iv) Any system for containing liquid is adequately sealed and has adequate space

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or other specified provision for expansion of the liquid;

(v) Any pressure relief device is operable and set in accordance with written procedures;

(vi) The package has been loaded and closed in accordance with written procedures;

(vii) Any structural part of the package that could be used to lift or tie down the package during transport is rendered inoperable for that purpose, unless it satisfies the design requirements of 10 CFR 71.45;

(viii) The level of non-fixed (removable) radioactive contamination on the external surfaces of each package offered for shipment is as low as reasonably achievable, and within the limits specified in USDOT regulations in 49 CFR 173.443;

(ix) External radiation levels around the package and around the vehicle, if applicable, will not exceed the limits specified in 10 CFR 71.47 at any time during transportation;

(x) Accessible package surface temperatures will not exceed the limits specified in 10 CFR 71.43(g) at any time during transportation; and

(xi) For fissile material, any moderator or neutron absorber, if required, is present and in proper condition.

(xii) When the isotopic abundance, mass, concentration, degree of irradiation, degree of moderation, or other pertinent property of fissile material in any package is not known, the licensee shall package the fissile material as if the unknown properties have credible values that will cause the maximum neutron multiplication.

(4) Air transport of plutonium.

(i) Notwithstanding the provisions of any general licenses and notwithstanding any exemptions stated directly in this section or included indirectly by citation of 49 CFR chapter I, as may be applicable, the licensee shall assure that plutonium in any form, whether for import, export, or domestic shipment, is not transported by air or delivered to a carrier for air transport unless:

(A) The plutonium is contained in a medical device designed for individual human application; or

(B) The plutonium is contained in a material in which the specific activity is less than or equal to the activity concentration values for plutonium specified in Appendix A, Table A-2, of this section and in which the radioactivity is essentially uniformly distributed; or

(C) The plutonium is shipped in a single package containing no more than an A2 quantity of plutonium in any isotope or form, and is shipped in accordance with § 175.105(a)(6); or

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(D) The plutonium is shipped in a package specifically authorized for the shipment of plutonium by air in the Certificate of Compliance for that package issued by the NRC.

(ii) Nothing in § 175.105(d)(4)(i) is to be interpreted as removing or diminishing the requirements of 10 CFR 73.24

(iii) For a shipment of plutonium by air which is subject to § 175.105(d)(4)(i)(D), the licensee shall, through special arrangement with the carrier, require compliance with 49 CFR 175.704, U.S. Department of Transportation regulations applicable to the air transport of plutonium.

(5) Opening instructions. Before delivery of a package to a carrier for transport, the licensee shall ensure that any special instructions needed to safely open the package have been sent to, or otherwise made available to, the consignee for the consignee's use in accordance with § 175.03(j)(6) of this Code.

(6) Records.

(i) Each licensee shall maintain, for a period of 3 years after shipment, a record of each shipment of licensed material not exempt under § 175.105(b)(2), showing where applicable--

(A) Identification of the packaging by model number and serial number;

(B) Verification that there are no significant defects in the packaging, as shipped;

(C) Volume and identification of coolant;

(D) Type and quantity of licensed material in each package, and the total quantity of each shipment;

(E) For each item of irradiated fissile material--

(a) Identification by model number and serial number;

(b) Irradiation and decay history to the extent appropriate to demonstrate that its nuclear and thermal characteristics comply with license conditions; and

(c) Any abnormal or unusual condition relevant to radiation safety;

(F) Date of the shipment;

(G) For fissile packages and for Type B packages, any special controls exercised;

(H) Name and address of the transferee;

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(I) Address to which the shipment was made; and

(J) Results of the determinations required by § 175.105(d)(3) and by the conditions of the package approval.

(ii) The licensee shall make available to the Department for inspection, upon reasonable notice, all records required by this section. Records are only valid if stamped, initialed, or signed and dated by authorized personnel or otherwise authenticated.

(iii) The licensee shall maintain sufficient written records to furnish evidence of the quality of packaging. The records to be maintained include results of the determinations required by § 175.105(d)(2); design, fabrication, and assembly records, results of reviews, inspections, tests, and audits; results of monitoring work performance and materials analyses; and results of maintenance, modification and repair activities. Inspection, test, and audit records must identify the inspector or data recorder, the type of observation, the results, the acceptability and the action taken in connection with any deficiencies noted. The records must be retained for 3 years after the life of the packaging to which they apply.

(7) Inspection and tests.

(i) The licensee or certificate holder shall permit the Department, at all reasonable times, to inspect the licensed material, packaging, premises, and facilities in which the licensed material or packaging is used, provided, constructed, fabricated, tested, stored, or shipped.

(ii) The licensee shall perform, and permit the Department to perform, any tests the Department deems necessary or appropriate for the administration of the requirements of this section.

(iii) The licensee shall notify the Department at least 45 days before fabrication of a package to be used for the shipment of licensed material having a decay heat load in excess of 5kW or with a maximum normal operating pressure in excess of 103kPa (15 lbf/in2) gauge.

(8) Reports. The licensee shall report to the Department within 30 days--

(i) Any instance in which there is significant reduction in the effectiveness of any approved Type B, or fissile, packaging during use;

(ii) Details of any defects with safety significance in Type B, or fissile, packaging after first use, with the means employed to repair the defects and prevent their recurrence; or

(iii) Instances in which the conditions of approval in the certificate of compliance were not observed in making a shipment.

(9) Advance notification of shipment of irradiated reactor fuel and nuclear waste.

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(i) As specified in §§ 175.105(d)(9)(ii), (iii) and (iv), each licensee shall provide advance notification to the governor of a State, or the governor's designee, and the Department, of the shipment of licensed material, through, or across the boundary of the State, before the transport, or delivery to a carrier, for transport, of licensed material outside the confines of the licensee's plant or other place of use or storage.

(ii) Advance notification is required under this subdivision for shipments of irradiated reactor fuel in quantities less than that subject to advance notification requirements of 10 CFR 73.37(f). Advance notification is also required under this subdivision for shipment of licensed material; other than irradiated fuel, meeting the following three conditions:

(A) The licensed material is required by this section to be in Type B packaging for transportation;

(B) The licensed material is being transported to or across a State boundary en route to a disposal facility or to a collection point for transport to a disposal facility; and

(C) The quantity of licensed material in a single package exceeds the least of the following:

(a) 3000 times the A1 value of the radionuclides as specified in appendix A. Table A-1 for special form radioactive material;

(b) 3000 times the A2 value of the radionuclides as specified in appendix A. Table A-1 for normal form radioactive material; or

(c) 1000 TBq (27,000 Ci).

(iii) Procedures for submitting advance notification.

(A) The notification must be made in writing to the office of each appropriate governor or governor's designee and to the Department.

(B) A notification delivered by mail must be postmarked at least 7 days before the beginning of the 7-day period during which departure of the shipment is estimated to occur.

(C) A notification delivered by any means other than mail must reach the office of the governor or of the governor's designee and the Department at least 4 days before the beginning of the 7-day period during which departure of the shipment is estimated to occur.

(a) A list of the names and mailing addresses of the governors' designees receiving advance notification of transportation of nuclear waste was published in the Federal Register on June 30, 1995 (60 FR 34306).

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(b) The list will be published annually in the Federal Register on or about June 30 to reflect any changes in information.

(c) A list of the names and mailing addresses of the governors' designees is available on request from the Director, Office of State Programs, U.S. Nuclear Regulatory Commission, Washington, DC 20555-0001.

(D) The licensee shall retain a copy of the notification as a record for 3 years.

(iv) Information to be furnished in advance notification of shipment. Each advance notification of shipment of irradiated reactor fuel or nuclear waste must contain the following information:

(A) The name, address, and telephone number of the shipper, carrier, and receiver of the irradiated reactor fuel or nuclear waste shipment;

(B) A description of the irradiated reactor fuel or nuclear waste contained in the shipment, as specified in the regulations of USDOT in 49 CFR 172.202 and 172.203(d);

(C) The point of origin of the shipment and the 7-day period during which departure of the shipment is estimated to occur;

(D) The 7-day period during which arrival of the shipment at State boundaries is estimated to occur;

(E) The destination of the shipment, and the 7-day period during which arrival of the shipment is estimated to occur; and

(F) A point of contact, with a telephone number, for current shipment information.

(v) Revision notice. A licensee who finds that schedule information previously furnished to a governor or governor's designee, or the Department, in accordance with this subdivision, will not be met, shall telephone a responsible individual in the office of the governor of the State or of the governor's designee, and the Department, and inform that individual of the extent of the delay beyond the schedule originally reported. The licensee shall maintain a record of the name of the individual contacted for 3 years.

(vi) Cancellation notice.

(A) Each licensee who cancels an irradiated reactor fuel or nuclear waste shipment for which advance notification has been sent shall send a cancellation notice to the governor of each State or to the governor's designee previously notified, and to the Department.

(B) The licensee shall state in the notice that it is a cancellation and identify the advance notification that is being canceled. The licensee shall retain a copy of the notice as a record for 3 years.

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(e) Quality Assurance. (1) Quality assurance requirements.

(i) Purpose. This subsection describes quality assurance requirements applying to design, purchase, fabrication, handling, shipping, storing, cleaning, assembly, inspection, testing, operation, maintenance, repair, and modification of components of packaging that are important to safety. As used in this subsection, "quality assurance" comprises all those planned and systematic actions necessary to provide adequate confidence that a system or component will perform satisfactorily in service. Quality assurance includes quality control, which comprises those quality assurance actions related to control of the physical characteristics and quality of the material or component to predetermined requirements.

(ii) Establishment of program. Each licensee shall establish, maintain, and execute a quality assurance program satisfying each of the applicable criteria of §§ 175.105(e)(1) through 175.105(e)(19) and satisfying any specific provisions that are applicable to the licensee's activities including procurement of packaging. The licensee shall apply each of the applicable criteria in a graded approach, i.e., to an extent that is consistent with its importance to safety.

(iii) Approval of program. Before the use of any package for the shipment of licensed material subject to this subsection each licensee shall obtain Department approval of its quality assurance program. Each licensee shall file a description of its quality assurance program, including a discussion of which requirements of this subsection are applicable and how they will be satisfied, with the Department.

(iv) Repealed.

(v) Repealed.

(vi) Previously approved programs. An NRC-approved quality assurance program that satisfies the applicable criteria of Appendix B of 10 CFR Part 50, and that is established, maintained, and executed with regard to transport packages, will be accepted as satisfying the requirements of § 175.105(e)(1)(ii) of this Code. Before first use, the licensee shall notify the NRC and the Department of its intent to apply its previously approved Appendix B program to transportation activities. The licensee shall identify the program by date of submittal to the NRC and date of NRC approval.

(2) Quality assurance organization.

(i) The licensee shall be responsible for the establishment and execution of the quality assurance program. The licensee may delegate to others, such as contractors, agents, or consultants, the work of establishing and executing the quality assurance program, or any part of the quality assurance program, but shall retain responsibility for the program. The licensee shall clearly establish and delineate, in writing, the authority and duties of persons and organizations performing activities affecting the safety-related functions of structures, systems, and components. These activities include performing the functions associated with attaining quality objectives and the quality assurance functions.

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(ii) The quality assurance functions are--

(A) Assuring that an appropriate quality assurance program is established and effectively executed; and

(B) Verifying, by procedures such as checking, auditing, and inspection, that activities affecting the safety-related functions have been performed correctly.

(iii) The persons and organizations performing quality assurance functions must have sufficient authority and organizational freedom to--

(A) Identify quality problems;

(B) Initiate, recommend, or provide solutions; and

(C) Verify implementation of solutions.

(iv) The persons and organizations performing quality assurance functions shall report to a management level that assures that the required authority and organizational freedom, including sufficient independence from cost and schedule, when opposed to safety considerations, are provided.

(v) Because of the many variables involved, such as the number of personnel, the type of activity being performed, and the location or locations where activities are performed, the organizational structure for executing the quality assurance program may take various forms, provided that the persons and organizations assigned the quality assurance functions have the required authority and organizational freedom.

(vi) Irrespective of the organizational structure, the individual(s) assigned the responsibility for assuring effective execution of any portion of the quality assurance program, at any location where activities subject to this subsection are being performed, must have direct access to the levels of management necessary to perform this function.

(3) Quality assurance program.

(i) The licensee shall establish, at the earliest practicable time consistent with the schedule for accomplishing the activities, a quality assurance program that complies with the requirements of §§ 175.105(e)(1) through 175.105(e)(19). The licensee shall document the quality assurance program by written procedures or instructions and shall carry out the program in accordance with those procedures throughout the period during which the packaging is used. The licensee shall identify the material and components to be covered by the quality assurance program, the major organizations participating in the program, and the designated functions of these organizations.

(ii) The licensee, through its quality assurance program, shall provide control over activities affecting the quality of the identified materials and components to

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an extent consistent with their importance to safety, and as necessary to assure conformance to the approved design of each individual package used for the shipment of radioactive material. The licensee shall assure that activities affecting quality are accomplished under suitably controlled conditions. Controlled conditions include the use of appropriate equipment; suitable environmental conditions for accomplishing the activity, such as adequate cleanliness; and assurance that all prerequisites for the given activity have been satisfied. The licensee shall take into account the need for special controls, processes, test equipment, tools, and skills to attain the required quality, and the need for verification of quality by inspection and test.

(iii) The licensee shall base the requirements and procedures of its quality assurance program on the following considerations concerning the complexity and proposed use of the package and its components:

(A) The impact of malfunction or failure of the item to safety;

(B) The design and fabrication complexity or uniqueness of the item;

(C) The need for special controls and surveillance over processes and equipment;

(D) The degree to which functional compliance can be demonstrated by inspection or test; and

(E) The quality history and degree of standardization of the item.

(iv) The licensee shall provide for indoctrination and training of personnel performing activities affecting quality, as necessary to assure that suitable proficiency is achieved and maintained. The licensee shall review the status and adequacy of the quality assurance program at established intervals. Management of other organizations participating in the quality assurance program shall review regularly the status and adequacy of that part of the quality assurance program which they are executing.

(4) Package design control.

(i) The licensee shall establish measures to assure that applicable requirements and the package design, as specified in the license for those materials and components to which this subdivision applies, are correctly translated into specifications, drawings, procedures, and instructions. These measures must include provisions to assure that appropriate quality standards are specified and included in design documents and that deviations from standards are controlled. Measures must be established for the selection and review for suitability of application of materials, parts, equipment, and processes that are essential to the safety-related functions of the materials, parts, and components of the packaging.

(ii) The licensee shall establish measures for the identification and control of design interfaces and for coordination among participating design organizations. These measures must include the establishment of written procedures, among participating design organizations, for the review, approval, release, distribution, and

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revision of documents involving design interfaces. The design control measures must provide for verifying or checking the adequacy of design, by methods such as design reviews, alternate or simplified calculational methods, or by a suitable testing program. For the verifying or checking process, the licensee shall designate individuals or groups other than those who were responsible for the original design, but who may be from the same organization. Where a test program is used to verify the adequacy of a specific design feature in lieu of other verifying or checking processes, the licensee shall include suitable qualification testing of a prototype or sample unit under the most adverse design conditions. The licensee shall apply design control measures to items such as the following:

(A) Criticality physics, radiation shielding, stress, thermal, hydraulic, and accident analyses;

(B) Compatibility of materials;

(C) Accessibility for inservice inspection, maintenance, and repair;

(D) Features to facilitate decontamination; and

(E) Delineation of acceptance criteria for inspections and tests.

(iii) The licensee shall subject design changes, including field changes, to design control measures commensurate with those applied to the original design. Changes in the conditions specified in the package approval require the Department's approval.

(5) Procurement document control. The licensee shall establish measures to assure that adequate quality is required in the documents for procurement of material, equipment, and services, whether purchased by the licensee or by its contractors or subcontractors. To the extent necessary, the licensee shall require contractors or subcontractors to provide a quality assurance program consistent with the applicable provisions of this section.

(6) Instructions, procedures, and drawings. The licensee shall prescribe activities affecting quality by documented instructions, procedures, or drawings of a type appropriate to the circumstances and shall require that these instructions, procedures, and drawings be followed. The instructions, procedures, and drawings must include appropriate quantitative or qualitative acceptance criteria for determining that important activities have been satisfactorily accomplished.

(7) Document control. The licensee shall establish measures to control the issuance of documents such as instructions, procedures, and drawings, including changes, which prescribe all activities affecting quality. These measures must assure that documents, including changes, are reviewed for adequacy, approved for release by authorized personnel, and distributed and used at the location where the prescribed activity is performed. These measures must assure that changes to documents are reviewed and approved.

(8) Control of purchased material, equipment, and services.

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(i) The licensee shall establish measures to assure that purchased material, equipment, and services, whether purchased directly or through contractors and subcontractors, conform to the procurement documents. These measures must include provisions, as appropriate, for source evaluation and selection, objective evidence of quality furnished by the contractor or subcontractor, inspection at the contractor or subcontractor source, and examination of products on delivery.

(ii) The licensee shall have available documentary evidence that material and equipment conform to the procurement specifications before installation or use of the material and equipment. The licensee shall retain, or have available, this documentary evidence for the life of the package to which it applies. The licensee shall assure that the evidence is sufficient to identify the specific requirements met by the purchased material and equipment.

(iii) The licensee shall assess the effectiveness of the control of quality by contractors and subcontractors at intervals consistent with the importance, complexity, and quantity of the product or services.

(9) Identification and control of materials, parts, and components. The licensee shall establish measures for the identification and control of materials, parts, and components. These measures must assure that identification of the item is maintained by heat number, part number, or other appropriate means, either on the item or on records traceable to the item, as required throughout fabrication, installation, and use of the item. These identification and control measures must be designed to prevent the use of incorrect or defective materials, parts, and components.

(10) Control of special processes. The licensee shall establish measures to assure that special processes, including welding, heat treating, and nondestructive testing, are controlled and accomplished by qualified personnel using qualified procedures in accordance with applicable codes, standards, specifications, criteria, and other special requirements.

(11) Internal inspection. The licensee shall establish and execute a program for inspection of activities affecting quality by or for the organization performing the activity, to verify conformance with the documented instructions, procedures, and drawings for accomplishing the activity. The inspection must be performed by individuals other than those who performed the activity being inspected. Examination, measurements, or tests of material or products processed must be performed for each work operation where necessary to assure quality. If direct inspection of processed material or products is not carried out, indirect control by monitoring processing methods, equipment, and personnel must be provided. Both inspection and process monitoring must be provided when quality control is inadequate without both. If mandatory inspection hold points, which require witnessing or inspecting by the licensee's designated representative and beyond which work should not proceed without the consent of its designated representative, are required, the specific hold points must be indicated in appropriate documents.

(12) Test control. The licensee shall establish a test program to assure that all testing required to demonstrate that the packaging components will perform satis-

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factorily in service is identified and performed in accordance with written test procedures that incorporate the requirements of this section and the requirements and acceptance limits contained in the package approval. The test procedures must include provisions for assuring that all prerequisites for the given test are met, that adequate test instrumentation is available and used, and that the test is performed under suitable environmental conditions. The licensee shall document and evaluate the test results to assure that test requirements have been satisfied.

(13) Control of measuring and test equipment. The licensee shall establish measures to assure that tools, gauges, instruments, and other measuring and testing devices used in activities affecting quality are properly controlled, calibrated, and adjusted at specified times to maintain accuracy within necessary limits.

(14) Handling, storage, and shipping control. The licensee shall establish measures to control, in accordance with instructions, the handling, storage, shipping, cleaning, and preservation of materials and equipment to be used in packaging to prevent damage or deterioration. When necessary for particular products, special protective environments, such as inert gas atmosphere, and specific moisture content and temperature levels must be specified and provided.

(15) Inspection, test, and operating status. (i) The licensee shall establish measures to indicate, by the use of markings such as stamps, tags, labels, routing cards, or other suitable means, the status of inspections and tests performed upon individual items of the packaging. These measures must provide for the identification of items that have satisfactorily passed required inspections and tests, where necessary to preclude inadvertent bypassing of the inspections and tests.

(ii) The licensee shall establish measures to identify the operating status of components of the packaging, such as tagging valves and switches, to prevent inadvertent operation.

(16) Nonconforming materials, parts, or components. The licensee shall establish measures to control materials, parts, or components that do not conform to the licensee's requirements to prevent their inadvertent use or installation. These measures must include, as appropriate, procedures for identification, documentation, segregation, disposition, and notification to affected organizations. Nonconforming items must be reviewed and accepted, rejected, repaired, or reworked in accordance with documented procedures.

(17) Corrective action. The licensee shall establish measures to assure that conditions adverse to quality, such as deficiencies, deviations, defective material and equipment, and nonconformances, are promptly identified and corrected. In the case of a significant condition adverse to quality, the measures must assure that the cause of the condition is determined and corrective action taken to preclude repetition. The identification of the significant condition adverse to quality, the cause of the condition, and the corrective action taken must be documented and reported to appropriate levels of management.

(18) Quality assurance records. The licensee shall maintain sufficient written records to describe the activities affecting quality. The records must include the instructions, procedures, and drawings required by subdivision (6) of this subsec-

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tion, to prescribe quality assurance activities and must include closely related specifications such as required qualifications of personnel, procedures, and equipment. The records must include the instructions or procedures which establish a records retention program that is consistent with applicable requirements and designates factors such as duration, location, and assigned responsibility. The licensee shall retain these records for 3 years beyond the date when the licensee last engages in the activity for which the quality assurance program was developed. If any portion of the written procedures or instructions is superseded, the licensee shall retain the superseded material for 3 years after it is superseded.

(19) Audits. The licensee shall carry out a comprehensive system of planned and periodic audits, to verify compliance with all aspects of the quality assurance program, and to determine the effectiveness of the program. The audits must be performed in accordance with written procedures or checklists by appropriately trained personnel not having direct responsibilities in the areas being audited. Audited results must be documented and reviewed by management having responsibility in the area audited. Follow-up action, including reaudit of deficient areas, must be taken where indicated.

HISTORICAL NOTE

Section repealed and added City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

<General Materials (GM) - References, Annotations, or Tables>

NOTES:

This section was repealed and reenacted on April 26, 1999 as a matter of compatibility to conform to NRC regulations found in 10 CFR Part 71 governing packaging and transportation of radioactive material.

I. Values of A1 and A2 for individual radionuclides, which are the bases for many activity limits elsewhere in this Code, are given in Table A-1. The curie (Ci) values specified are obtained by converting from the Terabecquerel (TBq) value. The Terabecquerel values are the regulatory standard. The curie values are for information only and are not intended to be the regulatory standard. Where values of A1 and A2 are unlimited, it is for radiation control purposes only. For nuclear criticality safety, some materials are subject to controls placed on fissile material.

II. a. For individual radionuclides whose identities are known, but which are not listed in Table A-1, the A1 and A2 values contained in Table A-3 may be used. Otherwise, the licensee shall obtain prior Department approval of the A1 and A2 values for radionuclides not listed in Table A-1, before shipping the material.

b. For individual radionuclides whose identities are known, but which are not listed in Table A-2, the exempt material activity concentration and exempt consign-

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ment activity values contained in Table A-3 may be used. Otherwise, the licensee shall obtain prior Department approval of the exempt material activity concentration and exempt consignment activity values for radionuclides not listed in Table A-2, before shipping the material.

c. The licensee shall submit requests for prior approval, described under paragraphs II(a) and II(b) of this Appendix, to the Department.

III. In the calculations of A1 and A2 for a radionuclide not in Table A-1, a single radioactive decay chain, in which radionuclides are present in their naturally occurring proportions, and in which no daughter radionuclide has a half-life either longer than 10 days, or longer than that of the parent radionuclide, shall be considered as a single radionuclide, and the activity to be taken into account, and the A1 or A2 value to be applied, shall be those corresponding to the parent radionuclide of that chain. In the case of radioactive decay chains in which any daughter radionuclide has a half-life either longer than 10 days, or greater than that of the parent radionuclide, the parent and those daughter radionuclides shall be considered as mixtures of different radionuclides.

IV. For mixtures of radionuclides whose identities and respective activities are known, the following conditions apply:

a. For special form radioactive material, the maximum quantity transported in a Type A package is as follows:

<<SIGMA>>,10i 0 B(i) _____ A1(i) #1 where B(i) is the activity of radionuclide i, and A1(i) is the A1 value for radionuclide I.

b. For normal form radioactive material, the maximum quantity transported in a Type A package is as follows: <<SIGMA>>B(i)/A2(i)<1 where B(i) is the activity of radionuclide i, and A2(i) is the A2 value for radionuclide i.

c. Alternatively, the A1 value for mixtures of special form material may be determined as follows: A1 for mixture = _____61

<<SIGMA>>,10i 0 f(i) _____ A1(i) where f(i) is the fraction of activity for radionuclide I in the mixture, and

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A1(i) is the appropriate A1 value for radionuclide I.

d. Alternatively, the A2 value for mixtures of normal form material may be determined as follows: A2 for mixture = _____61
<<SIGMA>>,10i 0 f(i) _____ A2(i) where f(i) is the fraction of activity for radionuclide I in the mixture, and A2(i) is the appropriate A2 value for radionuclide I.

e. The exempt activity concentration for mixtures of nuclides may be determined as follows: Exempt activity concentration for mixture = _____61
<<SIGMA>>,10i 0 f(i) _____ (i) where f(i) is the fraction of activity concentration of radionuclide I in the mixture, and [A] is the activity concentration for exempt material containing radionuclide I.

f. The activity limit for an exempt consignment for mixtures of radionuclides may be determined as follows: Exempt activity concentration for mixture = _____61
<<SIGMA>>,10i 0 f(i) _____ A(i) where f(i) is the fraction of activity of radionuclide I in the mixture, and A is the activity limit for exempt consignments for radionuclide I.

V. When the identity of each radionuclide is known, but the individual activities of some of the radionuclides are not known, the

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radionuclides may be grouped, and the lowest A1 or A2 value, as appropriate, for the radionuclides in each group may be used in applying the formulas in paragraph IV. Groups may be based on the total alpha activity and the total beta/gamma activity when these are known, using the lowest A1 or A2 values for the alpha emitters and beta/gamma emitters.

<<SIGMA>>,10i 0 B(i) _____ A1(i) #1 where B(i)

is the

activity of

radionuclide

i, and A1(i)

is the A1

value for

radionuclide

I.

b. For normal form radioactive material,

the maximum quantity transported in a

Type A package is as follows:

<<SIGMA>>B(i)/A2(i)<1 where B(i) is

the activity of radionuclide i, and

A2(i) is the A2 value for radionuclide

i.

c. Alternatively, the A1 value for

mixtures of special form material may

be determined as follows: A1 for

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mixture = _____61

<<SIGMA>>,10i 0 f(i) _____ A1(i)

where f(i) is the fraction of activity for radionuclide I in the mixture, and A1(i) is the appropriate A1 value for radionuclide I.

d. Alternatively, the A2 value for mixtures of normal form material may be determined as follows: A2 for

mixture = _____61

<<SIGMA>>,10i 0 f(i) _____ A2(i)

where f(i) is the fraction of activity for radionuclide I in the mixture, and A2(i) is the appropriate A2

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value for

radionuclide I.

e. The exempt activity concentration for

mixtures of nuclides may be determined

as follows: Exempt activity

concentration for mixture = _____61

<<SIGMA>>,10i 0 f(i) _____ (i)

where f(i) is the

fraction of

activity

concentration of

radionuclide I in

the mixture, and

[A] is the activity

concentration for

exempt material

containing

radionuclide I.

f. The activity limit for an exempt

consignment for mixtures of

radionuclides may be determined as

follows: Exempt activity concentration

for mixture = _____61

<<SIGMA>>,10i 0 f(i) _____ A(i)

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where f(i) is the fraction of activity of radionuclide I in the mixture, and A is the activity limit for exempt consignments for radionuclide I.

V. When the identity of each radionuclide is known, but the individual activities of some of the radionuclides are not known, the radionuclides may be grouped, and the lowest A1 or A2 value, as appropriate, for the radionuclides in each group may be used in applying the formulas in paragraph IV. Groups may be based on the total alpha activity and the total beta/gamma activity when these are known, using the lowest A1 or A2 values for the alpha emitters and beta/gamma emitters.

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[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

Table A-1
A1 and A2 Values for Radionuclides

Symbol of radionuclide	Element and atomic number	A1 (TBq)
A1 and A2 Values for Radionuclides		
Symbol of radionuclide	Element and atomic number	A1 (TBq)
Ac-225 (a)	Actinium (89)	8.0X10-1
Ac-227 (a)		9.0X10-1
Ac-228		6.0X10-1
Ag-105	Silver (47)	2.0
Ag-108m (a)		7.0X10-1
Ag-110m (a)		4.0X10-1
Ag-111		2.0
Al-26	Aluminum (13)	1.0X10-1
Am-241	Americium (95)	1.0X101
Am-242m (a)		1.0X101
Am-243 (a)		5.0
Ar-37	Argon (18)	4.0X101
Ar-39		4.0X101
Ar-41		3.0X10-1
As-72	Arsenic (33)	3.0X10-1
As-73		4.0X101
As-74		1.0
As-76		3.0X10-1
As-77		2.0X101
At-211 (a)	Astatine (85)	2.0X101
Au-193	Gold (79)	7.0
Au-194		1.0
Au-195		1.0X101
Au-198		1.0
Au-199		1.0X101
Ba-131 (a)	Barium (56)	2.0
Ba-133		3.0
Ba-133m		2.0X101
Ba-140 (a)		5.0X10-1
Be-7	Beryllium (4)	2.0X101
Be-10		4.0X101
Bi-205	Bismuth (83)	7.0X10-1
Bi-206		3.0X10-1
Bi-207		7.0X10-1

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Bi-210		1.0
Bi-210m (a)		6.0X10-1
Bi-212 (a)		7.0X10-1
Bk-247	Berkelium (97)	8.0
Bk-249 (a)		4.0X101
Br-76	Bromine (35)	4.0X10-1
Br-77		3.0
Br-82		4.0X10-1
C-11	Carbon (6)	1.0
C-14		4.0X101
Ca-41	Calcium (20)	Unlimited
Ca-45		4.0X101
Ca-47 (a)		3.0
Cd-109	Cadmium (48)	3.0X101
Cd-113m		4.0X101
Cd-115 (a)		3.0
Cd-115m		5.0X10-1
Ce-139	Cerium (58)	7.0
Ce-141		2.0X101
Ce-143		9.0X10-1
Ce-144 (a)		2.0X10-1
Cf-248	Californium (98)	4.0X101
Cf-249		3.0
Cf-250		2.0X101
Cf-251		7.0
Cf-252 (h)		5.0X10-2
Cf-253 (a)		4.0X101
Cf-254		1.0X10-3
Cl-36	Chlorine (17)	1.0X101
Cl-38		2.0X10-1
Cm-240	Curium (96)	4.0X101
Cm-241		2.0
Cm-242		4.0X101
Cm-243		9.0
Cm-244		2.0X101
Cm-245		9.0
Cm-246		9.0
Cm-247 (a)		3.0
Cm-248		2.0X10-2
Co-55	Cobalt (27)	5.0X10-1
Co-56		3.0X10-1
Co-57		1.0X101
Co-58		1.0
Co-58m		4.0X101
Co-60		4.0X10-1
Cr-51	Chromium (24)	3.0X101
Cs-129	Cesium (55)	4.0
Cs-131		3.0X101
Cs-132		1.0
Cs-134		7.0X10-1
Cs-134m		4.0X101
Cs-135		4.0X101
Cs-136		5.0X10-1

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Cs-137 (a)		2.0
Cu-64	Copper (29)	6.0
Cu-67		1.0X101
Dy-159	Dysprosium (66)	2.0X101
Dy-165		9.0X10-1
Dy-166 (a)		9.0X10-1
Er-169	Erbium (68)	4.0X101
Er-171		8.0X10-1
Eu-147	Europium (63)	2.0
Eu-148		5.0X10-1
Eu-149		2.0X101
Eu-150 (short lived)		2.0
Eu-150 (long lived)		7.0X10-1
Eu-152		1.0
Eu-152m		8.0X10-1
Eu-154		9.0X10-1
Eu-155		2.0X101
Eu-156		7.0X10-1
F-18	Fluorine (9)	1.0
Fe-52 (a)	Iron (26)	3.0X10-1
Fe-55		4.0X101
Fe-59		9.0X10-1
Fe-60 (a)		4.0X101
Ga-67	Gallium (31)	7.0
Ga-68		5.0X10-1
Ga-72		4.0X10-1
Gd-146 (a)	Gadolinium (64)	5.0X10-1
Gd-148		2.0X101
Gd-153		1.0X101
Gd-159		3.0
Ge-68 (a)	Germanium (32)	5.0X10-1

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+

****
******* This is piece 2. -- It begins at character 76 of table line 1. ********
****

A1 (Ci)	A2 (TBq)	A2 (Ci)	Specific activity	(TBq/g)	(Ci/g)
A1 (Ci)	A2 (TBq)	A2 (Ci)	Specific activity	(TBq/g)	(Ci/g)
2.2X101	6.0X10-3	1.6X10-1	2.1X103	5.8X104	
2.4X101	9.0X10-5	2.4X10-3	2.7	7.2X101	
1.6X101	5.0X10-1	1.4X101	8.4X104	2.2X106	
5.4X101	2.0	5.4X101	1.1X103	3.0X104	
1.9X101	7.0X10-1	1.9X101	9.7X10-1	2.6X101	
1.1X101	4.0X10-1	1.1X101	1.8X102	4.7X103	
5.4X101	6.0X10-1	1.6X101	5.8X103	1.6X105	
2.7	1.0X10-1	2.7	7.0X10-4	1.9X10-2	
2.7X102	1.0X10-3	2.7X10-2	1.3X10-1	3.4	
2.7X102	1.0X10-3	2.7X10-2	3.6X10-1	1.0X101	

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1.4X102	1.0X10-3	2.7X10-2	7.4X10-3	2.0X10-1
1.1X103	4.0X101	1.1X103	3.7X103	9.9X104
1.1X103	2.0X101	5.4X102	1.3	3.4X101
8.1	3.0X10-1	8.1	1.5X106	4.2X107
8.1	3.0X10-1	8.1	6.2X104	1.7X106
1.1X103	4.0X101	1.1X103	8.2X102	2.2X104
2.7X101	9.0X10-1	2.4X101	3.7X103	9.9X104
8.1	3.0X10-1	8.1	5.8X104	1.6X106
5.4X102	7.0X10-1	1.9X101	3.9X104	1.0X106
5.4X102	5.0X10-1	1.4X101	7.6X104	2.1X106
1.9X102	2.0	5.4X101	3.4X104	9.2X105
2.7X101	1.0	2.7X101	1.5X104	4.1X105
2.7X102	6.0	1.6X102	1.4X102	3.7X103
2.7X101	6.0X10-1	1.6X101	9.0X103	2.4X105
2.7X102	6.0X10-1	1.6X101	7.7X103	2.1X105
5.4X101	2.0	5.4X101	3.1X103	8.4X104
8.1X101	3.0	8.1X101	9.4	2.6X102
5.4X102	6.0X10-1	1.6X101	2.2X104	6.1X105
1.4X101	3.0X10-1	8.1	2.7X103	7.3X104
5.4X102	2.0X101	5.4X102	1.3X104	3.5X105
1.1X103	6.0X10-1	1.6X101	8.3X10-4	2.2X10-2
1.9X101	7.0X10-1	1.9X101	1.5X103	4.2X104
8.1	3.0X10-1	8.1	3.8X103	1.0X105
1.9X101	7.0X10-1	1.9X101	1.9	5.2X101
2.7X101	6.0X10-1	1.6X101	4.6X103	1.2X105
1.6X101	2.0X10-2	5.4X10-1	2.1X10-5	5.7X10-4
1.9X101	6.0X10-1	1.6X101	5.4X105	1.5X107
2.2X102	8.0X10-4	2.2X10-2	3.8X10-2	1.0
1.1X103	3.0X10-1	8.1	6.1X101	1.6X103
1.1X101	4.0X10-1	1.1X101	9.4X104	2.5X106
8.1X101	3.0	8.1X101	2.6X104	7.1X105
1.1X101	4.0X10-1	1.1X101	4.0X104	1.1X106
2.7X101	6.0X10-1	1.6X101	3.1X107	8.4X108
1.1X103	3.0	8.1X101	1.6X10-1	4.5
Unlimited	Unlimited	Unlimited	3.1X10-3	8.5X10-2
1.1X103	1.0	2.7X101	6.6X102	1.8X104
8.1X101	3.0X10-1	8.1	2.3X104	6.1X105
8.1X102	2.0	5.4X101	9.6X101	2.6X103
1.1X103	5.0X10-1	1.4X101	8.3	2.2X102
8.1X101	4.0X10-1	1.1X101	1.9X104	5.1X105
1.4X101	5.0X10-1	1.4X101	9.4X102	2.5X104
1.9X102	2.0	5.4X101	2.5X102	6.8X103
5.4X102	6.0X10-1	1.6X101	1.1X103	2.8X104
2.4X101	6.0X10-1	1.6X101	2.5X104	6.6X105
5.4	2.0X10-1	5.4	1.2X102	3.2X103
1.1X103	6.0X10-3	1.6X10-1	5.8X101	1.6X103
8.1X101	8.0X10-4	2.2X10-2	1.5X10-1	4.1
5.4X102	2.0X10-3	5.4X10-2	4.0	1.1X102
1.9X102	7.0X10-4	1.9X10-2	5.9X10-2	1.6
1.4	3.0X10-3	8.1X10-2	2.0X101	5.4X102
1.1X103	4.0X10-2	1.1	1.1X103	2.9X104
2.7X10-2	1.0X10-3	2.7X10-2	3.1X102	8.5X103
2.7X102	6.0X10-1	1.6X101	1.2X10-3	3.3X10-2

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5.4	2.0X10-1	5.4	4.9X106	1.3X108
1.1X103	2.0X10-2	5.4X10-1	7.5X102	2.0X104
5.4X101	1.0	2.7X101	6.1X102	1.7X104
1.1X103	1.0X10-2	2.7X10-1	1.2X102	3.3X103
2.4X102	1.0X10-3	2.7X10-2	1.9X10-3	5.2X101
5.4X102	2.0X10-3	5.4X10-2	3.0	8.1X101
2.4X102	9.0X10-4	2.4X10-2	6.4X10-3	1.7X10-1
2.4X102	9.0X10-4	2.4X10-2	1.1X10-2	3.1X10-1
8.1X101	1.0X10-3	2.7X10-2	3.4X10-6	9.3X10-5
5.4X10-1	3.0X10-4	8.1X10-3	1.6X10-4	4.2X10-3
1.4X101	5.0X10-1	1.4X101	1.1X105	3.1X106
8.1	3.0X10-1	8.1	1.1X103	3.0X104
2.7X102	1.0X101	2.7X102	3.1X102	8.4X103
2.7X101	1.0	2.7X101	1.2X103	3.2X104
1.1X103	4.0X101	1.1X103	2.2X105	5.9X106
1.1X101	4.0X10-1	1.1X101	4.2X101	1.1X103
8.1X102	3.0X101	8.1X102	3.4X103	9.2X104
1.1X102	4.0	1.1X102	2.8X104	7.6X105
8.1X102	3.0X101	8.1X102	3.8X103	1.0X105
2.7X101	1.0	2.7X101	5.7X103	1.5X105
1.9X101	7.0X10-1	1.9X101	4.8X101	1.3X103
1.1X103	6.0X10-1	1.6X101	3.0X105	8.0X106
1.1X103	1.0	2.7X101	4.3X10-5	1.2X10-3
1.4X101	5.0X10-1	1.4X101	2.7X103	7.3X104
5.4X101	6.0X10-1	1.6X101	3.2	8.7X101
1.6X102	1.0	2.7X101	1.4X105	3.9X106
2.7X102	7.0X10-1	1.9X101	2.8X104	7.6X105
5.4X102	2.0X101	5.4X102	2.1X102	5.7X103
2.4X101	6.0X10-1	1.6X101	3.0X105	8.2X106
2.4X101	3.0X10-1	8.1	8.6X103	2.3X105
1.1X103	1.0	2.7X101	3.1X103	8.3X104
2.2X101	5.0X10-1	1.4X101	9.0X104	2.4X106
5.4X101	2.0	5.4X101	1.4X103	3.7X104
1.4X101	5.0X10-1	1.4X101	6.0X102	1.6X104
5.4X102	2.0X101	5.4X102	3.5X102	9.4X103
5.4X101	7.0X10-1	1.9X101	6.1X104	1.6X106
1.9X101	7.0X10-1	1.9X101	6.1X104	1.6X106
2.7X101	1.0	2.7X101	6.5	1.8X102
2.2X101	8.0X10-1	2.2X101	8.2X104	2.2X106
2.4X101	6.0X10-1	1.6X101	9.8	2.6X102
5.4X102	3.0	8.1X101	1.8X101	4.9X102
1.9X101	7.0X10-1	1.9X101	2.0X103	5.5X104
2.7X101	6.0X10-1	1.6X101	3.5X106	9.5X107
8.1	3.0X10-1	8.1	2.7X105	7.3X106
1.1X103	4.0X101	1.1X103	8.8X101	2.4X103
2.4X101	9.0X10-1	2.4X101	1.8X103	5.0X104
1.1X103	2.0X10-1	5.4	7.4X10-4	2.0X10-2
1.9X102	3.0	8.1X101	2.2X104	6.0X105
1.4X101	5.0X10-1	1.4X101	1.5X106	4.1X107
1.1X101	4.0X10-1	1.1X101	1.1X105	3.1X106
1.4X101	5.0X10-1	1.4X101	6.9X102	1.9X104
5.4X102	2.0X10-3	5.4X10-2	1.2	3.2X101
2.7X102	9.0	2.4X102	1.3X102	3.5X103

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 8.1X101     6.0X10-1    1.6X101     3.9X104              1.1X106
 1.4X101     5.0X10-1    1.4X101     2.6X102              7.1X103
76.......+...90....+....0....+...10....+...20....+...30....+...40....+...5
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[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

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*******************************************************************************

******** This is piece 1. -- It begins at character 1 of table line 1. ********

*******************************************************************************
```

Table A-1

A1 and A2 Values for Radionuclides

Symbol of radionuclide Element and atomic number A1 (TBq)

A1 and A2 Values for Radionuclides

Symbol of radionuclide Element and atomic number A1 (TBq)

Ac-225 (a) Actinium (89) 8.0X10-1

Ac-227 (a) 9.0X10-1

Ac-228 6.0X10-1

Ag-105 Silver (47) 2.0

Ag-108m (a) 7.0X10-1

Ag-110m (a) 4.0X10-1

Ag-111 2.0

Al-26 Aluminum (13) 1.0X10-1

Am-241 Americium (95) 1.0X101

Am-242m (a) 1.0X101

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Am-243 (a)	5.0		
Ar-37		Argon (18)	4.0X101
Ar-39	4.0X101		
Ar-41	3.0X10-1		
As-72		Arsenic (33)	3.0X10-1
As-73	4.0X101		
As-74	1.0		
As-76	3.0X10-1		
As-77	2.0X101		
At-211 (a)		Astatine (85)	2.0X101
Au-193		Gold (79)	7.0
Au-194	1.0		
Au-195	1.0X101		
Au-198	1.0		
Au-199	1.0X101		
Ba-131 (a)		Barium (56)	2.0
Ba-133	3.0		
Ba-133m	2.0X101		
Ba-140 (a)	5.0X10-1		
Be-7		Beryllium (4)	2.0X101
Be-10	4.0X101		
Bi-205		Bismuth (83)	7.0X10-1
Bi-206	3.0X10-1		

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Bi-207 7.0X10-1

Bi-210 1.0

Bi-210m (a) 6.0X10-1

Bi-212 (a) 7.0X10-1

Bk-247 Berkelium (97) 8.0

Bk-249 (a) 4.0X101

Br-76 Bromine (35) 4.0X10-1

Br-77 3.0

Br-82 4.0X10-1

C-11 Carbon (6) 1.0

C-14 4.0X101

Ca-41 Calcium (20) Unlimited

Ca-45 4.0X101

Ca-47 (a) 3.0

Cd-109 Cadmium (48) 3.0X101

Cd-113m 4.0X101

Cd-115 (a) 3.0

Cd-115m 5.0X10-1

Ce-139 Cerium (58) 7.0

Ce-141 2.0X101

Ce-143 9.0X10-1

Ce-144 (a) 2.0X10-1

Cf-248 Californium (98) 4.0X101

Cf-249 3.0

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Cf-250	2.0X101		
Cf-251	7.0		
Cf-252 (h)	5.0X10-2		
Cf-253 (a)	4.0X101		
Cf-254	1.0X10-3		
Cl-36		Chlorine (17)	1.0X101
Cl-38	2.0X10-1		
Cm-240		Curium (96)	4.0X101
Cm-241	2.0		
Cm-242	4.0X101		
Cm-243	9.0		
Cm-244	2.0X101		
Cm-245	9.0		
Cm-246	9.0		
Cm-247 (a)	3.0		
Cm-248	2.0X10-2		
Co-55		Cobalt (27)	5.0X10-1
Co-56	3.0X10-1		
Co-57	1.0X101		
Co-58	1.0		
Co-58m	4.0X101		
Co-60	4.0X10-1		
Cr-51		Chromium (24)	3.0X101

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Cs-129 Cesium (55) 4.0

Cs-131 3.0X101

Cs-132 1.0

Cs-134 7.0X10-1

Cs-134m 4.0X101

Cs-135 4.0X101

Cs-136 5.0X10-1

Cs-137 (a) 2.0

Cu-64 Copper (29) 6.0

Cu-67 1.0X101

Dy-159 Dysprosium (66) 2.0X101

Dy-165 9.0X10-1

Dy-166 (a) 9.0X10-1

Er-169 Erbium (68) 4.0X101

Er-171 8.0X10-1

Eu-147 Europium (63) 2.0

Eu-148 5.0X10-1

Eu-149 2.0X101

Eu-150 (short lived) 2.0

Eu-150 (long lived) 7.0X10-1

Eu-152 1.0

Eu-152m 8.0X10-1

Eu-154 9.0X10-1

Eu-155 2.0X101

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Eu-156 7.0X10-1

F-18 Fluorine (9) 1.0

Fe-52 (a) Iron (26) 3.0X10-1

Fe-55 4.0X101

Fe-59 9.0X10-1

Fe-60 (a) 4.0X101

Ga-67 Gallium (31) 7.0

Ga-68 5.0X10-1

Ga-72 4.0X10-1

Gd-146 (a) Gadolinium (64) 5.0X10-1

Gd-148 2.0X101

Gd-153 1.0X101

Gd-159 3.0

Ge-68 (a) Germanium (32) 5.0X10-1

1...k...10....k...20....k...30....k...40....k...50....k...60....k...70....k

****

******* This is piece 2. -- It begins at character 76 of table line 1. ********

****

A1 (Ci) A2 (TBq) A2 (Ci) Specific activity (TBq/g) (Ci/g)

A1 (Ci) A2 (TBq) A2 (Ci) Specific activity (TBq/g) (Ci/g)

2.2X101 6.0X10-3 1.6X10-1 2.1X103 5.8X104

2.4X101 9.0X10-5 2.4X10-3 2.7 7.2X101

1.6X101 5.0X10-1 1.4X101 8.4X104 2.2X106

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5.4X101	2.0	5.4X101	1.1X103	3.0X104
1.9X101	7.0X10-1	1.9X101	9.7X10-1	2.6X101
1.1X101	4.0X10-1	1.1X101	1.8X102	4.7X103
5.4X101	6.0X10-1	1.6X101	5.8X103	1.6X105
2.7	1.0X10-1	2.7	7.0X10-4	1.9X10-2
2.7X102	1.0X10-3	2.7X10-2	1.3X10-1	3.4
2.7X102	1.0X10-3	2.7X10-2	3.6X10-1	1.0X101
1.4X102	1.0X10-3	2.7X10-2	7.4X10-3	2.0X10-1
1.1X103	4.0X101	1.1X103	3.7X103	9.9X104
1.1X103	2.0X101	5.4X102	1.3	3.4X101
8.1	3.0X10-1	8.1	1.5X106	4.2X107
8.1	3.0X10-1	8.1	6.2X104	1.7X106
1.1X103	4.0X101	1.1X103	8.2X102	2.2X104
2.7X101	9.0X10-1	2.4X101	3.7X103	9.9X104
8.1	3.0X10-1	8.1	5.8X104	1.6X106
5.4X102	7.0X10-1	1.9X101	3.9X104	1.0X106
5.4X102	5.0X10-1	1.4X101	7.6X104	2.1X106
1.9X102	2.0	5.4X101	3.4X104	9.2X105
2.7X101	1.0	2.7X101	1.5X104	4.1X105
2.7X102	6.0	1.6X102	1.4X102	3.7X103
2.7X101	6.0X10-1	1.6X101	9.0X103	2.4X105
2.7X102	6.0X10-1	1.6X101	7.7X103	2.1X105
5.4X101	2.0	5.4X101	3.1X103	8.4X104
8.1X101	3.0	8.1X101	9.4	2.6X102

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5.4X102	6.0X10-1	1.6X101	2.2X104	6.1X105
1.4X101	3.0X10-1	8.1	2.7X103	7.3X104
5.4X102	2.0X101	5.4X102	1.3X104	3.5X105
1.1X103	6.0X10-1	1.6X101	8.3X10-4	2.2X10-2
1.9X101	7.0X10-1	1.9X101	1.5X103	4.2X104
8.1	3.0X10-1	8.1	3.8X103	1.0X105
1.9X101	7.0X10-1	1.9X101	1.9	5.2X101
2.7X101	6.0X10-1	1.6X101	4.6X103	1.2X105
1.6X101	2.0X10-2	5.4X10-1	2.1X10-5	5.7X10-4
1.9X101	6.0X10-1	1.6X101	5.4X105	1.5X107
2.2X102	8.0X10-4	2.2X10-2	3.8X10-2	1.0
1.1X103	3.0X10-1	8.1	6.1X101	1.6X103
1.1X101	4.0X10-1	1.1X101	9.4X104	2.5X106
8.1X101	3.0	8.1X101	2.6X104	7.1X105
1.1X101	4.0X10-1	1.1X101	4.0X104	1.1X106
2.7X101	6.0X10-1	1.6X101	3.1X107	8.4X108
1.1X103	3.0	8.1X101	1.6X10-1	4.5
Unlimited	Unlimited	Unlimited	3.1X10-3	8.5X10-2
1.1X103	1.0	2.7X101	6.6X102	1.8X104
8.1X101	3.0X10-1	8.1	2.3X104	6.1X105
8.1X102	2.0	5.4X101	9.6X101	2.6X103
1.1X103	5.0X10-1	1.4X101	8.3	2.2X102
8.1X101	4.0X10-1	1.1X101	1.9X104	5.1X105

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1.4X101	5.0X10-1	1.4X101	9.4X102	2.5X104
1.9X102	2.0	5.4X101	2.5X102	6.8X103
5.4X102	6.0X10-1	1.6X101	1.1X103	2.8X104
2.4X101	6.0X10-1	1.6X101	2.5X104	6.6X105
5.4	2.0X10-1	5.4	1.2X102	3.2X103
1.1X103	6.0X10-3	1.6X10-1	5.8X101	1.6X103
8.1X101	8.0X10-4	2.2X10-2	1.5X10-1	4.1
5.4X102	2.0X10-3	5.4X10-2	4.0	1.1X102
1.9X102	7.0X10-4	1.9X10-2	5.9X10-2	1.6
1.4	3.0X10-3	8.1X10-2	2.0X101	5.4X102
1.1X103	4.0X10-2	1.1	1.1X103	2.9X104
2.7X10-2	1.0X10-3	2.7X10-2	3.1X102	8.5X103
2.7X102	6.0X10-1	1.6X101	1.2X10-3	3.3X10-2
5.4	2.0X10-1	5.4	4.9X106	1.3X108
1.1X103	2.0X10-2	5.4X10-1	7.5X102	2.0X104
5.4X101	1.0	2.7X101	6.1X102	1.7X104
1.1X103	1.0X10-2	2.7X10-1	1.2X102	3.3X103
2.4X102	1.0X10-3	2.7X10-2	1.9X10-3	5.2X101
5.4X102	2.0X10-3	5.4X10-2	3.0	8.1X101
2.4X102	9.0X10-4	2.4X10-2	6.4X10-3	1.7X10-1
2.4X102	9.0X10-4	2.4X10-2	1.1X10-2	3.1X10-1
8.1X101	1.0X10-3	2.7X10-2	3.4X10-6	9.3X10-5
5.4X10-1	3.0X10-4	8.1X10-3	1.6X10-4	4.2X10-3
1.4X101	5.0X10-1	1.4X101	1.1X105	3.1X106

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8.1	3.0X10-1	8.1	1.1X103	3.0X104
2.7X102	1.0X101	2.7X102	3.1X102	8.4X103
2.7X101	1.0	2.7X101	1.2X103	3.2X104
1.1X103	4.0X101	1.1X103	2.2X105	5.9X106
1.1X101	4.0X10-1	1.1X101	4.2X101	1.1X103
8.1X102	3.0X101	8.1X102	3.4X103	9.2X104
1.1X102	4.0	1.1X102	2.8X104	7.6X105
8.1X102	3.0X101	8.1X102	3.8X103	1.0X105
2.7X101	1.0	2.7X101	5.7X103	1.5X105
1.9X101	7.0X10-1	1.9X101	4.8X101	1.3X103
1.1X103	6.0X10-1	1.6X101	3.0X105	8.0X106
1.1X103	1.0	2.7X101	4.3X10-5	1.2X10-3
1.4X101	5.0X10-1	1.4X101	2.7X103	7.3X104
5.4X101	6.0X10-1	1.6X101	3.2	8.7X101
1.6X102	1.0	2.7X101	1.4X105	3.9X106
2.7X102	7.0X10-1	1.9X101	2.8X104	7.6X105
5.4X102	2.0X101	5.4X102	2.1X102	5.7X103
2.4X101	6.0X10-1	1.6X101	3.0X105	8.2X106
2.4X101	3.0X10-1	8.1	8.6X103	2.3X105
1.1X103	1.0	2.7X101	3.1X103	8.3X104
2.2X101	5.0X10-1	1.4X101	9.0X104	2.4X106
5.4X101	2.0	5.4X101	1.4X103	3.7X104
1.4X101	5.0X10-1	1.4X101	6.0X102	1.6X104

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5.4X102	2.0X101	5.4X102	3.5X102	9.4X103
5.4X101	7.0X10-1	1.9X101	6.1X104	1.6X106
1.9X101	7.0X10-1	1.9X101	6.1X104	1.6X106
2.7X101	1.0	2.7X101	6.5	1.8X102
2.2X101	8.0X10-1	2.2X101	8.2X104	2.2X106
2.4X101	6.0X10-1	1.6X101	9.8	2.6X102
5.4X102	3.0	8.1X101	1.8X101	4.9X102
1.9X101	7.0X10-1	1.9X101	2.0X103	5.5X104
2.7X101	6.0X10-1	1.6X101	3.5X106	9.5X107
8.1	3.0X10-1	8.1	2.7X105	7.3X106
1.1X103	4.0X101	1.1X103	8.8X101	2.4X103
2.4X101	9.0X10-1	2.4X101	1.8X103	5.0X104
1.1X103	2.0X10-1	5.4	7.4X10-4	2.0X10-2
1.9X102	3.0	8.1X101	2.2X104	6.0X105
1.4X101	5.0X10-1	1.4X101	1.5X106	4.1X107
1.1X101	4.0X10-1	1.1X101	1.1X105	3.1X106
1.4X101	5.0X10-1	1.4X101	6.9X102	1.9X104
5.4X102	2.0X10-3	5.4X10-2	1.2	3.2X101
2.7X102	9.0	2.4X102	1.3X102	3.5X103
8.1X101	6.0X10-1	1.6X101	3.9X104	1.1X106
1.4X101	5.0X10-1	1.4X101	2.6X102	7.1X103

76.....k...90....k....0....k...10....k...20....k...30....k...40....k...5

WIDETABLE

NOTE--Some parts of this form are wider than one screen. To view

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material that exceeds the width of this screen, use the right arrow

key. To return to the original screen, use the left arrow key.

Table A-1

A1 and A2 Values for Radionuclides

Symbol of radionuclide Element and atomic number A1 (TBq) A1 (Ci) A2 (TBq) A2 (Ci) Specific activity (TBq/g) (Ci/g)

A1 and A2 Values for Radionuclides

Symbol of radionuclide Element and atomic number A1 (TBq) A1 (Ci) A2 (TBq) A2 (Ci) Specific activity (TBq/g) (Ci/g)

Ac-225 (a) Actinium (89) 8.0X10-1 2.2X101 6.0X10-3 1.6X10-1 2.1X103 5.8X104

Ac-227 (a) 9.0X10-1 2.4X101 9.0X10-5 2.4X10-3 2.7 7.2X101

Ac-228 6.0X10-1 1.6X101 5.0X10-1 1.4X101 8.4X104 2.2X106

Ag-105 Silver (47) 2.0 5.4X101 2.0 5.4X101 1.1X103 3.0X104

Ag-108m (a) 7.0X10-1 1.9X101 7.0X10-1 1.9X101 9.7X10-1 2.6X101

Ag-110m (a) 4.0X10-1 1.1X101 4.0X10-1 1.1X101 1.8X102 4.7X103

Ag-111 2.0 5.4X101 6.0X10-1 1.6X101 5.8X103 1.6X105

Al-26 Aluminum (13) 1.0X10-1 2.7 1.0X10-1 2.7 7.0X10-4 1.9X10-2

Am-241 Americium (95) 1.0X101 2.7X102 1.0X10-3 2.7X10-2 1.3X10-1 3.4

Am-242m (a) 1.0X101 2.7X102 1.0X10-3 2.7X10-2 3.6X10-1 1.0X101

Am-243 (a) 5.0 1.4X102 1.0X10-3 2.7X10-2 7.4X10-3 2.0X10-1

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Ar-37 Argon (18) 4.0X101 1.1X103 4.0X101 1.1X103 3.7X103 9.9X104

Ar-39 4.0X101 1.1X103 2.0X101 5.4X102 1.3 3.4X101

Ar-41 3.0X10-1 8.1 3.0X10-1 8.1 1.5X106 4.2X107

As-72 Arsenic (33) 3.0X10-1 8.1 3.0X10-1 8.1 6.2X104 1.7X106

As-73 4.0X101 1.1X103 4.0X101 1.1X103 8.2X102 2.2X104

As-74 1.0 2.7X101 9.0X10-1 2.4X101 3.7X103 9.9X104

As-76 3.0X10-1 8.1 3.0X10-1 8.1 5.8X104 1.6X106

As-77 2.0X101 5.4X102 7.0X10-1 1.9X101 3.9X104 1.0X106

At-211 (a) Astatine (85) 2.0X101 5.4X102 5.0X10-1 1.4X101 7.6X104 2.1X106

Au-193 Gold (79) 7.0 1.9X102 2.0 5.4X101 3.4X104 9.2X105

Au-194 1.0 2.7X101 1.0 2.7X101 1.5X104 4.1X105

Au-195 1.0X101 2.7X102 6.0 1.6X102 1.4X102 3.7X103

Au-198 1.0 2.7X101 6.0X10-1 1.6X101 9.0X103 2.4X105

Au-199 1.0X101 2.7X102 6.0X10-1 1.6X101 7.7X103 2.1X105

Ba-131 (a) Barium (56) 2.0 5.4X101 2.0 5.4X101 3.1X103 8.4X104

Ba-133 3.0 8.1X101 3.0 8.1X101 9.4 2.6X102

Ba-133m 2.0X101 5.4X102 6.0X10-1 1.6X101 2.2X104 6.1X105

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Ba-140 (a) 5.0X10-1 1.4X101 3.0X10-1 8.1 2.7X103
7.3X104

Be-7 Beryllium (4) 2.0X101 5.4X102
2.0X101 5.4X102 1.3X104 3.5X105

Be-10 4.0X101 1.1X103 6.0X10-1 1.6X101 8.3X10-4
2.2X10-2

Bi-205 Bismuth (83) 7.0X10-1 1.9X101
7.0X10-1 1.9X101 1.5X103 4.2X104

Bi-206 3.0X10-1 8.1 3.0X10-1 8.1 3.8X103
1.0X105

Bi-207 7.0X10-1 1.9X101 7.0X10-1 1.9X101 1.9 5.2X101

Bi-210 1.0 2.7X101 6.0X10-1 1.6X101 4.6X103
1.2X105

Bi-210m (a) 6.0X10-1 1.6X101 2.0X10-2 5.4X10-1 2.1X10-
5 5.7X10-4

Bi-212 (a) 7.0X10-1 1.9X101 6.0X10-1 1.6X101 5.4X105
1.5X107

Bk-247 Berkelium (97) 8.0 2.2X102
8.0X10-4 2.2X10-2 3.8X10-2 1.0

Bk-249 (a) 4.0X101 1.1X103 3.0X10-1 8.1 6.1X101
1.6X103

Br-76 Bromine (35) 4.0X10-1 1.1X101
4.0X10-1 1.1X101 9.4X104 2.5X106

Br-77 3.0 8.1X101 3.0 8.1X101 2.6X104
7.1X105

Br-82 4.0X10-1 1.1X101 4.0X10-1 1.1X101 4.0X104
1.1X106

C-11 Carbon (6) 1.0 2.7X101
6.0X10-1 1.6X101 3.1X107 8.4X108

C-14 4.0X101 1.1X103 3.0 8.1X101 1.6X10-1 4.5

Ca-41 Calcium (20) Unlimited Unlimited
Unlimited Unlimited 3.1X10-3 8.5X10-2

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Ca-45 4.0X101 1.1X103 1.0 2.7X101 6.6X102
1.8X104

Ca-47 (a) 3.0 8.1X101 3.0X10-1 8.1 2.3X104
6.1X105

Cd-109 Cadmium (48) 3.0X101 8.1X102
2.0 5.4X101 9.6X101 2.6X103

Cd-113m 4.0X101 1.1X103 5.0X10-1 1.4X101 8.3 2.2X102

Cd-115 (a) 3.0 8.1X101 4.0X10-1 1.1X101 1.9X104
5.1X105

Cd-115m 5.0X10-1 1.4X101 5.0X10-1 1.4X101 9.4X102
2.5X104

Ce-139 Cerium (58) 7.0 1.9X102
2.0 5.4X101 2.5X102 6.8X103

Ce-141 2.0X101 5.4X102 6.0X10-1 1.6X101 1.1X103
2.8X104

Ce-143 9.0X10-1 2.4X101 6.0X10-1 1.6X101 2.5X104
6.6X105

Ce-144 (a) 2.0X10-1 5.4 2.0X10-1 5.4 1.2X102
3.2X103

Cf-248 Californium (98) 4.0X101
1.1X103 6.0X10-3 1.6X10-1 5.8X101 1.6X103

Cf-249 3.0 8.1X101 8.0X10-4 2.2X10-2 1.5X10-1 4.1

Cf-250 2.0X101 5.4X102 2.0X10-3 5.4X10-2 4.0 1.1X102

Cf-251 7.0 1.9X102 7.0X10-4 1.9X10-2 5.9X10-2 1.6

Cf-252 (h) 5.0X10-2 1.4 3.0X10-3 8.1X10-2 2.0X101
5.4X102

Cf-253 (a) 4.0X101 1.1X103 4.0X10-2 1.1 1.1X103
2.9X104

Cf-254 1.0X10-3 2.7X10-2 1.0X10-3 2.7X10-2 3.1X102
8.5X103

Cl-36 Chlorine (17) 1.0X101 2.7X102

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6.0X10-1 1.6X101 1.2X10-3 3.3X10-2

Cl-38 2.0X10-1 5.4 2.0X10-1 5.4 4.9X106
1.3X108

Cm-240 Curium (96) 4.0X101 1.1X103
2.0X10-2 5.4X10-1 7.5X102 2.0X104

Cm-241 2.0 5.4X101 1.0 2.7X101 6.1X102
1.7X104

Cm-242 4.0X101 1.1X103 1.0X10-2 2.7X10-1 1.2X102
3.3X103

Cm-243 9.0 2.4X102 1.0X10-3 2.7X10-2 1.9X10-3
5.2X101

Cm-244 2.0X101 5.4X102 2.0X10-3 5.4X10-2 3.0 8.1X101

Cm-245 9.0 2.4X102 9.0X10-4 2.4X10-2 6.4X10-3
1.7X10-1

Cm-246 9.0 2.4X102 9.0X10-4 2.4X10-2 1.1X10-2
3.1X10-1

Cm-247 (a) 3.0 8.1X101 1.0X10-3 2.7X10-2 3.4X10-
6 9.3X10-5

Cm-248 2.0X10-2 5.4X10-1 3.0X10-4 8.1X10-3 1.6X10-4
4.2X10-3

Co-55 Cobalt (27) 5.0X10-1 1.4X101
5.0X10-1 1.4X101 1.1X105 3.1X106

Co-56 3.0X10-1 8.1 3.0X10-1 8.1 1.1X103
3.0X104

Co-57 1.0X101 2.7X102 1.0X101 2.7X102 3.1X102
8.4X103

Co-58 1.0 2.7X101 1.0 2.7X101 1.2X103
3.2X104

Co-58m 4.0X101 1.1X103 4.0X101 1.1X103 2.2X105
5.9X106

Co-60 4.0X10-1 1.1X101 4.0X10-1 1.1X101 4.2X101
1.1X103

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Cr-51 Chromium (24) 3.0X101 8.1X102
3.0X101 8.1X102 3.4X103 9.2X104

Cs-129 Cesium (55) 4.0 1.1X102
4.0 1.1X102 2.8X104 7.6X105

Cs-131 3.0X101 8.1X102 3.0X101 8.1X102 3.8X103
1.0X105

Cs-132 1.0 2.7X101 1.0 2.7X101 5.7X103
1.5X105

Cs-134 7.0X10-1 1.9X101 7.0X10-1 1.9X101 4.8X101
1.3X103

Cs-134m 4.0X101 1.1X103 6.0X10-1 1.6X101 3.0X105
8.0X106

Cs-135 4.0X101 1.1X103 1.0 2.7X101 4.3X10-5
1.2X10-3

Cs-136 5.0X10-1 1.4X101 5.0X10-1 1.4X101 2.7X103
7.3X104

Cs-137 (a) 2.0 5.4X101 6.0X10-1 1.6X101 3.2
8.7X101

Cu-64 Copper (29) 6.0 1.6X102
1.0 2.7X101 1.4X105 3.9X106

Cu-67 1.0X101 2.7X102 7.0X10-1 1.9X101 2.8X104
7.6X105

Dy-159 Dysprosium (66) 2.0X101 5.4X102
2.0X101 5.4X102 2.1X102 5.7X103

Dy-165 9.0X10-1 2.4X101 6.0X10-1 1.6X101 3.0X105
8.2X106

Dy-166 (a) 9.0X10-1 2.4X101 3.0X10-1 8.1 8.6X103
2.3X105

Er-169 Erbium (68) 4.0X101 1.1X103
1.0 2.7X101 3.1X103 8.3X104

Er-171 8.0X10-1 2.2X101 5.0X10-1 1.4X101 9.0X104
2.4X106

Eu-147 Europium (63) 2.0 5.4X101

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2.0 5.4X101 1.4X103 3.7X104

Eu-148 5.0X10-1 1.4X101 5.0X10-1 1.4X101 6.0X102
1.6X104

Eu-149 2.0X101 5.4X102 2.0X101 5.4X102 3.5X102
9.4X103

Eu-150 (short lived) 2.0 5.4X101 7.0X10-1 1.9X101
6.1X104 1.6X106

Eu-150 (long lived) 7.0X10-1 1.9X101 7.0X10-1 1.9X101
6.1X104 1.6X106

Eu-152 1.0 2.7X101 1.0 2.7X101 6.5 1.8X102

Eu-152m 8.0X10-1 2.2X101 8.0X10-1 2.2X101 8.2X104
2.2X106

Eu-154 9.0X10-1 2.4X101 6.0X10-1 1.6X101 9.8 2.6X102

Eu-155 2.0X101 5.4X102 3.0 8.1X101 1.8X101
4.9X102

Eu-156 7.0X10-1 1.9X101 7.0X10-1 1.9X101 2.0X103
5.5X104

F-18 Fluorine (9) 1.0 2.7X101
6.0X10-1 1.6X101 3.5X106 9.5X107

Fe-52 (a) Iron (26) 3.0X10-1 8.1
3.0X10-1 8.1 2.7X105 7.3X106

Fe-55 4.0X101 1.1X103 4.0X101 1.1X103 8.8X101
2.4X103

Fe-59 9.0X10-1 2.4X101 9.0X10-1 2.4X101 1.8X103
5.0X104

Fe-60 (a) 4.0X101 1.1X103 2.0X10-1 5.4 7.4X10- 4
2.0X10-2

Ga-67 Gallium (31) 7.0 1.9X102
3.0 8.1X101 2.2X104 6.0X105

Ga-68 5.0X10-1 1.4X101 5.0X10-1 1.4X101 1.5X106
4.1X107

Ga-72 4.0X10-1 1.1X101 4.0X10-1 1.1X101 1.1X105

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3.1X106

Gd-146 (a) Gadolinium (64) 5.0X10-1 1.4X101
5.0X10-1 1.4X101 6.9X102 1.9X104

Gd-148 2.0X101 5.4X102 2.0X10-3 5.4X10-2 1.2 3.2X101

Gd-153 1.0X101 2.7X102 9.0 2.4X102 1.3X102
3.5X103

Gd-159 3.0 8.1X101 6.0X10-1 1.6X101 3.9X104
1.1X106

Ge-68 (a) Germanium (32) 5.0X10-1 1.4X101
5.0X10-1 1.4X101 2.6X102 7.1X103

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

Ge-71		4.0X101	1.1X103	4.0X101
Ge-77		3.0X10-1	8.1	3.0X10-1
Hf-172 (a)	Hafnium (72)	6.0X10-1	1.6X101	6.0X10-1
Hf-175		3.0	8.1X101	3.0
Hf-181		2.0	5.4X101	5.0X10-1
Hf-182		Unlimited	Unlimited	Unlimited
Hg-194 (a)	Mercury (80)	1.0	2.7X101	1.0
Hg-195m (a)		3.0	8.1X101	7.0X10-1
Hg-197		2.0X101	5.4X102	1.0X101
Hg-197m		1.0X101	2.7X102	4.0X10-1
Hg-203		5.0	1.4X102	1.0
Ho-166	Holmium (67)	4.0X10-1	1.1X101	4.0X10-1
Ho-166m		6.0X10-1	1.6X101	5.0X10-1
I-123	Iodine (53)	6.0	1.6X102	3.0
I-124		1.0	2.7X101	1.0
I-125		2.0X101	5.4X102	3.0
I-126		2.0	5.4X101	1.0
I-129		Unlimited	Unlimited	Unlimited
I-131		3.0	8.1X101	7.0X10-1
I-132		4.0X10-1	1.1X101	4.0X10-1
I-133		7.0X10-1	1.9X101	6.0X10-1
I-134		3.0X10-1	8.1	3.0X10-1

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I-135 (a)		6.0X10-1	1.6X101	6.0X10-1
In-111	Indium (49)	3.0	8.1X101	3.0
In-113m		4.0	1.1X102	2.0
In-114m (a)		1.0X101	2.7X102	5.0X10-1
In-115m		7.0	1.9X102	1.0
Ir-189 (a)	Iridium (77)	1.0X101	2.7X102	1.0X101
Ir-190		7.0X10-1	1.9X101	7.0X10-1
Ir-192 (c)		1.0	2.7X101	6.0X10-1
Ir-194		3.0X10-1	8.1	3.0X10-1
K-40	Potassium (19)	9.0X10-1	2.4X101	9.0X10-1
K-42		2.0X10-1	5.4	2.0X10-1
K-43		7.0X10-1	1.9X101	6.0X10-1
Kr-81	Krypton (36)	4.0X101	1.1X103	4.0X101
Kr-85		1.0X101	2.7X102	1.0X101
Kr-85m		8.0	2.2X102	3.0
Kr-87		2.0X10-1	5.4	2.0X10-1
La-137	Lanthanum (57)	3.0X101	8.1X102	6.0
La-140		4.0X10-1	1.1X101	4.0X10-1
Lu-172	Lutetium (71)	6.0X10-1	1.6X101	6.0X10-1
Lu-173		8.0	2.2X102	8.0
Lu-174		9.0	2.4X102	9.0
Lu-174m		2.0X101	5.4X102	1.0X101
Lu-177		3.0X101	8.1X102	7.0X10-1
Mg-28 (a)	Magnesium (12)	3.0X10-1	8.1	3.0X10-1
Mn-52	Manganese (25)	3.0X10-1	8.1	3.0X10-1
Mn-53		Unlimited	Unlimited	Unlimited
Mn-54		1.0	2.7X101	1.0
Mn-56		3.0X10-1	8.1	3.0X10-1
Mo-93	Molybdenum (42)	4.0X101	1.1X103	2.0X101
Mo-99 (a) (i)		1.0	2.7X101	6.0X10-1
N-13	Nitrogen (7)	9.0X10-1	2.4X101	6.0X10-1
Na-22	Sodium (11)	5.0X10-1	1.4X101	5.0X10-1
Na-24		2.0X10-1	5.4	2.0X10-1
Nb-93m	Niobium (41)	4.0X101	1.1X103	3.0X101
Nb-94		7.0X10-1	1.9X101	7.0X10-1
Nb-95		1.0	2.7X101	1.0
Nb-97		9.0X10-1	2.4X101	6.0X10-1
Nd-147	Neodymium (60)	6.0	1.6X102	6.0X10-1
Nd-149		6.0X10-1	1.6X101	5.0X10-1
Ni-59	Nickel (28)	Unlimited	Unlimited	Unlimited
Ni-63		4.0X101	1.1X103	3.0X101
Ni-65		4.0X10-1	1.1X101	4.0X10-1
Np-235	Neptunium (93)	4.0X101	1.1X103	4.0X101
Np-236 (short-lived)		2.0X101	5.4X102	2.0
Np-236 (long-lived)		9.0X100	2.4X102	2.0X10-2
Np-237		2.0X101	5.4X102	2.0X10-3
Np-239		7.0	1.9X102	4.0X10-1
Os-185	Osmium (76)	1.0	2.7X101	1.0
Os-191		1.0X101	2.7X102	2.0
Os-191m		4.0X101	1.1X103	3.0X101
Os-193		2.0	5.4X101	6.0X10-1
Os-194 (a)		3.0X10-1	8.1	3.0X10-1
P-32	Phosphorus (15)	5.0X10-1	1.4X101	5.0X10-1

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P-33		4.0X101	1.1X103	1.0
Pa-230 (a)	Protactinium (91)	2.0	5.4X101	7.0X10-2
Pa-231		4.0	1.1X102	4.0X10-4
Pa-233		5.0	1.4X102	7.0X10-1
Pb-201	Lead (82)	1.0	2.7X101	1.0
Pb-202		4.0X101	1.1X103	2.0X101
Pb-203		4.0	1.1X102	3.0
Pb-205		Unlimited	Unlimited	Unlimited
Pb-210 (a)		1.0	2.7X101	5.0X10-2
Pb-212 (a)		7.0X10-1	1.9X101	2.0X10-1
Pd-103 (a)	Palladium (46)	4.0X101	1.1X103	4.0X101
Pd-107		Unlimited	Unlimited	Unlimited
Pd-109		2.0	5.4X101	5.0X10-1
Pm-143	Promethium (61)	3.0	8.1X101	3.0
Pm-144		7.0X10-1	1.9X101	7.0X10-1
Pm-145		3.0X101	8.1X102	1.0X101
Pm-147		4.0X101	1.1X103	2.0
Pm-148m (a)		8.0X10-1	2.2X101	7.0X10-1
Pm-149		2.0	5.4X101	6.0X10-1
Pm-151		2.0	5.4X101	6.0X10-1
Po-210	Polonium (84)	4.0X101	1.1X103	2.0X10-2
Pr-142	Praseodymium (59)	4.0X10-1	1.1X101	4.0X10-1
Pr-143		3.0	8.1X101	6.0X10-1
Pt-188 (a)	Platinum (78)	1.0	2.7X101	8.0X10-1
Pt-191		4.0	1.1X102	3.0
Pt-193		4.0X101	1.1X103	4.0X101
Pt-193m		4.0X101	1.1X103	5.0X10-1
Pt-195m		1.0X101	2.7X102	5.0X10-1
Pt-197		2.0X101	5.4X102	6.0X10-1
Pt-197m		1.0X101	2.7X102	6.0X10-1
Pu-236	Plutonium (94)	3.0X101	8.1X102	3.0X10-3
Pu-237		2.0X101	5.4X102	2.0X101
Pu-238		1.0X101	2.7X102	1.0X10-3
Pu-239		1.0X101	2.7X102	1.0X10-3
Pu-240		1.0X101	2.7X102	1.0X10-3
Pu-241 (a)		4.0X101	1.1X103	6.0X10-2
Pu-242		1.0X101	2.7X102	1.0X10-3
Pu-244 (a)		4.0X10-1	1.1X101	1.0X10-3
Ra-223 (a)	Radium (88)	4.0X10-1	1.1X101	7.0X10-3
Ra-224 (a)		4.0X10-1	1.1X101	2.0X10-2
Ra-225 (a)		2.0X10-1	5.4	4.0X10-3
Ra-226 (a)		2.0X10-1	5.4	3.0X10-3
Ra-228 (a)		6.0X10-1	1.6X101	2.0X10-2
Rb-81	Rubidium (37)	2.0	5.4X101	8.0X10-1
Rb-83 (a)		2.0	5.4X101	2.0
Rb-84		1.0	2.7X101	1.0
Rb-86		5.0X10-1	1.4X101	5.0X10-1
Rb-87		Unlimited	Unlimited	Unlimited
Rb(nat)		Unlimited	Unlimited	Unlimited
Re-184	Rhenium (75)	1.0	2.7X101	1.0
Re-184m		3.0	8.1X101	1.0

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+...

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****
******* This is piece 2. -- It begins at character 79 of table line 1. ********
****

1.1X103	5.8X103	1.6X105
8.1	1.3X105	3.6X106
1.6X101	4.1X101	1.1X103
8.1X101	3.9X102	1.1X104
1.4X101	6.3X102	1.7X104
Unlimited	8.1X10-6	2.2X10-4
2.7X101	1.3X10-1	3.5
1.9X101	1.5X104	4.0X105
2.7X102	9.2X103	2.5X105
1.1X101	2.5X104	6.7X105
2.7X101	5.1X102	1.4X104
1.1X101	2.6X104	7.0X105
1.4X101	6.6X10-2	1.8
8.1X101	7.1X104	1.9X106
2.7X101	9.3X103	2.5X105
8.1X101	6.4X102	1.7X104
2.7X101	2.9X103	8.0X104
Unlimited	6.5X10-6	1.8X10-4
1.9X101	4.6X103	1.2X105
1.1X101	3.8X105	1.0X107
1.6X101	4.2X104	1.1X106
8.1	9.9X105	2.7X107
1.6X101	1.3X105	3.5X106
8.1X101	1.5X104	4.2X105
5.4X101	6.2X105	1.7X107
1.4X101	8.6X102	2.3X104
2.7X101	2.2X105	6.1X106
2.7X102	1.9X103	5.2X104
1.9X101	2.3X103	6.2X104
1.6X101	3.4X102	9.2X103
8.1	3.1X104	8.4X105
2.4X101	2.4X10-7	6.4X10-6
5.4	2.2X105	6.0X106
1.6X101	1.2X105	3.3X106
1.1X103	7.8X10-4	2.1X10-2
2.7X102	1.5X101	3.9X102
8.1X101	3.0X105	8.2X106
5.4	1.0X106	2.8X107
1.6X102	1.6X10-3	4.4X10-2
1.1X101	2.1X104	5.6X105
1.6X101	4.2X103	1.1X105
2.2X102	5.6X101	1.5X103
2.4X102	2.3X101	6.2X102
2.7X102	2.0X102	5.3X103
1.9X101	4.1X103	1.1X105
8.1	2.0X105	5.4X106
8.1	1.6X104	4.4X105

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Unlimited	6.8X10-5	1.8X10-3
2.7X101	2.9X102	7.7X103
8.1	8.0X105	2.2X107
5.4X102	4.1X10-2	1.1
1.6X101	1.8X104	4.8X105
1.6X101	5.4X107	1.5X109
1.4X101	2.3X102	6.3X103
5.4	3.2X105	8.7X106
8.1X102	8.8	2.4X102
1.9X101	6.9X10-3	1.9X10-1
2.7X101	1.5X103	3.9X104
1.6X101	9.9X105	2.7X107
1.6X101	3.0X103	8.1X104
1.4X101	4.5X105	1.2X107
Unlimited	3.0X10-3	8.0X10-2
8.1X102	2.1	5.7X101
1.1X101	7.1X105	1.9X107
1.1X103	5.2X101	1.4X103
5.4X101	4.7X10-4	1.3X10-2
5.4X10-1	4.7X10-4	1.3X10-2
5.4X10-2	2.6X10-5	7.1X10-4
1.1X101	8.6X103	2.3X105
2.7X101	2.8X102	7.5X103
5.4X101	1.6X103	4.4X104
8.1X102	4.6X104	1.3X106
1.6X101	2.0X104	5.3X105
8.1	1.1X101	3.1X102
1.4X101	1.1X104	2.9X105
2.7X101	5.8X103	1.6X105
1.9	1.2X103	3.3X104
1.1X10-2	1.7X10-3	4.7X10-2
1.9X101	7.7X102	2.1X104
2.7X101	6.2X104	1.7X106
5.4X102	1.2X10-4	3.4X10-3
8.1X101	1.1X104	3.0X105
Unlimited	4.5X10-6	1.2X10-4
1.4	2.8	7.6X101
5.4	5.1X104	1.4X106
1.1X103	2.8X103	7.5X104
Unlimited	1.9X10-5	5.1X10-4
1.4X101	7.9X104	2.1X106
8.1X101	1.3X102	3.4X103
1.9X101	9.2X101	2.5X103
2.7X102	5.2	1.4X102
5.4X101	3.4X101	9.3X102
1.9X101	7.9X102	2.1X104
1.6X101	1.5X104	4.0X105
1.6X101	2.7X104	7.3X105
5.4X10-1	1.7X102	4.5X103
1.1X101	4.3X104	1.2X106
1.6X101	2.5X103	6.7X104
2.2X101	2.5X103	6.8X104
8.1X101	8.7X103	2.4X105

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 1.1X103     1.4        3.7X101
 1.4X101     5.8X103    1.6X105
 1.4X101     6.2X103    1.7X105
 1.6X101     3.2X104    8.7X105
 1.6X101     3.7X105    1.0X107
 8.1X10-2    2.0X101    5.3X102
 5.4X102     4.5X102    1.2X104
 2.7X10-2    6.3X10-1   1.7X101
 2.7X10-2    2.3X10-3   6.2X10-2
 2.7X10-2    8.4X10-3   2.3X10-1
 1.6         3.8        1.0X102
 2.7X10-2    1.5X10-4   3.9X10-3
 2.7X10-2    6.7X10-7   1.8X10-5
 1.9X10-1    1.9X103    5.1X104
 5.4X10-1    5.9X103    1.6X105
 1.1X10-1    1.5X103    3.9X104
 8.1X10-2    3.7X10-2   1.0
 5.4X10-1    1.0X101    2.7X102
 2.2X101     3.1X105    8.4X106
 5.4X101     6.8X102    1.8X104
 2.7X101     1.8X103    4.7X104
 1.4X101     3.0X103    8.1X104
 Unlimited   3.2X10-9   8.6X10-8
 Unlimited   6.7X106    1.8X108
 2.7X101     6.9X102    1.9X104
 2.7X101     1.6X102    4.3X103
79....+...90....+....0....+...10
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[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

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 ******************************************************************************

 ******** This is piece 1. -- It begins at character 1 of table line 1. ********

 ******************************************************************************

 Ge-71                                        4.0X101  1.1X103     4.0X101

 Ge-77                                        3.0X10-1  8.1         3.0X10-1

 Hf-172 (a)              Hafnium (72)         6.0X10-1  1.6X101     6.0X10-1

 Hf-175                                       3.0  8.1X101     3.0
```

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Hf-181		2.0	5.4X101	5.0X10-1
Hf-182		Unlimited	Unlimited	Unlimited
Hg-194 (a)	Mercury (80)	1.0	2.7X101	1.0
Hg-195m (a)		3.0	8.1X101	7.0X10-1
Hg-197		2.0X101	5.4X102	1.0X101
Hg-197m		1.0X101	2.7X102	4.0X10-1
Hg-203		5.0	1.4X102	1.0
Ho-166	Holmium (67)	4.0X10-1	1.1X101	4.0X10-1
Ho-166m		6.0X10-1	1.6X101	5.0X10-1
I-123	Iodine (53)	6.0	1.6X102	3.0
I-124		1.0	2.7X101	1.0
I-125		2.0X101	5.4X102	3.0
I-126		2.0	5.4X101	1.0
I-129		Unlimited	Unlimited	Unlimited
I-131		3.0	8.1X101	7.0X10-1
I-132		4.0X10-1	1.1X101	4.0X10-1
I-133		7.0X10-1	1.9X101	6.0X10-1
I-134		3.0X10-1	8.1	3.0X10-1
I-135 (a)		6.0X10-1	1.6X101	6.0X10-1
In-111	Indium (49)	3.0	8.1X101	3.0
In-113m		4.0	1.1X102	2.0
In-114m (a)		1.0X101	2.7X102	5.0X10-1
In-115m		7.0	1.9X102	1.0
Ir-189 (a)	Iridium (77)	1.0X101	2.7X102	1.0X101

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Ir-190		7.0X10-1	1.9X101	7.0X10-1
Ir-192 (c)		1.0	2.7X101	6.0X10-1
Ir-194		3.0X10-1	8.1	3.0X10-1
K-40	Potassium (19)	9.0X10-1	2.4X101	9.0X10-1
K-42		2.0X10-1	5.4	2.0X10-1
K-43		7.0X10-1	1.9X101	6.0X10-1
Kr-81	Krypton (36)	4.0X101	1.1X103	4.0X101
Kr-85		1.0X101	2.7X102	1.0X101
Kr-85m		8.0	2.2X102	3.0
Kr-87		2.0X10-1	5.4	2.0X10-1
La-137	Lanthanum (57)	3.0X101	8.1X102	6.0
La-140		4.0X10-1	1.1X101	4.0X10-1
Lu-172	Lutetium (71)	6.0X10-1	1.6X101	6.0X10-1
Lu-173		8.0	2.2X102	8.0
Lu-174		9.0	2.4X102	9.0
Lu-174m		2.0X101	5.4X102	1.0X101
Lu-177		3.0X101	8.1X102	7.0X10-1
Mg-28 (a)	Magnesium (12)	3.0X10-1	8.1	3.0X10-1
Mn-52	Manganese (25)	3.0X10-1	8.1	3.0X10-1
Mn-53		Unlimited	Unlimited	Unlimited
Mn-54		1.0	2.7X101	1.0
Mn-56		3.0X10-1	8.1	3.0X10-1
Mo-93	Molybdenum (42)	4.0X101	1.1X103	2.0X101

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Mo-99 (a) (i)		1.0	2.7X101	6.0X10-1
N-13	Nitrogen (7)	9.0X10-1	2.4X101	6.0X10-1
Na-22	Sodium (11)	5.0X10-1	1.4X101	5.0X10-1
Na-24		2.0X10-1	5.4	2.0X10-1
Nb-93m	Niobium (41)	4.0X101	1.1X103	3.0X101
Nb-94		7.0X10-1	1.9X101	7.0X10-1
Nb-95		1.0	2.7X101	1.0
Nb-97		9.0X10-1	2.4X101	6.0X10-1
Nd-147	Neodymium (60)	6.0	1.6X102	6.0X10-1
Nd-149		6.0X10-1	1.6X101	5.0X10-1
Ni-59	Nickel (28)	Unlimited	Unlimited	Unlimited
Ni-63		4.0X101	1.1X103	3.0X101
Ni-65		4.0X10-1	1.1X101	4.0X10-1
Np-235	Neptunium (93)	4.0X101	1.1X103	4.0X101
Np-236 (short-lived)		2.0X101	5.4X102	2.0
Np-236 (long-lived)		9.0X100	2.4X102	2.0X10-2
Np-237		2.0X101	5.4X102	2.0X10-3
Np-239		7.0	1.9X102	4.0X10-1
Os-185	Osmium (76)	1.0	2.7X101	1.0
Os-191		1.0X101	2.7X102	2.0
Os-191m		4.0X101	1.1X103	3.0X101
Os-193		2.0	5.4X101	6.0X10-1
Os-194 (a)		3.0X10-1	8.1	3.0X10-1
P-32	Phosphorus (15)	5.0X10-1	1.4X101	5.0X10-1

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P-33		4.0X101	1.1X103	1.0
Pa-230 (a)	Protactinium (91)	2.0	5.4X101	7.0X10-2
Pa-231		4.0	1.1X102	4.0X10-4
Pa-233		5.0	1.4X102	7.0X10-1
Pb-201	Lead (82)	1.0	2.7X101	1.0
Pb-202		4.0X101	1.1X103	2.0X101
Pb-203		4.0	1.1X102	3.0
Pb-205		Unlimited	Unlimited	Unlimited
Pb-210 (a)		1.0	2.7X101	5.0X10-2
Pb-212 (a)		7.0X10-1	1.9X101	2.0X10-1
Pd-103 (a)	Palladium (46)	4.0X101	1.1X103	4.0X101
Pd-107		Unlimited	Unlimited	Unlimited
Pd-109		2.0	5.4X101	5.0X10-1
Pm-143	Promethium (61)	3.0	8.1X101	3.0
Pm-144		7.0X10-1	1.9X101	7.0X10-1
Pm-145		3.0X101	8.1X102	1.0X101
Pm-147		4.0X101	1.1X103	2.0
Pm-148m (a)		8.0X10-1	2.2X101	7.0X10-1
Pm-149		2.0	5.4X101	6.0X10-1
Pm-151		2.0	5.4X101	6.0X10-1
Po-210	Polonium (84)	4.0X101	1.1X103	2.0X10-2
Pr-142	Praseodymium (59)	4.0X10-1	1.1X101	4.0X10-1
Pr-143		3.0	8.1X101	6.0X10-1

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Pt-188 (a)	Platinum (78)	1.0	2.7X101	8.0X10-1
Pt-191		4.0	1.1X102	3.0
Pt-193		4.0X101	1.1X103	4.0X101
Pt-193m		4.0X101	1.1X103	5.0X10-1
Pt-195m		1.0X101	2.7X102	5.0X10-1
Pt-197		2.0X101	5.4X102	6.0X10-1
Pt-197m		1.0X101	2.7X102	6.0X10-1
Pu-236	Plutonium (94)	3.0X101	8.1X102	3.0X10-3
Pu-237		2.0X101	5.4X102	2.0X101
Pu-238		1.0X101	2.7X102	1.0X10-3
Pu-239		1.0X101	2.7X102	1.0X10-3
Pu-240		1.0X101	2.7X102	1.0X10-3
Pu-241 (a)		4.0X101	1.1X103	6.0X10-2
Pu-242		1.0X101	2.7X102	1.0X10-3
Pu-244 (a)		4.0X10-1	1.1X101	1.0X10-3
Ra-223 (a)	Radium (88)	4.0X10-1	1.1X101	7.0X10-3
Ra-224 (a)		4.0X10-1	1.1X101	2.0X10-2
Ra-225 (a)		2.0X10-1	5.4	4.0X10-3
Ra-226 (a)		2.0X10-1	5.4	3.0X10-3
Ra-228 (a)		6.0X10-1	1.6X101	2.0X10-2
Rb-81	Rubidium (37)	2.0	5.4X101	8.0X10-1
Rb-83 (a)		2.0	5.4X101	2.0
Rb-84		1.0	2.7X101	1.0
Rb-86		5.0X10-1	1.4X101	5.0X10-1

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Rb-87		Unlimited	Unlimited	Unlimited
Rb(nat)		Unlimited	Unlimited	Unlimited
Re-184	Rhenium (75)	1.0	2.7X101	1.0
Re-184m		3.0	8.1X101	1.0

1...k...10....k...20....k...30....k...40....k...50....k...60....k...70....k...

****

******* This is piece 2. -- It begins at character 79 of table line 1. ********

****

1.1X103	5.8X103	1.6X105
8.1	1.3X105	3.6X106
1.6X101	4.1X101	1.1X103
8.1X101	3.9X102	1.1X104
1.4X101	6.3X102	1.7X104
Unlimited	8.1X10-6	2.2X10-4
2.7X101	1.3X10-1	3.5
1.9X101	1.5X104	4.0X105
2.7X102	9.2X103	2.5X105
1.1X101	2.5X104	6.7X105
2.7X101	5.1X102	1.4X104
1.1X101	2.6X104	7.0X105
1.4X101	6.6X10-2	1.8
8.1X101	7.1X104	1.9X106
2.7X101	9.3X103	2.5X105

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8.1X101	6.4X102	1.7X104
2.7X101	2.9X103	8.0X104
Unlimited	6.5X10-6	1.8X10-4
1.9X101	4.6X103	1.2X105
1.1X101	3.8X105	1.0X107
1.6X101	4.2X104	1.1X106
8.1	9.9X105	2.7X107
1.6X101	1.3X105	3.5X106
8.1X101	1.5X104	4.2X105
5.4X101	6.2X105	1.7X107
1.4X101	8.6X102	2.3X104
2.7X101	2.2X105	6.1X106
2.7X102	1.9X103	5.2X104
1.9X101	2.3X103	6.2X104
1.6X101	3.4X102	9.2X103
8.1	3.1X104	8.4X105
2.4X101	2.4X10-7	6.4X10-6
5.4	2.2X105	6.0X106
1.6X101	1.2X105	3.3X106
1.1X103	7.8X10-4	2.1X10-2
2.7X102	1.5X101	3.9X102
8.1X101	3.0X105	8.2X106
5.4	1.0X106	2.8X107
1.6X102	1.6X10-3	4.4X10-2

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1.1X101	2.1X104	5.6X105
1.6X101	4.2X103	1.1X105
2.2X102	5.6X101	1.5X103
2.4X102	2.3X101	6.2X102
2.7X102	2.0X102	5.3X103
1.9X101	4.1X103	1.1X105
8.1	2.0X105	5.4X106
8.1	1.6X104	4.4X105
Unlimited	6.8X10-5	1.8X10-3
2.7X101	2.9X102	7.7X103
8.1	8.0X105	2.2X107
5.4X102	4.1X10-2	1.1
1.6X101	1.8X104	4.8X105
1.6X101	5.4X107	1.5X109
1.4X101	2.3X102	6.3X103
5.4	3.2X105	8.7X106
8.1X102	8.8	2.4X102
1.9X101	6.9X10-3	1.9X10-1
2.7X101	1.5X103	3.9X104
1.6X101	9.9X105	2.7X107
1.6X101	3.0X103	8.1X104
1.4X101	4.5X105	1.2X107
Unlimited	3.0X10-3	8.0X10-2

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8.1X102	2.1	5.7X101
1.1X101	7.1X105	1.9X107
1.1X103	5.2X101	1.4X103
5.4X101	4.7X10-4	1.3X10-2
5.4X10-1	4.7X10-4	1.3X10-2
5.4X10-2	2.6X10-5	7.1X10-4
1.1X101	8.6X103	2.3X105
2.7X101	2.8X102	7.5X103
5.4X101	1.6X103	4.4X104
8.1X102	4.6X104	1.3X106
1.6X101	2.0X104	5.3X105
8.1	1.1X101	3.1X102
1.4X101	1.1X104	2.9X105
2.7X101	5.8X103	1.6X105
1.9	1.2X103	3.3X104
1.1X10-2	1.7X10-3	4.7X10-2
1.9X101	7.7X102	2.1X104
2.7X101	6.2X104	1.7X106
5.4X102	1.2X10-4	3.4X10-3
8.1X101	1.1X104	3.0X105
Unlimited	4.5X10-6	1.2X10-4
1.4	2.8	7.6X101
5.4	5.1X104	1.4X106
1.1X103	2.8X103	7.5X104

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Unlimited	1.9X10-5	5.1X10-4
1.4X101	7.9X104	2.1X106
8.1X101	1.3X102	3.4X103
1.9X101	9.2X101	2.5X103
2.7X102	5.2	1.4X102
5.4X101	3.4X101	9.3X102
1.9X101	7.9X102	2.1X104
1.6X101	1.5X104	4.0X105
1.6X101	2.7X104	7.3X105
5.4X10-1	1.7X102	4.5X103
1.1X101	4.3X104	1.2X106
1.6X101	2.5X103	6.7X104
2.2X101	2.5X103	6.8X104
8.1X101	8.7X103	2.4X105
1.1X103	1.4	3.7X101
1.4X101	5.8X103	1.6X105
1.4X101	6.2X103	1.7X105
1.6X101	3.2X104	8.7X105
1.6X101	3.7X105	1.0X107
8.1X10-2	2.0X101	5.3X102
5.4X102	4.5X102	1.2X104
2.7X10-2	6.3X10-1	1.7X101
2.7X10-2	2.3X10-3	6.2X10-2

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2.7X10-2 8.4X10-3 2.3X10-1

1.6 3.8 1.0X102

2.7X10-2 1.5X10-4 3.9X10-3

2.7X10-2 6.7X10-7 1.8X10-5

1.9X10-1 1.9X103 5.1X104

5.4X10-1 5.9X103 1.6X105

1.1X10-1 1.5X103 3.9X104

8.1X10-2 3.7X10-2 1.0

5.4X10-1 1.0X101 2.7X102

2.2X101 3.1X105 8.4X106

5.4X101 6.8X102 1.8X104

2.7X101 1.8X103 4.7X104

1.4X101 3.0X103 8.1X104

Unlimited 3.2X10-9 8.6X10-8

Unlimited 6.7X106 1.8X108

2.7X101 6.9X102 1.9X104

2.7X101 1.6X102 4.3X103

79....k...90....k....0....k...10

WIDETABLE

NOTE--Some parts of this form are wider than one screen. To view material that exceeds the width of this screen, use the right arrow key. To return to the original screen, use the left arrow key.

Ge-71 4.0X101 1.1X103 4.0X101 1.1X103 5.8X103 1.6X105

Ge-77 3.0X10-1 8.1 3.0X10-

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1 8.1 1.3X105 3.6X106

Hf-172 (a) Hafnium (72) 6.0X10-1 1.6X101 6.0X10-
1 1.6X101 4.1X101 1.1X103

Hf-175 3.0 8.1X101 3.0
8.1X101 3.9X102 1.1X104

Hf-181 2.0 5.4X101 5.0X10-
1 1.4X101 6.3X102 1.7X104

Hf-182 Unlimited Unlimited Unlimited Un-
limited 8.1X10-6 2.2X10-4

Hg-194 (a) Mercury (80) 1.0 2.7X101 1.0
2.7X101 1.3X10-1 3.5

Hg-195m (a) 3.0 8.1X101 7.0X10-
1 1.9X101 1.5X104 4.0X105

Hg-197 2.0X101 5.4X102 1.0X101
2.7X102 9.2X103 2.5X105

Hg-197m 1.0X101 2.7X102 4.0X10-
1 1.1X101 2.5X104 6.7X105

Hg-203 5.0 1.4X102 1.0
2.7X101 5.1X102 1.4X104

Ho-166 Holmium (67) 4.0X10-1 1.1X101 4.0X10-
1 1.1X101 2.6X104 7.0X105

Ho-166m 6.0X10-1 1.6X101 5.0X10-
1 1.4X101 6.6X10-2 1.8

I-123 Iodine (53) 6.0 1.6X102 3.0
8.1X101 7.1X104 1.9X106

I-124 1.0 2.7X101 1.0
2.7X101 9.3X103 2.5X105

I-125 2.0X101 5.4X102 3.0
8.1X101 6.4X102 1.7X104

I-126 2.0 5.4X101 1.0
2.7X101 2.9X103 8.0X104

I-129 Unlimited Unlimited Unlimited Un-
limited 6.5X10-6 1.8X10-4

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I-131 3.0 8.1X101 7.0X10-
1 1.9X101 4.6X103 1.2X105

I-132 4.0X10-1 1.1X101 4.0X10-
1 1.1X101 3.8X105 1.0X107

I-133 7.0X10-1 1.9X101 6.0X10-
1 1.6X101 4.2X104 1.1X106

I-134 3.0X10-1 8.1 3.0X10-
1 8.1 9.9X105 2.7X107

I-135 (a) 6.0X10-1 1.6X101 6.0X10-
1 1.6X101 1.3X105 3.5X106

In-111 Indium (49) 3.0 8.1X101 3.0
8.1X101 1.5X104 4.2X105

In-113m 4.0 1.1X102 2.0
5.4X101 6.2X105 1.7X107

In-114m (a) 1.0X101 2.7X102 5.0X10-
1 1.4X101 8.6X102 2.3X104

In-115m 7.0 1.9X102 1.0
2.7X101 2.2X105 6.1X106

Ir-189 (a) Iridium (77) 1.0X101 2.7X102 1.0X101
2.7X102 1.9X103 5.2X104

Ir-190 7.0X10-1 1.9X101 7.0X10-
1 1.9X101 2.3X103 6.2X104

Ir-192 (c) 1.0 2.7X101 6.0X10-
1 1.6X101 3.4X102 9.2X103

Ir-194 3.0X10-1 8.1 3.0X10-
1 8.1 3.1X104 8.4X105

K-40 Potassium (19) 9.0X10-1 2.4X101 9.0X10-
1 2.4X101 2.4X10-7 6.4X10-6

K-42 2.0X10-1 5.4 2.0X10-
1 5.4 2.2X105 6.0X106

K-43 7.0X10-1 1.9X101 6.0X10-
1 1.6X101 1.2X105 3.3X106

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Kr-81 Krypton (36) 4.0X101 1.1X103 4.0X101
1.1X103 7.8X10-4 2.1X10-2

Kr-85 1.0X101 2.7X102 1.0X101
2.7X102 1.5X101 3.9X102

Kr-85m 8.0 2.2X102 3.0
8.1X101 3.0X105 8.2X106

Kr-87 2.0X10-1 5.4 2.0X10-
1 5.4 1.0X106 2.8X107

La-137 Lanthanum (57) 3.0X101 8.1X102 6.0
1.6X102 1.6X10-3 4.4X10-2

La-140 4.0X10-1 1.1X101 4.0X10-
1 1.1X101 2.1X104 5.6X105

Lu-172 Lutetium (71) 6.0X10-1 1.6X101 6.0X10-
1 1.6X101 4.2X103 1.1X105

Lu-173 8.0 2.2X102 8.0
2.2X102 5.6X101 1.5X103

Lu-174 9.0 2.4X102 9.0
2.4X102 2.3X101 6.2X102

Lu-174m 2.0X101 5.4X102 1.0X101
2.7X102 2.0X102 5.3X103

Lu-177 3.0X101 8.1X102 7.0X10-
1 1.9X101 4.1X103 1.1X105

Mg-28 (a) Magnesium (12) 3.0X10-1 8.1 3.0X10-
1 8.1 2.0X105 5.4X106

Mn-52 Manganese (25) 3.0X10-1 8.1 3.0X10-
1 8.1 1.6X104 4.4X105

Mn-53 Unlimited Unlimited Unlimited Un-
limited 6.8X10-5 1.8X10-3

Mn-54 1.0 2.7X101 1.0
2.7X101 2.9X102 7.7X103

Mn-56 3.0X10-1 8.1 3.0X10-
1 8.1 8.0X105 2.2X107

Mo-93 Molybdenum (42) 4.0X101 1.1X103 2.0X101

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5.4X102 4.1X10-2 1.1

Mo-99 (a) (i) 1.0 2.7X101 6.0X10-
1 1.6X101 1.8X104 4.8X105

N-13 Nitrogen (7) 9.0X10-1 2.4X101 6.0X10-
1 1.6X101 5.4X107 1.5X109

Na-22 Sodium (11) 5.0X10-1 1.4X101 5.0X10-
1 1.4X101 2.3X102 6.3X103

Na-24 2.0X10-1 5.4 2.0X10-
1 5.4 3.2X105 8.7X106

Nb-93m Niobium (41) 4.0X101 1.1X103 3.0X101
8.1X102 8.8 2.4X102

Nb-94 7.0X10-1 1.9X101 7.0X10-
1 1.9X101 6.9X10-3 1.9X10-1

Nb-95 1.0 2.7X101 1.0
2.7X101 1.5X103 3.9X104

Nb-97 9.0X10-1 2.4X101 6.0X10-
1 1.6X101 9.9X105 2.7X107

Nd-147 Neodymium (60) 6.0 1.6X102 6.0X10-
1 1.6X101 3.0X103 8.1X104

Nd-149 6.0X10-1 1.6X101 5.0X10-
1 1.4X101 4.5X105 1.2X107

Ni-59 Nickel (28) Unlimited Unlimited Unlimited Un-
limited 3.0X10-3 8.0X10-2

Ni-63 4.0X101 1.1X103 3.0X101
8.1X102 2.1 5.7X101

Ni-65 4.0X10-1 1.1X101 4.0X10-
1 1.1X101 7.1X105 1.9X107

Np-235 Neptunium (93) 4.0X101 1.1X103 4.0X101
1.1X103 5.2X101 1.4X103

Np-236 (short-lived) 2.0X101 5.4X102 2.0
5.4X101 4.7X10-4 1.3X10-2

Np-236 (long-lived) 9.0X100 2.4X102 2.0X10-
2 5.4X10-1 4.7X10-4 1.3X10-2

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Np-237 2.0X101 5.4X102 2.0X10-3 5.4X10-2 2.6X10-5 7.1X10-4

Np-239 7.0 1.9X102 4.0X10-1 1.1X101 8.6X103 2.3X105

Os-185 Osmium (76) 1.0 2.7X101 1.0 2.7X101 2.8X102 7.5X103

Os-191 1.0X101 2.7X102 2.0 5.4X101 1.6X103 4.4X104

Os-191m 4.0X101 1.1X103 3.0X101 8.1X102 4.6X104 1.3X106

Os-193 2.0 5.4X101 6.0X10-1 1.6X101 2.0X104 5.3X105

Os-194 (a) 3.0X10-1 8.1 3.0X10-1 8.1 1.1X101 3.1X102

P-32 Phosphorus (15) 5.0X10-1 1.4X101 5.0X10-1 1.4X101 1.1X104 2.9X105

P-33 4.0X101 1.1X103 1.0 2.7X101 5.8X103 1.6X105

Pa-230 (a) Protactinium (91) 2.0 5.4X101 7.0X10-2 1.9 1.2X103 3.3X104

Pa-231 4.0 1.1X102 4.0X10-4 1.1X10-2 1.7X10-3 4.7X10-2

Pa-233 5.0 1.4X102 7.0X10-1 1.9X101 7.7X102 2.1X104

Pb-201 Lead (82) 1.0 2.7X101 1.0 2.7X101 6.2X104 1.7X106

Pb-202 4.0X101 1.1X103 2.0X101 5.4X102 1.2X10-4 3.4X10-3

Pb-203 4.0 1.1X102 3.0 8.1X101 1.1X104 3.0X105

Pb-205 Unlimited Unlimited Unlimited Unlimited 4.5X10-6 1.2X10-4

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Pb-210 (a) 1.0 2.7X101 5.0X10-
2 1.4 2.8 7.6X101

Pb-212 (a) 7.0X10-1 1.9X101 2.0X10-
1 5.4 5.1X104 1.4X106

Pd-103 (a) Palladium (46) 4.0X101 1.1X103 4.0X101
1.1X103 2.8X103 7.5X104

Pd-107 Unlimited Unlimited Unlimited Un-
limited 1.9X10-5 5.1X10-4

Pd-109 2.0 5.4X101 5.0X10-
1 1.4X101 7.9X104 2.1X106

Pm-143 Promethium (61) 3.0 8.1X101 3.0
8.1X101 1.3X102 3.4X103

Pm-144 7.0X10-1 1.9X101 7.0X10-
1 1.9X101 9.2X101 2.5X103

Pm-145 3.0X101 8.1X102 1.0X101
2.7X102 5.2 1.4X102

Pm-147 4.0X101 1.1X103 2.0
5.4X101 3.4X101 9.3X102

Pm-148m (a) 8.0X10-1 2.2X101 7.0X10-
1 1.9X101 7.9X102 2.1X104

Pm-149 2.0 5.4X101 6.0X10-
1 1.6X101 1.5X104 4.0X105

Pm-151 2.0 5.4X101 6.0X10-
1 1.6X101 2.7X104 7.3X105

Po-210 Polonium (84) 4.0X101 1.1X103 2.0X10-
2 5.4X10-1 1.7X102 4.5X103

Pr-142 Praseodymium (59) 4.0X10-1 1.1X101 4.0X10-
1 1.1X101 4.3X104 1.2X106

Pr-143 3.0 8.1X101 6.0X10-
1 1.6X101 2.5X103 6.7X104

Pt-188 (a) Platinum (78) 1.0 2.7X101 8.0X10-
1 2.2X101 2.5X103 6.8X104

Pt-191 4.0 1.1X102 3.0

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8.1X101 8.7X103 2.4X105

Pt-193 4.0X101 1.1X103 4.0X101
1.1X103 1.4 3.7X101

Pt-193m 4.0X101 1.1X103 5.0X10-
1 1.4X101 5.8X103 1.6X105

Pt-195m 1.0X101 2.7X102 5.0X10-
1 1.4X101 6.2X103 1.7X105

Pt-197 2.0X101 5.4X102 6.0X10-
1 1.6X101 3.2X104 8.7X105

Pt-197m 1.0X101 2.7X102 6.0X10-
1 1.6X101 3.7X105 1.0X107

Pu-236 Plutonium (94) 3.0X101 8.1X102 3.0X10-
3 8.1X10-2 2.0X101 5.3X102

Pu-237 2.0X101 5.4X102 2.0X101
5.4X102 4.5X102 1.2X104

Pu-238 1.0X101 2.7X102 1.0X10-
3 2.7X10-2 6.3X10-1 1.7X101

Pu-239 1.0X101 2.7X102 1.0X10-
3 2.7X10-2 2.3X10-3 6.2X10-2

Pu-240 1.0X101 2.7X102 1.0X10-
3 2.7X10-2 8.4X10-3 2.3X10-1

Pu-241 (a) 4.0X101 1.1X103 6.0X10-
2 1.6 3.8 1.0X102

Pu-242 1.0X101 2.7X102 1.0X10-
3 2.7X10-2 1.5X10-4 3.9X10-3

Pu-244 (a) 4.0X10-1 1.1X101 1.0X10-
3 2.7X10-2 6.7X10-7 1.8X10-5

Ra-223 (a) Radium (88) 4.0X10-1 1.1X101 7.0X10-
3 1.9X10-1 1.9X103 5.1X104

Ra-224 (a) 4.0X10-1 1.1X101 2.0X10-
2 5.4X10-1 5.9X103 1.6X105

Ra-225 (a) 2.0X10-1 5.4 4.0X10-
3 1.1X10-1 1.5X103 3.9X104

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Ra-226 (a) 2.0X10-1 5.4 3.0X10-
3 8.1X10-2 3.7X10-2 1.0

Ra-228 (a) 6.0X10-1 1.6X101 2.0X10-
2 5.4X10-1 1.0X101 2.7X102

Rb-81 Rubidium (37) 2.0 5.4X101 8.0X10-
1 2.2X101 3.1X105 8.4X106

Rb-83 (a) 2.0 5.4X101 2.0
5.4X101 6.8X102 1.8X104

Rb-84 1.0 2.7X101 1.0
2.7X101 1.8X103 4.7X104

Rb-86 5.0X10-1 1.4X101 5.0X10-
1 1.4X101 3.0X103 8.1X104

Rb-87 Unlimited Unlimited Unlimited Un-
limited 3.2X10-9 8.6X10-8

Rb(nat) Unlimited Unlimited Unlimited Un-
limited 6.7X106 1.8X108

Re-184 Rhenium (75) 1.0 2.7X101 1.0
2.7X101 6.9X102 1.9X104

Re-184m 3.0 8.1X101 1.0
2.7X101 1.6X102 4.3X103

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

Re-186 2.0
Re-187 Unlimited
Re-188 4.0X10-1
Re-189 (a) 3.0
Re(nat) Unlimited
Rh-99 Rhodium (45) 2.0

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Rh-101		4.0
Rh-102		5.0X10-1
Rh-102m		2.0
Rh-103m		4.0X101
Rh-105		1.0X101
Rn-222 (a)	Radon (86)	3.0X10-1
Ru-97	Ruthenium (44)	5.0
Ru-103 (a)		2.0
Ru-105		1.0
Ru-106 (a)		2.0X10-1
S-35	Sulphur (16)	4.0X101
Sb-122	Antimony (51)	4.0X10-1
Sb-124		6.0X10-1
Sb-125		2.0
Sb-126		4.0X10-1
Sc-44	Scandium (21)	5.0X10-1
Sc-46		5.0X10-1
Sc-47		1.0X101
Sc-48		3.0X10-1
Se-75	Selenium (34)	3.0
Se-79		4.0X101
Si-31	Silicon (14)	6.0X10-1
Si-32		4.0X101
Sm-145	Samarium (62)	1.0X101
Sm-147		Unlimited
Sm-151		4.0X101
Sm-153		9.0
Sn-113 (a)	Tin (50)	4.0
Sn-117m		7.0
Sn-119m		4.0X101
Sn-121m (a)		4.0X101
Sn-123		8.0X10-1
Sn-125		4.0X10-1
Sn-126 (a)		6.0X10-1
Sr-82 (a)	Strontium (38)	2.0X10-1
Sr-85		2.0
Sr-85m		5.0
Sr-87m		3.0
Sr-89		6.0X10-1
Sr-90 (a)		3.0X10-1
Sr-91 (a)		3.0X10-1
Sr-92 (a)		1.0
T(H-3)	Tritium (1)	4.0X101
Ta-178 (long-lived)	Tantalum (73)	1.0
Ta-179		3.0X101
Ta-182		9.0X10-1
Tb-157	Terbium (65)	4.0X101
Tb-158		1.0
Tb-160		1.0
Tc-95m (a)	Technetium (43)	2.0
Tc-96		4.0X10-1
Tc-96m (a)		4.0X10-1
Tc-97		Unlimited

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Tc-97m		4.0X101
Tc-98		8.0X10-1
Tc-99		4.0X101
Tc-99m		1.0X101
Te-121	Tellurium (52)	2.0
Te-121m		5.0
Te-123m		8.0
Te-125m		2.0X101
Te-127		2.0X101
Te-127m (a)		2.0X101
Te-129		7.0X10-1
Te-129m (a)		8.0X10-1
Te-131m (a)		7.0X10-1
Te-132 (a)		5.0X10-1
Th-227	Thorium (90)	1.0X101
Th-228 (a)		5.0X10-1
Th-229		5.0
Th-230		1.0X101
Th-231		4.0X101
Th-232		Unlimited
Th-234 (a)		3.0X10-1
Th(nat)		Unlimited
Ti-44 (a)	Titanium (22)	5.0X10-1
Tl-200	Thallium (81)	9.0X10-1
Tl-201		1.0X101
Tl-202		2.0
Tl-204		1.0X101
Tm-167	Thulium (69)	7.0
Tm-170		3.0
Tm-171		4.0X101
U-230 (fast lung absorption) (a)(d)	Uranium (92)	4.0X101
U-230 (medium lung absorption) (a)(e)		4.0X101
U-230 (slow lung absorption) (a)(f)		3.0X101
U-232 (fast lung absorption) (d)		4.0X101
U-232 (medium lung absorption) (e)		4.0X101
U-232 (slow lung absorption) (f)		1.0X101
U-233 (fast lung absorption) (d)		4.0X101
U-233 (medium lung absorption) (e)		4.0X101
U-233 (slow lung absorption) (f)		4.0X101
U-234 (fast lung absorption) (d)		4.0X101
U-234 (medium lung absorption) (e)		4.0X101
U-234 (slow lung absorption) (f)		4.0X101
U-235 (all lung absorption types) (a),(d),(e),(f)		Unlimited
U-236 (fast lung absorption) (d)		Unlimited
U-236 (medium lung absorption) (e)		4.0X101
U-236 (slow lung absorption) (f)		4.0X101
U-238 (all lung absorption types) (d),(e),(f)		Unlimited
U (nat)		Unlimited
U (enriched to 20% or less) (g)		Unlimited
U (dep)		Unlimited
V-48	Vanadium (23)	4.0X10-1
V-49		4.0X101
W-178 (a)	Tungsten (74)	9.0

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W-181		3.0X101
W-185		4.0X101
W-187		2.0
W-188 (a)		4.0X10-1
Xe-122 (a)	Xenon (54)	4.0X10-1
Xe-123		2.0
Xe-127		4.0
Xe-131m		4.0X101
Xe-133		2.0X101
Xe-135		3.0
Y-87 (a)	Yttrium (39)	1.0
Y-88		4.0X10-1
Y-90		3.0X10-1

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

****
******* This is piece 2. -- It begins at character 80 of table line 1. ********
****

5.4X101	6.0X10-1	1.6X101	6.9X103	1.9X105
Unlimited	Unlimited	Unlimited	1.4X10-9	3.8X10-8
1.1X101	4.0X10-1	1.1X101	3.6X104	9.8X105
8.1X101	6.0X10-1	1.6X101	2.5X104	6.8X105
Unlimited	Unlimited	Unlimited	0.0	2.4X10-8
5.4X101	2.0	5.4X101	3.0X103	8.2X104
1.1X102	3.0	8.1X101	4.1X101	1.1X103
1.4X101	5.0X10-1	1.4X101	4.5X101	1.2X103
5.4X101	2.0	5.4X101	2.3X102	6.2X103
1.1X103	4.0X101	1.1X103	1.2X106	3.3X107
2.7X102	8.0X10-1	2.2X101	3.1X104	8.4X105
8.1	4.0X10-3	1.1X10-1	5.7X103	1.5X105
1.4X102	5.0	1.4X102	1.7X104	4.6X105
5.4X101	2.0	5.4X101	1.2X103	3.2X104
2.7X101	6.0X10-1	1.6X101	2.5X105	6.7X106
5.4	2.0X10-1	5.4	1.2X102	3.3X103
1.1X103	3.0	8.1X101	1.6X103	4.3X104
1.1X101	4.0X10-1	1.1X101	1.5X104	4.0X105
1.6X101	6.0X10-1	1.6X101	6.5X102	1.7X104
5.4X101	1.0	2.7X101	3.9X101	1.0X103
1.1X101	4.0X10-1	1.1X101	3.1X103	8.4X104
1.4X101	5.0X10-1	1.4X101	6.7X105	1.8X107
1.4X101	5.0X10-1	1.4X101	1.3X103	3.4X104
2.7X102	7.0X10-1	1.9X101	3.1X104	8.3X105
8.1	3.0X10-1	8.1	5.5X104	1.5X106
8.1X101	3.0	8.1X101	5.4X102	1.5X104
1.1X103	2.0	5.4X101	2.6X10-3	7.0X10-2
1.6X101	6.0X10-1	1.6X101	1.4X106	3.9X107
1.1X103	5.0X10-1	1.4X101	3.9	1.1X102
2.7X102	1.0X101	2.7X102	9.8X101	2.6X103
Unlimited	Unlimited	Unlimited	8.5X10-1	2.3X10-8
1.1X103	1.0X101	2.7X102	9.7X10-1	2.6X101
2.4X102	6.0X10-1	1.6X101	1.6X104	4.4X105

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1.1X102	2.0	5.4X101	3.7X102	1.0X104
1.9X102	4.0X10-1	1.1X101	3.0X103	8.2X104
1.1X103	3.0X101	8.1X102	1.4X102	3.7X103
1.1X103	9.0X10-1	2.4X101	2.0	5.4X101
2.2X101	6.0X10-1	1.6X101	3.0X102	8.2X103
1.1X101	4.0X10-1	1.1X101	4.0X103	1.1X105
1.6X101	4.0X10-1	1.1X101	1.0X10-3	2.8X10-2
5.4	2.0X10-1	5.4	2.3X103	6.2X104
5.4X101	2.0	5.4X101	8.8X102	2.4X104
1.4X102	5.0	1.4X102	1.2X106	3.3X107
8.1X101	3.0	8.1X101	4.8X105	1.3X107
1.6X101	6.0X10-1	1.6X101	1.1X103	2.9X104
8.1	3.0X10-1	8.1	5.1	1.4X102
8.1	3.0X10-1	8.1	1.3X105	3.6X106
2.7X101	3.0X10-1	8.1	4.7X105	1.3X107
1.1X103	4.0X101	1.1X103	3.6X102	9.7X103
2.7X101	8.0X10-1	2.2X101	4.2X106	1.1X108
8.1X102	3.0X101	8.1X102	4.1X101	1.1X103
2.4X101	5.0X10-1	1.4X101	2.3X102	6.2X103
1.1X103	4.0X101	1.1X103	5.6X10-1	1.5X101
2.7X101	1.0	2.7X101	5.6X10-1	1.5X101
2.7X101	6.0X10-1	1.6X101	4.2X102	1.1X104
5.4X101	2.0	5.4X101	8.3X102	2.2X104
1.1X101	4.0X10-1	1.1X101	1.2X104	3.2X105
1.1X101	4.0X10-1	1.1X101	1.4X106	3.8X107
Unlimited	Unlimited	Unlimited	5.2X10-5	1.4X10-3
1.1X103	1.0	2.7X101	5.6X102	1.5X104
2.2X101	7.0X10-1	1.9X101	3.2X10-5	8.7X10-4
1.1X103	9.0X10-1	2.4X101	6.3X10-4	1.7X10-2
2.7X102	4.0	1.1X102	1.9X105	5.3X106
5.4X101	2.0	5.4X101	2.4X103	6.4X104
1.4X102	3.0	8.1X101	2.6X102	7.0X103
2.2X102	1.0	2.7X101	3.3X102	8.9X103
5.4X102	9.0X10-1	2.4X101	6.7X102	1.8X104
5.4X102	7.0X10-1	1.9X101	9.8X104	2.6X106
5.4X102	5.0X10-1	1.4X101	3.5X102	9.4X103
1.9X101	6.0X10-1	1.6X101	7.7X105	2.1X107
2.2X101	4.0X10-1	1.1X101	1.1X103	3.0X104
1.9X101	5.0X10-1	1.4X101	3.0X104	8.0X105
1.4X101	4.0X10-1	1.1X101	3.1X104	3.0X105
2.7X102	5.0X10-3	1.4X10-1	1.1X103	3.1X104
1.4X101	1.0X10-3	2.7X10-2	3.0X101	8.2X102
1.4X102	5.0X10-4	1.4X10-2	7.9X10-3	2.1X10-1
2.7X102	1.0X10-3	2.7X10-2	7.6X10-4	2.1X10-2
1.1X103	2.0X10-2	5.4X10-1	2.0X104	5.3X105
Unlimited	Unlimited	Unlimited	4.0X10-9	1.1X10-7
8.1	3.0X10-1	8.1	8.6X102	2.3X104
Unlimited	Unlimited	Unlimited	8.1X10-9	2.2X10-7
1.4X101	4.0X10-1	1.1X101	6.4	1.7X102
2.4X101	9.0X10-1	2.4X101	2.2X104	6.0X105
2.7X102	4.0	1.1X102	7.9X103	2.1X105
5.4X101	2.0	5.4X101	2.0X103	5.3X104
2.7X102	7.0X10-1	1.9X101	1.7X101	4.6X102

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1.9X102	8.0X10-1	2.2X101	3.1X103	8.5X104
8.1X101	6.0X10-1	1.6X101	2.2X102	6.0X103
1.1X103	4.0X101	1.1X103	4.0X101	1.1X103
1.1X103	1.0X10-1	2.7	1.0X103	2.7X104
1.1X103	4.0X10-3	1.1X10-1	1.0X103	2.7X104
8.1X102	3.0X10-3	8.1X10-2	1.0X103	2.7X104
1.1X103	1.0X10-2	2.7X10-1	8.3X10-1	2.2X101
1.1X103	7.0X10-3	1.9X10-1	8.3X10-1	2.2X101
2.7X102	1.0X10-3	2.7X10-2	8.3X10-1	2.2X101
1.1X103	9.0X10-2	2.4	3.6X10-4	9.7X10-3
1.1X103	2.0X10-2	5.4X10-1	3.6X10-4	9.7X10-3
1.1X103	6.0X10-3	1.6X10-1	3.6X10-4	9.7X10-3
1.1X103	9.0X10-2	2.4	2.3X10-4	6.2X10-3
1.1X103	2.0X10-2	5.4X10-1	2.3X10-4	6.2X10-3
1.1X103	6.0X10-3	1.6X10-1	2.3X10-4	6.2X10-3
Unlimited	Unlimited	Unlimited	8.0X10-8	2.2X10-6
Unlimited	Unlimited	Unlimited	2.4X10-6	6.5X10-5
1.1X103	2.0X10-2	5.4X10-1	2.4X10-6	6.5X10-5
1.1X103	6.0X10-3	1.6X10-1	2.4X10-6	6.5X10-5
Unlimited	Unlimited	Unlimited	1.2X10-8	3.4X10-7
Unlimited	Unlimited	Unlimited	2.6X10-8	7.1X10-7
Unlimited	Unlimited	Unlimited	See Table A-4	See Table A-4
Unlimited	Unlimited	Unlimited	See Table A-4	(See Table A-3)
1.1X101	4.0X10-1	1.1X101	6.3X103	1.7X105
1.1X103	4.0X101	1.1X103	3.0X102	8.1X103
2.4X102	5.0	1.4X102	1.3X103	3.4X104
8.1X102	3.0X101	8.1X102	2.2X102	6.0X103
1.1X103	8.0X10-1	2.2X101	3.5X102	9.4X103
5.4X101	6.0X10-1	1.6X101	2.6X104	7.0X105
1.1X101	3.0X10-1	8.1	3.7X102	1.0X104
1.1X101	4.0X10-1	1.1X101	4.8X104	1.3X106
5.4X101	7.0X10-1	1.9X101	4.4X105	1.2X107
1.1X102	2.0	5.4X101	1.0X103	2.8X104
1.1X103	4.0X101	1.1X103	3.1X103	8.4X104
5.4X102	1.0X101	2.7X102	6.9X103	1.9X105
8.1X101	2.0	5.4X101	9.5X104	2.6X106
2.7X101	1.0	2.7X101	1.7X104	4.5X105
1.1X101	4.0X10-1	1.1X101	5.2X102	1.4X104
8.1	3.0X10-1	8.1	2.0X104	5.4X105

```
80..+...90....+....0....+...10....+...20....+...30....+...40....+...50
```

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****

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******** This is piece 1. -- It begins at character 1 of table line 1. ********

****

Re-186 2.0

Re-187 Unlimited

Re-188 4.0X10-1

Re-189 (a) 3.0

Re(nat) Unlimited

Rh-99 Rhodium (45) 2.0

Rh-101 4.0

Rh-102 5.0X10-1

Rh-102m 2.0

Rh-103m 4.0X101

Rh-105 1.0X101

Rn-222 (a) Radon (86) 3.0X10-1

Ru-97 Ruthenium (44) 5.0

Ru-103 (a) 2.0

Ru-105 1.0

Ru-106 (a) 2.0X10-1

S-35 Sulphur (16) 4.0X101

Sb-122 Antimony (51) 4.0X10-1

Sb-124 6.0X10-1

Sb-125 2.0

Sb-126 4.0X10-1

Sc-44 Scandium (21) 5.0X10-1

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Sc-46		5.0X10-1
Sc-47		1.0X101
Sc-48		3.0X10-1
Se-75	Selenium (34)	3.0
Se-79		4.0X101
Si-31	Silicon (14)	6.0X10-1
Si-32		4.0X101
Sm-145	Samarium (62)	1.0X101
Sm-147		Unlimited
Sm-151		4.0X101
Sm-153		9.0
Sn-113 (a)	Tin (50)	4.0
Sn-117m		7.0
Sn-119m		4.0X101
Sn-121m (a)		4.0X101
Sn-123		8.0X10-1
Sn-125		4.0X10-1
Sn-126 (a)		6.0X10-1
Sr-82 (a)	Strontium (38)	2.0X10-1
Sr-85		2.0
Sr-85m		5.0
Sr-87m		3.0
Sr-89		6.0X10-1

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Sr-90 (a) 3.0X10-1

Sr-91 (a) 3.0X10-1

Sr-92 (a) 1.0

T(H-3) Tritium (1) 4.0X101

Ta-178 (long-lived) Tantalum (73) 1.0

Ta-179 3.0X101

Ta-182 9.0X10-1

Tb-157 Terbium (65) 4.0X101

Tb-158 1.0

Tb-160 1.0

Tc-95m (a) Technetium (43) 2.0

Tc-96 4.0X10-1

Tc-96m (a) 4.0X10-1

Tc-97 Unlimited

Tc-97m 4.0X101

Tc-98 8.0X10-1

Tc-99 4.0X101

Tc-99m 1.0X101

Te-121 Tellurium (52) 2.0

Te-121m 5.0

Te-123m 8.0

Te-125m 2.0X101

Te-127 2.0X101

Te-127m (a) 2.0X101

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Te-129 7.0X10-1

Te-129m (a) 8.0X10-1

Te-131m (a) 7.0X10-1

Te-132 (a) 5.0X10-1

Th-227 Thorium (90) 1.0X101

Th-228 (a) 5.0X10-1

Th-229 5.0

Th-230 1.0X101

Th-231 4.0X101

Th-232 Unlimited

Th-234 (a) 3.0X10-1

Th(nat) Unlimited

Ti-44 (a) Titanium (22) 5.0X10-1

Tl-200 Thallium (81) 9.0X10-1

Tl-201 1.0X101

Tl-202 2.0

Tl-204 1.0X101

Tm-167 Thulium (69) 7.0

Tm-170 3.0

Tm-171 4.0X101

U-230 (fast lung absorption) (a)(d) Uranium (92) 4.0X101

U-230 (medium lung absorption) (a)(e) 4.0X101

U-230 (slow lung absorption) (a)(f) 3.0X101

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U-232 (fast lung absorption) (d) 4.0X101

U-232 (medium lung absorption) (e) 4.0X101

U-232 (slow lung absorption) (f) 1.0X101

U-233 (fast lung absorption) (d) 4.0X101

U-233 (medium lung absorption) (e) 4.0X101

U-233 (slow lung absorption) (f) 4.0X101

U-234 (fast lung absorption) (d) 4.0X101

U-234 (medium lung absorption) (e) 4.0X101

U-234 (slow lung absorption) (f) 4.0X101

U-235 (all lung absorption types) (a),(d),(e),(f) Unlimited

U-236 (fast lung absorption) (d) Unlimited

U-236 (medium lung absorption) (e) 4.0X101

U-236 (slow lung absorption) (f) 4.0X101

U-238 (all lung absorption types) (d),(e),(f) Unlimited

U (nat) Unlimited

U (enriched to 20% or less) (g) Unlimited

U (dep) Unlimited

V-48 Vanadium (23) 4.0X10-1

V-49 4.0X101

W-178 (a) Tungsten (74) 9.0

W-181 3.0X101

W-185 4.0X101

W-187 2.0

W-188 (a) 4.0X10-1

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Xe-122 (a) Xenon (54) 4.0X10-1

Xe-123 2.0

Xe-127 4.0

Xe-131m 4.0X101

Xe-133 2.0X101

Xe-135 3.0

Y-87 (a) Yttrium (39) 1.0

Y-88 4.0X10-1

Y-90 3.0X10-1

1...k...10....k...20....k...30....k...40....k...50....k...60....k... 70....k....

****

******* This is piece 2. -- It begins at character 80 of table line 1. ********

****

5.4X101	6.0X10-1	1.6X101	6.9X103	1.9X105
Unlimited	Unlimited	Unlimited	1.4X10-9	3.8X10-8
1.1X101	4.0X10-1	1.1X101	3.6X104	9.8X105
8.1X101	6.0X10-1	1.6X101	2.5X104	6.8X105
Unlimited	Unlimited	Unlimited	0.0	2.4X10-8
5.4X101	2.0	5.4X101	3.0X103	8.2X104
1.1X102	3.0	8.1X101	4.1X101	1.1X103
1.4X101	5.0X10-1	1.4X101	4.5X101	1.2X103
5.4X101	2.0	5.4X101	2.3X102	6.2X103
1.1X103	4.0X101	1.1X103	1.2X106	3.3X107

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2.7X102	8.0X10-1	2.2X101	3.1X104	8.4X105
8.1	4.0X10-3	1.1X10-1	5.7X103	1.5X105
1.4X102	5.0	1.4X102	1.7X104	4.6X105
5.4X101	2.0	5.4X101	1.2X103	3.2X104
2.7X101	6.0X10-1	1.6X101	2.5X105	6.7X106
5.4	2.0X10-1	5.4	1.2X102	3.3X103
1.1X103	3.0	8.1X101	1.6X103	4.3X104
1.1X101	4.0X10-1	1.1X101	1.5X104	4.0X105
1.6X101	6.0X10-1	1.6X101	6.5X102	1.7X104
5.4X101	1.0	2.7X101	3.9X101	1.0X103
1.1X101	4.0X10-1	1.1X101	3.1X103	8.4X104
1.4X101	5.0X10-1	1.4X101	6.7X105	1.8X107
1.4X101	5.0X10-1	1.4X101	1.3X103	3.4X104
2.7X102	7.0X10-1	1.9X101	3.1X104	8.3X105
8.1	3.0X10-1	8.1	5.5X104	1.5X106
8.1X101	3.0	8.1X101	5.4X102	1.5X104
1.1X103	2.0	5.4X101	2.6X10-3	7.0X10-2
1.6X101	6.0X10-1	1.6X101	1.4X106	3.9X107
1.1X103	5.0X10-1	1.4X101	3.9	1.1X102
2.7X102	1.0X101	2.7X102	9.8X101	2.6X103
Unlimited	Unlimited	Unlimited	8.5X10-1	2.3X10-8
1.1X103	1.0X101	2.7X102	9.7X10-1	2.6X101
2.4X102	6.0X10-1	1.6X101	1.6X104	4.4X105
1.1X102	2.0	5.4X101	3.7X102	1.0X104

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1.9X102	4.0X10-1	1.1X101	3.0X103	8.2X104
1.1X103	3.0X101	8.1X102	1.4X102	3.7X103
1.1X103	9.0X10-1	2.4X101	2.0	5.4X101
2.2X101	6.0X10-1	1.6X101	3.0X102	8.2X103
1.1X101	4.0X10-1	1.1X101	4.0X103	1.1X105
1.6X101	4.0X10-1	1.1X101	1.0X10-3	2.8X10-2
5.4	2.0X10-1	5.4	2.3X103	6.2X104
5.4X101	2.0	5.4X101	8.8X102	2.4X104
1.4X102	5.0	1.4X102	1.2X106	3.3X107
8.1X101	3.0	8.1X101	4.8X105	1.3X107
1.6X101	6.0X10-1	1.6X101	1.1X103	2.9X104
8.1	3.0X10-1	8.1	5.1	1.4X102
8.1	3.0X10-1	8.1	1.3X105	3.6X106
2.7X101	3.0X10-1	8.1	4.7X105	1.3X107
1.1X103	4.0X101	1.1X103	3.6X102	9.7X103
2.7X101	8.0X10-1	2.2X101	4.2X106	1.1X108
8.1X102	3.0X101	8.1X102	4.1X101	1.1X103
2.4X101	5.0X10-1	1.4X101	2.3X102	6.2X103
1.1X103	4.0X101	1.1X103	5.6X10-1	1.5X101
2.7X101	1.0	2.7X101	5.6X10-1	1.5X101
2.7X101	6.0X10-1	1.6X101	4.2X102	1.1X104
5.4X101	2.0	5.4X101	8.3X102	2.2X104
1.1X101	4.0X10-1	1.1X101	1.2X104	3.2X105

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1.1X101	4.0X10-1	1.1X101	1.4X106	3.8X107
Unlimited	Unlimited	Unlimited	5.2X10-5	1.4X10-3
1.1X103	1.0	2.7X101	5.6X102	1.5X104
2.2X101	7.0X10-1	1.9X101	3.2X10-5	8.7X10-4
1.1X103	9.0X10-1	2.4X101	6.3X10-4	1.7X10-2
2.7X102	4.0	1.1X102	1.9X105	5.3X106
5.4X101	2.0	5.4X101	2.4X103	6.4X104
1.4X102	3.0	8.1X101	2.6X102	7.0X103
2.2X102	1.0	2.7X101	3.3X102	8.9X103
5.4X102	9.0X10-1	2.4X101	6.7X102	1.8X104
5.4X102	7.0X10-1	1.9X101	9.8X104	2.6X106
5.4X102	5.0X10-1	1.4X101	3.5X102	9.4X103
1.9X101	6.0X10-1	1.6X101	7.7X105	2.1X107
2.2X101	4.0X10-1	1.1X101	1.1X103	3.0X104
1.9X101	5.0X10-1	1.4X101	3.0X104	8.0X105
1.4X101	4.0X10-1	1.1X101	3.1X104	3.0X105
2.7X102	5.0X10-3	1.4X10-1	1.1X103	3.1X104
1.4X101	1.0X10-3	2.7X10-2	3.0X101	8.2X102
1.4X102	5.0X10-4	1.4X10-2	7.9X10-3	2.1X10-1
2.7X102	1.0X10-3	2.7X10-2	7.6X10-4	2.1X10-2
1.1X103	2.0X10-2	5.4X10-1	2.0X104	5.3X105
Unlimited	Unlimited	Unlimited	4.0X10-9	1.1X10-7
8.1	3.0X10-1	8.1	8.6X102	2.3X104
Unlimited	Unlimited	Unlimited	8.1X10-9	2.2X10-7

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1.4X101	4.0X10-1	1.1X101	6.4	1.7X102
2.4X101	9.0X10-1	2.4X101	2.2X104	6.0X105
2.7X102	4.0	1.1X102	7.9X103	2.1X105
5.4X101	2.0	5.4X101	2.0X103	5.3X104
2.7X102	7.0X10-1	1.9X101	1.7X101	4.6X102
1.9X102	8.0X10-1	2.2X101	3.1X103	8.5X104
8.1X101	6.0X10-1	1.6X101	2.2X102	6.0X103
1.1X103	4.0X101	1.1X103	4.0X101	1.1X103
1.1X103	1.0X10-1	2.7	1.0X103	2.7X104
1.1X103	4.0X10-3	1.1X10-1	1.0X103	2.7X104
8.1X102	3.0X10-3	8.1X10-2	1.0X103	2.7X104
1.1X103	1.0X10-2	2.7X10-1	8.3X10-1	2.2X101
1.1X103	7.0X10-3	1.9X10-1	8.3X10-1	2.2X101
2.7X102	1.0X10-3	2.7X10-2	8.3X10-1	2.2X101
1.1X103	9.0X10-2	2.4	3.6X10-4	9.7X10-3
1.1X103	2.0X10-2	5.4X10-1	3.6X10-4	9.7X10-3
1.1X103	6.0X10-3	1.6X10-1	3.6X10-4	9.7X10-3
1.1X103	9.0X10-2	2.4	2.3X10-4	6.2X10-3
1.1X103	2.0X10-2	5.4X10-1	2.3X10-4	6.2X10-3
1.1X103	6.0X10-3	1.6X10-1	2.3X10-4	6.2X10-3
Unlimited	Unlimited	Unlimited	8.0X10-8	2.2X10-6
Unlimited	Unlimited	Unlimited	2.4X10-6	6.5X10-5
1.1X103	2.0X10-2	5.4X10-1	2.4X10-6	6.5X10-5

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1.1X103	6.0X10-3	1.6X10-1	2.4X10-6	6.5X10-5
Unlimited	Unlimited	Unlimited	1.2X10-8	3.4X10-7
Unlimited	Unlimited	Unlimited	2.6X10-8	7.1X10-7
Unlimited	Unlimited	Unlimited	See Table A-4	See Table A-4
Unlimited	Unlimited	Unlimited	See Table A-4	(See Table A-3)
1.1X101	4.0X10-1	1.1X101	6.3X103	1.7X105
1.1X103	4.0X101	1.1X103	3.0X102	8.1X103
2.4X102	5.0	1.4X102	1.3X103	3.4X104
8.1X102	3.0X101	8.1X102	2.2X102	6.0X103
1.1X103	8.0X10-1	2.2X101	3.5X102	9.4X103
5.4X101	6.0X10-1	1.6X101	2.6X104	7.0X105
1.1X101	3.0X10-1	8.1	3.7X102	1.0X104
1.1X101	4.0X10-1	1.1X101	4.8X104	1.3X106
5.4X101	7.0X10-1	1.9X101	4.4X105	1.2X107
1.1X102	2.0	5.4X101	1.0X103	2.8X104
1.1X103	4.0X101	1.1X103	3.1X103	8.4X104
5.4X102	1.0X101	2.7X102	6.9X103	1.9X105
8.1X101	2.0	5.4X101	9.5X104	2.6X106
2.7X101	1.0	2.7X101	1.7X104	4.5X105
1.1X101	4.0X10-1	1.1X101	5.2X102	1.4X104
8.1	3.0X10-1	8.1	2.0X104	5.4X105

80..k...90....k....0....k...10....k...20....k...30....k...40....k...50

WIDETABLE

NOTE--Some parts of this form are wider than one screen. To view

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material that exceeds the width of this screen, use the right arrow

key. To return to the original screen, use the left arrow key.

Re-186 2.0 5.4X101 6.0X10-1 1.6X101
6.9X103 1.9X105

Re-187 Unlimited Unlimited Unlimited Unlimited 1.4X10-
9 3.8X10-8

Re-188 4.0X10-1 1.1X101 4.0X10-1 1.1X101
3.6X104 9.8X105

Re-189 (a) 3.0 8.1X101 6.0X10-1 1.6X101
2.5X104 6.8X105

Re(nat) Unlimited Unlimited Unlimited Unlimited
0.0 2.4X10-8

Rh-99 Rhodium (45) 2.0 5.4X101 2.0 5.4X101
3.0X103 8.2X104

Rh-101 4.0 1.1X102 3.0 8.1X101
4.1X101 1.1X103

Rh-102 5.0X10-1 1.4X101 5.0X10-1 1.4X101
4.5X101 1.2X103

Rh-102m 2.0 5.4X101 2.0 5.4X101
2.3X102 6.2X103

Rh-103m 4.0X101 1.1X103 4.0X101 1.1X103
1.2X106 3.3X107

Rh-105 1.0X101 2.7X102 8.0X10-1 2.2X101
3.1X104 8.4X105

Rn-222 (a) Radon (86) 3.0X10-
1 8.1 4.0X10-3 1.1X10-1 5.7X103 1.5X105

Ru-97 Ruthenium (44) 5.0 1.4X102 5.0 1.4X102
1.7X104 4.6X105

Ru-103 (a) 2.0 5.4X101 2.0 5.4X101
1.2X103 3.2X104

Ru-105 1.0 2.7X101 6.0X10-1 1.6X101
2.5X105 6.7X106

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Ru-106 (a) 2.0X10-1 5.4 2.0X10-1 5.4
1.2X102 3.3X103

S-35 Sulphur (16) 4.0X101 1.1X103 3.0 8.1X101
1.6X103 4.3X104

Sb-122 Antimony (51) 4.0X10-1 1.1X101 4.0X10-1 1.1X101
1.5X104 4.0X105

Sb-124 6.0X10-1 1.6X101 6.0X10-1 1.6X101
6.5X102 1.7X104

Sb-125 2.0 5.4X101 1.0 2.7X101
3.9X101 1.0X103

Sb-126 4.0X10-1 1.1X101 4.0X10-1 1.1X101
3.1X103 8.4X104

Sc-44 Scandium (21) 5.0X10-1 1.4X101 5.0X10-1 1.4X101
6.7X105 1.8X107

Sc-46 5.0X10-1 1.4X101 5.0X10-1 1.4X101 1.3X103
3.4X104

Sc-47 1.0X101 2.7X102 7.0X10-1 1.9X101 3.1X104
8.3X105

Sc-48 3.0X10-1 8.1 3.0X10-1 8.1 5.5X104
1.5X106

Se-75 Selenium (34) 3.0 8.1X101 3.0 8.1X101
5.4X102 1.5X104

Se-79 4.0X101 1.1X103 2.0 5.4X101 2.6X10-
3 7.0X10-2

Si-31 Silicon (14) 6.0X10-1 1.6X101 6.0X10-1 1.6X101
1.4X106 3.9X107

Si-32 4.0X101 1.1X103 5.0X10-1 1.4X101
3.9 1.1X102

Sm-145 Samarium (62) 1.0X101 2.7X102 1.0X101 2.7X102
9.8X101 2.6X103

Sm-147 Unlimited Unlimited Unlimited Unlimited 8.5X10-
1 2.3X10-8

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Sm-151 4.0X101 1.1X103 1.0X101 2.7X102 9.7X10-1 2.6X101

Sm-153 9.0 2.4X102 6.0X10-1 1.6X101 1.6X104 4.4X105

Sn-113 (a) Tin (50) 4.0 1.1X102 2.0 5.4X101 3.7X102 1.0X104

Sn-117m 7.0 1.9X102 4.0X10-1 1.1X101 3.0X103 8.2X104

Sn-119m 4.0X101 1.1X103 3.0X101 8.1X102 1.4X102 3.7X103

Sn-121m (a) 4.0X101 1.1X103 9.0X10-1 2.4X101 2.0 5.4X101

Sn-123 8.0X10-1 2.2X101 6.0X10-1 1.6X101 3.0X102 8.2X103

Sn-125 4.0X10-1 1.1X101 4.0X10-1 1.1X101 4.0X103 1.1X105

Sn-126 (a) 6.0X10-1 1.6X101 4.0X10-1 1.1X101 1.0X10-3 2.8X10-2

Sr-82 (a) Strontium (38) 2.0X10-1 5.4 2.0X10-1 5.4 2.3X103 6.2X104

Sr-85 2.0 5.4X101 2.0 5.4X101 8.8X102 2.4X104

Sr-85m 5.0 1.4X102 5.0 1.4X102 1.2X106 3.3X107

Sr-87m 3.0 8.1X101 3.0 8.1X101 4.8X105 1.3X107

Sr-89 6.0X10-1 1.6X101 6.0X10-1 1.6X101 1.1X103 2.9X104

Sr-90 (a) 3.0X10-1 8.1 3.0X10-1 8.1 5.1 1.4X102

Sr-91 (a) 3.0X10-1 8.1 3.0X10-1 8.1 1.3X105 3.6X106

Sr-92 (a) 1.0 2.7X101 3.0X10-1 8.1

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4.7X105 1.3X107

T(H-3) Tritium (1) 4.0X101 1.1X103 4.0X101 1.1X103
3.6X102 9.7X103

Ta-178 (long-lived) Tantalum (73) 1.0 2.7X101 8.0X10-
1 2.2X101 4.2X106 1.1X108

Ta-179 3.0X101 8.1X102 3.0X101 8.1X102
4.1X101 1.1X103

Ta-182 9.0X10-1 2.4X101 5.0X10-1 1.4X101
2.3X102 6.2X103

Tb-157 Terbium (65) 4.0X101 1.1X103 4.0X101 1.1X103 5.6X10-
1 1.5X101

Tb-158 1.0 2.7X101 1.0 2.7X101 5.6X10-
1 1.5X101

Tb-160 1.0 2.7X101 6.0X10-1 1.6X101
4.2X102 1.1X104

Tc-95m (a) Technetium (43) 2.0 5.4X101 2.0 5.4X101
8.3X102 2.2X104

Tc-96 4.0X10-1 1.1X101 4.0X10-1 1.1X101 1.2X104
3.2X105

Tc-96m (a) 4.0X10-1 1.1X101 4.0X10-1 1.1X101
1.4X106 3.8X107

Tc-97 Unlimited Unlimited Unlimited Unlimited 5.2X10-
5 1.4X10-3

Tc-97m 4.0X101 1.1X103 1.0 2.7X101
5.6X102 1.5X104

Tc-98 8.0X10-1 2.2X101 7.0X10-1 1.9X101 3.2X10-
5 8.7X10-4

Tc-99 4.0X101 1.1X103 9.0X10-1 2.4X101 6.3X10-
4 1.7X10-2

Tc-99m 1.0X101 2.7X102 4.0 1.1X102
1.9X105 5.3X106

Te-121 Tellurium (52) 2.0 5.4X101 2.0 5.4X101
2.4X103 6.4X104

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Te-121m 5.0 1.4X102 3.0 8.1X101
2.6X102 7.0X103

Te-123m 8.0 2.2X102 1.0 2.7X101
3.3X102 8.9X103

Te-125m 2.0X101 5.4X102 9.0X10-1 2.4X101
6.7X102 1.8X104

Te-127 2.0X101 5.4X102 7.0X10-1 1.9X101
9.8X104 2.6X106

Te-127m (a) 2.0X101 5.4X102 5.0X10-1 1.4X101
3.5X102 9.4X103

Te-129 7.0X10-1 1.9X101 6.0X10-1 1.6X101
7.7X105 2.1X107

Te-129m (a) 8.0X10-1 2.2X101 4.0X10-1 1.1X101
1.1X103 3.0X104

Te-131m (a) 7.0X10-1 1.9X101 5.0X10-1 1.4X101
3.0X104 8.0X105

Te-132 (a) 5.0X10-1 1.4X101 4.0X10-1 1.1X101
3.1X104 3.0X105

Th-227 Thorium (90) 1.0X101 2.7X102 5.0X10-3 1.4X10-
1 1.1X103 3.1X104

Th-228 (a) 5.0X10-1 1.4X101 1.0X10-3 2.7X10-
2 3.0X101 8.2X102

Th-229 5.0 1.4X102 5.0X10-4 1.4X10-2 7.9X10-
3 2.1X10-1

Th-230 1.0X101 2.7X102 1.0X10-3 2.7X10-2 7.6X10-
4 2.1X10-2

Th-231 4.0X101 1.1X103 2.0X10-2 5.4X10-
1 2.0X104 5.3X105

Th-232 Unlimited Unlimited Unlimited Unlimited 4.0X10-
9 1.1X10-7

Th-234 (a) 3.0X10-1 8.1 3.0X10-1 8.1
8.6X102 2.3X104

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Th(nat) Unlimited Unlimited Unlimited Unlimited 8.1X10-
9 2.2X10-7

Ti-44 (a) Titanium (22) 5.0X10-1 1.4X101 4.0X10-1 1.1X101
6.4 1.7X102

Tl-200 Thallium (81) 9.0X10-1 2.4X101 9.0X10-1 2.4X101
2.2X104 6.0X105

Tl-201 1.0X101 2.7X102 4.0 1.1X102
7.9X103 2.1X105

Tl-202 2.0 5.4X101 2.0 5.4X101
2.0X103 5.3X104

Tl-204 1.0X101 2.7X102 7.0X10-1 1.9X101
1.7X101 4.6X102

Tm-167 Thulium (69) 7.0 1.9X102 8.0X10-1 2.2X101
3.1X103 8.5X104

Tm-170 3.0 8.1X101 6.0X10-1 1.6X101
2.2X102 6.0X103

Tm-171 4.0X101 1.1X103 4.0X101 1.1X103
4.0X101 1.1X103

U-230 (fast lung absorption) (a)(d) Uranium (92) 4.0X101 1.1X103
1.0X10-1 2.7 1.0X103 2.7X104

U-230 (medium lung absorption) (a)(e) 4.0X101 1.1X103 4.0X10-
3 1.1X10-1 1.0X103 2.7X104

U-230 (slow lung absorption) (a)(f) 3.0X101 8.1X102 3.0X10-
3 8.1X10-2 1.0X103 2.7X104

U-232 (fast lung absorption) (d) 4.0X101 1.1X103 1.0X10-
2 2.7X10-1 8.3X10-1 2.2X101

U-232 (medium lung absorption) (e) 4.0X101 1.1X103 7.0X10-
3 1.9X10-1 8.3X10-1 2.2X101

U-232 (slow lung absorption) (f) 1.0X101 2.7X102 1.0X10-
3 2.7X10-2 8.3X10-1 2.2X101

U-233 (fast lung absorption) (d) 4.0X101 1.1X103 9.0X10-
2 2.4 3.6X10-4 9.7X10-3

U-233 (medium lung absorption) (e) 4.0X101 1.1X103 2.0X10-

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2 5.4X10-1 3.6X10-4 9.7X10-3

U-233 (slow lung absorption) (f) 4.0X101 1.1X103 6.0X10-3 1.6X10-1 3.6X10-4 9.7X10-3

U-234 (fast lung absorption) (d) 4.0X101 1.1X103 9.0X10-2 2.4 2.3X10-4 6.2X10-3

U-234 (medium lung absorption) (e) 4.0X101 1.1X103 2.0X10-2 5.4X10-1 2.3X10-4 6.2X10-3

U-234 (slow lung absorption) (f) 4.0X101 1.1X103 6.0X10-3 1.6X10-1 2.3X10-4 6.2X10-3

U-235 (all lung absorption types) (a),(d),(e),(f) Unlimited Unlimited Unlimited Unlimited 8.0X10-8 2.2X10-6

U-236 (fast lung absorption) (d) Unlimited Unlimited Unlimited Unlimited 2.4X10-6 6.5X10-5

U-236 (medium lung absorption) (e) 4.0X101 1.1X103 2.0X10-2 5.4X10-1 2.4X10-6 6.5X10-5

U-236 (slow lung absorption) (f) 4.0X101 1.1X103 6.0X10-3 1.6X10-1 2.4X10-6 6.5X10-5

U-238 (all lung absorption types) (d),(e),(f) Unlimited Unlimited Unlimited Unlimited 1.2X10-8 3.4X10-7

U (nat) Unlimited Unlimited Unlimited Unlimited 2.6X10-8 7.1X10-7

U (enriched to 20% or less) (g) Unlimited Unlimited Unlimited Unlimited See Table A-4 See Table A-4

U (dep) Unlimited Unlimited Unlimited Unlimited See Table A-4 (See Table A-3)

V-48 Vanadium (23) 4.0X10-1 1.1X101 4.0X10-1 1.1X101 6.3X103 1.7X105

V-49 4.0X101 1.1X103 4.0X101 1.1X103 3.0X102 8.1X103

W-178 (a) Tungsten (74) 9.0 2.4X102 5.0 1.4X102 1.3X103 3.4X104

W-181 3.0X101 8.1X102 3.0X101 8.1X102 2.2X102 6.0X103

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W-185 4.0X101 1.1X103 8.0X10-1 2.2X101 3.5X102
9.4X103

W-187 2.0 5.4X101 6.0X10-1 1.6X101 2.6X104
7.0X105

W-188 (a) 4.0X10-1 1.1X101 3.0X10-1 8.1
3.7X102 1.0X104

Xe-122 (a) Xenon (54) 4.0X10-
1 1.1X101 4.0X10-1 1.1X101 4.8X104 1.3X106

Xe-123 2.0 5.4X101 7.0X10-1 1.9X101
4.4X105 1.2X107

Xe-127 4.0 1.1X102 2.0 5.4X101
1.0X103 2.8X104

Xe-131m 4.0X101 1.1X103 4.0X101 1.1X103
3.1X103 8.4X104

Xe-133 2.0X101 5.4X102 1.0X101 2.7X102
6.9X103 1.9X105

Xe-135 3.0 8.1X101 2.0 5.4X101
9.5X104 2.6X106

Y-87 (a) Yttrium (39) 1.0 2.7X101 1.0 2.7X101
1.7X104 4.5X105

Y-88 4.0X10-1 1.1X101 4.0X10-1 1.1X101 5.2X102
1.4X104

Y-90 3.0X10-1 8.1 3.0X10-1 8.1 2.0X104
5.4X105

a A1 and/or A2 values include contributions from daughter nuclides with half-lives less than 10 days.

b The values of A1 and A2 in Curies (Ci) are approximate and for information only; the regulatory standard units are Terabecquerels (TBq), (see Appendix A to § 175.105--Determination of A1 and A2, Section I.).

c The quantity may be determined from a measurement of the rate of decay or a measurement of the radiation level at a prescribed distance from the source.

d These values apply only to compounds of uranium that take the chemical form of UF6, UO2F2 and UO2(NO3)2 in both normal and accident conditions of transport.

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e These values apply only to compounds of uranium that take the chemical form of UO3, UF4, UCl4 and hexavalent compounds in both normal and accident conditions of transport.

f These values apply to all compounds of uranium other than those specified in notes (d) and (e) of this table.

g These values apply to unirradiated uranium only.

h A1 = 0.1 TBq (2.7 Ci) and A2 = 0.001 TBq (0.027 Ci) for Cf-252 for domestic use.

i A2 = 0.74 TBq (20 Ci) for Mo-99 for domestic use.

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

```
*******************************************************************************
******** This is piece 1. -- It begins at character 1 of table line 1. ********
*******************************************************************************

Y-91                          6.0X10-1    1.6X101     6.0X10-1    1.6X101
Y-91m                         2.0         5.4X101     2.0         5.4X101
Y-92                          2.0X10-1    5.4         2.0X10-1    5.4
Y-93                          3.0X10-1    8.1         3.0X10-1    8.1
Yb-169       Ytterbium (70)   4.0         1.1X102     1.0         2.7X101
Yb-175                        3.0X101     8.1X102     9.0X10-1    2.4X101
Zn-65        Zinc (30)        2.0         5.4X101     2.0         5.4X101
Zn-69                         3.0         8.1X101     6.0X10-1    1.6X101
Zn-69m (a)                    3.0         8.1X101     6.0X10-1    1.6X101
Zr-88        Zirconium (40)   3.0         8.1X101     3.0         8.1X101
Zr-93                         Unlimited   Unlimited   Unlimited   Unlimited
Zr-95 (a)                     2.0         5.4X101     8.0X10-1    2.2X101
Zr-97 (a)                     4.0X10-1    1.1X101     4.0X10-1    1.1X101
1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+..

*******************************************************************************
******* This is piece 2. -- It begins at character 78 of table line 1. ********
*******************************************************************************

 9.1X102    2.5X104
 1.5X106    4.2X107
 3.6X105    9.6X106
 1.2X105    3.3X106
```

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```
 8.9X102    2.4X104
 6.6X103    1.8X105
 3.0X102    8.2X103
 1.8X106    4.9X107
 1.2X105    3.3X106
 6.6X102    1.8X104
 9.3X10-5   2.5X10-3
 7.9X102    2.1X104
 7.1X104    1.9X106
78.....+...90....+..
```

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

```
*******************************************************************************

******** This is piece 1. -- It begins at character 1 of table line 1. ********

*******************************************************************************
```

Y-91		6.0X10-1	1.6X101	6.0X10-1	1.6X101
Y-91m		2.0	5.4X101	2.0	5.4X101
Y-92		2.0X10-1	5.4	2.0X10-1	5.4
Y-93		3.0X10-1	8.1	3.0X10-1	8.1
Yb-169	Ytterbium (70)	4.0	1.1X102	1.0	2.7X101
Yb-175		3.0X101	8.1X102	9.0X10-1	2.4X101
Zn-65	Zinc (30)	2.0	5.4X101	2.0	5.4X101
Zn-69		3.0	8.1X101	6.0X10-1	1.6X101
Zn-69m (a)		3.0	8.1X101	6.0X10-1	1.6X101
Zr-88	Zirconium (40)	3.0	8.1X101	3.0	8.1X101
Zr-93		Unlimited	Unlimited	Unlimited	Unlimited
Zr-95 (a)		2.0	5.4X101	8.0X10-1	2.2X101

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```
Zr-97 (a)                      4.0X10-1    1.1X101  4.0X10-1    1.1X101

1...k...10....k...20....k...30....k...40....k...50....k...60....k...70....k..

******************************************************************************

******* This is piece 2. -- It begins at character 78 of table line 1. ********

******************************************************************************

 9.1X102    2.5X104

 1.5X106    4.2X107

 3.6X105    9.6X106

 1.2X105    3.3X106

 8.9X102    2.4X104

 6.6X103    1.8X105

 3.0X102    8.2X103

 1.8X106    4.9X107

 1.2X105    3.3X106

 6.6X102    1.8X104

 9.3X10-5   2.5X10-3

 7.9X102    2.1X104

 7.1X104    1.9X106

78.....k...90....k..

 WIDETABLE

    NOTE--Some parts of this form are wider than one screen.  To view

    material that exceeds the width of this screen, use the right arrow

    key.  To return to the original screen, use the left arrow key.
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Y-91		6.0X10-1	1.6X101	6.0X10-1	1.6X101	9.1X102	2.5X104
Y-91m		2.0	5.4X101	2.0	5.4X101	1.5X106	4.2X107
Y-92		2.0X10-1	5.4	2.0X10-1	5.4	3.6X105	9.6X106
Y-93		3.0X10-1	8.1	3.0X10-1	8.1	1.2X105	3.3X106
Yb-169	Ytterbium (70)	4.0	1.1X102	1.0	2.7X101	8.9X102	2.4X104
Yb-175		3.0X101	8.1X102	9.0X10-1	2.4X101	6.6X103	1.8X105
Zn-65	Zinc (30)	2.0	5.4X101	2.0	5.4X101	3.0X102	8.2X103
Zn-69		3.0	8.1X101	6.0X10-1	1.6X101	1.8X106	4.9X107
Zn-69m (a)		3.0	8.1X101	6.0X10-1	1.6X101	1.2X105	3.3X106
Zr-88	Zirconium (40)	3.0	8.1X101	3.0	8.1X101	6.6X102	1.8X104
Zr-93		Unlimited	Unlimited	Unlimited	Unlimited	9.3X10-5	2.5X10-3
Zr-95 (a)		2.0	5.4X101	8.0X10-1	2.2X101	7.9X102	2.1X104
Zr-97 (a)		4.0X10-1	1.1X101	4.0X10-1	1.1X101	7.1X104	1.9X106

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

********** This is piece 1. -- It begins at character 1 of table line 1. ********

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****

TABLE A-2
EXEMPT MATERIAL ACTIVITY
CONCENTRATIONS AND EXEMPT
CONSIGNMENT ACTIVITY LIMITS
FOR RADIONUCLIDES

Symbol of radionuclide	Element and atomic number	Activity concentration for exempt material (Bq/g)

EXEMPT MATERIAL ACTIVITY
CONCENTRATIONS AND EXEMPT
CONSIGNMENT ACTIVITY LIMITS
FOR RADIONUCLIDES

Symbol of radionuclide	Element and atomic number	Activity concentration for exempt material (Bq/g)
Ac-225	Actinium (89)	1.0X101
Ac-227		1.0X10-1
Ac-228		1.0X101
Ag-105	Silver (47)	1.0X102
Ag-108m (b)		1.0X101
Ag-110m		1.0X101
Ag-111		1.0X103
Al-26	Aluminum (13)	1.0X101
Am-241	Americium (95)	1.0
Am-242m (b)		1.0
Am-243 (b)		1.0
Ar-37	Argon (18)	1.0X106
Ar-39		1.0X107
Ar-41		1.0X102
As-72	Arsenic (33)	1.0X101
As-73		1.0X103
As-74		1.0X101
As-76		1.0X102
As-77		1.0X103
At-211	Astatine (85)	1.0X103
Au-193	Gold (79)	1.0X102
Au-194		1.0X101
Au-195		1.0X102
Au-198		1.0X102
Au-199		1.0X102
Ba-131	Barium (56)	1.0X102
Ba-133		1.0X102
Ba-133m		1.0X102
Ba-140 (b)		1.0X101
Be-7	Beryllium (4)	1.0X103
Be-10		1.0X104
Bi-205	Bismuth (83)	1.0X101
Bi-206		1.0X101
Bi-207		1.0X101
Bi-210		1.0X103

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Bi-210m		1.0X101
Bi-212 (b)		1.0X101
Bk-247	Berkelium (97)	1.0
Bk-249		1.0X103
Br-76	Bromine (35)	1.0X101
Br-77		1.0X102
Br-82		1.0X101
C-11	Carbon (6)	1.0X101
C-14		1.0X104
Ca-41	Calcium (20)	1.0X105
Ca-45		1.0X104
Ca-47		1.0X101
Cd-109	Cadmium (48)	1.0X104
Cd-113m		1.0X103
Cd-115		1.0X102
Cd-115m		1.0X103
Ce-139	Cerium (58)	1.0X102
Ce-141		1.0X102
Ce-143		1.0X102
Ce-144 (b)		1.0X102
Cf-248	Californium (98)	1.0X101
Cf-249		1.0
Cf-250		1.0X101
Cf-251		1.0
Cf-252		1.0X101
Cf-253		1.0X102
Cf-254		1.0
Cl-36	Chlorine (17)	1.0X104
Cl-38		1.0X101
Cm-240	Curium (96)	1.0X102
Cm-241		1.0X102
Cm-242		1.0X102
Cm-243		1.0
Cm-244		1.0X101
Cm-245		1.0
Cm-246		1.0
Cm-247		1.0
Cm-248		1.0
Co-55	Cobalt (27)	1.0X101
Co-56		1.0X101
Co-57		1.0X102
Co-58		1.0X101
Co-58m		1.0X104
Co-60		1.0X101
Cr-51	Chromium (24)	1.0X103
Cs-129	Cesium (55)	1.0X102
Cs-131		1.0X103
Cs-132		1.0X101
Cs-134		1.0X101
Cs-134m		1.0X103
Cs-135		1.0X104
Cs-136		1.0X101

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Cs-137 (b)		1.0X101
Cu-64	Copper (29)	1.0X102
Cu-67		1.0X102
Dy-159	Dysprosium (66)	1.0X103
Dy-165		1.0X103
Dy-166		1.0X103
Er-169	Erbium (68)	1.0X104
Er-171		1.0X102
Eu-147	Europium (63)	1.0X102
Eu-148		1.0X101
Eu-149		1.0X102
Eu-150 (short lived)		1.0X103
Eu-150 (long lived)		1.0X101
Eu-152		1.0X101
Eu-152m		1.0X102
Eu-154		1.0X101
Eu-155		1.0X102
Eu-156		1.0X101
F-18	Fluorine (9)	1.0X101
Fe-52	Iron (26)	1.0X101
Fe-55		1.0X104
Fe-59		1.0X101
Fe-60		1.0X102
Ga-67	Gallium (31)	1.0X102
Ga-68		1.0X101
Ga-72		1.0X101
Gd-146	Gadolinium (64)	1.0X101
Gd-148		1.0X101
Gd-153		1.0X102
Gd-159		1.0X103
Ge-68	Germanium (32)	1.0X101
Ge-71		1.0X104

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

****
******* This is piece 2. -- It begins at character 80 of table line 1. ********
****

Activity concentration for exempt material (Ci/g)	Activity limit for exempt consignment (Bq)	Activity limit for exempt consignment (Ci)

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Activity concentration for exempt material (Ci/g)	Activity limit for exempt consignment (Bq)	Activity limit for exempt consignment (Ci)
2.7X10-10	1.0X104	2.7X10-7
2.7X10-12	1.0X103	2.7X10-8
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-11	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-5	1.0X108	2.7X10-3
2.7X10-4	1.0X104	2.7X10-7
2.7X10-9	1.0X109	2.7X10-2
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-8	1.0X106	2.7X10-5
2.7X10-8	1.0X107	2.7X10-4
2.7X10-9	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-6	1.0X107	2.7X10-4
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5

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2.7X10-8	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X105	2.7X10-6
2.7X10-11	1.0X103	2.7X10-8
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-11	1.0X103	2.7X10-8
2.7X10-11	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-9	1.0X105	2.7X10-6
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-8	1.0X105	2.7X10-6
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-8	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5

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2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-7	1.0X108	2.7X10-3

80..+...90....+....0....+...10....+...20....+...30....+...40....+...50....+....

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****

******** This is piece 1. -- It begins at character 1 of table line 1. ********

****

TABLE A-2

EXEMPT MATERIAL ACTIVITY

CONCENTRATIONS AND EXEMPT

CONSIGNMENT ACTIVITY LIMITS

FOR RADIONUCLIDES

Symbol of radionuclide Element and Activity concentration for

atomic exempt material (Bq/g)

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number

EXEMPT MATERIAL ACTIVITY

CONCENTRATIONS AND EXEMPT

CONSIGNMENT ACTIVITY LIMITS

FOR RADIONUCLIDES

Symbol of radionuclide	Element and atomic number	Activity concentration for exempt material (Bq/g)
Ac-225	Actinium (89)	1.0X101
Ac-227		1.0X10-1
Ac-228		1.0X101
Ag-105	Silver (47)	1.0X102
Ag-108m (b)		1.0X101
Ag-110m		1.0X101
Ag-111		1.0X103
Al-26	Aluminum (13)	1.0X101
Am-241	Americium (95)	1.0
Am-242m (b)		1.0
Am-243 (b)		1.0
Ar-37	Argon (18)	1.0X106
Ar-39		1.0X107
Ar-41		1.0X102
As-72	Arsenic (33)	1.0X101
As-73		1.0X103

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As-74		1.0X101
As-76		1.0X102
As-77		1.0X103
At-211	Astatine (85)	1.0X103
Au-193	Gold (79)	1.0X102
Au-194		1.0X101
Au-195		1.0X102
Au-198		1.0X102
Au-199		1.0X102
Ba-131	Barium (56)	1.0X102
Ba-133		1.0X102
Ba-133m		1.0X102
Ba-140 (b)		1.0X101
Be-7	Beryllium (4)	1.0X103
Be-10		1.0X104
Bi-205	Bismuth (83)	1.0X101
Bi-206		1.0X101
Bi-207		1.0X101
Bi-210		1.0X103
Bi-210m		1.0X101
Bi-212 (b)		1.0X101
Bk-247	Berkelium (97)	1.0
Bk-249		1.0X103

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Br-76	Bromine (35)	1.0X101
Br-77		1.0X102
Br-82		1.0X101
C-11	Carbon (6)	1.0X101
C-14		1.0X104
Ca-41	Calcium (20)	1.0X105
Ca-45		1.0X104
Ca-47		1.0X101
Cd-109	Cadmium (48)	1.0X104
Cd-113m		1.0X103
Cd-115		1.0X102
Cd-115m		1.0X103
Ce-139	Cerium (58)	1.0X102
Ce-141		1.0X102
Ce-143		1.0X102
Ce-144 (b)		1.0X102
Cf-248	Californium (98)	1.0X101
Cf-249		1.0
Cf-250		1.0X101
Cf-251		1.0
Cf-252		1.0X101
Cf-253		1.0X102
Cf-254		1.0

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Cl-36	Chlorine (17)	1.0X104
Cl-38		1.0X101
Cm-240	Curium (96)	1.0X102
Cm-241		1.0X102
Cm-242		1.0X102
Cm-243		1.0
Cm-244		1.0X101
Cm-245		1.0
Cm-246		1.0
Cm-247		1.0
Cm-248		1.0
Co-55	Cobalt (27)	1.0X101
Co-56		1.0X101
Co-57		1.0X102
Co-58		1.0X101
Co-58m		1.0X104
Co-60		1.0X101
Cr-51	Chromium (24)	1.0X103
Cs-129	Cesium (55)	1.0X102
Cs-131		1.0X103
Cs-132		1.0X101
Cs-134		1.0X101
Cs-134m		1.0X103

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Cs-135		1.0X104
Cs-136		1.0X101
Cs-137 (b)		1.0X101
Cu-64	Copper (29)	1.0X102
Cu-67		1.0X102
Dy-159	Dysprosium	1.0X103
(66)		
Dy-165		1.0X103
Dy-166		1.0X103
Er-169	Erbium (68)	1.0X104
Er-171		1.0X102
Eu-147	Europium (63)	1.0X102
Eu-148		1.0X101
Eu-149		1.0X102
Eu-150 (short lived)		1.0X103
Eu-150 (long lived)		1.0X101
Eu-152		1.0X101
Eu-152m		1.0X102
Eu-154		1.0X101
Eu-155		1.0X102
Eu-156		1.0X101
F-18	Fluorine (9)	1.0X101
Fe-52	Iron (26)	1.0X101
Fe-55		1.0X104

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Fe-59		1.0X101
Fe-60		1.0X102
Ga-67	Gallium (31)	1.0X102
Ga-68		1.0X101
Ga-72		1.0X101
Gd-146 (64)	Gadolinium	1.0X101
Gd-148		1.0X101
Gd-153		1.0X102
Gd-159		1.0X103
Ge-68	Germanium (32)	1.0X101
Ge-71		1.0X104

1...k...10....k...20....k...30....k...40....k...50....k...60....k... 70....k....

****

******* This is piece 2. -- It begins at character 80 of table line 1. ********

****

Activity concentration for exempt material (Ci/g)	Activity limit for exempt consignment (Bq)	Activity limit for exempt consignment (Ci)
Activity concentration for exempt material (Ci/g)	Activity limit for exempt consignment (Bq)	Activity limit for exempt consignment (Ci)
2.7X10-10	1.0X104	2.7X10-7

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2.7X10-12	1.0X103	2.7X10-8
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-11	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-5	1.0X108	2.7X10-3
2.7X10-4	1.0X104	2.7X10-7
2.7X10-9	1.0X109	2.7X10-2
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-8	1.0X106	2.7X10-5
2.7X10-8	1.0X107	2.7X10-4
2.7X10-9	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5

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2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-6	1.0X107	2.7X10-4
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X106	2.7X10-5

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2.7X10-8	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X105	2.7X10-6
2.7X10-11	1.0X103	2.7X10-8
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-11	1.0X104	2.7X10-7
2.7X10-10	1.0X104	2.7X10-7
2.7X10-11	1.0X103	2.7X10-8
2.7X10-11	1.0X103	2.7X10-8
2.7X10-11	1.0X104	2.7X10-7

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2.7X10-11	1.0X103	2.7X10-8
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X105	2.7X10-6
2.7X10-8	1.0X107	2.7X10-4
2.7X10-9	1.0X105	2.7X10-6
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-8	1.0X105	2.7X10-6
2.7X10-7	1.0X107	2.7X10-4
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-8	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-8	1.0X106	2.7X10-5
2.7X10-7	1.0X107	2.7X10-4
2.7X10-9	1.0X106	2.7X10-5

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2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X107	2.7X10-4
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-7	1.0X106	2.7X10-5
2.7X10-10	1.0X106	2.7X10-5
2.7X10-9	1.0X105	2.7X10-6
2.7X10-9	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X105	2.7X10-6
2.7X10-10	1.0X106	2.7X10-5
2.7X10-10	1.0X104	2.7X10-7
2.7X10-9	1.0X107	2.7X10-4
2.7X10-8	1.0X106	2.7X10-5
2.7X10-10	1.0X105	2.7X10-6
2.7X10-7	1.0X108	2.7X10-3

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80..k...90....k....0....k...10....k...20....k...30....k...40....k... 50....k....

WIDETABLE

NOTE--Some parts of this form are wider than one screen. To view

material that exceeds the width of this screen, use the right arrow

key. To return to the original screen, use the left arrow key.

TABLE A-2

EXEMPT MATERIAL ACTIVITY

CONCENTRATIONS AND EXEMPT

CONSIGNMENT ACTIVITY LIMITS

FOR RADIONUCLIDES

Symbol of radionuclide Element and Activity concentration for Activity concentration for Activity limit for Activity limit for

atomic exempt material (Bq/g) exempt material (Ci/g) exempt consignment exempt consignment

number (Bq) (Ci)

EXEMPT MATERIAL ACTIVITY

CONCENTRATIONS AND EXEMPT

CONSIGNMENT ACTIVITY LIMITS

FOR RADIONUCLIDES

Symbol of radionuclide Element and Activity concentration for Activity concentration for Activity limit for Activity limit for

atomic exempt material (Bq/g) exempt material (Ci/g) exempt consignment exempt consignment

number (Bq) (Ci)

Ac-225 Actinium (89)
1.0X101 2.7X10-10 1.0X104 2.7X10-7

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Ac-227 1.0X10-1 2.7X10-12 1.0X103
2.7X10-8

Ac-
228 1.0X101 2.7X1
0-10 1.0X106 2.7X10-5

Ag-105 Silver (47)
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Ag-108m (b)
1.0X101 2.7X10-10 1.0X106 2.7X10-5

Ag-110m
1.0X101 2.7X10-10 1.0X106 2.7X10-5

Ag-
111 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Al-26 Aluminum (13)
1.0X101 2.7X10-10 1.0X105 2.7X10-6

Am-241 Americium (95) 1.0
2.7X10-11 1.0X104 2.7X10-7

Am-242m (b) 1.0
2.7X10-11 1.0X104 2.7X10-7

Am-243 (b) 1.0
2.7X10-11 1.0X103 2.7X10-8

Ar-37 Argon
(18) 1.0X106 2.7X10-5 1.0X108
2.7X10-3

Ar-39
1.0X107 2.7X10-4 1.0X104 2.7X10-7

Ar-41
1.0X102 2.7X10-9 1.0X109 2.7X10-2

As-72 Arsenic
(33) 1.0X101 2.7X10-10 1.0X105
2.7X10-6

As-73
1.0X103 2.7X10-8 1.0X107 2.7X10-4

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As-74
1.0X101 2.7X10-10 1.0X106 2.7X10-5

As-76
1.0X102 2.7X10-9 1.0X105 2.7X10-6

As-77
1.0X103 2.7X10-8 1.0X106 2.7X10-5

At-211 Astatine (85)
1.0X103 2.7X10-8 1.0X107 2.7X10-4

Au-193 Gold (79)
1.0X102 2.7X10-9 1.0X107 2.7X10-4

Au-
194 1.0X101 2.7X1
0-10 1.0X106 2.7X10-5

Au-
195 1.0X102 2.7X1
0-9 1.0X107 2.7X10-4

Au-
198 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Au-
199 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Ba-131 Barium (56)
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Ba-
133 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Ba-133m
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Ba-140
(b) 1.0X101 2.7X10-
10 1.0X105 2.7X10-6

Be-7 Beryllium (4)
1.0X103 2.7X10-8 1.0X107 2.7X10-4

Be-10
1.0X104 2.7X10-7 1.0X106 2.7X10-5

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Bi-205 Bismuth
(83) 1.0X101 2.7X10-10 1.0X106
2.7X10-5

Bi-
206 1.0X101 2.7X1
0-10 1.0X105 2.7X10-6

Bi-
207 1.0X101 2.7X1
0-10 1.0X106 2.7X10-5

Bi-
210 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Bi-210m
1.0X101 2.7X10-10 1.0X105 2.7X10-6

Bi-212
(b) 1.0X101 2.7X10-
10 1.0X105 2.7X10-6

Bk-247 Berkelium (97) 1.0
2.7X10-11 1.0X104 2.7X10-7

Bk-
249 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Br-76 Bromine
(35) 1.0X101 2.7X10-10 1.0X105
2.7X10-6

Br-77
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Br-82
1.0X101 2.7X10-10 1.0X106 2.7X10-5

C-11 Carbon
(6) 1.0X101 2.7X10-10 1.0X106
2.7X10-5

C-
14 1.0X104 2.7X
10-7 1.0X107 2.7X10-4

Ca-41 Calcium

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(20) 1.0X105 2.7X10-6 1.0X107
2.7X10-4

Ca-45
1.0X104 2.7X10-7 1.0X107 2.7X10-4

Ca-47
1.0X101 2.7X10-10 1.0X106 2.7X10-5

Cd-109 Cadmium
(48) 1.0X104 2.7X10-7 1.0X106
2.7X10-5

Cd-113m
1.0X103 2.7X10-8 1.0X106 2.7X10-5

Cd-
115 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Cd-115m
1.0X103 2.7X10-8 1.0X106 2.7X10-5

Ce-139 Cerium (58)
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Ce-
141 1.0X102 2.7X1
0-9 1.0X107 2.7X10-4

Ce-
143 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Ce-144
(b) 1.0X102 2.7X10-
9 1.0X105 2.7X10-6

Cf-248 Californium
1.0X101 2.7X10-10 1.0X104 2.7X10-7

(98)

Cf-249 1.0
2.7X10-11 1.0X103 2.7X10-8

Cf-
250 1.0X101 2.7X1
0-10 1.0X104 2.7X10-7

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Cf-251 1.0
2.7X10-11 1.0X103 2.7X10-8

Cf-
252 1.0X101 2.7X1
0-10 1.0X104 2.7X10-7

Cf-
253 1.0X102 2.7X1
0-9 1.0X105 2.7X10-6

Cf-254 1.0
2.7X10-11 1.0X103 2.7X10-8

Cl-36 Chlorine (17)
1.0X104 2.7X10-7 1.0X106 2.7X10-5

Cl-38
1.0X101 2.7X10-10 1.0X105 2.7X10-6

Cm-240 Curium (96)
1.0X102 2.7X10-9 1.0X105 2.7X10-6

Cm-
241 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Cm-
242 1.0X102 2.7X1
0-9 1.0X105 2.7X10-6

Cm-243 1.0
2.7X10-11 1.0X104 2.7X10-7

Cm-
244 1.0X101 2.7X1
0-10 1.0X104 2.7X10-7

Cm-245 1.0
2.7X10-11 1.0X103 2.7X10-8

Cm-246 1.0
2.7X10-11 1.0X103 2.7X10-8

Cm-247 1.0
2.7X10-11 1.0X104 2.7X10-7

Cm-248 1.0

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2.7X10-11 1.0X103 2.7X10-8

Co-55 Cobalt (27)
1.0X101 2.7X10-10 1.0X106 2.7X10-5

Co-56
1.0X101 2.7X10-10 1.0X105 2.7X10-6

Co-57
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Co-58
1.0X101 2.7X10-10 1.0X106 2.7X10-5

Co-
58m 1.0X104 2.7X1
0-7 1.0X107 2.7X10-4

Co-60
1.0X101 2.7X10-10 1.0X105 2.7X10-6

Cr-51 Chromium (24)
1.0X103 2.7X10-8 1.0X107 2.7X10-4

Cs-129 Cesium (55)
1.0X102 2.7X10-9 1.0X105 2.7X10-6

Cs-
131 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Cs-
132 1.0X101 2.7X1
0-10 1.0X105 2.7X10-6

Cs-
134 1.0X101 2.7X1
0-10 1.0X104 2.7X10-7

Cs-134m
1.0X103 2.7X10-8 1.0X105 2.7X10-6

Cs-
135 1.0X104 2.7X1
0-7 1.0X107 2.7X10-4

Cs-
136 1.0X101 2.7X1
0-10 1.0X105 2.7X10-6

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Cs-137
(b) 1.0X101 2.7X10-
10 1.0X104 2.7X10-7

Cu-64 Copper (29)
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Cu-67
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Dy-
159 Dysprosium 1.0X103 2.7X1
0-8 1.0X107 2.7X10-4

(66)

Dy-
165 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Dy-
166 1.0X103 2.7X1
0-8 1.0X106 2.7X10-5

Er-169 Erbium (68)
1.0X104 2.7X10-7 1.0X107 2.7X10-4

Er-
171 1.0X102 2.7X1
0-9 1.0X106 2.7X10-5

Eu-147 Europium (63)
1.0X102 2.7X10-9 1.0X106 2.7X10-5

Eu-
148 1.0X101 2.7X1
0-10 1.0X106 2.7X10-5

Eu-
149 1.0X102 2.7X1
0-9 1.0X107 2.7X10-4

Eu-150 (short
lived) 1.0X103 2.7X10-
8 1.0X106 2.7X10-5

Eu-150 (long lived)
1.0X101 2.7X10-10 1.0X106 2.7X10-5

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Eu-152 1.0X101 2.7X10-10 1.0X106 2.7X10-5

Eu-152m 1.0X102 2.7X10-9 1.0X106 2.7X10-5

Eu-154 1.0X101 2.7X10-10 1.0X106 2.7X10-5

Eu-155 1.0X102 2.7X10-9 1.0X107 2.7X10-4

Eu-156 1.0X101 2.7X10-10 1.0X106 2.7X10-5

F-18 Fluorine (9) 1.0X101 2.7X10-10 1.0X106 2.7X10-5

Fe-52 Iron (26) 1.0X101 2.7X10-10 1.0X106 2.7X10-5

Fe-55 1.0X104 2.7X10-7 1.0X106 2.7X10-5

Fe-59 1.0X101 2.7X10-10 1.0X106 2.7X10-5

Fe-60 1.0X102 2.7X10-9 1.0X105 2.7X10-6

Ga-67 Gallium (31) 1.0X102 2.7X10-9 1.0X106 2.7X10-5

Ga-68 1.0X101 2.7X10-10 1.0X105 2.7X10-6

Ga-72 1.0X101 2.7X10-10 1.0X105 2.7X10-6

Gd-146 Gadolinium 1.0X101 2.7X10-10 1.0X106 2.7X10-5

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(64)

Gd-148 1.0X101 2.7X10-10 1.0X104 2.7X10-7

Gd-153 1.0X102 2.7X10-9 1.0X107 2.7X10-4

Gd-159 1.0X103 2.7X10-8 1.0X106 2.7X10-5

Ge-68 Germanium (32) 1.0X101 2.7X10-10 1.0X105 2.7X10-6

Ge-71 1.0X104 2.7X10-7 1.0X108 2.7X10-3

Ge-77 1.0X101 2.7X10-10 1.0X105 2.7X10-6

Hf-172	Hafnium (72)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Hf-175		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Hf-181		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Hf-182		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Hg-194	Mercury (80)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Hg-195m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Hg-197		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Hg-197m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Hg-203		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Ho-166	Holmium (67)	1.0X103	2.7X10-8	1.0X105	2.7X10-6
Ho-166m		1.0X101	2.7X10-10	1.0X106	2.7X10-5
I-123	Iodine (53)	1.0X102	2.7X10-9	1.0X107	2.7X10-4
I-124		1.0X101	2.7X10-10	1.0X106	2.7X10-5
I-125		1.0X103	2.7X10-8	1.0X106	2.7X10-5
I-126		1.0X102	2.7X10-9	1.0X106	2.7X10-5
I-129		1.0X102	2.7X10-9	1.0X105	2.7X10-6
I-131		1.0X102	2.7X10-9	1.0X106	2.7X10-5
I-132		1.0X101	2.7X10-10	1.0X105	2.7X10-6
I-133		1.0X101	2.7X10-10	1.0X106	2.7X10-5
I-134		1.0X101	2.7X10-10	1.0X105	2.7X10-6
I-135		1.0X101	2.7X10-10	1.0X106	2.7X10-5
In-111	Indium (49)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
In-113m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
In-114m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
In-115m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Ir-189	Iridium (77)	1.0X102	2.7X10-9	1.0X107	2.7X10-4
Ir-190		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Ir-192		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Ir-194		1.0X102	2.7X10-9	1.0X105	2.7X10-6

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K-40	Potassium (19)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
K-42		1.0X102	2.7X10-9	1.0X106	2.7X10-5
K-43		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Kr-81	Krypton (36)	1.0X104	2.7X10-7	1.0X107	2.7X10-4
Kr-85		1.0X105	2.7X10-6	1.0X104	2.7X10-7
Kr-85m		1.0X103	2.7X10-8	1.0X1010	2.7X10-1
Kr-87		1.0X102	2.7X10-9	1.0X109	2.7X10-2
La-137	Lanthanum (57)	1.0X103	2.7X10-8	1.0X107	2.7X10-4
La-140		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Lu-172	Lutetium (71)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Lu-173		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Lu-174		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Lu-174m		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Lu-177		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Mg-28	Magnesium (12)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
Mn-52	Manganese (25)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
Mn-53		1.0X104	2.7X10-7	1.0X109	2.7X10-2
Mn-54		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Mn-56		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Mo-93	Molybdenum (42)	1.0X103	2.7X10-8	1.0X108	2.7X10-3
Mo-99		1.0X102	2.7X10-9	1.0X106	2.7X10-5
N-13	Nitrogen (7)	1.0X102	2.7X10-9	1.0X109	2.7X10-2
Na-22	Sodium (11)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Na-24		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Nb-93m	Niobium (41)	1.0X104	2.7X10-7	1.0X107	2.7X10-4
Nb-94		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Nb-95		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Nb-97		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Nd-147	Neodymium (60)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
Nd-149		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Ni-59	Nickel (28)	1.0X104	2.7X10-7	1.0X108	2.7X10-3
Ni-63		1.0X105	2.7X10-6	1.0X108	2.7X10-3
Ni-65		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Np-235	Neptunium (93)	1.0X103	2.7X10-8	1.0X107	2.7X10-4
Np-236 (short-lived)		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Np-236 (long-lived)		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Np-237 (b)		1.0	2.7X10-11	1.0X103	2.7X10-8
Np-239		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Os-185	Osmium (76)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Os-191		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Os-191m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Os-193		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Os-194		1.0X102	2.7X10-9	1.0X105	2.7X10-6
P-32	Phosphorus (15)	1.0X103	2.7X10-8	1.0X105	2.7X10-6
P-33		1.0X105	2.7X10-6	1.0X108	2.7X10-3

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Pa-230	Protactinium (91)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Pa-231		1.0	2.7X10-11	1.0X103	2.7X10-8
Pa-233		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Pb-201	Lead (82)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Pb-202		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Pb-203		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Pb-205		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Pb-210 (b)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Pb-212 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Pd-103	Palladium (46)	1.0X103	2.7X10-8	1.0X108	2.7X10-3
Pd-107		1.0X105	2.7X10-6	1.0X108	2.7X10-3
Pd-109		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Pm-143	Promethium (61)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
Pm-144		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Pm-145		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Pm-147		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Pm-148m		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Pm-149		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Pm-151		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Po-210	Polonium (84)	1.0X101	2.7X10-10	1.0X104	2.7X10-7
Pr-142	Praseodymium (59)	1.0X102	2.7X10-9	1.0X105	2.7X10-6
Pr-143		1.0X104	2.7X10-7	1.0X106	2.7X10-5
Pt-188	Platinum (78)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Pt-191		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Pt-193		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Pt-193m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Pt-195m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Pt-197		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Pt-197m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Pu-236	Plutonium (94)	1.0X101	2.7X10-10	1.0X104	2.7X10-7
Pu-237		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Pu-238		1.0	2.7X10-11	1.0X104	2.7X10-7
Pu-239		1.0	2.7X10-11	1.0X104	2.7X10-7
Pu-240		1.0	2.7X10-11	1.0X103	2.7X10-8
Pu-241		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Pu-242		1.0	2.7X10-11	1.0X104	2.7X10-7
Pu-244		1.0	2.7X10-11	1.0X104	2.7X10-7
Ra-223 (b)	Radium (88)	1.0X102	2.7X10-9	1.0X105	2.7X10-6
Ra-224 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Ra-225		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Ra-226 (b)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Ra-228 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Rb-81	Rubidium (37)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Rb-83		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Rb-84		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Rb-86		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Rb-87		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Rb(nat)		1.0X104	2.7X10-7	1.0X107	2.7X10-4

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Re-184	Rhenium (75)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Re-184m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Re-186		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Ge-77		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Hf-172	Hafnium (72)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Hf-175		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Hf-181		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Hf-182		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Hg-194	Mercury (80)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Hg-195m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Hg-197		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Hg-197m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Hg-203		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Ho-166	Holmium (67)	1.0X103	2.7X10-8	1.0X105	2.7X10- 6
Ho-166m		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
I-123	Iodine (53)	1.0X102	2.7X10-9	1.0X107	2.7X10- 4
I-124		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
I-125		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
I-126		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
I-129		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
I-131		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
I-132		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
I-133		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
I-134		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
I-135		1.0X101	2.7X10-10	1.0X106	2.7X10- 5

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In-111	Indium (49)	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
In-113m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
In-114m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
In-115m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Ir-189	Iridium (77)	1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Ir-190		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Ir-192		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Ir-194		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
K-40 (19)	Potassium	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
K-42		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
K-43		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Kr-81	Krypton (36)	1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Kr-85		1.0X105	2.7X10-6	1.0X104	2.7X10- 7
Kr-85m		1.0X103	2.7X10-8	1.0X1010	2.7X10- 1
Kr-87		1.0X102	2.7X10-9	1.0X109	2.7X10- 2
La-137 (57)	Lanthanum	1.0X103	2.7X10-8	1.0X107	2.7X10- 4
La-140		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Lu-172	Lutetium (71)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Lu-173		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Lu-174		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Lu-174m		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Lu-177		1.0X103	2.7X10-8	1.0X107	2.7X10- 4

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Mg-28	Magnesium	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
(12)					
Mn-52	Manganese	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
(25)					
Mn-53		1.0X104	2.7X10-7	1.0X109	2.7X10- 2
Mn-54		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Mn-56		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Mo-93	Molybdenum	1.0X103	2.7X10-8	1.0X108	2.7X10- 3
(42)					
Mo-99		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
N-13	Nitrogen (7)	1.0X102	2.7X10-9	1.0X109	2.7X10- 2
Na-22	Sodium (11)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Na-24		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Nb-93m	Niobium (41)	1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Nb-94		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Nb-95		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Nb-97		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Nd-147	Neodymium	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
(60)					
Nd-149		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Ni-59	Nickel (28)	1.0X104	2.7X10-7	1.0X108	2.7X10- 3
Ni-63		1.0X105	2.7X10-6	1.0X108	2.7X10- 3
Ni-65		1.0X101	2.7X10-10	1.0X106	2.7X10- 5

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Np-235	Neptunium	1.0X103	2.7X10-8	1.0X107	2.7X10- 4
(93)					
Np-236 (short-lived)		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Np-236 (long-lived)		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Np-237 (b)		1.0	2.7X10-11	1.0X103	2.7X10- 8
Np-239		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Os-185	Osmium (76)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Os-191		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Os-191m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Os-193		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Os-194		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
P-32	Phosphorus	1.0X103	2.7X10-8	1.0X105	2.7X10- 6
(15)					
P-33		1.0X105	2.7X10-6	1.0X108	2.7X10- 3
Pa-230	Protactinium	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
(91)					
Pa-231		1.0	2.7X10-11	1.0X103	2.7X10- 8
Pa-233		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Pb-201	Lead (82)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Pb-202		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Pb-203		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Pb-205		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Pb-210 (b)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Pb-212 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10- 6

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Pd-103	Palladium	1.0X103	2.7X10-8	1.0X108	2.7X10- 3
(46)					
Pd-107		1.0X105	2.7X10-6	1.0X108	2.7X10- 3
Pd-109		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Pm-143	Promethium	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
(61)					
Pm-144		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Pm-145		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Pm-147		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Pm-148m		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Pm-149		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Pm-151		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Po-210	Polonium (84)	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Pr-142	Praseodymium	1.0X102	2.7X10-9	1.0X105	2.7X10- 6
(59)					
Pr-143		1.0X104	2.7X10-7	1.0X106	2.7X10- 5
Pt-188	Platinum (78)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Pt-191		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Pt-193		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Pt-193m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Pt-195m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Pt-197		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Pt-197m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5

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Pu-236	Plutonium	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
(94)					
Pu-237		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Pu-238		1.0	2.7X10-11	1.0X104	2.7X10- 7
Pu-239		1.0	2.7X10-11	1.0X104	2.7X10- 7
Pu-240		1.0	2.7X10-11	1.0X103	2.7X10- 8
Pu-241		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Pu-242		1.0	2.7X10-11	1.0X104	2.7X10- 7
Pu-244		1.0	2.7X10-11	1.0X104	2.7X10- 7
Ra-223 (b)	Radium (88)	1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Ra-224 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Ra-225		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Ra-226 (b)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Ra-228 (b)		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Rb-81	Rubidium (37)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Rb-83		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Rb-84		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Rb-86		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Rb-87		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Rb(nat)		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Re-184	Rhenium (75)	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Re-184m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Re-186		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Re-187		1.0X106	2.7X10-5	1.0X109	2.7X10-2
Re-188		1.0X102	2.7X10-9	1.0X105	2.7X10-6

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Re-189		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Re(nat)		1.0X106	2.7X10-5	1.0X109	2.7X10-2
Rh-99	Rhodium (45)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Rh-101		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Rh-102		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Rh-102m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Rh-103m		1.0X104	2.7X10-7	1.0X108	2.7X10-3
Rh-105		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Rn-222 (b)	Radon (86)	1.0X101	2.7X10-10	1.0X108	2.7X10-3
Ru-97	Ruthenium (44)	1.0X102	2.7X10-9	1.0X107	2.7X10-4
Ru-103		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Ru-105		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Ru-106 (b)		1.0X102	2.7X10-9	1.0X105	2.7X10-6
S-35	Sulphur (16)	1.0X105	2.7X10-6	1.0X108	2.7X10-3
Sb-122	Antimony (51)	1.0X102	2.7X10-9	1.0X104	2.7X10-7
Sb-124		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Sb-125		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sb-126		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Sc-44	Scandium (21)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
Sc-46		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Sc-47		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sc-48		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Se-75	Selenium (34)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
Se-79		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Si-31	Silicon (14)	1.0X103	2.7X10-8	1.0X106	2.7X10-5
Si-32		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Sm-145	Samarium (62)	1.0X102	2.7X10-9	1.0X107	2.7X10-4
Sm-147		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Sm-151		1.0X104	2.7X10-7	1.0X108	2.7X10-3
Sm-153		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sn-113	Tin (50)	1.0X103	2.7X10-8	1.0X107	2.7X10-4
Sn-117m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sn-119m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Sn-121m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Sn-123		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Sn-125		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Sn-126		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Sr-82	Strontium (38)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
Sr-85		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sr-85m		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Sr-87m		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Sr-89		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Sr-90 (b)		1.0X102	2.7X10-9	1.0X104	2.7X10-7

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Sr-91		1.0X101	2.7X10-10	1.0X105	2.7X10-6
Sr-92		1.0X101	2.7X10-10	1.0X106	2.7X10-5
T(H-3)	Tritium (1)	1.0X106	2.7X10-5	1.0X109	2.7X10-2
Ta-178 (long-lived)	Tantalum (73)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Ta-179		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Ta-182		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Tb-157	Terbium (65)	1.0X104	2.7X10-7	1.0X107	2.7X10-4
Tb-158		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tb-160		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tc-95m	Techneti-um (43)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tc-96		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tc-96m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Tc-97		1.0X103	2.7X10-8	1.0X108	2.7X10-3
Tc-97m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Tc-98		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tc-99		1.0X104	2.7X10-7	1.0X107	2.7X10-4
Tc-99m		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Te-121	Tellurium (52)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Te-121m		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Te-123m		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Te-125m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Te-127		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Te-127m		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Te-129		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Te-129m		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Te-131m		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Te-132		1.0X102	2.7X10-9	1.0X107	2.7X10-4
Th-227	Thorium (90)	1.0X101	2.7X10-10	1.0X104	2.7X10-7
Th-228 (b)		1.0	2.7X10-11	1.0X104	2.7X10-7
Th-229 (b)		1.0	2.7X10-11	1.0X103	2.7X10-8
Th-230		1.0	2.7X10-11	1.0X104	2.7X10-7
Th-231		1.0X103	2.7X10-8	1.0X107	2.7X10-4
Th-232		1.0X101	2.7X10-10	1.0X104	2.7X10-7
Th-234 (b)		1.0X103	2.7X10-8	1.0X105	2.7X10-6
Th (nat) (b)		1.0	2.7X10-11	1.0X103	2.7X10-8
Ti-44	Titanium (22)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
Tl-200	Thallium (81)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Tl-201		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Tl-202		1.0X102	2.7X10-9	1.0X106	2.7X10-5
Tl-204		1.0X104	2.7X10-7	1.0X104	2.7X10-7
Tm-167	Thulium (69)	1.0X102	2.7X10-9	1.0X106	2.7X10-5
Tm-170		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Tm-171		1.0X104	2.7X10-7	1.0X108	2.7X10-3
U-230 (fast lung	Uranium	1.0X101	2.7X10-10	1.0X105	2.7X10-6

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absorption) (b),(d)	(92)				
U-230 (medium lung absorption) (e)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-230 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-232 (fast lung absorption) (b),(d)		1.0	2.7X10-11	1.0X103	2.7X10-8
U-232 (medium lung absorption) (e)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-232 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-233 (fast lung absorption) (d)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-233 (medium lung absorption) (e)		1.0X102	2.7X10-9	1.0X105	2.7X10-6
U-233 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X105	2.7X10-6
U-234 (fast lung absorption) (d)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-234 (medium lung absorption) (e)		1.0X102	2.7X10-9	1.0X105	2.7X10-6
U-234 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X105	2.7X10-6
U-235 (all lung absorption types) (b),(d),(e),(f)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-236 (fast lung absorption) (d)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-236 (medium lung absorption) (e)		1.0X102	2.7X10-9	1.0X105	2.7X10-6
U-236 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U-238 (all lung absorption types) (b),(d),(e),(f)		1.0X101	2.7X10-10	1.0X104	2.7X10-7
U (nat) (b)		1.0	2.7X10-11	1.0X103	2.7X10-8
U (enriched to 20% or less) (g)		1.0	2.7X10-11	1.0X103	2.7X10-8
U (dep)		1.0	2.7X10-11	1.0X103	2.7X10-8
V-48	Vanadium (23)	1.0X101	2.7X10-10	1.0X105	2.7X10-6
V-49		1.0X104	2.7X10-7	1.0X107	2.7X10-4
W-178	Tungsten (74)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
W-181		1.0X103	2.7X10-8	1.0X107	2.7X10-4
W-185		1.0X104	2.7X10-7	1.0X107	2.7X10-4
W-187		1.0X102	2.7X10-9	1.0X106	2.7X10-5
W-188		1.0X102	2.7X10-9	1.0X105	2.7X10-6
Xe-122	Xenon (54)	1.0X102	2.7X10-9	1.0X109	2.7X10-2
Xe-123		1.0X102	2.7X10-9	1.0X109	2.7X10-2
Xe-127		1.0X103	2.7X10-8	1.0X105	2.7X10-6
Xe-131m		1.0X104	2.7X10-7	1.0X104	2.7X10-7

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Xe-133		1.0X103	2.7X10-8	1.0X104	2.7X10-7
Xe-135		1.0X103	2.7X10-8	1.0X1010	2.7X10-1
Y-87	Yttrium (39)	1.0X101	2.7X10-10	1.0X106	2.7X10-5
Y-88		1.0X101	2.7X10-10	1.0X106	2.7X10-5
Y-90		1.0X103	2.7X10-8	1.0X105	2.7X10-6
Y-91		1.0X103	2.7X10-8	1.0X106	2.7X10-5
Re-187		1.0X106	2.7X10-5	1.0X109	2.7X10- 2
Re-188		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Re-189		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Re(nat)		1.0X106	2.7X10-5	1.0X109	2.7X10- 2
Rh-99 (45)	Rhodium	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Rh-101		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Rh-102		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Rh-102m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Rh-103m		1.0X104	2.7X10-7	1.0X108	2.7X10- 3
Rh-105		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Rn-222 (b) (86)	Radon	1.0X101	2.7X10-10	1.0X108	2.7X10- 3
Ru-97 (44)	Ruthenium	1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Ru-103		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Ru-105		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Ru-106 (b)		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
S-35 (16)	Sulphur	1.0X105	2.7X10-6	1.0X108	2.7X10- 3

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Sb-122	Antimony	1.0X102	2.7X10-9	1.0X104	2.7X10- 7
(51)					
Sb-124		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Sb-125		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sb-126		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Sc-44	Scandium	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
(21)					
Sc-46		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Sc-47		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sc-48		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Se-75	Selenium	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
(34)					
Se-79		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Si-31	Silicon	1.0X103	2.7X10-8	1.0X106	2.7X10- 5
(14)					
Si-32		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Sm-145	Samarium	1.0X102	2.7X10-9	1.0X107	2.7X10- 4
(62)					
Sm-147		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Sm-151		1.0X104	2.7X10-7	1.0X108	2.7X10- 3
Sm-153		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sn-113	Tin (50)	1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Sn-117m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sn-119m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4

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Sn-121m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Sn-123		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Sn-125		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Sn-126		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Sr-82 (38)	Strontium	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Sr-85		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sr-85m		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Sr-87m		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Sr-89		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Sr-90 (b)		1.0X102	2.7X10-9	1.0X104	2.7X10- 7
Sr-91		1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Sr-92		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
T(H-3) (1)	Tritium	1.0X106	2.7X10-5	1.0X109	2.7X10- 2
Ta-178 (long-lived) (73)	Tantalum	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Ta-179		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Ta-182		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Tb-157 (65)	Terbium	1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Tb-158		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Tb-160		1.0X101	2.7X10-10	1.0X106	2.7X10- 5

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Tc-95m	Techneti-	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
um (43)					
Tc-96		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Tc-96m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Tc-97		1.0X103	2.7X10-8	1.0X108	2.7X10- 3
Tc-97m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Tc-98		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Tc-99		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
Tc-99m		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Te-121	Tellurium	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
(52)					
Te-121m		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Te-123m		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Te-125m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Te-127		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Te-127m		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Te-129		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Te-129m		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Te-131m		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Te-132		1.0X102	2.7X10-9	1.0X107	2.7X10- 4
Th-227	Thorium	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
(90)					
Th-228 (b)		1.0	2.7X10-11	1.0X104	2.7X10- 7
Th-229 (b)		1.0	2.7X10-11	1.0X103	2.7X10- 8

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Th-230		1.0	2.7X10-11	1.0X104	2.7X10- 7
Th-231		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
Th-232		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
Th-234 (b)		1.0X103	2.7X10-8	1.0X105	2.7X10- 6
Th (nat) (b)		1.0	2.7X10-11	1.0X103	2.7X10- 8
Ti-44 (22)	Titanium	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
Tl-200 (81)	Thallium	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Tl-201		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Tl-202		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Tl-204		1.0X104	2.7X10-7	1.0X104	2.7X10- 7
Tm-167 (69)	Thulium	1.0X102	2.7X10-9	1.0X106	2.7X10- 5
Tm-170		1.0X103	2.7X10-8	1.0X106	2.7X10- 5
Tm-171		1.0X104	2.7X10-7	1.0X108	2.7X10- 3
U-230 (fast lung absorption) (b),(d)	Uranium (92)	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
U-230 (medium lung absorption) (e)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
U-230 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
U-232 (fast lung		1.0	2.7X10-11	1.0X103	2.7X10- 8

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absorption) (b),(d)				
U-232 (medium lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption) (e)				
U-232 (slow lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption) (f)				
U-233 (fast lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption) (d)				
U-233 (medium lung	1.0X102	2.7X10-9	1.0X105	2.7X10- 6
absorption) (e)				
U-233 (slow lung	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
absorption) (f)				
U-234 (fast lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption) (d)				
U-234 (medium lung	1.0X102	2.7X10-9	1.0X105	2.7X10- 6
absorption) (e)				
U-234 (slow lung	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
absorption) (f)				
U-235 (all lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption types)				
(b),(d),(e),(f)				
U-236 (fast lung	1.0X101	2.7X10-10	1.0X104	2.7X10- 7
absorption) (d)				
U-236 (medium lung	1.0X102	2.7X10-9	1.0X105	2.7X10- 6
absorption) (e)				

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U-236 (slow lung absorption) (f)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
U-238 (all lung absorption types) (b),(d),(e),(f)		1.0X101	2.7X10-10	1.0X104	2.7X10- 7
U (nat) (b)		1.0	2.7X10-11	1.0X103	2.7X10- 8
U (enriched to 20% or less) (g)		1.0	2.7X10-11	1.0X103	2.7X10- 8
U (dep)		1.0	2.7X10-11	1.0X103	2.7X10- 8
V-48 (23)	Vanadium	1.0X101	2.7X10-10	1.0X105	2.7X10- 6
V-49		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
W-178 (74)	Tungsten	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
W-181		1.0X103	2.7X10-8	1.0X107	2.7X10- 4
W-185		1.0X104	2.7X10-7	1.0X107	2.7X10- 4
W-187		1.0X102	2.7X10-9	1.0X106	2.7X10- 5
W-188		1.0X102	2.7X10-9	1.0X105	2.7X10- 6
Xe-122 (54)	Xenon	1.0X102	2.7X10-9	1.0X109	2.7X10- 2
Xe-123		1.0X102	2.7X10-9	1.0X109	2.7X10- 2
Xe-127		1.0X103	2.7X10-8	1.0X105	2.7X10- 6
Xe-131m		1.0X104	2.7X10-7	1.0X104	2.7X10- 7

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Xe-133		1.0X103	2.7X10-8	1.0X104	2.7X10- 7
Xe-135		1.0X103	2.7X10-8	1.0X1010	2.7X10- 1
Y-87 (39)	Yttrium	1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Y-88		1.0X101	2.7X10-10	1.0X106	2.7X10- 5
Y-90		1.0X103	2.7X10-8	1.0X105	2.7X10- 6
Y-91		1.0X103	2.7X10-8	1.0X106	2.7X10- 5

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****
******** This is piece 1. -- It begins at character 1 of table line 1. ********
****

Y-91m		1.0X102	2.7X10-9	1.0X106
Y-92		1.0X102	2.7X10-9	1.0X105
Y-93		1.0X102	2.7X10-9	1.0X105
Yb-169	Ytterbium (70)	1.0X102	2.7X10-9	1.0X107
Yb-175		1.0X103	2.7X10-8	1.0X107
Zn-65	Zinc (30)	1.0X101	2.7X10-10	1.0X106
Zn-69		1.0X104	2.7X10-7	1.0X106
Zn-69m		1.0X102	2.7X10-9	1.0X106
Zr-88	Zirconium (40)	1.0X102	2.7X10-9	1.0X106
Zr-93 (b)		1.0X103	2.7X10-8	1.0X107
Zr-95		1.0X101	2.7X10-10	1.0X106
Zr-97 (b)		1.0X101	2.7X10-10	1.0X105

[cf11,10a
[Reserved]
[cf11,10b
Parent nuclides and their progeny included in secular equilibrium are listed in the following:

Sr-90	Y-90

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Zr-93	Nb-93m
Zr-97	Nb-97
Ru-106	Rh-106
Cs-137	Ba-137m
Ce-134	La-134
Ce-144	Pr-144
Ba-140	La-140
Bi-212	Tl-208 (0.36), Po-212 (0.64)
Pb-210	Bi-210, Po-210
Pb-212	Bi-212, Tl-208 (0.36), Po-212 (0.64)
Rn-220	Po-216
Rn-222	Po-218, Pb-214, Bi-214, Po-214
Ra-223	Rn-219, Po-215, Pb-211, Bi-211, Tl-207
Ra-224	Rn-220, Po-216, Pb-212, Bi-212, Tl-208(0.36), Po-212 (0.64)
Ra-226	Rn-222, Po-218, Pb-214, Bi-214, Po-214, Pb-210, Bi-210, Po-210
Ra-228	Ac-228
Th-226	Ra-222, Rn-218, Po-214
Th-228	Ra-224, Rn-220, Po-216, Pb-212, Bi-212, Tl-208 (0.36), Po-212 (0.64)
Th-229	Ra-225, Ac-225, Fr-221, At-217, Bi-213, Po-213, Pb-209
Th-nat	Ra-228, Ac-228, Th-228, Ra-224, Rn-220, Po-216, Pb-212, Bi-212, Tl-208 (0.36), Po-212 (0.64)
Th-234	Pa-234m
U-230	Th-226, Ra-222, Rn-218, Po-214
U-232	Th-228, Ra-224, Rn-220, Po-216, Pb-212, Bi-212, Tl-208 (0.36), Po-212 (0.64)
U-235	Th-231
U-238	Th-234, Pa-234m
U-nat	Th-234, Pa-234m, U-234, Th-230, Ra-226, Rn-222, Po-218, Pb-214, Bi-214, Po-214, Pb-210, Bi-210, Po-210
U-240	Np-240m
Np-237	Pa-233
Am-242m	Am-242
Am-243	Np-239 [cf11,10c[Reserved] [cf11,10d These values

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apply only to compounds of uranium that take the chemical form of UF6, UO2F2 and UO2(NO3)2 in both normal and accident conditions of transport. [cf11,10e These values apply only to compounds of uranium that take the chemical form of UO3, UF4, UCl4 and hexavalent compounds in both normal and accident conditions of transport. [cf11,10f These values apply to all compounds of uranium other than those specified in notes (d) and (e) of this table. [cf11,10g These values apply to unirradiated uranium only.

TABLE A-3
GENERAL VALUES FOR A[cf171 AND A[cf172

Contents	A[cf17,101[cf3,4 ,3	(TBq)	(Ci)	A[cf17,10-2[cf3,4 ,3
Only beta or gamma emitting radionuclides are known to be present	1 x 10-1	2.7 x 100	2 x 10 -2	5.4 x 10-1
Only alpha emitting radionuclides are known to be present	2 x 10-1	5.4 x 100	9 x 10-5	2.4 x 10-3
No relevant data are available	1 x 10-3	2.7 x 10-2	9 x 10-5	2.4 x 10-3

TABLE A-4
ACTIVITY-MASS RELATIONSHIPS FOR URANIUM

Uranium Enrichment 1 wt % U-235 present	Specific Activity ,3	TBq/g	Ci/g
0.45	1.8 x 10-8	5.0 x 10-7	
0.72	2.6 x 10-8	7.1 x 10-7	
1	2.8 x 10-8	7.6 x 10-7	

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1.5	3.7 x 10-8	1.0 x 10-6
5	1.0 x 10-7	2.7 x 10-6
10	1.8 x 10-7	4.8 x 10-6
20	3.7 x 10-7	1.0 x 10-5
35	7.4 x 10-7	2.0 x 10-5
50	9.3 x 10-7	2.5 x 10-5
90	2.2 x 10-6	5.8 x 10-5
93	2.6 x 10-6	7.0 x 10-5
95	3.4 x 10-6	9.1 x 10-5

1 The figures for uranium include representative values for the activity of the uranium-234 that is concentrated during the enrichment process. (60 FR 50264, Sept. 28, 1995 as amended at 61 FR 28724, June 6, 1996; 69 FR 3800, Jan. 26, 2004).

1...+...10....+...20....+...30....+...40....+...50....+...60....+...70....+....

****
******* This is piece 2. -- It begins at character 80 of table line 1. ********
****

2.7X10-5
2.7X10-6
2.7X10-6
2.7X10-4
2.7X10-4
2.7X10-5
2.7X10-5
2.7X10-5
2.7X10-5
2.7X10-4
2.7X10-5
2.7X10-6a

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(TBq)	(Ci)	Activity concentration for exempt material (Bq/g)	Activity concentration for exempt material (Ci/g)	Activity limits for exempt consignments (Bq)
1 x 101	2.7 x1-0-10	1 x 104	2.7 x10-7	
1 x 10-1	2.7 x1-0-12	1 x 103	2.7 x10-8	
1 x 10-1	2.7 x 10-12	1 x 103	2.7 x 10-8	

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80..+...90....+....0....+...10....+...20....+...30....+...40....+

****
******* This is piece 3. -- It begins at character 145 of table line 1. *******
****

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Activity
limits for
exempt
consignments
(Ci)

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145.....+...6

[Note: The following TABLE/FORM is too wide to be displayed on one screen. You must print it for a meaningful review of its contents. The table has been divided into multiple pieces with each piece containing information to help you assemble a printout of the table. The information for each piece includes: (1) a three line message preceding the tabular data showing by line # and character # the position of the upper left-hand corner of the piece and the position of the piece within the entire table; and (2) a numeric scale following the tabular data displaying the character positions.]

****

******** This is piece 1. -- It begins at character 1 of table line 1. ********

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****

Y-91m		1.0X102	2.7X10-9	1.0X106
Y-92		1.0X102	2.7X10-9	1.0X105
Y-93		1.0X102	2.7X10-9	1.0X105
Yb-169	Ytterbium (70)	1.0X102	2.7X10-9	1.0X107
Yb-175		1.0X103	2.7X10-8	1.0X107
Zn-65	Zinc (30)	1.0X101	2.7X10-10	1.0X106
Zn-69		1.0X104	2.7X10-7	1.0X106
Zn-69m		1.0X102	2.7X10-9	1.0X106
Zr-88	Zirconium (40)	1.0X102	2.7X10-9	1.0X106
Zr-93 (b)		1.0X103	2.7X10-8	1.0X107
Zr-95		1.0X101	2.7X10-10	1.0X106
Zr-97 (b)		1.0X101	2.7X10-10	1.0X105

[cf11,10a

[Reserved]

[cf11,10b

Parent nuclides and their progeny included in secular equilibrium are listed in the following:

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Sr-90	Y-90
Zr-93	Nb-93m
Zr-97	Nb-97
Ru-106	Rh-106
Cs-137	Ba-137m
Ce-134	La-134
Ce-144	Pr-144
Ba-140	La-140
Bi-212	Tl-208 (0.36), Po-212 (0.64)
Pb-210	Bi-210, Po-210
Pb-212	Bi-212, Tl-208 (0.36), Po-212 (0.64)
Rn-220	Po-216
Rn-222	Po-218, Pb-214, Bi-214, Po-214
Ra-223	Rn-219, Po-215, Pb-211, Bi-211, Tl-207
Ra-224	Rn-220, Po-216, Pb-212, Bi-212, Tl-208(0.36), Po-212 (0.64)
Ra-226	Rn-222, Po-218, Pb-214, Bi-214, Po-214, Pb-210, Bi-210, Po-210

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Ra-228 Ac-228

Th-226 Ra-222, Rn-218, Po-214

Th-228 Ra-224, Rn-220, Po-216,

Pb-212, Bi-212, Tl-208

(0.36), Po-212 (0.64)

Th-229 Ra-225, Ac-225, Fr-221,

At-217, Bi-213, Po-213,

Pb-209

Th-nat Ra-228, Ac-228, Th-228,

Ra-224, Rn-220, Po-216,

Pb-212, Bi-212, Tl-208

(0.36), Po-212 (0.64)

Th-234 Pa-234m

U-230 Th-226, Ra-222, Rn-218,

Po-214

U-232 Th-228, Ra-224, Rn-220,

Po-216, Pb-212, Bi-212,

Tl-208 (0.36), Po-212

(0.64)

U-235 Th-231

U-238 Th-234, Pa-234m

U-nat Th-234, Pa-234m, U-234,

Th-230, Ra-226, Rn-222,

Po-218, Pb-214, Bi-214,

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Po-214, Pb-210, Bi-210,

Po-210

U-240 Np-240m

Np-237 Pa-233

Am-242m Am-242

Am-243 Np-239 [cf11,10c[Reserved]

[cf11,10d These values

apply only to compounds of

uranium that take the

chemical form of UF6,

UO2F2 and UO2(NO3)2 in

both normal and accident

conditions of transport.

[cf11,10e These values

apply only to compounds of

uranium that take the

chemical form of UO3, UF4,

UCl4 and hexavalent

compounds in both normal

and accident conditions of

transport. [cf11,10f These

values apply to all

compounds of uranium other

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than those specified in

notes (d) and (e) of this

table. [cf11,10g These

values apply to

unirradiated uranium only.

TABLE A-3

GENERAL VALUES

FOR A[cf171 AND

A[cf172

Contents	A[cf17,101[cf3,4 ,3	(TBq)	(Ci)	A[cf17,10-2[cf3,4 ,3
Only beta or gamma emitting radionuclides are known to be present	1 x 10-1	2.7 x 100	2 x 10 -2	5.4 x 10- 1
Only alpha emitting radionuclides are known to be present	2 x 10-1	5.4 x 100	9 x 10-5	2.4 x 10- 3
No relevant data are available	1 x 10-3	2.7 x 10-2	9 x 10-5	2.4 x 10- 3

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TABLE A-4

ACTIVITY-MASS

RELATIONSHIPS

FOR URANIUM

Uranium Enrichment 1 wt % U-235 present	Specific Activity ,3	TBq/g Ci/g
0.45	1.8 x 10-8	5.0 x 10-7
0.72	2.6 x 10-8	7.1 x 10-7
1	2.8 x 10-8	7.6 x 10-7
1.5	3.7 x 10-8	1.0 x 10-6
5	1.0 x 10-7	2.7 x 10-6
10	1.8 x 10-7	4.8 x 10-6
20	3.7 x 10-7	1.0 x 10-5
35	7.4 x 10-7	2.0 x 10-5
50	9.3 x 10-7	2.5 x 10-5
90	2.2 x 10-6	5.8 x 10-5
93	2.6 x 10-6	7.0 x 10-5
95	3.4 x 10-6	9.1 x 10-5

1 The figures for uranium include representative

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values for the

activity of the

uranium-234

that is

concentrated

during the

enrichment

process. (60 FR

50264, Sept.

28, 1995 as

amended at 61

FR 28724, June

6, 1996; 69 FR

3800, Jan. 26,

2004).

1...k...10....k...20....k...30....k...40....k...50....k...60....k... 70....k....

****

******* This is piece 2. -- It begins at character 80 of table line 1. ********

****

2.7X10-5

2.7X10-6

2.7X10-6

2.7X10-4

2.7X10-4

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2.7X10-5

2.7X10-5

2.7X10-5

2.7X10-5

2.7X10-4

2.7X10-5

2.7X10-6a

(TBq) (Ci) Activity Activity Activity

concentration concentration limits for

for exempt for exempt exempt

material material consignments

(Bq/g) (Ci/g) (Bq)

1 x 101 2.7 x1- 1 x 104 2.7 x10-7

0-10

1 x 10-1 2.7 x1- 1 x 103 2.7 x10-8

0-12

1 x 10-1 2.7 x 1 x 103 2.7 x 10-8

10-12

80..k...90....k....0....k...10....k...20....k...30....k...40....k

****

******* This is piece 3. -- It begins at character 145 of table line 1. *******

****

Activity

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limits for

exempt

consignments

(Ci)

145.....k...6

WIDETABLE

NOTE--Some parts of this form are wider than one screen. To view

material that exceeds the width of this screen, use the right arrow

key. To return to the original screen, use the left arrow key.

Y-91m 1.0X102 2.7X10-9
1.0X106 2.7X10-5

Y-92 1.0X102 2.7X10-9
1.0X105 2.7X10-6

Y-93 1.0X102 2.7X10-9
1.0X105 2.7X10-6

Yb-169 Ytterbium (70) 1.0X102 2.7X10-9
1.0X107 2.7X10-4

Yb-175 1.0X103 2.7X10-8
1.0X107 2.7X10-4

Zn-65 Zinc (30) 1.0X101 2.7X10-
10 1.0X106 2.7X10-5

Zn-69 1.0X104 2.7X10-7
1.0X106 2.7X10-5

Zn-69m 1.0X102 2.7X10-9
1.0X106 2.7X10-5

Zr-88 Zirconium (40) 1.0X102 2.7X10-9
1.0X106 2.7X10-5

Zr-93 (b) 1.0X103 2.7X10-8
1.0X107 2.7X10-4

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Zr-95 1.0X101 2.7X10-
10 1.0X106 2.7X10-5

Zr-97 (b) 1.0X101 2.7X10-
10 1.0X105 2.7X10-6a

[cf11,10a

[Reserved]

[cf11,10b

Parent nuclides

and their

progeny

included in

secular

equilibrium are

listed in the

following:

Sr-90	Y-90
Zr-93	Nb-93m
Zr-97	Nb-97
Ru-106	Rh-106
Cs-137	Ba-137m
Ce-134	La-134
Ce-144	Pr-144
Ba-140	La-140
Bi-212	Tl-208 (0.36), Po-212 (0.64)
Pb-210	Bi-210, Po-210

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Pb-212	Bi-212, Tl-208 (0.36), Po-212 (0.64)
Rn-220	Po-216
Rn-222	Po-218, Pb-214, Bi-214, Po-214
Ra-223	Rn-219, Po-215, Pb-211, Bi-211, Tl-207
Ra-224	Rn-220, Po-216, Pb-212, Bi-212, Tl-208(0.36), Po-212 (0.64)
Ra-226	Rn-222, Po-218, Pb-214, Bi-214, Po-214, Pb-210, Bi-210, Po-210
Ra-228	Ac-228
Th-226	Ra-222, Rn-218, Po-214
Th-228	Ra-224, Rn-220, Po-216, Pb-212, Bi-212, Tl-208 (0.36), Po-212 (0.64)
Th-229	Ra-225, Ac-225, Fr-221, At-217, Bi-213, Po-213, Pb-209
Th-nat	Ra-228, Ac-228, Th-228, Ra-224, Rn-220, Po-216, Pb-212, Bi-212, Tl-208

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(0.36), Po-212 (0.64)

Th-234 Pa-234m

U-230 Th-226, Ra-222, Rn-218,

Po-214

U-232 Th-228, Ra-224, Rn-220,

Po-216, Pb-212, Bi-212,

Tl-208 (0.36), Po-212

(0.64)

U-235 Th-231

U-238 Th-234, Pa-234m

U-nat Th-234, Pa-234m, U-234,

Th-230, Ra-226, Rn-222,

Po-218, Pb-214, Bi-214,

Po-214, Pb-210, Bi-210,

Po-210

U-240 Np-240m

Np-237 Pa-233

Am-242m Am-242

Am-243 Np-239 [cf11,10c[Reserved]

[cf11,10d These values

apply only to compounds of

uranium that take the

chemical form of UF6,

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UO2F2 and UO2(NO3)2 in

both normal and accident

conditions of transport.

[cf11,10e These values

apply only to compounds of

uranium that take the

chemical form of UO3, UF4,

UCl4 and hexavalent

compounds in both normal

and accident conditions of

transport. [cf11,10f These

values apply to all

compounds of uranium other

than those specified in

notes (d) and (e) of this

table. [cf11,10g These

values apply to

unirradiated uranium only.

TABLE A-3

GENERAL VALUES

FOR A[cf171 AND

A[cf172

Contents A[cf17,101[cf3,4 ,3 (TBq) (Ci) A[cf17,10-
(TBq) (Ci) Activity Activity Activity Activity Activity

2[cf3,4 concentration concentration limits for lim-

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its for

,3 for exempt for exempt exempt exempt

material material consignments consignments

(Bq/g) (Ci/g) (Bq) (Ci)

Only beta or 1 x 10-1 2.7 x 100 2 x 10 -2 5.4 x 10- 1 1 x 101 2.7 x1- 1 x 104 2.7 x10-7

gamma emitting 0-10

radionuclides

are known to be

present

Only alpha 2 x 10-1 5.4 x 100 9 x 10-5 2.4 x 10- 3 1 x 10-1 2.7 x1- 1 x 103 2.7 x10-8

emitting 0-12

radionuclides

are known to be

present

No relevant data 1 x 10-3 2.7 x 10-2 9 x 10-5 2.4 x 10- 3 1 x 10-1 2.7 x 1 x 103 2.7 x 10-8

are available 10-12

TABLE A-4

ACTIVITY-MASS

RELATIONSHIPS

FOR URANIUM

Uranium

Enrichment 1

wt % U-235 Specific Activity ,3 TBq/g Ci/g

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present

0.45	1.8 x 10-8	5.0 x 10-7
0.72	2.6 x 10-8	7.1 x 10-7
1	2.8 x 10-8	7.6 x 10-7
1.5	3.7 x 10-8	1.0 x 10-6
5	1.0 x 10-7	2.7 x 10-6
10	1.8 x 10-7	4.8 x 10-6
20	3.7 x 10-7	1.0 x 10-5
35	7.4 x 10-7	2.0 x 10-5
50	9.3 x 10-7	2.5 x 10-5
90	2.2 x 10-6	5.8 x 10-5
93	2.6 x 10-6	7.0 x 10-5
95	3.4 x 10-6	9.1 x 10-5

1 The figures for uranium include representative values for the activity of the uranium-234 that is concentrated during the enrichment

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process. (60 FR

50264, Sept.

28, 1995 as

amended at 61

FR 28724, June

6, 1996; 69 FR

3800, Jan. 26,

2004).

HISTORICAL NOTE

Subd. (a) par (1) subpar (ii) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (a) par (5) amended in part (various additions and renumbering) City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (a) par (6) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (b) pars (1), (2) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (b) par (3) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (c) par (3) subpar (v) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (c) par (5) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (d) par (3) subpar (xii) added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (d) par (4) subpar (i) clause (B) amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Subd. (d) par (9) subpar (iii) clause (C) opening par amended City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

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Subd. (e) par (1) subpars (iv), (v) repealed City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

Appendix A repealed and added City Record Sept. 25, 2008 eff. Oct. 25, 2008. [See Vol. 9 Statement of Basis and Purpose No. 30]

DERIVATION

Formerly Section repealed and added City Record June 30, 1999 eff. July 30, 1999. [See Vol. 9 Statements of Basis and Purpose No. 13]

NOTES:

On September 17, 2008, the Board of Health amended multiple provisions of Health Code § 175.105, including the repeal and reenactment of § 175.105 Appendix A, in order to maintain compatibility with applicable NRC regulations found in various provisions of 10 CFR Part 71 relating to the transportation and packaging of radioactive material.

24 RCNY Hlth. Code § 175.105
END OF DOCUMENT

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.201. Microwave Ovens.

(a) Applicability. The provisions of this Code relating to microwave ovens shall apply to such ovens sold, offered for sale, repaired or altered for use in homes, restaurants or other food vending establishments, hospitals or other medical care facilities, schools, and other establishments in the City where the public may be exposed.

(b) Definitions. As used in the sections of this Code relating to microwave ovens, the following definitions shall apply:

(1) "Microwave oven" means a device designed to heat, cook or dry food through the application of electromagnetic energy at frequencies assigned by the Federal Communications Commission in the normal industrial, scientific and medical heating bands ranging from 890 megahertz to 6,000 megahertz.

(2) "Cavity" means that portion of the microwave oven in which food may be heated, cooked or dried.

(3) "Door" means the movable barrier which prevents access to the cavity during operation and whose function is to prevent emission of microwave energy from the passage or opening which provides access to the cavity.

(4) "Safety interlock" means a device or system of devices which is intended to prevent generation of microwave energy when access to the cavity is possible.

(5) "Service adjustments or service procedures" means those servicing methods prescribed by the manufacturer for a specific product model.

(6) "Stirrer" means that feature of a microwave oven which is intended to provide uniform heating of the load by constantly changing the standing wave pattern within the cavity or moving the load.

(7) "External surface" means the outside surface of the cabinet or enclosure provided by the manufacturer as part of the microwave oven, including doors, door han-

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dles, latches and control knobs.

(c) Requirements for microwave ovens. (1) Power density limit. The power density of the microwave radiation emitted by a microwave oven shall not exceed 5 milliwatts (mW) per cm 2 at any point 5 cm (2 in.) or more from the external surface of the oven except that a microwave oven offered for sale or sold by its manufacturer shall not have a power density exceeding 1 mW per cm 2 at any point 5 cm (2 in.) or more from the external surface of the oven.

(2) Measurements and test conditions. (i) Microwave ovens shall be in compliance with the power density limits if the maximum reading obtained at the location of greatest microwave radiation emission does not exceed the limits specified in this section when the emission is measured through at least one stirrer cycle.

(ii) The emission shall not exceed the requirements of § 175.201(c)(1) when the microwave oven is operated at its maximum output and contains a load of 275 15 milliliters of tap water initially at 20.5 degrees Centigrade placed within the cavity at the center of the load-carrying surface provided by the manufacturer. The water container shall be a low form 600 milliliter beaker having an inside diameter of approximately 8.5 cm and made of an electrically non-conductive material such as glass or plastic.

(iii) Measurements shall be made with the door fully closed as well as with the door fixed in any other position which allows the oven to operate.

(3) Door and safety interlocks. (i) Microwave ovens manufactured prior to October 6, 1971 shall have one safety interlock.

(ii) Microwave ovens manufactured from October 6, 1971 through November 6, 1976:

(A) shall have a minimum of two operative safety interlocks, one of which shall be concealed. A concealed safety interlock on a fully assembled microwave oven must not be operable by any part of the body, or a rod 3 mm or greater in diameter and with a useful length of 10 cm. A magnetically operated interlock is considered to be concealed only if a test magnet, held in place on the oven by gravity or its own attraction, cannot operate the safety interlock. The test magnet shall have a pull at zero air gap of at least 4.5 kg and a pull at 1 cm air gap of at least 450 g when the face of the magnet which is toward the interlock switch when the magnet is in the test position is pulling against one of the large faces of a mild steel armature having dimensions of 80 mm by 50 mm by 8 mm.

(B) The insertion of an object into the oven cavity through any opening while the door is closed shall not cause microwave radiation emission from the oven to exceed the applicable power density limits specified in § 175.201(c)(1).

(iii) For microwave ovens manufactured on or after August 4, 1974:

(A) One (the primary) required safety interlock shall prevent microwave radiation emission in excess of the requirement of § 175.201(c)(1); the

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other (secondary) required safety interlock shall prevent microwave radiation emission in excess of 5 mW per cm 2 at any point 5 cm (2 in.) or more from the external surface of the oven. The two required safety interlocks shall be designated as primary or secondary in the service instructions for the oven.

(B) A means of monitoring one or both of the required safety interlocks shall be provided which shall cause the oven to become inoperable and remain so until repaired if the required safety interlock(s) should fail to perform required functions as specified in this section. Interlock failures shall not disrupt the monitoring function.

(iv) Microwave ovens manufactured on and after November 7, 1976:

(A) shall have a minimum of two operative safety interlocks. At least

one operative safety interlock on a fully assembled microwave oven shall not be operable by any part of the human body, or any object with a straight insertable length of 10 centimeters. Such interlock must also be concealed, unless its actuation is prevented when access to the interlock is possible. Any visible actuator or device to prevent actuation of this safety interlock must not be removable without disassembly of the oven or its door. A magnetically operated interlock is considered to be concealed, or its actuation is considered to be prevented, only if a test magnet held in place on the oven by gravity or its own attraction cannot operate the safety interlock. The test magnet shall be capable of lifting vertically at zero air gap at least 4.5 kilograms, and at 1 centimeter air gap at least 450 grams when the face of the magnet, which is toward the interlock when the magnet is in test position, is pulling against one of the large faces of a mild steel armature having dimensions of 80 millimeters by 50 millimeters by 8 millimeters.

(B) Microwave radiation emission from such ovens in excess of the limits specified in § 175.201(c)(1) shall not be caused by insertion of an insulated wire through any opening in the external surfaces of a fully assembled oven into the cavity, waveguide, or other microwave-energy-containing spaces while the door is closed, provided the wire, when inserted, could consist of two straight segments forming an obtuse angle of not less then 170 degrees.

(v) Failure of any single mechanical or electrical component of the microwave oven shall not cause all safety interlocks to be inoperative.

(vi) Service adjustments or service procedures on the microwave oven shall not cause the safety interlocks to become inoperative or the microwave radiation emission to exceed the power density limits of this section as a result of such service adjustments or procedures.

(4) Enforcement by the department, notice of repair and installation. (i) Any microwave oven found deficient in meeting the provisions of § 175.201(c) after survey by a Department representative shall be immediately taken out of service until such deficiencies have been corrected. The Bureau of Radiological Health shall be notified within 48 hours of the completion of such repairs.

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(ii) Within 48 hours of an installation of a microwave oven in any restaurant or other food vending establishment, hospital or other medical care facility, school, or other establishment in the City where the public may be exposed, a notification of such installation shall be made to the Bureau of Radiological Health.

(e) User instructions. (1) For microwave ovens manufactured prior to October 3, 1975 manufacturers thereof shall provide or cause to be provided, with each oven, adequate instructions for its safe use including clear warnings of precautions to be taken to avoid possible exposure to microwave radiation.

(2) For microwave ovens manufactured on or after October 3, 1975 manufacturers thereof shall provide or cause to be provided, with each oven, radiation safety instructions which:

(i) occupy a separate section and are an integral part of the regularly supplied user's manual and cookbook, if supplied separately, and are located so as to elicit the attention of the reader;

(ii) are as legible and durable as other instructions with the title emphasized to elicit the attention of the reader by such means as boldfaced type, contrasting color, a heavy-lined border, or similar means; and

(iii) contain the following wording:

"PRECAUTIONS TO AVOID POSSIBLE EXPOSURE TO EXCESSIVE MICROWAVE ENERGY
(A) Do not attempt to operate this oven with the door open since open-door operation can result in harmful exposure to microwave energy. It is important not to defeat or tamper with the safety interlocks.

(B) Do not place any object between the oven front face and the door or allow soil or cleaner residue to accumulate on sealing surfaces.

(C) Do not operate the oven if it is damaged. It is particularly important that the oven door close properly and that there is no damage to the:

(a) door (bent);

(b) hinges and latches (broken or loosened);

(c) door seals and sealing surfaces.

(D) The oven should not be adjusted or repaired by anyone except properly qualified service personnel."

(f) Service instructions. (1) For microwave ovens manufactured prior to October 3, 1975 manufacturers thereof shall provide or cause to be provided to servicing dealers and distributors and to others upon request, for each oven model, adequate instructions for service adjustment and service procedures including clear warnings of precautions to be taken to avoid possible exposure to microwave radiation.

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(2) For microwave ovens manufactured on or after October 3, 1975 manufacturers thereof shall provide or cause to be provided to servicing dealers and distributors and to others upon request, for each oven model, adequate instruction for service adjustments and service procedures, and, in addition, radiation safety instructions which:

(i) occupy a separate section and are an integral part of the regularly supplied service manual and are located so as to elicit the attention of the reader;

(ii) are as legible and durable as other instructions with the title emphasized so as to elicit the attention of the reader by such means as boldfaced type, contrasting color, a heavy-lined border, or by similar means; and

(iii) contain the following wording:

"PRECAUTIONS TO BE OBSERVED BEFORE AND DURING SERVICING TO AVOID POSSIBLE EXPOSURE TO EXCESSIVE MICROWAVE ENERGY

(A) Do not operate or allow the oven to be operated with the door open.

(B) Make the following safety checks on all ovens to be serviced before activating the magnetron or other microwave source, and make repairs as necessary:

(a) interlock operation;

(b) proper door closing;

(c) seal and sealing surfaces (arcing, wear, and other damage);

(d) damage to or loosening of hinges and latches;

(e) evidence of dropping or abuse.

(C) Before turning on microwave power for any service test or inspection within the microwave generating compartments, check the magnetron, waveguide or transmission line, and cavity for proper alignment, integrity, and connections.

(D) Any defective or misadjusted components in the interlock monitor, door seal, and microwave generation and transmission systems shall be repaired, replaced, or adjusted before the oven is released to the owner.

(E) A microwave leakage check to verify compliance with the Federal performance standard should be performed on each oven prior to release to the owner."

(iv) Include additional radiation safety precautions or instructions which may be necessary for particular oven designs or models.

(g) Warning labels on microwave ovens. (1) Microwave ovens manufactured on or after October 3, 1975 shall have the following warning labels:

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(i) A label, permanently attached to or inscribed on the oven, which shall be legible and readily viewable during normal oven use, which shall have the title emphasized and be so located as to elicit the attention of the user, and which shall bear the following warning statement:

"PRECAUTIONS FOR SAFE USE TO AVOID POSSIBLE EXPOSURE TO EXCESSIVE MICROWAVE ENERGY
DO NOT Attempt to Operate This Oven With:

(A) Object Caught in Door

(B) Door That Does Not Close Properly

(C) Damaged Door, Hinge, Latch, Sealing Surface"

(ii) A label permanently attached to or inscribed on the external surface of the oven which shall be legible and readily viewable during servicing and which shall have the word "CAUTION" emphasized and so located thereon as to elicit the attention of service personnel, and which shall bear the following warning statement:

"CAUTION: This Device is to be Serviced Only by Properly Qualified Service Personnel. Consult the Service Manual for Proper Service Procedures to Assure Continued Compliance with the Federal Performance Standard for Microwave Ovens and for Precautions to be Taken to Avoid Possible Exposure to Excessive Microwave Energy".

(iii) The labels provided in accordance with § 175.201(g)(1)(i) and (ii) shall bear only the statements specified therein, except for additional radiation safety warnings or instructions which may be necessary for particular oven designs or models.

(iv) A microwave oven model may be exempted from one or more of the radiation safety warnings specified in § 175.201(g)(1)(I) based upon a determination pursuant to the federal Radiation Control for Health and Safety Act of 1968 and the regulations promulgated thereunder that such model would continue to comply with the standards contained in § 175.201(c)(1), (2) and (3) under the adverse condition of use addressed by such precautionary statements.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.

Other
<General Materials (GM) - References, Annotations, or Tables>

24 RCNY Hlth. Code § 175.201
END OF DOCUMENT

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NEW YORK CITY CHARTER, CODE, AMENDMENTS & RULES
RULES & REGULATIONS OF THE CITY OF NEW YORK
NEW YORK CITY HEALTH CODE
TITLE IV. ENVIRONMENTAL SANITATION.
PART B. CONTROL OF ENVIRONMENT.
ARTICLE 175. RADIATION CONTROL.
RADIOACTIVE MATERIALS

The Rules & Regulations of the City of New York
is current through June 30, 2009.

§ 175.301. Television receivers and other electronic devices.

(a) No television receiver or other electronic device, whether used in the home or elsewhere, which emits radiation on application of high voltage, shall be offered, transferred or consigned for sale or use in the City of New York unless it is so constructed as to prevent radiation therefrom at a level greater than 1.29 E-7 C-kg-1 (0.5 milliroentgens per hour), measured five (5) cm (2 in.) from any accessible surface and averaged over an area of 10 cm 2 (1.55 in. 2).

(b) No replacement part shall be offered, transferred or consigned for sale or use in the City of New York which on being installed could cause the assembled unit for which it is intended to exceed the radiation limit allowed under this section.

(c) No person shall alter or adjust any television receiver or other electronic device, whether used in the home or elsewhere, which can emit radiation in such manner as to increase the radiation emission level thereof, unless the level thereby achieved be within the emission limit allowed under this section.

HISTORICAL NOTE

Section repealed and added City Record June 27, 1994 eff. July 27, 1994.

Section in original publication July 1, 1991.
<General Materials (GM) - References, Annotations, or Tables>

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