CHAPTER 33-10-04.1 STANDARDS FOR PROTECTION AGAINST RADIATION

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33-10-04.1-01. Purpose.

- 1. This chapter establishes standards for protection against ionizing radiation resulting from activities conducted pursuant to licenses or registrations issued by the department.
- 2. The requirements of this chapter are designed to control the receipt, 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 chapter. However, nothing in this chapter shall be construed as limiting actions that may be necessary to protect health and safety in an emergency.

History: Effective March 1, 1994; amended effective July 1, 1995. **General Authority:** NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.01-04

33-10-04.1-02. Scope. This chapter applies to persons licensed or registered by the department to receive, possess, use, transfer, or dispose of sources of radiation. The limits in this chapter do not apply to doses due to background radiation, to any medical administration or therapy the individual has received, to exposure from individuals administered radioactive material and released in accordance with subsection 12 of section 33-10-07-05, or to to exposure of patients to radiation for the purpose of medical diagnosis or therapy, to exposure from individuals administered radioactive material and released under section 33-10-07-05, or to exposure from the purpose of patients to radiation for the purpose of medical diagnosis or therapy, to exposure from individuals administered radioactive material and released under section 33-10-07.1-32 or to exposure from voluntary participation in medical research programs.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-03. Definitions. As used in this chapter:

- 1. <u>"Air-purifying respirator" means a respirator with an</u> <u>air-purifying filter, cartridge, or canister that removes</u> <u>specific air contaminants by passing ambient air through</u> <u>the air-purifying element.</u>
- 2. "Annual limit on intake" (ALL 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. Annual limit on intake 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 five-hundredths savored sievert [5 rem] or a committed dose equivalent of five-tenths savored sievert [50 rem] to any individual organ or tissue. Annual limit on intake values for intake by ingestion and by inhalation of selected radionuclides are given in table I, columns 1 and 2, of appendix B.
- 3. "Assigned protection factor (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 assigned protection factor.

- 4. 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 selfcontained breathing apparatus (SCBA) units.
- 25. "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 ten days, for class W, weeks, from ten to one hundred days, and for class Y, years, of greater than one hundred days. "Lung class" and "inhalation class" are equivalent terms.
- <u>36</u>. "Declared pregnant woman" means a woman who has voluntarily informed <u>her employer the licensee</u>, in writing, of her pregnancy and the estimated date of conception. <u>The declaration remains in effect until the declared pregnant woman withdraws the declaration in writing or is no longer pregnant.</u>
- 7. <u>"Demand respirator" means an atmosphere-supplying</u> respirator that admits breathing air to the facepiece only when a negative pressure is created inside the facepiece by inhalation.
- 48. "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 two thousand hours under conditions of light work, results in an intake of one annual limit on intake. The condition of light work is an inhalation rate of 1.2 cubic meters of air per hour for two thousand hours in a year. Derived air concentration values are given in table I, column 3, of appendix B.
- "Derived air concentration-hour" (DAC-hour) means the <u>59</u>. 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 thousand derived air registrant may take two concentration-hours to represent one annual limit on equivalent to a committed effective dose intake. equivalent of five-hundredths sievert [5 rem].
- 10. "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 half-mask respirator or a

<u>disposable escape-only self-contained breathing apparatus</u> (SCBA).

- 6<u>11</u>. "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.
- 12. "Filtering facepiece (dust mask)" means a negative pressure particulate respirator with a filter as an integral part of the facepiece or with the entire facepiece composed of the filtering medium, not equipped with elastomeric sealing surfaces and adjustable straps
- 13. "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.
- 14. "Fit test" means the use of a protocol to qualitatively or quantitatively evaluate the fit of a respirator on an individual.
- 15. <u>"Helmet" means a rigid respiratory inlet covering that</u> also provides head protection against impact and penetration.
- <u>16.</u> <u>"Hood" means a respiratory inlet covering that completely</u> <u>covers the head and neck and may also cover portions of</u> <u>the shoulders and torso.</u>
- 7<u>17</u>. "Inhalation class" [see "class"].
- 18. <u>"Loose-fitting facepiece" means a respiratory inlet</u> covering that is designed to form a partial seal with the face.
- 819. "Lung class" [see "class"].
- 20. "Negative pressure respirator (tight fitting)" means a respirator in which the air pressure inside the facepiece is negative during inhalation with respect to the ambient air pressure outside the repirator.
- 921. "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. "Deterministic effect" is an equivalent term.

- 1022. "Planned special exposure" means an infrequent exposure to radiation, separate from and in addition to the annual occupational dose limits.
- 23. Positive pressure respirator means a respirator in which the pressure inside the respiratory inlet covering exceeds the ambient air pressure outside the respirator.
- 24. Powered air-purifying respirator (PAPR) means an air-purifying respirator that uses a blower to force the ambient air through air-purifying elements to the inlet covering.
- 25. Pressure demand respirator means a positive pressure atmosphere-supplying respirator that admits breathing air to the facepiece when the positive pressure is reduced inside the facepiece by inhalation.
- 26. Qualitative fit test (OLFT) means a pass/fail fit test to assess the adequacy of respirator fit that relies on the individual's response to the test agent.
- 27. Quality Factor (Q) means the modifying factor (listed in tables I and II of section 33-10-01.14) that is used to derive dose equivalent from absorbed dose.
- 28. Quantitative fit test (QNFT) means an assessment of the adequacy of respirator fit by numerically measuring the amount of leakage into the respirator.
- 1129. "Quarter" means a period of time equal to onefourth of the year observed by the licensee, approximately thirteen consecutive weeks, providing that the beginning of the first quarter in a year coincides with the starting date of the year and that no day is omitted or duplicated in consecutive quarters.
- 1230. "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. A description of the reference man is contained in the International Commission on Radiological Protection report, ICRP Publication 23, "Report of the Task Group on Reference Man".
- 1331. "Respiratory protection equipment" means an apparatus, such as a respirator, used to reduce an individual's intake of airborne radioactive materials.

- 14<u>32</u>. "Sanitary sewerage" means a system of public sewers for carrying off wastewater and refuse, but excluding sewage treatment facilities, septic tanks, and leach fields owned or operated by the licensee or registrant.
- 33. <u>"Self-contained breathing apparatus (SCBA)" means an</u> <u>atmosphere-supplying respirator for which the breathing</u> <u>air source is designed to be carried by the user.</u>
- 1534. "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. "Probabilistic effect" is an equivalent term.
- <u>35.</u> "Supplied-air respirator (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.
- <u>36. "Tight-fitting facepiece" means a respiratory inlet</u> covering that forms a complete seal with the face.
- 37. "User seal check (fit check)" means an action conducted by the respirator user to determine if the respirator is properly seated to the face. Examples include negative pressure check, positive pressure check, irritant smoke check, or isoamyl acetate check.
- 1638. "Very high radiation area" means an area, accessible to individuals, in which radiation levels from radiation sources external to the body could result in an individual receiving an absorbed dose in excess of five gray [500 rad] in one hour at one meter 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.)
- 1739."Weighting factor" w_T for an organ or tissue (T)means the proportion of the risk of stochasticeffects resulting from irradiation of that organ ortissue to the total risk of stochastic effects whenthe whole body is irradiated uniformly.the whole body is effective dose equivalent, thevalues of w_T are:

ORGAN DOSE WEIGHTING	FACTORS
Organ or	
Tissue	$W_{ m T}$
Conside	0.25
Gonads	••=•
Breast	0.15
Red bone marrow	0.12
Lung	0.12
Thyroid	0.03
Bone surfaces	0.03
Remainder	0.30ª
Whole body	1.00 ^b

^a 0.30 results from 0.06 for each of five "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, $w_T = 1.0$, 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.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04, 28-32-02 Law Implemented: NDCC 23-20.1-03

33-10-04.1-04. Implementation. This chapter shall go into effect on became effective March 1, 1994, and all licensees and registrants must comply by that date except for the following:

- 1. Any existing license or registration condition that is in place prior to implementation of this chapter and is more restrictive than this chapter remains in force until there is an amendment or renewal of the license or registration.
- 2. If a license or registration condition exempts a licensee or registrant from a provision of this chapter in effect on or before March 1, 1994, it also exempts the licensee or registrant from the corresponding provision of this chapter.

3. If a license or registration condition cites provisions of this chapter in effect prior to March 1, 1994, which do not correspond to any provisions of this chapter the license or registration condition remains in force until there is an amendment or renewal of the license or registration that modifies or removes this condition.

History: Effective March 1, 1994. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-05. Radiation protection programs.

- Each licensee or registrant shall develop, document, and implement a radiation protection program sufficient to ensure compliance with the provisions of this chapter. See subsection 2 of section 33-10-04.1-15 for recordkeeping requirements relating to these programs.
- 2. To the extent practicable practical, the licensee or registrant shall use procedures and engineering controls based upon sound radiation protection principles to achieve occupational doses and public doses that are as low as is reasonably achievable (ALARA).
- 3. At intervals not to exceed twelve months, the licensee or registrant shall review the radiation protection program content and implementation.
- 4. To implement the as low as is reasonably achievable (ALARA) requirements of subsection 2, and notwithstanding the requirements of subsection 1 of section 33-10-04.1-07, a constraint on air emissions of radioactive material environment, to the excluding radon-222 and its daughters, shall be established by licensees, such that the individual member of the public likely to receive the highest dose will not be expected to receive a total effective dose equivalent in excess of one-tenth millisieverts [10 mrem] per year from these emissions. If a licensee subject to this requirement exceeds this dose constraint, the licensee shall report the exceedance as provided in subsection 3 of section 33-10-04.1-16 and promptly take appropriate corrective action to ensure against recurrence.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-06. Occupational dose limits.

- 1. Occupational dose limits for adults.
 - a. The licensee or registrant shall control the occupational dose to individual adults, except for planned special exposures pursuant to subsection 6, to the following dose limits:
 - (1) An annual limit, which is the more limiting of:
 - (a) The total effective dose equivalent being equal to five-hundredths sievert [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 being equal to fivetenths sievert [50 rem].
 - (2) The annual limits to the lens of the eye, to the skin <u>of the whole body</u>, and to the <u>skin of the</u> extremities, which are:
 - (a) An eye dose equivalent <u>A lens dose</u> <u>equivalent</u> of fifteen-hundredths sievert [15 rem], and
 - (b) A shallow dose equivalent of five-tenths sievert [50 rem] to the skin <u>of the whole</u> <u>body</u> or to <u>the skin of</u> any extremity.
 - b. 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. See paragraphs 1 and 2 of subdivision e of subsection 6.
 - c. The assigned deep dose equivalent and shallow dose equivalent shall <u>must</u> be for the <u>portion part</u> of the body receiving the highest exposure. determined as follows: The assigned shallow dose equivalent must be the dose averaged over the contiguous ten square centimeters of skin receiving the highest exposure.
 - (1) The deep dose equivalent, eye <u>lens</u> 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.

- (2) Reserved. When a protective apron is worn while working with medical fluoroscopic equipment and monitoring is conducted as specified in paragraph 5 of subdivision a of subsection 2 of section 33-10-04.1-09, the effective dose equivalent for external radiation shall be determined as follows:
 - (a) When only one individual monitoring device is used and it is located at the neck (collar) outside the protective apron, the reported deep dose equivalent shall be the effective dose equivalent for external radiation; or
 - (b) When only one individual monitoring device is used and it is located at the neck (collar) outside the protective apron, and the reported dose exceeds twenty five percent of the limit specified in subdivision a of subsection 1 of section 33-10-04.1-06, the reported deep dose equivalent value multiplied by three-tenths shall be the effective dose equivalent for external radiation; or
 - (c) When two individual monitoring devices are worn, one under the protective apron at the waist and the other 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 one and five-tenths and the deep dose equivalent reported for the individual monitoring device located at the neck outside the protective apron multiplied by fourhundredths.
 - (d) Subparagraphs b and c only apply when all of the following conditions are met:

- [1] The individual monitoring devices have not been exposed to radiation from radioactive material.
- [2] Leaded glasses, a thyroid shield, and a wrap-around protective apron have been worn whenever using the medical fluoroscopic equipment.
- [3] The area around the medical fluoroscopic equipment has been equipped with lead shielding or transparent protective barriers for control of scattered radiation.
- [4] The medical fluoroscopic procedures have been performed in a way that minimizes beam on time, such as utilizing last image hold.
- [5] Users of the medical fluoroscopic equipment must have had formal training in radiation safety and operation of medical fluoroscopic equipment.
- [6] Performance of the medical fluoroscopic equipment must be monitored and maintained via a quality assurance program.
- [7] Patient and staff radiation exposures from medical fluoroscopic equipment must be monitored and actions taken to correct problems.
- d. Derived air concentration and annual limit on intake values are presented in table I of appendix B and may be used to determine the individual's dose and to demonstrate compliance with the occupational dose limits. See subsection 7 of section 33-10-04.1-15.
- e. Notwithstanding the annual dose limits, the licensee shall limit the soluble uranium intake by an individual to ten milligrams in a week in consideration of chemical toxicity. See footnote 3 of appendix B.
- f. 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. See subdivision e of subsection 5.

2. Compliance with requirements for summation of external and internal doses.

- If the licensee or registrant is required to a. monitor pursuant to both subdivision a and subdivision b of subsection 2 of section 33-10-04.1-09, 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 subdivision a of subsection 2 of section 33-10-04.1-09 or only pursuant to subdivision b of subsection 2 of section 33-10-04.1-09, 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 subdivision b. subdivision c and subdivision d. 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.
- b. 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:
 - The sum of the fractions of the inhalation annual limit on intake for each radionuclide, or
 - (2) The total number of derived air concentrationhours (DAC-hours) for all radionuclides divided by two thousand, or
 - (3)The sum of the calculated committed effective dose equivalents to all significantly irradiated organs or tissues (T) calculated from bioassay data using appropriate 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, w_{T} , and the committed dose equivalent, $H_{T.50}$, per unit intake is greater than ten percent of the maximum weighted value of $H_{T,50}$,

that is, $w_{\scriptscriptstyle T} H_{\scriptscriptstyle T,50},$ per unit intake for any organ or tissue.

- c. Intake by oral ingestion. If the occupationally exposed individual receives an intake of radionuclides by oral ingestion greater than ten percent of the applicable oral annual limit on intake, the licensee or registrant shall account for this intake and include it in demonstrating compliance with the limits.
- d. Intake through wounds or absorption through skin. The licensee or registrant shall evaluate and, to the extent practical, account for intakes through wounds or skin absorption. The intake through intact skin has been included in the calculation of derived air concentration for hydrogen-3 and does not need to be evaluated or accounted for pursuant to this subdivision d.

3. Determination of external dose from airborne radioactive material.

- Licensees or registrants shall, when determining a. the dose from airborne radioactive material, the contribution include to the deep dose equivalent, eye dose equivalent lens dose and shallow dose equivalent from equivalent, external exposure to the radioactive cloud. See appendix B, footnotes 1 and 2.
- b. Airborne radioactivity measurements and derived air concentration 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.

- a. For purposes of assessing dose used to determine compliance with occupational dose equivalent limits, the licensee or registrant shall, when required pursuant to subsection 2 of section 33-10-04.1-09, take suitable and timely measurements of:
 - Concentrations of radioactive materials in air in work areas;

- (2) Quantities of radionuclides in the body;
- Quantities of radionuclides excreted from the body; or
- (4) Combinations of these measurements.
- b. Unless respiratory protection equipment is used, as provided in subsection 3 of section 33-10-04.1-11, or the assessment of intake is based on bioassays, the licensee or registrant shall assume that an individual inhales radioactive material at the airborne concentration in which the individual is present.
- c. 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 or registrant may:
 - 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;
 - (2) Upon prior approval of the department, adjust the derived air concentration or annual limit on intake values to reflect the actual physical and chemical characteristics of airborne radioactive material, for example, aerosol size distribution or density; and
 - (3) 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.
- d. If the licensee or registrant chooses to assess intakes of class Y material using the measurements given in paragraph 2 or 3 of subdivision a, the licensee or registrant may delay the recording and reporting of the assessments for periods up to seven months, unless otherwise required bv subsection 2 or 3 of section 33-10-04.1-16. This delay permits the licensee or registrant to make additional measurements basic to the assessments.
- e. If the identity and concentration of each radionuclide in a mixture are known, the fraction of the derived air concentration applicable to the

mixture for use in calculating derived air concentration-hours shall be either:

- (1) The sum of the ratios of the concentration to the appropriate derived air concentration value, that is, D, W, or Y, from appendix B for each radionuclide in the mixture; or
- (2) The ratio of the total concentration for all radionuclides in the mixture to the most restrictive derived air concentration value for any radionuclide in the mixture.
- f. 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 derived air concentration for the mixture shall be the most restrictive derived air concentration of any radionuclide in the mixture.
- g. When a mixture of radionuclides in air exists, a licensee or registrant may disregard certain radionuclides in the mixture if:
 - (1) The licensee or registrant uses the total activity of the mixture in demonstrating compliance with the dose limits in subsection 1 and in complying with the monitoring requirements in subdivision b of subsection 2 of section 33-10-04.1-09, and
 - (2) The concentration of any radionuclide disregarded is less than ten percent of its derived air concentration, and
 - (3) The sum of these percentages for all of the radionuclides disregarded in the mixture does not exceed thirty percent.
- h. When determining the committed effective dose equivalent, the following information may be considered:
 - (1) In order to calculate the committed effective dose equivalent, the licensee or registrant may assume that the inhalation of one annual limit on intake, or an exposure of two thousand derived air concentration-hours, results in a committed effective dose equivalent of five-hundredths sievert [5 rem] for radionuclides that have their annual limit on intakes or derived air concentrations based on the committed effective dose equivalent.

(2)For an annual limit on intake and the associated derived air concentration determined by the nonstochastic organ dose limit of five-tenths sievert [50 rem], the intake of radionuclides that would result in a committed effective dose equivalent of fivehundredths sievert [5 rem], that is, the stochastic annual limit on intake, is listed in parentheses in table I of appendix B. As a simplifying assumption, the licensee or registrant may use the stochastic annual limit on intake to determine committed effective dose equivalent. However, if the licensee or registrant uses the stochastic annual limit on intake, the licensee or registrant shall also demonstrate that the limit in subparagraph 2 of paragraph 1 of subdivision a of subsection 1 is met.

5. Determination of prior occupational dose.

- a. For each individual who is likely to receive, in a year, an occupational dose requiring monitoring pursuant to subsection 2 of section 33-10-04.1-09, the licensee or registrant shall:
 - (1) Determine the occupational radiation dose received during the current year; and
 - (2) Attempt to obtain the records of cumulative occupational radiation dose.
- b. Prior to permitting an individual to participate in a planned special exposure, the licensee or registrant shall determine:
 - The internal and external doses from all previous planned special exposures;
 - (2) All doses in excess of the limits, including doses received during accidents and emergencies, received during the lifetime of the individual; and
 - (3) All lifetime cumulative occupational radiation dose.
- c. In complying with the requirements of subdivision a of subsection 5, a licensee or registrant may:
 - (1) 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;

- (2) Accept, as the record of cumulative radiation dose, an up-to-date department's occupational radiation exposure history form (SFN 19443) 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
- Obtain reports of the individual's dose (3) equivalent from the most recent employer for work involving radiation exposure, or the if individual's current employer, the individual is not employed by the licensee or registrant, by telephone, telegram, facsimile, electronic media, or letter. The licensee or registrant shall request written а verification of the dose data if the authenticity of the transmitted report cannot be established.
- d. The licensee or registrant shall record the (1)exposure history, as required by subdivision a, on the department's occupational radiation exposure history form (SFN 19443), or other and legible record, of all the clear 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 For each period for which the exposure. licensee or registrant obtains reports, the licensee or registrant shall use the dose shown in the report in preparing the department's occupational radiation exposure history form (SFN 19443) 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 the department's occupational radiation exposure form (SFN 19443) equivalent historv or indicating the periods of time for which data are not available.

- (2)Licensees or registrants are not required to reevaluate the separate external dose equivalents and committed internal dose equivalents or intakes of radionuclides assessed pursuant to the rules in chapter 33-10-04 in effect before January 1, 1994. occupational Further. exposure histories obtained and recorded on the department's occupational radiation exposure history form (SFN 19443) or equivalent before January 1, 1994, 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.
- e. 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 shall assume:
 - (1)establishing administrative In controls pursuant to subdivision f of subsection 1 of section 33-10-04.1-06 for the current year, allowable dose that the limit for the individual is reduced by twelve and fivetenths millisieverts [1.25 rem] for each guarter for which records were unavailable and the individual was engaged in activities that could have resulted in occupational radiation exposure; and
 - (2) That the individual is not available for planned special exposures.
- f. The licensee or registrant shall retain the records on the department's occupational radiation exposure history form (SFN 19443) or equivalent until the department terminates each pertinent license or registration requiring this record. The licensee registrant shall retain or records used in preparing the department's occupational radiation exposure history form (SFN 19443) or equivalent for three 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 doses received under the limits specified in subsection 1 provided that each of the following conditions is are satisfied:
 - a. The licensee or registrant authorizes a planned special exposure only in an exceptional situation when alternatives that might avoid the higher dose

estimated to result from the planned special exposure are unavailable or impractical.

- b. The licensee or registrant, and employer if the employer is not the licensee or registrant, specifically authorizes the planned special exposure, in writing, before the exposure occurs.
- c. Before a planned special exposure, the licensee or registrant ensures that each individual involved is:
 - (1) Informed of the purpose of the planned operation; and
 - (2) 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
 - (3) Instructed in the measures to be taken to keep the dose as low as reasonably achievable considering other risks that may be present.
- d. Prior to permitting an individual to participate in a planned special exposure, the licensee or registrant ascertains prior doses as required by subdivision b of subsection 5 during the lifetime of the individual for each individual involved.
- e. Subject to subdivision b of subsection 1, 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:
 - The numerical values of any of the dose limits in subdivision a of subsection 1 in any year; and
 - (2) Five times the annual dose limits in subdivision a of subsection 1 during the individual's lifetime.
- f. The licensee or registrant maintains records of the conduct of a planned special exposure in accordance with subsection 6 of section 33-10-04.1-15 and submits a written report in accordance with subsection 4 of section 33-10-04.1-16.
- g. 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 thirty 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 subdivision a of subsection 1 but shall be included in evaluations required by subdivisions d and e.

- 7. Occupational dose limits for minors. The annual occupational dose limits for minors are ten percent of the annual occupational dose limits specified for adult workers in subsection 1.
- 8. Dose <u>equivalent</u> to an embryo or fetus.
 - a. The licensee or registrant shall ensure that the dose <u>equivalent</u> to an <u>the</u> embryo or fetus during the entire pregnancy, due to occupational exposure of a declared pregnant woman, does not exceed five millisieverts [0.5 rem]. See subsection 7 of section 33-10-04.1-15 for recordkeeping requirements.
 - b. The licensee or registrant shall make efforts to avoid substantial variation above a uniform monthly exposure rate to a declared pregnant woman so as to satisfy the limit in subdivision a (the national council on radiation protection and measurements recommended in NCRP report No. 91 "Recommendations on Limits for Exposure to Ionizing Radiation" (June 1, 1987) that no more than five-tenths millisievert [0.05 rem] to the embryo or fetus be received in any one month).
 - c. The dose <u>equivalent</u> to an <u>the</u> embryo or fetus shall be taken as the sum of:
 - (1) The deep dose equivalent to the declared pregnant woman; and
 - (2) The dose <u>equivalent</u> to the embryo or fetus from radionuclides in the embryo or fetus and radionuclides in the declared pregnant woman.
 - d. If by the time the woman declares pregnancy to the licensee or registrant, the dose <u>equivalent</u> to the embryo or fetus has exceeded four and five-tenths millisieverts [0.45 rem], the licensee or registrant shall be deemed to be in compliance with subdivision a of subsection 8 of section 33-10-04.1-06 if the additional dose <u>equivalent</u> to the

embryo or fetus does not exceed five-tenths millisievert [0.05 rem] during the remainder of the pregnancy.

History: Effective March 1, 1994; amended effective July 1, 1995; May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-07. Radiation dose limits for individual members of the public.

- 1. Dose limits for individual members of the public.
 - a. Each licensee or registrant shall conduct operations so that:
 - effective dose (1)The total equivalent to individual members of the public from the licensed or registered operation does not exceed one millisievert [0.1 rem] in a year, exclusive of the dose contribution from radiation, from background any medical administration the individual has received. from exposure to individuals administered radioactive material and released in accordance with subsection 12 of section 33-10-07-05 section 33-10-07.1-32 ("release of individuals containing unsealed radioactive material or implants containing radioactive material"), voluntary participation in medical research programs, and from the licensee's or registrant's disposal of radioactive material into sanitary sewerage in accordance with section 33-10-04.1-14. subsection 3 of Retrofit shall not be required for locations facilities where within only radiation machines existed prior to January 1, 1994, and the previous requirements of five met millisievert [0.5 rem] in a year; and
 - in any unrestricted area from (2)The dose sources exclusive of the external dose from contributions patients administered released radioactive material and in accordance with subsection 12 of section 33-10-07-05 does not exceed two-hundredths millisievert [0.002 rem] in any one-hour.
 - b. If the licensee or registrant permits members of the public to have access to restricted areas, the

limits for members of the public continue to apply to those individuals.

- C. Notwithstanding paragraph 1 of subdivision a of subsection 1 of this section, a licensee may permit visitors to an individual who cannot be released, under section 33-10-07.1-32 ("release of individuals containing unsealed radioactive material or implants containing radioactive material"), to receive a radiation dose greater than one millisievert [one hundred millirems] if:
 - (1) The radiation dose received does not exceed five millisieverts [five hundred millirems]; and
 - (2) The authorized user, as defined in chapter 33-10-07.1 ("medical use of radioactive material"), has determined before the visit that it is appropriate.
- ed. A licensee, registrant, or an applicant for a license or registration may apply for prior department authorization to operate up to an annual dose limit for an individual member of the public of five millisieverts [0.5 rem]. This application shall include the following information:
 - Demonstration of the need for and the expected duration of operations in excess of the limit in subdivision a;
 - (2) The licensee's or registrant's program to assess and control dose within the five millisieverts [0.5 rem] annual limit; and
 - (3) The procedures to be followed to maintain the dose as low as reasonably achievable.
- de. In addition to the requirements of this chapter a licensee or registrant subject to the provisions of the United States environmental protection agency's generally applicable environmental radiation standards in 40 CFR 190 shall comply with those standards.
- ef. The department may impose additional restrictions on radiation levels in unrestricted areas and on the total quantity of radionuclides that a licensee or registrant may release in effluents in order to restrict the collective dose.

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

- a. 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 subsection 1.
- b. A licensee or registrant shall show compliance with the annual dose limit in subsection 1 by:
 - (1) 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
 - (2) 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 II of appendix B; and
 - (b) If an individual were continuously present in an unrestricted area, the dose from external sources would not exceed two-hundredths millisievert [0.002 rem] in an hour and five-tenths millisievert [0.05 rem] in a year.
- c. Upon approval from the department, the licensee or registrant may adjust the effluent concentration values in appendix B, table II, 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.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-08. Testing for leakage or contamination of sealed sources.

1. Testing for leakage or contamination of sealed sources.

- a. The licensee or registrant in possession of any sealed source shall assure that:
 - (1) Each sealed source, except as specified in subdivision b of subsection 1, 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 six months before transfer to the licensee or registrant.
 - (2) Each sealed source that is not designed to emit alpha particles is tested for leakage or contamination at intervals not to exceed six months or at alternative intervals approved by the department, after evaluation of information specified by paragraphs 4 and 5 of subdivision k of subsection 5 of section 33-10-03-05, an agreement state, a licensing state, or the United States nuclear regulatory commission.
 - (3) Each sealed source that is designed to emit alpha particles is tested for leakage or contamination at intervals not to exceed three months or at alternative intervals approved by the department, after evaluation of information specified by paragraphs 4 and 5 of subdivision $k \pm 0$ of subsection 5 of section 33-10-03-05, an agreement state, a licensing state, or the United States nuclear regulatory commission.
 - (4) 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.
 - (5) Tests for leakage for all sealed sources, except brachytherapy sources manufactured to contain radium, shall be capable of detecting the presence of one hundred eighty-five becquerels [0.005 µCi] 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.

- (6) The test for leakage for brachytherapy sources manufactured to contain radium shall be capable of detecting an absolute leakage rate of thirty-seven becquerels $[0.001 \ \mu\text{Ci}]$ of radon-222 in a twenty-four-hour period when the collection efficiency for radon-222 and its daughters has been determined with respect to collection method, volume, and time.
- (7) 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 one hundred eighty-five becquerels [0.005 µCi] of a radium daughter which has a half-life greater than four days.
- b. A licensee or registrant need not perform test for leakage or contamination on the following sealed sources:
 - Sealed sources containing only radioactive material with a half-life of less than thirty days;
 - (2) Sealed sources containing only radioactive material as a gas;
 - (3) Sealed sources containing three and seventenths megabecquerels [100 μ Ci] or less of beta or photon-emitting material or three hundred seventy kilobecquerels [10 μ Ci] or less of alpha-emitting material;
 - (4) Sealed sources containing only hydrogen-3;
 - (5) Seeds of iridium-192 encased in nylon ribbon; and
 - (6) 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 leakage or contamination within six months before the date of use or transfer.

- c. Tests for leakage or contamination from sealed sources shall be performed by persons specifically authorized by the department, an agreement state, a licensing state, or the United States nuclear regulatory commission to perform such services.
- d. Test results shall be kept in units of becquerel or microcurie and maintained for inspection by the department. Records of test results for sealed sources shall be made pursuant to subsection 4 of section 33-10-04.1-15.
- e. The following shall be considered evidence that a sealed source is leaking:
 - (1) The presence of one hundred eighty-five becquerels $[0.005 \ \mu\text{Ci}]$ or more of removable contamination on any test sample.
 - (2) Leakage of thirty-seven becquerels [0.001 μCi] of radon-222 per twenty-hour hours for brachytherapy sources manufactured to contain radium.
 - (3) The presence of removable contamination resulting from the decay of one hundred eighty-five becquerels (0.005 µCi) or more of radium.
- f. 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 section.
- g. Reports of test results for leaking or contaminated sealed sources shall be made pursuant to subsection 8 of section 33-10-04.1-16.

History: Effective March 1, 1994; amended effective July 1, 1995. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-09. Survey and monitoring.

- 1. General
 - a. Each licensee or registrant shall make, or cause to be made, surveys that:

- (1) Are necessary for the licensee or registrant to comply with this chapter; and
- (2) Are <u>necessary</u> <u>reasonable</u> under the circumstances to evaluate:
 - (a) <u>The magnitude and extent of Rradiation</u> levels; <u>and</u>
 - (b) Concentrations or quantities of radioactive material; and
 - (c) The potential radiological hazards that could be present.
- b. 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 twelve months for the radiation measured except when a more frequent interval is specified in another applicable section of these rules or a license condition.
- c. 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 subsection 1 of section 33-10-04.1-06, with other provisions of this article, or with conditions specified in a license or registration shall be processed and evaluated by a dosimetry processor:
 - (1) Holding current personnel dosimetry accreditation from the national voluntary laboratory accreditation program (NVLAP) of the national institute of standards and technology; and
 - Approved in this accreditation process for the (2) type of radiation or radiations included in the national voluntary laboratorv closely accreditation program that most approximates the type of radiation or radiations for which the individual wearing the dosimeter is monitored.
- d. The licensee or registrant shall ensure that adequate precautions are taken to prevent a deceptive exposure of an individual monitoring device.

- 2. Conditions requiring individual monitoring of external and internal occupational dose. Each licensee or registrant shall monitor exposures from sources of radiation at levels sufficient to demonstrate compliance with the occupational dose limits of this chapter. At a minimum:
 - a. Each licensee or registrant shall monitor occupational exposure to radiation <u>from licensed</u> <u>and unlicensed radiation sources under the control</u> <u>of the licensee</u> and shall supply and require the use of individual monitoring devices by:
 - Adults likely to receive, in one year from sources external to the body, a dose in excess of ten percent of the limits in subdivision a of subsection 1 of section 33-10-04.1-06; and
 - (2) Minors and declared pregnant women likely to receive, in one year from sources external to the body, a dose in excess of ten percent of any of the applicable limits in subsections 7 or 8 of section 33-10-04.1-06; and Minors likely to receive, in one year from radiation sources external to the body, a deep dose equivalent in excess of one millisievert [one hundred millirem], a lens dose equivalent in excess of one and five-tenths millisieverts [one hundred fifty millirem], or a shallow dose equivalent to the skin of the whole body or to the skin of any extremity in excess of <u>five millisieverts [five hundred millirem]</u> (the assigned shallow dose equivalent must be the dose averaged over the contiguous ten square centimeters of skin receiving the highest exposure); and
 - (3) <u>Declared pregnant women likely to receive</u> <u>during the entire pregnancy, from radiation</u> <u>sources external to the body, a deep dose</u> <u>equivalent in excess of one millisievert [one</u> <u>hundred millirem]; and</u>
 - (<u>34</u>) Individuals entering a high or very high radiation area.
 - (4<u>5</u>) Reserved. Individuals working with medical fluoroscopic equipment.
 - (a) An individual monitoring device used for the dose to an embryo or fetus of a declared pregnant woman, pursuant to

<u>subdivision a of subsection 8 of section</u> <u>33-10-04.1-06, shall be located under the</u> <u>protective apron at the waist.</u>

- (b) An individual monitoring device used for <u>lens dose equivalent shall be located at</u> <u>the neck (collar), or an unshielded</u> <u>location closer to the eye, outside the</u> <u>protective apron.</u>
- (c) When only one individual monitoring device is used to determine the effective dose equivalent for external radiation pursuant to paragraph 2 of subdivision c of subsection 1 of section 33-10-04.1-06, it shall be located at the neck (collar) outside the protective apron. When a second individual monitoring device is used, for the same purpose, it shall be located under the protective apron at the waist. The second individual monitoring device is required for a declared pregnant woman.
- Each licensee or registrant shall monitor, to determine compliance with subsection 4 of section 33-10-04.1-06, the occupational intake of radioactive material by and assess the committed effective dose equivalent to:
 - Adults likely to receive, in one year, an intake in excess of ten percent of the applicable annual limit on intake in table I, columns 1 and 2, of appendix B; and
 - (2) Minors and declared pregnant women likely to receive, in one year, a committed effective dose equivalent in excess of five-tenths millisievert [0.05 rem]. Minors likely to receive, in one year, a committed effective dose equivalent in excess of one-tenth millisievert [ten millirem].
 - (3) Declared pregnant women likely to receive, during the entire pregnancy, a committed effective dose equivalent in excess of one millisievert [one hundred millirem].
- 3. Location of Individual Monitoring Devices. Each licensee or registrant shall ensure that individuals who are required to monitor occupational doses in accordance with subdivision a of subsection 2 of this section wear individual monitoring 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. When a protective apron is worn, the location of the individual monitoring device is typically at the neck (collar);
- b. An individual monitoring device used for monitoring the dose to an embryo or fetus of a declared pregnant woman, pursuant to subdivision a of subsection 8 of section 33-10-04.1-06, shall be located at the waist under any protective apron being worn by the woman;
- c. An individual monitoring device used for monitoring the lens dose equivalent, to demonstrate compliance with subparagraph a of paragraph 2 of subdivision a of subsection 1 of section 33-10-04.1-06, shall be located at the neck (collar), outside any protective apron being worn by the monitored individual, or at an unshielded location closer to the eye;
- d. An individual monitoring device used for monitoring the dose to the extremities, to demonstrate compliance with subparagraph b of paragraph 2 of subdivision a of subsection 1 of section 33-10-04.1-06, shall be worn on the extremity likely to receive the highest exposure. Each individual monitoring device shall be oriented to measure the highest dose to the extremity being monitored.

History: Effective March 1, 1994; amended effective July 1, 1995. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-10. Control of exposure from external sources in restricted areas.

- 1. Control of access to high radiation areas.
 - a. 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:
 - (1) 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 one millisievert [0.1 rem] in

one hour at thirty centimeters from the source of radiation or from any surface that the radiation penetrates;

- (2) 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
- (3) Entryways that are locked, except during periods when access to the areas is required, with positive control over each individual entry.
- b. In place of the controls required by subdivision a of subsection 1 for a high radiation area, the licensee or registrant may substitute continuous direct or electronic surveillance that is capable of preventing unauthorized entry.
- c. The licensee or registrant may apply to the department for approval of alternative methods for controlling access to high radiation areas.
- d. The licensee or registrant shall establish the controls required by subdivisions a and c of subsection 1 in a way that does not prevent individuals from leaving a high radiation area.
- e. 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 rules of the United States department of transportation provided that:
 - (1) The packages do not remain in the area longer than three days; and
 - (2) The dose rate at one meter from the external surface of any package does not exceed one tenth millisievert [0.01 rem] per hour.
- f. 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 chapter and to operate within the as low as is reasonably achievable provisions of the licensee's or registrant's radiation protection program.

q. 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 subsection 1 if the registrant has met all the specific requirements for access and control specified in other applicable parts of this article, such as, chapter 33-10-05 for industrial radiography, chapter 33-10-06 for x-ravs in the healing arts, and chapter 33-10-09 for particle accelerators.

2. Control of access to very high radiation areas.

- a. In addition to the requirements in subsection 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 five gray [500 rad] or more in one hour at one meter 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 nonself-shielded irradiators.
- b. 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 subdivision a if the registrant has met all the specific requirements for access and control specified in other applicable parts of this article, such as, chapter 33-10-05 for industrial radiography, chapter 33-10-06 for x-rays in the healing arts, and chapter 33-10-09 for particle accelerators.

3. Control of access to very high radiation areas -- irradiators.

a. This subsection applies to licensees or registrants with sources of radiation in nonself-shielded irradiators. This subsection does not apply to sources of radiation that are used in teletherapy, in industrial radiography, or in completely selfshielded 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.

- b. Each area in which there may exist radiation levels in excess of five gray [500 rad] in one hour at one meter from a source of radiation that is used to irradiate materials shall meet the following requirements:
 - (1) Each entrance or access point shall be equipped with entry control devices which:
 - (a) Function automatically to prevent any individual from inadvertently entering a very high radiation area;
 - (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 one millisievert [0.1 rem] in one 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 one millisievert [0.1 rem] in one hour.
 - (2) Additional control devices shall be provided so that, upon failure of the entry control devices to function as required by paragraph 1:
 - (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 one millisievert [0.1 rem] in one 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.

- (3) 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 one millisievert [0.1 rem] in one 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.
- (4) 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.
- (5) 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 paragraph 3 and 4.
- (6) Each area shall be equipped with devices that will automatically generate conspicuous visible and audible alarm signals to alert personnel in the area before 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.
- (7) 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.

- (8) 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 one millisievert [0.1 rem] in one hour.
- (9) The entry control devices required in paragraph 1 shall be tested for proper functioning. See subsection 10 of section 33-10-04.1-15 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;
 - (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.
- (10) 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.
- (11) 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.

- Licensees, registrants, or applicants for licenses c. or registrations for sources of radiation within the purview of subdivision b 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 subdivision b, 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 subdivision b. 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 hiah radiation levels before an individual can gain access to the area where such sources of radiation are used.
- d. The entry control devices required by subdivisions b and c shall be established in such a way that no individual will be prevented from leaving the area.

History: Effective March 1, 1994. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-11. 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 practical, 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.
 - <u>a.</u> When it is not practical 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, consistent with maintaining the total effective dose equivalent as low as is reasonably achievable (ALARA), shall

increase monitoring and limit intakes by one or more of the following means:

- a.(1) Control of access;
- b.(2) Limitation of exposure times;
- c.(3) Use of respiratory protection equipment; or
- d.(4) Other controls.
- b. If the licensee performs an as low as reasonably achievable (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.

- a. If the licensee or registrant <u>assigns or permits</u> <u>the</u> uses <u>of</u> respiratory protection equipment to limit intakes <u>of radioactive material</u> pursuant to subsection 2:
 - (1) Except as <u>otherwise</u> provided in paragraph 2 <u>this section</u>, 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 <u>(NIOSH)</u> and the <u>mine safety</u> and health administration.
 - The licensee or registrant may use respiratory (2)protection equipment that has not been tested or certified by the national institute for occupational safety and health and the mine safety and health administration, 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, provided the licensee or registrant has submitted to the department and the department has approved an application for authorized use of that respiratory protection equipment, including a demonstration by testing, or a demonstration on the basis of test information, that the material and performance characteristics of the respiratory protection equipment are capable of providing

the proposed degree of protection under anticipated conditions of use.

- (3) 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 respiratory protection equipment selection, and estimate exposures doses;
 - (b) Surveys and bioassays, as appropriate, to evaluate actual intakes;
 - (c) Testing of respiratory protection equipment for operability, including user seal check for face sealing devices and functional check for others, immediately prior to each use;
 - (d) Written procedures regarding selection, fitting, issuance, maintenance, and testing of respiratory protection equipment, including testing for operability immediately prior to each use; supervision and training of personnel; monitoring, including air sampling and bioassays; and recordkeeping; and
 - [1] Monitoring, including air sampling and bioassays;
 - [2] <u>Supervision and training of</u> respirator users;
 - [3] Fit testing;
 - [4] <u>Respirator selection;</u>
 - [5] Breathing air quality;
 - [6] Inventory and control;
 - [7] Storage, issuance, maintenance, repair, testing, and quality assurance of respiratory protection equipment;
 - [8] <u>Recordkeeping;</u>

- [9] Limitations on periods of respirator use and relief from respirator use;
- [10] The use of process or other engineering controls, instead of respiratory protection equipment;
- [11] The routine, nonroutine, and emergency use of respiratory protection equipment.
- (e) Determination by a physician prior to <u>the</u> initial fitting of respiratory protection equipment, and either every twelve months thereafter or periodically at a frequency determined by a physician, that the individual user is medically fit to use the respiratory protection equipment.
- (f) Fit testing, with fit factor greater than or equal to ten times the assigned protection factor for negative pressure devices, and a fit factor greater than or equal to five hundred 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 not to exceed one year. Fit testing must be performed with the facepiece operating in the negative pressure mode.
- (4) The licensee or registrant shall issue a written policy statement on respiratory protection equipment usage covering:
 - (a) The use of process or other engineering controls, instead of respiratory protection equipment;
 - (b) The routine, nonroutine, and emergency use of respiratory protection equipment; and
 - (c) The length of periods of respiratory protection equipment use and relief from respiratory protection equipment use.
- (54) The licensee or registrant shall advise each respiratory protection equipment user that the user may leave the area at any time for relief from respiratory protection equipment 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.

- (65) The licensee or registrant shall use respiratory protection equipment within the equipment manufacturer's expressed limitations for type and mode of use. The licensee or registrant shall also consider limitations appropriate to the type and mode of use. When selecting respiratory devices the licensee or registrant and shall provide proper visual for vision correction, adequate communication, low temperature work environments, and the concurrent use of other safety or radiological protection equipment. The licensee or registrant shall use equipment in such a way as not to interfere with the proper operation of the respirator and shall consider other special capabilities, such as adequate skin protection, when needed.
- (6) <u>Standby rescue persons are required whenever</u> 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 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.
- (7) Atmosphere-supplying respirators must be supplied with respirable air of grade D quality or better as defined by the Compressed Gas Association in publication G-7.1, "Commodity Specification for Air," 1997 and included in the regulations of the United

<u>States occupational safety and health</u> <u>administration (29 CFR 1910.134(i)(1)(ii)(A)</u> <u>through (E). Grade D quality air criteria</u> <u>including:</u>

- (a) Oxygen content (v/v) of 19.5-23.5%;
- (b) Hydrocarbon (condensed) content of 5 milligrams per cubic meter of air or less;
- (c) Carbon monoxide (CO) content of 10 ppm or less;
- (d) Carbon dioxide content of 1,000 ppm or less; and
- (e) Lack of noticable odor.
- (8) The licensee shall ensure that no objects, materials or substances, such as facial hair, or any conditions that interfere with the face--facepiece 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 tight-fitting respirator facepiece.
- b.(9) When estimating exposure of <u>dose to</u> individuals to <u>from intake of</u> airborne radioactive materials, the licensee or registrant may make allowance for respiratory protection equipment used to limit intakes pursuant to subsection 2, provided that the following conditions, in addition to those in subdivision a, are satisfied:
- (1) The licensee or registrant selects respiratory protection equipment that provides a protection factor, specified in appendix A, greater than the multiple by which peak concentrations of airborne radioactive materials in the working area are expected to exceed the values specified in appendix B, table I, column 3. However, if the selection of respiratory protection equipment with a protection factor greater than the multiple defined in the preceding sentence is inconsistent with the goal specified in subsection 2 of keeping the total effective dose equivalent as low as is reasonably achievable, the licensee or registrant may

select respiratory protection equipment with a lower protection factor only if such a selection would result in keeping the total effective dose equivalent as low as is reasonably achievable. The concentration of radioactive material in the air that is inhaled when respiratorys protection equipment are worn may be is initially estimated by dividing the average concentration in air, during each period of uninterrupted use, by the assumed to be the ambient concentration in air without respiratory protection, divided by the assigned protection factor. If the exposure dose is later found to be greater than initially estimated, the corrected value shall be used; however, if the exposure is later found to be less than initially estimated, the corrected value may be used.

- (2) The licensee or registrant shall obtain authorization from the department before assigning respiratory protection factors in excess of those specified in appendix A. 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.
- c. 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.
- d. The licensee or registrant shall notify the department in writing at least thirty days before the date that respiratory protection equipment is first used pursuant to either subdivision a.
 - 4. Further restrictions on the use of respiratory protection equipment. The department may impose restrictions in addition to those in subsection 2, subsection 3, and appendix A to:

- a. Ensure that the respiratory protection program of the licensee or registrant is adequate to limit exposures of doses to individuals to from intakes of airborne radioactive materials consistent with maintaining total effective dose equivalent as low as reasonably achievable; and
- b. Limit the extent to which a licensee <u>or registrant</u> may use respiratory protection equipment instead of process controls or other engineering controls.
- 5. Application for use of higher assigned protection factors. The licensee or registrant shall obtain authorization from the department before using assigned protection factors in excess of those specified in appendix A. The department may authorize a licensee or registrant to use higher assigned protection factors on receipt of an application that:
 - a. <u>Describes the situation for which a need exists for</u> <u>higher protection factors; and</u>
 - <u>b.</u> <u>Demonstrates that the respiratory protection</u> <u>equipment provides these higher protection factors</u> <u>under the proposed conditions of use.</u>

History: Effective March 1, 1994; amended effective July 1, 1995; May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-12. Storage <u>Security</u> 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.
- <u>2. Control of sources of radiation not in storage.</u>
- a. The licensee or registrant shall control and maintain constant surveillance of licensed or registered radioactive material that is in an unrestricted area and that is not in storage or in a patient who has been released in accordance with the patient release criteria in subsection 12 of section 33-10-07-05.

- b. The registrant shall maintain control of radiation machines that are in a controlled or unrestricted area and that are not in storage.
- 1. The licensee or registrant shall secure radioactive material from unauthorized removal or access.
- 2. The licensee or registrant shall maintain constant surveillance, and use devices or administrative procedures to prevent unauthorized use of radioactive material that is in an unrestricted area and that is not in storage.
- <u>3.</u> The registrant shall secure registered radiation machines from unauthorized removal.
- <u>4.</u> The registrant shall use devices or administrative procedures to prevent unauthorized use of registered radiation machines.

History: Effective March 1, 1994. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-13. Precautionary procedures.

- 1. Caution signs.
 - a. Standard radiation symbol. Unless otherwise authorized by the department, the symbol prescribed by this subsection 1 shall use the colors magenta, or purple, or black on yellow background. The symbol prescribed is the three-bladed design as shown in appendix H.
 - b. Exception to color requirements for standard radiation symbol. Notwithstanding the requirements of subdivision a, 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.
 - c. Additional information on signs and labels. In addition to the contents of signs and labels prescribed in this chapter, 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.

- a. 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 "CAUTION7: RADIATION AREA".
- b. 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 "CAUTION7: HIGH RADIATION AREA" or "DANGER7: HIGH RADIATION AREA".
- c. 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 DANGER7: VERY HIGH RADIATION AREA".
- d. 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 "CAUTION7: AIRBORNE RADIOACTIVITY AREA" or "DANGER7: AIRBORNE RADIOACTIVITY AREA".
- e. Posting of areas or rooms in which licensed or registered material is used or stored. The licensee or registrant shall post each area or room in which there is used or stored an amount of licensed or registered material exceeding ten times the quantity of such material specified in appendix C with a conspicuous sign or signs bearing the radiation symbol and the words "CAUTION7: RADIOACTIVE MATERIAL(S)" or "DANGER7: RADIOACTIVE MATERIAL(S)".

3. Exceptions to posting requirements.

- a. A licensee or registrant is not required to post caution signs in areas or rooms containing sources of radiation for periods of less than eight hours, if each of the following conditions is met:
 - (1) 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 chapter; and
 - (2) The area or room is subject to the licensee's or registrant's control.

- b. Rooms or other areas in hospitals that are occupied by patients are not required to be posted with caution signs pursuant to subsection 2 provided that the patient could be released from control pursuant to subsection 12 of section 33-10-07-05.
- c. 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 thirty centimeters from the surface of the sealed source container or housing does not exceed five hundredths millisievert [0.005 rem] per hour.
- d. 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.

- a. The licensee or registrant shall ensure that each container of licensed or registered material bears durable, clearly visible label bearing a the radiation symbol and "CAUTION7: the words RADIOACTIVE MATERIAL" or "DANGER7: 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 usinq the containers, or working in the vicinity of the containers, to take precautions to avoid or minimize exposures.
- b. Each licensee or registrant 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.
- c. 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.
- 5. **Exemptions to labeling requirements.** A licensee or registrant is not required to label:
 - Containers holding licensed or registered material in quantities less than the quantities listed in appendix C;

- Containers holding licensed or registered material in concentrations less than those specified in table III of appendix B;
- c. Containers attended by an individual who takes the precautions necessary to prevent the exposure of individuals in excess of the limits established by this chapter;
- Containers when they are in transport and packaged đ. and labeled in accordance with the rules of the States department of transportation United containing radioactive (Labeling of packages materials is required by the United States department of transportation if the amount and type of radioactive material exceeds the limits for an excepted quantity or article as defined and limited by United States department of transportation rules 49 CFR 173.403(m) and (w) and 173.421-424.);
- e. 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
- f. Installed manufacturing or process equipment, such as piping and tanks.

6. Procedures for receiving and opening packages.

- a. Each licensee or registrant who expects to receive a package containing quantities of radioactive material in excess of a type A quantity, as defined in section 33-10-13-02 and appendix A of chapter 33-10-13, shall make arrangements to receive:
 - (1) The package when the carrier offers it for delivery; or
 - (2) The notification of the arrival of the package at the carrier's terminal and to take possession of the package expeditiously.
- b. Each licensee or registrant shall:
 - (1) Monitor the external surfaces of a labeled package for radioactive contamination unless

the package contains only radioactive material in the form of gas or in special form as defined in section 33-10-01-04. Labeled package means posted with a radioactive white I, yellow II, or yellow III label as specified in United States department of transportation rules 49 CFR 172.403 and 172.436-440;

- Monitor the external surfaces of a labeled (2)package 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 section 33-10-13-02 and appendix A of chapter 33-10-13. Labeled package means posted with а radioactive white I, yellow II, or yellow III label as specified in United States department of transportation rules 49 CFR 172.403 and 172.436-440; and
- (3) 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.
- с. The licensee or registrant shall perform the monitoring required by subdivision b as soon as practical after receipt of the package, but not later than three 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 if there is evidence of degradation of package integrity, such as a package that is crushed, wet, or damaged. Ιf a package is received after working hours, and has no evidence of degradation of package integrity, the package shall be monitored no later than three hours from the beginning of the next working day.
- d. The licensee or registrant shall immediately notify the final delivery carrier and, by telephone and telegram, mailgram, or facsimile, the department by telephone in accordance with contact information in section 33-10-01-13 when:
 - Removable radioactive surface contamination exceeds the limits of subsection 8 of section 33-10-13-15; or
 - (2) External radiation levels exceed the limits of subsections 9 and 10 of section 33-10-13-15.

- e. Each licensee or registrant shall:
 - (1) Establish, maintain, and retain written procedures for safely opening packages in which radioactive material is received; and
 - (2) Ensure that the procedures are followed and that due consideration is given to special instructions for the type of package being opened.
- f. Licensees or registrants transferring transporting special form sources in vehicles owned or operated by the licensee or registrant to and from a worksite are exempt from the contamination monitoring requirements of subdivision b, but are not exempt from the monitoring requirement in subdivision b for measuring radiation levels that ensures that the source is still properly lodged in its shield.

History: Effective March 1, 1994; amended effective July 1, 1995; May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04

33-10-04.1-14. Waste disposal.

- 1. General requirements.
 - a. A licensee or registrant shall dispose of licensed or registered material only:
 - By transfer to an authorized recipient as provided in subsection 6 or in chapter 33-10-03, or to the United States department of energy;
 - (2) By decay in storage;
 - (3) By release in effluents within the limits in subsection 1 of section 33-10-04.1-07; or
 - (4) As authorized pursuant to subsection 2, 3, 4, or 5.
 - b. A person shall be specifically licensed or registered to receive waste containing licensed or registered material from other persons for:

- (1) Treatment prior to disposal;
- (2) Treatment or disposal by incineration;
- (3) Decay in storage;
- (4) Disposal at a land disposal facility licensed pursuant to 10 CFR 61; or
- (5) Storage until transferred to a storage or disposal facility authorized to receive the waste.
- 2. Method for obtaining approval of proposed disposal procedures. A licensee or registrant or applicant for a license or registration may apply to the department for approval of proposed procedures, not otherwise authorized in this article, to dispose of licensed or registered material generated in the licensee's or registrant's operations. Each application shall include:
 - a. A description of the waste containing licensed or registered 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;
 - b. An analysis and evaluation of pertinent information on the nature of the environment;
 - c. The nature and location of other potentially affected facilities; and
 - d. Analyses and procedures to ensure that doses are maintained as low as is reasonably achievable and within the dose limits in this chapter.

3. Disposal by release into sanitary sewerage.

- a. A licensee or registrant may discharge licensed or registered material into sanitary sewerage if each of the following conditions is satisfied:
 - The material is readily soluble, or is readily dispersible biological material, in water;
 - (2) The quantity of licensed or registered radioactive material that the licensee or registrant releases into the sewer in one month divided by the average monthly volume of water released into the sewer by the licensee or registrant does not exceed the

concentration listed in table III of appendix B;

- (3) If more than one radionuclide is released, the following conditions must also be satisfied:
 - registrant shall (a) The licensee or determine the fraction of the limit in table III of appendix B represented by discharges into sanitary sewerage by dividing the actual monthly average each radionuclide concentration of released by the licensee or registrant into the sewer by the concentration of that radionuclide listed in table III of appendix B: and
 - (b) The sum of the fractions for each radionuclide required by subparagraph a does not exceed unity; and
- (4) The total quantity of licensed or registered radioactive material that the licensee or registrant releases into the sanitary sewerage in a year does not exceed one hundred eightyfive gigabecquerels [5 Ci] of hydrogen-3, thirty-seven gigabecquerels [1 Ci] of carbon-14, and 37 gigabecquerels [1 Ci] of all other radioactive materials combined.
- b. Excreta from individuals undergoing medical diagnosis or therapy with radioactive material are not subject to the limitations contained in subdivision a.
- 4. **Treatment or disposal by incineration**. A licensee or registrant may treat or dispose of licensed or registered material by incineration only in the form and concentration specified in subsection 5 or as specifically approved by the department pursuant to subsection 2.

5. Disposal of specific wastes.

- a. A licensee or registrant may dispose of the following licensed or registered material as if it were not radioactive:
 - (1) One and eighty-five one-hundredths kilobecquerels $[0.05 \ \mu Ci]$, or less, of hydrogen-3 or carbon-14 per gram of medium used for liquid scintillation counting; and

- (2) One and eighty-five one-hundredths kilobecquerels $[0.05 \ \mu Ci]$, or less, of hydrogen-3 or carbon-14 per gram of animal tissue, averaged over the weight of the entire animal.
- b. A licensee or registrant shall not dispose of tissue pursuant to paragraph 2 of subdivision a in a manner that would permit its use either as food for humans or as animal feed.
- c. The licensee or registrant shall maintain records in accordance with subsection 9 of section 33-10-04.1-15.

6. Transfer for disposal and manifests.

- a. The requirements of this subsection and appendix D and appendix G are designed to:
 - (1) <u>cControl</u> transfers of low-level radioactive waste by any waste generator, waste collector, or waste processor licensee, as defined in appendix G, who ships low-level waste either directly, or indirectly through a waste collector or waste processor, to a licensed low-level radioactive waste <u>land</u> disposal facility,
 - (2) eEstablish a manifest tracking system, and
 - (3) <u>sSupplement</u> existing requirements concerning transfers and recordkeeping for those wastes.
- b. Beginning March 1, 1998, all affected licensees must use appendix G. Prior to March 1, 1998, a low-level radioactive waste disposal facility operator or its regulatory authority may require the shipper to use appendix D or appendix G. Licensees using appendix D shall comply with paragraph 1 of subdivision b of this subsection. Licensees using appendix G shall comply with paragraph 2 of subdivision b.
 - (1) Each shipment of radioactive waste intended 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 D.
 - (2) Any licensee shipping radioactive waste intended for ultimate disposal at a licensed land disposal facility must document the

information required on the uniform low-level radioactive waste manifest and transfer this recorded manifest information to the intended consignee in accordance with appendix G.

- c. Each shipment manifest shall include a certification by the waste generator as specified in section II of appendix D or appendix G, as appropriate.
- d. 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 D or appendix G, as appropriate.
- 7. Compliance with environmental and health protection rules. Nothing in subsection 1, 2, 3, 4, 5, or 6 relieves the licensee or registrant from complying with other applicable federal, state, and local rules governing any other toxic or hazardous properties of materials that may be disposed of in accordance with subsection 1, 2, 3, 4, 5, or 6.

History: Effective March 1, 1994; amended effective July 1, 1995; May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-04.1

33-10-04.1-15. Records.

- 1. General provisions.
 - a. Each licensee or registrant shall use the international system units becquerel, gray, sievert, and coulomb per kilogram, or the 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 chapter.
 - b. Notwithstanding the requirements of subdivision a, when recording information on shipment manifests, as required in paragraph 2 of subdivision b of subsection 6 of section 33-10-04.1-14, information must be recorded in the international system of units or in the international system of units and units as specified in subdivision a.

c. The licensee or registrant shall make a clear distinction among the quantities entered on the records required by this chapter, such as, total effective dose equivalent, total organ dose equivalent, shallow dose equivalent, eye lens dose equivalent, deep dose equivalent, or committed effective dose equivalent.

2. Records of radiation protection programs.

- a. Each licensee or registrant shall maintain records of the radiation protection program, including:
 - (1) The provisions of the program; and
 - (2) Audits and other reviews of program content and implementation.
- b. The licensee or registrant shall retain the records required by paragraph 1 of subdivision a until the department terminates each pertinent license or registration requiring the record. The licensee or registrant shall retain the records required by paragraph 2 of subdivision a for three years after the record is made.

3. Records of surveys.

- a. Each licensee or registrant shall maintain records showing the results of surveys and calibrations required by subsection 1 of section 33-10-04.1-09 and subdivision b of subsection 6 of section 33-10-04.1-13. The licensee or registrant shall retain these records for three years after the record is made.
- b. The licensee or registrant shall retain each of the following records until the department terminates each pertinent license or registration requiring the record:
 - Records of the results of surveys to determine the dose from external sources of radiation, and used, in the absence of or in combination with individual monitoring data, in the assessment of individual dose equivalents;
 - (2) Records of the results of measurements and calculations used to determine individual intakes of radioactive material and used in the assessment of internal dose;

- (3) Records showing the results of air sampling, surveys, and bioassays required pursuant to subparagraphs a and b of paragraph 3 of subdivision a of subsection 3 of section 33-10-04.1-11; and
- (4) Records of the results of measurements and calculations used to evaluate the release of radioactive effluents to the environment. This includes those records of the results of measurements and calculations used to evaluate the release of radioactive effluents to the environment required under the standards for protection against radiation in effect prior to March 1, 1994.
- c. Upon termination of the license or registration, the licensee or registrant shall permanently store records on the department's occupational radiation exposure history form (SFN 19443) or equivalent, or shall make provision with the department for transfer to the department.
- 4. Records of tests for leakage or contamination of sealed sources. Records of tests for leakage or contamination of sealed sources (required by subsection 1 of section 33-10-04.1-08) shall be kept in units of becquerel or microcurie and maintained for inspection by the department for five years after the records are made.

5. Records of prior occupational dose.

- a. The licensee or registrant shall retain the records of prior occupational dose and exposure history as specified in subsection 5 of section 33-10-04.1-06 on the department's occupational radiation exposure history form (SFN 19443) or equivalent until the department terminates each pertinent license or registration requiring this record. The licensee or registrant shall retain records used in preparing the department's occupational radiation exposure history form (SFN 19443) or equivalent for three years after the record is made.
- b. Upon termination of the license or registration, the licensee or registrant shall permanently store records on the department's occupational radiation exposure history form (SFN 19443) or equivalent, or shall make provision with the department for transfer to the department.
- 6. Records of planned special exposures.

- a. For each use of the provisions of subsection 6 of section 33-10-04.1-06 for planned special exposures, the licensee or registrant shall maintain records that describe:
 - The exceptional circumstances requiring the use of a planned special exposure;
 - (2) The name of the management official who authorized the planned special exposure and a copy of the signed authorization;
 - (3) What actions were necessary;
 - (4) Why the actions were necessary;
 - (5) What precautions were taken to assure that doses were maintained as low as is reasonably achievable;
 - (6) What individual and collective doses were expected to result; and
 - (7) The doses actually received in the planned special exposure.
- b. The licensee or registrant shall retain the records until the department terminates each pertinent license or registration requiring these records.
- c. Upon termination of the license or registration, the licensee or registrant shall permanently store records on the department's occupational radiation exposure history form (SFN 19443) or equivalent, or shall make provision with the department for transfer to the department.

7. Records of individual monitoring results.

- a. Recordkeeping requirement. Each licensee or registrant shall maintain records of doses received by all individuals for whom monitoring was required pursuant to subsection 2 of section 33-10-04.1-09, and records of doses received during planned special exposures, accidents, and emergency Assessments of dose equivalent and conditions. records made using units in effect before January 1, 1994, need not be changed. These records shall include, when applicable:
 - (1) The deep dose equivalent to the whole body, eye lens dose equivalent to the eye, shallow

dose equivalent to the skin, and shallow dose equivalent to the extremities;

- (2) The estimated intake of radionuclides, see subsection 2 of section 33-10-04.1-06;
- (3) The committed effective dose equivalent assigned to the intake of radionuclides;
- (4) The specific information used to calculate the committed effective dose equivalent pursuant to subdivisions a and c of subsection 4 of section 33-10-04.1-06 and when required by subsection 2 of section 33-10-04.1-09;
- (5) The total effective dose equivalent when required by subsection 2 of section 33-10-04.1-06; and
- (6) The total of the deep dose equivalent and the committed dose to the organ receiving the highest total dose.
- b. Recordkeeping frequency. The licensee or registrant shall make entries of the records specified in subdivision a at intervals not to exceed one year.
- c. Recordkeeping format. The licensee or registrant shall maintain the records specified in subdivision a on the department's current occupational radiation exposure form (SFN 8416), in accordance with the instructions for the department's current occupational radiation exposure form (SFN 8416), or in clear and legible records containing all the information required by the department's current occupational radiation exposure form (SFN 8416).
- d. The licensee or registrant shall maintain the records of dose <u>equivalent</u> to an <u>the</u> embryo or 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.
- e. The licensee or registrant shall retain each required form or record until the department terminates each pertinent license or registration requiring the record.
- f. Upon termination of the license or registration, the licensee or registrant shall permanently store

records on the department's occupational radiation exposure history form (SFN 19443) or equivalent, or shall make provision with the department for transfer to the department.

8. Records of dose to individual members of the public.

- a. Each licensee or registrant shall maintain records sufficient to demonstrate compliance with the dose limit for individual members of the public. See subsection 1 of section 33-10-04.1-07.
- b. The licensee or registrant shall retain the records required by subdivision a until the department terminates each pertinent license or registration requiring the record.

9. Records of waste disposal.

- a. Each licensee or registrant shall maintain records of the disposal of licensed or registered materials made pursuant to subsection 2, 3, 4, or 5 of section 33-10-04.1-14, chapter 33-10-03, or disposal by burial in soil, including burials authorized before October 1, 1982.
- b. The licensee or registrant shall retain the records required by subdivision a until the department terminates each pertinent license or registration requiring the record.

Requirements for disposition of these records, prior to license termination, are located in subsection 14 of section 33-10-03-05 and in sections 33-10-04.1-14 and 33-15-04.1-15 for activities licensed or registered under this article.

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

a. Each licensee or registrant shall maintain records of tests made pursuant to paragraph 9 of subdivision b of subsection 3 of section 33-10-04.1-10 on entry control devices for very high radiation areas. These records must include the date, time, and results of each such test of function.

- b. The licensee or registrant shall retain the records required by subdivision a for three years after the record is made.
- Form of records. 11. Each record required by this chapter 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 or 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. The licensee shall maintain adequate safeguards against tampering with and loss of records.
- 12. Prior to license termination, each licensee authorized to possess radioactive material with a half-life greater than one hundred twenty days, in an unsealed form, shall forward the following records to the department:
 - a. Records of disposal of licensed material made under subsection 2 of section 33-10-04.1-14 (including records of burials made before the effective date of this section), subsections 3, 4, and 5 of section 33-10-04.1-14; and
 - b. Records required by paragraph 4 of subdivision b of subsection 3 of section 33-10-04.1-15.
- 13. If licensed activities are transferred or assigned in accordance with subdivision b of subsection 7 of section 33-10-03-05, each licensee authorized to possess radioactive material, with a half-life greater than one hundred twenty days, in an unsealed form, shall transfer the following records to the new licensee and the new licensee will be responsible for maintaining these records until the licensee is terminated:
 - a. Records of disposal of licensed material made under subsection 2 of section 33-10-04.1-14 (including burials made before the effective date of this section), subsections 3, 4, and 5 of section 33-10-04.1-14, and
 - b. Records required by paragraph 4 of subdivision b of subsection 3 of section 33-10-04.1-15.

14. Prior to license termination, each licensee shall forward the records required by subdivision g of subsection 14 of section 33-10-03-05 to the department.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04, 23-20.1-09.1

33-10-04.1-16. Reports.

- 1. Reports of stolen, lost, or missing licensed or registered sources of radiation.
 - a. Telephone reports. Each licensee or registrant shall report to the department by telephone as follows:
 - (1) 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 one thousand times the quantity specified in appendix C under such circumstances that it appears to the licensee or registrant that an exposure could result to individuals in unrestricted areas; or
 - (2)Within thirty days after its occurrence becomes known to the licensee or registrant, lost, stolen, or missing licensed or radioactive registered material in an aggregate quantity greater than ten times the quantity specified in appendix C that is still missing.
 - (3) Immediately after its occurrence becomes known to the registrant, a stolen, lost, or missing radiation machine.
 - b. Written reports. Each licensee or registrant required to make a report pursuant to subdivision a, within thirty days after making the telephone report, shall make a written report to the department setting forth the following information:
 - (1) 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;

- (2) A description of the circumstances under which the loss or theft occurred;
- (3) A statement of disposition, or probable disposition, of the licensed or registered source of radiation involved;
- (4) Exposures of individuals to radiation, circumstances under which the exposures occurred, and the possible total effective dose equivalent to persons in unrestricted areas;
- (5) Actions that have been taken, or will be taken, to recover the source of radiation; and
- (6) 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.
- c. Subsequent to filing the written report, the licensee or registrant shall also report additional substantive information on the loss or theft within thirty days after the licensee or registrant learns of such information.
- d. The licensee or registrant shall prepare any report filed with the department pursuant to this subsection 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.

- a. 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:
 - (1) An individual to receive:
 - (a) A total effective dose equivalent of twenty-five one-hundredths sievert [25 rem] or more;

- (b) An eye <u>A lens</u> dose equivalent of seventyfive one-hundredths sievert [75 rem] or more; or
- (c) A shallow dose equivalent to the skin or extremities or a total organ dose equivalent of two and five-tenths gray [250 rad] or more; or
- (2) The release of radioactive material, inside or outside of a restricted area, so that, had an individual been present for twenty-four hours, the individual could have received an intake five times the annual limit on intake. This provision does not apply to locations where personnel are not normally stationed during routine operations, such as hot-cells or process enclosures.
- b. Twenty-four-hour notification. Each licensee or registrant, within twenty-four hours of discovery of the event, shall 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:
 - (1) An individual to receive, in a period of twenty-four hours:
 - (a) A total effective dose equivalent exceeding five-hundredths sievert [5 rem];
 - (b) An eye <u>A lens</u> dose equivalent exceeding fifteen hundredths sievert [15 rem]; or
 - (c) A shallow dose equivalent to the skin or extremities or a total organ dose equivalent exceeding five-tenths sievert [50 rem]; or
 - (2) The release of radioactive material, inside or outside of a restricted area, so that, had an individual been present for twenty-four hours, the individual could have received an intake in excess of one annual limit on intake. This provision does not apply to locations where personnel are not normally stationed during routine operations, such as hot-cells or process enclosures.

- c. The licensee or registrant shall prepare each report filed with the department pursuant to this subsection so that names of individuals who have received exposure to sources of radiation are stated in a separate and detachable portion of the report.
- d. Licensees or registrants shall make the reports required by subdivisions a and b to the department by telephone, telegram, mailgram, or facsimile to the department in accordance with contact information contained in section 33-10-01-13.
- e. The provisions of this subsection 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 subsection 4.

3. Reports of exposures, radiation levels, and concentrations of radioactive material exceeding the constraints or limits.

- a. Reportable events. In addition to the notification required by subsection 2, each licensee or registrant shall submit a written report within thirty days after learning of any of the following occurrences:
 - (1) Incidents for which notification is required by subsection 2; or
 - (2) Doses in excess of any of the following:
 - (a) The occupational dose limits for adults in subsection 1 or section 33-10-04.1-06;
 - (b) The occupational dose limits for a minor in subsection 7; of section 33-10-04.1-06;
 - (c) The limits for an embryo or fetus of a declared pregnant woman in subsection 8 of section 33-10-04.1-06;
 - (d) The limits for an individual member of the public in subsection 1 of section 33-10-04.1-07; or
 - (e) Any applicable limit in the license or registration; or

- (f) The as low as is reasonably achievable (ALARA) constraints for air emissions established under subsection 2 of section 33-10-04.1-05.
- (3) 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 ten times the applicable limit set forth in this chapter or in the license or registration, whether or not involving exposure of any individual in excess of the limits in subsection 1 of section 33-10-04.1-07; or
- (4) For licensees subject to the provisions of United States environmental protection agency's generally applicable environmental radiation standards in 40 CFR 190, levels of radiation or releases of radioactive material in excess of those standards, or of license conditions related to those standards.
- b. Contents of reports.
 - Each report required by subdivision a shall describe the extent of exposure of individuals to radiation and radioactive material, including, as appropriate:
 - (a) Estimates of each individual's dose;
 - (b) The levels of radiation and concentrations of radioactive material involved;
 - (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, as low as is reasonably achievable (ALARA) constraints, generally applicable environmental standards, and associated license or registration conditions.

- Each report filed pursuant to subdivision a (2)shall include for each occupationally overexposed individual: the name, social security account number, and date of birth. With respect to the limit for the embrvo or fetus in subsection 8 of section 33-10-04.1-06, 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.
- c. All licensees or registrants who make reports pursuant to subdivision a 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 thirty days following any planned special exposure conducted in accordance with subsection 6 of section 33-10-04.1-06, informing the department that a planned special exposure was conducted and indicating the date the planned special exposure occurred and the information required by subsection 6 of section 33-10-04.1-05.

5. Reporting requirements.

- a. Immediate report. Each licensee shall notify the department as soon as possible but not later than four hours after the discovery of an event that prevents immediate protective actions necessary to avoid exposures to radiation or radioactive materials that could exceed regulatory limits or releases of licensed material that could exceed regulatory limits (events may include fires, explosions, toxic gas releases, etc.).
- b. Twenty-four-hour report. Each licensee shall notify the department within twenty-four hours after the discovery of any of the following events involving licensed material:
 - (1) An unplanned contamination event that:
 - (a) Requires access to the contaminated area, by workers or the public, to be restricted for more than twenty-four hours by imposing additional radiological controls or by prohibiting entry into the area;

- (b) Involves a quantity of material greater than five times the lowest annual limit on intake specified in appendix B of this chapter 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 twenty-four hours to decay prior to decontamination.
- (2) An event in which equipment is disabled or fails to function as designed when:
 - (a) The equipment is required by rule or license condition to prevent releases exceeding regulatory limits, to prevent exposures to radiation and radioactive materials 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.
- (3) 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.
- (4) An unplanned fire or explosion damaging any licensed material or any device, container, or equipment containing licensed material when:
 - (a) The quantity of material involved is greater than five times the lowest annual limit on intake specified in appendix B of this chapter for the material; and
 - (b) The damage affects the integrity of the licensed material or its container.
- c. Preparation and submission of reports. Reports made by licensees in response to the requirements of this section must be made as follows:
 - (1) Licensees shall make reports required by subdivisions a and b by telephone to the

department. To the extent that the information is available at the 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.
- (2) Written report. Each licensee who makes a report required by subdivisions a and b shall submit a written followup report within thirty days of the initial report. Written reports prepared pursuant to other rules may be submitted to fulfill this requirement if the reports contain all of the necessary information and the appropriate distribution is made.
 - (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 exact location of the event;
 - (c) The isotopes, quantities, and chemical and physical form of the licensed material involved;
 - (d) Date and time of the event;
 - (e) Corrective actions taken or planned and the results of any evaluations or assessments; and
 - (f) The extent of exposure of individuals to radiation or to radioactive materials without identification of individuals by name.
- 6. Reports of individual monitoring.

- a. This section applies to each person licensed or registered by the department to:
 - Possess or use sources of radiation for purposes of industrial radiography pursuant to chapter 33-10-03 and 33-10-05; or
 - (2) Receive radioactive waste from other persons for disposal pursuant to chapter 33-10-03; or
 - (3) Possess or use at any time, for processing or manufacturing for distribution pursuant to chapter 33-10-03 or 33-10-07, radioactive material in quantities exceeding any one of the following quantities:

Radionuclide

Activity^a

	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, or order pursuant to section 33-10-01-09, 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.
- b. Each licensee or registrant in a category listed in subdivision a 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 subsection 2 of section 33-10-04.1-09 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 the department's current occupational radiation exposure form (SFN 8416) or equivalent or electronic media containing all the information required by the department's current occupational radiation exposure form (SFN 8416).

c. The licensee or registrant shall file the report required by subdivision b, covering the preceding year, on or before April thirtieth of each year. The licensee or registrant shall submit the report to the department.

7. Notifications and reports to individuals.

- a. Requirements for notification and reports to individuals of exposure to radiation or radioactive material are specified in subsection 3 of section 33-10-10-02.
- b. When a licensee or registrant is required pursuant to this section to report to the department any exposure of an individual to radiation or radioactive material, the licensee or registrant shall also provide the individual a copy of the report submitted to the department. Such reports shall be transmitted at a time not later than the transmittal to the department.
- 8. Reports of leaking or contaminated sealed sources. The licensee or registrant shall file a report within five days with the department if the test for leakage or contamination required pursuant to subsection 1 of section 33-10-04.1-08 indicates a sealed source is leaking or contaminated. The report shall include the equipment involved, the test results, and the corrective action taken.

History: Effective March 1, 1994; amended effective July 1, 1995; May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23,20.1-03, 23-20.1-04, 23-20.1-09.1

33-10-04.1-17. Additional requirements - Vacating premises. Each specific licensee or registrant shall, no less than thirty days before vacating or relinquishing possession or control of premises which may have been contaminated with radioactive material as a result of his activities, notify the department in writing of intent to vacate. When deemed necessary by the department, the licensee shall decontaminate the premises in accordance with the following or in such other manner as the department may specify.

1. Premises. Each licensee before vacating any premise, or transferring the premise shall permanently decontaminate such premises to meet the criteria for decommissioning in section 18. A survey shall be made after such decontamination and the department and the landlord or subsequent tenant or transferee shall be provided with a copy of such survey no less than thirty days before vacating or relinquishing possession or control of premises. No such premise may be vacated, sold, or transferred until the decontamination survey has been verified and accepted by the department.

2. Equipment. No machinery, instruments, laboratory equipment, or any other property used in contact with, or close proximity to radioactive material at a licensed premise may be assigned, sold, leased, or transferred to an unlicensed person unless such property has been permanently decontaminated below or equal to the standards specified in appendix F. A survey shall be made after such decontamination and the department and subsequent transferee or owner shall be provided with a copy of such survey. No such equipment may be assigned, sold, leased, or transferred until such documentation survey has been verified and accepted by the department.

History: Effective March 1, 1994; amended effective May 1, 1998. General Authority: NDCC 23-20.1-04 Law Implemented: NDCC 23-20.1-03, 23-20.1-04, 23-20.1-04.1

33-10-04.1-18. Radiological criteria for decommissioning <u>license termination</u>.

- 1. General provisions.
 - a. The criteria in this section apply to the decommissioning of licensed facilities.
 - b. The criteria in this section do not apply to sites which:
 - (1) Have been decommissioned prior to January 1, 1997, and met the August 20, 1997 in accordance with criteria identified in the United States nuclear regulatory commission's site decommissioning management plan action plan to ensure timely cleanup of site decommissioning management plan sites in 57 FR 13389; April 16,1992 of April 16, 1992 (57 FR 13389);
 - (2) Have previously submitted and received department approval on a <u>license termination</u> <u>plan or</u> decommissioning plan that is compatible with the criteria identified in the United States nuclear regulatory commission's <u>site decommissioning management plan</u> action

plan to ensure timely cleanup of site decommissioning management plan sites in 57 FR 13389; April 16, 1992; or of April 16, 1992 (57 FR 13389); or

- Submit a sufficient license termination plan (3) or decommissioning plan before January 1, 1999 August 20, 1998, and such license termination plan or decommissioning plan is approved by department before January 1, 2000 the August 20, 1999, and in accordance with the criteria identified in the United States nuclear regulatory commission's site decommissioning management plan action plan to ensure timely cleanup of site decommissioning management plan sites in 57 FR 13389; April 16, 1992. If an environmental impact statement is required in the submittal, and if, because of the environmental impact statement, the department cannot approve the plan before January 1, 2000, then the department may grant an extension of April 16, 1992 (57 FR 13389), except that if an environmental impact statement is required in the submittal, there will be a provision for dav-to-day extension.
- After a site has been decommissioned and the c. license terminated in accordance with the criteria in this section, the department will require on cleanup only if, based new additional information, it determines that the criteria of section met and residual this were not radioactivity remaining at the site could result in significant threat to public health and safety.
- d. When calculating total effective dose equivalent to the average member of the critical group, the licensee shall base estimates on the greatest determine the peak annual total effective dose equivalent dose expected within the first one thousand years after decommissioning. Estimates must be substantiated using actual measurements to the maximum extent practical.
- 2. Radiological criteria for unrestricted use. A site will be considered acceptable for unrestricted use if the residual radioactivity that is distinguishable from background radiation results in a total effective dose equivalent to an average member of the critical group that does not exceed twenty-five hundredths millisievert [25 millirem] per year, including that from groundwater sources of drinking water, and the residual radioactivity

has been reduced to levels that are as low as reasonably achievable. Determination of the levels which are as low as reasonably achievable shall take into account consideration of any detriments, such as loss <u>deaths</u> from transportation accidents, expected to potentially result from decontamination and waste disposal.

- 3. Criteria for license termination under restricted conditions. A site will be considered acceptable for license termination under restricted conditions if:
 - The licensee can demonstrate a. that further reductions in residual radioactivity necessary to comply with the provisions of subsection 2 would result in net public or environmental harm or were not being made because the residual levels associated with restricted conditions are as low as reasonably achievable. Determination of the levels which are as low as reasonably achievable shall take into account consideration of any detriments, such as loss from transportation traffic accidents, expected to potentially result from decontamination and waste disposal;
 - b. The licensee has made provisions for legally enforceable institutional controls that provide reasonable assurance that the total effective dose equivalent from residual radioactivity distinguishable from background to the average member of the critical group will not exceed twenty-five hundredths millisieverts [25 millirem] per year;
 - c. The licensee has provided sufficient financial assurance to enable an independent third party, including a governmental custodian of a site, to assume and carry out responsibilities for any necessary control and maintenance of the site. Acceptable financial assurance mechanisms are:
 - Funds placed into an account segregated from the licensee's assets and outside the licensee's administrative control as described in chapter 33-10-03;
 - (2) Surety method, insurance, or other guarantee method as described in chapter 33-10-03;
 - (3) A statement of intent in the case of federal, state, or local government licensees, as described in chapter 33-10-03; or

- (4) When a governmental entity is assuming custody and ownership of a site, an arrangement that is deemed acceptable by such governmental entity7.
- The licensee has submitted a decommissioning plan d. license termination plan to the department or indicating the licensee's intent to decommission in accordance with chapter 33-10-03, and specifying that the licensee intends to decommission by restricting use of the site. The licensee shall document in the license termination plan or decommissioning plan how the advice of individuals and institutions in the community who may be affected by the decommissioning has been sought and incorporated, as appropriate, following analysis of that advice. Licensees proposing to decommission by restricting use of the site shall seek advice from such affected parties regarding the following matters concerning the proposed decommissioning:
 - (1) <u>Licensees proposing to decommission by</u> restricting use of the site shall seek advice from such affected parties regarding the following matters concerning the proposed decommissioning:
 - (a) Whether provisions for institutional controls proposed by the licensee;
 - (a) [1] Will provide reasonable assurance that the total effective dose equivalent from residual radioactivity distinguishable from background to the average member of the critical group will not exceed twenty-five hundredths millisievert [25 millirem] total effective dose equivalent per year;
 - (b) [2] Will be enforceable; and
 - (c) [3] Will not impose undue burdens on the local community or other affected parties 7.
 - (2) (b) Whether the licensee has provided sufficient financial assurance to enable an independent third party, including a governmental custodian of a site, to assume and carry out responsibilities for any necessary control and maintenance of the site;

- (32) In seeking advice on the issues identified in this subdivision paragraph 1, the licensee shall provide for:
 - (a) Participation by representatives of a broad cross section of community interests who may be affected by the decommissioning;
 - (b) An opportunity for a comprehensive, collective discussion on the issues by the participants represented; and
 - (c) A publicly available summary of the results of all such discussions, including a description of the individual viewpoints of the participants on the issues and the extent of agreement and disagreement among the participants on the issues; and
- e. Residual radioactivity at the site has been reduced so that if the institutional controls were no longer in effect, there is reasonable assurance that the total effective dose equivalent from residual radioactivity distinguishable from background to the average member of the critical group is as low as reasonably achievable and would not exceed either:
 - (1) one millisievert [100 millirem] per year; or
 - (2) five millisieverts [500 millirem] per year
 provided the licensee:
 - (a) Demonstrates that further reductions in residual radioactivity necessary to comply with the one millisievert [100 millirem] per year value of paragraph 1 are not technically achievable, would be prohibitively expensive, or would result in net public or environmental harm;
 - (b) Makes provisions for durable institutional controls; and
 - (c) Provides sufficient financial assurance to enable a responsible government entity or independent third party, including a governmental custodian of a site, both to carry out periodic rechecks of the site no less frequently than every five years to assure that the institutional controls

remain in place as necessary to meet the criteria of subdivision b and to assume and carry out responsibilities for any necessary control and maintenance of those controls. Acceptable financial assurance mechanisms are those in subdivision c.

4. Alternate criteria for license termination.

- <u>a.</u> The department may terminate a license using alternate criteria greater than the dose criterion of subsection 2, subdivision b of subsection 3, or <u>item 1 of subparagraph a of</u> paragraph 1 of subdivision d of subsection 3, if the licensee:
- Provides assurance that public health and (1)a. safety would continue to be protected, and that it is unlikely that the total dose from all manmade sources combined, other than than medical, would be more the one millisievert [100 millirem] per year limit of section 33-10-04.1-07 would be unlikely, by submitting an analysis of possible sources of exposure;
- b. (2) Has employed to the extent practical restrictions on site use according to the provisions of subsection 3 in minimizing exposures at the site;
- (3) Reduced doses to as low as is reasonably c. achievable levels. <u>Determination of the</u> levels which are as low as reasonably achievable shall take taking into account consideration of any detriments, such as loss traffic from transportation accidents, potentially result from expected to decontamination and waste disposal;
- submitted a decommissioning plan or d. (4) Has license termination plan to the department the licensee's intent indicating to decommission in accordance with subsection 8 of section 33-10-03-05 and specifying that the licensee proposes to decommission by use of alternate criteria. The licensee shall document in the decommissioning plan or the license termination plan how the advice of individuals and institutions in the community who may be affected by the decommissioning has been sought and incorporated, as appropriate, following analysis of that advise advice. In

seeking such advise <u>advice</u>, the licensee shall provide for:

- (1) (a) Participation by representatives of a broad cross section of community interests who may be affected by the decommissioning;
- (2) (b) An opportunity for a comprehensive, collective discussion on the issues by the participants represented; and
- (3) (c) A publicly available summary of the results of all such discussions, including a description of the individual viewpoints of the participants on the issues and the extent of agreement and disagreement among the participants on the issues; and.
- eb. The use of alternate criteria to terminate a license requires the approval of the department after addressing any comments provided by the United States environmental protection agency, the United States nuclear regulatory commission, and any public comments submitted pursuant to subsection 5.
- 5. **Public notification and public participation.** Upon the receipt of a license termination plan or decommissioning plan from the licensee, or a proposal by the licensee for release of a site pursuant to subsection 3 or 4, or whenever the department deems such notice to be in the public interest, the department shall provide opportunity for public comment. Public comment procedures shall include the following:
 - a. Notice shall be given by publication in a newspaper of general circulation in the area where the license<u>e</u> is located or in a state publication designed to give public notice; to persons on a mailing list developed by the department, including those who request in writing to be on the list; and by other means if necessary to assure adequate notice of the affected public. This shall include publishing a notice in a forum, such as local newspapers, letters to state or local organizations, or other appropriate forum, that is readily accessible to individuals in the vicinity of the site, and soliciting comments from affected parties.

- <u>b.</u> Notice shall be made to, and comments solicited from, the United States environmental protection agency and United States nuclear regulatory commission for cases where the licensee proposes to release a site pursuant to subsection 4.
- c. Notice shall be made to, and comments solicited from, local and state governments in the vicinity of the site and any Indian Nation or other indigenous people that have treaty or statutory rights that could be affected by the decommissioning.
- bd. The notice shall identify the affected facility; the name and address of the licensee; the name and address of the department; a brief description of the plan; the name, address, and telephone number of a person from whom interested persons may obtain additional information, including copies of the plan, all relevant supporting materials, and all other materials available to the department that are relevant to the decision; a brief description of the comment procedures required by this subsection; and the time and place of any hearing that may be held, including a statement of procedures to request a hearing, unless a hearing has already been scheduled;
- ce. The department shall provide at least thirty days for public comment and shall give notice of any public hearing at least thirty days in advance of the hearing; and
- df. The department shall keep a record of the commenters and also of the issues raised during the public participation process. These records shall be available to the public.
- 6. Minimization of contamination. Applicants for licenses, other than renewals, shall describe in the application how facility design and procedures for operation will minimize, to the extent <u>practical practicable</u>, contamination of the facility and the environment, facilitate eventual decommissioning, and minimize, to the extent <u>practical practicable</u>, the generation of radioactive waste.

History: Effective May 1, 1998. General Authority: 23-20.1-04 Law Implemented: 23-20.1-03, 23-20.1-04, 23-20.1-04.1

APPENDIX A ASSIGNED PROTECTION FACTORS FOR RESPIRATORS¹

				igned on Factors
Descript	ion	Modes ³	Particulates only	Particulates, gases & vapors
(1) 2	AIR-PURIFYING RESPIRATORS ⁶			
I I I I I I I I I I I I I	Filtering facepiece ⁴ disposable Facepiece, half -mask⁷ Facepiece, full Facepiece, half Facepiece full Helmet or hood Facepiece, loose-fitting	NP NP PP PP PP PP PP	4 10 100 50 1000 1000 25	
(2) 4	ATMOSPHERE-SUPPLYING RESPIRATORS⁵			
1	1. Air-line respirator			
	Facepiece, half Facepiece, half Facepiece, half Facepiece, full Facepiece, full Helment or hood Facepiece, loose- fitting Suit	CF D CF D PD CF CF CF		50 10 50 1000 100 1000 25 2

- Self-contained breathing apparatus[®] 2. (SCBA) Faceoiece. full Facepiece, full Facepiece, full Facepiece, full
 - D 10.000^{8} PD RD $10,000^{8}$ RP
- COMBINATION RESPIRATORS <u>3.</u> Any combination of air-purifying and atmosphere-supplying respirators

Facepiece, full

<u>Assigned</u>	prot	<u>cection</u>	factor
for type	and	mode o	<u>f</u>
operation	n as	listed	above.

100⁹

100²

FOOTNOTES

1. For use in the selection of respiratory protection equipment to be used only where the contaminants have been identified and the concentrations, or possible These assigned protection factors apply only in a concentrations, are known. respiratory protection program that meets the requirements of this chapter. They are applicable only to airborne radiological hazards and may not be appropriate to circumstances when chemical or other respiratory hazards exist instead of, or in addition to, radioactive hazards. Selection and use of respirators for such circumstances must also comply with United States Department of Labor regulations.

Radioactive contaminants for which the concentration values in Table 1, Column 3 of Appendix B of Chapter 33-10-04.1 are based on internal dose due to inhalation may. in addition, present external exposure hazards at higher concentrations. Under these circumstances, limitations on occupancy may have to be governed by external dose limits.

2. Only for shaven faces and where nothing interferes with the seal of tight-fitting facepieces against the skin. Hoods and suits are excepted. No NIOSH approval schedule is currently available for atmosphere supplying suits. This equipment may be used in an acceptable respiratory protection program as long as all the other

minimum program requirements in section 33-10-04.1-11, with the exception of fit testing, are met.

- 3. The mode symbols are defined as follows:
 - CF = continuous flow
 - D = demand
 - NP = negative pressure, that is, negative phase during inhalation
 - PD = pressure demand, that is, always positive pressure
 - PP = positive pressure powered air-purifying
 - RD = demand, recirculating or closed circuit
 - RP = <u>positive</u> 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 protection 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: The concentration inhaled is the ambient airborne concentration divided by the protection factor.
 - b. The protection factors apply:
 - (1) Only for individuals trained in using respirators and wearing properly fitted respirators that are used and maintained under supervision in a well-planned respiratory protection program.
 - (2) For air-purifying respirators only when high efficiency particulate filters, above ninety-nine and ninety-seven hundredths percent removal efficiency by thermally generated three-tenths micron dioctyl phthalate (DOP) test or equivalent, are used in atmospheres not deficient in oxygen and not containing radioactive gas or vapor respiratory hazards.
- (3) No adjustment is to be made for the use of sorbents against radioactive material in the form of gases or vapors.

- (4) 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. Licensees may permit individuals to use this type of respirator who have not been medically screened or fit tested on the device provided that no credit be taken for their use in estimating intake or dose. It is also recognized that it is difficult to perform an effective positive or negative pressure pre-use user seal check on this type of device. All other respiratory protection program requirements listed in subsection 33-10-04.1-11.3 apply. An assigned protection factor has not been assigned for these devices. However, an assigned protection factor equal to 10 may be used if the licensee can demonstrate a fit factor of at least 100 by use of a validated or evaluated, qualitative or quantitative fit test.
- 5. Excluding The assigned protection factors for gases and vapors are not applicable to radioactive contaminants that present an absorption or submersion hazard. For tritium oxide vapor, approximately one-third of the intake occurs by absorption through the skin so that an overall protection factor of less than two three is appropriate when atmosphere-supplying respirators are used to protect against tritium Exposure to radioactive noble gases is not considered a significant oxide. respiratory hazard, and protective actions for these contaminants should be based on external (submersion) dose considerations. If the protection factor for respiratory protection equipment is five, the effective protection factor for tritium is about one and four tenths; with protection factors of ten; the effective factor for tritium oxide is about one and seven tenths; and with protection factors of one hundred or more, the effective factor for tritium oxide is about one and nine tenths. Airpurifying 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. <u>Air</u> purifying respirators with assigned protection factors <100 must be equipped with

particulate filters that are at least 95 percent efficient. Air purifying respirators with APF = 100 must be equipped with particulate filters that are at least 99 percent efficient. Air purifying respirators with assigned protection factors >100 must be equipped with particulate filters that are at least 99.97 percent efficient. The licensee may apply to the department for the use of an assigned protection factor greater than 1 for sorbent cartridges as protection against airborne radioactive gases and vapors (e.g., radioiodine).

- 7. Under-chin type only. No distinction is made in this appendix between elastomeric half-masks with replaceable cartridges and those designed with the filter medium as an integral part of the facepiece (e.g., disposable or reusable disposable). Both types are acceptable so long as the seal area of the latter contains some substantial type of seal-enhancing material such as rubber or plastic, the two or more suspension straps are adjustable, the filter medium is at least 95 percent efficient and all other requirements of this chapter are met. 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 I, column 3 of appendix B of chapter 33-10-04.1.1. 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 flow-rates are maintained. A protection factor of no more than one thousand may be utilized for tested-and-certified supplied-air hoods when a minimum air flow of six 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 two thousand 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 six 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 six cubic 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. There shall be a standby rescue person equipped with a respirator or other apparatus appropriate for the potential hazards and communications equipment whenever suppliedair suits are used.
- 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.
- 118. 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. This device may not be used by any individual who experiences perceptible outward leakage of breathing gas while wearing the device.
- 9. The licensee should implement institutional controls to assure that these devices are not used in areas immediately dangerous to life or health (IDLH).
- 12. Quantitative fit testing shall be performed on each individual, and no more than two hundredths percent leakage is allowed with this type of apparatus. Perceptible outward leakage of gas from this or any positive pressure selfcontained 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 I, column 3 of appendix B of chapter 33-10-04.1.1 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.

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

Introduction

For each radionuclide, table I indicates the chemical form which is to be used for selecting the appropriate annual limit on intake or derived air concentration value. The annual limit on intakes and derived air concentrations for inhalation are given for an aerosol with an activity median aerodynamic diameter (AMAD) of one µm, 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 of less than ten days for D, from ten to one hundred days for W, and of greater than one hundred days for Y. The class (D, W, Y) given in the column headed "class" applies only to the inhalation annual limit on intakes and derived air concentrations given in table I columns 2 and 3. Table II provides concentration limits for airborne and liquid effluents released to the general environment. Table III provides concentration limits for discharges to sanitary sewerage.

Note:

The values in tables I, II, and III are presented in the computer "E" notation. In this notation a value of 6E-02 represents a value of 6×10^{-2} or six hundredths, 6E+2 represents 6×10^{2} or six hundred, and 6E+0 represents 6×10^{0} or six.

Table I "Occupational Values"

Note that the columns in table I of this appendix captioned "oral ingestion annual limit on intake," "inhalation annual limit on intake," and "derived air concentration," are applicable to occupational exposure to radioactive material.

The annual limit on intakes 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 five hundredths sieverts (five rem), stochastic annual limit on intake, or (2) a committed dose equivalent of five tenths sieverts (fifty rem) to an organ or tissue, non-stochastic annual limit on intake. The stochastic annual limit on intakes 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 five hundredths sieverts (five rem). The derivation includes multiplying the committed dose equivalent to an organ or tissue by a weighting factor, w_m . 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 effects when the whole body is irradiated The values of w_{π} are listed under the definition of weighting factor in uniformly. section 33-10-04.1.1-03. The non-stochastic annual limit on intakes were derived to avoid non-stochastic effects, such as prompt damage to tissue or reduction in organ function.

A value of $w_T = 0.06$ 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 gastro-intestinal 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 (hands and forearms, feet, and lower legs), 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 annual limit on intake is defined by the stochastic dose limit, this value alone is given. When an annual limit on intake is determined by the non-stochastic dose limit to an organ, the organ or tissue to which the limit applies is shown, and the annual limit on intake for the stochastic limit is shown in parentheses. Abbreviated organ or tissue designations are used:

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LLI wall = lower large intestine wall;
St wall = stomach wall;
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Blad wall = bladder wall; and Bone surf = bone surface.

The use of the annual limit on intakes listed first, the more limiting of the stochastic and non-stochastic annual limit on intakes, 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 nonstochastic annual limit on intake is limiting, use of that non-stochastic annual limit on intake is considered unduly conservative, the licensee may use the stochastic annual limit on intake to determine the committed effective dose equivalent. However, the licensee shall also ensure that the five tenths sievert (fifty 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 For the case where there is no external dose contribution, this would be dose. demonstrated if the sum of the fractions of the nonstochastic annual limit on intakes (ALIne) that contribute to the committed dose equivalent to the organ receiving the highest dose does not exceed unity, that is, Σ (intake (in μ Ci) of each radionuclide/ALI_{ns}) < one. If there is an external deep dose equivalent contribution of H_a, then this sum must be less than 1 - (H_a/50), instead of \leq one.

The derived air concentration (DAC) values are derived limits intended to control chronic occupational exposures. The relationship between the derived air concentration and the annual limit on intake is given by:

DAC = ALI(in μ Ci)/(2000 hours per working year x 60 minutes/hour x 2 x 10⁴ ml per minute) = [ALI/2.4 x 10⁹] μ Ci/ml,

where 2×10^4 ml is the volume of air breathed per minute at work by reference man under working conditions of light work.

The derived air concentration values relate to one of two modes of exposure: either external submersion or the internal committed dose equivalents resulting from inhalation

of radioactive materials. Derived air concentrations based upon submersion are for immersion in a semi-infinite cloud of uniform concentration and apply to each radionuclide separately.

The annual limit on intake and derived air concentration 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 annual limit on intake and derived air concentration 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 subsection 2 of section 33-10-04.1.1-06. 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 II "Effluent Concentrations"

The columns in table II of this appendix captioned "effluent concentrations," "air" and "water" are applicable to the assessment and control of dose to the public, particularly in the implementation of the provisions of subsection 2 of section 33-10-04.1.1-07. The concentration values given in columns 1 and 2 of table II 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 five tenths millisievert (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 derived air concentration, the stochastic annual limit on intake was used in deriving the corresponding airborne effluent limit in table II. For this reason, the derived air concentration and airborne effluent limits are not always proportional as was the case in appendix A of the 1992 revision of chapter 33-10-04.1.1.

The air concentration values listed in table II, column 1 were derived by one of two methods. For those radionuclides for which the stochastic limit is governing, the occupational stochastic inhalation annual limit on intake was divided by 2.4×10^9 (ml), relating the inhalation annual limit on intake to the derived air concentration, as explained above, and then divided by a factor of three hundred. The factor of three hundred includes the following components: a factor of fifty to relate the five hundredths sievert (5 rem) annual occupational dose limit to the one millisievert (0.1 rem) limit for members of the public, a factor of three 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 two 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 derived air concentration in table I, column 3 was divided by two hundred nineteen. The factor of two hundred nineteen is composed of a factor of fifty, as described above, and a factor of four and thirty-eight hundredths relating occupational exposure for two thousand hours per year to full-time exposure (8,760 hours per year). Note that an additional factor of two for age considerations is not warranted in the submersion case.

The water concentrations were derived by taking the most restrictive occupational stochastic oral ingestion annual limit on intake and dividing by 7.3×10^7 . The factor of 7.3×10^7 (ml) includes the following components: the factors of fifty and two described above and a factor of 7.3×10^5 (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 annual limit on intakes and derived air concentrations, 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 either from knowledge of the radionuclide composition of the source or from actual measurements.

Table III "Releases to Sewers"

The monthly average concentrations for release to sanitary sewerage are applicable to the provisions in subsection 3 of section 33-10-04.1.1-14. The concentration values were derived by taking the most restrictive occupational stochastic oral ingestion annual limit on intake and dividing by 7.3×10^6 (ml). The factor of 7.3×10^6 (ml) is composed of a factor of 7.3×10^5 (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 five millisieverts (0.5 rem).

LIST OF ELEMENTS

Name	Symbol	Atomic Number	Name	Symbol	Atomic Number
Actinium Aluminum Antimony Argon Arsenic Astatine Barium Berkelium Berkelium Bismuth Bromine Cadium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Calcium Cobalt Copper Curium Dysprosium Einsteinium Einsteinium Fruncium Francium Gadolinium Gadolinium Gallium Gallium Holmium Holmium Holmium Indium Indium Indine Iridium Indium Lanthanum Lead Lutetium Magnesium Manganese Mendelevium	AAlmbrstakeirdat esiroumysrum rdaeuto nirkabugaM MMMM	$\begin{array}{c} 89\\ 13\\ 95\\ 51\\ 18\\ 33\\ 85\\ 97\\ 43\\ 55\\ 97\\ 43\\ 55\\ 12\\ 29\\ 66\\ 99\\ 63\\ 10\\ 97\\ 64\\ 12\\ 57\\ 26\\ 57\\ 27\\ 6\\ 71\\ 25\\ 125\\ 101 \end{array}$	Mercury Molybdenum Neodymium Neotymium Nickel Nicbium Palladium Phosphorus Platinum Plutonium Potassium Praseodymium Promethium Promethium Radon Rhenium Radon Rhenium Rubidium Ruthenium Scandium Scandium Scandium Scandium Scandium Scandium Stilicon Silicon Silicon Silicon Silicon Silicon Silicon Tantalum Technetium Tellurium Tellurium Thallium Thorium Thulium Tin Titanium Yungsten Uranium Yttrium Zinc Zirconium	god pibsd tuo rmaanenbumceigar aceblhfini NNOPPPPPRRRRRRRSSSANSsTTTTTTTSTWUVXYYZz	8469381665844991186555742144718633251090242340900 88924765844991186555742144718633251090242340900 88733251090242340900

			Occı	Table I pational Val	lues	Tabl Efflu Concent	ent	Table III Releases to Sewers
			<u>Col. 1</u> Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhala ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
	Hydrogen-3	Water, DAC includes skin absorption	8E+4	8E+4	2E-5	1E-7	1E-3	1E-2
		Gas (HT or T_2) Submersion ¹ : Use	e above valu	es as HT an	nd T ₂ oxidize	e in air and	in the body t	O HTO.
1	Beryllium-7	W, all compounds except those given for Y Y, oxides, halides, and	4E+4	2E+4	9E-6	3E-8	6E-4	6E-3
		nitrates	-	2E+4	8E-6	3E-8	-	-
ł	Beryllium-10	W, see ⁷ Be	1E+3 LLI wall	2E+2	6E-8	2E-10	-	-
		Y, see ⁷ Be	(1E+3)	_ 1E+1	- 6E~9	_ 2E-11	2E-5	2E-4
5	Carbon-11 ²	Monoxide Dioxide Compounds	- - 4E+5	1E+6 6E+5 4E+5	5E-4 3E-4 2E-4	2E-6 9E-7 6E-7	- 6E-3	- - 6E-2
i	Carbon-14	Monoxide Dioxide Compounds	- 2E+3	2E+6 2E+5 2E+3	7E-4 9E-5 1E-6	2E-6 3E-7 3E-9	- 3E-5	- - 3E-4
)	Fluorine-18 ²	D, fluorides of H, Li, Na, K, Rb, Cs, and Fr	5E+4 St wall	7E+4	3E-5	1E-7	-	-
		W, fluorides of Be, Mg, Ca, Sr, Ba, Ra, Al, 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	(5E+4) - -	- 9E+4 8E+4	- 4E-5 3E-5	- 1E-7 1E-7	7E-4 -	7E-3 - -
.1	Sodium-22	D, all compounds	4E+2	6E+2	3E-7	9E-10	6E-6	6E-5
1	Sodium-24	D, all compounds	4E+3	5E+3	2E-6	7E-9	5E-5	5E-4

			Occu	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
tomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
2	Magnesium-28	D all compounds except							
2	Magnesium-26	D, all compounds except those given for W W, oxides, hydroxides, carbides, halides, and	7E+2	2E+3	7E-7	2E-9	9E-6	9E-5	
		nitrates	-	1E+3	5E-7	2E-9	-	-	
3	Aluminum-26	D, all compounds except those given for W	4E+2	6E+1	3E-8	9E-11	6E-6	6E-5	
		W, oxides, hydroxides, carbides, halides, and nitrates	_	9E+1	4E-8	1E-10	-	-	
4	Silicon-31	D, all compounds except those given for W and Y	9E+3	3E+4	1E-5	4E-8	1E-4	1E-3	
		D, all compounds except those given for W and Y W, oxides, hydroxides, carbides, and nitrates Y, aluminosilicate glass	-	3E+4 3E+4	1E-5 1E-5	5E-8 4E-8	-	-	
4	Silicon-32	D, see ³¹ Si	2E+3 LLI wall	2E+2	1E-7	3E-10	-	-	
		W, see ³¹ Si Y, see ³¹ Si	(3E+3) - -	- 1E+2 5E+0	- 5E-8 2E-9	- 2E-10 7E-12	4E-5 - -	4E-4 - -	
5	Phosphorus-32	D, all compounds except phosphates given for W W, phosphates of Zn [*] , S [*] , Mg [*] , Fe [*] , Bi [*] , and lanthanides	6E+2	9E+2	4E-7	1E-9	9E-6	9E-5	
		S , Mg ^{*-} , Fe , Bi , and lanthanides	-	4E+2	2E-7	5E-10	-	-	
5	Phosphorus-33	D, see ${}^{32}_{32}P$ W, see ${}^{32}P$	6E+3	8E+3 3E+3	4E-6 1E-6	1E-8 4E-9	8E-5 -	8E-4 -	

			Occi	Table I upational Va	lues	Tabl Efflu Concent	ent	Table I Release Sewers	s to
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly	,
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concent (µCi/ml	ration)
16	Sulfur-35	Vapor D, sulfides and sulfates	1E+4	6E-6	2E-8	-	-		
		except those given for W	1E+4 LLI wall	2E+4	7E-6	2E-8	-	-	
			(8E+3)	-	-	-	1E-4	1E-3	W eleme ntal sulfu r 6E+3
		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	_	2E+3	9 E -7	3E-9	_ ·	_	01.5
17	Chlorine-36	D, chlorides of H, Li, Na, K, Rb, Cs, and Fr W, chlorides of lantha- nides, 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	2E+3	2E+3	1E-6	3E-9	2E-5	2E-4	
			-	2E+2	1E-7	3E-10	-	-	
17	Chlorine-38 ²	D, see ³⁶ Cl	2E+4 St wall	4E+4	2E-5	6E-8	-	-	
		W, see ³⁶ Cl	(3E+4)	- 5E+4	_ 2E-5		3E-4	3E-3	
17	Chlorine-39 ²	D, see ³⁶ Cl	2E+4 St wall	5E+4	2E-5	7E-8	-	-	
		W, see ³⁶ Cl	(4E+4)	- 6E+4	_ 2E-5	- 8E-8	5E-4 -	5E-3	
18	Argon-37	Submersion ¹	-	-	1E+0	6E-3	-	-	
18	Argon-39	Submersion ¹	-	-	2E-4	8E-7	-	-	
18	Argon-41	Submersion ¹	-	-	3E-6	1E-8	-	-	
19	Potassium-40	D, all compounds	3E+2	4E+2	2E-7	6E-10	4E-6	4E-5	
19	Potassium-42	D, all compounds	5E+3	5E+3	2E-6	7E-9	6E-5	6E-4	
19	Potassium-43	D, all compounds	6E+3	9E+3	4E-6	1E-8	9E-5	9E-4	

			Occu	Table I pational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral Ingestion	Col. 2 Inhalat	Col. 3	Col. 1	Col. 2	Monthly Average Concentration (µCi/ml)
tomic Radionuclic	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	
9	Potassium-44 ²	D, all compounds	2E+4	7E+4	3E-5	9E-8	-	-
			St wall (4E+4)	-	-	-	5E-4	5E-3
.9	Potassium-45 ²	D, all compounds	3E+4 St wall	1E+5	5E-5	2E~7	-	-
			(5E+4)	-	-	-	7E-4	7E-3
0	Calcium-41	W, all compounds	3E+3 Bone surf	4E+3	2E-6	-	-	-
			(4E+3)	Bone surf (4E+3)	-	5E-9	6E-5	6E-4
0	Calcium-45	W, all compounds	2E+3	8E+2	4E-7	1E-9	2E-5	2E-4
0	Calcium-47	W, all compounds	8E+2	9E+2	4E-7	1E-9	1E-5	1E-4
1	Scandium-43	Y, all compounds	7E+3	2E+4	9E-6	3E-8	1E-4	1E-3
1	Scandium-44m	Y, all compounds	5E+2	7E+2	3E-7	1E-9	7E-6	7E-5
1	Scandium-44	Y, all compounds	4E+3	1E+4	5E-6	2E-8	5E-5	5E-4
1	Scandium-46	Y, all compounds	9E+2	2E+2	1E-7	3E-10	1E-5	1E-4
1	Scandium-47	Y, all compounds	2E+3	3E+3	1E-6	4E-9	-	-
			LLI wall (3E+3)	-	-	-	4E-5	4E-4
1	Scandium-48	Y, all compounds	8E+2	1E+3	6E-7	2E-9	1E-5	1E-4
1	Scandium-49 ²	Y, all compounds	2E+4	5E+4	2E-5	8E-8	3E-4	3E-3

				Table I Ipational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalat ALI (µCi)	ion DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
22	Titanium-44	D, all compounds except those given for W and Y W, oxides, hydroxides, carbides, halides, and	3 E +2	1E+1	5E-9	2E-11	4E-6	4E-5
		carbides, halides, and nitrates Y, SrTiO ₃	Ξ	3E+1 6E+0	1E-8 2E-9	4E-11 8E-12	-	-
22	Titanium-45	D, see ⁴⁴ Ti W, see ⁴⁴ Ti Y, see ⁴⁴ Ti	9E+3 _	3E+4 4E+4 3E+4	1E-5 1E-5 1E-5	3E-8 5E-8 4E-8	1E-4 -	1E-3
3	Vanadium-47 ²	Y, see 441 D, all compounds except those given for W	3E+4 St wall	8E+4	3E-5	1E-7	_	_
		W oxides hydroxides	(3E+4)	-	-	-	4E-4	4E-3
		W, oxides, hydroxides, carbides, and halides	-	1E+5	4E-5	1E-7	-	-
3	Vanadium-48	D, see ⁴⁷ V W, see ⁴⁷ V	6E+2 -	1E+3 6E+2	5E-7 3E-7	2E-9 9E-10	9E-6 -	9E-5 -
3	Vanadium-49	D, see ⁴⁷ V	7E+4 LLI wall	3E+4 Bone surf	1E-5	-	-	-
		W, see ⁴⁷ V	(9E+4) -	(3E+4) 2E+4	- 8E-6	5E-8 2E-8	1E-3 -	1E-2 _
:4	Chromium-48	D, all compounds except those given for W and Y W, halides and nitrates Y, oxides and hydroxides	6E+3 - -	1E+4 7E+3 7E+3	5E-6 3E-6 3E-6	2E-8 1E-8 1E-8	8E-5 -	8E-4 -
4	Chromium-49 ²	D, see ⁴⁸ Cr W, see ⁴⁸ Cr Y, see ⁴⁸ Cr	3E+4 _ _	8E+4 1E+5 9E+4	4E-5 4E-5 4E-5	1E-7 1E-7 1E-7	4E-4 	4E-3 -
24	Chromium-51	D, see ⁴⁸ Cr W, see ⁴⁸ Cr Y, see ⁴⁸ Cr	4E+4 - -	5E+4 2E+4 2E+4	2E-5 1E-5 8E-6	6E-8 3E-8 3E-8	5E-4 -	5E-3 -

			0000	Table I pational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral Ingestion	Col. 2 Inhalat	Col. 3	Col. 1	Col. 2	Monthly Average
Atomic 1 No.	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentration (µCi/ml)
5	Manganese-51 ²	D, all compounds except	07.4	5E+4	2E-5	7E-8	3E4	3E-3
		D, all compounds except those given for W W, oxides, hydroxides, halides, and nitrates	2E+4 -	5E+4 6E+4	2E-5 3E-5	7E-8	5E-4 -	5E-5
5	Manganese-52m ²	D, see 51 Mn	3E+4	9E+4	4E-5	1E-7	-	-
		W, see ⁵¹ Mn	St wall (4E+4) -	- 1E+5	_ 4E-5	- 1E-7	5E-4	5E-3
5	Manganese-52	D, see ⁵¹ Mn W, see ⁵¹ Mn	7E+2	1E+3 9E+2	5E-7 4E-7	2E-9 1E-9	1E-5	1E-4
5	Manganese-53	D, see ⁵¹ Mn	5E+4	1E+4	5E-6	-	7E-4	7E-3
		W, see ⁵¹ Mn	-	Bone surf (2E+4) 1E+4	_ 5E-6	3E-8 2E-8	-	-
5	Manganese-54	D, see ⁵¹ Mn W, see ⁵¹ Mn	2E+3	9E+2 8E+2	4E-7 3E-7	1E-9 1E-9	3E-5 -	<u>3</u> E-4
5	Manganese-56	D, see ⁵¹ Mn W, see ⁵¹ Mn	5E+3 -	2E+4 2E+4	6E-6 9E-6	2E-8 3E-8	7E-5 -	7E-4 -
5	Iron-52	D, all compounds except those given for W	9E+2	3E+3	1E-6	4E-9	1E-5	1E-4
		W, oxides, hydroxides, and halides	-	2E+3	1E-6	3E-9	-	-
6	Iron-55	D, see ⁵² Fe W, see ⁵² Fe	9E+3	2E+3 4E+3	8E-7 2E-6	3E-9 6E-9	1E-4	1E-3
6	Iron-59	D, see ⁵² Fe W, see ⁵² Fe	8E+2 -	3E+2 5E+2	1E-7 2E-7	5E-10 7E-10	1E-5	1E-4
5	Iron-60	D, see ⁵² Fe W, see ⁵² Fe	3E+1 -	6E+0 2E+1	3E-9 8E-9	9E-12 3E-11	4E-7	4E-6

	** * · · · · · · · · · · · · · · · · ·			Table I upational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentratior (µCi/ml)
?7	Cobalt-55	W, all compounds except those given for Y	1E+3	3E+3	1E-6	4E-9	2E-5	2E-4
		Y, oxides, hydroxides, halides, and nitrates	-	3E+3	1E-6	4E-9	-	_
:7	Cobalt-56	W, see ⁵5Co Y, see ⁵5Co	5E+2 4E+2	3E+2 2E+2	1E-7 8E-8	4E-10 3E-10	6E-6 -	6E-5
:7	Cobalt-57	W, see ⁵5Co Y, see ⁵5Co	8E+3 4E+3	3E+3 7E+2	1E-6 3E-7	4E-9 9E-10	6E-5	6E-4
7	Cobalt-58m	W, see ⁵5Co Y, see ⁵5Co	<u>6</u> E+4	9E+4 6E+4	4E-5 3E-5	1E-7 9E-8	8E-4	8E-3
7	Cobalt-58	W, see ⁵⁵ Co Y, see ⁵⁵ Co	2E+3 1E+3	1E+3 7E+2	5E-7 3E-7	2E-9 1E-9	2E-5	2E-4
7	$Cobalt-60m^2$	W, see ⁵⁵ Co	1E+6 St wall	4E+6	2E-3	6E-6	-	-
		Y, see ⁵⁵ Co	(1E+6) _	- 3E+6	- 1E-3	- 4E-6	2E-2 -	2E-1 -
7	Cobalt-60	W, see ⁵⁵ Co Y, see ⁵⁵ Co	5E+2 2E+2	2E+2 3E+1	7E-8 1E-8	2E-10 5E-11	3E-6	3E-5
7	Cobalt-61 ²	W, see ⁵⁵ Co Y, see ⁵⁵ Co	2E+4 2E+4	6E+4 6E+4	3E-5 2E-5	9E-8 8E-8	3E-4	3E-3
7	$Cobalt-62m^2$	W, see ⁵⁵ Co	4E+4 St wall	2E+5	7E-5	2E-7	-	-
		Y, see ⁵⁵ Co	(5E+4) -	- 2E+5	- 6E-5	- 2E-7	7E-4	7E-3
8	Nickel-56	D, all compounds except those given for W	1E+3	2E+3	8E-7	3E-9	2E-5	2E-4
		W, oxides, hydroxides, and carbides Vapor	-	1E+3 1E+3	5E-7 5E-7	2E-9 2E-9	-	-

			0cc1	Table I Ipational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhal: ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
8	Nickel-57	D, see ⁵⁶ Ni W, see ⁵⁶ Ni Vapor	2E+3 _	5E+3 3E+3 6E+3	2E-6 1E-6 3E-6	7E-9 4E-9 9E-9	2E-5 -	2E-4 -	
8	Nickel-59	D, see ⁵⁶ Ni W, see ⁵⁶ Ni Vapor	2E+4 	4E+3 7E+3 2E+3	2E-6 3E-6 8E-7	5E-9 1E-8 3E-9	3E-4 -	3E-3 _	
8	Nickel-63	D, see ⁵ ⁶ Ni W, see ⁵ ^N i Vapor	9E+3 _ _	2E+3 3E+3 8E+2	7E-7 1E-6 3E-7	2E-9 4E-9 1E-9	1E-4 -	1E-3 -	
8	Nickel-65	D, see ⁵⁶ Ni W, see ⁵⁶ Ni Vapor	8E+3 - -	2E+4 3E+4 2E+4	1E-5 1E-5 7E-6	3E-8 4E-8 2E-8	1E-4 - -	1E-3 - -	
8	Nickel-66	D, see ⁵⁶ Ni	4E+2	2E+3	7E-7	2E-9	-	-	
		W, see ⁵⁶ Ni Vapor	LLI wall (5E+2) - -	- 6E+2 3E+3	- 3E-7 1E-6	- 9E-10 4E-9	6E-6 - -	6E-5 - -	
9	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 Y, oxides and hydroxides	-	1E+5 1E+5	5E-5 4E-5	2E-7 1E-7	-		
9	Copper-61	D, see ⁶⁰ Cu W, see ⁶⁰ Cu Y, see ⁶⁰ Cu	1E+4 - -	3E+4 4E+4 4E+4	1E-5 2E-5 1E-5	4E-8 6E-8 5E-8	2E-4 -	2E-3 _	
9	Copper-64	D, see ⁶⁰ Cu W, see ⁶⁰ Cu Y, see ⁶⁰ Cu	1E+4 - -	3E+4 2E+4 2E+4	1E-5 1E-5 9E-6	4E-8 3E-8 3E-8	2E-4 -	2E-3 -	
9	Copper-67	D, see 60 Cu W, see 60 Cu Y, see 60 Cu	5E+3 - -	8E+3 5E+3 5E+3	3E-6 2E-6 2E-6	1E-8 7E-9 6E-9	6E-5 -	6E-4 -	
0	Zinc-62	Y, all compounds	1E+3	3E+3	1E-6	4E-9	2E-5	2E-4	
0	Zinc-63 ²	Y, all compounds	2E+4 St wall (3E+4)	7E+4 -	3E-5	9E-8	- 3E-4	- 3E-3	

			Occi	Table I upational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
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	2E+5	7E-5	2E-7	-	-
		W, oxides, hydroxides,	(6E+4)	-		-	9E-4	9E-3
		carbides, halides, and nitrates	-	2E+5	8E-5	3E-7	-	-
31	Gallium-66	D, see ⁶⁵ Ga W, see ⁶⁵ Ga	1E+3 -	4E+3 3E+3	1E-6 1E-6	5E-9 4E-9	1E-5 -	1E-4
31	Gallium-67	D, see ⁶⁵ Ga W, see ⁶⁵ Ga	7E+3 -	1E+4 1E+4	6E-6 4E-6	2E-8 1E-8	1E-4	1E-3
31	Gallium-68 ²	D, see ⁶⁵ Ga W, see ⁶⁵ Ga	2E+4 -	4E+4 5E+4	2E-5 2E-5	6E-8 7E-8	2E-4	2E-3
31	Gallium-70 ²	D, see ⁶⁵ Ga	5E+4	2E+5	7E-5	2E-7	-	-
		W, see ⁶⁵ Ga	St wall (7E+4) -	- 2E+5	- 8E-5	- 3E-7	1E-3	1E-2 -
31	Gallium-72	D, see ⁶⁵ Ga W, see ⁶⁵ Ga	1E+3 -	4E+3 3E+3	1E-6 1E-6	5E-9 4E-9	2E-5	2E-4
31	Gallium-73	D, see ⁶⁵ Ga W, see ⁶⁵ Ga	5E+3	2E+4 2E+4	6E-6 6E-6	2E-8 2E-8	7E-5	7E-4
32	Germanium-66	D, all compounds except those given for W	_ 2E+4	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 66 Ge	3E+4	9E+4	4E-5	1E-7	-	_
			St wall (4E+4)	_	_	_	6E-4	6E-3
		W, see ⁶⁶ Ge		1E+5	4E-5	1E-7	_	-
32	Germanium-68	D, see ⁶⁶ Ge	5E+3	4E+3	2E-6	5E-9	6E-5	6E-4

			Table I Occupational Values			Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1 Oral Ingestion	Col. 2 Inhala	Col. 3 ation	Col. 1	Col. 2	Monthly Average	
Atomic No.	Radionuclide	Class	ALÍ (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentration (µCi/ml)	
		W, see ⁶⁶ Ge	-	1E+2	4E-8	1E-10	_	-	
2	Germanium-69	D, see ⁶⁶ Ge W, see ⁶⁶ Ge	1E+4 -	2E+4 8E+3	6E-6 3E-6	2E-8 1E-8	2E-4	2E-3 -	
2	Germanium-71	D, see ⁶⁶ Ge	5E+5	4E+5	2E-4	6E-7	7E-3	7E-2	
		W, see ⁶⁶ Ge	-	4E+4	2E-5	6E-8	-	-	
32	Germanium-75 ²	D, see ⁶⁶ Ge	4E+4	8E+4	3E-5	1E-7	-	-	
		W, see ⁶⁶ Ge	St wall (7E+4) -	- 8E+4	- 4E-5	 1E-7	9E-4	<u>9</u> E-3	
2	Germanium-77	D, see ⁶⁶ Ge W, see ⁶⁶ Ge	9E+3	1E+4 6E+3	4E-6 2E-6	1E-8 8E-9	1E-4	1E-3	
2	Germanium-78 ²	D, see ⁶⁶ Ge	2E+4	2E+4	9E-6	3E-8	-	-	
			St wall (2E+4)	-	-		3E-4	3E-3	
_		W, see ⁶⁶ Ge	-	2E+4	9E-6	3E-8	~	-	
3	Arsenic-69 ²	W, all compounds	3E+4 St wall (4E+4)	1E+5 -	5E-5	2E-7	- 6E-4	- 6E-3	
3	$Arsenic-70^{2}$	W, all compounds	1E+4	5E+4	2E-5	7E-8	2E-4	2E-3	
3	Arsenic-71	W, all compounds	4E+3	5E+3	2E-6	6E-9	5E-5	5E-4	
3	Arsenic-72	W, all compounds	9E+2	1E+3	6E-7	2E-9	1E-5	1E-4	
3	Arsenic-73	W, all compounds	8E+3	2E+3	7E-7	2E-9	1E-4	1E-3	
3	Arsenic-74	W. all compounds	1E+3	8E+2	3E-7	1E-9	2E-5	2E-4	
3	Arsenic-76	W, all compounds	1E+3	1E+3	6E-7	2E-9	1E-5	1E-4	
3	Arsenic-77	W, all compounds	4E+3	5E+3	2E-6	7E-9	-	_	
			LLI wall (5E+3)	-	-	-	6E-5	6E-4	
3	Arsenic-78 ²	W, all compounds	8E+3	2E+4	9E-6	3E-8	1E-4	1E-3	
4	Selenium-70 ²	D, all compounds except those given for W W, oxides, hydroxides,	2E+4	4E+4	2E-5	5E-8	1E-4	1E-3	
		carbides, and elemental Se	1E+4	4E+4	2E-5	6E-8	_	_	

			Occi	Table I Ipational Va	lues	Efflu	e II ent rations	Table III Releases to Sewers Monthly Average Concentration
Atomic	Radionuclide	Class	Col. 1 Oral Ingestion ALI	Col. 2 Inhal ALI	Col. 3 ation DAC	Col. 1 Air	Col. 2	
No.			(µĈi)	(µĈi)	(µCi/ml)	(µCi/ml)	Water (µCi/ml)	(µCi/ml)
		、						
34	Selenium-73m ²	D, see ⁷⁰ Se W, see ⁷⁰ Se	6E+4 3E+4	2E+5 1E+5	6E-5 6E-5	2E-7 2E-7	4E-4 -	4E-3
34	Selenium-73	D, see ⁷⁰ Se W, see ⁷⁰ Se	3E+3	1E+4 2E+4	5E-6 7E-6	2E-8 2E-8	4E-5	4E-4
34	Selenium-75	D, see ⁷⁰ Se W, see ⁷⁰ Se	5E+2	7E+2 6E+2	3E-7 3E-7	1E-9 8E-10	7E-6	7E-5
34	Selenium-79	D, see ⁷⁰ Se W, see ⁷⁰ Se	6E+2	8E+2 6E+2	3E-7 2E-7	1E-9 8E-10	8E-6	8E-5
34	Selenium-81m ²	D, see ⁷⁰ Se W, see ⁷⁰ Se	4E+4 2E+4	7E+4 7E+4	3E-5 3E-5	9E-8 1E-7	3E-4	<u>3</u> E-3
34	Selenium-81 ²	D, see ⁷⁰ Se	6E+4 St wall	2 E +5	9E-5	3E-7	-	-
		W, see ⁷⁰ Se	(8E+4) -	- 2E+5	_ 1E-4	- 3E-7	1E-3 -	1E-2 -
34	Selenium-83 ²	D, see ⁷⁰ Se W, see ⁷⁰ Se	4E+4 3E+4	1E+5 1E+5	5E-5 5E-5	2E-7 2E-7	4E-4	4E-3
35	$Bromine-74m^2$	D, bromides of H, Li, Na, K, Rb, Cs, and Fr	1E+4	4E+4	2E-5	5E-8	-	-
		W, bromides of lantha- nides, 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	St wall (2E+4)	- 4E+4	- 2E-5	- 6E-8	3E-4	3E-3
35	Bromine-74 ²	D, see ^{74m} Br	2E+4 St wall	7E+4	3E-5	1E-7	-	-
		W, see ^{74m} Br	(4E+4)	- 8E+4	- 4E-5	- 1E-7	5E-4	5E-3 -
5	Bromine-75 ²	D, see ^{74m} Br	3E+4 St wall	5E+4	2E-5	7E-8	-	-
		W, see ^{74m} Br	(4E+4)	 5E+4	- 2E-5	- 7E-8	5E-4 -	5E-3 -
5	Bromine-76	D, see ^{74m} Br W, see ^{74m} Br	4E+3	5E+3 4E+3	2E-6 2E-6	7E-9 6E-9	5E-5	5E-4

			Occi	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Oral Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
35	Bromine-77	D, see ^{74m} Br W, see ^{74m} Br	2E+4 -	2E+4 2E+4	1E-5 8E-6	3E-8 3E-8	2E-4 -	2E-3	
35	Bromine-80m	D, see ^{74m} Br W, see ^{74m} Br	2E+4 -	2E+4 1E+4	7E-6 6E-6	2E-8 2E-8	3E-4 -	3E-3 -	
35	Bromine-80 ²	D, see ^{74m} Br	5E+4 St wall	2E+5	8E-5	3E-7	-	-	
		W, see ^{74m} Br	(9E+4)	- 2E+5	- 9E-5	- 3E-7	1E-3 -	1E-2 -	
35	Bromine-82	D, see ^{74m} Br W, see ^{74m} Br	3E+3	4E+3 4E+3	2E-6 2E-6	6E-9 5E-9	4E-5	4E-4	
85	Bromine-83	D, see ^{74m} Br	5E+4 St wall	6E+4	3E-5	9E-8	-	-	
		W, see ^{74m} Br	(7E+4)	- 6E+4	- 3E-5		9E-4	9E-3	
5	Bromine-84 ²	D, see ^{74m} Br	2E+4	6E+4	2E-5	8E-8	-	-	
		W, see ^{74m} Br	St wall (3E+4) -	- 6E+4	- 3E-5	- 9E-8	4E-4	4E-3	
36	Krypton-74 ²	Submersion ¹	-	-	3E-6	1E-8	-	-	
86	Krypton-76	Submersion ¹	-	-	9E-6	4E-8	~	-	
86	Krypton-77 ²	Submersion ¹	-	-	4E-6	2E-8	-	-	
36	Krypton-79	Submersion ¹	-	-	2E-5	7E-8	-	-	
86	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	-	-	
86	Krypton-85	Submersion ¹	-	-	1E-4	7E-7	-	-	
6	Krypton-87 ²	Submersion	-	-	5E-6	2E-8	-	-	
6	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	- 8E-4	- 8E-3	
37	Rubidium-81m ²	D, all compounds	2E+5	3E+5	1E-4	5E-7	-	-	

				Table I Occupational Values			e II ent rations	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
			St wall (3E+5)	-	-		4E-3	4E-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
37	Rubidium-88 ²	D, all compounds	2E+4	6E+4	3E-5	9E-8	-	-
		(3E+4)	St wall	-	-	4E-4	4E-3	
37	Rubidium-89 ²	D, all compounds	4E+4	1E+5	6E-5	2E-7	-	-
			St wall (6E+4)	-	-	-	9E-4	9E-3
38	Strontium-80 ²	D, all soluble compounds except SrTiO, Y, all insoluble com-	4E+3	1E+4	5E-6	2E-8	6E-5	6E-4
		pounds and SrTi03	-	1E+4	5E-6	2E-8	-	-
38	Strontium-81 ²	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	3E+4 2E+4	8E+4 8E+4	3E-5 3E-5	1E-7 1E-7	3E-4	3E-3
38	Strontium-82	D, see ⁸⁰ Sr	3E+2	4E+2	2E-7	6E-10	-	
		Y, see ⁸⁰ Sr	LLI wall (2E+2) 2E+2	- 9E+1	- 4E-8	 1E-10	3E-6	3E-5 -
38	Strontium-83	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	3E+3 2E+3	7E+3 4E+3	3E-6 1E-6	1E-8 5E-9	3E-5	3E-4
38	Strontium-85m ²	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	2E+5	6E+5 8E+5	3E-4 4E-4	9E-7 1E-6	3E-3	3E-2
38	Strontium-85	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	3E+3	3E+3 2E+3	1E-6 6E-7	4E-9 2E-9	4E-5	4E-4
38	Strontium-87m	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	5E+4 4E+4	1E+5 2E+5	5E-5 6E-5	2E-7 2E-7	6E-4	6E-3
38	Strontium-89	D, see ⁸⁰ Sr	6E+2	8E+2	4E-7	1E-9	_	-

Atomic No.			Occu	Table I Occupational Values			e II ent rations	Table III Releases to Sewers	
	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
		Y, see ⁸⁰ Sr	LLI wall (6E+2) 5E+2	_ 1E+2	- 6E-8	_ 2E-10	8E-6 -	8E-5 -	
8	Strontium-90	D, see ⁸⁰ Sr	3E+1 Bone surf	2E+1 Bone surf	8E-9	-	-		
		Y, see ⁸⁰ Sr	(4E+1) -	(2E+1) 4E+0	_ 2E-9	3E-11 6E-12	5E-7 -	5E-6 -	
8	Strontium-91	D, see ${}^{80}_{0}$ Sr Y, see ${}^{80}_{0}$ Sr	2E+3	6E+3 4E+3	2E-6 1E-6	8E-9 5E-9	2E-5	2E-4	
8	Strontium-92	D, see ⁸⁰ Sr Y, see ⁸⁰ Sr D, see ⁸⁰ Sr Y, see ⁸⁰ Sr	3E+3	9E+3 7E+3	4E-6 3E-6	1E-8 9E-9	4E-5 -	4E-4 -	
9	Yttrium-86m²	W, all compounds except those given for Y Y, oxides and hydroxides	2E+4 -	6E+4 5E+4	2E-5 2E-5	8E-8 8E-8	3E-4	3E-3	
9	Yttrium-86	W, see ^{86m} Y Y, see ^{86m} Y	1E+3	3E+3 3E+3	1E-6 1E-6	5E-9 5E-9	2E-5 -	2E-4	
9	Yttrium-87	W, see ^{86m} Y Y, see ^{86m} Y	2E+3 -	3E+3 3E+3	1E-6 1E-6	5E-9 5E-9	3E-5 -	<u>3</u> E-4	
9	Yttrium-88	W, see ^{86m} Y Y, see ^{86m} Y	1E+3 -	3E+2 2E+2	1E-7 1E-7	3E-10 3E-10	1E-5 -	$\frac{1E-4}{-}$	
9	Yttrium-90m	W, see ^{86m} Y Y, see ^{86m} Y	8E+3 -	1E+4 1E+4	5E-6 5E-6	2E-8 2E-8	1E-4	1E-3 -	
9	Yttrium-90	W, see ^{86m} Y	4E+2 LLI wall	7E+2	3E-7	9E-10	-	-	
		Y, see ^{86m} Y	(5E+2) -	- 6E+2	- 3E-7	_ 9E-10	7E-6 -	7E-5 -	
9	Yttrium-91m ²	W, see ^{86m} Y Y, see ^{86m} Y	1 E +5	2E+5 2E+5	1E-4 7E-5	3E-7 2E-7	2E-3	2E-2	
9	Yttrium-91	W, see ^{86m} Y	5E+2 LLI wall	2E+2	7E-8	2E-10		-	
		Y, see ^{86m} Y	(6E+2)	- 1E+2	- 5E-8		8E-6 -	8E-5	
9	Yttrium-92	W, see ^{86m} Y Y, see ^{86m} Y	3E+3	9E+3 8E+3	4E-6 3E-6	1E-8 1E-8	4E-5	4E-4 -	
9	Yttrium-93	W, see ^{86m} Y Y, see ^{86m} Y	1E+3 _	3E+3 2E+3	1E-6 1E-6	4E-9 3E-9	2E-5	2E-4	

				Table I pational Valu	les	Table II Effluent Concentrations		Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 Lion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
39	Yttrium-94 ²	W, see ^{86m} Y	2E+4 St_wall	8E+4	3E-5	1E-7	-	-
		Y, see ^{86m} Y	(3E+4) _	- 8E+4	_ 3E~5	- 1E-7	4E-4 -	4E-3
9	Yttrium-95 ²	W, see ^{86m} Y	4E+4 St wall	2E+5	6E-5	2E-7	-	-
		Y, see ^{86m} Y	(5E+4) -	_ 1E+5	- 6E-5	- 2E-7	7E-4 -	7E-3
0	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 Y, carbide	-	3E+3 2E+3	1E-6 1E-6	4E-9 3E-9	-	Ξ
0	Zirconium-88	D, see ⁸⁶ Zr W, see ⁸⁶ Zr Y, see ⁸⁶ Zr	4E+3 -	2E+2 5E+2	9E-8 2E-7	3E-10 7E-10	5E-5	5E-4 -
0	Zirconium-89			3E+2	1E-7	4E-10	-	~
0	2110011100-89	D, see ⁸⁶ Zr W, see ⁸⁶ Zr Y, see ⁸⁶ Zr	2E+3 	4E+3 2E+3 2E+3	1E-6 1E-6 1E-6	5E-9 3E-9 3E-9	2E-5 - -	2E-4 -
0	Zirconium-93	D, see ⁸⁶ Zr	1E+3 Bone surf	6E+0 Bone surf	3E-9	-	-	_
		W, see ⁸⁶ Zr	(3E+3)	(2E+1) 2E+1	- 1E-8	2E-11 -	4E-5 -	4E-4
		Y, see ⁸⁶ Zr	-	Bone surf (6E+1) 6E+1	- 2E-8	9E-11	-	-
			-	Bone surf (7E+1)	_	9E-11	_	-
0	Zirconium-95	D, see ⁸⁶ Zr	1E+3	1E+2 Bone surf	5E-8	-	2E-5	2E-4
		W, see ⁹⁶ Zr Y, see ⁸⁶ Zr	-	(3E+2) 4E+2 3E+2	- 2E-7 1E-7	4E-10 5E-10 4E-10	-	-
0	Zirconium-97	D, see ${}^{86}Zr$ W, see ${}^{86}Zr$ Y, see ${}^{86}Zr$	6E+2	2E+3 1E+3 1E+3	8E-7 6E-7 5E-7	3E-9 2E-9 2E-9 2E-9	9E-6 -	9E-5 -
1	Niobium-88 ²	W, all compounds except those given for Y	5E+4	2E+5	9E-5	3E-7	-	_
		Y, oxides and hydroxides	St wall (7E+4)	_ 2E+5	- 9E-5	- 3E-7	1E-3	1E-2

	Radionuclide		0001	Table I upational Values		Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic		Class	Col. 1 Oral Ingestion ALI	Col. 2 Inhal ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration
No.			(µCi)	(µC1)	(µC1/m1)	(µC1/m1)	(µC1/m1)	(µCi/ml)
11	Niobium-89 ²	W, see ⁸⁸ Nb	1E+4	4E+4	2E-5	6E-8	1E-4	1E-3
±	(66 min)	Y, see 88 Nb	~	4E+4	2E-5	5E-8	-	-
1	Niobium-89	W, see ⁸⁸ Nb	5E+3	2E+4	8E-6	3E-8	7E-5	7E-4
	(122 min)	Y, see ⁸⁸ Nb	-	2E+4	6E-6	2E-8	-	-
1	Niobium-90	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	1E+3	3E+3 2E+3	1E-6 1E-6	4E-9 3E-9	1E-5 -	1E-4 _
1	Niobium-93m	W, see ⁸⁸ Nb	9E+3 LLI wall	2E+3	8E-7	3E-9	-	-
		Y, see ⁸⁸ Nb	(1E+4) -	- 2E+2	- 7E-8	_ 2E-10	2E-4	2E-3
1	Niobium-94	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	9E+2	2E+2 2E+1	8E-8 6E-9	3E-10 2E-11	1E-5 -	1E-4
1	Niobium-95m	W, see ⁸⁸ Nb	2E+3	3E+3	1E-6	4E-9	-	-
		Y, see ⁸⁸ Nb	LLI wall (2E+3) -	- 2E+3	- 9E-7	- 3E-9	3E-5	3E-4
1	Niobium-95	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	2E+3	1E+3 1E+3	5E-7 5E-7	2E-9 2E-9	3E-5	<u>3</u> E-4
1	Niobium-96	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	1E+3 -	3E+3 2E+3	1E-6 1E-6	4E-9 3E-9	2E-5	<u>2</u> E-4
1	Niobium-97 ²	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	2E+4	8E+4 7E+4	3E-5 3E-5	1E-7 1E-7	3E-4	<u>3</u> E-3
1	Niobium-98 ²	W, see ⁸⁸ Nb Y, see ⁸⁸ Nb	1E+4	5E+4 5E+4	2E-5 2E-5	8E-8 7E-8	2E-4	2E-3
2	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 ₂	2E+3	5E+3	2E-6	6E-9	-	-
2	Molybdenum-93m	D, see ⁹⁰ Mo Y, see ⁹⁰ Mo	9E+3 4E+3	2E+4 1E+4	7E-6 6E-6	2E-8 2E-8	6E-5 -	6E-4
2	Molybdenum-93	D, see ⁹⁰ Mo Y, see ⁹⁰ Mo	4E+3 2E+4	5E+3 2E+2	2E-6 8E-8	8E-9 2E-10	5E-5 -	5E-4
2	Molybdenum~99	D, see ⁹⁰ Mo	2E+3 LLI wall	3E+3	1E-6	4E-9	-	-

			Table I Occupational Values			Table II Effluent Concentrations		Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly
No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
		Y, see ⁹⁰ Mo	(1E+3) 1E+3	_ 1E+3	- 6E-7	- 2E~9	2E-5 -	2E-4
2	Molybdenum-101 ²	D, see ⁹⁰ Mo	4E+4 St wall	1E+5	6E-5	2E-7	-	-
		Y, see ⁹⁰ Mo	(5E+4) -	- 1E+5	- 6E-5	_ 2E-7	7E-4	7E-3 -
13	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	-	-
3	Technetium-93	D, see 9^{3m} TC W, see 9^{3m} TC	3E+4 -	7E+4 1E+5	3E-5 4E-5	1E-7 1E-7	4E-4 -	4E-3
3	Technetium-94m ²	D, see 9^{3m} TC W, see 9^{3m} TC	2E+4	4E+4 6E+4	2E-5 2E-5	6E-8 8E-8	3E-4	3E-3
3	Technetium-94	D, see ^{93m} Tc W, see ^{93m} Tc	9E+3	2E+4 2E+4	8E-6 1E-5	3E-8 3E-8	1E-4	1E-3
3	Technetium-95m	D, see ^{93m} Tc W, see ^{93m} Tc	4E+3	5E+3 2E+3	2E-6 8E-7	8E-9 3E-9	5E-5	5E-4
3	Technetium-95	D, see ^{93m} Tc W, see ^{93m} Tc	1E+4 -	2E+4 2E+4	9E-6 8E-6	3E-8 3E-8	1E-4	1E-3
3	Technetium-96m ²	D, see ${}^{93m}_{93m}$ Tc W, see 93m Tc	2E+5	3E+5 2E+5	1E-4 1E-4	4E-7 3E-7	2E-3	2E-2
3	Technetium-96	D, see $93m$ Tc W, see $93m$ Tc	2E+3	3E+3 2E+3	1E-6 9E-7	5E-9 3E-9	3E-5	3E-4
3	Technetium-97m	D, see ^{93m} Tc	5E+3	7E+3 St wall	3E-6	-	6E-5	6E-4
		W, see ^{93m} TC	-	(7E+3) 1E+3	- 5E-7	1E-8 2E-9	Ξ	-
3	Technetium-97	D, see ^{93m} Tc W, see ^{93m} Tc	4E+4	5E+4 6E+3	2E-5 2E-6	7E-8 8E-9	5E-4	5E-3
3	Technetium-98	D, see ^{93m} Tc W, see ^{93m} Tc	1E+3 -	2E+3 3E+2	7E-7 1E-7	2E-9 4E-10	1E-5 -	1E-4
3	Technetium-99m	D, see ^{93m} Tc W, see ^{93m} Tc	8E+4 -	2E+5 2E+5	6E~5 1E-4	2E-7 3E-7	1E-3 -	1E-2
3	Technetium-99	D, see ^{93m} Tc	4E+3	5E+3	2E-6	-	6E-5	6E-4

			Occu	Table I upational Val	ues.	Tabl Efflu Concent	ent	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Sewers Monthly Average Concentration (µCi/ml) - - 2E-2 - 4E-3 - 2E-3 - 1E-3 - 3E-4 - 7E-4 - 3E-5 - 2E-3 - 2E-2 - - 2E-3 - - 2E-3 - - 2E-3 - - 2E-3 - - - - - - - - - - - - -
		W, see ^{93m} TC	-	St wall (6E+3) 7E+2	- 3E-7	8E-9 9E-10	-	-
43	Technetium-101 ²	D, see ^{93m} Tc	9E+4 St wall	3E+5	1E-4	5E-7	-	-
		W, see ^{93m} Tc	(1E+5) -	- 4E+5	- 2E-4	- 5E-7	2E~3	2E-2
43	Technetium-104 ²	D, see ^{93m} TC	2E+4	7E+4	3E-5	1E-7	-	-
		W, see ^{93m} Tc	St wall (3E+4) -	_ 9E+4	- 4E-5	- 1E-7	4E-4 -	4E-3
44	Ruthenium-94 ²	D, all compounds except those given for W and Y W, halides Y, oxides and hydroxides	2E+4 _ _	4E+4 6E+4 6E+4	2E-5 3E-5 2E-5	6E-8 9E-8 8E-8	2E-4 -	
14	Ruthenium-97	D, see ⁹⁴ Ru W, see ⁹⁴ Ru Y, see ⁹⁴ Ru	8E+3 - -	2E+4 1E+4 1E+4	8E-6 5E-6 5E-6	3E-8 2E-8 2E-8	1E-4 - -	
14	Ruthenium-103	D, see ⁹⁴ Ru W, see ⁹⁴ Ru Y, see ⁹⁴ Ru	2E+3 _ _	2E+3 1E+3 6E+2	7E-7 4E-7 3E-7	2E-9 1E-9 9E-10	3E-5 -	-
	Ruthenium-105	D, see ⁹⁴ Ru W, see ⁹⁴ Ru Y, see ⁹⁴ Ru	5E+3 _ _	1E+4 1E+4 1E+4	6E-6 6E-6 5E-6	2E-8 2E-8 2E-8	7E-5 	-
14	Ruthenium-106	D, see ⁹⁴ Ru	2E+2 LLI wall	9E+1	4E-8	1E-10	-	-
		W, see ⁹⁴ Ru Y, see ⁹⁴ Ru	(2E+2) - -	- 5E+1 1E+1	- 2E-8 5E-9	- 8E-11 2E-11	3E-6 -	~
15	Rhodium-99m	D, all compounds except those given for W and Y W, halides Y, oxides and hydroxides	2E+4 	6E+4 8E+4 7E+4	2E-5 3E-5 3E-5	8E-8 1E-7 9E-8	2E-4 - -	
15	Rhodium-99	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	2E+3 -	3E+3 2E+3 2E+3	1E-6 9E-7 8E-7	4E-9 3E-9 3E-9	3E-5 -	3E-4 -
45	Rhodium-100	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	2E+3 	5E+3 4E+3 4E+3	2E-6 2E-6 2E-6	7E-9 6E-9 5E-9	2E-5 - -	2E-4

			Occi	Table I upational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhal ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
15	Rhodium-101m	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	6E+3 -	1E+4 8E+3 8E+3	5E-6 4E-6 3E-6	2E-8 1E-8 1E-8	8E-5 - -	8E-4 - -
15	Rhodium-101	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	2E+3 _ _	5E+2 8E+2 2E+2	2E-7 3E-7 6E-8	7E-10 1E-9 2E-10	3E-5 -	3E-4 -
45	Rhodium-102m	D, see ^{99m} Rh	1E+3 LLI wall	5E+2	2E-7	7E-10	-	-
		W, see ^{99m} Rh Y, see ^{99m} Rh	(1E+3) _ _	- 4E+2 1E+2	- 2E-7 5E-8	- 5E-10 2E-10	2E-5 - -	2E-4 _
15	Rhodium-102	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	6E+2 - -	9E+1 2E+2 6E+1	4E-8 7E-8 2E-8	1E-10 2E-10 8E-11	8E-6 -	8E-5 - -
45	Rhodium-103m ²	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	4E+5 -	1E+6 1E+6 1E+6	5E-4 5E-4 5E-4	2E-6 2E-6 2E-6	6E-3 	6E-2
5	Rhodium-105	D, see ^{99m} Rh	4E+3 LLI wall	1E+4	5E-6	2E-8	-	
		W, see ^{99m} Rh Y, see ^{99m} Rh	(4E+3) - -	- 6E+3 6E+3	- 3E-6 2E-6	- 9E-9 8E-9	5E~5 - -	5E-4 - -
15	Rhodium-106m	D, see ^{99m} Rh W, see ^{99m} Rh Y, see ^{99m} Rh	8E+3 - -	3E+4 4E+4 4E+4	1E-5 2E-5 1E-5	4E-8 5E-8 5E-8	1E-4 - -	1E-3 -
15	Rhodium-107 ²	D, see ^{99m} Rh	7E+4 St wall	2E+5	1E-4	3E-7	-	-
		W, see ^{99m} Rh Y, see ^{99m} Rh	(9E+4) - -	- 3E+5 3E+5	- 1E-4 1E-4	- 4E-7 3E-7	1E-3 _	1E-2 -
16	Palladium-100	D, all compounds except those given for W and Y W, nitrates Y, oxides and hydroxides	1E+3 	1E+3 1E+3 1E+3	6E-7 5E-7 6E-7	2E-9 2E-9 2E-9	2E-5 -	2E-4 - -
46	Palladium-101	D, see ¹⁰⁰ Pd W, see ¹⁰⁰ Pd Y, see ¹⁰⁰ Pd	1E+4 _ _	3E+4 3E+4 3E+4	1E-5 1E-5 1E-5	5E-8 5E-8 4E-8	2E-4 	2E-3
16	Palladium-103	D, see ¹⁰⁰ Pd	6E+3 LLI wall	6E+3	3E-6	9E-9	-	-

			0001	Table I Ipational Val	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 1E-3 - - 5E-3 - 3E-4 - 9E-3 - 5E-3 - 5E-3 -
tomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Averagē Concentration
		W, see ¹⁰⁰ Pd Y, see ¹⁰⁰ Pd	(7E+3) _ _	- 4E+3 4E+3	- 2E-6 1E-6	- 6E-9 5E-9	1E-4 -	-
6	Palladium-107	D, see ¹⁰⁰ Pd	3E+4 LLI wall	2E+4 Kidneys	9E-6	-	-	-
		W, see 100 Pd Y, see 100 Pd	(4E+4) - -	(2E+4) 7E+3 4E+2	- 3E-6 2E-7	3E-8 1E-8 6E-10	5E-4 - -	
6	Palladium-109	D, see 100 Pd W, see 100 Pd Y, see 100 Pd Y, see 100 Pd	2E+3 - -	6E+3 5E+3 5E+3	3E-6 2E-6 2E-6	9E-9 8E-9 6E-9	3E-5 -	-
7	Silver-102 ²	D, all compounds except those given for W and Y	5E+4 St wall	2E+5	8E-5	2E-7	-	
		W, nitrates and sulfides Y, oxides and hydroxides	(6E+4) - -	- 2E+5 2E+5	- 9E-5 8E-5	- 3E-7 3E-7	9E-4 - -	
7	Silver-103 ²	D, see 102 Ag W, see 102 Ag Y, see 102 Ag	4E+4 	1E+5 1E+5 1E+5	4E-5 5E-5 5E-5	1E-7 2E-7 2E-7	5E-4 - -	
7	Silver-104m ²	D, see 102 Ag W, see 102 Ag Y, see 102 Ag	3E+4 -	9E+4 1E+5 1E+5	4E-5 5E-5 5E-5	1E-7 2E-7 2E-7	4E-4 - -	4E-3
7	Silver-104 ²	D, see ${}^{102}_{102}$ Ag W, see ${}^{102}_{102}$ Ag Y, see 102 Ag	2E+4 - -	7E+4 1E+5 1E+5	3E-5 6E-5 6E-5	1E-7 2E-7 2E-7	3E-4 _	3E-3 _
7	Silver-105	D, see ${}^{102}_{102}$ Ag W, see ${}^{102}_{102}$ Ag Y, see 102 Ag	3E+3 _	1E+3 2E+3 2E+3	4E-7 7E-7 7E-7	1E-9 2E-9 2E-9	4E-5 -	4E-4 _
7	Silver-106m	D, see 102 Ag W, see 102 Ag Y, see 102 Ag	8E+2 -	7E+2 9E+2 9E+2	3E-7 4E-7 4E-7	1E-9 1E-9 1E-9	1E-5 -	1E-4 -
7	Silver-106 ²	D, see 102 Ag	6E+4	2E+5	8E~5	3E-7	-	-
		W, see ¹⁰² Ag Y, see ¹⁰² Ag	St. wall (6E+4) -	- 2E+5 2E+5	- 9E-5 8E-5	- 3E-7 3E-7	9E-4 - -	9E-3 - -
7	Silver-108m	D, see 10^{2} Ag W, see 10^{2} Ag	<u>6</u> E+2	2E+2 3E+2	8E-8 1E-7	3E-10 4E-10	9E-6	9E-5

			Occi	Table I upational Val	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml) - 6E-5 - - 2E-4 - 4E-4 - - 4E-4 - - 3E-3 - 3E-3 - - 3E-3 - - 6E-5 -
		Y, see ¹⁰² Ag	_	2E+1	1E-8	3E-11	-	-
17	Silver-110m	D, see 102 Ag W, see 102 Ag Y, see 102 Ag	5E+2 -	1E+2 2E+2 9E+1	5E-8 8E-8 4E-8	2E-10 3E-10 1E-10	6E-6 	-
47	Silver-111	D, see ¹⁰² Ag	9E+2 LLI wall	2E+3 Liver	6E-7	-	-	-
		W, see 102Ag Y, see 102Ag	(1E+3) - -	(2E+3) 9E+2 9E+2	- 4E-7 4E-7	2E-9 1E-9 1E-9	2E-5 - -	-
47	Silver-112	D, see 102 Ag W, see 102 Ag Y, see 102 Ag	3E+3 	8E+3 1E+4 9E+3	3E-6 4E-6 4E-6	1E-8 1E-8 1E-8	4E-5 -	-
17	Silver-115 ²	D, see ¹⁰² Ag	3E+4	9E+4	4E-5	1E-7	-	-
		W, see 102 Ag Y, see 102 Ag	St wall (3E+4) -	- 9E+4 8E+4	- 4E-5 3E-5	- 1E-7 1E-7	4E-4 - -	-
8	Cadmium-104 ²	D, all compounds except those given for W and Y W, sulfides, halides,	2E+4	7E+4	3E-5	9E-8	3E-4	3E-3
		Y, oxides and hydroxides	Ξ	1E+5 1E+5	5E-5 5E-5	2E-7 2E-7	-	
18	Cadmium-107	D, see 104 Cd W, see 104 Cd Y, see 104 Cd	2E+4 	5E+4 6E+4 5E+4	2E-5 2E-5 2E-5	8E-8 8E-8 7E-8	3E-4 _	-
8	Cadmium-109	D, see ¹⁰⁴ Cd	3E+2	4E+1	1E-8	-	-	-
		W, see 104 Cd	Kidneys (4E+2) -	Kidneys (5E+1) 1E+2 Kidneys	- 5E-8	7E-11 -	6E-6 -	6E-5 -
		Y, see ¹⁰⁴ Cd	-	(1E+2) 1E+2	- 5E-8	2E-10 2E-10	-	-
8	Cadmium-113m	D, see ¹⁰⁴ Cđ	2E+1	2E+0	1E-9	-		-
		W, see ¹⁰⁴ Cd	Kidneys (4E+1) -	Kidneys (4E+0) 8E+0 Kidneys	- 4E-9	5E-12	5E-7 -	5E-6 -
		Y, see 104 Cd	-	(1E+1) 1E+1	- 5E-9	2E-11 2E-11	-	-
8	Cadmium-113	D, see ¹⁰⁴ Cd	2E+1	2E+0	9E-10	-	-	-

			0000	Table I pational Val	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral Ingestion	Col. 2 Inhala	Col. 3	Col. 1	Col. 2	Monthly Average
Atomic No.	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentration (µCi/ml)
		W, see ¹⁰⁴ Cđ	Kidneys (3E+1)	Kidneys (3E+0) 8E+0 Kidneys	- 3E-9	5E-12	4E-7 -	4E-6 -
		Y, see ¹⁰⁴ Cd	-	(1E+1) 1E+1	- 6E-9	2E-11 2E-11	Ξ	-
48	Cadmium-115m	D, see ¹⁰⁴ Cd	3E+2	5E+1 Kidneys	2E-8	-	4E-6	4E-5
		W, see ¹⁰⁴ Cd Y, see ¹⁰⁴ Cd	- -	(8E+1) 1E+2 1E+2	- 5E-8 6E-8	1E-10 2E-10 2E-10	-	
48	Cadmium-115	D, see ¹⁰⁴ Cd	9E+2 LLI wall	1E+3	6E-7	2E-9	-	-
		W, see 104 Cd Y, see 104 Cd	(1E+3) _ _	- 1E+3 1E+3	- 5E-7 6E-7	- 2E-9 2E-9	1E-5 -	1E-4 - -
48	Cadmium-117m	D, see ¹⁰⁴ Cd W, see ¹⁰⁴ Cd Y, see ¹⁰⁴ Cd	5E+3 - -	1E+4 2E+4 1E+4	5E-6 7E-6 6E-6	2E-8 2E-8 2E-8	6E-5 -	6E-4 - -
48	Cadmium-117	D, see 104 Cd W, see 104 Cd Y, see 104 Cd	5E+3 - -	1E+4 2E+4 1E+4	5E-6 7E-6 6E-6	2E-8 2E-8 2E-8	6E-5 _	6E-4 - -
49	Indium-109	D, all compounds except those given for W W, oxides, hydroxides, halides, and nitrates	2E+4	4E+4	2E-5	6E-8	3E-4	3E-3
			-	6E+4	3E-5	9E-8	-	-
49	Indium-110 ² (69.1 min)	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	2E+4 -	4E+4 6E+4	2E-5 2E-5	6E-8 8E-8	2E-4 -	2E-3
49	Indium-110 (4.9 h)	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	5E+3	2E+4 2E+4	7E-6 8E-6	2E-8 3E-8	7E-5	7E-4 -
49	Indium-111	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	4E+3	6E+3 6E+3	3E-6 3E-6	9E-9 9E-9	6E-5	6E-4 -
49	Indium-112 ²	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	2E+5 -	6E+5 7E+5	3E-4 3E-4	9E-7 1E-6	2E-3	2E-2 -
49	Indium-113m ²	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	5E+4 ~	1E+5 2E+5	6E-5 8E-5	2E-7 3E-7	7E-4 -	7E-3
49	Indium-114m	D, see ¹⁰⁹ In	3E+2 LLI wall	6E+1	3E-8	9E-11	-	-

			Occu	Tablë I upational Valu	ues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
		W, see ¹⁰⁹ In	(4E+2) -	_ 1E+2	- 4E-8	_ 1E-10	5E-6	5E-5
.9	Indium-115m	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	1E+4	4E+4 5E+4	2E-5 2E-5	6E-8 7E-8	2E-4	2E-3
9	Indium-115	D, see 109 In W, see 109 In	4E+1	1E+0 5E+0	6E-10 2E-9	2E-12 8E-12	5E-7 -	5E-6
9	Indium-116m ²	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	2E+4	8E+4 1E+5	3E-5 5E-5	1E-7 2E-7	3E-4	3E-3
9	Indium-117m ²	D, see ¹⁰⁹ In W, see ¹⁰⁹ In	1E+4 _	3E+4 4E+4	1E-5 2E-5	5E-8 6E-8	2E-4	2E-3 -
9	Indium-117 ²	D, see ¹⁰⁹ In W, see In	6E+4 -	2E+5 2E+5	7E-5 9E~5	2E-7 3E-7	8E-4	8E-3
9	Indium-119m ²	D, see ¹⁰⁹ In	4E+4 St_wall	1E+5	5E-5	2E-7	-	-
		W, see ¹⁰⁹ In	(5E+4) _	_ 1E+5	- 6E-5	- 2E-7	7E-4	7E-3 -
0	Tin-110	D, all compounds except those given for W W, sulfides, oxides, hydroxides, halides, nitrates, and stannic	4E+3	1E+4	5E-6	2E-8	5E~5	5E-4
•	mt . 1112	phosphate	-	1E+4	5E-6	2E-8	-	-
0	Tin-111 ²	D, see 110 Sn W, see 110 Sn	7E+4 -	2E+5 3E+5	9E-5 1E-4	3E-7 4E-7	1E-3 -	1E-2 -
D	Tin-113	D, see ¹¹⁰ Sn	2E+3 LLI wall	1E+3	5E-7	2E-9	-	-
		W, see ¹¹⁰ Sn	(2E+3) -	- 5E+2	- 2E-7	- 8E-10	3E-5 -	3E-4 -
0	Tin-117m	D, see ¹¹⁰ Sn	2E+3 LLI wall	1E+3 Bone surf	5E-7	-	-	-
		W, see ¹¹⁰ Sn	(2E+3) -	(2E+3) 1E+3	- 6E-7	3E-9 2E-9	3E-5	3E-4
0	Tin-119m	D, see ¹¹⁰ Sn	3E+3 LLI wall	2E+3	1E-6	3E-9	-	-
		W, see ¹¹⁰ Sn	(4E+3)	- 1E+3	- 4E-7	_ 1E-9	6E-5	6E-4
D	Tin-121m	D, see ¹¹⁰ Sn	3E+3	9E+2	4E-7	1E-9	-	-

			Occu	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 5E-4 - 8E-4 - 9E-5 - 6E-5 9E-4 - 1E-3 1E-2
omic	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala	Col. 3 ation DAC (µCi/ml)	Col. 1	Col. 2 Water	Average Concentration
lo .			(µC1)	(µCi)	(µC1/m1)	(µĈi/ml)	(µĈi/ml)	(µC1/m1)
		W, see ¹¹⁰ Sn	LLI wall (4E+3)	-		- 8E-10	5E-5	5E-4
0	Tin-121	W, See Sn D, see ¹¹⁰ Sn	- 6E+3	5E+2 2E+4	2E-7 6E-6	2E-8	-	-
,	1111-121	W, see 110 Sn	LLI wall (6E+3)	- 1E+4	5E-6	2E-8	8E-5 -	8E-4
D	Tin-123m ²	D, see 110 Sn W, see 110 Sn	5E+4 -	1E+5 1E+5	5E-5 6E-5	2E-7 2E-7	7E-4	
)	Tin-123	D, see ¹¹⁰ Sn	5E+2 LLI wall	6E+2	3E-7	9E-10	-	-
		W, see ¹¹⁰ Sn	(6E+2)	_ 2E+2	- 7E-8	 2E-10	9E~6 -	9E~5 -
)	Tin-125	D, see ¹¹⁰ Sn	4E+2 LLI wall	9E+2	4E-7	1E-9	-	-
		W, see ¹¹⁰ Sn	(5E+2) -	- 4E+2	- 1E-7	- 5E~10	6E-6	
D	Tin-126	D, see 110 Sn W, see 110 Sn	3E+2 -	6E+1 7E+1	2E-8 3E-8	8E-11 9E-11	4E-6 -	
C	Tin-127	D, see ¹¹⁰ Sn W, see ¹¹⁰ Sn	7E+3 -	2E+4 2E+4	8E-6 8E-6	3E-8 3E-8	9E-5 -	9E-4
0	Tin-128 ²	D, see ¹¹⁰ Sn W, see ¹¹⁰ Sn	9E+3	3E+4 4E+4	1E-5 1E-5	4E-8 5E-8	1E-4 -	1E-3
1	Antimony-115 ²	D, all compounds except those given for W W, oxides, hydroxides, halides, sulfides,	8E+4	2E+5	1E-4	3E-7	1E-3	1E-2
		sulfates, and nitrates	-	3E+5	1E-4	4E-7	-	-
-	Antimony-116m ²	D, see 115 Sb W, see 115 Sb	2E+4 -	7E+4 1E+5	3E-5 6E-5	1E-7 2E-7	3E-4	3E-3
1	Antimony-116 ²	D, see ¹¹⁵ Sb	7E+4 St wall	3E+5	1E-4	4E-7	-	-
		W, see ¹¹⁵ Sb	(9E+4)		_ 1E-4	- 5E-7	1E-3 -	1E-2
	Antimony-117	D, see 115 Sb W, see 115 Sb	7E+4	2E+5 3E+5	9E-5 1E-4	3E-7 4E-7	9E-4	9E-3
	Antimony-118m	D, see ¹¹⁵ Sb	6E+3	2E+4	8E-6	3E-8	7E-5	7E-4

		······································	0000	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Releases to Sewers Monthly Average Concentration (µCi/ml) - 2E-3 - - 2E-2 - 1E-4 - 1E-4 - 3E-2 - 7E-5 3E-4 - 9E-3 - 7E-5 - 3E-4 - 1E-4 - 1E-4 - 1E-4 - 1E-4 - 1E-4 - 1E-5 - 3E-2 - 7E-5 - 3E-2 - 7E-5 - 3E-2 - 7E-5 - 3E-2 - 7E-5 - 3E-2 - 7E-5 - 7E-7 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-5 - 7E-7 -7E-7 -7 -7E-7 -7 -7E-7 -7 -7E	
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)		
		W, see ¹¹⁵ Sb	5E+3	2E+4	9E-6	3E-8	-	-	
1	Antimony-119	D, see ¹¹⁵ Sb W, see ¹¹⁵ Sb	2E+4 2E+4	5E+4 3E+4	2E-5 1E-5	6E-8 4E-8	2E-4	2E-3	
1	Antimony-120 ² (16 min)	D, see ¹¹⁵ Sb	1E+5 St wall	4E+5	2E-4	6E-7	-		
	(10 min)	W, see ¹¹⁵ Sb	(2E+5)	- 5E+5	_ 2E-4	- 7E-7	2E-3		
51	Antimony-120 (5.76 d)	D, see ¹¹⁵ Sb W, see ¹¹⁵ Sb	1E+3 9E+2	2E+3 1E+3	9E-7 5E-7	3E-9 2E-9	1E-5		
1	Antimony-122	D, see ¹¹⁵ Sb	8E+2	2E+3	1E-6	3E-9	-	_	
		W, see ¹¹⁵ Sb	LLI wall (8E+2) 7E+2	_ 1E+3	- 4E-7	- 2E-9	1E-5 -		
1	Antimony-124m ²	D, see 115 Sb W, see 115 Sb	3E+5 2E+5	8E+5 6E+5	4E-4 2E-4	1E-6 8E-7	3E-3	3E-2	
1	Antimony-124	D, see 115 Sb W, see 115 Sb	6E+2 5E+2	9E+2 2E+2	4E-7 1E-7	1E-9 3E-10	<u>7</u> E-6	7E-5	
1	Antimony-125	D, see 115 Sb W, see 115 Sb	2E+3	2E+3 5E+2	1E-6 2E-7	3E-9 7E-10	3E-5		
1	Antimony-126m ²	D, see ¹¹⁵ Sb	5E+4 St wall	2E+5	8E-5	3E-7	-	-	
		W, see ¹¹⁵ Sb	(7E+4) -	- 2E+5	- 8E-5	- 3E-7	9E-4		
1	Antimony-126	D, see ¹¹⁵ Sb W, see ¹¹⁵ Sb	6E+2 5E+2	1E+3 5E+2	5E-7 2E-7	2E-9 7E-10	7E-6		
1	Antimony-127	D, see ¹¹⁵ Sb	8E+2 LLI wall	2E+3	9E-7	3E-9	-	-	
		W, see ¹¹⁵ Sb	(8E+2) 7E+2	- 9E+2	 4E-7	- 1E-9	1E-5 -		
1	Antimony-128 ² (10.4 min)	D, see ¹¹⁵ Sb	8E+4 St wall	4E+5	2E-4	5E-7	-		
	,	W, see ¹¹⁵ Sb	(1E+5) -	_ 4E+5	- 2E-4	- 6E~7	1E-3 -		
1	Antimony-128 (9.01 h)	D, see 115 Sb W, see 115 Sb	1E+3	4E+3 3E+3	2E-6 1E-6	6E-9 5E-9	2E-5	2E-4	

			Occu	Table I pational Valu	ies	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 4E-4 - - 2E-3 - - 1E-3 - 1E-4 - 4E-4 -
tomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 ion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentratior (µCi/ml)
1	Antimony-129	D, see 115 Sb W, see 115 Sb	3E+3	9E+3 9E+3	4E-6 4E-6	1E-8 1E-8	4E-5	4E-4
1	Antimony-130 ²	D, see 115 Sb W, see 115 Sb	2E+4	6E+4 8E+4	3E-5 3E-5	9E-8 1E-7	3E-4	3E~3 -
1	Antimony-131 ²	D, see ¹¹⁵ Sb	1E+4 Thyroid	2E+4 Thyroid	1E-5	-	-	-
		W, see ¹¹⁵ Sb	(2E+4) -	(4E+4) 2E+4 Thyroid	- 1E-5	6E-8	2E-4 -	2E-3
			-	(4E+4)	-	6E-8	~	-
2	Tellurium-116	D, all compounds except those given for W W, oxides, hydroxides,	8E+3	2E+4	9E-6	3E-8	1E-4	1E-3
		and nitrates	-	3E+4	1E-5	4E-8	-	-
2	Tellurium-121m	D, see ¹¹⁶ Te	5E+2 Bone surf	2E+2 Bone surf	8E-8	-	-	-
		W, see ¹¹⁶ Te	(7E+2) -	(4E+2) 4E+2	- 2E-7	5E-10 6E-10	1E-5	
2	Tellurium-121	D, see ¹¹⁶ Te W, see ¹¹⁶ Te	3E+3 -	4E+3 3E+3	2E-6 1E-6	6E-9 4E-9	4E~5 -	4E-4 -
2	Tellurium-123m	D, see ¹¹⁶ Te	6E+2	2E+2	9E-8	-	-	-
		W, see ¹¹⁶ Te	Bone surf (1E+3) -	Bone surf (5E+2) 5E+2	- 2E-7	8E-10 8E-10	1E-5 -	1E-4 -
2	Tellurium-123	D, see ¹¹⁶ Te	5E+2 Bone surf	2E+2 Bone surf	8E-8	-	-	-
		W, see ¹¹⁶ Te	(1E+3) -	(5E+2) 4E+2	- 2E-7	7E-10 -	2E-5 -	2E-4
			-	Bone surf (1E+3)	-	2E-9	-	-
2	Tellurium-125m	D, see ¹¹⁶ Te	1E+3 Bone surf	4E+2 Bone surf	2E-7	-	-	-
		W, see ¹¹⁶ Te	(1E+3)	(1E+3) 7E+2	- 3E-7	1E-9 1E-9	2E-5	2E-4
2	Tellurium-127m	D, see ¹¹⁶ Te	6E+2	3E+2 Bone_surf	1E-7	-	9E-6	9E-5
		W, see ¹¹⁶ Te	-	(4E+2) 3E+2	- 1E-7	6E-10 4E-10	Ξ	-
2	Tellurium-127	D, see ¹¹⁶ Te	7E+3	2E+4	9E-6	3E-8	1E-4	1E-3

			0001	Table I upational Val	lues	Efflu	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral Ingestion	Col. 2 Inhala	Col. 3	Col. 1	Col. 2	Monthly Average
Atomic No.	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentration (µCi/ml)
		W, see ¹¹⁶ Te	-	2E+4	7E-6	2E-8	-	-
2	Tellurium-129m	D, see 116 Te W, see 116 Te	5E+2 -	6E+2 2E+2	3E-7 1E-7	9E-10 3E-10	7E-6 -	7E-5 -
52	Tellurium-129 ²	D, see 116 Te W, see 116 Te	3E+4	6E+4 7E+4	3E-5 3E-5	9E-8 1E-7	4E-4	4E-3
2	Tellurium-131m	D, see ¹¹⁶ Te	3E+2	4E+2	2E-7	-	-	-
		W, see ¹¹⁶ Te	Thyroid (6E+2)	Thyroid (1E+3) 4E+2	_ 2E-7	2E-9	8E-6	8E-5
		.,	_	Thyroid (9E+2)	~	1E-9	_	_
2	Tellurium-131 ²	D, see ¹¹⁶ Te	3E+3	5E+3	2E-6	-	-	-
		W, see ¹¹⁶ Te	Thyroid (6E+3) -	Thyroid (1E+4) 5E+3	_ 2E-6	2E-8	8E-5	8E-4
		,		Thyroid (1E+4)	-	2E-8	-	_
2	Tellurium-132	D, see ¹¹⁶ Te	2E+2	2E+2	9E-8	-	-	-
		W, see ¹¹⁶ Te	Thyroid (7E+2)	Thyroid (8E+2) 2E+2	- 9E-8	1E-9	9E-6	9E-5
		W, SEE 1E	-	Thyroid (6E+2)	-	- 9E-10	-	-
2	Tellurium-133m ²	D, see ¹¹⁶ Te	3E+3	(0 <u>1</u> ,2) 5E+3	2E-6	-	_	_
			Thyroid (6E+3)	Thyroid (1E+4)		2E-8	9E-5	9E-4
		W, see ¹¹⁶ Te	-	5E+3 Thyroid	2E-6	_	-	-
2	Tellurium-133 ²	D, see ¹¹⁶ Te	- 1E+4 Thvroid	(1Ē+4) 2E+4 Thyroid	- 9E-6	2E-8 -		
		W, see ¹¹⁶ Te	(3E+4)	(6E+4) 2E+4	_ 9E-6	8E-8	$\frac{4E-4}{-4}$	4E-3
			-	Thyroid (6E+4)	-	8E-8	_	_
2	Tellurium-134 ²	D, see ¹¹⁶ Te	2E+4	2E+4	1E-5	-	-	-
			Thyroid	Thyroid (5E+4)	-	7E-8	3E-4	3E-3
		W, see ¹¹⁶ Te	-	2E+4 Thyroid	1E-5	-	-	-
			-	(5Ē+4)	-	7E-8	-	-

			Occi	Table I upational Val	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Releases to
NO.		······································	(µCi)	(µCi)	(µCi/ml)	(µCi/ml)	(µCi/ml)	(µCi/ml)
53	Iodine-120m ²	D, all compounds	1E+4 Thyroid (1E+4)	2E+4 -	9E-6 -	3E-8 -	- 2E-4	
53	Iodine-120 ²	D, all compounds	4E+3 Thyroid (8E+3)	9E+3 Thyroid (1E+4)	4E-6 -	- 2E-8	- 1E-4	-
53	Iodine-121	D, all compounds	1E+4 Thyroid (3E+4)	2E+4 Thyroid (5E+4)	8E-6	- 7E-8	- 4E-4	
53	Iodine-123	D, all compounds	3E+3 Thyroid (1E+4)	6E+3 Thyroid (2E+4)	3E-6 -	- 2E-8	- 1E-4	- 1E-3
53	Iodine-124	D, all compounds	5E+1 Thyroid (2E+2)	8E+1 Thyroid (3E+2)	3E-8 -	- 4E-10	- 2E-6	
53	Iodine-125	D, all compounds	4E+1 Thyroid (1E+2)	6E+1 Thyroid (2E+2)	3E-8 -	- 3E-10	- 2E-6	
3	Iodine-126	D, all compounds	2E+1 Thyroid (7E+1)	4E+1 Thyroid (1E+2)	1E-8	- 2E-10	- 1E-6	
53	Iodine-128 ²	D, all compounds	4E+4 St wall (6E+4)	1E+5	5E-5 -	2E-10 2E-7 -	- 8E-4	_
3	Iodine-129	D, all compounds	5E+0 Thyroid (2E+1)	9E+0 Thyroid (3E+1)	4E-9	- 4E-11	- 2E-7	
3	Iodine-130	D, all compounds	4E+2 Thyroid (1E+3)	7E+2 Thyroid (2E+3)	3E-7	- 3E-9	- 2E-5	
3	Iodine-131	D, all compounds	3E+1 Thyroid (9E+1)	5E+1 Thyroid (2E+2)	2E-8	- 2E-10	- 1E-6	- 1E-5
53	$Iodine-132m^2$	D, all compounds	4E+3 Thyroid (1E+4)	8E+3 Thyroid (2E+4)	4E-6	- 3E-8	- 1E-4	- 1E-3
53	Iodine-132	D, all compounds	4E+3 Thyroid	8E+3 Thyroid	3E-6	-	-	-

			Occu	Table I Ipational Val	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhala ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Monthly Average Concentration (µCi/ml) 1E-3 - 7E-5 - 4E-3 - 3E-4 - - - - - - - - - - - - - - - - - -	
								·	
			(9E+3)	(1E+4)	-	2E-8	1E-4	1E-3	
3	Iodine-133	D, all compounds	1E+2	3E+2	1E-7	_	-	-	
			Thyroid (5E+2)	Thyroid (9E+2)	-	1E-9	7E-6	7E-5	
3	Iodine-134 ²	D, all compounds	2E+4	5E+4	2E-5	6E-8	-	-	
			Thyroid (3E+4)	-	-		4E-4	4E-3	
3	Iodine-135	D, all compounds	8E+2	2E+3	7E-7	_	-	-	
			Thyroid (3E+3)	Thyroid (4E+3)	-	6E-9	3E-5	3E-4	
4	Xenon-120 ²	Submersion ¹	-	-	1E-5	4E-8	-	-	
4	Xenon-121 ²	Submersion ¹	-	-	2E-6	1E-8	-	-	
4	Xenon-122	Submersion ¹	-	-	7E-5	3E-7	-	-	
4	Xenon-123	Submersion ¹	~	-	6E-6	3E-8	-	-	
4	Xenon-125	Submersion ¹	-	-	2E-5	7E-8	-	-	
4	Xenon-127	Submersion ¹	-	-	1E-5	6E-8	-	-	
4	Xenon-129m	Submersion ¹	-		2E-4	9E-7	-	-	
4	Xenon-131m	Submersion ¹	-	-	4E-4	2E-6	-	-	
4	Xenon-133m	Submersion ¹	-	-	1E-4	6E-7	-	-	
4	Xenon-133	Submersion ¹	-	-	1E-4	5E-7	-	-	
4	Xenon-135m ²	Submersion ¹	-	-	9E-6	4E-8	-	-	
4	Xenon-135	Submersion ¹	-	-	1E-5	7E-8	-	-	
4	Xenon-138 ²	Submersion ¹	-	-	4E-6	2E-8	-	-	
5	Cesium-125 ²	D, all compounds	5E+4 St wall	1E+5	6E-5	2E-7	-	-	
			(9E+4)	-	-	-	1E-3	1E-2	
5	Cesium-127	D, all compounds	6E+4	9E+4	4E-5	1E-7	9E-4	9E-3	
5	Cesium-129	D, all compounds	2E+4	3E+4	1E-5	5E-8	3E-4	3E-3	
5	Cesium-130 ²	D, all compounds	6E+4	2E+5	8E-5	3E-7	-	-	

			Occu	Table I upational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly	
tomic No.	Radionuclide	Class	Indestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
			St wall (1E+5)	-	-	_	1E-3	1E-2	
5	Cesium-131	D, all compounds	2E+4	3E+4	1E-5	4E-8	3E-4	3E-3	
5	Cesium-132	D, all compounds	3E+3	4E+3	2E-6	6E-9	4E-5	4E-4	
5	Cesium-134m	D, all compounds	1E+5	1E+5	6E-5	2E-7	-	-	
			St wall (1E+5)	-	-	-	2E-3	2E-2	
5	Cesium-134	D, all compounds	7E+1	1E+2	4E-8	2E-10	9E-7	9E-6	
5	Cesium-135m ²	D, all compounds	1E+5	2E+5	8E-5	3E-7	1E-3	1E-2	
5	Cesium-135	D, all compounds	7E+2	1E+3	5E-7	2E-9	1E-5	1E-4	
5	Cesium-136	D, all compounds	4E+2	7E+2	3E-7	9E-10	6E-6	6E-5	
5	Cesium-137	D, all compounds	1E+2	2E+2	6E-8	2E-10	1E-6	1E-5	
5	Cesium-138 ²	D, all compounds	2E+4	6E+4	2E-5	8E-8	-	-	
			St wall (3E+4)	-	-	-	4E - 4	4E-3	
6	Barium-126 ²	D, all compounds	6E+3	2E+4	6E-6	2E-8	8E-5	8E-4	
6	Barium-128	D, all compounds	5E+2	2E+3	7E-7	2E-9	7E-6	7E-5	
6	$Barium-131m^2$	D, all compounds	4E+5 St wall	1E+6	6E-4	2E-6	-	-	
			(5E+5)	-	-	-	7E-3	7E-2	
6	Barium-131	D, all compounds	3E+3	8E+3	3E-6	1E-8	4E-5	4E-4	
6	Barium~133m	D, all compounds	2E+3 LLI wall	9E+3	4E-6	1E-8	-	-	
			(3E+3)	-	-	-	4E-5	4E-4	
6	Barium-133	D, all compounds	2E+3	7E+2	3E-7	9E-10	2E-5	2E-4	
6	Barium-135m	D, all compounds	3E+3	1E+4	5E-6	2E-8	4E-5	4E-4	
6	Barium-139 ²	D, all compounds	1E+4	3E+4	1E-5	4E-8	2E-4	2E-3	
6	Barium-140	D, all compounds	5E+2 LLI wall	1E+3	6E-7	2E-9	-	-	
			(6E+2)	-	-	-	8E-6	8E-5	

			Occi	Table I upational Va	lues	Tabl Efflu Concent	e II lent rations	Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
56	Barium-141 ²	D, all compounds	2E+4	7E+4	3E-5	1E-7	3E-4	3E-3	
6	Barium-142 ²	D, all compounds	5E+4	1E+5	6E-5	2E-7	7E-4	7E-3	
7	Lanthanum-131 ²	D, all compounds except those given for W W, oxides and hydroxides	5E+4 -	1E+5 2E+5	5E-5 7E-5	2E-7 2E-7	6E-4	6E-3	
57	Lanthanum-132	D, see ¹³¹ La W, see ¹³¹ La	3E+3 -	1E+4 1E+4	4E-6 5E-6	1E-8 2E-8	4E-5	4E-4	
7	Lanthanum-135	D, see ¹³¹ La W, see ¹³¹ La	4E+4 -	1E+5 9E+4	4E-5 4E-5	1E-7 1E-7	5E-4	5E-3	
7	Lanthanum-137	D, see ¹³¹ La	1E+4	6E+1	3E-8	-	2E-4	2E-3	
		W, see ¹³¹ La	-	Liver (7E+1) 3E+2 Liver	_ 1E-7	1E-10 -	-	- -	
			-	(3E+2)	-	4E-10	-	-	
7	Lanthanum-138	D, see $^{131}_{131}$ La W, see 131 La	9E+2 -	4E+0 1E+1	1E-9 6E-9	5E-12 2E-11	1E-5 -	1E-4	
7	Lanthanum-140	D, see ¹³¹ La W, see ¹³¹ La	6E+2 -	1E+3 1E+3	6E-7 5E-7	2E-9 2E-9	9E-6 -	9E-5 -	
7	Lanthanum-141	D, see ¹³¹ La W, see ¹³¹ La	4E+3 -	9E+3 1E+4	4E-6 5E-6	1E-8 2E-8	5E-5 -	5E-4	
7	Lanthanum-142 ²	D, see ¹³¹ La W, see ¹³¹ La	8E+3 -	2E+4 3E+4	9E-6 1E-5	3E-8 5E-8	$\frac{1}{2}E-4$	1E-3 -	
7	Lanthanum-143 ²	D, see ¹³¹ La	4E+4 St_wall	1E+5	4E-5	1E-7	-	-	
		W, see ¹³¹ La	(4E+4)	- 9E+4	- 4E-5	- 1E-7	5E-4 -	5E-3 -	
8	Cerium-134	W, all compounds except those given for Y	5E+2 LLI wall	7E+2	3E-7	1E-9	-	-	
			(6E+2)	-	-	-	8E-6	8E~5	

			0cc1	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	Col, 1	Col. 2	Monthly	
omic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentratio (µCi/ml)	
		Y, oxides, hydroxides, and fluorides	-	7E+2	3E-7	9E-10	-	-	
3	Cerium-135	W, see ¹³⁴ Ce Y, see ¹³⁴ Ce	2E+3 -	4E+3 4E+3	2E-6 1E-6	5E-9 5E-9	2E-5	2E-4 -	
3	Cerium-137m	W, see ¹³⁴ Ce	2E+3 LLI wall	4E+3	2E-6	6E-9	-	-	
		Y, see ¹³⁴ Ce	(2Ē+3)	- 4E+3	- 2E-6	- 5E-9	3E-5	3E-4	
3	Cerium-137	W, see 134 Ce Y, see 134 Ce	5E+4	1E+5 1E+5	6E-5 5E-5	2E-7 2E-7	7E-4	7E-3	
3	Cerium-139	W, see ¹³⁴ Ce Y, see ¹³⁴ Ce	5E+3	8E+2 7E+2	3E-7 3E-7	1E-9 9E-10	7E-5	7E-4	
3	Cerium-141	W, see ¹³⁴ Ce	2E+3 LLI wall	7E+2	3E-7	1E-9	-	-	
		Y, see ¹³⁴ Ce	(2E+3) -	- 6E+2	- 2E-7	- 8E-10	3E-5	3E-4	
3	Cerium-143	W, see ¹³⁴ Ce	1E+3 LLI wall	2E+3	8E-7	3E-9	-	-	
		Y, see ¹³⁴ Ce	(1E+3) -	- 2E+3	- 7E-7	- 2E-9	2E-5 -	2E-4	
3	Cerium-144	W, see ¹³⁴ Ce	2E+2 LLI wall	3E+1	1E-8	4E-11	-	-	
		Y, see ¹³⁴ Ce	(3Ē+2) -	- 1E+1	- 6E-9	_ 2E-11	3E-6 -	3E-5 -	
Ð	Praseodymium-1	36 ² those given for Y	W, all cor 5E+4	mpounds exc 2E+5	ept 1E-4	3E-7	-	-	
		Y	St wall (7E+4)	-	-	-	1E-3	1E-2	
		Y, oxides, hydroxides, carbides, and fluorides	-	2E+5	9E-5	3E-7	-	-	
Ð	Praseodymium-1	37 ² Y, see ¹³⁶ Pr	W, see ¹³⁶	?r 1E+5	4E+4 6E-5	2E+5 2E-7	6E-5 -	2E-7 5	
Ð	Praseodymium-1	38m Y, see ¹³⁶ Pr	W, see ¹³⁶ H	?r 4E+4	1E+4 2E-5	5E+4 6E-8	2E-5 -	8E-8 11	
Ð	Praseodymium-1	³⁹ Y, see ¹³⁶ Pr	W, see ¹³⁶ H	?r 1E+5	4E+4 5E-5	1E+5 2E-7	5E-5 -	2E-7 6	
Э	Praseodymium-1	.42m ²	W, see ¹³⁶ H	Pr	8E+4	2E+5	7E-5	2E-7 1	

	•		Occu	Table I pational Va	alues	Tabl Efflu Concent	e II lent rations	Table I Release Sewers	s to
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal ALI (µCi)	Col. 3 DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concenti (µCi/ml	ration
		Y, see ¹³⁶ Pr	-	1E+5	6E-5	2E-7	-	-	
59	Praseodymium-14	2 Y, see ¹³⁶ Pr	W, see ¹³⁶ P.	r 2E+3	1E+3 8E-7	2E+3 3E-9	9E-7	<u>3</u> E-9	1E-51E-4
59	Praseodymium-14	3	W, see ¹³⁶ P	r	9E+2	8E+2	3E-7	1E-9	
		Y, see ¹³⁶ Pr	LLI wall (1E+3) -	- 7E+2		- 9E-10	2E-5 -	2E-4 -	
59	Praseodymium-14	1 ²	W, see ¹³⁶ P St wall	r	3E+4	1E+5	5E-5	2E-7	
		Y, see ¹³⁶ Pr	(4E+4)	- 1E+5	- 5E-5	- 2E-7	6E-4	6E-3 -	
59	Praseodymium-14	5 Y, see ¹³⁶ Pr	W, see ¹³⁶ P: -	r 8E+3	3E+3 3E-6	9E+3 1E-8	4E-6	1E-8 -	4E-54E-4
59	Praseodymium-14	7 ²	W, see ¹³⁶ P: St wall	r	5E+4	2E+5	8E-5	3E-7	
		Y, see ¹³⁶ Pr	(8E+4) -	- 2E+5		- 3E-7	1E-3 -	1E-2 -	
60	Neodymium-136 ²	W, all compounds except those given for Y Y, oxides, hydroxides, carbides, and fluorides	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 ¹³⁶ Nd Y, see ¹³⁶ Nd	2E+3	6E+3 5E+3	3E-6 2E-6	9E-9 7E-9	3E-5	<u>3</u> E-4	
60	Neodymium-139m	W, see 136 Nd Y, see 136 Nd	5E+3 -	2E+4 1E+4	7E-6 6E-6	2E-8 2E-8	7E-5 .	7E-4	
60	Neodymium~139²	W, see 136 Nd Y, see 136 Nd	9E+4 -	3E+5 3E+5	1E-4 1E-4	5E-7 4E-7	1E-3 -	1E-2	
60	Neodymium-141	W, see 136 Nd Y, see 136 Nd	2E+5	7E+5 6E+5	3E-4 3E-4	1E-6 9E-7	2E-3	2E-2	

			Occu	Table I pational Valu	les	Tabl Efflu Concent	ent	Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
tomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalat ALI (µCi)	ion DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
0	Neodymium-147	W, see ¹³⁶ Nd	1E+3 LLI wall	9E+2	4E-7	1E-9	-	-	
		Y, see ¹³⁶ Nd	(1E+3) _	- 8E+2	_ 4E-7	_ 1E-9	2E-5 -	2E-4 -	
50	Neodymium-149 ²	W, see ¹³⁶ Nd Y, see ¹³⁶ Nd	1E+4 -	3E+4 2E+4	1E-5 1E-5	4E-8 3E-8	1E-4	1E-3 -	
50	Neodymium-151 ²	W, see 136 Nd Y, see 136 Nd	7E+4	2E+5 2E+5	8E-5 8E-5	3E-7 3E-7	9E-4	9E-3	
1	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	-	-	
1	Promethium-143	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	5E+3	6E+2 7E+2	2E-7 3E-7	8E-10 1E-9	7E-5 -	7E-4	
51	Promethium-144	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	1E+3 -	1E+2 1E+2	5E-8 5E-8	2E-10 2E-10	2E-5 -	2E-4	
51	Promethium-145	W, see ¹⁴¹ Pm	1E+4	2E+2	7E-8	-	1E-4	1E-3	
		Y, see ¹⁴¹ Pm	-	Bone surf (2E+2) 2E+2	- 8E-8	3E-10 3E-10	_	-	
1	Promethium-146	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	2E+3	5E+1 4E+1	2E-8 2E-8	7E-11 6E-11	2E-5	<u>2</u> E-4	
1	Promethium-147	W, see ¹⁴¹ Pm	4E+3	1E+2	5E-8	-	-	-	
		Y, see ¹⁴¹ Pm	LLI wall (5E+3) -	Bone surf (2E+2) 1E+2	- 6E-8	3E-10 2E-10	7E-5 -	7E-4	
1	Promethium-148m	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	7E+2 -	3E+2 3E+2	1E-7 1E-7	4E-10 5E-10	1E-5 _	1E-4 -	
1	Promethium-148	W, see ¹⁴¹ Pm	4E+2	5E+2	2E-7	8E-10	-	-	
		Y, see ¹⁴¹ Pm	LLI wall (5E+2) -	- 5E+2	- 2E-7	- 7E-10	7E-6	7E-5 -	

			Occu	Table I pational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 tion DAC (µCi/ml)	Col. 1	Col. 2 Water	Monthly Average Concentration
			(µCI)	(µст)	(µC1/m1)	(µCi/ml)	(µCi/ml)	(µCi/ml)
61	Promethium-149	W, see ¹⁴¹ Pm	1E+3 LLI wall	2E+3	8E-7	3E-9	-	-
		Y, see ¹⁴¹ Pm	<u>(</u> 1E+3)	_ 2E+3	- 8E-7	- 2E-9	2E-5 -	2E-4
61	Promethium-150	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	5E+3 -	2E+4 2E+4	8E-6 7E-6	3E-8 2E-8	7E-5 -	7E-4 -
61	Promethium-151	W, see ¹⁴¹ Pm Y, see ¹⁴¹ Pm	2E+3 _	4E+3 3E+3	1E-6 1E-6	5E-9 4E-9	2E-5	2E-4
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	2E+5	8E-5	2E-7	-	-
			(6E+4)	-	-	-	8E-4	8E-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	бЕ+3	5E+2	2E-7	7E-10	8E-5	8E-4
62	Samarium-146	W, all compounds	1E+1 Bone surf (3E+1)	4E-2 Bone surf (6E-2)	1E-11 -	- 9E-14	 3E-7	- 3E-6
62	Samarium-147	W, all compounds	2E+1	4E-2	2E-11	-	-	-
			Bone surf (3E+1)	Bone surf (7E-2)	-	1E-13	4E-7	4E-6
62	Samarium-151	W, all compounds	1E+4 LLI wall	1E+2 Bone surf	4E-8	-	-	-
			(1E+4)	(2E+2)	-	2E-10	2E-4	2E-3
62	Samarium-153	W, all compounds	2E+3	3E+3	1E-6	4E-9	-	-
			LLI wall (2E+3)	-	-	-	3E-5	3E-4
62	Samarium-155 ²	W, all compounds	6E+4	2E+5	9E-5	3E-7	-	-
			St wall (8E+4)	-	-	-	1E-3	1E-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
53	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

			Occu	Table I pational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
			<u>Col. 1</u>	Col. 2	Col. 3	Col. 1	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Oral Ingestion ALI (µCi)	Inhalation ALI DAC (µCi) (µCi/ml		Air (µCi/ml)	Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
53	Europium-148	W, all compounds	1E+3	4E+2	1E-7	5E-10	1E-5	1E-4	
53	Europium-149	W, all compounds	1E+4	3E+3	1E-6	4E-9	2E-4	2E-3	
53	Europium-150 (12.62 h)	W, all compounds	3E+3	8E+3	4E-6	1E-8	4E-5	4E-4	
53	Europium-150 (34.2 y)	W, all compounds	8E+2	2E+1	8E-9	3E-11	1E-5	1E-4	
53	Europium-152m	W, all compounds	3E+3	6E+3	3E-6	9E-9	4E-5	4E-4	
53	Europium-152	W, all compounds	8E+2	2E+1	1E-8	3E-11	1E-5	1E-4	
53	Europium-154	W, all compounds	5E+2	2E+1	8E-9	3E-11	7E-6	7E-5	
53	Europium-155	W, all compounds	4E+3	9E+1	4E-8	-	5E-5	5E-4	
			-	Bone surf (1E+2)	-	2E-10	-	-	
53	Europium-156	W, all compounds	6E+2	5E+2	2E-7	6E-10	8E-6	8E-5	
53	Europium-157	W, all compounds	2E+3	5E+3	2E-6	7E-9	3E-5	3E-4	
53	Europium-158 ²	W, all compounds	2E+4	6E+4	2E-5	8E-8	3E-4	3E-3	
54	Gadolinium-145 ²	D, all compounds except those given for W	5E+4 St wall	2E+5	6E-5	2E-7	-	-	
		W, oxides, hydroxides,	(5E+4)	-	-	-	6E-4	6E-3	
		and fluorides	-	2E+5	7E-5	2E-7	-	-	
54	Gadolinium-146	D, see 145 Gd W, see 145 Gd	1E+3 -	1E+2 3E+2	5E-8 1E-7	2E-10 4E-10	2E-5 -	2E-4 -	
54	Gadolinium-147	D, see 145 Gd W, see 145 Gd	2E+3 -	4E+3 4E+3	2E-6 1E-6	6E-9 5E-9	3E-5 -	3E-4	
54	Gadolinium-148	D, see ¹⁴⁵ Gd	1E+1 Bone surf	8E+3 Bono gurf	3E-12	-	-	-	
		W, see ¹⁴⁵ Gđ	Bone surr (2E+1) -	Bone surf (2E-2) 3E-2 Bone surf	_ 1E-11	2E-14 -	3E-7 -	3E-6	
				(6E-2)	-	8E-14		-	
54	Gadolinium-149	D, see 145 Gd W, see 145 Gd	3E+3	2E+3 2E+3	9E-7 1E-6	3E-9 3E-9	4E-5 -	4E-4	

			Occu	Table I pational Valu	les	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat (µCi)	Col. 3 tion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
64	Gadolinium-151	D, see ¹⁴⁵ Gd	6E+3	4E+2	2E-7	-	9E-5	9E-4	
		W, see ¹⁴⁵ Gd	-	Bone surf (6E+2) 1E+3	- 5E-7	9E-10 2E-9	-	-	
54	Gadolinium-152	D, see ¹⁴⁵ Gd	2E+1	1E-2	4E-12	_	-	-	
		W, see ¹⁴⁵ Gd	Bone surf (3E+1) -	Bone surf (2E-2) 4E-2	_ 2E-11	3E-14	4E-7	4E-6	
			-	Bone surf (8E-2)	-	1E-13	-	-	
54	Gadolinium-153	D, see ¹⁴⁵ Gd	5E+3	1E+2	6E-8	-	6E-5	6E-4	
		W, see ¹⁴⁵ Gd	-	Bone surf (2E+2) 6E+2	_ 2E-7	3E-10 8E-10	-		
54	Gadolinium-159	D, see ¹⁴⁵ Gd W, see ¹⁴⁵ Gd	3E+3	8E+3 6E+3	3E-6 2E-6	1E-8 8E-9	4E-5	$\frac{4E-4}{-}$	
55	Terbium-147 ²	W, all compounds	9E+3	3E+4	1E-5	5E-8	1E-4	1E-3	
5	Terbium-149	W, all compounds	5E+3	7E+2	3E-7	1E-9	7E-5	7E-4	
55	Terbium-150	W, all compounds	5E+3	2E+4	9E-6	3E-8	7E-5	7E-4	
55	Terbium-151	W, all compounds	4E+3	9E+3	4E-6	1E-8	5E-5	5E-4	
55	Terbium-153	W, all compounds	5E+3	7E+3	3E-6	1E-8	7E-5	7E-4	
5	Terbium-154	W, all compounds	2E+3	4E+3	2E-6	6E-9	2E-5	2E-4	
5	Terbium-155	W, all compounds	6E+3	8E+3	3E-6	1E-8	8E-5	8E-4	
5	Terbium-156m (5.0 h)	W, all compounds	2E+4	3E+4	1E-5	4E-8	2E-4	2E-3	
5	Terbium-156m (24.4 h)	W, all compounds	7E+3	8E+3	3E-6	1E-8	1E-4	1E-3	
55	Terbium-156	W, all compounds	1E+3	1E+3	6E-7	2E-9	1E-5	1E-4	
5	Terbium-157	W, all compounds	5E+4	3E+2	1E-7	-	-	-	
			LLI wall (5E+4)	Bone surf (6E+2)	-	8E-10	7E-4	7E-3	
5	Terbium-158	W, all compounds	1E+3	2E+1	8E-9	3E-11	2E-5	2E-4	
5	Terbium-160	W, all compounds	8E+2	2E+2	9E-8	3E-10	1E-5	1E-4	

			Occi	Table I Ipational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalation ALI DAC (µCi) (µCi/ml)		Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
55	Terbium-161	W, all compounds	2E+3 LLI wall (2E+3)	2E+3 -	7E-7 -	2E-9 -	- 3E-5	- 3E-4	
56	Dysprosium-155	W, all compounds	9E+3	3E+4	1E-5	4E-8	1E-4	1E-3	
6	Dysprosium-157	W, all compounds	2E+4	6E+4	3E-5	9E-8	3E-4	3E-3	
56	Dysprosium-159	W, all compounds	1E+4	2E+3	1E~6	3E-9	2E-4	2E-3	
56	Dysprosium-165	W, all compounds	1E+4	5E+4	2E-5	6E-8	2E-4	2E-3	
56	Dysprosium-166	W, all compounds	6E+2	7E+2	3E-7	1E-9		-	
			LLI wall (8E+2)	-	-	_	1E-5	1E-4	
7	Holmium-155 ²	W, all compounds	4E+4	2E+5	6E-5	2E-7	6E-4	6E-3	
7	Holmium-157 ²	W, all compounds	3E+5	1E+6	6E-4	2E-6	4E-3	4E-2	
7	Holmium-159 ²	W, all compounds	2E+5	1E+6	4E~4	1E-6	3E-3	3E-2	
7	Holmium-161	W, all compounds	1E+5	4E+5	2E-4	6E-7	1E-3	1E-2	
57	Holmium-162m ²	W, all compounds	5E+4	3E+5	1E-4	4E-7	7E-4	7E-3	
57	Holmium-162 ²	W, all compounds	5E+5 St wall	2E+6	1E-3	3E-6	-	-	
			(8E+5)	-	-	-	1E-2	1E-1	
57	$Holmium-164m^2$	W, all compounds	1E+5	3E+5	1E-4	4E-7	1E-3	1E-2	
57	Holmium-164 ²	W, all compounds	2E+5 St wall	6E+5	3E-4	9E-7	- .	-	
			(2E+5)	-	-	-	3E-3	3E-2	
57	Holmium-166m	W, all compounds	6E+2	7E+0	3E-9	9E-12	9E-6	9E-5	
7	Holmium-166	W, all compounds	9E+2 LLI wall	2E+3	7E-7	2E-9	-	-	
			(9E+2)	-	-	-	1E-5	1E-4	
7	Holmium-167	W, all compounds	2E+4	6E+4	2E-5	8E-8	2E-4	2E-3	
8	Erbium-161	W, all compounds	2E+4	6E+4	3E-5	9E-8	2E-4	2E-3	
58	Erbium-165	W, all compounds	6E+4	2E+5	8E-5	3E-7	9E-4	9E-3	
8	Erbium-169	W, all compounds	3E+3	3E+3	1E-6	4E-9	-	-	

			Table I Occupational Values			Table II Effluent Concentrations		Table III Releases to Sewers	
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
							<u> </u>		
			LLI wall (4E+3)	-	-	-	5E-5	5E-4	
58	Erbium-171	W, all compounds	4E+3	1E+4	4E-6	1E-8	5E-5	5E-4	
58	Erbium-172	W, all compounds	1E+3	1E+3	6E-7	2E-9	-		
			LLI wall (1E+3)	-	-	-	2E-5	2E-4	
59	Thulium-162 ²	W, all compounds	7E+4	3E+5	1E-4	4E-7	-	-	
			St wall (7E+4)	-	-	-	1E-3	1E-2	
59	Thulium-166	W, all compounds	4E+3	1E+4	6E-6	2E-8	6E-5	6E-4	
59	Thulium-167	W, all compounds	2E+3 LLI wall	2E+3	8E-7	3E-9	-	-	
			(2E+3)	-	-	-	3E-5	3E-4	
59	Thulium-170	W, all compounds	8E+2 LLI wall	2E+2	9E-8	3E-10	-	-	
			(1E+3)	-	-	-	1E-5	1E-4	
59	Thulium-171	W, all compounds	1E+4 LLI wall	3E+2 Bone surf	1E-7		-	-	
			(1E+4)	(6E+2)	-	8E-10	2E-4	2E-3	
59	Thulium-172	W, all compounds	7E+2 LLI wall	1E+3	5E-7	2E-9	-	-	
			(8E+2)	-	-	-	1E-5	1E-4	
59	Thulium-173	W, all compounds	4E+3	1E+4	5E-6	2E-8	6E-5	6E-4	
59	Thulium-175 ²	W, all compounds	7E+4 St wall	3E+5	1E-4	4E-7	-	-	
			(9E+4)	-	-	-	1E-3	1E-2	
0	Ytterbium-162 ²	W, all compounds except those given for Y	7E+4	3E+5	1E-4	4E-7	1E-3	1E-2	
		Y, oxides, hydroxides, and fluorides	-	3E+5	1E-4	4E-7	-	-	
0	Ytterbium-166		1E+3				2E-5	2E-4	
		-	_	2E+3 2E+3	8E-7 8E-7	3E-9 3E-9		-	
0	Ytterbium-167 ²	W, see 162 Yb Y, see 162 Yb	3E+5 -	8E+5 7E+5	3E-4 3E-4	1E-6 1E-6	4E-3	4E-2	
0	Ytterbium-169	W, see 162 Yb	2E+3	8E+2	4E-7	1E-9	2E-5	2E-4	

			0001	Table I Ipational Valu	ies	Table II Effluent Concentrations		Table III Releases to Sewers	
comic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
		Y, see ¹⁶² Yb	-	7E+2	3E-7	1E-9	-	-	
0	Ytterbium-175	W, see ¹⁶² Yb	3E+3 LLI wall	4E+3	1E-6	5E-9	-	-	
		Y, see ¹⁶² Yb	(3E+3)	- 3E+3	- 1E-6	- 5E-9	4E~5 -	4E-4	
0	Ytterbium-177 ²	W, see 162 Yb Y, see 162 Yb	2E+4 _	5E+4 5E+4	2E-5 2E-5	7E-8 6E-8	2E-4	2E-3	
D	Ytterbium-178 ²	W, see 162 Yb Y, see 162 Yb	1E+4 -	4E+4 4E+4	2E-5 2E-5	6E-8 5E-8	2E-4	2E-3	
1	Lutetium-169	W, all compounds except those given for Y	3E+3	4E+3	2E-6	6E-9	3E-5	3E-4	
		Y, oxides, hydroxides, and fluorides	-	4E+3	2E-6	6E-9	-	-	
1	Lutetium-170	W, see 169 Lu Y, see 169 Lu	1E+3 -	2E+3 2E+3	9E-7 8E-7	3E-9 3E-9	2E-5	2E-4 -	
1	Lutetium-171	W, see ¹⁶⁹ Lu Y, see ¹⁶⁹ Lu	2E+3	2E+3 2E+3	8E-7 8E-7	3E-9 3E-9	3E-5 -	3E-4	
1	Lutetium-172	W, see ¹⁶⁹ Lu Y, see ¹⁶⁹ Lu	1E+3 -	1E+3 1E+3	5E-7 5E-7	2E-9 2E-9	1E-5 -	1E-4 -	
1	Lutetium-173	W, see ¹⁶⁹ Lu	5E+3	3E+2 Bone surf	1E-7	-	7E-5	7E-4	
		Y, see ¹⁶⁹ Lu	-	(5E+2) 3E+2	_ 1E-7	6E-10 4E-10	-	-	
1	Lutetium-174m	W, see ¹⁶⁹ Lu	2E+3 LLI wall	2E+2 Bone surf	1E-7	-	-	-	
		Y, see ¹⁶⁹ Lu	(3E+3)	(3E+2) 2E+2	- 9E-8	5E-10 3E-10	4E-5	4E-4	
1	Lutetium-174	W, see ¹⁶⁹ Lu	5E+3	1E+2	5E-8	-	7E-5	7E-4	
		Y, see ¹⁶⁹ Lu	-	Bone surf (2E+2) 2E+2	- 6E-8	3E-10 2E-10	-	-	
1	Lutetium-176m	W, see ¹⁶⁹ Lu Y, see ¹⁶⁹ Lu	8E+3 -	3E+4 2E+4	1E-5 9E-6	3E-8 3E-8	1E-4	1E-3	
1	Lutetium-176	W, see ¹⁶⁹ Lu	7E+2	5E+0	2E-9	-	1E-5	1E-4	
		Y, see ¹⁶⁹ Lu	-	Bone surf (1E+1) 8E+0	_ 3E-9	2E-11 1E-11	-	-	

			Table I Occupational Values			Table II Effluent Concentrations		Table III Releases to Sewers
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 tion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
71	Lutetium-177m	W, see ¹⁶⁹ Lu	7E+2	1E+2 Bone_surf	5E-8	-	1E-5	1E-4
		Y, see ¹⁶⁹ Lu	-	(1E+2) 8E+1	- 3E-8	2E-10 1E-10	-	-
1	Lutetium-177	W, see ¹⁶⁹ Lu	2E+3 LLI wall	2E+3	9E-7	3E-9		-
		Y, see ¹⁶⁹ Lu	(3Ē+3) -	_ 2E+3	- 9E-7	- 3E-9	4E-5	$\frac{4E-4}{-}$
1	$Lutetium-178m^2$	W, see ¹⁶⁹ Lu	5E+4 St. wall	2E+5	8E-5	3E-7	-	-
		Y, see ¹⁶⁹ Lu	(6±+4)	_ 2E+5	- 7E-5	_ 2E-7	8E-4	8E-3
1	Lutetium-178 ²	W, see ¹⁶⁹ Lu	4E+4 St wall	1E+5	5E-5	2E-7	-	-
		Y, see ¹⁶⁹ Lu	(4E+4)	- 1E+5	- 5E-5	- 2E-7	6E-4	6E-3
1	Lutetium-179	W, see ¹⁶⁹ Lu Y, see ¹⁶⁹ Lu	6E+3	2E+4 2E+4	8E-6 6E-6	3E-8 3E-8	9E-5	9E-4
2	Hafnium-170	D, all compounds except those given for W	3E+3	6E+3	2 <u>E</u> -6	8E-9	4E-5	4E-4
		W, oxides, hydroxides, carbides, and nitrates	-	5E+3	2E-6	6E-9	_	-
2	Hafnium-172	D, see ¹⁷⁰ Hf	1E+3	9E+0 Bone surf	4E-9	-	2E-5	2E-4
		W, see ¹⁷⁰ Hf	Ξ	(2E+1) 4E+1	- 2E-8	3E-11	-	-
			-	Bone surf (6E+1)	-	8E-11	-	-
2	Hafnium-173	D, see ¹⁷⁰ Hf W, see ¹⁷⁰ Hf	5E+3 -	1E+4 1E+4	5E-6 5E-6	2E-8 2E-8	7E-5	7E-4
2	Hafnium-175	D, see ¹⁷⁰ Hf	3E+3	9E+2 Bone surf	4E-7	-	4E-5	4E-4
		W, see ¹⁷⁰ Hf	-	(1E+3) 1E+3	- 5E-7	1E-9 2E-9	-	-
2	Hafnium-177m ²	D, see ¹⁷⁰ Hf W, see ¹⁷⁰ Hf	2E+4	6E+4 9E+4	2E-5 4E-5	8E-8 1E-7	3E-4	3E-3
2	Hafnium-178m	D, see ¹⁷⁰ Hf	3E+2	1E+0	5E-10	-	3E-6	3E-5
			-	Bone surf (2E+0)	-	3E-12	-	-

			Occu	Table I pational Valu	ıes	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
tomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
		W, see ¹⁷⁰ Hf	-	5E+0 Bone surf (9E+0)	2E-9 -	- 1E-11	-	-
2	Hafnium-179m	D, see ¹⁷⁰ Hf	1E+3	3E+2 Borno guraf	1E-7	-	1E-5	1E-4
		W, see ¹⁷⁰ Hf	-	Bone surf (6E+2) 6E+2	_ 3E-7	8E-10 8E-10	-	
2	Hafnium-180m	D, see $^{170}_{170}$ Hf W, see 170 Hf	7E+3	2E+4 3E+4	9E-6 1E-5	3E-8 4E-8	1E-4	1E-3 -
2	Hafnium-181	D, see ¹⁷⁰ Hf	1E+3	2E+2 Bone surf	7E-8	-	2E-5	2E-4
		W, see ¹⁷⁰ Hf	-	(4E+2) 4E+2	- 2E-7	6E-10 6E-10		
2	Hafnium-182m ²	D, see $^{170}_{170}$ Hf W, see 170 Hf	4E+4 -	9E+4 1E+5	4E-5 6E-5	1E-7 2E-7	5E-4	5E-3
2	Hafnium-182	D, see ¹⁷⁰ Hf	2E+2 Bone surf	8E-1 Bone surf	3E-10	-	-	-
		W, see ¹⁷⁰ Hf	(4E+2)	(2E+0) 3E+0	_ 1E-9	2E-12 -	5E-6 -	5E-5 -
				Bone surf (7E+0)	-	1E-11	-	
2	Hafnium-183 ²	D, see 170 Hf W, see 170 Hf	2E+4	5E+4 6E+4	2E-5 2E-5	6E-8 8E-8	3E-4	<u>3</u> E-3
2	Hafnium-184	D, see ¹⁷⁰ Hf W, see ¹⁷⁰ Hf	2E+3 -	8E+3 6E+3	3E-6 3E-6	1E-8 9E-9	3E-5	<u>3</u> E-4
3	Tantalum-172 ²	W, all compounds except those given for Y Y, elemental Ta, oxides, hydroxides, halides,	4E+4	1E+5	5E-5	2E-7	5E-4	5 E- 3
		carbides, nitrates, and nitrides	-	1E+5	4E-5	1E-7	-	-
3	Tantalum-173	W, see 172 Ta Y, see 172 Ta	7E+3	2E+4 2E+4	8E-6 7E-6	3E-8 2E-8	9E-5 -	9E-4
3	Tantalum-174 ²	W, see ${}^{172}_{172}$ Ta Y, see 172 Ta	3E+4	1E+5 9E+4	4E-5 4E-5	1E-7 1E-7	$\frac{4E-4}{-}$	4E-3
3	Tantalum-175	W, see 172 Ta Y, see 172 Ta	6E+3 -	2E+4 1E+4	7E-6 6E-6	2E-8 2E-8	8E-5 -	8E-4

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			Occu	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 5E-4 - 2E-3 - 3E-3 - 3E-3 - 3E-4 - 2E-4 - 3E-3 - 2E-4 - 3E-2 1E-4 - 2E-4 - 1E-2 - 1E-2
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhal: ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Averagê Concentration
73	Tantalum-176	W, see 172 Ta Y, see 172 Ta	4E+3	1E+4 1E+4	5E-6 5E-6	2E-8 2E-8	5E-5 -	5E-4 -
73	Tantalum-177	W, see ¹⁷² Ta Y, see ¹⁷² Ta	1E+4 _	2E+4 2E+4	8E-6 7E-6	3E-8 2E-8	2E-4	2E-3
73	Tantalum-178	W, see ¹⁷² Ta Y, see ¹⁷² Ta	2E+4	9E+4 7E+4	4E-5 3E-5	1E-7 1E-7	2E-4	2E-3
73	Tantalum-179	W, see ¹⁷² Ta Y, see ¹⁷² Ta	2E+4	5E+3 9E+2	2E-6 4E-7	8E-9 1E-9	3E-4	3E-3
73	Tantalum-180m	W, see ¹⁷² Ta Y, see ¹⁷² Ta	2E+4 -	7E+4 6E+4	3E-5 2E-5	9E-8 8E-8	3E-4	3E-3
'3	Tantalum-180	W, see ¹⁷² Ta Y, see ¹⁷² Ta	1E+3 -	4E+2 2E+1	2E-7 1E-8	6E-10 3E-11	2E-5 -	2E-4
3	Tantalum-182m ²	W, see ¹⁷² Ta	2E+5 St wall	5E+5	2E-4	8E-7	-	-
		Y, see ¹⁷² Ta	(2E+5)	- 4E+5	- 2E-4	- 6E-7	3E-3 -	3E-2
13	Tantalum-182	W, see ¹⁷² Ta Y, see ¹⁷² Ta	8E+2 -	3E+2 1E+2	1E-7 6E-8	5E-10 2E-10	1E-5 -	1E-4
3	Tantalum-183	W, see ¹⁷² Ta	9E+2 LLI wall	1E+3	5E-7	2E-9	-	-
		Y, see ¹⁷² Ta	(1E+3)	_ 1E+3	- 4E-7	_ 1E-9	2E-5	2E-4
3	Tantalum-184	W, see ¹⁷² Ta Y, see ¹⁷² Ta	2E+3	5E+3 5E+3	2E-6 2E-6	8E~9 7E-9	3E-5	3E-4
3	Tantalum-185 ²	W, see ¹⁷² Ta Y, see ¹⁷² Ta	3E+4	7E+4 6E+4	3E-5 3E-5	1E-7 9E-8	$\frac{4E-4}{-}$	
'3	Tantalum-186 ²	W, see ¹⁷² Ta	5E+4 St wall	2E+5	1E-4	3E-7	-	-
		Y, see ¹⁷² Ta	(7E+4)	_ 2E+5	- 9E-5	- 3E-7	1E-3	1E-2
4	Tungsten-176	D, all compounds	1E+4	5E+4	2E-5	7E-8	1E-4	1E-3
4	Tungsten-177	D, all compounds	2E+4	9E+4	4E-5	1E-7	3E-4	3E-3
4	Tungsten-178	D, all compounds	5E+3	2E+4	8E-6	3E~8	7E-5	7E-4
4	Tungsten-179 ²	D, all compounds	5E+5	2E+6	7E-4	2E-6	7E-3	7E-2

			0ccu	Table I pational Val	lues	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 2E-3 - 4E-4 3E-4 - 2E-5 - 2E-2 - 1E-2 7E-4 9E-4 - 3E-4 - 3E-4 - 3E-4 - 3E-4 -
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Releases to Sewers Monthly Average Concentration (µCi/ml) 2E-3 - 4E-4 3E-4 - 7E-5 - 2E-2 - 1E-2 - 7E-4 - 9E-4 - 2E-4 - 3E-4
No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhal; ALI (µCi)	ation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	
4	Tungsten-181	D, all compounds	2E+4	3E+4	1E-5	5E-8	2E-4	2E-3
4	Tungsten-185	D, all compounds	2E+3 LLI wall	7E+3	3E-6	9E-9	-	-
			(3E+3)	-	-	-	4E-5	4E-4
4	Tungsten-187	D, all compounds	2E+3	9E+3	4E-6	1E-8	3E-5	3E-4
4	Tungsten-188	D, all compounds	4E+2	1E+3	5E-7	2E-9	-	-
			LLI wall (5E+2)	-		-	7E-6	7E-5
5	Rhenium-177 ²	D, all compounds except those given for W	9E+4	3E+5	1E-4	4E-7	-	-
			St wall (1E+5)	-	-	-	2E-3	Average Concentration (µCi/ml) 2E-3 - 4E-4 3E-4 - 7E-5 - 2E-2 - 2E-2 - - 1E-2 - 7E-4 - 9E-4 - 2E-4 - 3E-4 - 3E-4 - 2E-4 - 2E-4 -
		W, oxides, hydroxides, and nitrates	-	4E+5	1E-4	5E-7	_	-
5	Rhenium-178 ²	D, see ¹⁷⁷ Re	7E+4	3E+5	1E-4	4E-7	_	-
		W, see ¹⁷⁷ Re	St wall (1E+5) -	- 3E+5	_ 1E-4	- 4E-7	1E-3	1E-2
5	Rhenium-181	D, see 177 Re W, see 177 Re	5E+3	9E+3 9E+3	4E-6	1E-8	7E-5	7E-4
			-		4E-6	1E-8	-	-
5	Rhenium-182 (12.7 h)	D, see ¹⁷⁷ Re W, see ¹⁷⁷ Re	7E+3 -	1E+4 2E+4	5E-6 6E-6	2E-8 2E-8	9E-5	9E-4
5	Rhenium-182 (64.0 h)	D, see ¹⁷⁷ Re W, see ¹⁷⁷ Re	1E+3 -	2E+3 2E+3	1E-6 9E-7	3E-9 3E-9	2E-5	2E-4
5	Rhenium-184m	D, see $^{177}_{177}$ Re W, see 177 Re	2E+3	3E+3 4E+2	1E-6 2E-7	4E-9 6E-10	3E-5	3E-4
5	Rhenium-184	D, see $^{177}_{177}$ Re W, see 177 Re	2E+3	4E+3 1E+3	1E-6 6E-7	5E-9 2E-9	3E-5	3E-4
5	Rhenium-186m	D, see 177 Re	1E+3	2E+3	7E-7	-	-	-
		W, see ¹⁷⁷ Re	St wall (2E+3)	St wall (2E+3) 2E+2	- 6E-8	3E-9 2E-10	2E-5	2E-4
5	Rhenium-186	W, See ¹⁷⁷ Re W, see ¹⁷⁷ Re	- 2E+3	3E+3	1E-6	4E-9 2E-9	- 3E-5	- 3E-4
			-	2E+3	7E-7	2E-9	-	-
5	Rhenium-187	D, see ¹⁷⁷ Re	6E+5	8E+5 St wall	4E - 4	-	8E-3	8E-2

			Occi	Table I upational Va	lues	Tabl Efflu Concent	e II lent trations	Table III Releases to Sewers Monthly Average Concentration (µC1/ml) - 1E-2 - 2E-4 - 4E-4 - 1E-2 -
Atomic	Radionuclide	Class	Col. 1 Oral Ingestion ALI	Col. 2 Inhal ALI	Col. 3 .ation DAC	Col. 1 Air	Col. 2 Water	Average
No.			(µĈi)	(µĈi)	DAC (µCi/ml)	(µĈi/ml)	(µCi/ml)	Concentration (µCi/ml)
		W, see ¹⁷⁷ Re	Ξ	(9E+5) 1E+5	- 4E-5	1E-6 1E-7	-	
75	Rhenium-188m ²	D, see ¹⁷⁷ Re W, see ¹⁷⁷ Re	8E+4	1E+5 1E+5	6E-5 6E-5	2E-7 2E-7	1E-3 -	1E-2 -
75	Rhenium-188	D, see ¹⁷⁷ Re W, see ¹⁷⁷ Re	2E+3 -	3E+3 3E+3	1E-6 1E-6	4E-9 4E-9	2E-5	2E-4
75	Rhenium-189	D, see ¹⁷⁷ Re W, see ¹⁷⁷ Re	3E+3 -	5E+3 4E+3	2E-6 2E-6	7E-9 6E-9	4E-5 -	
76	Osmium-180 ²	D, all compounds except those given for W and Y W, halides and nitrates Y, oxides and hydroxides	1E+5 - -	4E+5 5E+5 5E+5	2E-4 2E-4 2E-4	5E-7 7E-7 6E-7	1E-3 -	-
76	Osmium-181 ²	D, see ¹⁸⁰ Os W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	1E+4 - -	4E+4 5E+4 4E+4	2E-5 2E-5 2E-5	6E-8 6E-8 6E-8	2E-4 - -	-
76	Osmium-182	D, see ¹⁸⁰ Os W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	2E+3 	6E+3 4E+3 4E+3	2E-6 2E-6 2E-6	8E-9 6E-9 6E-9	3E-5 - -	-
76	Osmium-185	D, see ¹⁸⁰ Os W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	2E+3 	5E+2 8E+2 8E+2	2E-7 3E-7 3E-7	7E-10 1E-9 1E-9	3E-5 -	-
76	Osmium-189m	D, see ¹⁸⁰ Os W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	8E+4 -	2E+5 2E+5 2E+5	1E-4 9E-5 7E-5	3E-7 3E-7 2E-7	1E-3 -	-
76	Osmium-191m	D, see ¹⁸⁰ Os W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	1E+4 _ _	3E+4 2E+4 2E+4	1E-5 8E-6 7E-6	4E-8 3E-8 2E-8	2E-4 -	-
76	Osmium-191	D, see ¹⁸⁰ Os	2E+3 LLI wall	2E+3	9E-7	3E-9	-	-
		W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	(3E+3) _ _	- 2E+3 1E+3	- 7E-7 6E-7	- 2E-9 2E-9	3E-5 -	3E-4 -
76	Osmium-193	D, see ¹⁸⁰ Os	2E+3 LLI wall	5E+3	2E-6	6E-9	-	-
		W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	(2E+3) _ _	- 3E+3 3E+3	- 1E-6 1E-6	- 4E-9 4E-9	2E-5 - -	2E-4 -

		······································	Occi	Table I upational Va	lues	Tabl Efflu Concent	ent	Table III Releases to Sewers
tomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhala ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
6	Osmium-194	D, see ¹⁸⁰ Os	4E+2 LLI wall	4E+1	2E-8	6E-11	- 8E-6	- 8E-5
		W, see ¹⁸⁰ Os Y, see ¹⁸⁰ Os	(6E+2) - -	- 6E+1 8E+0	- 2E-8 3E-9	8E-11 1E-11	- -	- -
7	Iridium-182 ²	D, all compounds except those given for W and Y	4E+4 St wall	1E+5	6E-5	2E-7	-	
		W, halides, nitrates, and metallic iridium	(4E+4) -	- 2E+5 1E+5	- 6E-5 5E-5	- 2E-7 2E-7	6E-4	6E-3 -
7	Iridium-184	Y, oxides and hydroxides D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	- 8E+3 -	2E+4 3E+4	1E-5 1E-5	3E-8 5E-8	_ 1E-4	1E-3
7	Iridium-185	D, see $\frac{182}{182}$ Ir W, see $\frac{182}{182}$ Ir	 5E+3 	3E+4 1E+4 1E+4	1E-5 5E-6 5E-6	4E-8 2E-8 2E-8	- 7E-5 -	- 7E-4 -
7	Iridium-186	D, see $\frac{182}{182}$ Ir W, see $\frac{182}{102}$ Ir	- 2E+3 -	1E+4 8E+3 6E+3 6E+3	4E-6 3E-6 3E-6 2E-6	1E-8 1E-8 9E-9 8E-9	- 3E-5 -	- 3E-4 -
7	Iridium-187	Y, see ¹⁸² Ir D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	- 1E+4 -	3E+4 3E+4 3E+4 3E+4	1E-5 1E-5 1E-5	5E-8 4E-8 4E-8	1E-4 -	1E-3 _
7	Iridium-188	1, see 182 Ir W, see 182 Ir W, see 182 Ir Y, see 182 Ir	2E+3 	5E+3 4E+3 3E+3	2E-6 1E-6 1E-6	6E-9 5E-9 5E-9	3E-5 _	3E-4 _
7	Iridium-189	D, see ¹⁸² Ir	5E+3	5E+3	2E-6	7E-9	-	-
		W, see 102 Ir Y, see 102 Ir	LLI wall (5E+3) - -	- 4E+3 4E+3	- 2E-6 1E-6	- 5E-9 5E-9	7E-5 - -	7E-4
7	Iridium-190m ²	D, see 182 Ir W, see 182 Ir Y, see 182 Ir	2E+5 _ _	2E+5 2E+5 2E+5	8E-5 9E-5 8E-5	3E-7 3E-7 3E-7	2E-3 _	2E-2 ~
7	Iridium-190	D, see 162 Ir W, see 162 Ir Y, see 182 Ir	1E+3 _	9E+2 1E+3 9E+2	4E-7 4E-7 4E-7	1E-9 1E-9 1E-9	1E-5 -	1E-4 -
7	Iridium-192m	D, see 182 Ir	3E+3	9E+1	4E-8	1E-10	4E-5	4E-4

			Occi	Table I upational Val	lues	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) - 1E-4 - 9E-5 - 1E-4 - 2E-3 2E-3 2E-3 2E-4 1E-3 - 2E-3 2E-4 1E-3 - 2E-3 2E-4 1E-3 - 2E-3 2E-4 - 6E-3
	m - 31 1 / 3 -	-	Col. 1 Oral Ingestion ALI	Col. 2 Inhala	Col. 3 ation	<u>Col. 1</u>	Col. 2	Average
Atomic No.	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentration (µCi/ml)
		W, see ¹⁸² Ir Y, see ¹⁸² Ir	-	2E+2 2E+1	9E-8 6E-9	3E-10 2E-11	-	
77	Iridium-192	D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	9E+2 -	3E+2 4E+2 2E+2	1E-7 2E-7 9E-8	4E-10 6E-10 3E-10	1E-5 -	-
7	Iridium-194m	D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	6E+2 - -	9E+1 2E+2 1E+2	4E-8 7E-8 4E-8	1E-10 2E-10 1E-10	9E-6 - -	-
7	Iridium-194	D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	1E+3 _ _	3E+3 2E+3 2E+3	1E-6 9E-7 8E-7	4E-9 3E-9 3E-9	1E-5 	-
7	Iridium-195m	D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	8E+3 -	2E+4 3E+4 2E+4	1E-5 1E-5 9E-6	3E-8 4E-8 3E-8	1E-4 -	-
7	Iridium-195	D, see ¹⁸² Ir W, see ¹⁸² Ir Y, see ¹⁸² Ir	1E+4 -	4E+4 5E+4 4E+4	2E-5 2E-5 2E-5	6E-8 7E-8 6E-8	2E-4 	-
8	Platinum-186	D, all compounds	1E+4	4E+4	2E-5	5E-8	2E-4	2E-3
8	Platinum-188	D, all compounds	2E+3	2E+3	7E-7	2E-9	2E-5	2E-4
8	Platinum-189	D, all compounds	1E+4	3E+4	1E-5	4E-8	1E-4	1E-3
8	Platinum-191	D, all compounds	4E+3	8E+3	4E-6	1E-8	5E-5	5E-4
8	Platinum-193m	D, all compounds	3E+3 LLI wall	6E+3	3E-6	8E-9	-	-
			(3E+4)	-	-	-	4E~5	4E - 4
8	Platinum-193	D, all compounds	4E+4 LLI wall	2E+4	1E-5	3E-8	-	-
			(5E+4)	-	-	-	6E-4	6E-3
8	Platinum-195m	D, all compounds	2E+3 LLI wall	4E+3	2E-6	6E-9	-	-
	_		(2E+3)	-	-	-	3E-5	3E-4
8	Platinum-197m ²	D, all compounds	2E+4	4E+4	2E-5	6E-8	2E-4	2E-3
8	Platinum-197	D, all compounds	3E+3	1E+4	4E-6	1E-8	4E-5	4E-4
8	Platinum-199 ²	D, all compounds	5E+4	1E+5	6E-5	2E-7	7E-4	7E-3

			Occi	Table I pational Va	lues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 2E-4 1E-3 - 4E-4 - 1E-4 - 2E-4 - 4E-4 - - 4E-4 - - 4E-4 - - 4E-4 - - - 4E-4 - - - 4E-4 - - - -
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhal ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Averagē Concentration
		· · · · · · · · · · · · · · · · · · ·						
78	Platinum-200	D, all compounds	1E+3	3E+3	1E-6	5E-9	2E-5	2E-4
'9	Gold-193	D, all compounds except those given for W and Y W, halides and nitrates Y, oxides and hydroxides	9E+3 _ _	3E+4 2E+4 2E+4	1E-5 9E-6 8E-6	4E-8 3E-8 3E-8	1E-4 _	-
9	Gold-194	D, see ¹⁹³ Au W, see ¹⁹³ Au Y, see ¹⁹³ Au	3E+3 -	8E+3 5E+3 5E+3	3E-6 2E-6 2E-6	1E-8 8E-9 7E-9	4E-5 - -	-
9	Gold-195	D, see ¹⁹³ Au W, see ¹⁹³ Au Y, see ¹⁹³ Au	5E+3 -	1E+4 1E+3 4E+2	5E-6 6E-7 2E-7	2E-8 2E-9 6E-10	7E-5 - -	-
'9	Gold~198m	D, see ¹⁹³ Au W, see ¹⁹³ Au Y, see ¹⁹³ Au	1E+3 _	3E+3 1E+3 1E+3	1E-6 5E-7 5E-7	4E-9 2E-9 2E-9	1E-5 - -	-
'9	Gold-198	D, see ¹⁹³ Au W, see ¹⁹³ Au Y, see ¹⁹³ Au	1E+3 _	4E+3 2E+3 2E+3	2E-6 8E-7 7E-7	5E-9 3E-9 2E-9	2E-5 -	-
9	Gold-199	D, see ¹⁹³ Au	3E+3 LLI wall	9E+3	4E-6	1E-8	-	-
		W, see ¹⁹³ Au Y, see ¹⁹³ Au	(3E+3) -	- 4E+3 4E+3	- 2E-6 2E-6	- 6E-9 5E-9	4E-5 - -	_
9	Gold-200m	D, see 193 Au W, see 193 Au Y, see 193 Au	1E+3 _	4E+3 3E+3 2E+4	1E-6 1E-6 1E-6	5E-9 4E-9 3E-9	2E-5 -	-
9	Gold-200 ²	D, see ¹⁹³ Au W, see ¹⁹³ Au Y, see ¹⁹³ Au	3E+4 	6E+4 8E+4 7E+4	3E-5 3E-5 3E-5	9E-8 1E-7 1E-7	4E-4 	_
'9	Gold-201 ²	D, see ¹⁹³ Au	7E+4 St wall	2E+5	9E-5	3E-7	-	-
		W, see ¹⁹³ Au Y, see ¹⁹³ Au	(9E+4) - -	- 2E+5 2E+5	- 1E-4 9E-5	- 3E-7 3E-7	1E-3 -	1E-2 - -
30	Mercury-193m	Vapor Organic D D, sulfates W ovides budrovidos	- 4E+3 3E+3	8E+3 1E+4 9E+3	4E-6 5E-6 4E-6	1E-8 2E-8 1E-8	- 6E-5 4E-5	- 6E-4 4E-4
		W, oxides, hydroxides, halides, nitrates, and sulfides	-	8E+3	3E-6	1E-8	-	_

			Occi	Table I upational V	alues	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers Monthly Average Concentration (μ Ci/ml)
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inha ALI (µCi)	Col. 3 lation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	
30	Mercury-193	Vapor Organic D D, see ^{193m} Hg W, see ^{193m} Hg	- 2E+4 2E+4	3E+4 6E+4 4E+4 4E+4	1E-5 3E-5 2E-5 2E-5	4E-8 9E-8 6E-8 6E-8		2E-3
10	Mercury-194	Vapor Organic_D D, see ^{193m} Hg W, see ^{193m} Hg	- 2E+1 8E+2	3E+1 3E+1 4E+1 1E+2	1E-8 1E-8 2E-8 5E-8	4E-11 4E-11 6E-11 2E-10	- 2E~7 1E-5 -	1E-4
80	Mercury-195m	Vapor Organic_D D, see ^{193m} Hg W, see ^{193m} Hg	- 3E+3 2E+3 -	4E+3 6E+3 5E+3 4E+3	2E-6 3E-6 2E-6 2E-6	6E-9 8E-9 7E-9 5E-9	- 4E-5 3E-5 -	4E-4 3E-4
0	Mercury-195	Vapor Organic_D D, see ^{193m} Hg W, see ^{193m} Hg	- 2E+4 1E+4 -	3E+4 5E+4 4E+4 3E+4	1E-5 2E-5 1E-5 1E-5	4E-8 6E-8 5E-8 5E-8 5E-8	- 2E-4 2E-4	2E-3 2E-3
0	Mercury-197m	Vapor Organic_D D, see ^{193m} Hg W, see ^{193m} Hg	- 4E+3 3E+3 -	5E+3 9E+3 7E+3 5E+3	2E-6 4E~6 3E-6 2E-6	7E-9 1E-8 1E-8 7E-9	- 5E-5 4E-5 -	5E-4 4E-4
0	Mercury-197	Vapor Organic_D D, see ^{193m} Hg W, see ^{193m} Hg	- 7E+3 6E+3 -	8E+3 1E+4 1E+4 9E+3	4E-6 6E-6 5E-6 4E-6	1E-8 2E-8 2E-8 1E-8	- 9E-5 8E-5 -	8E-4
0	Mercury-199m ²	Vapor Organic D D, see ^{193m} Hg W, see ^{193m} Hg	- 6E+4 St wall (1E+5) 6E+4	8E+4 2E+5 - 1E+5 2E+5	3E-5 7E-5 - 6E-5 7E~5	1E-7 2E-7 - 2E-7 2E-7 2E-7	- - 1E-3 8E-4	- 1E-2
0	Mercury-203	W, see ^{193m} Hğ Vapor Organic D D, see ^{193m} Hg W, see ^{193m} Hg	- 5E+2 2E+3 -	2E+5 8E+2 8E+2 1E+3 1E+3	7E~5 4E-7 3E~7 5E-7 5E-7 5E-7	2E-7 1E-9 1E-9 2E-9 2E-9 2E-9	- 7E-6 3E-5	-
1	Thallium-194m ²	D, all compounds	5E+4 St wall (7E+4)	2E+5 -	6E~5	2E-7	- 1E-3	- 1E-2
1	Thallium-194 ²	D, all compounds	3E+5 St wall (3E+5)	6E+5 -	2E-4	8E-7 -	- 4E-3	- 4E-2

			Table I Occupational Values		les	Table II Effluent Concentrations		Table III Releases to Sewers Monthly Average Concentration (µCi/ml) 9E-3 1E-2 4E-3 3E-3 9E-3 1E-2 4E-3 3E-3 9E-3 1E-3 2E-4 8E-3 4E-4 1E-3 1E-3 2E-5 7E-4 5E-4 3E-3 4E-3 3E-3 4E-3 3E-3 4E-4 1E-3 2E-5 7E-4 5E-4 3E-3 2E-5 7E-4 5E-4 3E-3 2E-5 7E-4 5E-4 3E-3 -3 -4 5E-4 3E-3 -4
			$\frac{\text{Col. 1}}{\text{Col. 1}}$	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Releases to Sewers Monthly Average Concentration (µCi/ml) 9E-3 1E-2 4E-3 3E-3 9E-3 1E-3 2E-3 5E-4 2E-4 8E-3 4E-4 2E-4 8E-3 4E-3 3E-3 4E-4 1E-3 2E-5 7E-4 5E-4 3E-3
Atomic No.	Radionuclide	Class	Oral Ingestion ALI (µCi)	Inhalation ALI DAC (µCi) (µCi/ml)		Air (µCi/ml)	Water (µCi/ml)	Average Concentration
31	Thallium-195 ²	D, all compounds	6E+4	1E+5	5E-5	2E-7	9E-4	9E-3
31	Thallium-197	D, all compounds	7E+4	1E+5	5E-5	2E-7	1E-3	1E-2
31	Thallium-198 m^2	D, all compounds	3E+4	5E+4	2E-5	8E-8	4E-4	4E-3
31	Thallium-198	D, all compounds	2E+4	3E+4	1E-5	5E-8	3E-4	3E-3
31	Thallium-199	D, all compounds	6E+4	8E+4	4E-5	1E-7	9E-4	9E-3
31	Thallium-200	D, all compounds	8E+3	1E+4	5E-6	2E-8	1E-4	1E-3
31	Thallium-201	D, all compounds	2E+4	2E+4	9E-6	3E-8	2E-4	2E-3
81	Thallium-202	D, all compounds	4E+3	5E+3	2E-6	7E-9	5E-5	5E-4
1	Thallium-204	D, all compounds	2E+3	2E+3	9E-7	3E-9	2E-5	2E-4
2	Lead-195m ²	D, all compounds	6E+4	2E+5	8E-5	3E-7	8E-4	8E-3
2	Lead-198	D, all compounds	3E+4	6E+4	3E-5	9E-8	4E-4	4E-3
32	Lead-199 ²	D, all compounds	2E+4	7E+4	3E-5	1E-7	3E-4	3E-3
32	Lead-200	D, all compounds	3E+3	6E+3	3E-6	9E-9	4E-5	4E-4
32	Lead-201	D, all compounds	7E+3	2E+4	8E-6	3E-8	1E-4	1E-3
32	Lead-202m	D, all compounds	9E+3	3E+4	1E-5	4E-8	1E-4	1E-3
32	Lead-202	D, all compounds	1E+2	5E+1	2E-8	7E-11	2E-6	2E-5
32	Lead-203	D, all compounds	5E+3	9E+3	4E-6	1E-8	7E-5	7E-4
32	Lead-205	D, all compounds	4E+3	1E+3	6E-7	2E-9	5E-5	5E-4
32	Lead-209	D, all compounds	2E+4	6E+4	2E-5	8E-8	3E-4	3E-3
32	Lead-210	D, all compounds	6E-1	2E-1	1E-10	-	-	-
			Bone surf (1E+0)	Bone surf (4E-1)	-	6E-13	1E-8	1E-7
12	Lead-211 ²	D, all compounds	1E+4	6E+2	3E-7	9E-10	2E-4	2E-3
32	Lead-212	D, all compounds	8E+1	3E+1	1E-8	5E-11	-	-
			Bone surf (1E+2)	-	-	-	2E-6	2E-5
32	Lead-214 ²	D, all compounds	9E+3	8E+2	3E-7	1E-9	1E-4	1E-3

			Occi	Table I upational Va	lues	Table II Effluent Concentrations		Table III Releases to Sewers	
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhal ALI (µCi)	Col. 3 ation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)	
33	Bismuth-200 ²	D, nitrates W, all other compounds	3E+4 -	8E+4 1E+5	4E-5 4E-5	1E-7 1E-7	4E-4	4E-3	
33	Bismuth-201 ²	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	1E+4 -	3E+4 4E+4	1E-5 2E-5	4E-8 5E-8	2E-4	2E-3	
33	Bismuth-202 ²	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	1E+4 -	4E+4 8E+4	2E-5 3E-5	6E-8 1E-7	2E-4	2E-3	
33	Bismuth-203	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	2E+3 -	7E+3 6E+3	3E-6 3E-6	9E-9 9E-9	3E-5	3E-4	
33	Bismuth-205	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	1E+3 ~	3E+3 1E+3	1E-6 5E-7	3E-9 2E-9	2E-5	2E-4 -	
33	Bismuth-206	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	6E+2 -	1E+3 9E+2	6E-7 4E-7	2E-9 1E-9	9E-6	9E-5	
33	Bismuth-207	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	1E+3 -	2E+3 4E+2	7E-7 1E-7	2E-9 5E-10	1E-5 -	1E-4	
33	Bismuth-210m	D, see ²⁰⁰ Bi	4E+1 Kidneys	5E+0 Kidneys	2E-9	-	-	-	
		W, see ²⁰⁰ Bi	(6E+1) -	(6E+0) 7E-1	_ 3E-10	9E-12 9E-13	8E-7 -	8E-6	
33	Bismuth-210	D, see ²⁰⁰ Bi	8E+2	2E+2 Kidneys	1E-7	-	1E-5	1E-4	
		W, see ²⁰⁰ Bi	-	(4E+2) 3E+1	- 1E-8	5E-10 4E-11	-	-	
33	Bismuth-212 ²	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	5E+3	2E+2 3E+2	1E-7 1E-7	3E-10 4E-10	7E-5	7E-4	
33	Bismuth-213 ²	D, see ²⁰⁰ Bi W, see ²⁰⁰ Bi	7E+3	3E+2 4E+2	1E-7 1E-7	4E-10 5E-10	1E-4	1E-3 -	
33	Bismuth-214 ²	D, see ²⁰⁰ Bi	2E+4 St wall	8E+2	3E-7	1E-9	-	-	
		W, see ²⁰⁰ Bi	(2E+4) -	- 9E-2	- 4E-7	_ 1E-9	3E-4	3E-3	
4	Polonium-203 ²	D, all compounds except those given for W W, oxides, hydroxides,	3E+4	6E+4	3E-5	9E-8	3E-4	3E-3	
		and nitrates	-	9E+4	4E-5	1E-7	-	-	
4	Polonium-205 ²	D, see ²⁰³ Po W, see ²⁰³ Po	2E+4 -	4E+4 7E+4	2E-5 3E-5	5E-8 1E-7	3E-4	3E-3	

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			Occu	Table I pational Valu	ies	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Releases to
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalat ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	
84	Polonium-207	D, see ²⁰³ Po W, see ²⁰³ Po	8E+3 -	3E+4 3E+4	1E-5 1E-5	3E-8 4E-8	1E-4	1E-3 -
34	Polonium-210	D, see ²⁰³ Po W, see ²⁰³ Po	3E+0	6E-1 6E-1	3E-10 3E-10	9E-13 9E-13	4E-8 -	4E-7 -
85	Astatine-207 ²	D, halides W	6E+3	3E+3 2E+3	1E-6 9E-7	4E-9 3E-9	8E-5	8E-4
85	Astatine-211	D, halides W	1E+2 -	8E+1 5E+1	3E-8 2E-8	1E-10 8E-11	2E-6	2E-5 -
86	Radon-220	With daughters removed With daughters present	Ξ	2E+4 2E+1 (or 12 working level months)	7E-6 9E-9 (or 1.0 working level)	2E-8 3E-11	-	Ξ
36	Radon~222	With daughters removed With daughters present	-	1E+4 1E+2 (or 4 working level months)	4E-6 3E-8 (or 0.33 working level)	1E-8 1E-10	-	Ξ
87	Francium-222 ²	D, all compounds	2E+3	5E+2	2E-7	6E-10	3E-5	3E-4
37	Francium-223 ²	D, all compounds	6E+2	8E+2	3E-7	1E-9	8E-6	8E-5
88	Radium-223	W, all compounds	5E+0	7E-1	3E-10	9E-13	-	-
			Bone surf (9E+0)	-	-	- .	1E-7	1E-6
38	Radium-224	W, all compounds	8E+0 Bone surf	2E+0	7E-10	2E-12	-	-
			(2E+1)	-		-	2E-7	2E-6
38	Radium-225	W, all compounds	8E+0 Bone surf	7E-1	3E-10	9E-13	-	-
			(2E+1)	-	-	-	2E-7	2E-6
38	Radium-226	W, all compounds	2E+0 Bone surf	6E-1	3E-10	9E-13	-	-
		····	(5E+0)	-	-	-	6E-8	6E-7
38	Radium-227 ²	W, all compounds	2E+4 Bone surf	1E+4 Bone surf	6E-6	-	-	-
-	Ruttum 227	W, all composition	(2E+4)	(2E+4)	-	3E-8	3E-4	3E-3

			Occu	Table I pational Valu	les	Tabl Efflu Concent	e II lent rations	Table III Releases to Sewers Monthly Average Concentration (µCi/ml) - 6E-7 - 3E-4 - - 7E-6 - - - 2E-5 - - 5E-8 - - 3E-4 - - - 3E-4 - - - - - - - - - - - - - - - - - -
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	
								·····
88	Radium-228	W, all compounds	2E+0 Bone surf (4E+0)	1E+0 	5E-10 -	2E-12 -	- 6E-8	
89	Actinium-224	D, all compounds except those given for W and Y	2E+3 LLI wall	3E+1 Bone surf	1E-8	-	-	-
		W, halides and nitrates Y, oxides and hydroxides	(2E+3) 	(4E+1) 5E+1 5E+1	- 2E-8 2E-8	5E-11 7E-11 6E-11	3E-5 -	3E-4 _
89	Actinium-225	D, see ²²⁴ Ac	5E+1 LLI wall	3E-1 Bone surf	1E-10	-	-	-
		W, see 224 Ac Y, see 224 Ac	(5E+1) -	(5E-1) 6E-1 6E-1 6E-1	- 3E-10 3E-10	7E-13 9E-13 9E-13	7E-7 - -	-
89	Actinium-226	D, see ²²⁴ Ac	1E+2 LLI wall	3E+0 Bone surf	1E-9	-	-	-
		W, see 224 Ac Y, see 224 Ac	(1E+2) 	(4E+0) 5E+0 5E+0 5E+0	- 2E-9 2E-9	5E-12 7E-12 6E-12	2E~6 	-
89	Actinium-227	D, see ²²⁴ Ac	2E-1 Bone surf	4E-4 Bone surf	2E-13	-	-	-
		W, see ²²⁴ Ac	(4E-1)	(8E-4) 2E-3 Bone surf	- 7E-13	1E-15 _	5E-9 -	5E-8 -
		Y, see ²²⁴ Ac	-	(3E-3) 4E-3	_ 2E-12	4E-15 6E-15		
39	Actinium-228	D, see ²²⁴ Ac	2E+3	9E+0 Bone surf	4E-9	-	3E-5	3E-4
		W, see ²²⁴ Ac	-	(2E+1) 4E+1	- 2E-8	2E-11 -	-	-
		Y, see ²²⁴ Ac	-	Bone surf (6E+1) 4E+1	_ 2E-8	8E-11 6E-11	-	2
90	Thorium-226 ²	W, all compounds except those given for Y	5E+3 St wall	2E+2	6E-8	2E-10	-	-
		Y, oxides and hydroxides	(5E+3) -	_ 1E+2	- 6E-8	 2E-10	7E-5 -	7E-4
90	Thorium-227	W, see 226 Th Y, see 226 Th	1E+2 -	3E-1 3E-1	1E-10 1E-10	5E-13 5E-13	2E-6	2E-5
90	Thorium-228	W, see ²²⁶ Th	6E+0	1E-2	4E-12	-	-	-

			Occu	Table I pational Valu	165	Tabl Efflu Concent	e II ent rations	Table III Releases Sewers	to
Atomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 ion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentratio (µCi/ml)	tion
		Y, see ²²⁶ Th	Bone surf (1E+1)	Bone surf (2E-2) 2E-2	- 7E-12	3E-14 2E-14	2E-7	2E-6	
90	Thorium-229	W, see ²²⁶ Th	6E-1 Bone surf	9E-4 Bone surf	4E-13	-	-	-	
		Y, see ²²⁶ Th	(1E+0)	(2E-3) 2E-3	- 1E-12	3E-15	2E-8	2E-7	
		1, 500 111	-	Bone surf (3E-3)	_	4E-15		-	
90	Thorium-230	W, see ²²⁶ Th	4E+0	6E-3	3E-12	-	-	-	
		226m1	Bone surf (9E+0)	Bone surf (2E-2) 2E-2	- 6E-12	2E-14	1E-7	1E-6	
	·	Y, see ²²⁶ Th	-	Bone surf	0E-12 -	- 3E-14	-	-	
90	Thorium-231	W, see $^{226}_{226}$ Th Y, see 226 Th	4E+3	6E+3 6E+3	3E-6 3E-6	9E-9 9E-9	5E-5	5E~4 -	
90	Thorium-232	W, see ²²⁶ Th	7E-1	1E-3	5E-13	-	-	-	
		Y, see ²²⁶ Th	Bone surf (2E+0) -	Bone surf (3E-3) 3E-3	_ 1E-12	4E-15 -	3E-8	3E-7 -	
			-	Bone surf (4E-3)	-	6E-15	-	-	
90	Thorium-234	W, see ²²⁶ Th	3E+2 LLI wall	2E+2	8E-8	3E-10	-	-	
		Y, see ²²⁶ Th	(4E+2) -	- 2E+2	- 6E-8		5E-6 -	5E-5 ~	
91	Protactinium-2	27 ² those given for Y Y, oxides and hydroxides	W, all com 4E+3 -	npounds exce 1E+2 1E+2	pt 5E-8 4E-8	2E-10 1E-10	5E-5	5E-4	
91	Protactinium-2	28	W, see 227 P	a	1E+3	1E+1	5E-9	-	2E-52
		Y, see ²²⁷ Pa	-	Bone surf (2E+1) 1E+1	- 5E-9	3E-11 2E-11	-	Ξ	
91	Protactinium-2	30	W, see 227 P	'a	6E+2	5E+0	2E-9	7E-12	
		Y, see ²²⁷ Pa	Bone surf (9E+2)	_ 4E+0	- 1E-9	- 5E-12	1E-5 -	1E-4 -	
91	Protactinium-2		W, see ²²⁷ P Bone surf	a Bone surf	2E-1	2E-3	6E-13	-	

			Occupa	Table I ational Valu	165	Tabl Efflu Concent	e II ent rations	Table II Releases Sewers	
Atomic No.	Radionuclide	Class	Oral Ingestion	Col. 2 Inhalat ALI (µCi)	Col. 3 :ion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentr (µCi/ml)	ation
		Y, see ²²⁷ Pa	-	(4E-3) 4E-3 Bone surf	_ 2E-12	6E-15 -	6E-9	6E-8	
				(6E-3)	-	8E-15	-	-	
91	Protactinium-232	2	W, see ²²⁷ Pa	Dama aurof	1E+3	2E+1	9E-9	-	2E-52E-
		Y, see ²²⁷ Pa	-	Bone surf (6E+1) 6E+1 Bone surf	_ 2E-8	8E-11 -	-	-	
			-	(7E+1)	-	1E - 10	-	-	
91	Protactinium-233	3	W, see ²²⁷ Pa LLI wall		1E+3	7E+2	3E-7	1E-9	
		Y, see ²²⁷ Pa	(2E+3) -	- 6E+2	_ 2E-7	- 8E-10	2E-5	2E-4	
91	Protactinium-234	Y, see ²²⁷ Pa	W, see ²²⁷ Pa	7E+3	2E+3 3E-6	8E+3 9E-9	3E-6	1E-8	3E-53E~
2	Uranium-230	D, UF ₆ , UO ₂ F ₂ , UO ₂ (NO_3) ₂	Bone surf	4E-1 Bone_surf	2E-10	-	-	-	
		W, UO ₃ , UF ₄ , UCl ₄ Y, UO ₂ , U ₃ O ₈	(6E+0)	(6E-1) 4E-1 3E-1	- 1E-10 1E-10	8E-13 5E-13 4E-13	8E-8 - -	8E-7 	
2	Uranium-231	D, see ²³⁰ U	5E+3 LLI wall	8E+3	3E-6	1E-8	-	-	
		W, see $^{230}_{230}$ U Y, see 230 U	(4E+3)	- 6E+3 5E+3	- 2E-6 2E-6	- 8E-9 6E-9	6E-5	6E-4 - -	
2	Uranium-232	D, see ²³⁰ U		2E-1	9E-11	-	-		
		W, see 230 U Y, see 230 U	(4E+0)	Bone surf (4E-1) 4E-1 8E-3	- 2E-10 3E-12	6E-13 5E-13 1E-14	6E-8 -	6E-7 -	
92	Uranium-233	D, see ²³⁰ U		1E+0	5E-10		_	_	
		W, see 230 U Y, see 230 U	(2E+1)	Bone surf (2E+0) 7E-1 4E-2	- 3E-10 2E-11	3E-12 1E-12 5E-14	3E-7	3E-6	
2	Uranium-234 ³	D, see ²³⁰ U		1E+0	5E-10		_	_	
		W, see ²³⁰ U Y, see ²³⁰ U	Bone surf $(2E+1)$	Bone surf (2E+0) 7E-1 4E-2	- 3E-10 2E-11	3E-12 1E-12 5E-14	3E-7 _	3E-6 -	

			Occur	Table I Dational Valu	ies	Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers
tomic No.	Radionuclide	Class	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inhalat ALI (µCi)	Col. 3 Lion DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
2	Uranium-235 ³	D, see ²³⁰ U	1E+1 Bone surf	1E+0 Bone surf	6E-10	-	-	-
		W, see ²³⁰ U Y, see ²³⁰ U	(2E+1) -	(2E+0) 8E-1 4E-2	- 3E-10 2E-11	3E-12 1E-12 6E-14	3E-7 -	3E-6 -
2	Uranium-236	D, see ²³⁰ U	1E+1 Bone surf	1E+0 Bono gurf	5E-10	-	-	-
		W, see 230 U Y, see 230 U	(2E+1) -	Bone surf (2E+0) 8E-1 4E-2	- 3E-10 2E-11	3E-12 1E-12 6E-14	3E-7 - -	3E-6 - -
2	Uranium-237	D, see ²³⁰ U	2E+3	3E+3	1E-6	4E-9	-	-
		W, see 230 U Y, see 230 U	LLI wall (2E+3) - -	- 2E+3 2E+3	- 7E-7 6E-7	- 2E-9 2E-9	3E-5 -	3E-4 _
2	Uranium-238 ³	D, see ²³⁰ U	1E+1 Dono surf	1E+0 Bone surf	6E-10	-	-	-
		W, see ²³⁰ U Y, see ²³⁰ U	Bone surf (2E+1) - -	(2E+0) 8E-1 4E-2	- 3E-10 2E-11	3E-12 1E-12 6E-14	3E-7 -	3E-6 -
2	Uranium-239 ²	D, see $^{230}_{230}$ U W, see $^{230}_{230}$ U Y, see 230 U	7E+4 - -	2E+5 2E+5 2E+5	8E-5 7E-5 6E-5	3E-7 2E-7 2E-7	9E-4 - -	9E-3 -
2	Uranium-240	D, see $^{230}_{230}$ U W, see $^{230}_{230}$ U Y, see 230 U	1E+3 	4E+3 3E+3 2E+3	2E-6 1E-6 1E-6	5E-9 4E-9 3E-9	2E-5 -	2E-4 - -
2	Uranium-natural ³	·	D, see 230 U	1E+1	1E+0	5E-10	-	
		W, see ²³⁰ U Y, see ²³⁰ U	Bone surf (2E+1) -	Bone surf (2E+0) 8E-1 5E-2	- 3E-10 2E-11	3E-12 9E-13 9E-14	3E-7 -	3E-6 -
3	Neptunium-232 ²	W, all compounds	1E+5	2E+3 Bone surf	7E-7	-	2E-3	2E-2
				(5E+2)	-	6E-9	-	-
3	Neptunium-233 ²	W, all compounds	8E+5	3E+6	1E-3	4E-6	1E-2	1E-1
3	Neptunium-234	W, all compounds	2E+3	3E+3	1E-6	4E-9	3E-5	3E-4
3	Neptunium-235	W, all compounds	2E+4 LLI wall	8E+2 Bone surf	3E-7	-	-	-

			Occu	Table I pational Valu	les	Efflu	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly
Atomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalat ALI (µCi)	ion DAC (μCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
			(2E+4)	(1E+3)	-	2E-9	3E-4	3E-3
93	Neptunium-236 (1.15E+5 y)	W, all compounds	3E+0 Bone surf	2E-2 Bone surf	9E~12	-	-	-
	(T.T.2E+2 Å)		(6E+0)	(5E~2)	-	8E-14	9E-8	9E-7
93	Neptunium-236 (22.5 h)	W, all compounds	3E+3 Bone surf	3E+1 Bone surf	1E-8	-	-	-
	(22.5 11)		(4E+3)	(7E+1)	-	1E-10	5E-5	5E-4
93	Neptunium-237	W, all compounds	5E-1 Bone surf	4E-3 Bone surf	2E-12	-	-	-
			(1E+0)	(1E-2)	-	1E-14	2E-8	2E-7
)3	Neptunium-238	W, all compounds	1E+3	6E+1 Bone surf	3E-8	-	2E-5	2E-4
			-	(2E+2)	-	2E-10	-	-
93	Neptunium-239	W, all compounds	2E+3 LLI wall	2E+3	9E-7	3E-9	-	-
			(2E+3)	-	-	-	2E-5	2E-4
93	Neptunium-240 ²	W, all compounds	2E+4	8E+4	3E-5	1E-7	3E-4	3E-3
94	Plutonium-234	W, all compounds except PuO ₂ Y, PuO ₂	8E+3 -	2E+2 2E+2	9E-8 8E-8	3E-10 3E-10	1E-4	1E-3
94	Plutonium-235 ²	W, see ²³⁴ Pu Y, see ²³⁴ Pu	9E+5 -	3E+6 3E+6	1E-3 1E-3	4E-6 3E-6	1E-2	1E-1 -
94	Plutonium-236	W, see ²³⁴ Pu	2E+0	2E-2	8E-12	-		-
		Y, see ²³⁴ Pu	Bone surf (4E+0) -	Bone surf (4E-2) 4E-2	- 2E-11	5E-14 6E-14	6E-8	6E-7
94	Plutonium-237	W, see ²³⁴ Pu Y, see ²³⁴ Pu	1 E+4 -	3E+3 3E+3	1E-6 1E-6	5E-9 4E-9	2E-4	2E-3
94	Plutonium-238	W, see ²³⁴ Pu	9E-1 Bono curf	7E-3 Bone surf	3E-12		-	-
	Y, see ²³⁴ Pu -	(1E-2) 2E-2	- 8E-12	2E-14 2E-14	2E-8	2E-7		

			Occu	Table I pational Valu	les	Table Efflu Concent	e II ent rations	Table III Releases to Sewers
			Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly
tomic No.	Radionuclide	Class	Ingestion ALI (µCi)	Inhalat ALI (µCi)	ion DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
4	Plutonium-239	W, see ²³⁴ Pu	8E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-
		Y, see ²³⁴ Pu	(1E+0) -	(1E-2) 2E-2	- 7E-12	2E-14	2E-8	2E-7
			-	Bone surf (2E-2)	-	2E-14	-	-
4	Plutonium-240	W, see ²³⁴ Pu	8E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-
		Y, see ²³⁴ Pu	(1E+0)	(1E-2) 2E-2	_ 7E-12	2E-14 -	2E-8	2E-7
			-	Bone surf (2E-2)	-	2E-14		-
4	Plutonium-241	W, see ²³⁴ Pu	4E+1 Bone surf	3E-1 Bone surf	1E-10	-	-	-
		Y, see ²³⁴ Pu	(7E+1)	(6E-1) 8E-1		8E-13	1E-6	1E-5 -
			-	Bone surf (1E+0)	-	1E-12	-	-
4	Plutonium-242	W, see ²³⁴ Pu	8E-1 Bone surf	7E-3 Bone surf	3E-12	-	-	-
		Y, see ²³⁴ Pu	(1E+0) -	(1E-2) 2E-2	- 7E-12	2E-14	2E-8	2E-7
			-	Bone surf (2E-2)	-	2E-14	-	
4	Plutonium-243	W, see ²³⁴ Pu Y, see ²³⁴ Pu	2E+4 -	4E+4 4E+4	2E-5 2E-5	5E-8 5E-8	2E-4	2E-3
4	Plutonium-244	W, see ²³⁴ Pu	8E-1 Bone surf	7E-3 Bone surf	3E-12	-	-	-
		Y, see ²³⁴ Pu	(2E+0)	(1E-2) 2E-2	- 7E-12	2E-14 -	2E-8	2E-7 -
			-	Bone surf (2E-2)	-	2E-14	-	-
4	Plutonium-245	W, see ²³⁴ Pu Y, see ²³⁴ Pu	2E+3 -	5E+3 4E+3	2E-6 2E-6	6E-9 6E-9	3E-5	3E-4
4	Plutonium-246	W, see ²³⁴ Pu	4E+2 LLI wall	3E+2	1E-7	4E-10	-	-
		Y, see ²³⁴ Pu	(4E+2) -	- 3E+2	- 1E-7	- 4E-10	6E-6	6E-5 -
5	Americium-237 ²	W, all compounds	8E+4	3E+5	1E-4	4E-7	1E-3	1E-2

			Table I Occupational Values			Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
	D- 4/		Col. 1 Oral Ingestion ALI	Col. 2 Inhalat	Col. 3	<u>Col. 1</u>	Col. 2	Monthly Average Concentration	
NO.	Radionuclide	Class	ALI (µCi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Concentratio (µCi/ml)	
95	Americium-238 ²	W, all compounds	4E+4	3E+3	1E-6	_	5E-4	5E-3	
•	11.10220101.1 200		-	Bone surf (6E+3)	-	9E-9	-	-	
5	Americium-239	W, all compounds	5E+3	1E+4	5E-6	2E-8	7E-5	7E-4	
5	Americium-240	W, all compounds	2E+3	3E+3	1E-6	4E-9	3E-5	3E-4	
)5	Americium-241	W, all compounds	8E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-	
			(1E+0)	(1E-2)	-	2E-14	2E-8	2E-7	
5	Americium-242m	W, all compounds	8E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-	
			(1E+0)	(1E-2)	-	2E-14	2E-8	2E-7	
5	Americium-242	W, all compounds	4E+3	8E+1 Bone surf	4E-8	-	5E-5	5E-4	
		-	-	(9E+1)	-	1E-10	-	-	
5	Americium-243	W, all compounds	8E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-	
			(1E+0)	(1E-2)	-	2E-14	2E-8	2E-7	
5	Americium-244m ²	W, all compounds	6E+4 St wall	4E+3 Bone surf	2E-6	-	-	-	
_			(8E+4)	(7E+3)	-	1E-8	1E-3	1E-2	
5	Americium-244	W, all compounds	3E+3	2E+2 Bone_surf	8E-8	-	4E-5	4E - 4	
-	2	**	-	(3E+2)	-	4E-10	-	-	
5 5	Americium-245	W, all compounds	3E+4	8E+4	3E-5	1E-7	4E-4	4E-3	
5	Americium-246m ²	W, all compounds	5E+4 St wall	2E+5 -	8E-5	3E-7	-	~	
5	Americium-246 ²	W, all compounds	(6E+4) 3E+4	- 1E+5	- 4E-5	-	8E-4	8E-3	
6	Curium-238	W, all compounds	3E+4 2E+4	1E+5 1E+3	4E-5 5E-7	1E-7 2E-9	4E-4 2E-4	4E-3	
о б		· -	2E+4 6E+1	1E+3 6E-1	5E-7 2E-10	2E-9	2E-4	2E-3	
U I	Curium-240 W, all compounds	w, arr compounds	Bone surf (8E+1)	Bone surf $(6E-1)$	26-1V -	- 9E-13	- 1E-6	- 1E-5	
6	Curium-241	W, all compounds	(8E+1) 1E+3	(0E-1) 3E+1	- 1E-8		1E-0 2E-5	2E-4	
~	CALLON LIL	., all compounds	-	Bone surf (4E+1)		- 5E-11	2E-5	25-4	

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			Table I Occupational Values			Tabl Efflu Concent	e II ent rations	Table III Releases to Sewers	
			$\frac{\text{Col. 1}}{1}$	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Oral Ingestion ALI (µCi)	Inhalat ALI (µCi)	ion DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
6	Curium-242	W, all compounds	3E+1	3E-1	1E-10	-	-	-	
			Bone surf (5E+1)	Bone surf (3E-1)	-	4E-13	7E-7	7E-6	
6	Curium-243	W, all compounds	1E+0 Bone surf	9E-3 Bone surf	4E-12		-	-	
			(2E+0)	(2E-2)	-	2E-14	3E-8	3E-7	
6	Curium-244	W, all compounds	1E+0 Bone_surf	1E-2 Bone surf	5E-12	-	-	-	
			(3E+0)	(2E-2)	-	3E-14	3E-8	3E-7	
6	Curium-245	W, all compounds	7E-1 Bone surf	6E-3 Bone surf	3E-12	-	-	-	
		(1E+0)		(1E-2)	-	2E-14	2E-8	2E-7	
6	Curium-246	W, all compounds	7E-1 Bone_surf	6E-3 Bone surf	3E-12	-	-	-	
			(1E+0)	(1E-2)	-	2E-14	2E-8	2E-7	
6	Curium-247	W, all compounds	8E-1 Bone_surf	6E-3 Bone surf	3E-12		-	-	
			(1E+0)	(1E-2)	-	2E-14	2E-8	2E-7	
6	Curium-248	W, all compounds	2E-1 Bone surf	2E-3 Bone surf	7E-13	-	-	-	
			(4E-1)	(3E-3)	-	4E-15	5E-9	5E-8	
6	Curium-249 ²	W, all compounds	5E+4	2E+4 Bone surf	7E-6	-	7E-4	7E-3	
			-	(3E+4)	-	4E-8	-	-	
6	Curium-250	W, all compounds	4E-2 Bone surf	3E-4 Bone surf	1E-13		-	-	
				(6E-2)	(5E-4)	-	8E-16	9E-10	9E-9
7	Berkelium-245	W, all compounds	2E+3	1E+3	5E-7	2E-9	3E-5	3E-4	
7	Berkelium-246	W, all compounds	3E+3	3E+3	1E-6	4E-9	4E-5	4E-4	
7	Berkelium-247	W, all compounds	5E-1 Bone surf	4E-3 Bone surf	2E-12	-	-	-	
			(1E+0)	(9E-3)		1E - 14	2E-8	2E-7	
7	Berkelium-249	W, all compounds	2E+2 Bone surf	2E+0 Bone surf	7E-10	-	-	-	
,			(5E+2)	(4E+0)	-	5E-12	6E-6	6E-5	

			0cc1	Table I pational Valu	les	Efflu	e II ent rations	Table III Releases to Sewers
Atomic	Radionuclide	Class	Col. 1 Oral Ingestion ALI	Col. 2 Inhalat	Col. 3	Col. 1	Col. 2 Water	Monthly Average
No			(µĈi)	ALI (µCi)	DAC (µCi/ml)	Air (µCi/ml)	(µCi/ml)	Concentration (µCi/ml)
97	Berkelium-250	W, all compounds	9E+3 -	3E+2 Bone surf (7E+2)	1E-7	- 1E~9	1E-4 -	1E-3 -
98	Californium-244 ²	W, all compounds except those given for Y	3E+4 St wall	6E+2	2E-7	8E-10		-
		Y, oxides and hydroxides	(3E+4) -	- 6E+2	- 2E-7	- 8E-10	4E-4 -	4E-3
98	Californium-246	W, see ²⁴⁴ Cf Y, see ²⁴⁴ Cf	4E+2 -	9E+0 9E+0	4E-9 4E-9	1E-11 1E~11	5E-6	5E-5
98	Californium-248	W, see ²⁴⁴ Cf	8E+0 Bone surf	6E-2 Bone surf	3E-11	-	-	-
		Y, see ²⁴⁴ Cf	(2E+1) -	(1E-1) 1E-1	- 4E-11	2E-13 1E-13	2E-7 -	2E-6
98	Californium-249	W, see ²⁴⁴ Cf	5E-1 Bone_surf	4E-3 Bone_surf	2E-12	-	-	-
		Y, see ²⁴⁴ Cf	(1E+0) -	(9E-3) 1E-2 Bone_surf	- 4E-12	1E-14 -	2E-8 -	2E-7 -
98	Californium-250	W, see ²⁴⁴ Cf	- 1E+0	(1E-2) 9E-3	- 4E-12	2E-14	-	-
		Y, see ²⁴⁴ Cf	Bone surf (2E+0)	Bone surf (2E-2) 3E-2	 1E-11	3E-14 4E-14	3E-8	3E-7
98	Californium-251		5E-1	4E-3	2E-12		-	-
		Y, see ²⁴⁴ Cf	Bone surf (1E+0) -	Bone surf (9E-3) 1E-2	_ 4E-12	1E-14	2E-8	2E-7
			-	Bone surf (1E-2)	-	2E-14	-	-
98	Californium-252	W, see ²⁴⁴ Cf	2E+0 Bone_surf	2E-2 Bone_surf	8E-12	-	-	-
		Y, see ²⁴⁴ Cf	(5E+0) -	(4E-2) 3E-2	- 1E-11	5E-14 5E-14	7E-8 -	7E-7 -
98	Californium-253	W, see ²⁴⁴ Cf	2E+2 Bone surf (4E+2)	2E+0	8E-10	3E-12	-	-
		Y, see ²⁴⁴ Cf	(4些+ <i>2)</i> 一	- 2E+0	- 7E-10	- 2E-12	5E-6	5E-5
98	Californium-254	W, see ²⁴⁴ Cf Y, see ²⁴⁴ Cf	2E+0 -	2E~2 2E-2	9E-12 7E-12	3E-14 2E-14	3E-8	3E-7

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			Occu	Table I pational Valu	es	Tabl Efflu Concent	ent	Table III Releases to Sewers	
			Col. 1 Oral	Col. 2	Col. 3	<u>Col. 1</u>	Col. 2	Monthly	
Atomic No.	Radionuclide	Class	Ingestion Inhalat ALI ALI (µCi) (µCi)			Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)	
99	Einsteinium-250	W, all compounds	4E+4	5E+2 Bone surf	2E-7	-	6E-4	6E-3	
			-	(1E+3)	-	2E-9	-	-	
9	Einsteinium-251	W, all compounds	7E+3	9E+2	4E-7	-	1E-4	1E-3	
			-	Bone surf (1E+3)	-	2E-9	-	-	
9	Einsteinium-253	W, all compounds	2E+2	1E+0	6E-10	2E-12	2E-6	2E-5	
9	Einsteinium-254r	n	W,_all_com	pounds	3E+2	1E+1	4E-9	1E-11	
			LĹI wall (3E+2)	-	-	-	4E-6	4E-5	
9	Einsteinium-254	W, all compounds	8E+0	7E-2	3E-11	-	-	-	
			Bone surf (2E+1)	Bone surf (1E-1)	-	2E-13	2E-7	2E-6	
00	Fermium-252	W, all compounds	5E+2	1E+1	5E-9	2E-11	6E-6	6E-5	
.00	Fermium-253	W, all compounds	1E+3	1E+1	4E-9	1E-11	1E-5	1E-4	
.00	Fermium-254	W, all compounds	3E+3	9E+1	4E-8	1E-10	4E-5	4E-4	
.00	Fermium-255	W, all compounds	5E+2	2E+1	9E-9	3E-11	7E-6	7E-5	
.00	Fermium-257	W, all compounds	2E+1	2E-1	7E-11	-	-	-	
			Bone surf (4E+1)	Bone surf (2E-1)	-	3E-13	5E-7	5E-6	
.01	Mendelevium-257	W, all compounds	7E+3	8E+1	4E-8	-	1E-4	1E-3	
			-	Bone surf (9E+1)	-	1E-10	-	-	
01	Mendelevium-258	W, all compounds	3E+1	2E-1	1E-10	-	-	-	
			Bone surf (5E+1)	Bone surf (3E-1)	-	5E-13	6E-7	6E-6	
	Any single radio above with decay alpha emission of sion and with ra- life less than of	onuclide not listed y mode other than or spontaneous fis- adioactive half- 2 hours Submersion ¹	_	2E+2	1E-7	1E-9		_	

		Occ	Table I Occupational Values			e II ent rations	Table III Releases to Sewers
		Col. 1 Oral			<u>Col. 1</u>	Col. 2	Monthly
Atomic No.	Radionuclide Class	Ingestion ALI (µCi)		lation DAC (µCi/ml)	Air Water (µCi/ml) (µCi/ml)		Average Concentration (µCi/ml)
	Any single radionuclide not listed above with decay mode other than alpha emission or spontaneous fis- sion and with radioactive half- life greater than 2 hours	_	2E-1	1E-10	1E-12	1E-8	1E-7
•	Any single radionuclide not listed above that decays by alpha emission or spontaneous fission, or any mix- ture for which either the identity or the concentration of any radio-						

FOOTNOTES: "Submersion" means that values given are for submersion in a hemispherical semi-infinite cloud of airborne material.

²These radionuclides have radiological half-lives of less than two hours. The total effective dose equivalent received during operations with these radionuclides might include a significant contribution from external exposure. The derived air concentration values for all radionuclides, other than those designated class "submersion," are based upon the committed effective dose equivalent due to the intake of the radionuclide into the body and do NOT include potentially significant contributions to dose equivalent from external exposures. The licensee may substitute 1E-7 µCi/ml for the listed derived air concentration to account for the submersion dose prospectively, but should use individual monitoring devices or other radiation measuring instruments that measure external exposure to demonstrate compliance with the limits (see subsection 3 of section 33-10-04.1-06).

³For soluble mixtures of U-238, U-234, and U-235 in air, chemical toxicity may be the limiting factor (see subdivision e of subsection 1 of section 33-10-04.1.-06). If the percent by weight (enrichment) of U-235 is not greater than five, the concentration value for a forty-hour workweek is two tenths milligrams uranium per cubic meter of air average. For any enrichment, the product of the average concentration and time of exposure during a forty-hour workweek shall not exceed 8E-3 (SA) µCi-hr/ml, where SA is the specific activity of the uranium inhaled. The specific activity for natural uranium is 6.77E-7 curies per gram U. The specific activity for other mixtures of U-238, U-235, and U-234, if not known, shall be:

SA = 3.6E-7 curies/gram U U-depleted

 $SA = [0.4 + 0.38 \text{ (enrichment)} + 0.0034 \text{ (enrichment)}^2] E-6$, enrichment > 0.72

where enrichment is the percentage by weight of U-235, expressed as percent.

NOTE :

- 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 derived air concentration for the mixture shall be the most restrictive derived air concentration of any radionuclide in the mixture. 1.
- 2. If the identity of each radionuclide in the mixture is not known, but it is known that certain radionuclides specified in this appendix are not present in the mixture, the inhalation annual limit on intake, derived air concentration, and effluent and sewage concentrations for the mixture are the lowest values specified in this appendix for any radionuclide that is not known to be absent from the mixture; or

	Table I Efflu		Tab Releases	le II	Table III	
Occupational Values	EILIU	lenc	Releases	Concen	trations	Sewers
	Col. 1 Oral	Col. 2	Col. 3	Col. 1	Col. 2	Monthly
Atomic Radionuclide Class No.	Ingestion ALI (µCi)	Inh ALI (µCi)	alation DAC (µCi/ml)	Air (µCi/ml)	Water (µCi/ml)	Average Concentration (µCi/ml)
If it is known that Ac-227-D and Cm-250-W are not present	-	7E-4	3E-13	-	-	~
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	-	7E-3	3E-12	-	-	
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	_	7E-2	3E-11	_		-
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	-	7E-1	3E-10	_	-	-
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			7E+0	3E-9	_	
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 $-$	_	-		1E-14	_	-

	Occ	Table I cupational	Values	Effl	le II uent trations	Table III Releases to Sewers
Atomic Radionuclide Class No.	Col. 1 Oral Ingestion ALI (µCi)	Col. 2 Inh ALI (µCi)	Col. 3 alation DAC (µCi/ml)	Col. 1 Air (µCi/ml)	Col. 2 Water (µCi/ml)	Monthly Average Concentration (µCi/ml)
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	_	_	_	1E-13	_	-
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	-	-	-	1E-12	_	
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, 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	-	_	_	-1E-6	1E-5	

3. If a mixture of radionuclides consists of uranium and its daughters in ore dust (10 μm activity median aerodynamic diameter particle distribution assumed) prior to chemical separation of the uranium from the ore, the following values may be used for the derived air concentration of the mixture: 6E-11 μCi of gross alpha activity from uranium-238, uranium-234, thorium-230, and radium-226 per milliliter of air; 3E-11 μCi of natural uranium per milliliter of air; or forty-five micrograms of natural uranium per cubic meter of air.

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 this appendix 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 "one" (i.e., "unity").

Example: If radionuclides "A," "B," and "C" are present in concentrations C_{λ} , C_{B} , and C_{c} , and if the applicable derived air concentrations are DAC_{λ} , DAC_{B} , and DAC_{c} , respectively, then the concentrations shall be limited so that the following relationship exists:

$$\begin{array}{cccc} C_{A} & + & C_{B} & + & C_{C} \\ \hline DAC_{A} & & \overline{DAC_{B}} & & \overline{DAC_{C}} \end{array} \leq 1$$

4.

QUANTITIES1 OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

_		~	
Radionuclide	Quantity $(\mu Ci) *$	Radionuclide	Quantity $(\mu Ci) *$
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	100	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	
Chlorine-39	1,000	Cobalt-58	1,000 100
Argon-39	1,000	Cobalt-60m	1,000
Argon-41	1,000	Cobalt-60	1,000
Potassium-40	100	Cobalt-60 Cobalt-61	
Potassium-42	1,000	Cobalt-62m	1,000
Potassium-43	1,000	Nickel-56	1,000 100
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	
Scandium-44	100	Copper-64	1,000 1,000
Scandium-46	10	Copper-67	
Scandium-47	100	Zinc-62	1,000 100
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
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	
Germanium-78	1,000	Rubidium-87	100 100
Arsenic-69	1,000	Rubidium-87 Rubidium-88	
Arsenic-70	1,000	Rubidium-88 Rubidium-89	1,000
Arsenic-71		· · · ·	1,000
Arsenic-72	100 100	Strontium-80	100
AT SCHIC- / 2	TOO	Strontium-81	1,000

QUANTITIES1 OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

Radionuclide	Quantity	Radionuclide	Quantity
	(µCi)*		(µCi)*
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
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-110	
Technetium-99m	1,000	(69.1 min)	1,000
Technetium-99	100	Indium-110	
Technetium-101	1,000	(4.9 h)	1,000
Technetium-104	1,000	Indium-111	100
	·		

QUANTITIES ¹ OF LICENSED OR REGISTERED MATERIAL REQUIRING	OF LICENSED	OR	REGISTERED	MATERIAL	REOUIRING	LABELING

Radionuclide	Quantity (µCi)*	Radionuclide	Quantity $(\mu Ci) *$
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		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	1 000
Antimony-117 Antimony-118m	1,000 1,000	Iodine-128	1,000
Antimony-119	1,000	Iodine-129 Iodine-130	1 10
Antimony-120	1,000	Iodine-131	10
(16 min)	1,000	Iodine-131	100
Antimony-120	1,000	Iodine-132	100
(5.76 d)	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
(10.4 min)	1,000	Xenon-129m	1,000
Antimony-128		Xenon-131m	1,000
(9.01 h)	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

QUANTITIES1 OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

$\begin{array}{c ccccccccccccccccccccccccccccccccccc$	Radionuclide	Quantity	Radionuclide	Quantity
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	Radionaciide	$(\mu Ci) *$	Radionactive	
$\begin{array}{llllllllllllllllllllllllllllllllllll$	Barium-126	1,000	Promethium-141	
$\begin{array}{llllllllllllllllllllllllllllllllllll$		100		
$\begin{array}{c c c c c c c c c c c c c c c c c c c $				
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$\begin{array}{c} {\rm Cerium-134} & 100 & {\rm Samarium-156} & 1,000 \\ {\rm Cerium-137m} & 100 & {\rm Europium-145} & 100 \\ {\rm Cerium-137m} & 1,000 & {\rm Europium-146} & 100 \\ {\rm Cerium-137} & 1,000 & {\rm Europium-146} & 100 \\ {\rm Cerium-139} & 100 & {\rm Europium-148} & 100 \\ {\rm Cerium-141} & 100 & {\rm Europium-148} & 100 \\ {\rm Cerium-143} & 100 & {\rm Europium-150} & \\ {\rm Cerium-144} & 1 & (12.62 h) & 100 \\ {\rm Praseodymium-136} & 1,000 & {\rm Europium-150} & \\ {\rm Praseodymium-138m} & 1,000 & {\rm Europium-152} & 1 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-152} & 1 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 100 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143m} & 1,000 & {\rm Gadolinium-145} & 1,000 \\ {\rm Neodymium-136m} & 1,000 & {\rm Gadolinium-147m} & 100 \\ {\rm Neodymium-139m} & 1,000 & {\rm Gadolinium-148m} & 0.001 \\ {\rm Neodymium-139m} & 1,000 & {\rm Gadolinium-151m} & 10 \\ {\rm Neodymium-147m} & 100 & {\rm Gadolinium-151m} & 10 \\ {\rm Neodymium-147m} & 100 & {\rm Gadolinium-151m} & 10 \\ {\rm Neodymium-151m} & 1,000 & {\rm Gadolinium-151m} & 10 \\ {\rm Neodymium-151m} & 1,000 & {\rm Gadolinium-151m} & 10 \\ {\rm Neodymium-151m} & 1,000 & {\rm Ytterbium-166m} & 1,000 \\ {\rm Terbium-151m} & 1,000 & {\rm Ytterbium-167m} & 1,000 \\ {\rm Terbium-151m} & 1,000 & {\rm Ytterbium-177m} & 100 \\ {\rm Terbium-155m} & 1,000 & {\rm Ytterbium-177m} & 100 \\ {\rm Terbium-156m} & {\rm Iutetium-177m} & 100 \\ {\rm Terbium-156$			Samarium-155	
$\begin{array}{c} \mbox{Cerium-137m} & 100 & Europium-146 & 100 \\ \mbox{Cerium-137} & 1,000 & Europium-147 & 100 \\ \mbox{Cerium-141} & 100 & Europium-148 & 10 \\ \mbox{Cerium-141} & 100 & Europium-148 & 10 \\ \mbox{Cerium-143} & 100 & Europium-148 & 10 \\ \mbox{Cerium-144} & 1 & (12.62 h) & 100 \\ \mbox{Praseodymium-136} & 1,000 & Europium-150 \\ \mbox{Praseodymium-137m} & 1,000 & Europium-152 & 1 \\ \mbox{Praseodymium-139m} & 1,000 & Europium-152 & 1 \\ \mbox{Praseodymium-142m} & 1,000 & Europium-154 & 1 \\ \mbox{Praseodymium-142m} & 1,000 & Europium-155 & 10 \\ \mbox{Praseodymium-142m} & 1,000 & Europium-155 & 10 \\ \mbox{Praseodymium-143} & 100 & Europium-156 & 100 \\ \mbox{Praseodymium-144} & 1,000 & Europium-155 & 10 \\ \mbox{Praseodymium-145} & 1,000 & Europium-156 & 100 \\ \mbox{Praseodymium-145} & 1,000 & Europium-156 & 100 \\ \mbox{Praseodymium-145} & 1,000 & Europium-157 & 100 \\ \mbox{Praseodymium-145} & 1,000 & Gadolinium-146 & 10 \\ \mbox{Neodymium-138} & 100 & Gadolinium-148 & 0.001 \\ \mbox{Neodymium-139m} & 1,000 & Gadolinium-148 & 0.001 \\ \mbox{Neodymium-147} & 1,000 & Gadolinium-152 & 100 \\ \mbox{Neodymium-147} & 1,000 & Gadolinium-152 & 100 \\ \mbox{Neodymium-147} & 1,000 & Gadolinium-153 & 10 \\ \mbox{Neodymium-147} & 1,000 & Gadolinium-151 & 10 \\ \mbox{Neodymium-149} & 1,000 & Gadolinium-151 & 10 \\ \mbox{Neodymium-147} & 1,000 & Ytterbium-166 & 100 \\ \mbox{Terbium-151} & 1,000 & Ytterbium-167 & 1,000 \\ \mbox{Terbium-152} & 1,000 & Ytterbium-177 & 1,000 \\ \mbox{Terbium-155} & 1,000 & Ytterbium-177 & 1,000 \\ \mbox{Terbium-156} & 1,000 & Ytterbium-177 & 1,000 \\ \mbox{Terbium-156} & 100 & Ytterbium-177 & 1,000 \\ \mbox{Terbium-156} & 100 & Iutetium-173 & 10 \\ \mbox{Terbium-156} & 100 & Iutetium-173 & 10 \\ \mbox{Terbium-156} & 100 & Iutetium-173 & 10 \\ \mbox{Terbium-156} & 100 & Iutetium-174 & 10 \\ \mbox{Terbium-158} & 1 & 00 \\ \mbox{Terbium-156} & 100 & Iutetium-174 & 10 \\ \mbox{Terbium-158} & 1 & 00 \\ \mbo$	Cerium-134	100		
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$\begin{array}{cccc} {\rm Cerium-144} & 1 & (12.62 \ h) & 100 \\ {\rm Praseodymium-136} & 1,000 & {\rm Europium-150} \\ {\rm Praseodymium-137} & 1,000 & {\rm Europium-152} & 1 \\ {\rm Praseodymium-138m} & 1,000 & {\rm Europium-152m} & 100 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 1 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-142m} & 1,000 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-143} & 100 & {\rm Europium-155} & 10 \\ {\rm Praseodymium-144} & 1,000 & {\rm Europium-157} & 100 \\ {\rm Praseodymium-144} & 1,000 & {\rm Europium-157} & 100 \\ {\rm Praseodymium-145} & 100 & {\rm Europium-158} & 1,000 \\ {\rm Praseodymium-145} & 100 & {\rm Gadolinium-145} & 1,000 \\ {\rm Neodymium-138} & 100 & {\rm Gadolinium-147} & 100 \\ {\rm Neodymium-139m} & 1,000 & {\rm Gadolinium-147} & 100 \\ {\rm Neodymium-139m} & 1,000 & {\rm Gadolinium-151} & 10 \\ {\rm Neodymium-141} & 1,000 & {\rm Gadolinium-151} & 10 \\ {\rm Neodymium-147} & 100 & {\rm Gadolinium-151} & 10 \\ {\rm Neodymium-147} & 1,000 & {\rm Gadolinium-152} & 100 \\ {\rm Neodymium-147} & 1,000 & {\rm Gadolinium-159} & 100 \\ {\rm Terbium-151} & 1,000 & {\rm Gadolinium-159} & 100 \\ {\rm Terbium-151} & 1,000 & {\rm Ytterbium-162} & 1,000 \\ {\rm Terbium-153} & 1,000 & {\rm Ytterbium-167} & 1,000 \\ {\rm Terbium-153} & 1,000 & {\rm Ytterbium-177} & 1,000 \\ {\rm Terbium-154} & 100 & {\rm Ytterbium-177} & 1,000 \\ {\rm Terbium-155} & 1,000 & {\rm Ytterbium-177} & 1,000 \\ {\rm Terbium-156m} & {\rm Lutetium-170} & 100 \\ {\rm Terbium-156m} & {\rm Lutetium-171} & 100 \\ {\rm C24.4 \ h} & 1,000 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetium-174} & 10 \\ {\rm Terbium-156} & 10 & {\rm Lutetiu$				100
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QUANTITIES1 OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

Radionuclide	Quantity	Radionuclide	Quantity
	<u>(µCi)*</u>		<u>(µCi)*</u>
Dysprosium-155	1,000	Lutetium-177m	10
Dysprosium-157	1,000	Lutetium-177	100
Dysprosium-159	100	Lutetium-178m	1,000
Dysprosium-165	1,000	Lutetium-178	1,000
Dysprosium-166	100	Lutetium-179	1,000
Holmium-155	1,000	Hafnium-170	100
Holmium-157	1,000	Hafnium-172	1
Holmium-159	1,000	Hafnium-173	1,000
Holmium-161	1,000	Hafnium-175	100
Holmium-162m	1,000	Hafnium-177m	1,000
Holmium-162	1,000	Hafnium-178m	0.1
Holmium-164m	1,000	Hafnium-179m	10
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	
Tantalum-185	1,000	Iridium-190	1,000 100
Tantalum-186	1,000	Iridium-190	100
Tungsten-176	1,000		10
Tungsten-177	1,000	(1.4 min) Iridium-192	10
Tungsten-178	1,000	(73.8 d)	1
Tungsten-179	1,000	Iridium-194m	10
Tungsten-181	1,000	Iridium-194	100
Tungsten-185	100	Iridium-195m	
Tungsten-187	100	Iridium-195	1,000 1,000
Tungsten-188	10	Platinum-186	
Rhenium-177	1,000		1,000
Rhenium-178	1,000	Platinum-188	100
Rhenium-181		Platinum-189	1,000
	1,000	Platinum-191	100
Rhenium-182 (12.7 h)	1 000	Platinum-193m	100
	1,000	Platinum-193	1,000
Rhenium-182 $(64, 0, b)$	100	Platinum-195m	100
(64.0 h) Rhenium-184m	100	Platinum-197m	1,000
· · · · · · · · · · · · · · · · · · ·	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

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RadionuclideQuantityRadionuclideQuantity $(\mu')^*$ (ui)^*(ui)^*(ui)^*Osmium-1801,000Gold-199100Osmium-182100Gold-200100Osmium-182100Gold-2011,000Osmium-183100Gold-2011,000Osmium-1841,000Mercury-193100Osmium-1891,000Mercury-1951,000Osmium-1811,000Mercury-1951,000Osmium-1841Mercury-1951,000Iridium-1851,000Mercury-1971,000Iridium-186100Mercury-1971,000Iridium-186100Mercury-1971,000Iridium-1871,000Radium-2230.1Iridium-1861,000Radium-2230.1Iridium-1871,000Radium-2240.1Thallium-1941,000Radium-2250.1Thallium-1951,000Radium-2260.1Thallium-1961,000Radium-2271,000Thallium-2011,000Radium-2271,000Thallium-2011,000Radium-2270.001Thallium-203100Radium-2270.001Thallium-204100Actinium-2270.01Italium-2051,000Thorium-23610Lead-1981,000Thorium-2381Lead-1991,000Thorium-232100Lead-2051,000Thorium-233100Lead-2051,000 </th <th></th> <th></th> <th></th> <th><u> </u></th>				<u> </u>
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$\begin{array}{cccccccccccccccccccccccccccccccccccc$	Osmium-180		Gold-199	
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QUANTITIES¹ OF LICENSED OR REGISTERED MATERIAL REQUIRING LABELING

Radionuclide	Quantity (µCi)*	Radionuclide	Quantity (µCi)*
Neptunium-236		Curium-242	0.01
(22.5 h)	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 - I
Americium-244m	100	Einsteinium-254	0.01
Americium-244	10	Fermium-252	1
Americium-245	1,000	Fermium-253	ī
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		Any radionuclide	
radionuclide not		other than alpha-	
listed above or		emitting radionuclides	
mixtures of alpha		not listed above, or	
emitters of unknown		mixtures of beta	
composition	0.001	emitters of unknown	
COMPOSICION	0.001		0 01
		composition	0.01

NOTE: For purposes of subdivision e of subsection 2 of section 33-10-04.1.1-13, subdivision a of subsection 5 of section 33-10-04.1.1-13, and subdivision a of subsection 1 of section 33-10-04.1.1-16 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 radionuclides in the combination may not exceed "1" -- that is, unity.

¹The 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 Chapter 33-10-04.1.1, rounding to the nearest factor of 10, and constraining the values listed between 37 Bq and 37 MBq (0.001 and 1,000 μ Ci). Values of 3.7 MBq (100 μ Ci) have been assigned for radionuclides having a radioactive half-life in excess of E+9 years, except rhenium, 37 MBq (1,000 μ Ci), to take into account their low specific activity.

APPENDIX D (Reserved)

REQUIREMENTS FOR TRANSFER OF LOW-LEVEL RADIOACTIVE WASTE FOR DISPOSAL AT LAND DISPOSAL FACILITIES AND MANIFESTS

1. 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 United States 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 one-tenth percent 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 subsection 1 of appendix E 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 may be shipping papers used to meet United States department of transportation or United States environmental protection agency rules or requirements of the receiver, provided all the required information is included. Copies of manifests may be legible carbon copies or legible photocopies.

2. 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 rules of the United States department of transportation and the department. An authorized representative of the waste generator shall sign and date the manifest.

3. 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 paragraphs 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 paragraphs 4 through 8. A licensee shall:
- (1) Prepare all wastes so that the waste is classified according to subsection 1 of appendix E and meets the waste characteristics requirements in subsection 2 of appendix E;

- (2) Label each package of waste to identify whether it is class A waste, class B waste, or class C waste, in accordance with subsection 1 of appendix E;
- (3) Conduct a quality control program to ensure compliance with subsections 1 and 2 of appendix E; the program shall include management evaluation of audits;
 - (4) Prepare shipping manifests to meet the requirements of subsections 1 and 2;
 - (5) Forward a copy of the manifest to the intended recipient, 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 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 subsection 12 of section 33-10-03-05; 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 subdivision e.
- b. Any waste collector licensee who handles only prepackaged waste shall:
 - (1) Acknowledge receipt of the waste from the generator within one 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 subsection 1. 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 subsection 12 of section 33-10-03-05, and retain information from generator manifest until the license is terminated; 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 subdivision e.
- c. Any licensed waste processor who treats or repackages wastes shall:
 - (1) Acknowledge receipt of the waste from the generator within one week of receipt by returning a signed copy of the manifest or equivalent documentation;
 - (2) Prepare a new manifest that meets the requirements of subsections 1 and 2. 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 subsection 1 of appendix E and meets the waste characteristics requirements in subsection 2 of appendix E;
 - (4) Label each package of waste to identify whether it is class A waste, class B waste, or class C waste, in accordance with subsections 1 and 3 of appendix E;
 - (5) Conduct a quality control program to ensure compliance with subsections 1 and 2 of appendix E. 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 subsection 12 of section 33-10-03-05; 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 subdivision e.
- d. The land disposal facility operator shall:
 - (1) Acknowledge receipt of the waste within one 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 license is terminated; and
 - (3) Notify the shipper, that is, the generator, the collector, or processor, and the department when any shipment or portion of a shipment has not arrived within sixty 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 twenty days after transfer; and
- (2) Be traced and reported to whom. The 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 two weeks of completion of the investigation.

APPENDIX E CLASSIFICATION AND CHARACTERISTICS OF LOW-LEVEL RADIOACTIVE WASTE

- 1. Classification of radioactive waste for land disposal
 - Considerations. Determination of the classification of a. radioactive waste involves two considerations. First, consideration must be given to the concentration of longlived 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 subdivision a of subsection 2. If class A waste also meets the stability requirements set forth in subdivision b of subsection 2, 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 subsection 2.
 - (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 subsection 2.

- c. Classification determined by long-lived radionuclides. If the radioactive waste contains only radionuclides listed in table I, classification shall be determined as follows:
 - (1) If the concentration does not exceed one-tenth times the value in table I, the waste is class A.
 - (2) If the concentration exceeds one-tenth 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 subdivision g.

TABLE I

	Concentration	Concentration
Radionuclide	curie/cubic meter ^a	nanocurie/gram ^b

C-14	8	
C-14 in activated		
metal	80	
Ni-59 in activated		
metal	220	
Nb-94 in activated		
metal	0.2	
Тс-99	3	
I-129	0.08	
Alpha emitting transur	anic	
radionuclides with ha		
life greater than five	e	
years		100
Pu-241		3,500
Cm-242		20,000
Ra-226		100
		200

^aTo convert the curie per cubic meter values to gigabecquerel per cubic meter, multiply the curie per cubic meter value by thirty-seven.

^bTo convert the nanocurie per gram values to becquerel per gram, multiply the nanocurie per gram value by thirty-seven.

- 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 subdivision 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 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 subdivision g.

TABLE II

Radionuclide	Concentration, Column 1		cubic meter* Column 3
Total of all radio- nuclides 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 curie per cubic meter value to gigabecquerel per cubic meter, multiply the curie per cubic meter value by thirty-seven. There are no limits established for these radionuclides in class в class or С 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-lived and shortlived 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 one-tenth 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 one-tenth times the value listed in table I, but does not exceed the value in table I, 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.
- The sum of the g. 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 one if the waste class is to be determined by that column. Example: А waste contains strontium-90 in а of concentration and one eighty-five-hundredths terabecquerels per cubic meter (50 Ci/ m^3) and cesium-137 in concentration of eight а hundred fourteen

gigabecquerels per cubic meter (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 strontium-90 fraction, fifty divided by one hundred fifty is one-third, for cesium-137 fraction, twenty-two divided by forty-four is one-half; the sum of the fractions is eighty-three hundredths. Since the sum is less than one, the waste is class B.

- Determination of concentrations in wastes. The h. 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 be correlated with actual indirect methods can 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.
- 2. 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 chapter 33-10-04.1.1, 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 freestanding and noncorrosive liquid as is reasonably achievable, but in no case shall the liquid exceed one percent 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 paragraph 8.
- (7) Waste must not be pyrophoric material. Pyrophoric materials contained in wastes shall be treated, prepared, and packaged to be nonflammable. (See section 33-10-01-04 for the definition of pyrophoric.)
- (8) Wastes in a gaseous form shall be packaged at an absolute pressure that does not exceed one and onehalf atmospheres at twenty degrees Celsius. Total activity shall not exceed three and seven tenths terabecquerels (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 nonradiological 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.
 - Waste (1)shall have structural stability. А 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 paragraphs 3 and 4 of subdivision a of subsection 2, liquid wastes, or wastes containing liquid, shall be converted into a form that contains as little freestanding

and noncorrosive liquid as is reasonably achievable, but in no case shall the liquid exceed one percent of the volume of the waste when the waste is in a disposal container designed to ensure stability, or 0.5 percent 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.

3. 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 subsection 1.

APPENDIX F

Standards for Unrestricted Areas

(a) <u>Surface contamination limits</u>

(1) Alpha emitters

<u>0.555 Bq</u> 100 cm ²	=	<u>15 pCi</u> 100 cm ²	$= \frac{33 \text{ dpm}}{100 \text{ cm}^2}$	average over any one surface
<u>1.665 Bq</u> 100 cm ²	=	<u>45 pCi</u> 100 cm ²	$= \frac{100 \text{ dpm}}{100 \text{ cm}^2}$	maximum
<u>166.5 Bq</u> 100 cm ²	=	<u>450 pCi</u> 100 cm ²	$= \frac{1000 \text{ dpm}}{100 \text{ cm}^2}$	average over any one surface
<u>832.5 Bq</u> 100 cm ²	=	<u>2250 pCi</u> 100 cm ²	= <u>5000 dpm</u> 100 cm ²	maximum or
	<u>1.665 Bq</u> 100 cm ² <u>166.5 Bq</u> 100 cm ²	$\frac{1.665 \text{ Bq}}{100 \text{ cm}^2} = \frac{1.665 \text{ Bq}}{100 \text{ cm}^2} = \frac{166.5 \text{ Bq}}{100 \text{ cm}^2} = \frac{166.5 \text{ Bq}}{100 \text{ cm}^2}$	$\frac{1.665 \text{ Bg}}{100 \text{ cm}^2} = \frac{45 \text{ pCi}}{100 \text{ cm}^2}$ $\frac{166.5 \text{ Bg}}{100 \text{ cm}^2} = \frac{450 \text{ pCi}}{100 \text{ cm}^2}$	$\frac{1.665 \text{ Bq}}{100 \text{ cm}^2} = \frac{45 \text{ pCi}}{100 \text{ cm}^2} = \frac{100 \text{ dpm}}{100 \text{ cm}^2}$ $\frac{166.5 \text{ Bq}}{100 \text{ cm}^2} = \frac{450 \text{ pCi}}{100 \text{ cm}^2} = \frac{1000 \text{ dpm}}{100 \text{ cm}^2}$

 $\frac{2.5 \ \mu \text{Sv}}{\text{hr}} = \frac{(0.25 \text{ mrem})}{\text{hr}} \text{ maximum at 1 cm from surface}$

(2) Beta-Gamma emitters

<pre>(i) Removable: (all beta- gamma emitters except hydrogen-3)</pre>	<u>3.7 Bq</u> 100 cm ²	=	<u>100 pCi</u> 100 cm ²	average over any one surface
	<u>18.5 Bc</u> 100 cm ²	I	$= \frac{500 \text{ pCi}}{100 \text{ cm}^2}$	maximum
Removable: (hydrogen-3)	<u>37 Bq</u> 100 cm ²	=	<u>1000 pCi</u> 100 cm ²	average over any one surface
	<u>185 Bg</u> 100 cm ²	=	<u>5000 pCi</u> 100 cm ²	maximum

(ii) Total (fixed): $\frac{2.5 \ \mu Sv}{hr} = (\frac{0.25 \ mrem}{hr})$ maximum at 1 cm from surface

- (b) <u>Concentration in air and water</u>: appendix B, table II of chapter 33-10-04.1.1.
- (c) <u>Concentrations in soil and other materials except water</u>:
 - Radioactive material except source material and radium: Schedule A, column II of chapter 33-10-03.
 - (2) Source material and radium in soil: Concentration of radionuclides above background concentrations for total radium, averaged over areas of 100 square meters, shall not exceed:
 - (i) 5 picocuries per gram of dry soil, averaged over the first 15 centimeters below the surface; and
 - (ii) 5 picocuries per gram of dry soil, averaged over layers of 15 centimeters thickness more than 15 centimeters below the surface.
 - (3) Source material and radium in other materials: Concentration of radionuclides above background concentrations for total radium shall not exceed 5 picocuries per gram.
- (d) The level of gamma radiation measured at a distance of 100 centimeters from the surface shall not exceed background.

APPENDIX G REQUIREMENTS FOR TRANSFERS OF LOW-

LEVEL RADIOACTIVE WASTE INTENDED FOR DISPOSAL AT LICENSED LAND DISPOSAL FACILITIES AND MANIFESTS

I. Manifest

A waste generator, collector, or processor who transports, or offers for transportation, low-level radioactive waste intended for ultimate disposal at a licensed low-level radioactive waste land disposal facility must prepare a Manifest (Federal OMB Control Numbers 3150-0164, -0165, and -0166) reflecting information requested on applicable U.S. Nuclear Regulatory Commission (NRC) Forms 540 (Uniform Low-Level Radioactive Waste 541 (Uniform Low-Level and (Shipping Paper)) Manifest Radioactive Waste Manifest (Container and Waste Description)) and, if necessary, on an applicable NRC Form 542 (Uniform Low-Level Radioactive Waste Manifest (manifest Index and Regional Compact Tabulation)). NRC Forms 540 and 540A must be completed and must physically accompany the pertinent low-level waste shipment. Upon agreement between shipper and consignee, NRC Forms 541 and 541A and 542 and 542A may be completed, transmitted, and stored in electronic media with the capability for producing legible, accurate, and complete records on the respective forms. Licensees are not required by the department to comply with the manifesting requirements of this part chapter 33-10-04.1 when they ship:

- (a) LLW for processing and expect its return (i.e., for storage under their license) prior to disposal at a licensed land disposal facility;
- (b) LLW that is being returned to the licensee who is the "waste generator" or "generator," as defined in this part <u>appendix</u>; or
- (c) Radioactively contaminated material to a "waste processor" that becomes the processor's "residual waste."

For guidance in completing these forms, refer to the instructions that accompany the forms. Copies of manifests required by this appendix may be legible carbon copies, photocopies, or computer printouts that reproduce the data in the format of the uniform manifest. NRC Forms 540, 540A, 541, 541A, 542 and 542A, and the accompanying instructions, in hard copy, may be obtained from the Information and Records Management Branch, Office of Information Resources Management, U.S. Nuclear Regulatory Commission, Washington, DC 20555, telephone (301) 415-7232.

This appendix includes information requirements of the Department of Transportation, as codified in 49 CFR part 172. Information on hazardous, medical, or other waste, required to

meet Environmental Protection Agency regulations, as codified in 40 CFR parts 259, 261 or elsewhere, is not addressed in this section, and must be provided on the required EPA forms. However, the required EPA forms must accompany the Uniform Low-Level Radioactive Waste Manifest required by this chapter.

As used in this appendix, the following definitions apply:

Chelating agent has the same meaning as that given in Chapter 33-10-01.

Chemical description means a description of the principal chemical characteristics of a low-level radioactive waste.

Computer-readable medium means that the regulatory agency's computer can transfer the information from the medium into its memory and process the data.

Consignee means the designated receiver of the shipment of low-level radioactive waste.

Decontamination facility means a facility operating under a Commission or Agreement State license whose principal purpose is decontamination of equipment or materials to accomplish recycle, reuse, or other waste management objectives, and, for purposes of this <u>part chapter</u>, is not considered to be a consignee for LLW shipments.

Disposal container means a container principally used to confine low-level radioactive waste during disposal operations at a land disposal facility (also see "high integrity container"). Note that for some shipments, the disposal container may be the transport package.

EPA identification number means the number received by a transporter following application to the Administrator of EPA as required by 40 CFR part 263.

Generator means a licensee operating under a Commission or Agreement State license who (1) is a waste generator as defined in this chapter, or (2) is the licensee to whom waste can be attributed within the context of the Low-Level Radioactive Waste Policy Amendments Act of 1985 (e.g., waste generated as a result of decontamination or recycle activities).

High integrity container (HIC) means a container commonly designed to meet the structural stability requirements of the U.S. Nuclear Regulatory Commission Requirements in 10 CFR part 61 section 56 61.56, and to meet Department of Transportation requirements for a Type A package.

Land disposal facility means the land, buildings and structures, and equipment which are intended to be used for the disposal of radioactive waste. For purposes of this chapter, a "geologic repository" as defined in 10 CFR part 60 or 63 is not considered a "land disposal facility".

NRC Forms 540, 540A, 541, 541A, 542, and 542A are official NRC Forms referenced in this appendix. Licensees need not use originals of these NRC Forms as long as any substitute forms are equivalent to the original documentation in respect to content, clarity, size, and location of information. Upon agreement between the shipper and consignee, NRC Forms 541 (and 541A) and NRC Forms 542 (and 542A) may be completed, transmitted, and stored in electronic media. The electronic media must have the capability for producing legible, accurate, and complete records in the format of the uniform manifest.

Package has the same meaning as that given in Chapter 33-10-01 means the assembly of components necessary to ensure compliance with the packaging requirements of United States department of transportation regulations, together with its radioactive contents, as presented for transport.

Physical description means the items called for on NRC Form 541 to describe a low-level radioactive waste.

Residual waste means low-level radioactive waste resulting from processing or decontamination activities that cannot be easily separated into distinct batches attributable to specific waste generators. This waste is attributable to the processor or decontamination facility, as applicable.

Shipper means the licensed entity (i.e., the waste generator, waste collector, or waste processor) who offers low-level radioactive waste for transportation, typically consigning this type of waste to a licensed waste collector, waste processor, or land disposal facility operator.

Shipping paper means NRC Form 540 <u>(or equivalent)</u> and, if required, NRC Form 540A <u>(or equivalent)</u> which includes the information required by DOT in 49 CFR part 172.

Source material has the same meaning as that given in Chapter 33-10-01.

Special nuclear material has the same meaning as that given in Chapter 33-10-01.

Uniform Low-Level Radioactive Waste Manifest or uniform manifest means the combination of NRC Forms 540, 541, and, if necessary,

542, and their respective continuation sheets as needed, or equivalent.

Waste collector means an entity, operating under a Commission or Agreement State license, whose principal purpose is to collect and consolidate waste generated by others, and to transfer this waste, without processing or repackaging the collected waste, to another licensed waste collector, licensed waste processor, or licensed land disposal facility.

Waste description means the physical, chemical and radiological description of a low-level radioactive waste as called for on NRC Form 541.

Waste generator means an entity, operating under a Commission or Agreement State license, who (1) possesses any material or component that contains radioactivity or is radioactively contaminated for which the licensee foresees no further use, and (2) transfers this material or component to a licensed land disposal facility or to a licensed waste collector or processor for handling or treatment prior to disposal. A licensee performing processing or decontamination services may be a "waste generator" if the transfer of low-level radioactive waste from its facility is defined as "residual waste."

Waste processor means an entity, operating under a Commission or Agreement State license, whose principal purpose is to process, repackage, or otherwise treat low-level radioactive material or waste generated by others prior to eventual transfer of waste to a licensed low-level radioactive waste land disposal facility.

Waste type means a waste within a disposal container having a unique physical description (i.e., a specific waste descriptor code or description; or a waste sorbed on or solidified in a specifically defined media).

Information Requirements

A. General Information

The shipper of the radioactive waste, shall provide the following information on the uniform manifest:

- 1. The name, facility address, and telephone number of the licensee shipping the waste;
- 2. An explicit declaration indicating whether the shipper is acting as a waste generator, collector, processor, or a combination of these identifiers for purposes of the manifested shipment; and

- 3. The name, address, and telephone number, or the name and EPA identification number for the carrier transporting the waste.
- B. Shipment Information

The shipper of the radioactive waste shall provide the following information regarding the waste shipment on the uniform manifest:

- 1. The date of the waste shipment;
- 2. The total number of packages/disposal containers;
- 3. The total disposal volume and disposal weight in the shipment;
- 4. The total radionuclide activity in the shipment;
- 5. The activity of each of the radionuclides H-3, C-14, Tc-99, and I-129 contained in the shipment; and
- 6. The total masses of U-233, U-235, and plutonium in special nuclear material, and the total mass of uranium and thorium in source material.
- C. Disposal Container and Waste Information

The shipper of the radioactive waste shall provide the following information on the uniform manifest regarding the waste and each disposal container of waste in the shipment:

- 1. An alphabetic or numeric identification that uniquely identifies each disposal container in the shipment;
- 2. A physical description of the disposal container, including the manufacturer and model of any high integrity container;
- 3. The volume displaced by the disposal container;
- 4. The gross weight of the disposal container, including the waste;
- 5. For waste consigned to a disposal facility, the maximum radiation level at the surface of each disposal container;
- 6. A physical and chemical description of the waste;

- 7. The total weight percentage of chelating agent for any waste containing more than 0.1% chelating agent by weight, plus the identity of the principal chelating agent;
- 8. The approximate volume of waste within a container;
- The sorbing or solidification media, if any, and the identity of the solidification media vendor and brand name;
- 10. The identities and activities of individual radionuclides contained in each container, the masses of U-233, U-235, and plutonium in special nuclear material, and the masses of uranium and thorium in source material. For discrete waste types (i.e., activated materials, contaminated equipment, mechanical filters, sealed source/devices, and wastes in solidification/stabilization media), the identities and activities of individual radionuclides associated with or contained in these waste types within a disposal container shall be reported;
- 11. The total radioactivity within each container; and
- 12. For wastes consigned to a disposal facility, the classification of the waste pursuant to 10 CFR part 61 section 55 61.55. Waste not meeting the structural stability requirements of 10 CFR par 61 section 56 subsection (b) 61.56(b) must be identified.
- D. Uncontainerized Waste Information

The shipper of the radioactive waste shall provide the following information on the uniform manifest regarding a waste shipment delivered without a disposal container:

- 1. The approximate volume and weight of the waste;
- 2. A physical and chemical description of the waste;
- 3. The total weight percentage of chelating agent if the chelating agent exceeds 0.1% by weight, plus the identity of the principal chelating agent;
- 4. For waste consigned to a disposal facility, the classification of the waste pursuant to 10 CFR part 61 section 55 61.55. Waste not meeting the structural stability requirements of 10 CFR part 61 section 56 subsection (b) 61.56(b) must be identified;

- 5. The identities and activities of individual radionuclides contained in the waste, the masses of U-233, U-235, and plutonium in special nuclear material, and the masses of uranium and thorium in source material; and
- 6. For wastes consigned to a disposal facility, the maximum radiation levels at the surface of the waste.
- E. Multi-Generator Disposal Container Information

This section applies to disposal containers enclosing mixtures of waste originating from different generators. (Note: The origin of the LLW resulting from a processor's activities may be attributable to one or more "generators" (including "waste generators") as defined in this part <u>appendix</u>). It also applies to mixtures of wastes shipped in an uncontainerized form, for which portions of the mixture within the shipment originate from different generators.

- 1. For homogeneous mixtures of waste, such as incinerator ash, provide the waste description applicable to the mixture and the volume of the waste attributed to each generator.
- For heterogeneous mixtures of waste, such as the 2. combined products from a large compactor, identify each generator contributing waste to the disposal container, for discrete waste types (i.e., activated and, materials, contaminated equipment, mechanical filters, source/devices, and wastes in sealed solidification/stabilization media), the identities and activities of individual radionuclides contained in these waste types within the disposal container. For each generator, provide the following:
 - (a) The volume of waste within the disposal container;
 - (b) A physical and chemical description of the waste, including the solidification agent, if any;
 - (c) The total weight percentage of chelating agents for any disposal container containing more than 0.1% chelating agent by weight, plus the identity of the principal chelating agent;
 - (d) The sorbing or solidification media, if any, and the identity of the solidification media vendor and brand name if the media is claimed to meet stability requirements in 10 CFR part 61 section 56 subsection (b) 61.56(b); and

(e) Radionuclide identities and activities contained in the waste, the masses of U-233, U-235, and plutonium in special nuclear material, and the masses of uranium and thorium in source material if contained in the waste.

II. Certification

An authorized representative of the waste generator, processor, or collector shall certify by signing and dating the shipment manifest 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 Department of Transportation and the Commission. A collector in signing the certification is certifying that nothing has been done to the collected waste which would invalidate the waste generator's certification.

III. Control and Tracking

- $\pm \underline{A}$. Any licensee who transfers radioactive waste to a land disposal facility or a licensed waste collector shall comply with the requirements in paragraphs $\underline{A.1}$ $\underline{A.(a)}$ through $\underline{9}$ $\underline{A.(i)}$ of this section. Any licensee who transfers waste to a licensed waste processor for waste treatment or repackaging shall comply with the requirements of paragraphs $\underline{A.4}$ $\underline{A.(d)}$ through $\underline{9}$ $\underline{A.(i)}$ of this section. A licensee shall:-
 - (a) Prepare all wastes so that the waste is classified according to 10 CFR part 61 section 55 61.55 and meets the waste characteristics requirements in 10 CFR part 61 section 56 61.56.
 - (b) Label each disposal container (or transport package if potential radiation hazards preclude labeling of the individual disposal container) of waste to identify whether it is Class A waste, Class B waste, Class C waste, or greater then than Class C waste, in accordance with 10 CFR part 61 section 55 61.55;
 - (c) Conduct a quality assurance program to assure compliance with 10 CFR part 61 section 55 61.55 and 10 CFR part 61 section 56 61.56 (the program must include management evaluation of audits);
 - (d) Prepare the NRC Uniform Low-Level Radioactive Waste Manifest as required by this appendix;

- (e) Forward a copy or electronically transfer the Uniform Low-Level Radioactive Waste Manifest to the intended consignee so that either $(\underline{\tau_i})$ receipt of the manifest precedes the LLW shipment or (ii) the manifest is delivered to the consignee with the waste at the time the waste is transferred to the consignee. Using both $(\underline{\tau_i})$ and (ii) is also acceptable;
- (f) Include NRC Form 540 (and NRC Form 540A, if required) with the shipment regardless of the option chosen in paragraph A.5 <u>A.(e)</u> of this section;
- (g) Receive acknowledgment of the receipt of the shipment in the form of a signed copy of NRC Form 540;
- (h) Retain a copy of or electronically store the Uniform Low-Level Radioactive Waste Manifest and documentation of acknowledgment of receipt as the record of transfer of licensed material as required by 10 CFR Parts 30, 40, and 70; and
- (i) For any shipments or any part of a shipment for which acknowledgment of receipt has not been received within the times set forth in this appendix, conduct an investigation in accordance with paragraph E of this appendix.
- B. Any waste collector licensee who handles only prepackaged waste shall:
 - (a) Acknowledge receipt of the waste from the shipper within one week of receipt by returning a signed copy of NRC Form 540;
 - (b) Prepare a new manifest to reflect consolidated shipments that meet the requirements of this appendix. The waste collector shall ensure that, for each container of waste in the shipment, the manifest identifies the generator of that container of waste;
 - (c) Forward a copy or electronically transfer the Uniform Low-Level Radioactive Waste Manifest to the intended consignee so that either:
 - (1) Receipt of the manifest precedes the LLW shipment or

- (2) The manifest is delivered to the consignee with the waste at the time the waste is transferred to the consignee. Using both (± 1) and (± 2) is also acceptable;
- (d) Include NRC Form 540 (and NRC Form 540A, if required) with the shipment regardless of the option chosen in paragraph B.3 <u>B.(c)</u> of this section;
- (e) Receive acknowledgment of the receipt of the shipment in the form of a signed copy of NRC Form 540;
- (f) Retain a copy of or electronically store the Uniform Low-Level Radioactive Waste Manifest and documentation of acknowledgment of receipt as the record of transfer of licensed material as required by 10 CFR parts 30, 40, and 70;
- (g) For any shipments or any part of a shipment for which acknowledgment of receipt has not been received within the times set forth in this appendix, conduct an investigation in accordance with paragraph E of this appendix; and
- (h) Notify the shipper, the department, and the Administrator of the nearest Commission Regional Office when any shipment, or part of a shipment, has not arrived within 60 days after receipt of an advance manifest, unless notified by the shipper that the shipment has been canceled.
- C. Any licensed waste processor who treats or repackages waste shall:
 - (a) Acknowledge receipt of the waste from the shipper within one week of receipt by returning a signed copy of NRC Form 540;
 - (b) Prepare a new manifest that meets the requirements of this appendix. Preparation of the new manifest reflects that the processor is responsible for meeting these requirements. For each container of waste in the shipment, the manifest shall identify the waste generators, the preprocessed waste volume, and the other information as required in paragraph I.E. of this appendix;
 - (c) Prepare all wastes so that the waste is classified according to 10 CFR part 61 section 55 61.55 and

meets the waste characteristics requirements in 10 CFR part 61 section 56 61.56;

- (d) Label each package of waste to identify whether it is Class A waste, Class B waste, or Class C waste, in accordance with 10 CFR part 61 section 55 61.55 and 10 CFR part 61 section 57 61.57;
- (e) Conduct a quality assurance program to assure compliance with 10 CFR part 61 section 55 61.55 and 10 CFR part 61 section 56 61.56 (the program shall include management evaluation of audits);
- (f) Forward a copy or electronically transfer the Uniform Low-Level Radioactive Waste Manifest to the intended consignee so that either: $(\underline{\dagger i})$ Receipt of the manifest precedes the LLW shipment or (ii) the manifest is delivered to the consignee with the waste at the time the waste is transferred to the consignee. Using both $(\underline{\dagger i})$ and (ii) is also acceptable;
- (g) Include NRC Form 540 (and NRC Form 540A, if required) with the shipment regardless of the option chosen in paragraph C.6 <u>C.(f)</u> of this section;
- (h) Receive acknowledgment of the receipt of the shipment in the form of a signed copy of NRC Form 540;
- Retain a copy of or electronically store the Uniform Low-Level Radioactive Waste Manifest and documentation of acknowledgment of receipt as the record of transfer of licensed material as required by 10 CFR parts 30, 40, and 70;
- (j) For any shipment or any part of a shipment for which acknowledgment of receipt has not been received within the times set forth in this appendix, conduct an investigation in accordance with paragraph E of this appendix; and
- (k) Notify the shipper, the department, and the Administrator of the nearest Commission Regional Office when any shipment, or part of a shipment, has not arrived within 60 days after receipt of an advance manifest, unless notified by the shipper that the shipment has been canceled.
- D. The land disposal facility operator shall:

- (a) Acknowledge receipt of the waste within one week of receipt by returning, as a minimum, a signed copy of NRC Form 540 to the shipper. The shipper to be notified is the licensee who last possessed the waste and transferred the waste to the operator. If any discrepancy exists between materials listed on the Uniform Low-Level Radioactive Waste Manifest and materials received, copies or electronic transfer of the affected forms must be returned indicating the discrepancy;
- (b) Maintain copies of all completed manifests and electronically store the information required by 10 CFR part 61 section 80 subsection (1) 61.80(1) until the <u>department or</u> Commission terminates the license; and
- (c) Notify the shipper, the department, and the Administrator of the nearest Commission Regional Office when any shipment, or part of a shipment, has not arrived within 60 days after receipt of an advance manifest, unless notified by the shipper that the shipment has been canceled.
- E. Any shipment or part of a shipment for which acknowledgment is not received within the times set forth in this section must:
 - (a) Be investigated by the shipper if the shipper has not received notification or receipt within twenty days after transfer; and
 - (b) Be traced and reported. The investigation shall include tracing the shipment and filing a report with <u>the department and</u> the nearest commission regional office. Each licensee who conducts a trace investigation shall file a written report with <u>the department and</u> the appropriate nuclear regulatory commission regional office within two weeks of completion of the investigation.