

QUAD CITIES NUCLEAR POWER STATION

RADIOACTIVE WASTE AND ENVIRONMENTAL MONITORING

ANNUAL REPORT 1985

TELEDYNE ISOTOPES MIDWEST LABORATORY

NORTHBROOK, ILLINOIS

MARCH 1986

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INTRODUCTION

Units 1 and 2 of the Quad Cities Station located near Cordova, Illinois next to the Mississippi River, are 800 MWe boiling water reactors, similar in design to Dresden Units 2 and 3. The plant has been designed to keep releases to the environment at levels below those specified in the regulations.

Liquid effluents from Quad Cities are released to the Mississippi River in controlled batches after radioassay of each batch. Gaseous effluents are released to the atmosphere after delay to permit decay of short half-life gases. Releases to the atmosphere are calculated on the basis of analyses of daily grab samples of noble gases and continuously collected composite samples of iodine and particulate matter. The results of effluent analyses are summarized on a monthly basis and reported to the Nuclear Regulatory Commission as required per Technical Specifications. Airborne concentrations of noble gases, I-131 and particulate radioactivity in off-site areas are calculated using effluent and meteorological data on isotopic composition of effluents.

Environmental monitoring is conducted by sampling at indicator and reference (background) locations in the vicinity of the Quad Cities plant to measure changes in radiation or radioactivity levels that may be attributable to plant operations. If significant changes attributable to Quad Cities are measured, these changes are correlated with effluent releases. External gamma radiation exposure from noble gases and I-131 in milk are the most critical pathways at this site; however, an environmental monitoring program is conducted which includes other pathways of less importance.

SUMMARY

Gaseous and liquid effluents for the period remained at a fraction of the Technical Specification limits. Calculations of environmental concentrations based on effluent, Mississippi River flow, and meteorological data for the period indicate that consumption by the public of radionuclides attributable to the plant are unlikely to exceed the regulatory limits. Gamma radiation exposure from noble gases released to the atmosphere represented the critical pathway for the period with a maximum individual dose estimated to be 0.04 mrem for the year, when a shielding and occupancy factor of 0.7 is assumed. The assessment of radiation doses are performed in accordance with the Off-Site Dose Calculation Manual (ODCM). The results of analysis confirm that the station is operating in compliance with 10CFR50 Appendix I and 40 CFR 190.

1.0 EFFLUENTS

1.1 Gaseous Effluents to the Atmosphere

Measured concentrations and isotopic composition of noble gases, radioiodine, and particulate radioactivity released to the atmosphere during the year, are listed in Table 1.1-1. A total of $2.95\text{E}+03$ curies of fission and activation gases was released with an average release rate of $9.33\text{E}+01$ $\mu\text{Ci}/\text{sec}$.

A total of 0.049 curies of I-131 was released during the year, with an average release rate of $7.20\text{E}-03$ $\mu\text{Ci}/\text{sec}$.

A total of 0.557 curies of beta-gamma emitters and $4.91\text{E}-05$ curies of alpha emitters was released as airborne particulate matter, with an average release rate of $1.74\text{E}-02$ $\mu\text{Ci}/\text{sec}$.

A total of 52.5 curies of tritium was released, with an average release rate of 1.68 $\mu\text{Ci}/\text{sec}$.

1.2 Liquids Released to the Mississippi

A total of $1.82\text{E}+06$ liters of radioactive liquid waste (prior to dilution) containing 1.46 curies (excluding tritium, gases, and alpha) were discharged after dilution with a total of $1.35\text{E}+12$ liters of water. These wastes were released at a quarterly average concentration of $9.6\text{E}-09$ $\mu\text{Ci}/\text{ml}$ during the first and second quarters, discharged on an unidentified nuclide basis; and $2.03\text{E}-07$ $\mu\text{Ci}/\text{ml}$ during the third and fourth quarters, which is 33.4% of the Technical Specification release limits for unidentified radioactivity. A total of $1.5\text{E}-05$ curies of alpha radioactivity and 3.41 curies of tritium were released. Quarterly release estimates and principal radio-nuclides in liquid effluents are given in Table 1.2-1.

2.0 SOLID RADIOACTIVE WASTE

Solid radioactive wastes were shipped to U. S. Ecology; Chem Nuclear Company, Tri-State; and Barnwell Nuclear Center, South Carolina. The record of waste shipments is summarized in Table 2.0-1.

3.0 DOSE TO MAN

3.1 Gaseous Effluent Pathways

Gamma Dose Rates

Gamma air and whole body dose rates off-site were calculated based on measured release rates, isotopic composition of the noble gases, and meteorological data for the period (Table 3.1-1). Isodose

contours of whole body dose are shown in Figure 3.1-1 for the year. Based on measured effluents and meteorological data, the maximum dose to an individual would be 0.04 mrem for the year, with an occupancy or shielding factor of 0.7 included. The maximum gamma air dose was 0.075 mrad.

Beta Air and Skin Rates

The range of beta particles in air is relatively small (on the order of a few meters or less); consequently, plumes of gaseous effluents may be considered "infinite" for purpose of calculating the dose from beta radiation incident on the skin. However, the actual dose to sensitive skin tissues is difficult to calculate because this depends on the beta particle energies, thickness of inert skin, and clothing covering sensitive tissues. For purposes of this report the skin is taken to have a thickness of 7 mg/cm^2 and an occupancy factor of 1.0 is used. The skin dose from beta and gamma radiation for the year was 0.089 mrem.

The air concentrations of radioactive noble gases at the off-site receptor locations are given in Figure 3.1-2. The maximum off-site beta air dose for the year was 0.044 mrad.

Radioactive Iodine

The human thyroid exhibits a significant capacity to concentrate ingested or inhaled iodine, and the radioiodine, I-131, released during routine operation of the plant, may be made available to man thus resulting in a dose to the thyroid. The principal pathway of interest for this radionuclide is ingestion of radioiodine in milk by an infant. Calculation made in previous years indicate that contributions to doses from inhalation of I-131 and I-133, and I-133 in milk are negligible.

Iodine-131 Concentrations in Air

The calculated concentration contours for I-131 in air are shown in Figure 3.1-3. Included in these calculations is an iodine cloud depletion factor which accounts for the phenomenon of elemental iodine deposition on the ground. The maximum off-site average concentration is estimated to be $1.64\text{E-}03 \text{ pCi/m}^3$ for the year.

Dose to Infant's Thyroid

The hypothetical thyroid dose to an infant living near the plant via ingestion of milk was calculated. The radionuclide considered was I-131 and the source of milk was taken to be the nearest dairy farm with the cows pastured from May to October. The maximum infant's thyroid dose was 0.29 mrem during the year (Table 3.1-1).

Concentrations of Particulates in Air

Concentration contours of radioactive airborne particulates are shown in Figure 3.1-4. The maximum off-site average level is estimated to be $1.26E-03$ pCi/m³.

Summary of Doses

Table 3.1-1 summarizes the doses resulting from releases of airborne radioactivity via the different exposure pathways.

3.2 Liquid Effluent Pathways

The three principal pathways through the aquatic environment for potential doses to man from liquid waste are ingestion of potable water, eating aquatic foods, and exposure while walking on the shoreline. Not all of these pathways are applicable at a given time or station but a reasonable approximation of the dose can be made by adjusting the dose formula for season of the year or type and degree of use of the aquatic environment. NRC* developed equations were used to calculate the doses to the whole body, lower GI tract, thyroid, bone and skin; specific parameters for use in the equations are given in the Commonwealth Edison Off-site Dose Calculation Manual. The maximum whole body dose for the year was $9.97E-01$ mrem and no organ dose exceeded 1.45 mrem.

4.0 SITE METEOROLOGY

A summary of the site meteorological measurements taken during each quarter of the year is given in Appendix II. The data are presented as cumulative joint frequency distributions of 296' level wind direction and wind speed class by atmospheric stability class determined from the temperature difference between the 296' and 33' levels. Data recovery for all measurements on the tower was about 99.7%.

5.0 ENVIRONMENTAL MONITORING

Tables 5.0-1 and 5.0-2 provide an outline of the radiological environmental monitoring program as required in current Technical Specifications. This program went into effect in November 1977 and differs from previous programs in the number and types of analyses performed. Tables 5.0-3 to 5.0-6 summarize data for the year.

Except for tables of special interest, tables listing all data are no longer included in the annual report. All data tables are available for inspection at the Station or in the Corporate offices.

Specific findings for various environmental media are discussed in Sections 5.1 through 5.5.

* Nuclear Regulatory Commission, Regulatory Guide 1.109 (Rev. 1).

5.1 Gamma Radiation

External radiation dose from on-site sources and noble gases released to the atmosphere was measured at six indicator and ten reference (background) locations using solid lithium fluoride thermoluminescent dosimeters (TLD). A comparison of the TLD results for reference stations with on-site and off-site indicator stations is included in Table 5.1-1. A total of 61 additional TLDs were installed on June 1, 1980 such that each sector was covered at both five miles and the site boundary. Locations of the TLDs are shown in Figures 5.0-1 and 5.0-2.

5.2 Airborne I-131 and Particulate Radioactivity

Concentrations of airborne I-131 and particulate radioactivity at monitoring locations are summarized in Tables 5.0-2 through 5.0-5. Locations of the samplers are shown in figure 5.0-1. Airborne I-131 remained below the LLD of 0.1 pCi/m^3 throughout the year.

Gross beta concentrations ranged from 0.006 to 0.071 pCi/m^3 at indicator locations with an average concentration of 0.024 pCi/m^3 for the year. No radioactivity attributable to station operation was detected in any sample.

5.3 Aquatic Radioactivity

Surface water samples were collected daily and composited for analysis weekly from the Inlet Canal, Blowdown Diffuser Pipe, East Moline Water Works, and Davenport Water Works. The cooling water samples were analyzed weekly for gross beta concentrations. A composite sample from each quarter from the blowdown diffuser pipe did not indicate measurable radioactivity attributable to station operation except for the last three weeks in December, 1985. During these three weeks, the mean gross beta concentration in the inlet canal samples measured 19.8 pCi/l and 1566 pCi/l in the samples from the diffuser pipe blowdown. For the rest of the period, annual mean gross beta concentration in the blowdown diffuser and inlet canal pipe water samples measured 4.1 and 3.9 pCi/l , respectively.

Samples from the two water works were composited monthly and analyzed for gamma emitters. All samples analyzed were below the limits of detection for the program indicating that there was no measurable amount of radioactivity due to station operation present.

Levels of gamma radioactivity in fish collected were measured and found in all cases to be below the lower limits of detection for the program.

A sediment sample was analyzed by gamma spectrometry. Gamma-emitters were below the limits of detection, indicating the presence of no radioactivity due to station operation.

5.4 Milk

Milk samples were collected monthly from November through April and weekly from May through October and analyzed for iodine-131. Sampled locations were the Hansen Dairy Farm located about 6.0 miles northeast of the Station, and Musal Dairy Farm located 5.5 miles southwest of the Station, both being the closest dairies to the station. Radioiodine was below the limits of detection of 0.5 pCi/l during the grazing period (May to October) and 5.0 pCi/l during the non-grazing period (November to April).

5.5 Special Collection

No special collections were made during the period.

6.0 ANALYTICAL PROCEDURES

A description of the procedures used for analyzing radioactivity in environmental samples is given in Appendix III of this report.

7.0 MILCH ANIMAL CENSUS

A census of milch animals was conducted within five miles of the Station. The survey was conducted by "door-to-door" canvas and by information from Illinois and Iowa Agricultural Agents. The census was conducted by G. Kreuder on August 16, 1985.

There were no dairy farms within a five mile radius of the Quad Cities Nuclear Power Station. The findings of the survey follow.

Within 2 Miles of the Plant

None

Within 2 - 6 Miles of the Plant

Illinois* -- Mel Hanson (Jeff) -- (309) 887-4568

6.0 miles NE -- Type: Holstein
Number: 35
Feed: Hay, corn silage,
commercial feed

Gene Dornbush -- (309) 887-4986

5-1/2 miles NE -- Type: Holstein
Number: 20
Feed: Hay, green chopped
hay, corn

Within 2 - 6 Miles of the Plant (continued)

Iowa* -- Alan Musal, Rural Route, Princeton, Iowa -
(319) 289-4786

5-1/2 miles SW -- Type: Holstein
Number: 55
Feed: Hay, chopped corn

Carl Otte, Rural Route, Princeton, Iowa

5-1/2 miles W -- Number: 30

8.0 NEAREST RESIDENT CENSUS

A census of the nearest residences within a five (5) mile radius was conducted on August 16, 1985 by G. Kreuder. The location of residences remained unchanged from the previous census. The nearest residences are listed below.

<u>Direction</u>	<u>Distance</u>
N	0.6 miles
NNE	1.0 miles
NE	1.3 miles
ENE	2.8 miles
E	2.3 miles
ESE	2.0 miles
SE	1.0 miles
SSE	1.1 miles
S	0.8 miles
SSW	3.0 miles
SW	2.8 miles
WSW	2.0 miles
W	2.5 miles
WNW	2.5 miles
NW	2.0 miles
NNW	2.0 miles

9.0 INTERLABORATORY COMPARISON PROGRAM RESULTS

Teledyne Isotopes Midwest Laboratory (formerly Hazleton Environmental Sciences) has participated in interlaboratory comparison (crosscheck) programs since the formulation of its quality control program in December 1971. These programs are operated by agencies which supply environmental-type samples (e.g., milk or water) containing concentrations of radionuclides known to the issuing agency but not to participant laboratories. The purpose of such a program is to provide an independent check on the laboratory's analytical procedures and to alert it to any possible problems.

Participant laboratories measure the concentrations of specified radionuclides and report them to the issuing agency. Several months later, the agency reports the known values to the participant laboratories and specifies control limits. Results consistently higher or lower than the known values or outside the control limits indicate a need to check the instruments or procedures used.

The results in Table A-1 were obtained through participation in the environmental sample crosscheck program for milk, water, air filters, and food samples during the period 1982 through 1985. This program has been conducted by the U. S. Environmental Protection Agency Intercomparison and Calibration Section, Quality Assurance Branch, Environmental Monitoring and Support Laboratory, Las Vegas, Nevada.

The results in Table A-2 were obtained for thermoluminescent dosimeters (TLD's) during the period 1976, 1977, 1979, 1980, and 1981 through participation in the Second, Third, Fourth, and Fifth International Intercomparison of Environmental Dosimeters under the sponsorships listed in Table A-2.

Table A-1. U.S. Environmental Protection Agency's crosscheck program, comparison of EPA and Teledyne Isotopes Midwest Laboratory results for milk, water, air filters, and food samples, 1982 through 1985.^a

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STW-270	Water	Jan. 1982	Sr-89	24.3 \pm 2.0	21.0 \pm 5.0
			Sr-90	9.4 \pm 0.5	12.0 \pm 1.5
STW-273	Water	Jan. 1982	I-131	8.6 \pm 0.6	8.4 \pm 1.5
STW-275	Water	Feb. 1982	H-3	1580 \pm 147	1820 \pm 342
STW-276	Water	Feb. 1982	Cr-51	<61	0
			Co-60	26.0 \pm 3.7	20 \pm 5
			Zn-65	<13	15 \pm 5
			Ru-106	<46	20 \pm 5
			Cs-134	26.8 \pm 0.7	22 \pm 5
			Cs-137	29.7 \pm 1.4	23 \pm 5
STW-277	Water	Mar. 1982	Ra-226	11.9 \pm 1.9	11.6 \pm 1.7
STW-278	Water	Mar. 1982	Gross alpha	15.6 \pm 1.9	19 \pm 5
			Gross beta	19.2 \pm 0.4	19 \pm 5
STW-280	Water	Apr. 1982	H-3	2690 \pm 80	2860 \pm 360
STW-281	Water	Apr. 1982	Gross alpha	75 \pm 7.9	85 \pm 21
			Gross beta	114.1 \pm 5.9	106 \pm 5.3
			Sr-89	17.4 \pm 1.8	24 \pm 5
			Sr-90	10.5 \pm 0.6	12 \pm 1.5
			Ra-226	11.4 \pm 2.0	10.9 \pm 1.5
			Co-60	<4.6	0
STW-284	Water	May 1982	Gross alpha	31.5 \pm 6.5	27.5 \pm 7
			Gross beta	25.9 \pm 3.4	29 \pm 5
STW-285	Water	June 1982	H-3	1970 \pm 1408	1830 \pm 340
STW-286	Water	June 1982	Ra-226	12.6 \pm 1.5	13.4 \pm 3.5
			Ra-228	11.1 \pm 2.5	8.7 \pm 2.3
STW-287	Water	June 1982	I-131	6.5 \pm 0.3	4.4 \pm 0.7
STW-290	Water	Aug. 1982	H-3	3210 \pm 140	2890 \pm 619

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STW-291	Water	Aug. 1982	I-131	94.6 \pm 2.5	87 \pm 15
STW-292	Water	Sept. 1982	Sr-89	22.7 \pm 3.8	24.5 \pm 8.7
			Sr-90	10.9 \pm 0.3	14.5 \pm 2.6
STW-296	Water	Oct. 1982	Co-60	20.0 \pm 1.0	20 \pm 8.7
			Zn-65	32.3 \pm 5.1	24 \pm 8.7
			Cs-134	15.3 \pm 1.5	19.0 \pm 8.7
			Cs-137	21.0 \pm 1.7	20.0 \pm 8.7
STW-297	Water	Oct. 1982	H-3	2470 \pm 20	2560 \pm 612
STW-298	Water	Oct. 1982	Gross alpha	32 \pm 30	55 \pm 24
			Gross beta	81.7 \pm 6.1	81 \pm 8.7
			Sr-89	<2	0
			Sr-90	14.1 \pm 0.9	17.2 \pm 2.6
			Cs-134	<2	1.8 \pm 8.7
			Cs-137	22.7 \pm 0.6	20 \pm 8.7
			Ra-226	13.6 \pm 0.3	12.5 \pm 3.2
			Ra-228	3.9 \pm 1.0	3.6 \pm 0.9
STW-301	Water	Nov. 1982	Gross alpha	12.0 \pm 1.0	19.0 \pm 8.7
			Gross beta	34.0 \pm 2.7	24.0 \pm 8.7
STW-302	Water	Dec. 1982	I-131	40.0 \pm 0.0	37.0 \pm 10
STW-303	Water	Dec. 1982	H-3	1940 \pm 20	1990 \pm 345
STW-304	Water	Dec. 1982	Ra-226	11.7 \pm 0.6	11.0 \pm 1.7
			Ra-228	<3	0
STW-306	Water	Jan. 1983	Sr-89	20.0 \pm 8.7	29.2 \pm 5
			Sr-90	21.7 \pm 8.4	17.2 \pm 1.5
STW-307	Water	Jan. 1983	Gross alpha	29.0 \pm 4.09	29.0 \pm 13
			Gross beta	29.3 \pm 0.6	31.0 \pm 8.7
STM-309	Milk	Feb. 1983	Sr-89	35 \pm 2.0	37 \pm 8.7
			Sr-90	13.7 \pm 0.6	18 \pm 2.6
			I-131	55.7 \pm 3.2	55 \pm 10.4
			Cs-137	29 \pm 1.0	26 \pm 8.7
			Ba-140	<27	0
			K-40	1637 \pm 5.8	1512 \pm 131

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STW-310	Water	Feb. 1983	H-3	2470 \pm 80	2560 \pm 612
STW-311	Water	March 1983	Ra-226 Ra-228	11.9 \pm 1.3 <2.7	12.7 \pm 3.3 0
STW-312	Water	March 1983	Gross alpha Gross beta	31.6 \pm 4.59 27.0 \pm 2.0	31 \pm 13.4 28 \pm 8.7
STW-313	Water	April 1983	H-3	3240 \pm 80	3330 \pm 627
STW-316	Water	May 1983	Gross alpha Gross beta Sr-89 Sr-90 Ra-226 Co-60 Cs-134 Cs-137	94 \pm 7 133 \pm 5 19 \pm 1 12 \pm 1 7.9 \pm 0.4 30 \pm 2 27 \pm 2 29 \pm 1	64 \pm 19.9 149 \pm 12.4 24 \pm 8.7 13 \pm 2.6 8.5 \pm 2.25 30 \pm 8.7 33 \pm 8.7 27 \pm 8.7
STW-317	Water	May 1983	Sr-89 Sr-90	59.7 \pm 2.1 33.7 \pm 1.5	57 \pm 8.7 38 \pm 3.3
STW-318 ^f	Water	May 1983	Gross alpha Gross beta	12.8 \pm 1.5 49.4 \pm 3.9	11 \pm 8.7 57 \pm 8.7
STM-320	Milk	June 1983	Sr-89 Sr-90 I-131 Cs-137 K-40	20 \pm 0 10 \pm 1 30 \pm 1 52 \pm 2 1553 \pm 57	25 \pm 8.7 16 \pm 2.6 30 \pm 10.4 47 \pm 8.7 1486 \pm 129
STW-321	Water	June 1983	H-3	1470 \pm 89	1529 \pm 583
STW-322	Water	June 1983	Ra-226 Ra-228	4.3 \pm 0.2 <2.5	4.8 \pm 1.24 0
STW-323	Water	July 1983	Gross alpha Gross beta	3 \pm 1 21 \pm 0	7 \pm 8.7 22 \pm 8.7
STW-324	Water	August 1983	I-131	13.3 \pm 0.6	14 \pm 10.4

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STAF-326	Air Filter	August 1983	Gross beta	42 \pm 2	36 \pm 8.7
			Sr-90	14 \pm 2	10 \pm 2.6
			Cs-137	19 \pm 1	15 \pm 8.7
STW-328	Water	Sept. 1983	Gross alpha	2.3 \pm 0.6	5 \pm 8.7
			Gross beta	10.7 \pm 1.2	9 \pm 8.7
STW-329	Water	Sept. 1983	Ra-226	3.0 \pm 0.2	3.1 \pm 0.81
			Ra-228	3.2 \pm 0.7	2.0 \pm 0.52
STW-331	Water	Oct. 1983	H-3	1300 \pm 30	1210 \pm 570
STW-335	Water	Dec. 1983	I-131	19.6 \pm 1.9	20 \pm 10.4
STW-336	Water	Dec. 1983	H-3	2870 \pm 100	2389 \pm 608
STAF-337	Air Filter	Nov. 1983	Gross alpha	18.0 \pm 0.2	19 \pm 8.7
			Gross beta	58.6 \pm 1.2	50 \pm 8.7
			Sr-90	10.9 \pm 0.1	15 \pm 2.6
			Cs-137	30.1 \pm 2.5	20 \pm 8.7
STW-339	Water	Jan. 1984	Sr-89	47.2 \pm 1.9	36 \pm 8.7
			Sr-90	22.5 \pm 4.0	24 \pm 2.6
STW-343	Water	Feb. 1984	H-3	2487 \pm 76	2383 \pm 607
STM-347	Milk	March 1984	I-131	5.3 \pm 1.1	6 \pm 1.6
STW-349	Water	March 1984	Ra-226	4.0 \pm 0.2	4.1 \pm 1.06
			Ra-228	3.6 \pm 0.3	2.0 \pm 0.52
STW-350	Water	March 1984	Gross alpha	3.8 \pm 1.1	5 \pm 8.7
			Gross beta	24.2 \pm 2.0	20 \pm 8.7
STW-354	Water	April 1984	H-3	3560 \pm 50	3508 \pm 630
STW-355	Water	April 1984	Gross alpha	21.0 \pm 4.1	35 \pm 15.2
			Gross beta	127.8 \pm 4.1	147 \pm 12.7
			Sr-89	29.3 \pm 2.0	23 \pm 8.7
			Sr-90	16.6 \pm 0.7	26 \pm 2.6
			Ra-226	4.0 \pm 1.0	4.0 \pm 1.04
			Co-60	32.3 \pm 1.4	30 \pm 8.7
			Cs-134	33.6 \pm 3.1	30 \pm 8.7
			Cs-137	33.3 \pm 2.2	26 \pm 8.7

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/lb	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STW-358	Water	May 1984	Gross alpha Gross beta	3.0 \pm 0.6 6.7 \pm 1.2	3 \pm 8.7 6 \pm 8.7
STM-366	Milk	June 1984	Sr-89 Sr-90 I-131 Cs-137 K-40	21 \pm 3.1 13 \pm 2.0 46 \pm 5.3 38 \pm 4.0 1577 \pm 172	25 \pm 8.7 17 \pm 2.6 43 \pm 10.4 35 \pm 8.7 1496 \pm 130
STW-368	*Water	July 1984	Gross alpha Gross beta	5.1 \pm 1.1 11.9 \pm 2.4	6 \pm 8.7 13 \pm 8.7
STW-369	Water	August 1984	I-131	34.3 \pm 5.0	34.0 \pm 10.4
STW-370	Water	August 1984	H-3	3003 \pm 253	2817 \pm 617
STF-371	Food	July 1984	Sr-89 Sr-90 I-131 Cs-137 K-40	22.0 \pm 5.3 14.7 \pm 3.1 <172 24.0 \pm 5.3 2503 \pm 132	25.0 \pm 8.7 20.0 \pm 2.6 39.0 \pm 10.4 25.0 \pm 8.7 2605 \pm 226.0
STAF-372	Air Filter	August 1984	Gross alpha Gross beta Sr-90 Cs-137	15.3 \pm 1.2 56.0 \pm 0.0 14.3 \pm 1.2 21.0 \pm 2.0	17 \pm 8.7 51 \pm 8.7 18 \pm 2.4 15 \pm 8.7
STW-375	Water	Sept. 1984	Ra-226 Ra-228	5.1 \pm 0.4 2.2 \pm 0.1	4.9 \pm 1.27 2.3 \pm 0.60
STW-377	Water	Sept. 1984	Gross alpha Gross beta	3.3 \pm 1.2 12.7 \pm 2.3	5.0 \pm 8.7 16.0 \pm 8.7
STW-379	Water	Oct. 1984	H-3	2860 \pm 312	2810 \pm 356
STW-380	Water	Oct. 1984	Cr-51 Co-60 Zn-65 Ru-106 Cs-134 Cs-137	<36 20.3 \pm 1.2 150 \pm 8.1 <30 31.3 \pm 7.0 26.7 \pm 1.2	40 \pm 8.7 20 \pm 8.7 147 \pm 8.7 47 \pm 8.7 31 \pm 8.7 24 \pm 8.7

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/lb	
				TIML Result ±2 ^c	EPA Result ±3, n=1 ^d
STM-382	Milk	Oct. 1984	Sr-89	15.7±4.2	22±8.7
			Sr-90	12.7±1.2	16±2.6
			I-131	41.7±3.1	42±10.4
			Cs-137	31.3±6.1	32±8.7
			K-40	1447±66	1517±131
STW-384	Water (Blind)	Oct. 1984 Sample A	Gross alpha	9.7±1.2	14±8.7
			Ra-226	3.3±0.2	3.0±0.8
			Ra-228	3.4±1.6	2.1±0.5
			Uranium	NA ^e	5±10.4
	Sample B	Gross beta	48.3±5.0	64±8.7	
		Sr-89	10.7±4.6	11±8.7	
		Sr-90	7.3±1.2	12±2.6	
		Co-60	16.3±1.2	14±8.7	
		Cs-134	<2	2±8.7	
Cs-137	16.7±1.2	14±8.7			
STAF-387	Air Filter	Nov. 1984	Gross alpha	18.7±1.2	15±8.7
			Gross beta	59.0±5.3	52±8.7
			Sr-90	18.3±1.2	21±2.6
			Cs-137	10.3±1.2	10±8.7
STW-388	Water	Dec. 1984	I-131	28.0±2.0	36±10.4
STW-389	Water	Dec. 1984	H-3	3583±110	3182±624
STW-391	Water	Dec. 1984	Ra-226	8.4±1.7	8.6±2.2
			Ra-228	3.1±0.2	4.1±1.1
STW-392	Water	Jan. 1985	Sr-89	<3.0	3.0±8.7
			Sr-90	27.3±5.2	30.0±2.6
STW-393	Water	Jan. 1985	Gross alpha	3.3±1.2	5±8.7
			Gross beta	17.3±3.0	15±8.7
STS-395	Food	Jan. 1985	Sr-89	25.3±6.4	34.0±5.0
			Sr-90	27.0±8.8	26.0±1.5
			I-131	38.0±2.0	35.0±6.0
			Cs-137	32.7±2.4	29.0±5.0
			K-40	1410±212	1382±120

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b		
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$	
STW-397	Water	Feb. 1985	Cr-51	<29	48±8.7	
			Co-60	21.3±3.0	20±8.7	
			Zn-65	53.7±5.0	55±8.7	
			Ru-106	<23	25±8.7	
			Cs-134	32.3±1.2	35±8.7	
			Cs-137	25.3±3.0	25±8.7	
STW-398	Water	Feb. 1985	H-3	3869±319	3796±634	
STM-400	Milk	March 1985	I-131	7.3±2.4	9.0±0.9	
STW-402	Water	March 1985	Ra-226	4.6±0.6	5.0±1.3	
			Ra-228	<0.8	9.0±2.3	
			Reanalysis Ra-228	9.0±0.4		
STW-404	Water	March 1985	Gross alpha	4.7±2.3	6±8.7	
			Gross beta	11.3±1.2	15±8.7	
STAF-405	Air Filter	March 1985	Gross alpha	9.3±1.0	10.0±8.7	
			Gross beta	42.0±1.1	36.0±8.7	
			Sr-90	13.3±1.0	15.0±2.6	
			Cs-137	6.3±1.0	6.0±8.7	
STW-407	Water	April 1985	I-131	8.0±0.0	7.5±1.3	
STW-408	Water	April 1985	H-3	3399±150	3559±630	
STW-409	Water	April 1985				
			(Blind)			
			Sample A	Gross alpha	29.7±1.8	32.0±5.0
				Ra-226	4.4±0.2	4.1±0.6
				Ra-228	NA ^e	6.2±0.9
				Uranium	NA ^e	7.0±6.0
			Sample B	Gross beta	74.3±11.8	72.0±5.0
				Sr-89	12.3±7.6	10.0±5.0
				Sr-90	14.7±2.4	15.0±1.5
				Co-60	14.7±2.4	15.0±5.0
				Cs-134	12.0±2.0	15.0±5.0
				Cs-137	14.0±2.0	12.0±5.0

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/lb	
				TIML Result $\pm 2^c$	EPA Result $\pm 3, n=1^d$
STW-413	Water	May 1985	Sr-89 Sr-90	36.0 \pm 12.4 14.3 \pm 4.2	39.0 \pm 5.0 15.0 \pm 1.5
STW-414	Water	May 1985	Gross alpha Gross beta	8.3 \pm 4.1 8.7 \pm 1.2	12.0 \pm 5.0 11.0 \pm 5.0
STW-416	Water	June 1985	Cr-51 Co-60 Zn-65 Ru-106 Cs-134 Cs-137	44.7 \pm 6.0 14.3 \pm 1.2 50.3 \pm 7.0 55.3 \pm 5.8 32.7 \pm 1.2 22.7 \pm 2.4	44.0 \pm 5.0 14.0 \pm 5.0 47.0 \pm 5.0 62.0 \pm 5.0 35.0 \pm 5.0 20.0 \pm 5.0
STW-418	Water	June 1985	H-3	2446 \pm 132	2416 \pm 351
STM-421	Milk	June 1985	Sr-89 Sr-90 I-131 Cs-137 K-40	10.3 \pm 4.6 9.0 \pm 2.0 11.7 \pm 1.2 12.7 \pm 1.2 1512 \pm 62	11.0 \pm 8.7 11.0 \pm 2.6 11.0 \pm 10.4 11.0 \pm 8.7 1525 \pm 132
STW-423	Water	July 1985	Gross alpha Gross beta	5.0 \pm 0.0 5.0 \pm 2.0	11.0 \pm 8.7 8.0 \pm 8.7
STW-425	Water	August 1985	I-131	25.7 \pm 3.0	33.0 \pm 10.4
STW-426	Water	August 1985	H-3	4363 \pm 83	4480 \pm 776
STAF-427	Air Filter	August 1985	Gross alpha Gross beta Sr-90 Cs-137	11.3 \pm 0.6 46.0 \pm 1.0 17.7 \pm 0.6 10.3 \pm 0.6	13.0 \pm 8.7 44.0 \pm 8.7 18.0 \pm 2.6 8.0 \pm 8.7
STW-429	Water	Sept. 1985	Sr-89 Sr-90	15.7 \pm 0.6 7.0 \pm 0.0	20.0 \pm 8.7 7.0 \pm 2.6
STW-430	Water	Sept. 1985	Ra-226 Ra-228	8.2 \pm 0.3 4.1 \pm 0.3	8.9 \pm 2.3 4.6 \pm 1.2
STW-431	Water	Sept. 1985	Gross alpha Gross beta	4.7 \pm 0.6 4.7 \pm 1.2	8.0 \pm 8.7 8.0 \pm 8.7

Table A-1. (continued)

Lab Code	Sample Type	Date Collected	Analysis	Concentration in pCi/l ^b	
				TIML Result $\pm 2\sigma^c$	EPA Result $\pm 3\sigma, n=1^d$
STW-433	Water	Oct. 1985	Cr-51	<13	21.0 \pm 8.7
			Co-60	19.30.6	20.0 \pm 8.7
			Zn-65	19.7 \pm 0.6	19.0 \pm 8.7
			Ru-106	<19	20.0 \pm 8.7
			Cs-134	17.0 \pm 1.0	20.0 \pm 8.7
			Cs-137	19.3 \pm 1.2	20.0 \pm 8.7
STW-435	Water	Oct. 1985	H-3	1957 \pm 50	1974 \pm 598

^a Results obtained by Teledyne Isotopes Midwest Laboratory as a participant in the environmental sample crosscheck program operated by the Intercomparison and Calibration Section, Quality Assurance Branch, Environmental Monitoring and Support Laboratory, U.S. Environmental Protection Agency, (EPA), Las Vegas, Nevada.

^b All results are in pCi/l, except for elemental potassium (K) data, which are in mg/l; air filter samples, which are in pCi/filter; and food, which is in pCi/kg.

^c Unless otherwise indicated, the TIML results are given as the mean ± 2 standard deviations for three determinations.

^d USEPA results are presented as the known values \pm control limits of 3σ for $n = 1$.

^e NA = Not analyzed.

^f Analyzed but not reported to the EPA.

^g Results after calculations corrected (error in calculations when reported to EPA).

Table A-2. Crosscheck program results, thermoluminescent dosimeters (TLDs).

Lab Code	TLD Type	Measurement	mR		Average $\pm 2\sigma^d$ (all participants)
			Teledyne Result $\pm 2\sigma^a$	Known Value	
<u>2nd International Intercomparison^b</u>					
115-2 ^b	CaF ₂ :Mn Bulb	Gamma-Field	17.0 \pm 1.9	17.1 ^c	16.4 \pm 7.7
		Gamma-Lab	20.8 \pm 4.1	21.3 ^c	18.8 \pm 7.6
<u>3rd International Intercomparison^e</u>					
115-3 ^e	CaF ₂ :Mn Bulb	Gamma-Field	30.7 \pm 3.2	34.9 \pm 4.8 ^f	31.5 \pm 3.0
		Gamma-Lab	89.6 \pm 6.4	91.7 \pm 14.6 ^f	86.2 \pm 24.0
<u>4th International Intercomparison^g</u>					
115-49	CaF ₂ :Mn Bulb	Gamma-Field	14.1 \pm 1.1	14.1 \pm 1.4 ^f	16.0 \pm 9.0
		Gamma-Lab (Low)	9.3 \pm 1.3	12.2 \pm 2.4 ^f	12.0 \pm 7.6
		Gamma-Lab (High)	40.4 \pm 1.4	45.8 \pm 9.2 ^f	43.9 \pm 13.2
<u>5th International Intercomparison^h</u>					
115-5A ^h	CaF ₂ :Mn Bulb	Gamma-Field	31.4 \pm 1.8	30.0 \pm 6.0 ⁱ	30.2 \pm 14.6
		Gamma-Lab at beginning	77.4 \pm 5.8	75.2 \pm 7.6 ⁱ	75.8 \pm 40.4
		Gamma-Lab at the end	96.6 \pm 5.8	88.4 \pm 8.8 ⁱ	90.7 \pm 31.2

Table A-2. (Continued)

Lab Code	TLD Type	Measurement	mR		
			Teledyne Result $\pm 2\sigma^a$	Known Value	Average $\pm 2\sigma^d$ (all participants)
115-58 ^h	LiF-100 Chips	Gamma-Field	30.3 \pm 4.8	30.0 \pm 6 ⁱ	30.2 \pm 14.6
		Gamma-Lab at beginning	81.1 \pm 7.4	75.2 \pm 7.6 ⁱ	75.8 \pm 40.4
		Gamma-Lab at the end	85.4 \pm 11.7	88.4 \pm 8.8 ⁱ	90.7 \pm 31.2

^a Lab result given is the mean ± 2 standard deviations of three determinations.

^b Second International Intercomparison of Environmental Dosimeters conducted in April of 1976 by the Health and Safety Laboratory (GASL), New York, New York, and the School of Public Health of the University of Texas, Houston, Texas.

^c Value determined by sponsor of the intercomparison using continuously operated pressurized ion chamber.

^d Mean ± 2 standard deviations of results obtained by all laboratories participating in the program.

^e Third International Intercomparison of Environmental Dosimeters conducted in summer of 1977 by Oak Ridge National Laboratory and the School of Public Health of the University of Texas, Houston, Texas.

^f Value ± 2 standard deviations as determined by sponsor of the intercomparison using continuously operated pressurized ion chamber.

^g Fourth International Intercomparison of Environmental Dosimeters conducted in summer of 1979 by the School of Public Health of the University of Texas, Houston, Texas.

^h Fifth International Intercomparison of Environmental Dosimeter conducted in fall of 1980 at Idaho Falls, Idaho and sponsored by the School of Public Health of the University of Texas, Houston, Texas and Environmental Measurements Laboratory, New York, New York, U.S. Department of Energy.

ⁱ Value determined by sponsor of the intercomparison using continuously operated pressurized ion chamber.

APPENDIX I

DATA TABLES AND FIGURES

Table 1.1-1

QCP 100-S25
Revision 3
May 1985

EFFLUENT AND WASTE DISPOSAL
SEMI-ANNUAL REPORT Jan-June 1985
GASEOUS EFFLUENTS - SUMMATION OF ALL RELEASES

	Unit	Quarter First	Quarter Second	Est. Total Error, %
A. FISSION & ACTIVATION GASES				
1. Total Release	CI	3.7E02	7.3E02	
2. Average release rate for period	μ CI/sec	4.8E01	9.3E01	
3. Percent of Tech Spec limit * Chimney & stack	%	3.0E-01 1.0E-1	4.1E-1 1.7E-1	
B. IODINE				
1. Total Iodine-131	CI	1.4E-02	5.7E-03	
2. Average release rate for period	μ CI/sec	1.8E-03	7.2E-04	
C. PARTICULATES				
1. Particulates with half-lives > 8 days	CI	3.2E-02	3.5E-02	
2. Average release rate for period	μ CI/sec	4.1E-03	4.5E-03	
3. Gross alpha radioactivity	CI	9.6E-06	1.1E-05	
D. TRITIUM				
1. Total Release	CI	21.5	7.07	
2. Average release rate for period	μ CI/sec	2.80	0.90	
E. Iodine 131 & 133, Tritium and Particulates				
Percent of Tech spec Limit Chimney & stack	%	1.7E-01	9.0E-01	

Nobel Gas Gamma/Beta

Table 1.1-1 (continued)

QCP 100-S25
Revision 3
May 1985

EFFLUENT AND WASTE DISPOSAL
SEMI-ANNUAL REPORT July-December 1985
GASEOUS EFFLUENTS - SUMMATION OF ALL RELEASES

	Unit	Quarter Third	Quarter Fourth	Est. Total Error, %
A. FISSION & ACTIVATION GASES				
1. Total Release	CI	6.5E02	1.2E03	
2. Average release rate for period	$\mu\text{Ci}/\text{sec}$	8.4E01	1.5E02	
3. Percent of Tech Spec limit * Chimney & stack	%	$\frac{2.3E-01}{6.0E-2}$	$\frac{5.6E-1}{9.8E-2}$	
B. IODINE				
1. Total Iodine-131	CI	1.7E-02	1.2E-02	
2. Average release rate for period	$\mu\text{Ci}/\text{sec}$	2.1E-03	1.5E-03	
C. PARTICULATES				
1. Particulates with half-lives > 8 days	CI	2.2E-01	2.7E-01	
2. Average release rate for period	$\mu\text{Ci}/\text{sec}$	2.7E-02	3.4E-02	
3. Gross alpha radioactivity	CI	1.34E-05	1.51E-05	
D. TRITIUM				
1. Total Release	CI	7.34E00	1.63E01	
2. Average release rate for period	$\mu\text{Ci}/\text{sec}$	9.2E-01	2.1E00	
E. Iodine 131 & 133, Tritium and Particulates				
Percent of Tech spec Limit Chimney & stack	%	2.3E00	5.1E-01	

*Noble Gas Gamma Radiation/Noble Gas Beta Radiation

Table 1.1-1 (continued)

QCP 100-525
Revision 3

MAIN CHIMNEY

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter First	Quarter Second	Quarter	Quarter
1. Fission gases					
Kr-85	CI	< LLD	< LLD		
Kr-85m	CI	5.7E00	3.4E01		
Kr-87	CI	5.1E00	3.5E01		
Kr-88	CI	7.7E00	3.9E01		
Xe-133	CI	3.1E01	1.7E02		
X3-135	CI	7.3E01	2.4E02		
Xe-135m	CI	5.9E01	4.3E01		
Xe-138	CI	1.7E02	1.3E02		
	CI	--	--		
	CI	--	--		
	CI	--	--		
Unidentified	CI	--	--		
Total for Period	CI	2.5E02	6.9E02		

2. Iodines

I-131	CI	1.11E-02	4.63E-03		
I-133	CI	6.70E-02	4.07E-02		
I-135	CI	6.20E-02	7.21E-02		
Total for Period	CI	1.40E-01	1.17E-01		

Table 1.1-1 (continued)

QCP 100-525
Revision 3Main Chimney
GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter Third	Quarter Fourth	Quarter	Quarter
1. Fission gases					
Kr-85	CI	<LLD	<LLD		
Kr-85m	CI	5.1E01	8.1E01		
Kr-87	CI	3.4E00	5.9E01		
Kr-88	CI	1.7E01	1.1E02		
Xe-133	CI	4.1E02	4.8E02		
Xe-135	CI	4.5E01	3.0E02		
Xe-135m	CI	3.8E+1	7.6E01		
Xe-138	CI	9.0E01	6.8E01		
	CI				
	CI				
	CI				
Unidentified	CI	<LLD	<LLD		
Total for Period	CI	6.5E02	1.2E03		

2. Iodines

I-131	CI	1.5E-02	9.0E-03		
I-133	CI	8.9E-02	4.4E-02		
I-135	CI	1.3E-01	8.7E-02		
Total for Period	CI	2.3E-01	1.4E-01		

Table 1.1-1 (continued)

OCP 100-S25
Revision 3

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter First	Quarter Second	Quarter ---	Quarter ---
3. Particulates		Main Chimney			
SR-89	CI	1.22E-03	1.45E-03		
SR-90	CI	3.65E-05	3.08E-05		
Cs-134	CI	7.10E-05	1.17E-04		
Cs-137	CI	4.34E-04	8.78E-04		
Ba-140	CI	3.24E-03	2.54E-03		
La-140	CI	8 Day	Half Line		
Cr-51	CI	<LLD	<LLD		
Mn-54	CI	<LLD	<LLD		
Co-58	CI	<LLD	<LLD		
Co-60	CI	9.60E-04	1.02E-03		
I-131	CI	7.50E-04	3.95E-04		
Ag-110m	CI	<LLD	2.1E-05		
	CI	7.38E-03	6.10E-03		
	CI	9.8 E-03	8.20E-03		
	CI				
	CI				
	CI				
Unidentified	CI				

* Sr-89, Sr-90 Values for May and June are projected

Table 1.1-1 (continued)

QCP 100-S25
Revision 3

Main Chimney

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter Third	Quarter Fourth	Quarter	Quarter

3. Particulates

SR-89	CI	1.4E-03	3.3E-03		
SR-90	CI	2.5E-05	1.3E-05		
Cs-134	CI	1.39E-05	<LLD		
Cs-137	CI	6.4E-04	8.1E-04		
Ba-140	CI	4.7E-03	3.1E-03		
La-140	CI	<LLD	<LLD		
Cr-51	CI	<LLD	<LLD		
Mn-54	CI	<LLD	<LLD		
Co-58	CI	<LLD	<LLD		
Co-60	CI	1.2E-03	9.1E-04		
I-131	CI	1.1E-03	1.6E-03		
Ag-110m	CI	<LLD	<LLD		
I-133	CI	1.1E-02	9.7E-03		
I-135	CI	1.8E-01	2.4E-01		
Ru103	CI	4.8E-4	<LLD		
	CI				
	CI				
Unidentified	CI	<LLD	<LLD		
TOTAL	CI	2.1E-01	2.6E-01		

Table 1.1-1 (continued)

QCP 100-S25
Revision 3

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter First	Quarter Second	Quarter	Quarter

1. Fission gases

Reactor Vent

Kr-85	CI	<LLD	<LLD		
Kr-85m	CI	<LLD	<LLD		
Kr-87	CI	<LLD	3.01		
Kr-88	CI	<LLD	<LLD		
Xe-133	CI	1.2	24.8		
X3-135	CI	17.6	9.13		
Xe-135m	CI	3.01	<LLD		
Xe-138	CI	<LLD	<LLD		
	CI	---	---		
	CI	---	---		
Total for Period	CI	21.8	36.9		

2. Iodines

I-131	CI	1.99E-03	4.93E-04		
I-133	CI	4.33E-03	3.02E-03		
I-135	CI	4.38E-03	4.61E-03		
Total for Period	CI	8.81E-03	8.17E-03		

Table 1.1-1 (continued)

QCP 100-S25
Revision 3Reactor Vents
GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter Third	Quarter Fourth	Quarter	Quarter

1. Fission gases

Kr-85	CI	<LLD	<LLD		
Kr-85m	CI	<LLD	<LLD		
Kr-87	CI	<LLD	<LLD		
Kr-88	CI	<LLD	<LLD		
Xe-133	CI	5.4E-01	9.0E-02		
Xe-135	CI	9.8E00	8.7E00		
Xe-135m	CI	<LLD	<LLD		
Xe-138	CI	3.2E00	<LLD		
	CI				
	CI				
Total for Period	CI	1.4E01	8.8E00		

2. Iodines

I-131	CI	6.5E-04	8.4E-04		
I-133	CI	2.5E-03	2.9E-03		
I-135	CI	5.5E-03	5.4E-03		
Total for Period	CI	8.7E-03	9.1E-03		

Table 1.1-1 (continued)

QCP 100-S25
Revision 3

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter First	Quarter Second	Quarter ---	Quarter ---
3. Particulates					
Reactor Vents					
Sr-89 *	CI	5.38E-05	3.47E-04		
Sr-90	CI	4.53E-05	3.48E-05		
Cs-134	CI	7.5E-05	5.03E-05		
Cs-137	CI	3.26E-04	1.06E-03		
Ba-140	CI	3.42E-04	4.45E-04		
La-140	CI	<8 Day	Half Line		
Cr-51	CI	5.90E-04	8.78E-04		
Mn-54	CI	<LLD	3.7E-05		
Co-58	CI	<LLD	3.6E-05		
Co-60	CI	2.97E-03	2.01E-03		
I-131	CI	1.36E-04	1.44E-04		
Ag-110m	CI	<LLD	4.56E-05		
I-133	CI	1.80E-03	1.54E-03		
I-135	CI	1.76E-03	7.65E-03		
Ru-103	CI	1.87E-04	<LLD		
	CI	---	---		
	CI	---	---		
Unidentified	CI	---	---		

* Sr-89, Sr-90 values for May and June are projected

Table 1.i-1 (continued)

QCP 100-S25
Revision 3

GASEOUS EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter Third	Quarter Fourth	Quarter	Quarter
3. Particulates					
SR-89	CI	4.0E-04	2.5E-04		
SR-90	CI	3.2E-05	5.1E-06		
Cs-134	CI	3.8E-06	2.7E-05		
Cs-137	CI	4.4E-04	5.0E-04		
Ba-140	CI	1.2E-04	2.5E-05		
La-140	CI	<LLD	<LLD		
Cr-51	CI	4.8E-04	1.3E-03		
Mn-54	CI	4.9E-05	1.3E-05		
Co-58	CI	5.5E-05	8.4E-06		
Co-60	CI	1.8E-03	2.1E-03		
I-131	CI	1.1E-04	1.9E-04		
Ag-110m	CI	<LLD	3.3E-05		
I-133	CI	1.1E-03	2.5E-03		
I-135	CI	2.6E-03	2.3E-03		
Ru103	CI	1.1E-04	6.8E-05		
As-76	CI	<LLD	1.4E-05		
	CI				
Unidentified	CI				
TOTAL	CI	7.3E-03	9.3E-03		

Table 1.2-1

QCP 100-S25
Revision 3

APPROVED

LIQUID EFFLUENTS - SUMMATION OF ALL RELEASES

JUN 03 1985

O.C.O.S.R.

	Unit	Quarter First	Quarter Second	Est. Total Error, %
A. FISSION & ACTIVATION PRODUCTS				
1. Total release (not including tritium, gases, alpha)	CI	4.4E-03	3.8E-02	
2. Average diluted concentration during batch discharges period	$\mu\text{Ci/ml}$	2.2E-09	1.7E-08	
3. Percent of applicable limit *	%	.018 .024	.086 .079	
4. Maximum diluted concentration during batch discharges	$\mu\text{Ci/ml}$	3.2E-09	5.6E-09	
B. TRITIUM				
1. Total Release	CI	.86	1.35	
2. Average diluted concentration during batch discharges	$\mu\text{Ci/ml}$	4.4E-07	6.0E-07	
3. Percent of applicable limit	%	1.5E-02	2.0E-02	
C. DISSOLVED AND ENTRAINED GASES				
1. Total release	CI	3.42E-04	2.18E-04	
2. Average diluted concentration during batch discharges	$\mu\text{Ci/ml}$	1.7E-10	9.6E-11	
3. Percent of applicable limit	%	5.4E-05	1.9E-05	
D. GROSS ALPHA RADIOACTIVITY				
1. Total Release	CI	<LLD	<LLD	
2. Average concentration released during batch discharges	$\mu\text{Ci/ml}$	---	---	
E. VOLUME OF WASTE RELEASED (prior to dilution)				
	Liters	4.8E+05	5.3E+05	
F. VOLUME OF DILUTION WATER USED DURING BATCH DISCHARGES				
	Liters	1.96E+09	2.26E+09	
G. TOTAL VOLUME OF DILUTION WATER DURING PERIOD (QUARTER)				
	Liters	2.8E+11	2.0E+11	

*WHOLE BODY ORGAN

Table 1.2-1 (continued)

OCP 100-S25
Revision 3

APPROVED

LIQUID EFFLUENTS - SUMMATION OF ALL RELEASES

JUN 03 1985

O.C.O.S.R.

	Unit	Quarter Third	Quarter Fourth	Est. Total Error, %
A. FISSION & ACTIVATION PRODUCTS				
1. Total release (not including tritium, gases, alpha)	Cl	1.6E-02	1.4E00	
2. Average diluted concentration during batch discharges period	$\mu\text{Cl/ml}$	6.2E-09	4.0E-07	
3. Percent of applicable limit *	%	1.5E-1 1.7E-1	3.7E01 1.5E01	
4. Maximum diluted concentration during batch discharges	$\mu\text{Cl/ml}$	1.10E-08	1.0E-06	
B. TRITIUM				
1. Total Release	Cl	3.9E-01	8.1E-01	
2. Average diluted concentration during batch discharges	$\mu\text{Cl/ml}$	1.5E-07	2.3E-07	
3. Percent of applicable limit	%	5.0E-03	7.7E-03	
C. DISSOLVED AND ENTRAINED GASES				
1. Total release	Cl	2.3E-05	1.7E-02	
2. Average diluted concentration during batch discharges	$\mu\text{Cl/ml}$	8.8E-12	4.9E-09	
3. Percent of applicable limit	%	1.5E-06	2.4E-03	
D. GROSS ALPHA RADIOACTIVITY				
1. Total Release	Cl	<LLD	1.5E-05	
2. Average concentration released during batch discharges	$\mu\text{Cl/ml}$	---	4.3E-12	
E. VOLUME OF WASTE RELEASED (prior to dilution)				
	Liters	3.1E05	4.94E05	
F. VOLUME OF DILUTION WATER USED DURING BATCH DISCHARGES				
	Liters	2.6E09	3.5E09	
G. TOTAL VOLUME OF DILUTION WATER DURING PERIOD (QUARTER)				
	Liters	5.02E11	3.70E11	

Table 1.2-1 (continued)

OCP 100-S25
Revision J

LIQUID EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter First	Quarter Second	Quarter	Quarter
Sr-89 *	CI			5.14E-05	3.17E-05
Sr-90 *	CI			8.03E-06	2.88E-02
Cs-134	CI			4.58E-05	3.68E-05
Cs-137	CI			7.70E-04	1.17E-03
I-131	CI			8.30E-06	5.91E-05
Co-58	CI			4.50E-05	1.82E-04
Co-60	CI			1.99E-03	5.17E-03
Fe-59	CI			<LLD	6.43E-05
Zn-65	CI			6.40E-05	6.11E-05
Mn-54	CI			1.92E-04	3.63E-04
Cr-51	CI			7.60E-04	1.33E-03
Zr-95	CI			<LLD	<LLD
Nb-95	CI			<LLD	<LLD
Mo-99	CI			<LLD	<LLD
Ag-110m	CI			<LLD	3.60E-04
Ba-104	CI			<LLD	<LLD
Cs-136	CI			<LLD	<LLD
La-140	CI			<8 Day	Half Line
Fe-55*	CI			4.18E-04	7.57E-04
Unidentified	CI			---	---
Total for Period (above)	CI			4.35E-03	3.84E-02
Xe-133	CI			1.95E-04	1.98E-04
Xe-135	CI			1.47E-04	1.98E-05

Prepared by Robert J. LyonsApproved by Joseph S. [Signature]

APPROVED

Rad-Cehm Supervisor

*Sr-89, Sr-90, Fe-55 for May and June are projected

Table 1.2-1 (continued)

QCP 100-S25
Revision 3

LIQUID EFFLUENTS

Nuclides Released	Unit	Continuous Mode		Batch Mode	
		Quarter Third	Quarter Fourth	Quarter Third	Quarter Fourth
SR-89	C1	<LLD	<LLD ^a	4.3E-05	6.0E-02
SR-90	C1	<LLD	<LLD ^a	1.4E-02	2.2E-02
Cs-134	C1	<LLD	<LLD	2.9E-05	1.7E-01
Cs-137	C1	<LLD	<LLD	4.8E-04	1.1E00
I-131	C1	<LLD	<LLD	1.3E-05	1.2E-03
Co-58	C1	<LLD	<LLD	4.4E-05	6.5E-04
Co-60	C1	<LLD	<LLD	1.2E-03	3.1E-02
Fe-59	C1	<LLD	<LLD	<LLD	<LLD
Zn-65	C1	<LLD	<LLD	<LLD	3.9E-05
Mn-54	C1	<LLD	<LLD	8.6E-05	1.2E-03
Cr-51	C1	<LLD	<LLD	4.1E-05	2.7E-03
Zr-95	C1	<LLD	<LLD	<LLD	<LLD
Nb-95	C1	<LLD	<LLD	<LLD	<LLD
Mo-99	C1	<LLD	<LLD	<LLD	<LLD
Ag-110m	C1	<LLD	<LLD	5.2E-05	8.4E-05
Ba-140	C1	<LLD	<LLD	<LLD	<LLD
Cs-136	C1	<LLD	<LLD	<LLD	<LLD
La-140	C1	<LLD	<LLD	<LLD	<LLD
Fe55	C1	<LLD	<LLD ^a	3.5E-04	2.0E-02
I-133	C1	<LLD	<LLD	3.3E-06	<LLD
Total for Period (above)	C1	<LLD	<LLD	1.6E-02	1.4E00
Xe-133	C1	<LLD	<LLD	2.3E-05	<LLD
Xe-135	C1	<LLD	<LLD	<LLD	1.7E-02

Prepared by

Robert Sizer
APPROVED

Approved by

Frank Sizer
Rad-Cem Supervisor

(Final)

JUN 05 1935

JUN 05 1935

^aProjected Values Based on Prior Analysis

Table 2.0-1

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

1/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
01/03/85	CN	BSC	178.00	8280.00
01/03/85	TRI-STATE	BSC	26.20	13183000.00
01/07/85	CN	BSC	178.00	7603.00
01/11/85	CN	BSC	178.00	8216.00
01/11/85	HACHE	USE	872.00	408.62
01/12/85	TRI-STATE	BSC	26.20	11069000.00
01/14/85	CN	BSC	178.00	10390.00
01/16/85	CN	BSC	178.00	9894.00
01/16/85	TRI-STATE	BSC	26.20	10457000.00
01/18/85	CN	BSC	178.00	9549.00
01/19/85	TRI-STATE	BSC	26.20	10616000.00
01/21/85	CN	BSC	178.00	9566.00
01/23/85	CN	BSC	105.00	81.80
01/23/85	CN	BSC	178.00	7275.00
01/23/85	HACHE	USE	999.50	151.82
01/23/85	TRI-STATE	BSC	26.20	7482000.00
01/24/85	CN	USE	105.00	937.00
01/30/85	CN	BSC	178.00	9986.00

MONTHLY TOTALS

3814.50

52889338.24

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

ROAD-CITIES STATION

2/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	HILLCURIES
-----	-----	-----	-----	-----
02/04/85	CN	BSC	178.00	8875.00
02/08/85	CN	BSC	178.00	6718.60
02/11/85	CN	BSC	178.00	6241.00
02/13/85	CN	BSC	178.00	5928.60
02/21/85	CN	BSC	178.00	5076.00
02/22/85	HACKE	USE	1289.75	56.17
02/25/85	CN	BSC	178.00	4736.10
02/27/85	CN	BSC	178.00	5689.60
02/28/85	HACKE	USE	505.00	451.60

MONTHLY TOTALS

3120.75

43772.67

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

3/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
03/04/85	CN	BSC	178.00	5000.00
03/06/85	CN	BSC	178.00	5339.00
03/08/85	CN	BSC	178.00	3096.00
03/13/85	CN	BSC	178.00	7441.00
03/14/85	HACKE	USE	1289.75	93.49
03/18/85	CN	BSC	178.00	19151.00
03/21/85	CN	BSC	178.00	13547.00
03/26/85	CN	BSC	178.00	7528.00
03/27/85	CN	BSC	178.00	10644.00
03/29/85	CN	USE	110.50	1256.90

MONTHLY TOTALS

2824.25

73096.39

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

4/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
-----	-----	-----	-----	-----
04/01/85	CN	BSC	178.00	9949.00
04/02/85	HACKE	USE	1289.75	412.50
04/03/85	CN	BSC	178.00	6380.10
04/10/85	CN	BSC	178.00	6111.00
04/11/85	HACKE	USE	570.00	500.21
04/15/85	CN	BSC	178.00	5298.70
04/17/85	CN	BSC	178.00	1551.00
04/19/85	HACKE	USE	1289.75	202.55
04/19/85	CN	USE	105.00	1747.80
04/22/85	CN	BSC	178.00	4113.00
04/29/85	CN	BSC	178.00	2648.00

MONTHLY TOTALS

4500.50

41913.86

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

5/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
05/02/85	HACKE	USE	532.50	653.80
05/02/85	CN	BSC	178.00	3668.00
05/06/85	CN	BSC	178.00	3708.10
05/08/85	CN	BSC	178.00	2965.00
05/08/85	HACKE	USE	1289.75	80.03
05/13/85	CN	BSC	178.00	10243.20
05/15/85	HACKE	USE	1289.75	106.58
05/16/85	CN	BSC	178.00	17762.00
05/20/85	CN	BSC	178.00	13265.00
05/23/85	CN	BSC	178.00	9893.00
05/30/85	CN	BSC	178.00	11069.00
05/31/85	HACKE	USE	1289.75	55.67
05/31/85	CN	USE	105.00	514.93

MONTHLY TOTALS

5930.75

73984.31

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

6/1985

DATE	CO TRANS	BURIAL SITE	VOLUME	HILLCURIES
-----	-----	-----	-----	-----
06/03/85	CN	BSC	178.00	8995.00
06/06/85	CN	BSC	178.00	11894.00
06/10/85	CN	BSC	178.00	10177.00
06/12/85	CN	BSC	178.00	7244.00
06/12/85	HACKE	USE	1289.75	81.88
06/17/85	CN	BSC	178.00	7529.00
06/19/85	CN	BSC	178.00	7533.00
06/20/85	HACKE	USE	728.50	352.19
06/24/85	CN	BSC	178.00	7600.00
06/26/85	CN	BSC	178.00	9017.40
06/27/85	HACKE	USE	1063.80	72.81

MONTHLY TOTALS

4506.05

70496.28

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

7/85

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
07/01/85	CN	BSC	178.00	5445.00
07/05/85	CN	BSC	178.00	5843.00
07/08/85	CN	BSC	178.00	5764.00
07/12/85	CN	BSC	178.00	5076.40
07/15/85	CN	BSC	178.00	5076.40
07/18/85	CN	BSC	178.00	2913.00
07/18/85	HACKE	USE	502.50	993.78
07/22/85	CN	BSC	178.00	4557.60
07/25/85	CN	BSC	178.00	7336.70
07/29/85	CN	BSC	178.00	8930.00
07/29/85	CN	BSC	178.00	7337.00
07/31/85	CN	BSC	178.00	10060.00

MONTHLY TOTALS

2460.50

69332.88

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

8/85

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
08/01/85	HACKE	USE	1041.25	131.78
08/05/85	CN	BSC	178.00	8884.80
08/08/85	CN	BSC	178.00	8334.00
08/12/85	CN	BSC	178.00	7967.00
08/15/85	CN	BSC	178.00	8115.00
08/16/85	HACKE	USE	1026.25	120.38
08/19/85	CN	BSC	178.00	6241.90
08/22/85	HACKE	USE	611.75	641.06
08/22/85	CN	BSC	178.00	10634.00
08/26/85	CN	BSC	178.00	11106.00
08/28/85	CN	BSC	178.00	10412.00
08/30/85	HACKE	USE	792.75	48.00
08/30/85	CN	USE	105.00	1476.67

MONTHLY TOTALS 5001.00 74112.09

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

9/85

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
09/03/85	CN	BSC	178.00	8226.00
09/06/85	CN	BSC	178.00	5580.50
09/06/85	HACKE	USE	845.25	433.51
09/09/85	CN	BSC	178.00	5984.40
09/12/85	CN	BSC	178.00	8482.20
09/12/85	HACKE	USE	770.25	50.84
09/16/85	CN	BSC	178.00	7855.00
09/18/85	CN	BSC	178.00	13558.00
09/20/85	CN	BSC	178.00	11190.00
09/23/85	CN	BSC	178.00	9949.00
09/26/85	HACKE	USE	1289.75	66.11
09/26/85	CN	BSC	178.00	8391.00

MONTHLY TOTALS

4507.25

79766.59

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

10/85

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLCURIES
-----	-----	-----	-----	-----
10/02/85	HACKE	USE	611.75	1619.80
10/03/85	CN	BSC	200.00	9379.00
10/07/85	CN	BSC	200.00	7067.00
10/09/85	CN	BSC	178.00	5677.00
10/11/85	CN	BSC	178.00	6679.00
10/15/85	CN	BSC	178.00	7026.30
10/18/85	CN	USE	105.00	1535.14
10/21/85	CN	BSC	178.00	6247.00
10/23/85	CN	BSC	178.00	8252.40
10/25/85	CN	BSC	178.00	8840.00
10/28/85	CN	BSC	178.00	7615.00
10/30/85	CN	BSC	178.00	9199.00

MONTHLY TOTALS

2540.75

79136.64

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

11/85

DATE	CO TRANS	BURIAL SITE	VOLUME	MILLICURIES
11/01/85	CN	BSC	200.00	10349.00
11/04/85	CN	BSC	126.00	159713.00
11/06/85	CN	BSC	101.00	121162.00
11/08/85	CN	BSC	101.00	121162.00
11/12/85	CN	BSC	126.00	159713.00
11/13/85	HACKE	USE	585.00	142.70
11/14/85	CN	BSC	126.00	321864.00
11/18/85	CN	BSC	101.00	244173.00
11/19/85	HACKE	USE	800.25	215.11
11/20/85	CN	BSC	101.00	244173.00
11/25/85	CN	BSC	200.00	63645.00
11/26/85	CN	BSC	200.00	40437.00

MONTHLY TOTALS

2767.25

1486748.81

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TPI STATE

Table 2.0-1 (continued)

SOLID RADIOACTIVE WASTE SUMMARY

UNITS 1/2

QUAD-CITIES STATION

12/85

DATE	CO TRANS	BURIAL SITE	VOLUME	HILLICUFIES
12/03/85	CN	BSC	200.00	32125.00
12/04/85	CN	BSC	200.00	35211.00
12/05/85	HACKE	USE	1093.75	183.09
12/06/85	CN	BSC	200.00	22304.00
12/09/85	CN	BSC	200.00	28589.00
12/11/85	HACKE	USE	502.50	1262.50
12/12/85	CN	BSC	200.00	30698.00
12/16/85	CN	BSC	178.00	35054.00
12/18/85	CN	BSC	200.00	47181.00
12/19/85	HACKE	USE	1289.75	243.12
12/20/85	CN	BSC	200.00	36577.00
12/23/85	CN	BSC	178.00	28116.00
12/27/85	CN	BSC	200.00	27975.00
12/30/85	CN	BSC	200.00	21594.00

MONTHLY TOTALS 5042.00 347112.71

USE - U.S. ECOLOGY
 BSC - BARNWELL SOUTH CAROLINA
 CN - CHEM NUCLEAR CO.
 TS - TRI STATE

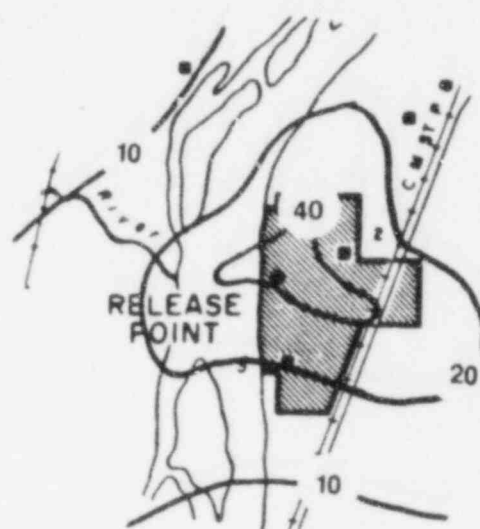
Figure 3.1-1

Estimated Cumulative Gamma Dose (mrem)
from the Quad Cities Station for the
period January-December 1985.

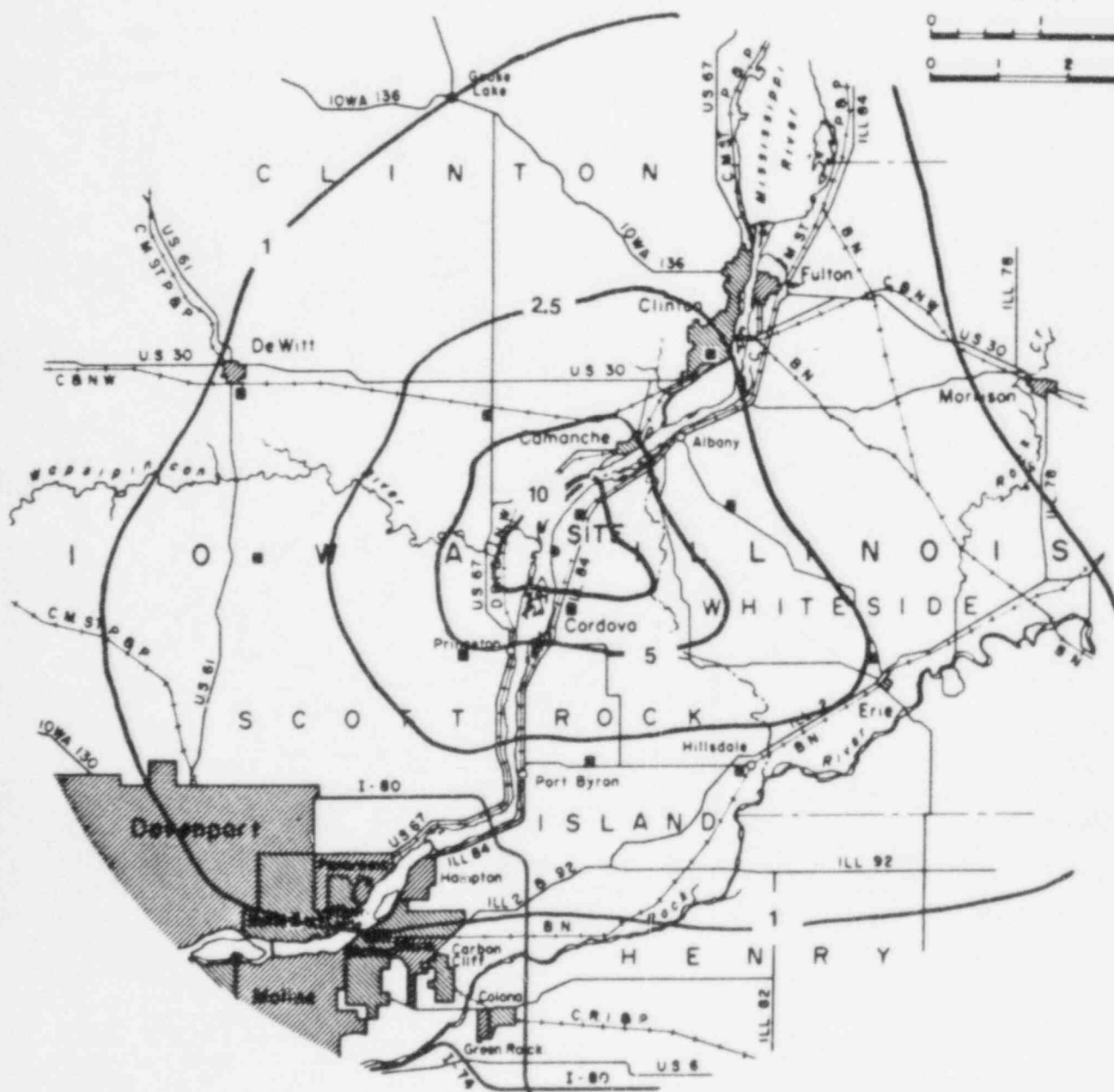
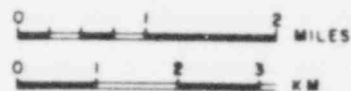
Isopleth Labels

Small figure - multiply by 10^{-3}

Large figure - multiply by 10^{-3}



SCALE



SCALE

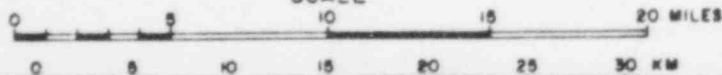


Figure 3.1-2

Estimated Total Concentration (pCi/m^3) of Noble Gases from the Quad Cities Station for the period January-December 1985.

Isopleth Labels

Small figure - multiply by 10^{-1}

Large figure - multiply by 10^{-1}

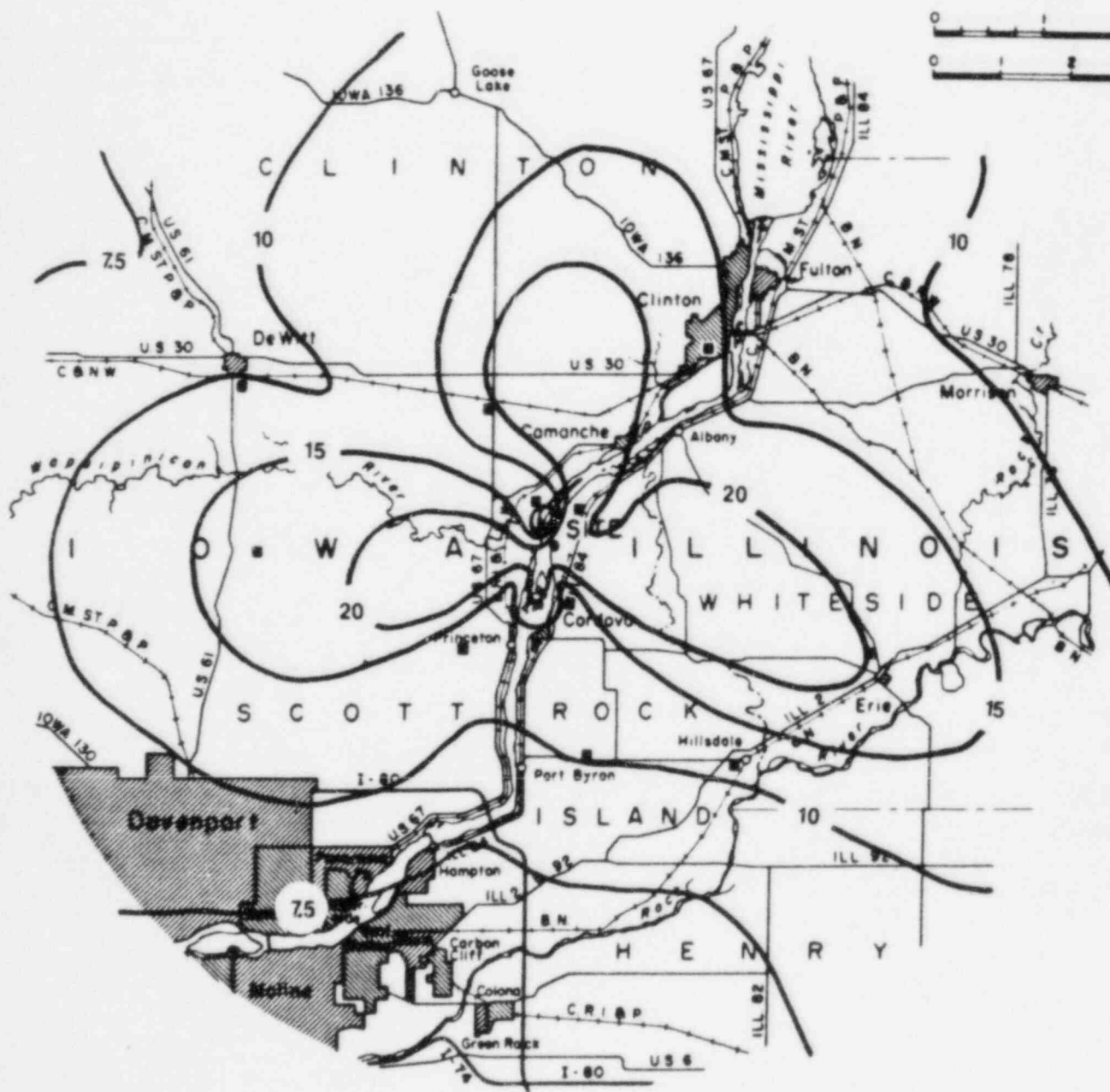
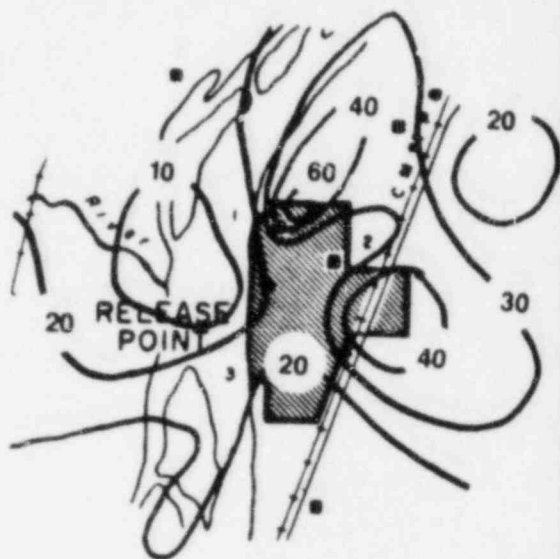


Figure 3.1-3

Estimated Total Concentration (pCi/m³) of Iodine from the Quad Cities Station for the period January-December 1985.

Isopleth Labels
 Small figure - multiply by 10⁻⁴
 Large figure - multiply by 10⁻⁴

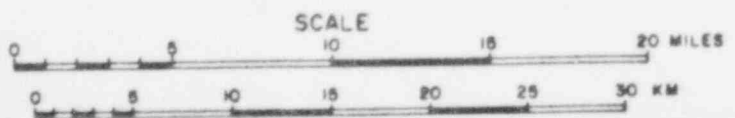
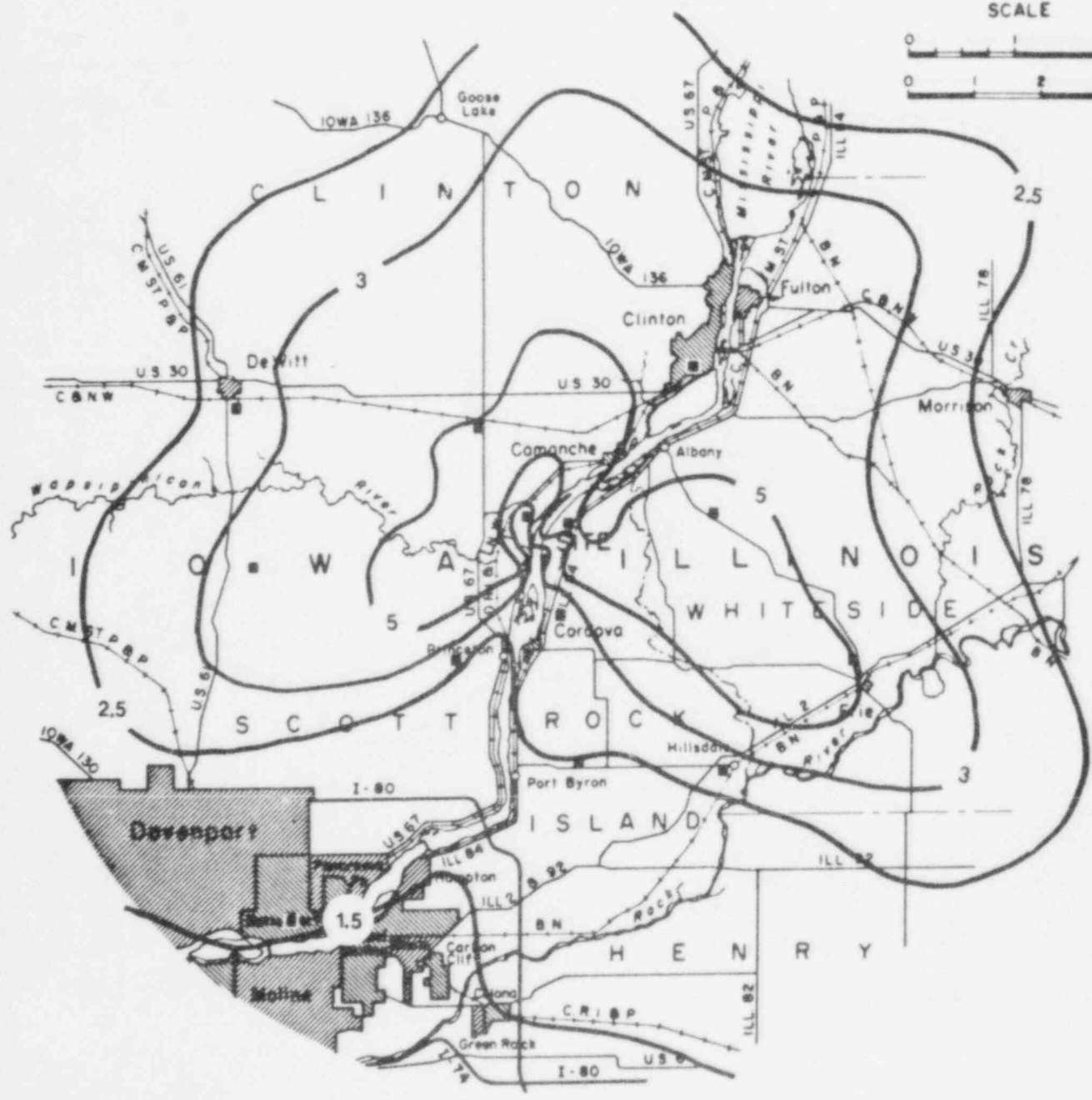
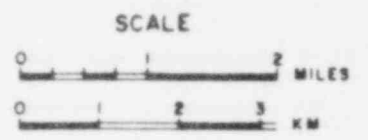
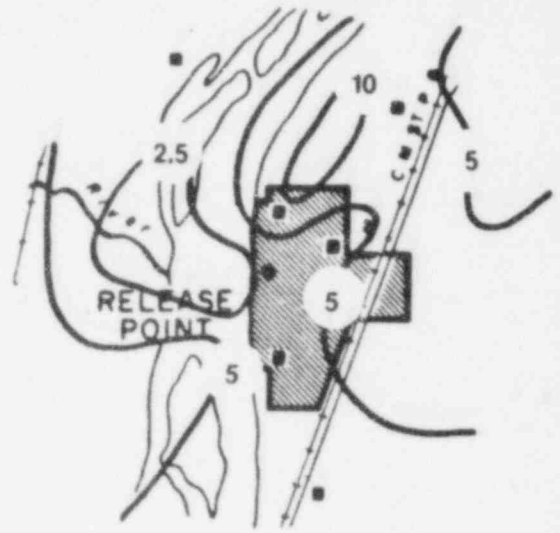


Figure 3.1-4

Estimated Total Concentration (pCi/m³) of
Particulate Matter from the Quad Cities Station
for the period January-December 1985.

Isopleth Labels

Small figure - multiply by 10⁻⁴

Large figure - multiply by 10⁻⁴

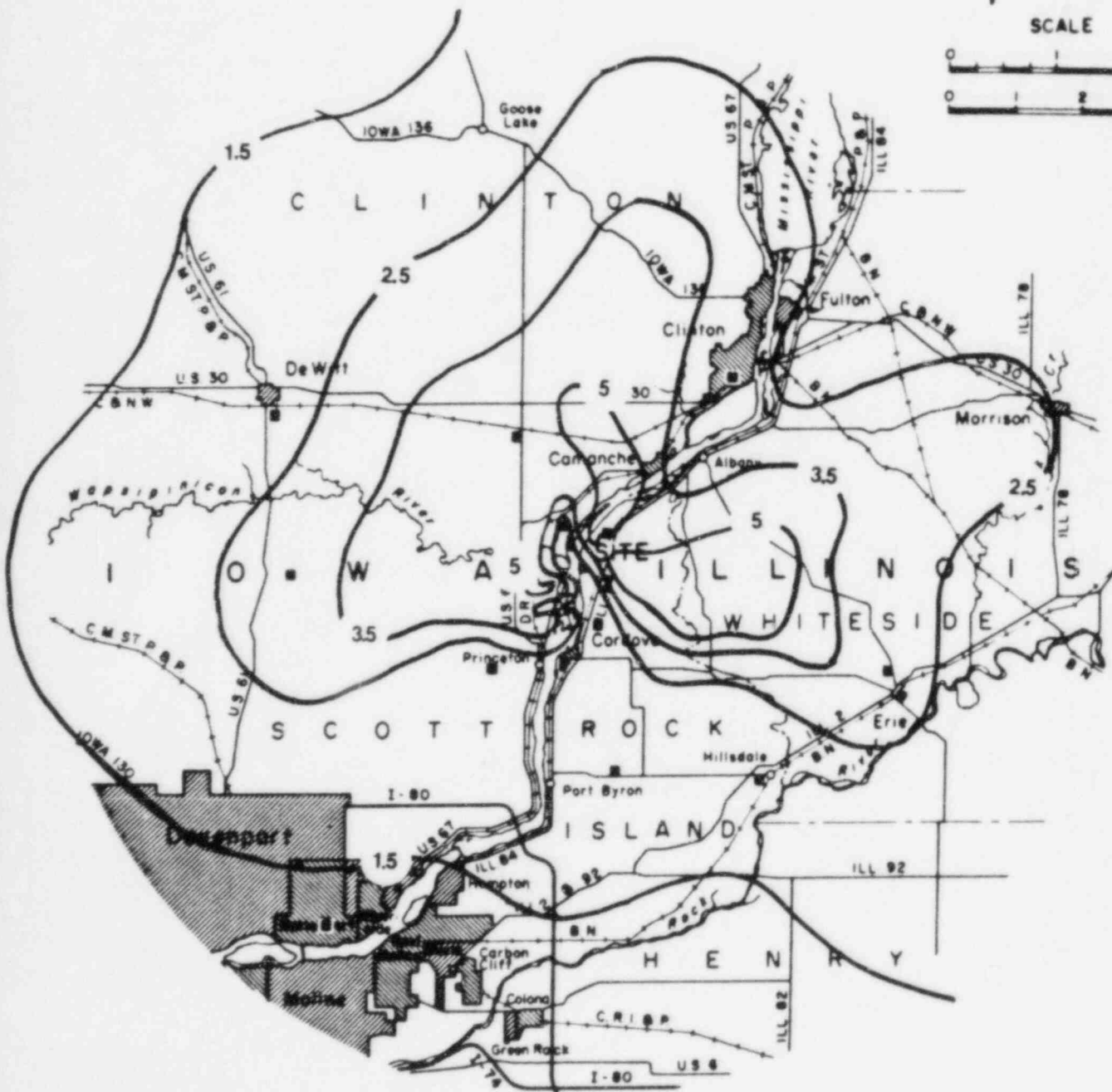
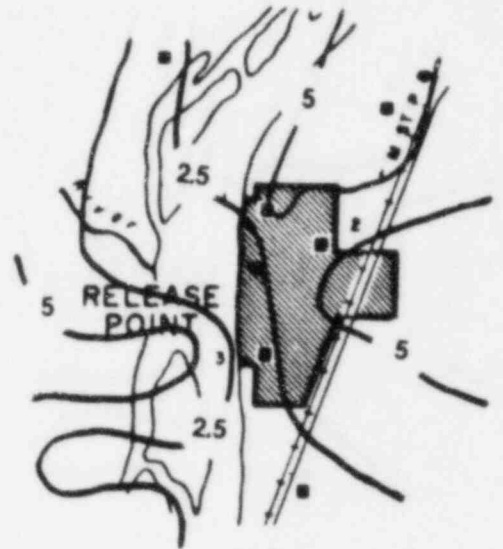


Table 3.1-1

WAOU CITIES UNIT ONE
 MAXIMUM DOSES RESULTING FROM AIRBORNE RELEASES
 PERIOD OF RELEASE - 1/ 1/85 TO 12/31/85 CALCULATED 02/06/86

TYPE	1ST QUARTER 1/85- 3/85	2ND QUARTER 4/85- 6/85	3RD QUARTER 7/85- 9/85	4TH QUARTER 10/85-12/85	ANNUAL
GAMMA AIR (MRAD)	7.44E-03 (MVM)	1.05E-02 (MVM)	5.39E-03 (M)	1.42E-02 (M)	3.78E-02 (M)
BETA AIR (MRAD)	6.24E-03 (NM)	1.02E-02 (VM)	3.33E-03 (NM)	5.19E-03 (NM)	2.49E-02 (NM)
TOT. BODY (MREM)	4.13E-03 (MVM)	5.39E-03 (M)	2.80E-03 (M)	7.63E-03 (M)	2.00E-02 (M)
SKIN (MREM)	1.04E-02 (MVM)	1.41E-02 (VM)	6.35E-03 (NM)	1.43E-02 (NM)	4.56E-02 (NM)
ORGAN (MREM)	6.61E-03 (MVM)	3.69E-02 (S)	9.24E-02 (S)	2.10E-02 (S)	1.56E-01 (S)
	THYROID	THYROID	THYROID	THYROID	THYROID

THIS IS A REPORT FOR THE CALENDAR YEAR 1985

COMPLIANCE STATUS - 10 CFR 50 APP. 1

	LIMIT	% OF APP 1					YRLY	2 OF APP. 1
		1ST QTR	2ND QTR	3RD QTR	4TH QTR	YRLY		
	OBJ	1/85- 3/85	4/85- 6/85	7/85- 9/85	10/85- 12/85	OBJ		
GAMMA AIR (MRAD)	5.0	0.16	0.21	0.11	0.25	10.0	0.33	
BETA AIR (MRAD)	10.0	0.06	0.10	0.03	0.03	20.0	0.12	
TOT. BODY (MREM)	2.0	0.17	0.22	0.11	0.31	5.0	0.44	
SKIN (MREM)	7.0	0.15	0.19	0.08	0.19	15.0	0.30	
ORGAN (MREM)	7.0	0.09	0.49	1.23	0.28	15.0	1.04	
		THYROID	THYROID	THYROID	THYROID		THYROID	

Table 3.1-1 (continued)

QUAD CITIES UNIT TWO
 MAXIMUM DOSES RESULTING FROM AIRBORNE RELEASES
 PERIOD OF RELEASE - 1/1/85 TO 12/31/85 CALCULATED 02/06/86

TYPE	1ST QUARTER 1/85- 3/85	2ND QUARTER 4/85- 6/85	3RD QUARTER 7/85- 9/85	4TH QUARTER 10/85-12/85	ANNUAL
GAMMA AIR (MRAD)	7.20E-03 (W)	9.93E-03 (W)	5.76E-03 (WNW)	1.41E-02 (W)	3.70E-02 (W)
BETA AIR (MRAD)	3.82E-03 (NW)	7.40E-03 (NW)	3.39E-03 (NW)	4.68E-03 (NW)	1.93E-02 (NW)
TOT. BODY (MREM)	3.87E-03 (W)	5.32E-03 (W)	2.96E-03 (W)	7.59E-03 (W)	1.97E-02 (W)
SKIN (MREM)	8.37E-03 (WNW)	1.40E-02 (NW)	6.92E-03 (WNW)	1.39E-02 (NW)	4.31E-02 (NW)
ORGAN (MREM)	5.78E-03 (S)	2.76E-02 (S)	8.31E-02 (S)	1.71E-02 (S)	1.33E-01 (S)
	LIVER	THYROID	THYROID	THYROID	THYROID

THIS IS A REPORT FOR THE CALENDAR YEAR 1985

COMPLIANCE STATUS - 10 CFR 50 APP. I

	DAILY	% OF APP. I.				YRLY	% OF APP. I
		1ST QTR 1/85- 3/85	2ND QTR 4/85- 6/85	3RD QTR 7/85- 9/85	4TH QTR 10/85- 12/85		
GAMMA AIR (MRAD)	5.0	0.14	0.20	0.12	0.28	10.0	0.37
BETA AIR (MRAD)	10.0	0.04	0.07	0.03	0.05	20.0	0.10
TOT. BODY (MREM)	2.5	0.15	0.21	0.12	0.30	5.0	0.34
SKIN (MREM)	7.5	0.11	0.19	0.09	0.18	15.0	0.29
ORGAN (MREM)	7.5	0.08	0.37	1.11	0.23	15.0	0.89
		LIVER	THYROID	THYROID	THYROID		THYROID

Table 3.2-1

DUAD CITIES UNIT ONE
 MAXIMUM DOSES (MREM) RESULTING FROM LIQUID EFFLUENTS
 PERIOD OF RELEASE - 1/ 1/85 TO 12/31/85 CALCULATED 02/06/86 *

DOSE TYPE	1ST QUARTER 1/85- 3/85	2ND QUARTER 4/85- 6/85	3RD QUARTER 7/85- 9/85	4TH QUARTER 10/85-12/85	ANNUAL
TOTAL BODY	2.25E-04	1.25E-03	2.23E-03	9.93E-01	9.97E-01
INTERNAL ORGAN	9.34E-04	3.88E-03	8.59E-03	1.45E+00	1.45E+00
	LIVER	BONE	BONE	LIVER	LIVER

* THIS IS A REPORT FOR THE CALENDAR YEAR 1985

COMPLIANCE STATUS - 10 CFR 50 APP. I

QTRLY OBJ	----- % OF APP. I. -----				YRLY OBJ	% OF APP. I	
	1ST QTR 1/85- 3/85	2ND QTR 4/85- 6/85	3RD QTR 7/85- 9/85	4TH QTR 10/85- 12/85			
TOTAL BODY (MREM)	1.5	0.02	0.08	0.15	66.21	3.0	33.23
CRIT. ORGAN (MREM)	5.0	0.02	0.08	0.17	28.91	10.0	14.48
	LIVER	BONE	BONE	LIVER	LIVER	LIVER	LIVER

Table 3.2-1 (continued)

DUAD CITIES UNIT TWO
 MAXIMUM DOSES (MREM) RESULTING FROM LIQUID EFFLUENTS
 PERIOD OF RELEASE - 1/ 1/85 TO 12/31/85 CALCULATED 02/06/86 *

DOSE TYPE	1ST	2ND	3RD	4TH	ANNUAL
	QUARTER 1/85- 3/85	QUARTER 4/85- 6/85	QUARTER 7/85- 9/85	QUARTER 10/85-12/85	
TOTAL	2.48E-04	1.32E-03	2.23E-03	1.23E-04	3.97E-03
BODY					
INTERNAL	1.51E-03	3.95E-03	8.59E-03	1.82E-04	1.39E-02
ORGAN	LIVER	BONE	BONE	LIVER	BONE

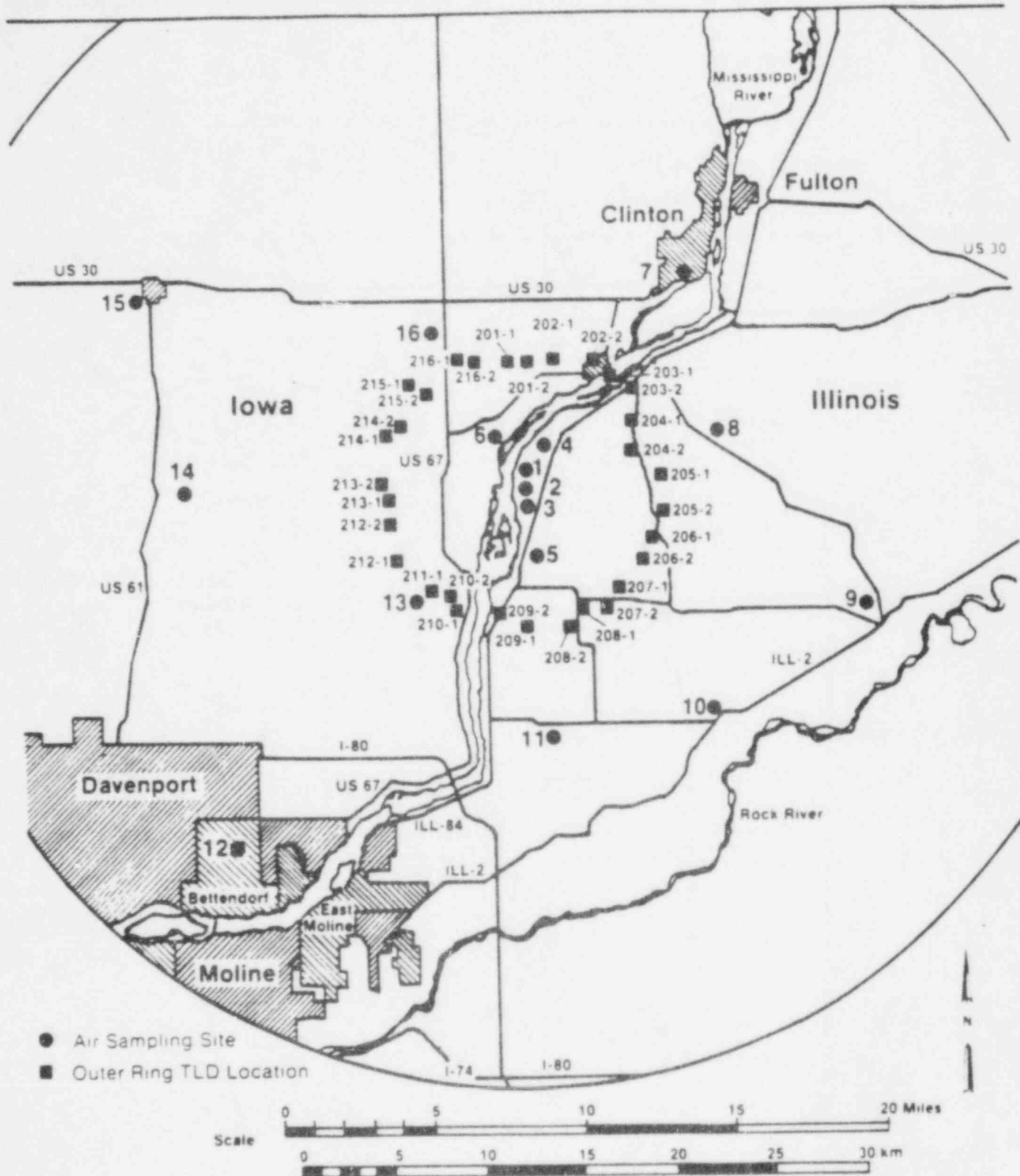
* THIS IS A REPORT FOR THE CALENDAR YEAR 1985

COMPLIANCE STATUS - 10 CFR 50 APP. I

	YRLY OBJ	% OF APP. I				YRLY OBJ	% OF APP. I
		1ST QTR 1/85- 3/85	2ND QTR 4/85- 6/85	3RD QTR 7/85- 9/85	4TH QTR 10/85- 12/85		
TOTAL BODY (MREM)	1.5	0.02	0.09	0.15	0.01	3.0	0.13
CRIT. ORGAN (MREM)	5.0	0.03	0.08	0.17	0.00	10.0	0.14
		LIVER	BONE	BONE	LIVER		BONE

Figure 5.0-1

DECEMBER 1985

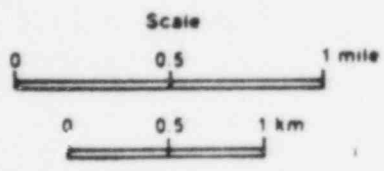
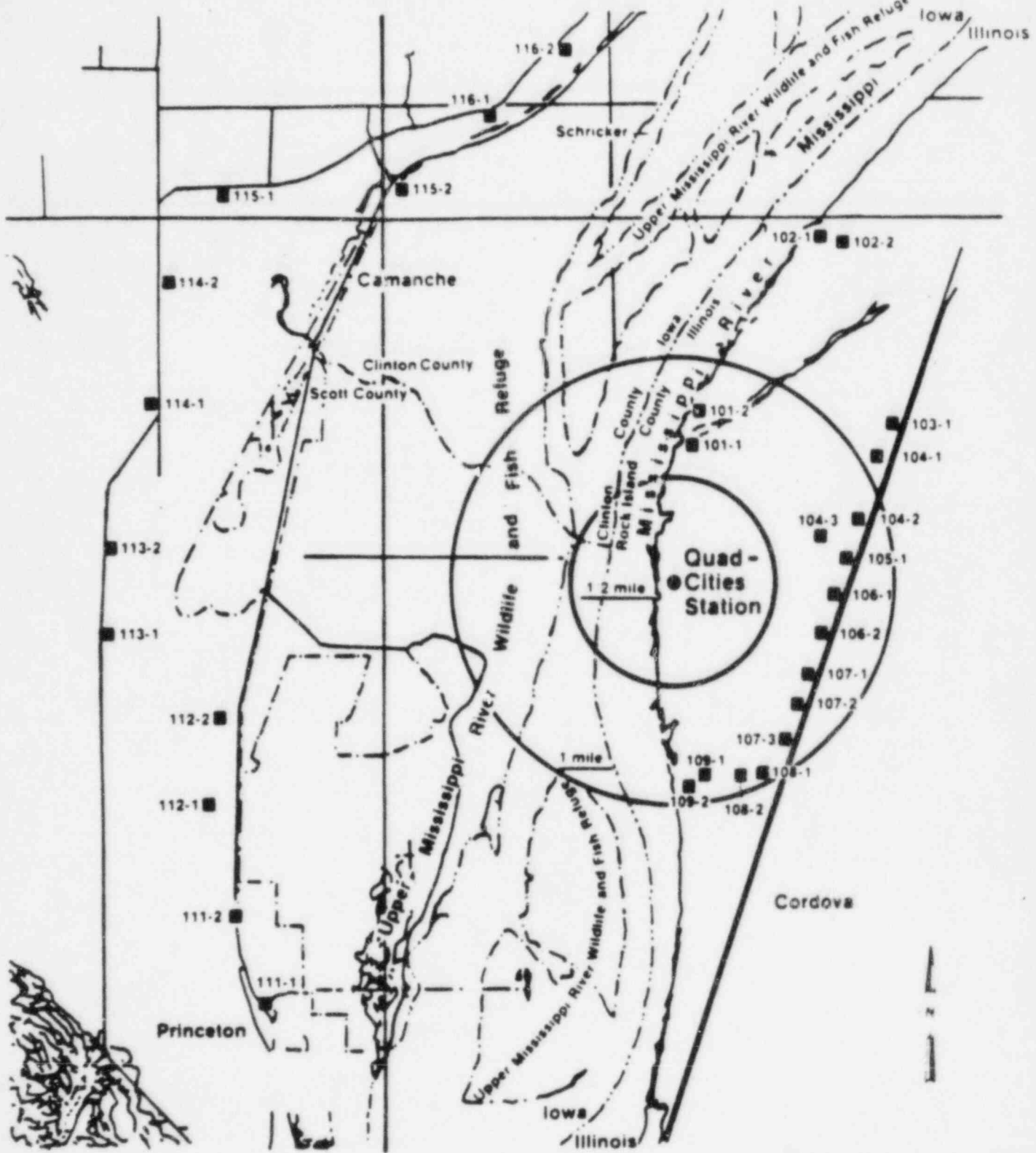


**QUAD-CITIES STATION
Units 1 & 2**

FIXED AIR SAMPLING SITES
AND OUTER RING TLD LOCATIONS

Figure 5.0-2

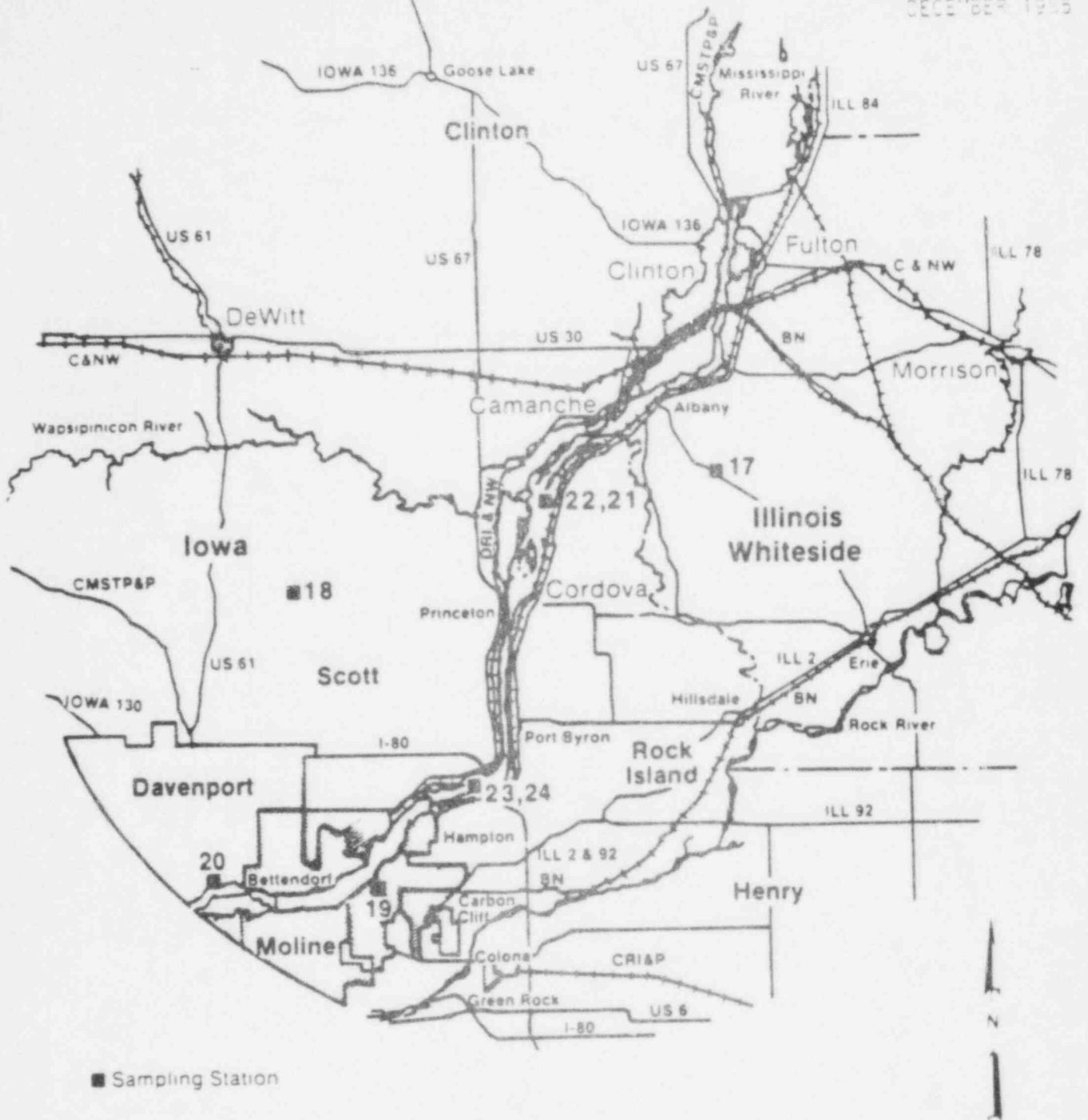
DECEMBER 1985



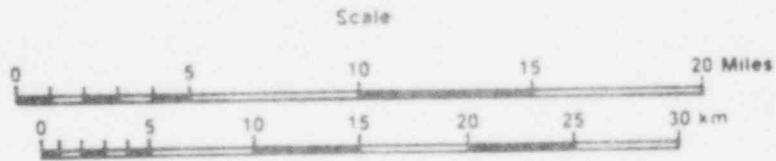
QUAD-CITIES STATION Units 1 & 2
INNER RING TLD LOCATIONS

Figure 5.0-3

DECEMBER 1995



■ Sampling Station



QUAD-CITIES STATION
Units 1 & 2

MILK, FISH, WATER, AND
 SEDIMENT SAMPLE LOCATIONS

Table 5.0-1
 QUAD CITIES NUCLEAR POWER STATION
 STANDARD RADIOLOGICAL SAMPLING PROGRAM

Location Code	Location Type ^a	Location Description	Media							
			Air Samples	TLDs	Milk	Public Water	Cooling Water	Sediment	Fish	Census
Q-01		On-site No. 1	X	X						
-02		On-site No. 2	X	X						
-03		On-site No. 3	X	X						
-04		Nitrin	X	X						
-05		Saddle Club	X	X						
-06		Hanson's Boat Landing	X	X						
-07		Clinton	X	X						
-08		Sikkema Farm	X	X						
-09	C	Erie	X	X						
-10	C	Hillsdale	X	X						
-11		Port Byron	X	X						
-12		Bettendorf	X	X						
-13		Princeton	X	X						
-14	C	Utica Ridge Road	X	X						
-15	C	Dewitt	X	X						
-16	C	Low Moor	X	X						
-17	C	Hansen Dairy Farm			X					X
-18		Musal Farm			X					X
-19		East Moline Water Works				X				
-20	C	Davenport Water Works				X				
-21	C	Inlet Canal					X			
-22		Discharge Canal					X			
-23		Lock and Dam No. 14 (Mississippi River)						X		
-24		Davenport Fish Market (Pool No. 14, Mississippi River)							X	

^a Control (background) locations are indicated by a "C" in this column. All other locations are indicators.

Table 5.0-2

QUAD CITIES STATION

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLING LOCATIONS

1. AIR SAMPLERS

<u>Site Code</u> ^a	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-01	a. On-site Station No. 1	0.5	0
Q-02	b. On-site Station No. 2	0.5	70
Q-03	c. On-site Station No. 3	0.6	170
Q-04	d. Nitrite No. 4	1.5	40
Q-05	e. Saddle Club Dairy Farm	1.8	160
Q-06	f. Hanson's Boat Landing	1.8	340
Q-07	g. Clinton	9.0	40
Q-08	h. Sikkema Farm	7.0	70
Q-09 (C)	i. Erie	13.0	110
Q-10 (C)	j. Hillsdale	10.0	130
Q-11	k. Port Byron	8.0	170
Q-12	l. Bettendorf	13.0	218
Q-13	m. Princeton	4.8	220
Q-14 (C)	n. Utica Ridge Road	11.0	270
Q-15 (C)	o. DeWitt	13.0	300
Q-16 (C)	p. Low Moor	6.0	330

2. TLDs

a. Same as No. 1.

b. Special TLD Samplers

<u>Revised Site Code</u> ^b	<u>Previous Site Code</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-101 1,2	Same	0.7	4
Q-102 1,2	Same	1.7	21
Q-103 1	Same	1.2	58
Q-104 1	Q-103 2	1.1	60
Q-104 2	Q-104 1	0.95	77
Q-104 3	Q-104 2	0.63	77
Q-105 1	Same	0.75	91

^a Control (reference) locations are denoted by a "C" after site code. All other locations are indicators.

^b Effective August 1, 1985.

Table 5.0-2 (continued)

QUAD CITIES STATION

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLING LOCATIONS

2. TLDsb. Special TLD Samplers (continued)

<u>Revised Site Code</u>	<u>Previous Site Code</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-106 1	Q-105 2	0.71	109
Q-106 2	Q-106 1	0.71	118
Q-107 1	Q-106 2	0.71	128
Q-107 2	Q-107 1	0.73	137
Q-107 3	Q-107-2	0.78	146
Q-108 1,2	Same	0.9	155
Q-109 1,2	Same	0.95	176
Q-111 1,2	Same	2.6	230
Q-112 1,2	Same	2.4	246
Q-113 1,2	Same	2.5	264
Q-114 1,2	Same	2.6	286
Q-115 1,2	Same	2.3	310
Q-116 1,2	Same	2.2	339
Q-201 1,2	Same	4.0	356
Q-202 1,2	Same	4.4	17
Q-203 1,2	Same	5.5	34
Q-204 1,2	Same	4.5	61
Q-205 1,2	Same	4.5	83
Q-206 1,2	Same	4.8	113
Q-207 1,2	Same	4.8	133
Q-208 1,2	Same	4.4	158
Q-209 1,2	Same	4.8	179
Q-210 1,2	Same	4.4	210
Q-211 1	Same	5.0	223
Q-212 1,2	Same	4.8	242
Q-213 1,2	Same	4.7	265
Q-214 1,2	Same	4.8	310
Q-215 1,2	Same	4.8	316
Q-216 1,2	Same	4.5	333

3. MILK

<u>Site Code^a</u>	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-17 (C)	a. Hansen Dairy Farm	6.0	70
Q-18	b. Musal Farm	5.5	225

^a Control (reference) locations are denoted by a "C" after site code. All other locations are indicators.

Table 5.0-2 (continued)

QUAD CITIES STATION
ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLING LOCATIONS

4. PUBLIC WATER SUPPLY

<u>Site Code</u> ^a	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-19	a. East Moline Water Works	16.0	206
Q-20 (C)	b. Davenport Water Works	18.0	219

5. COOLING WATER

<u>Site Code</u> ^a	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-21 (C)	a. Inlet	At Station	
Q-22	b. Discharge		

6. FISH

<u>Site Code</u> ^a	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-24	Davenport Fish Market	Pool No. 14 of Mississippi River	

7. BOTTOM SEDIMENTS

<u>Site Code</u> ^a	<u>Location</u>	<u>Distance (miles)</u>	<u>Direction (°)</u>
Q-23	Lock and Dam No. 14	15.0	210

^a Control (reference) locations are denoted by a "C" after site code. All other locations are indicators.

Table 5.0-2 (continued)

QUAD CITIES STATION
ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLE COLLECTION AND ANALYSES

Sample Media	Location		Collection Frequency	Type of Analysis	Frequency of Analysis	Remarks
	Code ^a	Site				
1. Airborne Particulates	a. Onsite and Near Field		Continuous operation for one week.	Gross beta	Weekly	On all samples. Gamma Isot if gross beta in a sample exceeds by 5X the average concentration of the preceding calendar quarter for the sample location. <u>Non-routine Reporting Levels^b</u> Cs-134 10 Cs-137 20 pCi/m ³ .
	Q-1	Onsite No. 1				
	Q-2	Onsite No. 2				
	Q-3	Onsite No. 3				
	Q-4	Nitroin				
	Q-5	Saddle Club Dairy Farm				
	Q-6	Hanson's Boat Landing	Same as 1a.	Filter Exchange	Weekly	<u>Non-routine Reporting Levels^b</u> Same as 1(a) when analyses are made.
	b. Far Field					
	Q-7	Clinton				
	Q-8	Sikkoma Farm				
	Q-9 (C)	Erie				
	Q-10 (C)	Hillsdale				
	Q-11	Port Byron				
	Q-12 (C)	Bettendorf				
	Q-13	Princeton				
	Q-14 (C)	Utica Ridge Road				
Q-15 (C)	DeWitt					
Q-16	Low Moor					
2. Airborne Iodine	Same as 1.		Continuous operation for two weeks	I-131	Biweekly	Biweekly = Every two weeks. On all samples. <u>Non-routine Reporting Level^b</u> 0.9 pCi/m ³
3. Air Sampling Train	Same as 1.		--	Test and Maintenance	Weekly	On all samplers.

^a Control (reference) locations are denoted by a "C" in this column. All other locations are indicators.

^b Average concentration over calendar quarter.

Table 5.0-2 (continued)

QUAD CITIES STATION
ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLE COLLECTION AND ANALYSES

Sample Media	Location		Collection Frequency	Type of Analysis	Frequency of Analysis	Remarks
	Code ^a	Site				
4. TLD	Q-101 1,2	Inner Ring	Quarterly	Gamma	Quarterly	Two sets at all AP locations. One set read quarterly. Second set read if required by Commonwealth Edison. At other locations, all sets read quarterly. Minimum of two TLDs per set.
	102 1,2					
103 1						
104 1,2,3						
105 1						
106 1,2						
107 1,2,3						
108 1,2						
109 1,2						
111 1,2						
through						
116 1,2	Outer Ring					
L-201 1,2						
through						
210 1,2						
211						
212 1,2						
through						
215 1,2						
5. Milk	Q-17 (C)	Hansen Dairy Musal Farm	Weekly: May through October	I-131	Weekly: May through October	On all samples. LLD: 0.5 pCi/l.
	Q-18		Monthly: November through April	I-131	Monthly: November through April	On all samples. LLD: 5.0 pCi/l.
						<u>Non-routine Reporting Levels^b</u> I-131 3; Cs-134 60; Cs-137 70; Ba-La-140 300 pCi/l

^a Control (reference) locations are denoted by a "C" in this column. All other locations are indicators.
^b Average concentration over calendar year.

Table 5.0-2 (continued)

QUAD CITIES STATION
ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLE COLLECTION AND ANALYSES

Sample Media	Location		Collection Frequency	Type of Analysis	Frequency of Analysis	Remarks
	Code ^a	Site				
6. Public Water	Q-19 Q-20 (C)	East Moline Water Works Davenport Water Works	Weekly	Gamma Isot	Monthly	On monthly composite from each location. <u>Non-routine Reporting Levels^b</u> (See footnote "c.")
7. Cooling Water	Q-21 (C) Q-22	Inlet Canal Discharge Canal	Weekly	Gross beta	Weekly	On notification provided by station personnel.
8. Fish	Q-24	Pool 14 of Mississippi River	Semiannually	Gamma isot	Semiannually	On edible portions only. At least two species. <u>Non-routine Reporting Levels^b</u> Mn-54 3×10^4 ; Fe-59 1×10^4 ; Co-58 3×10^4 ; Co-60 1×10^4 ; Zn-65 2×10^4 ; Cs-134 1×10^3 ; Cs-137 1×10^3 pCi/kg wet weight.
9. Bottom Sediments	Q-23	Lock and Dam No. 14	Annually	Gamma Isot	Annually	

^a Control (reference) locations are denoted by a "C" in this column. All other locations are indicators.

^b Average concentration over calendar year.

^c H-3 2×10^4 , Mn-54 1×10^3 , Fe-59 4×10^2 , Co-58 1×10^2 , Co-60 3×10^2 , Zn-65 3×10^2 , Zr-Nb-95 4×10^2 , I-131 2, Cs-134 30, Cs-137 50, Ba-La-140 2×10^2 pCi/l.

Table 5.0-2 (continued)

QUAD CITIES STATION

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM, SAMPLE COLLECTION AND ANALYSES (continued)

Sample Media	Location		Collection Frequency	Type of Analysis	Frequency of Analysis	Remarks
	Code	Site				
10. Dairy Census	a.	Site boundary to 2 miles	--	a. Enumeration by a door-to-door or equivalent counting technique.	Annually	During grazing season.
	b.	2 miles to 5 miles	--	b. Enumeration by using referenced information from county agricultural agents or other reliable sources.	Annually	During grazing season.
	c.	At dairies listed in Item 5.	--	c. Inquire as to feeding practices: (1) Pasture only. (2) Feed and chop only. (3) Pasture and feed; if both, ask farmer to estimate fraction of food from pasture: <25%, 25-50%, 50-75%, or >75%.	Annually	During grazing season.
11. Nearest Residence Census	In all 16 sectors				Annually	

Table 5.0-3

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM QUARTERLY SUMMARY

Name of Facility Quad Cities Nuclear Power Station Docket No. 50-254, 50-265
 Location of Facility Rock Island, Illinois Reporting Period 1st Quarter 1985
 (County, State)

Sample Type (Units)	Type and Number of Analyses	LLD	Indicator Locations Mean ^a Range	Location with Highest Quarterly Mean		Control Locations Mean ^a Range	Number of Non-routine Results
				Location	Mean Range		
Air Particulates (pCi/m ³)	Gross Beta 77	0.01	0.025 (77/77) (0.009-0.056)	Q-05, Saddle Club Dairy, 1.8 mi @ 160°	0.027 (13/13) (0.009-0.055)	None	0
Airborne Iodine (pCi/m ³)	I-131 36	0.10	<LLD	-	-	None	0
Gamma Background (TLDs) (mR/Qtr.)	Gamma Dose 16	3.0	12.0 (6/6) (9.3-15.5)	Q-10, Hillsdale 10.0 mi @ 130°	30.0 (1/1) -	14.3 (10/10) (4.6-30.0)	0
Milk (pCi/l)	I-131 6	5.0	<LLD	-	-	None	0
Cooling Water (pCi/l)	Gross Beta 26	2.0	4.2 (13/13) (3.1-6.1)	Q-22A, Diffuser Pipe Blowdown at Station	4.4 (13/13) (3.1-6.1)	4.2 (13/13) (2.9-5.5)	0
	Tritium 1	200	240 (1/1)		240 (1/1)	None	0
Public Water (pCi/l)	Gamma Spec. 6						
	Cs-134 10	10.0	<LLD	-	-	None	0
	Cs-137 10	10.0	<LLD	-	-	None	0
	Other Gammas 20.0		<LLD	-	-	None	0

^a Mean and range based on detectable measurements only. Fractions indicated in parentheses.

Table 5.0-4

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM QUARTERLY SUMMARY

Name of Facility Quad Cities Nuclear Power Station Docket No. 50-254, 50-265
 Location of Facility Rock Island, Illinois Reporting Period 2nd Quarter 1985
 (County, State)

Sample Type (Units)	Type and Number of Analyses	LLD	Indicator Locations Mean ^a Range	Location with Highest Quarterly Mean		Control Locations Mean ^a Range	Number of Non-routine Results
				Location	Mean Range		
Air Particulates (pCi/m ³)	Gross Beta 77	0.01	0.019 (75/77) (0.008-0.071)	Q-04, Nitrin 1.5 mi @ 40*	0.029 (11/12) (0.015-0.073)	None	0
Airborne Iodine (pCi/m ³)	I-131 42	0.10	<LLD	-	-	None	0
Gamma Background (TLDs) (mR/Qtr.)	Gamma Dose 16	3.0	11.6 (6/6) (10.3-13.9)	Q-10, Hillsdale 10.0 mi @ 130* Q-12, Bettendorf 13.0 mi @ 218*	16.5 (1/1) - 16.5 (1/1) -	11.9 (10/10) (8.1-16.5)	0
Milk (pCi/l)	I-131 20	5/0.5 ^b	<LLD	-	-	None	0
Cooling Water (pCi/l)	Gross Beta 26	1.0	4.5 (13/13) (3.5-7.5)	Q-21, Inlet Canal at Station	4.9 (13/13) (3.4-6.2)	4.5 (13/13) (3.9-5.3)	0
	Tritium 1	200	<LLD	-	-	None	0
Public Water (pCi/l)	Gamma Spec. 6						
	Cs-134 10	10.0	<LLD	-	-	None	0
	Cs-137 10	10.0	<LLD	-	-	None	0
	Other Gammas	20.0	<LLD	-	-	None	0
Fish (pCi/g wet)	Gamma Spec. 4						
	Cs-134	0.1	<LLD	-	-	None	0
	Cs-137	0.1	<LLD	-	-	None	0
	Other Gammas	0.2	<LLD	-	-	None	0

^a Mean and range based on detectable measurements only. Fractions indicated in parentheses.
^b November - April LLD = 5.0; May - October LLD = 0.5.

Table 5.0-5

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM QUARTERLY SUMMARY

Name of Facility Quad Cities Nuclear Power Station Docket No. 50-254, 50-265
 Location of Facility Rock Island, Illinois Reporting Period 3rd Quarter 1985
 (County, State)

Sample Type (Units)	Type and Number of Analyses	LLD	Indicator Locations Mean ^a Range	Location with Highest Quarterly Mean		Control Locations Mean ^a Range	Number of Non-routine Results
				Location	Mean Range		
Air Particulates (pCi/m ³)	Gross Beta 78	0.01	0.023 (78/78) (0.006-0.039)	Q-05, Saddle Club Dairy 1.8 mi @ 160° Q-06, Hanson's Boat Dock 1.8 mi @ 340°	0.025 (12/12) (0.015-0.039) 0.025 (12/12) (0.016-0.035)	None	0
Airborne Iodine (pCi/m ³)	I-131 36	0.10	<LLD	-	-	None	0
Gamma Background (TLDs) (mR/Qtr.)	Gamma Dose 16	3.0	10.6 (6/6) (7.0-13.0)	Q-11, Port Byron 8.0 mi @ 170°	14.3 (1/1) -	9.9 (10/10) (7.6-14.3)	0
Milk (pCi/l)	I-131 20	5/0.5 ^b	<LLD	-	-	None	0
Cooling Water (pCi/l)	Gross Beta 26	1.0	3.4 (13/13) (3.0-4.4)	Q-22A, Diffuser Pipe Blowdown at Station	3.6 (13/13)	3.6 (13/13) (3.1-4.0)	0
	Tritium 1	200	<LLD	-	-	None	0
Public Water (pCi/l)	Gamma Spec. 6						
	Cs-134	10.0	<LLD	-	-	None	0
	Cs-137	10.0	<LLD	-	-	None	0
	Other Gammas	20.0	<LLD	-	-	None	0
Bottom Sediments (pCi/g dry)	Gamma Spec. 1						
	Cs-134	0.1	<LLD	-	-	None	0
	Cs-137	0.1	<LLD	-	-	None	0
	Other Gammas	0.2	<LLD	-	-	None	0

^a Mean and range based on detectable measurements only. Fractions indicated in parentheses.

Table 5.0-6

ENVIRONMENTAL RADIOLOGICAL MONITORING PROGRAM QUARTERLY SUMMARY

Name of Facility Quad Cities Nuclear Power Station Docket No. 50-254, 50-265
 Location of Facility Rock Island, Illinois Reporting Period 4th Quarter 1985
 (County, State)

Sample Type (Units)	Type and Number of Analyses	LLD	Indicator Locations Mean ^a Range	Location with Highest Quarterly Mean		Control Locations Mean ^a Range	Number of Non-routine Results
				Location	Mean Range		
Air Particulates (pCi/m ³)	Gross Beta 75	0.01	0.027 (75/75) (0.006-0.068)	Q-05, Saddle Club Dairy 1.8 mi @ 160°	0.029 (14/14) (0.014-0.061)	None	0
Airborne Iodine (pCi/m ³)	I-131 39	0.10	<LLD	-	-	None	0
Gamma Background (TLDs) (mR/Qtr.)	Gamma Dose 16	3.0	18.3 (6/6) (13.2-22.5)	Q-04, Nitrin 1.5 mi @ 40°	22.5 (1/1) -	16.1 (10/10) (11.6-21.3)	0
Milk (pCi/l)	I-131 20	5/0.5 ^b	<LLD	-	-	None	0
Cooling Water (pCi/l)	Gross Beta 26	1.0	364.5 (13/13) (3.6-4242.9)	Q-22A, Diffuser Pipe Blowdown	364.5 (13/13) (3.6-4242.9)	7.1 (13/13) (1.1-35.7)	4
	Tritium 1	200	380 (1/1)	Q-22A, Diffuser Pipe Blowdown	380 (1/1)	None	0
Public Water (pCi/l)	Gamma Spec. 6						
	Cs-134	10.0	<LLD	-	-	None	0
	Cs-137	10.0	<LLD	-	-	None	0
	Other Gammas	20.0	<LLD	-	-	None	0
Fish (pCi/g wet)	Gamma Spec. 8						
	Cs-134	0.1	<LLD	-	-	None	0
	Cs-137	0.1	<LLD	-	-	None	0
	Other Gammas	0.2	<LLD	-	-	None	0

^a Mean and range based on detectable measurements only. Fractions indicated in parentheses.

^b November - April LLD = 5.0; May - October LLD = 0.5.

Table 5.1-1

GAMMA RADIATION AS MEASURED BY THERMOLUMINESCENT DOSIMETERS (TLDs)

STANDARD RADIOLOGICAL MONITORING PROGRAM					
		<u>1st Quarter</u>	<u>2nd Quarter</u>	<u>3rd Quarter</u>	<u>4th Quarter</u>
Date Placed:		12-28-84	03-28-85	06-29-85	09-28-85
Date Removed:		03-28-85	06-29-85	09-28-85	12-27-85
Days in the Field:		90	93	91	90
Location		Average mR/Quarter			
<u>On-Site Indicator Locations</u>					
Q-01	On-Site No. 1	9.3±2.3	10.4±1.7	12.9±3.1	17.5±6.1
Q-02	On-Site No. 2	10.0±1.9	11.5±2.1	11.0±2.1	21.7±1.1
Q-03	On-Site No. 3	14.9±1.9	12.8±1.9	9.8±2.1	14.5±1.1
Mean ± s.d.		11.4±3.0	11.6±1.2	11.2±1.6	17.9±3.6
<u>Off-Site Indicator Locations</u>					
Q-04	Nitrin	11.3±2.9	10.3±2.2	9.7±1.8	22.5±1.4
Q-05	Saddle Club Dairy	11.4±2.8	10.4±1.0	7.0±2.0	20.5±1.2
Q-06	Hanson's Dock	15.5±2.4	13.9±1.7	13.0±1.3	13.2±0.9
Mean ± s.d.		12.7±2.4	11.5±2.0	9.9±3.0	18.7±4.9
<u>Background Locations</u>					
Q-07	Clinton	4.6±2.4	9.7±2.3	8.5±3.7	11.6±0.9
Q-08	Sitkema Farm	9.8±2.5	10.3±1.8	7.9±1.7	19.0±3.3
Q-09	Erie	8.0±2.6	11.2±1.2	12.0±1.7	16.0±1.3
Q-10	Hillsdale	30.0±4.5	16.5±1.5	8.4±1.1	18.3±1.5
Q-11	Port Byron	21.9±3.2	12.9±1.3	14.3±1.9	21.3±1.7
Q-12	Bettendorf	12.0±2.4	16.5±2.0	10.8±1.5	15.5±1.3
Q-13	Princeton	12.6±2.1	11.5±2.0	10.1±1.8	13.0±0.8
Q-14	Utica Ridge Road	21.4±3.6	10.9±1.2	8.5±1.5	16.2±1.2
Q-15	DeWitt	12.0±2.9	8.1±1.8	7.6±2.2	11.9±1.0
Q-16	Low Moor	10.4±3.2	11.0±1.6	11.3±4.2	18.6±1.0
Mean ± s.d.		14.3±7.7	11.9±2.8	9.9±2.2	16.1±3.2

Table 5.1-1 (continued)

GAMMA RADIATION AS MEASURED BY TLDs (continued)

SPECIAL PROGRAM					
Inner Ring, Near Site Boundary, Indicator Locations					
		1st Quarter	2nd Quarter	3rd Quarter	4th Quarter
Date Placed:		12-28-84	03-28-85	06-29-85	09-28-85
Date Removed:		03-28-84	06-29-85	09-28-85	12-27-85
Days in the Field:		90	93	91	90
Previous Location Code	Revised Location Code ^a	Average mR/Quarter			
Q-101-1	same	7.1±2.0	9.9±1.7	9.2±3.1	17.8±0.7
Q-101-2	same	15.5±2.3	15.7±3.1	14.9±4.3	18.1±3.2
Q-102-1	same	18.9±2.2	15.0±2.7	14.6±1.0	44.7±2.0 ^b
Q-102-2	same	6.5±2.1	11.9±2.2	13.0±1.3	NDC ^c
Q-103-1	same	5.8±2.4	10.9±1.9	12.7±2.1	11.5±1.1
Q-103-2	104-1	2.2±2.4	8.9±1.9	14.8±1.4	11.3±0.8
Q-104-1	104-2	8.1±3.1	9.6±1.8	9.2±1.8	13.2±2.8
Q-104-2	104-3	3.8±1.8	12.2±2.1	10.1±1.4	15.7±1.3
Q-105-1	same	4.3±2.0	15.7±3.9	11.0±1.2	13.7±1.0
Q-105-2	106-1	2.0±1.9	10.1±1.5	12.4±1.3	13.4±1.0
Q-106-1	106-2	5.2±2.3	11.8±2.3	13.1±1.8	19.8±16.7 ^b
Q-106-2	107-1	3.9±2.3	14.6±1.7	12.0±3.9	14.7±1.4
Q-107-1	107-2	7.6±2.5	11.5±1.8	9.0±2.2	13.9±1.4
Q-107-2	107-3	4.6±2.6	10.9±1.7	9.5±1.1	14.9±0.6
Q-108-1	same	4.0±2.4	10.3±3.0	9.4±1.4	12.3±1.0
Q-108-2	same	9.5±3.4	11.1±1.2	9.7±1.5	11.5±0.5
Q-109-1	same	2.7±2.0	9.1±1.2	9.8±1.2	14.8±1.0
Q-109-2	same	11.0±2.0	11.9±1.3	11.4±2.0	11.8±1.2
Q-111-1	same	2.8±2.0	12.9±2.8	10.5±1.5	13.0±3.8
Q-111-2	same	3.6±2.2	14.0±1.5	13.7±1.9	13.4±1.2
Q-112-1	same	13.6±2.6	11.7±1.4	6.8±1.9	11.7±0.8
Q-112-2	same	10.3±2.7	15.1±5.8	20.3±3.0	14.3±0.9
Q-113-1	same	13.4±3.1	10.6±2.0	9.5±3.2	17.9±0.8
Q-113-2	same	10.0±2.9	11.5±2.1	13.9±2.0	17.6±1.6
Q-114-1	same	11.0±2.0	11.3±1.1	10.7±1.6	10.0±0.7
Q-114-2	same	8.4±1.8	10.6±1.7	8.4±3.3	8.9±1.1
Q-115-1	same	18.3±2.6	14.6±1.9	9.7±3.2	13.2±1.0
Q-115-2	same	5.7±2.6	8.9±4.2	12.6±4.5	10.9±2.5
Q-116-1	same	9.5±1.9	10.0±1.5	12.2±4.7	12.1±0.9
Q-116-2	same	3.9±2.8	15.0±1.7	10.6±1.5	13.2±0.7
Mean ± s.d.		7.8±4.7	12.2±2.4	11.5±2.6	13.5±2.4

^a Effective 3rd quarter 1985.^b Chips damaged (white); not included in quarterly mean.^c ND = No data; TLD lost in the field.

Table 5.1-1 (continued)

GAMMA RADIATION AS MEASURED BY TLDs (continued)

SPECIAL PROGRAM				
Outer Ring, Near 5 Mile Radius, Indicator Locations				
	1st Quarter	2nd Quarter	3rd Quarter	4th Quarter
Date Placed:	12-28-84	03-28-85	06-29-85	09-28-85
Date Removed:	03-28-85	06-29-85	09-28-85	12-27-85
Days in the Field:	90	93	91	90
Location	Average mR/Qtr			
Q-201-1	10.8±2.2	15.3±1.8	13.4±1.8	14.4±1.5
Q-201-2	12.0±2.3	17.0±1.7	11.8±1.9	14.7±0.6
Q-202-1	10.6±4.9	9.0±2.8	7.8±2.6	11.2±0.8
Q-202-2	11.8±2.1	12.2±2.3	8.2±1.5	10.5±1.0
Q-203-1	8.8±5.7	9.3±2.8	10.8±3.2	14.6±0.6
Q-203-2	13.8±5.5	17.8±1.1	10.7±1.4	27.4±2.4 ^a
Q-204-1	13.2±2.3	11.1±1.1	15.6±2.1	15.7±3.4
Q-204-2	13.7±3.2	13.9±5.1	12.1±1.8	13.9±1.2
Q-205-1	12.6±2.8	15.0±1.6	12.5±1.8	13.3±2.8
Q-205-2	8.3±3.0	14.7±1.6	11.9±1.3	16.4±3.4
Q-206-1	11.2±2.2	16.3±1.5	ND ^b	16.4±3.9
Q-206-2	11.9±2.7	13.6±2.2	12.0±1.8	46.1±3.1 ^a
Q-207-1	11.9±2.4	15.4±2.1	11.0±1.4	16.4±1.4
Q-207-2	9.5±2.9	12.4±2.5	13.4±2.1	ND ^b
Q-208-1	11.9±2.7	14.4±2.6	14.4±4.6	27.1±1.3 ^a
Q-208-2	12.9±2.4	13.5±1.6	10.6±1.6	15.2±3.6
Q-209-1	12.4±2.2	17.7±1.4	12.9±1.8	14.3±3.1
Q-209-2	10.2±2.0	16.6±2.2	10.0±2.7	13.8±2.8
Q-210-1	12.7±1.9	13.2±3.4	9.4±1.7	14.4±4.8
Q-210-2	9.6±2.3	17.2±6.1	10.2±1.4	13.8±1.0
Q-211-1	10.6±2.6	14.3±2.2	13.0±2.9	18.6±1.6
Q-212-1	12.9±2.1	13.3±2.0	10.8±2.4	12.7±0.7
Q-212-2	12.9±2.1	11.1±1.5	10.5±1.3	25.2±1.5 ^a
Q-213-1	13.6±1.9	11.2±2.7	9.2±1.2	14.9±1.2
Q-213-2	11.9±1.9	12.3±1.5	10.3±2.9	34.4±1.6 ^a
Q-214-1	12.7±2.0	11.4±1.2	9.4±1.2	27.0±1.7
Q-214-2	14.3±2.3	15.1±2.2	13.7±1.0	28.2±4.9
Q-215-1	10.8±2.7	13.7±1.1	16.7±1.9	12.1±0.9
Q-215-2	14.3±2.9	15.4±1.9	19.3±2.4	26.6±0.6 ^a
Q-216-1	10.9±2.9	ND ^a	17.7±1.2	14.0±1.2
Q-216-2	11.1±2.5	15.9±1.8	15.3±1.0	28.1±1.0 ^a
Mean ± s.d.	11.8±1.6	14.0±2.4	12.2±2.8	15.5±4.2

^a Chips damaged (white); not included in quarterly mean.

^b ND = No data; TLDs lost in the field.

APPENDIX II

METEOROLOGICAL DATA

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH 1935
 STABILITY CLASS - EXTREMELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	0-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	0	0	0	0
ESE	0	0	0	1	0	0	1
SE	0	0	0	0	0	0	0
SSE	0	0	0	0	1	0	1
S	0	0	0	3	4	3	10
SSW	0	0	0	1	0	0	1
SW	0	0	0	0	0	0	0
WSW	0	0	1	0	0	0	1
W	0	0	1	3	0	0	4
WNW	0	0	0	6	0	0	6
NW	0	0	1	0	2	0	3
NNW	0	0	0	0	0	0	0
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	0	3	14	7	3	27

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH 1985
 STABILITY CLASS - MODERATELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	0-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	0	1	0	0	1
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	0	0	0	0
ESE	0	0	0	3	0	0	3
SE	0	0	0	0	1	0	1
SSE	0	0	0	0	1	0	1
S	0	0	0	1	1	1	3
SSW	0	2	3	0	1	0	6
SW	0	0	0	0	0	0	0
WSW	0	0	0	0	0	0	0
W	0	0	0	0	0	0	0
WNW	0	0	1	3	0	0	4
Ww	0	0	4	3	3	0	10
Nw	0	0	2	1	0	0	3
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	2	10	11	7	1	32

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH
 STABILITY CLASS - SLIGHTLY UNSTABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-03 P)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	0	1	0	0	1
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	1	2	1	4
E	0	0	0	0	0	0	0
ESE	0	0	0	0	1	0	1
SE	0	0	0	0	1	0	1
SSE	0	0	1	3	0	0	4
S	0	0	0	0	1	1	2
SSW	0	1	0	1	0	0	2
SW	0	0	1	0	0	0	1
WSW	0	1	0	0	0	0	1
W	0	0	0	0	0	0	0
WNW	0	0	0	2	2	0	4
NW	0	0	5	2	4	1	12
NNW	0	0	2	6	0	0	8
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	2	9	16	11	3	41

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH
 STABILITY CLASS - NEUTRAL
 WINDS MEASURED AT 296 FEET

1985
 (DELTA 1 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	0-3	4-7	8-12	13-18	19-24	GT 24	
N	1	4	7	12	6	0	30
NNE	0	1	4	3	3	0	11
NE	0	3	12	6	6	2	31
ENE	0	0	17	29	11	9	67
E	0	2	15	25	12	10	64
ESE	0	1	13	16	8	1	39
SE	0	5	14	21	11	1	52
SSE	0	2	9	14	3	4	32
S	0	0	5	19	7	2	33
SSW	0	5	7	23	7	0	47
SW	0	6	12	19	6	0	43
WSW	1	2	19	41	25	13	101
W	0	6	13	24	39	21	110
WNW	1	2	12	36	150	46	299
NW	0	2	17	44	43	28	134
NW	0	1	10	22	18	4	55
VARIABLE	0	0	0	0	0	0	0
TOTAL	9	45	106	413	355	143	1150

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH
 STABILITY CLASS - SLIGHTLY STABLE
 WINDS MEASURED AT 296 FEET

1985

(DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	2	6	14	9	0	0	31
NNE	0	3	7	0	0	0	10
NE	2	6	7	5	0	0	20
ENE	6	5	15	13	0	0	39
E	2	1	8	21	6	1	39
ESE	2	2	4	11	6	0	25
SE	1	5	7	7	2	0	22
SSE	1	5	3	13	5	1	28
S	0	2	15	24	9	14	64
SSW	0	3	9	44	28	6	90
SW	1	4	22	14	6	0	47
WSW	0	1	4	11	11	2	29
W	1	5	6	23	24	1	60
WNW	3	8	8	45	22	1	87
NW	2	5	11	37	29	1	85
NNW	5	3	15	11	0	0	34
VARIABLE	0	0	0	0	0	0	0
TOTAL	28	64	155	288	148	27	710

HOURS OF CALM IN THIS STABILITY CLASS - 0

HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0

HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH 1985
 STABILITY CLASS - MODERATELY STABLE (DELTA T 296-30 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	0-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	1	2	0	0	3
NNE	1	1	0	1	0	0	3
NE	0	0	1	3	0	0	4
ENE	0	1	0	2	0	0	3
E	0	2	0	1	0	0	3
ESE	0	0	3	5	4	0	12
SE	0	0	1	1	2	0	4
SSE	0	1	1	7	1	0	10
S	0	0	1	7	0	0	8
SSW	1	1	3	5	0	0	10
SW	1	3	4	0	0	0	8
WSW	0	1	7	2	1	0	11
W	1	1	4	3	0	0	9
WNW	1	3	2	11	0	0	17
NW	1	2	2	5	0	0	10
NNW	1	6	6	2	1	0	16
VARIABLE	0	0	0	0	0	0	0
TOTAL	7	22	36	57	9	0	131

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JANUARY-MARCH
 STABILITY CLASS - EXTREMELY STABLE
 WINDS MEASURED AT 296 FEET

1985

(DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4- 7	8-12	13-18	19-24	GT 24	
N	2	4	0	0	0	0	6
NNE	3	4	2	0	0	0	9
NE	0	2	0	1	0	0	3
ENE	0	1	0	0	0	0	1
E	1	1	0	0	0	0	2
ESE	2	1	0	2	0	0	5
SE	0	1	0	0	2	0	3
SSE	0	0	1	1	1	0	3
S	0	0	2	7	0	0	9
SSW	0	2	3	4	0	0	9
SW	1	0	3	2	0	0	6
WSW	0	0	0	0	0	0	0
W	0	1	1	0	0	0	2
WNW	0	0	0	0	0	0	0
NW	0	0	2	1	2	0	5
NNW	0	2	0	3	0	0	5
VARIABLE	0	0	0	0	0	0	0
TOTAL	9	19	14	21	5	0	68

HOURS OF CALM IN THIS STABILITY CLASS - 0

HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0

HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 1

DUAL CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE 1985
 STABILITY CLASS - EXTREMELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	0-3	4-7	8-12	13-18	19-24		
N	0	0	0	4	0	0	4
NNE	0	0	1	0	0	0	1
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	1	0	0	1
ESE	0	0	0	0	0	0	0
SE	0	0	1	1	0	0	2
SSE	0	0	2	4	4	2	12
S	0	1	1	4	2	1	9
SSW	0	0	10	12	24	5	71
SW	0	0	4	6	1	0	13
WSW	0	2	0	4	3	1	10
W	0	0	12	5	2	1	20
WSW	0	1	8	10	2	3	24
NW	9	1	0	2	4	1	11
NNW	0	0	0	1	1	1	3
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	5	69	77	45	15	181

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE 1985
 STABILITY CLASS - MODERATELY UNSTABLE (DELTA T 296-33 F)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	0	0	3	5	4	0	12
NNE	0	0	3	4	0	0	7
NE	0	0	6	2	0	0	8
ENE	0	0	1	2	0	0	3
E	0	0	0	3	0	0	3
ESE	0	0	0	1	0	0	1
SE	0	0	3	4	0	0	7
SSE	0	0	5	2	2	2	11
S	0	1	1	4	2	3	11
SSW	0	2	4	6	4	4	20
SW	0	4	2	5	0	0	11
WSW	0	6	0	3	4	1	14
W	0	3	1	4	0	4	15
WNW	0	2	7	4	5	6	25
NW	0	0	1	5	6	4	16
NNW	0	2	1	0	2	0	5
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	21	39	54	32	24	169

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE 1985
 STABILITY CLASS - SLIGHTLY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 27.6 FEET

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	0-3	4-7	8-12	13-18	19-24		
N	0	5	4	2	0	0	11
NNE	0	0	9	7	1	0	17
NE	0	1	8	5	1	1	16
ENE	0	0	7	4	1	0	12
E	0	0	7	6	0	0	13
ESE	0	0	2	4	2	0	8
SE	0	0	1	2	0	0	3
SSE	0	1	5	2	0	2	10
S	0	1	2	4	1	6	14
SSW	0	0	3	5	3	2	13
SW	0	5	2	2	0	0	12
WSW	0	2	2	2	1	1	8
W	0	3	2	0	3	5	13
WNW	0	6	1	1	11	0	27
NW	0	4	5	7	1	1	18
NNW	0	1	4	4	0	0	9
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	34	64	60	30	26	214

HOURS OF CALM IN THIS STABILITY CLASS - 9
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE
 STABILITY CLASS - NEUTRAL
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 F)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.3-3	4-7	8-12	13-18	19-24	GT 24	
N	0	9	32	10	10	1	62
NNE	0	6	14	20	4	0	44
NE	0	7	29	33	16	0	85
ENE	0	3	33	19	14	0	69
E	0	2	13	11	4	4	34
ESE	0	1	2	10	6	2	21
SE	1	2	12	7	0	0	22
SSE	1	1	8	9	6	8	33
S	1	1	7	14	19	14	56
SSW	1	7	10	18	11	21	68
SW	5	14	23	33	9	0	84
WSW	1	3	8	28	6	1	47
W	2	3	10	19	15	10	59
WNW	0	13	12	28	34	16	103
NW	4	8	16	22	14	1	65
NNW	1	3	17	5	3	0	29
VARIABLE	0	0	0	0	0	0	0
TOTAL	17	63	246	266	171	78	881

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

DUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE
 STABILITY CLASS - SLIGHTLY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	1	10	2	0	0	13
NNE	1	4	7	6	1	0	19
NE	0	1	6	14	5	0	26
ENE	0	1	6	14	2	0	23
E	0	3	9	15	3	0	30
ESE	0	2	6	17	2	0	27
SE	0	2	3	3	0	0	6
SSE	1	2	4	9	6	5	27
S	1	4	6	28	24	13	76
SSW	0	1	12	39	23	11	66
SW	0	1	19	25	2	0	47
WSW	0	3	7	7	1	2	20
W	0	1	5	14	8	0	28
WNW	1	7	13	11	11	0	43
NW	0	2	8	13	9	0	26
NNW	0	0	10	5	0	0	15
VARIABLE	0	0	0	0	0	0	0
TOTAL	4	35	133	222	91	31	516

HOURS OF CALM IN THIS STABILITY CLASS - 0

HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0

HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE
 STABILITY CLASS - MODERATELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	1	0	0	0	1
NNE	0	1	9	2	0	0	12
NE	0	3	5	5	0	0	13
ENE	0	0	2	3	0	0	5
E	0	2	5	7	1	0	15
ESE	0	5	6	6	2	0	19
SE	0	3	4	6	1	0	14
SSE	2	6	7	6	0	0	21
S	0	2	7	6	5	0	20
SSW	0	1	2	3	0	0	6
SW	1	0	8	2	0	0	11
WSW	0	0	5	2	0	0	7
W	0	1	4	5	1	0	11
WNW	1	1	0	14	4	0	20
NW	0	2	1	2	1	0	6
NNW	0	0	1	1	0	0	2
VARIABLE	0	0	0	0	0	0	0
TOTAL	4	27	67	70	15	0	183

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

DUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - APRIL-JUNE
 STABILITY CLASS - EXTREMELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	1	1	0	0	2
E	0	1	1	0	0	0	2
ESE	0	2	4	0	1	0	7
SE	0	1	4	1	0	0	6
SSE	0	0	1	2	0	0	3
S	0	0	3	0	0	0	3
SSW	1	1	0	0	0	0	2
SW	0	0	2	0	0	0	2
WSW	1	0	2	1	0	0	4
W	0	1	2	2	0	0	5
WSW	1	1	2	0	0	0	4
NW	0	0	0	0	0	0	0
NNW	0	0	0	0	0	0	0
VARIABLE	0	0	0	0	0	0	0
TOTAL	3	7	22	7	1	0	40

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER 1985
 STABILITY CLASS - EXTREMELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	0	0	0	0
ESE	0	0	0	0	0	0	0
SE	0	0	0	4	0	0	4
SSE	0	0	4	3	0	0	7
S	0	0	0	5	7	1	13
SSW	0	0	11	20	14	0	45
SW	0	0	1	1	1	0	3
WSW	0	1	3	1	1	0	6
W	0	0	8	6	1	0	15
WNW	0	0	2	6	1	0	9
NW	0	0	6	6	0	0	12
NNW	0	0	0	3	0	0	3
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	1	35	55	25	1	117

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER 1985
 STABILITY CLASS - MODERATELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	6	0	0	0	6
NNE	0	0	1	0	0	0	1
NE	0	0	0	2	0	0	2
ENE	0	0	0	3	0	0	3
E	0	0	0	0	0	0	0
ESE	0	0	0	2	1	0	3
SE	0	0	0	2	1	0	3
SSE	0	1	5	0	1	0	7
S	0	0	3	2	6	0	11
SSW	0	0	13	9	5	1	28
SW	0	5	3	1	0	0	9
WSW	0	1	1	2	3	0	7
W	0	3	1	1	1	0	6
WNW	0	1	2	2	1	0	6
NW	0	1	4	1	0	0	6
NNW	0	0	3	4	0	0	7
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	12	42	31	19	1	105

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER
 STABILITY CLASS - SLIGHTLY UNSTABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	3	3	3	0	0	9
NNE	0	0	0	1	0	0	1
NE	0	2	1	2	0	0	5
ENE	0	0	3	4	0	0	7
E	0	0	0	4	0	0	4
ESE	0	1	8	4	0	0	13
SE	0	0	5	3	0	0	8
SSE	0	0	4	2	0	0	6
S	0	1	11	4	0	3	19
SSW	0	4	18	6	1	1	30
SW	0	3	5	3	1	0	12
WSW	0	3	1	4	1	0	9
W	1	3	3	4	0	1	12
WNW	0	4	0	5	2	0	11
NW	0	3	6	1	0	0	10
NNW	0	1	4	6	0	0	11
VARIABLE	0	0	0	0	0	0	0
TOTAL	1	28	72	56	5	5	167

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER
 STABILITY CLASS - NEUTRAL
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	1	16	13	7	0	0	37
NNE	0	9	8	2	0	0	19
NE	0	10	21	32	2	0	65
ENE	0	9	13	13	1	0	36
E	0	12	16	23	1	0	52
ESE	0	13	32	19	0	0	64
SE	1	6	31	16	1	1	56
SSE	1	6	10	15	5	0	37
S	2	6	17	15	16	4	60
SSW	0	14	24	30	9	1	78
SW	5	22	23	16	5	0	71
WSW	6	14	14	12	8	1	55
W	2	5	13	21	19	6	66
WNW	4	9	8	11	4	0	36
NW	6	15	20	16	2	1	60
NNW	0	9	15	17	1	0	42
VARIABLE	0	0	0	0	0	0	0
TOTAL	28	175	278	265	74	14	834

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER
 STABILITY CLASS - SLIGHTLY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	7	14	10	1	0	32
NNE	0	7	12	8	0	0	27
NE	0	1	10	6	1	0	18
ENE	2	4	9	9	0	0	24
E	2	5	7	15	1	0	30
ESE	0	2	12	12	2	0	28
SE	0	6	12	23	1	0	42
SSE	0	3	13	21	3	0	40
S	1	4	15	28	35	1	84
SSW	1	2	17	48	30	1	99
SW	0	9	19	3	0	0	31
WSW	2	11	15	8	1	0	37
W	2	2	11	9	0	0	24
WNW	0	6	10	11	1	0	28
NW	1	7	17	12	1	0	38
NNW	1	6	7	7	0	0	21
VARIABLE	0	0	0	0	0	0	0
TOTAL	12	82	200	230	77	2	603

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER
 STABILITY CLASS - MODERATELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4- 7	8-12	13-18	19-24		
N	0	0	8	3	0	0	11
NNE	0	2	3	0	0	0	5
NE	0	2	8	1	0	0	11
ENE	0	1	4	7	0	0	12
E	1	4	4	7	1	0	17
ESE	0	1	7	22	3	0	33
SE	1	3	9	32	1	0	46
SSE	1	1	6	12	4	0	24
S	0	1	4	16	5	0	26
SSW	0	5	13	10	1	0	29
SW	0	15	2	0	0	0	17
WSW	1	3	7	3	0	0	14
W	0	3	0	7	0	0	10
WNW	0	3	5	3	2	0	13
NW	0	2	3	3	0	0	8
NNW	1	2	3	2	1	0	9
VARIABLE	0	0	0	0	0	0	0
TOTAL	5	48	86	128	18	0	285

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - JULY-SEPTEMBER
 STABILITY CLASS - EXTREMELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4- 7	8-12	13-18	19-24		
N	1	1	0	0	0	0	2
NNE	2	0	0	0	0	0	2
NE	0	0	0	0	0	0	0
ENE	0	1	0	0	0	0	1
E	0	2	1	1	0	0	4
ESE	0	0	2	0	0	0	2
SE	0	0	4	11	1	0	16
SSE	2	3	1	19	2	0	27
S	0	3	1	8	0	0	12
SSW	1	6	3	6	0	0	16
SW	1	4	1	0	0	0	6
WSW	0	1	3	0	0	0	4
W	1	0	0	0	0	0	1
WNW	0	1	0	1	0	0	2
NW	0	0	0	0	0	0	0
NNW	1	1	0	0	0	0	2
VARIABLE	0	0	0	0	0	0	0
TOTAL	9	23	16	46	3	0	97

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER 1985
 STABILITY CLASS - EXTREMELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	0	0	0	0
ESE	0	0	0	0	0	0	0
SE	0	0	0	0	0	0	0
SSE	0	0	0	0	0	0	0
S	0	0	0	0	0	0	0
SSW	0	0	3	6	0	0	9
SW	0	0	1	0	0	0	1
WSW	0	0	0	0	1	0	1
W	0	0	0	2	2	0	4
WNW	0	0	0	0	1	0	1
NW	0	0	0	0	0	0	0
NNW	0	0	0	0	0	0	0
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	0	4	8	4	0	16

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER 1985
 STABILITY CLASS - MODERATELY UNSTABLE (DELTA T 296-33 FT)
 WINDS MEASURED AT 296 FEET

WIND DIRECTION	WIND SPEED (IN MPH)						TOTAL
	.8-3	4-7	8-12	13-18	19-24	GT 24	
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	1	0	1
ENE	0	0	0	0	0	0	0
E	0	0	0	0	0	0	0
ESE	0	0	0	0	0	0	0
SE	0	0	0	0	0	0	0
SSE	0	0	0	0	0	0	0
S	0	0	1	1	0	0	2
SSW	0	1	2	1	2	0	6
SW	0	1	1	1	0	0	3
WSW	0	0	3	7	0	1	11
W	0	0	2	10	8	1	21
WNW	0	0	0	2	2	1	5
NW	0	0	0	0	1	0	1
NNW	0	0	0	0	0	0	0
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	2	9	22	14	3	50

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 5
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER
 STABILITY CLASS - SLIGHTLY UNSTABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	0	0	0	0	0	0	0
NNE	0	0	1	0	0	0	1
NE	0	0	1	1	3	0	5
ENE	0	2	5	4	0	0	11
E	0	1	1	1	0	0	3
ESE	0	0	1	0	0	0	1
SE	0	0	0	0	0	0	0
SSE	0	0	4	4	2	0	10
S	0	0	3	4	2	0	9
SSW	0	0	3	1	3	1	8
SW	0	3	5	1	0	2	11
WSW	0	1	0	3	0	7	11
W	0	2	3	11	10	8	34
WNW	0	0	6	17	18	6	47
NW	0	0	1	8	6	0	15
NNW	0	0	0	4	0	0	4
VARIABLE	0	0	0	0	0	0	0
TOTAL	0	9	34	59	44	24	170

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 9
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER
 STABILITY CLASS - NEUTRAL
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 F)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	1	6	21	21	1	0	50
NNE	3	16	50	22	0	0	91
NE	1	17	42	31	25	1	117
ENE	0	11	40	43	6	0	100
E	2	15	42	32	16	1	108
ESE	0	12	21	25	11	2	71
SE	2	16	19	22	9	0	68
SSE	3	8	13	19	13	5	61
S	0	1	5	18	14	10	48
SSW	0	6	5	18	17	6	52
SW	2	6	14	19	4	0	45
WSW	4	5	10	29	32	5	85
W	0	5	18	58	29	6	116
WNW	1	8	25	65	36	32	167
NW	0	6	26	57	24	2	115
NNW	1	3	29	27	13	4	77
VARIABLE	0	0	0	0	0	0	0
TOTAL	20	141	380	506	250	74	1371

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 59
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER
 STABILITY CLASS - SLIGHTLY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4- 7	8-12	13-18	19-24		
N	0	0	6	5	0	0	11
NNE	0	1	9	8	0	0	18
NE	1	4	11	3	1	0	20
ENE	0	4	9	7	8	0	28
E	1	3	7	10	6	0	27
ESE	0	3	6	8	1	0	18
SE	0	6	1	6	4	0	17
SSE	1	0	3	4	5	2	15
S	0	1	5	25	8	1	40
SSW	0	4	17	29	4	0	54
SW	0	4	7	6	0	0	17
WSW	1	1	5	6	0	0	13
W	0	2	7	15	7	0	31
WNW	0	0	4	18	0	0	22
NW	0	0	1	3	0	0	4
NNW	2	2	6	6	1	0	17
VARIABLE	0	0	0	0	0	0	0
TOTAL	6	35	104	159	45	3	352

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 17
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER
 STABILITY CLASS - MODERATELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)						GT 24	TOTAL
	.8-3	4- 7	8-12	13-18	19-24			
N	0	2	2	3	0	0	7	
NNE	0	0	5	2	0	0	7	
NE	0	0	2	6	0	0	8	
ENE	0	0	0	5	0	0	5	
E	0	1	1	3	0	0	5	
ESE	0	0	1	4	0	0	5	
SE	0	1	1	2	7	0	11	
SSE	0	1	4	3	0	0	8	
S	0	0	2	19	4	0	25	
SSW	0	2	1	4	0	0	7	
SW	0	1	2	0	0	0	3	
WSW	0	0	0	4	0	0	4	
W	0	1	2	1	0	0	4	
WNW	0	0	0	8	0	0	8	
NW	0	0	2	2	0	0	4	
NNW	0	1	1	1	2	0	5	
VARIABLE	0	0	0	0	0	0	0	
TOTAL	0	10	26	67	13	0	116	

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 3
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

QUAD CITIES NUCLEAR POWER STATION
 PERIOD OF RECORD - OCTOBER-DECEMBER
 STABILITY CLASS - EXTREMELY STABLE
 WINDS MEASURED AT 296 FEET

1985
 (DELTA T 296-33 FT)

WIND DIRECTION	WIND SPEED (IN MPH)					GT 24	TOTAL
	.8-3	4-7	8-12	13-18	19-24		
N	0	0	0	0	0	0	0
NNE	0	0	0	0	0	0	0
NE	0	0	0	0	0	0	0
ENE	0	0	0	0	0	0	0
E	0	0	0	1	1	0	2
ESE	0	0	0	0	0	0	0
SE	0	0	0	0	0	0	0
SSE	0	1	1	0	0	0	2
S	0	4	3	9	2	0	18
SSW	1	2	5	2	2	0	12
SW	0	0	1	0	0	0	1
WSW	0	0	0	0	0	0	0
W	0	0	0	0	0	0	0
WNW	0	1	0	2	0	0	3
NW	0	1	1	0	0	0	2
NNW	0	0	0	0	0	0	0
VARIABLE	0	0	0	0	0	0	0
TOTAL	1	9	11	14	5	0	40

HOURS OF CALM IN THIS STABILITY CLASS - 0
 HOURS OF MISSING WIND MEASUREMENTS IN THIS STABILITY CLASS - 0
 HOURS OF MISSING STABILITY MEASUREMENTS IN ALL STABILITY CLASSES - 0

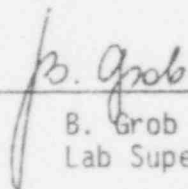
APPENDIX III

ANALYTICAL PROCEDURES

ANALYTICAL PROCEDURES MANUAL
TELEDYNE ISOTOPES MIDWEST LABORATORY
PREPARED FOR
COMMONWEALTH EDISON COMPANY

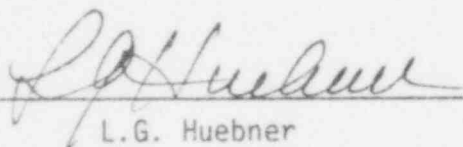
Note: These procedures are taken from the complete Procedures Manual. Only procedures applicable to the CECO Radiological Environmental Monitoring Programs are included in this manual.

Compiled by:



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L.G. Huebner
General Manager

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SECTION 1.0

SAMPLE PREPARATION

Different classes of samples require different preparations. In general, food products are prepared as for home use, while others are dried and ashed as received.

1.1 Fish

1. Wash the fish.
2. Fillet and place the flesh immediately (to prevent moisture loss) in a 500 cc plastic container. Add a few cc of formaldehyde. Seal and record wet weight.

NOTE: If bones are to be analyzed, boil remaining fish in water for about 1 hour. Clean the bones. Air dry, weigh and record as wet weight. Dry at 125° C. Record dry weight. Ash at 800° C, cool, weigh, and record the ash weight. Grind to a homogeneous sample. The sample is ready for analysis.

3. Gamma scan fillet without delay or store in a freezer.
4. After gamma spectroscopic analysis is completed transfer the sample to a drying pan and dry at 125° C.
5. Cool, weigh, and record dry weight.
6. Ash by gradually increasing the temperature to 450° C. If considerable amounts of carbon remain after overnight ashing, the sample should be brushed and placed back in the muffle furnace until ashing is completed. Record ash weight. The sample is now ready for analysis.

NOTE: If there is sufficient quantity, use surplus flesh for drying and ashing, instead of waiting for gamma scanning to be completed.

L.G. Huebner
L.G. Huebner

1.2 Bottom Sediments and Soil

1. Air dry the entire sample. Grind or pulverize the sample and sieve through a #20 mesh screen.
2. For gamma-spectroscopic analysis, seal 500 cc of the ground sample in a Marinelli beaker. Record dry weight.
3. Seal the remaining sample (up to 1 kg) in a plastic container and save for other analyses or for possible future rechecking.

1.3 Drinking (clear) water (EPA Method 900.0)

A representative sample must be collected from a free-flowing source of drinking water, and should be large enough so that adequate aliquots can be taken to obtain the required sensitivity.

It is recommended that samples be preserved at the time of collection by adding enough 1N HNO₃ to the sample to bring it to pH 2 (15 ml 1N HNO₃ per liter of sample is usually sufficient). If samples are to be collected without preservation, they should be brought to the laboratory within 5 days, then preserved and held in the original container for a minimum of 16 hours before analysis or transfer of the sample.

The container choice should be plastic over glass to prevent loss due to breakage during transportation and handling.

If the sample was not acidified at the time of collection, use the following procedure:

Procedure

1. Remove 100 ml of sample for tritium analysis, if required.

NOTE: Water should not be acidified for tritium analysis. If samples are acidified in the field, an additional aliquot should be collected.

2. Add 15 ml of 1N HNO₃ per liter of sample in the original container.
3. Hold the sample in the original container for a minimum of 16 hours before analysis or transfer of the sample.
4. When taking an aliquot for analysis, take acid addition into account. For example:

<u>Sample volume to be analyzed</u>	<u>Volume of aliquot required</u>
200 ml	203 ml
400 ml	406 ml
600 ml	609 ml
800 ml	812 ml
1000 ml	1015 ml
2000 ml	2030 ml
3000 ml	3045 ml
3500 ml	3552 ml

For other volumes, adjust aliquots correspondingly, at the rate of 1.5 ml per 100 ml of sample.

2.1 Airborne Particulates

2.1.1. Gross Alpha and/or Gross Beta Activity

Procedure

1. Store the sample for 5 days from the day of collection to allow for decay of short-lived radon and thoron daughters.
2. Place a 47 mm filter on a stainless steel planchet and count the sample in a proportional counter.
3. Calculate the activity in pCi/m³ using computer program AIRPAT.

Calculations

Gross alpha (beta) concentration:

$$(\text{pCi/m}^3) = \frac{A}{B \times C \times 2.22} + \frac{2 \sqrt{E_{sb}^2 + E_b^2}}{B \times C \times 2.22}$$

Where:

- A = net alpha (beta) count rate (cpm)
- B = efficiency for counting alpha (beta) activity (cpm/dpm)
- C = volume of sample (m³)
- E_{sb} = counting error of sample plus background
- E_b = counting error of background

2.2.2 Gross Alpha and/or Gross Beta Activity in Dissolved Solids (see note)Principle of Method

Water samples containing suspended matter are filtered through a membrane filter and the filtrate is analyzed. The filtered water sample is evaporated and the residue is transferred to a tared planchet for counting gross alpha and/or gross beta activity.

Reagents

Lucite: 0.5 mg/ml in acetone
Nitric acid, HNO₃: 3N
Nitric acid, HNO₃: concentrated

Apparatus

Filters; Millipore, membrane Type AA, 0.8 μ
Filtration equipment
Planchets (Standard 2" x 1/8" Beckman planchet)
Proportional counter

Procedure

1. Filter a volume of sample containing not more than 100 mg of dissolved solids for alpha assay, or not more than 200 mg of dissolved solids for beta assay.

Note: For gross alpha and gross beta assay in the same sample limit amount of solids to 100 mg.
2. Wash the non-filterable solids on the filter. (Save the filters with suspended matter for separate analyses. See Section 2.2.1).
3. Evaporate the filtrate to NEAR dryness on a hot plate. Add 25 ml concentrated HNO₃ and evaporate to NEAR dryness.

Note: For analysis of total residue (for clear water) proceed as described above but do not filter the water. Measure out the appropriate amount and proceed with step 3.

Section 2.2.2.(continued)

4. With distilled water and a few drops of 3N HNO₃, transfer the residue to a 50 ml beaker. Evaporate to NEAR dryness.
5. Transfer quantitatively the residue to a TARED PLANCHET, using an eye dropper.
6. Wash the beaker with distilled water and combine the washing and the residue in the planchet. Evaporate to dryness.
7. Bake in muffle furnace at 500° C for 45 min., cool and weigh.
8. Add a few drops (6-7 drops) of lucite solution and dry under the infrared lamp for 10-20 minutes.
9. Store the sample in a desiccator until it is to be counted.
10. Count the gross alpha and/or the gross beta activity in a low background proportional counter.
11. Calculate the activity in pCi/l using computer program OWATAB.

Calculations:

Gross alpha (beta) concentration:

$$(\text{pCi/liter}) = \frac{A}{B \times C \times D \times 2.22} + \frac{2 \sqrt{E_{sb}^2 + E_b^2}}{B \times C \times D \times 2.22}$$

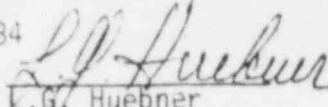
Where:

- A = net alpha (beta) count rate (cpm)
- B = efficiency for counting alpha (beta) activity (cpm/dpm)
- C = volume of sample (liters)
- D = correction factor for self-absorption in the sample
- E_{sb} = counting error of sample plus background
- E_b = counting error of background

Reference: Radioassay Procedures for Environmental Samples, U.S. Department of Health, Education and Welfare. Environmental Health Series, January 1967.

3.1 Airborne Particulates - Gamma Spectroscopic Analyses by Germanium Detector

1. Put the air filter in a filter cup container.
2. Place the filter cup inside the shield on the detector.
3. Count long enough to meet the LLD requirements.
4. Store the spectra on the disc.
5. After storing, calculate gamma activities using computer and corresponding calibrated geometry.
6. Return the filters to the original envelopes for storage or further analyses.



E.G. Huebner

3.2 Airborne Iodine Gamma Spectroscopic Analysis by Germanium Detector

NOTE: Because of the short half-life of I-131, count the samples as soon as possible after receipt and no later than 48 hours.

1. Load the charcoal cartridges in a specially designed holder or transfer charcoal from each cartridge to individual plastic bags. Seal the bags.
2. Label each bag with corresponding project ID, locations ID, and date of collection.
3. Place the bags in a standard geometry container, cap the container and secure the cap with a tape.
4. Place the holder or container on the detector and count for a period of time that will meet the required Lower Limit of Detection (LLD).

Calculation:

$$A_1 = \text{I-131 activity (pCi/sample)} = \frac{A}{2.22 \times B} \quad (\text{at counting time}) \quad (1)$$

Where:

A = net count rate of I-131 in the 0.36 MeV peak (cpm)
 B = efficiency for the I-131 in 0.36 MeV peak (cpm/dpm)

Correction for Equilibrium (assuming constant concentration over the sampling period) and Decay:

$$C = \frac{\lambda A_1 e^{\lambda t_1}}{F (1 - e^{-\lambda t_2})} \quad (2)$$

Where:

C = equilibrium concentration of I-131 (pCi/m³)
 A₁ = activity of I-131 at the time of counting (pCi/sample)
 e = the base of the natural logarithm = 2.71828
 λ = 0.693/half life (days) = 0.693/8.04 = 0.0862/day
 t₁ = elapsed time between the end of sampling and mid-counting point (in days)
 t₂ = duration of collection (in days)
 F = m³/day

3.3 Water - Gamma Spectroscopic Analyses by Germanium Detector

Procedure

1. Measure 3.5 liters of water into a Marinelli beaker.
2. Place the beaker inside the shield on the detector.
3. Count long enough to meet LLD requirements.
4. Store the spectrum on a disc.
5. After storing, calculate gamma activities, using computer program and corresponding calibrated geometry.
6. Transfer the sample back to the original container for further analyses.

3.4 Soil and Bottom Sediments - Gamma Spectroscopic Analyses by Germanium Detector

Procedure

1. Transfer the portion of the ground sample set aside for gamma scanning into a 500 ml Marinelli container.
2. Record the dry weight.
3. Place the container inside the shield on the detector.
4. Count the gamma activity long enough to meet the LLD requirements.
5. Store the spectrum on a disc.
6. After storing, calculate gamma activities using computer and corresponding calibrated geometry.
7. Transfer the sample back to the original container for further analyses.

3.5 Fish and Wildlife - Gamma Spectroscopic Analyses by Ge(Li) Detector

Procedure

1. Transfer a portion of the clean wet flesh of fish or animal into a 500 ml Marinelli container.
2. Record wet weight.
3. Add a few cc of formaldehyde and seal the container.
4. Place the container inside the shield on the detector.
5. Count long enough to meet the LLD requirements.
6. Store the spectrum on a disc.
7. After storing, calculate gamma activities using computer and corresponding calibrated geometry.
8. Transfer the sample back to the original container for further analyses.

3.6 Ambient Gamma Radiation

A. Thermoluminescent Dosimeters (TLD) - Light Response (Efficiency)

Harshaw Lithium Fluoride TLD-100 chips, 1/8" x 1/8" x 0.035".

Procedure

1. Rinse the chips with warm trichloroethylene followed by the methanol rinse. Dry.
2. Place the chips in a platinum crucible.
3. Anneal for 1 hour at 400°C.
4. Cool quickly by placing the crucible on a metal plate.
5. Anneal for 2 hours at 100°C.
Note: Avoid exposing the chips to the fluorescent light.
6. Seal 5 chips each in black plastic.
7. Mount the packs on the turntable.
8. Position the Ra-226 needle in the middle of the turntable and start rotation (appr. 60 revolutions per minute). Record the time.
9. Irradiate the chips for 2-6 hrs.
10. Remove the packages from the turntable. Return the Ra-226 needle to the lead container. Record the time.
11. Take the chips out of the plastic bag and place them in the vial.
12. Postanneal the chips for 10 minutes at 100°C.
13. Read each chip in the TLD Reader (For test procedure see "Performance Test Procedure for TLD Reader").
14. Calculate mean \pm one sigma deviation of five chips.
15. Calculate light response of TLD's (correction factor) by the following equation:

Section 3.6 (continued)Calculations

$$\text{C.F. (nanocoulombs/mR)} = \frac{A}{B}$$

Where:

C.F = correction factor (efficiency) to be applied in calculating exposure of field TLDs

A = Net reading in nanocoulombs

B = known exposure to TLDs

The exposure to the TLDs (B) is calculated as follows:

$$\text{mR/hr} = \frac{8400 \times \text{mg Ra-226}}{r^2}$$

For our setup use the following parameters:

$$\text{Ra-226} = 0.0933 \pm 1.5\%$$

$$r = 19.6 \text{ cm}$$

Thus:

$$\text{mR/hr} = \frac{8400 \times 0.0933}{384.16} = 2.040$$

The total exposure (B) is equal to:

$$B \text{ (mR)} = 2.040 \times \text{hours of exposure to the Ra-226 needle.}$$

3.7 Procedure for Preparation and Readout of TLD Chips

Materials

Harshaw Lithium Fluoride TLD-100 chips, 1/8" x 1/8" x 0.035".
Black plastic bags or boxes
Plastic sealer
Vacuum needle (for handling the chips)
TLD reader

Note: Never handle the chips with bare hands. Use plastic-covered forceps or vacuum needle. Handle them gently, e.g. do not drop them into the vial or on the table. They chip off easily, resulting in efficiency change.

Procedure

1. Rinse the chips with warm trichloroethylene followed by the methanol rinse. Dry.
2. Place the chips in a platinum crucible.
3. Anneal for 1 hour at 400°C.
4. Cool quickly by placing the crucible on a metal plate.
5. Anneal for 2 hours at 100°C.
6. Seal 3 to 5 chips (depending on the specifications) in black plastic or plastic boxes.
7. Label and send out by U.S. Mail.
8. Upon arrival at the lab, store TLDs in the big shield until readout day. Do not store longer than a few days.
9. Connect chips reader one day prior to readout.
10. Turn on gas for a few minutes before readout. Adjust to the mark.
11. Set parameter on the 2000P as follows:

HV - 470 V (It is 970 V, internal volts = 500).

Readout time: 20"

T₁ - 140° C (Preset)

T₂ - 250° C (Preset)

Rise time: -12°/sec (Preset)

Preheat - 100° C (Preset)

Start reading - 90° C

Section 3.7 (continued)

12. Prepare the chips as follows (do this before proceeding to the next step).
 - 12.1 Turn on small muffler furnace or drying oven and adjust to 100°C. Use glass thermometer. Muffler's indicator is not accurate. Let furnace stabilize.
 - 12.2 Unpack the chips (under reduced incandescent light) and gently place them in the glass vials marked with appropriate location numbers.
 - 12.3 Place the vials in the furnace. Preanneal for 10 min. at 100°C.
13. Open the drawer and read the standard. It should read 5.70±0.04. Adjust HV, if needed. Take 3 readings after final adjustment. Record.
14. Close the drawer.
15. Check bkg. It should read about 0.7-0.8 in 20". If it is higher, adjust the knob in the back of 2000 P (on left side when facing the instrument).

Note: Adjust bkg as low as possible but do not let the needle hit zero. The instrument will not record below zero.
16. Make 10 bkg readings (no chip in). Record. Read (do not record) at least 2 dummies to stabilize the temperature.
17. Place the chip in, wait until temperature goes down to 90° C and press "read" button. Make sure the chip is in the cavity of the heating plate.
18. After readout is completed, record the reading, open the drawer, and place next chip.
19. Repeat Steps 17 and 18 until all chips are read out.

Note: If reading will last longer than 1.5-2.0 hrs., check the standard and bkg about every 2.0 hrs.
20. After readout is completed, turn off the gas.
21. For calculations, use computer program OGTLD.PUB.

3.8 Tritium in Water (Direct Method)

Principle of Method

The water sample is purified by distillation, and portion of the distillate is transferred to a counting vial containing a scintillation fluid. The contents of the vial are then mixed and counted in a liquid scintillation counter.

Reagents

Scintillation medium, insta-gel scintillator
Tritium standard solution

Apparatus

Condenser
Distillation flask, 250-ml capacity
Liquid scintillation counter
Liquid scintillation counting vials

Procedure

1. Distill a 30 ml aliquot of the sample in a 250-ml distillation flask. Add a boiling chip to the flask. Connect a side arm adapter and a condenser to the outlet of the flask. Place a glass vial at the outlet of the condenser. Heat the sample to 100 - 150° C to distill, just to dryness. Collect the distillate for tritium analysis.
2. Dispense 13 ml of the distillate to a low potassium glass vial.
3. Prepare background and standard tritium-water solutions for counting, using the same amount as the sample. Use low tritium background distilled water for these preparations (distillate of most deep well water sources is acceptable, but each source should be checked for tritium activity before using).
4. Dark-adapt all samples, backgrounds, and standards. Add 10 ml of insta-gel scintillator. Count the samples, backgrounds and standards. Count samples containing less than 200 pCi/l for 300 minutes and samples containing more than 200 pCi/l for 200 minutes.

Section 3.8 (continued)

5. Counting efficiency:

$$\text{Eff} = \frac{\text{cpm of Standard} - \text{cpm of background}}{\text{dpm Standard}}$$

6. Sample Concentration:

$$\text{pCi/ml} = \frac{A}{2.22 \times E \times V \times e^{-\lambda t}}$$

Where:

A = net count rate (cpm)

E = efficiency (cpm/dpm)

V = volume (ml)

$$\lambda = \frac{0.693}{12.26} = 0.05652$$

t = elapsed time from the time of collection to the counting time (in years)

7. Calculate tritium activity using computer program H3.

3.9 Iodine-131 in Milk by Ion Exchange on Anion Exchange Column

After samples have been treated to convert all iodine in the sample to a common oxidation state, the iodine is isolated by solvent extraction or a combination of ion exchange and solvent extraction steps.

Iodine, as the iodide, is concentrated by adsorption on an anion exchanged column. Following a NaCl wash, the iodine is eluted with sodium hypochlorite. Iodine in the iodate form is reduced to I_2 and the elemental iodine extracted into CCl_4 , back-extracted into water then finally precipitated as palladium iodide.

Chemical recovery of the added carrier is determined gravimetrically from the PdI_2 precipitate. I-131 is determined by beta counting the PdI_2 .

Reagents

Anion exchange resin, Dowex 1-X8 (50-100 mesh) chloride form.

Carbon tetrachloride, CCl_4 - reagent grade.

Hydrochloric acid, HCl, 1N.

Hydrochloric acid, HCl, 3N.

H_2O - HNO_3 - NH_2OH HCl wash solution: 50 ml H_2O ; 10 ml 1M - NH_2OH -HCl; 10 ml conc. HNO_3 .

Hydroxylamine hydrochloride, NH_2OH HCl - 1 M.

Nitric acid, HNO_3 - concentrated.

Palladium chloride, PdI_2 , 20 mg Pd^{++}/ml . (1.2 g $PdCl_2/100$ ml 6N HCl).

Sodium bisulfite, $NaHSO_3$ - 1 M

Sodium chloride, NaCl - 2M

Sodium hypochlorite, NaOCl - 5% (Clorox).

Section 3.9 (continued)Special Apparatus

Chromatographic column, 20 mm x 150 mm (Reliance Glass Cat.#R2725T).

Vacuum filter holder, 2.5 cm² filter area

Filter paper, Whatman #42, 21 mm

Mylar

Polyester gummed tape, 1 1/2", Scotch #853

Drying oven

A. Ion Exchange Procedure

1. Set up an ion exchange column of 20 mm diameter and 150 mm length.
2. Pour 20 ml of a slurry of Dowex 1-X8, Cl⁻ form (50-100 mesh) into the column and wash down sides with water. Add 2 ml of I⁻ carrier to 2 liters milk, stir for 20 minutes.
3. Pass the sample through the ion exchange column at a flow rate of 20 ml/min. Save the effluent for other analyses and label it "iodine effluent".
4. Wash column with 500 ml of hot distilled water for milk samples or 200 ml of distilled water for water samples. Discard wash.
5. Wash column with 100 ml of 2 M NaCl at a flow rate of 4 ml/min. Discard wash.
6. Drain the solution from the column.
7. Measure 50 ml 5% sodium hypochlorite in a graduated cylinder. Add sodium hypochlorite to column in 10-20 ml increments, stirring resin as needed to eliminate gas bubbles and maintain flow rate of 2 ml/min. Collect eluate in 250-ml beaker and discard the resin.

B. Iodine Extraction Procedure

1. Acidify the eluate from step 7 using concentrated HNO₃ to make the sample 2-3 N in HNO₃, and transfer to 250 ml separatory funnel. (Add the acid slowly with stirring until the vigorous reaction subsides.) Volume of concentrated HNO₃ required will depend on eluate volume as follows):

Section 3.9 (continued)B. Iodine Extraction Procedure (continued)

eluate volume (ml)	concentrated HNO ₃ (ml)
50-60	10
60-70	12
70-80	14
80-90	16

2. Add 50 ml of CCl₄ and 10 ml of 1 M hydroxylamine hydrochloride (freshly prepared). Extract iodine into organic phase (about 2 minutes equilibration). Draw off the organic phase (lower phase) into another separatory funnel.
3. Add 25 ml of CCl₄ and 5 ml of 1 M hydroxylamine hydrochloride to the first separatory funnel and again equilibrate for 2 minutes. Combine the organic phases. Discard the aqueous phase (Upper phase) if no other analyses are required. If Pu, U or Sr is required on the same sample aliquot, submit the aqueous phase and data sheet to the appropriate laboratory section.
4. Add 20 ml H₂O-HNO₃-NH₂OH HCl wash solution to the separatory funnel containing the CCl₄. Equilibrate 2 minutes. Allow phases to separate and transfer CCl₄ (lower phase) to a clean separatory funnel. Discard the wash solution.
5. Add 25 ml H₂O and 10 drops of 1 M sodium bisulfite (freshly prepared) to the separatory funnel containing the CCl₄. Equilibrate for 2 minutes. Discard the organic phase (lower phase). Drain aqueous phase (upper phase) into a 100-ml beaker. Proceed to the Precipitation of PdI₂.

C. Precipitation of Palladium Iodide

CAUTION: AMMONIUM HYDROXIDE INTERFERES WITH THIS PROCEDURE

1. Add 10 ml of 3 N HCl to the aqueous phase from the iodine extraction procedure in step 5.
2. Place the beaker on a stirrer-hot plate. Using the magnetic stirrer, boil and stir the sample until it evaporates to 30 ml or begins to turn yellow.
3. Add 1.0 ml of 20 mg Pd⁺⁺/ml palladium chloride per liter of milk used dropwise, to the solution.

Section 3.9 (continued)C. Precipitation of Palladium Iodide (continued)

4. Turn the heat off, but continue to stir the sample until it cools to room temperature. Place the beaker in a stainless steel tray and put in the refrigerator overnight.
5. Weigh a clean 21 mm Whatman #42 filter which has been stored over silica gel in a desiccator.
6. Place the weighed filter in the filter holder. Filter the sample and wash the residue with water and then with absolute alcohol.
7. Remove filter from filter holder and place it on a stainless steel planchet.
8. Dry under the lamp for 20 minutes.
9. Cut a 1 1/2" strip of polyester tape and lay it on a clean surface, gummed side up. Place the filter, precipitate side up, in the center of the tape.
10. Cut a 1 1/2" wide piece of mylar. Using a spatula to press it in place, put it directly over the precipitate and seal the edges to the polyester tape. Trim to about 5 mm from the edge of the filter with scissors.
11. Mount the sample on the plastic disc and write the sample number on the back side of the disc.
12. Count the sample on a proportional beta counter.

Calculations

Calculate the sample activity using computer program I131.

Reference: "Determination of 1-131 by Beta-Gamma coincidence Counting of PdI₂". Radiological Science Laboratory. Division of Laboratories and Research, New York State Department of Health, March 1975, Revised February 1977.

Section 8.1

8.1 Strontium-89 and Strontium-90 in Milk by Ion Exchange

Principle of Method

A citrate complex of yttrium, strontium, and barium carriers at the pH of milk is added to the milk sample. The mixture is then passed successively through cation- and anion-exchange resin columns. Strontium, barium, and calcium are absorbed on the cation-exchange resin, and the yttrium carrier with the yttrium 90 daughter of strontium 90 is retained on the anion-exchange resin.

The yttrium is eluted from the anion resin with hydrochloric acid and precipitated as the oxalate. Lanthanum 140, which may be a contaminant, is removed by dissolving yttrium oxalate in concentrated nitric acid and extracting yttrium from the solution into an equal volume of pre-equilibrated tributyl phosphate. The lanthanum 140 remains in the concentrated nitric acid to be discarded. Yttrium is re-extracted from the organic phase with dilute nitric acid and precipitated as the oxalate. The precipitate is weighed to determine recovery of yttrium carrier, then counted for yttrium 90 activity.

Strontium, barium, and calcium are eluted from the cation-exchange resin with sodium chloride solution. Following dilution of the eluate, the alkaline earths are precipitated as carbonates. The carbonates are then converted to nitrates, and strontium and barium nitrate are precipitated. The nitrate precipitate is dissolved, and barium is precipitated as the chromate, purified as the chloride, and then counted to determine the barium 140. From the supernate, strontium is precipitated as the nitrate, dissolved in water, and reprecipitated as strontium nitrate. The nitrate is converted to the carbonate, which is filtered, weighed to determine strontium carrier recovery, and counted for "total radiostrontium".

The concentration of strontium-89 is calculated as the difference between the activity for "total radiostrontium" and the activity due to strontium-90.

Reagents

Ammonium acetate buffer: pH 5.0

Ammonium hydroxide, NH₄OH: concentrated (15N)

Ammonium oxalate, (NH₄)₂C₂O₄.H₂O: 1N

Anion-exchange resin: Dowex 1-X8 (Cl⁻ form, 50-100 mesh)

Carrier solutions:

Ba⁺² as barium nitrate, Ba(NO₃)₂: 20 mg Ba⁺² per ml

Sr⁺² as strontium nitrate, Sr(NO₃)₂: 20 mg Sr⁺² per ml

Y⁺³ as yttrium nitrate, Y(NO₃): 10 mg Y⁺³ per ml

Cation-exchange resin: Dowex 50W-X8 (Na⁺ form, 50-100 mesh)

Citrate solution: 3N (pH 6.5)

Section 8.1 (Continued)Diethyl ether, $(C_2H_5)_2$:anhydrousEthyl alcohol, C_2H_5OH : absolute (100%), 95%Hydrochloric acid, HCl : concentrated (12N, 6N*, 2N*)Hydrochloric acid-diethyl ether, $HCl-(C_2H_5)_2O$:5.1 v/vNitric acid, HNO_3 :fuming (90%), concentrated (16N)*, 14N, 6N, 0.1 N*Oxalic acid, $H_2C_2O_4 \cdot 2H_2O$:2N*Sodium carbonate, Na_2CO_3 :3N, 0.1 NSodium chloride, $NaCl$:4NSodium chromate, Na_2CrO_4 :3NTri-n-butyl phosphate (TBP), $(C_4H_9)_3PO_4$:pre-equilibrated with 14N HNO_3 *

* Starred reagents are used only in processing the anion column effluent to determine strontium-90 concentration (Part A).

Apparatus

Ion-exchange system: The apparatus for this system is illustrated in Figure 8.1-1. It consists of three glass components connected one above the other for gravity flow. At the top is a graduated, 1-liter glass separatory funnel which serves as the reservoir. Below it is connected a 250 ml glass column, 5 cm in diameter and 25 cm long, which services as the cation column. Below this is connected the anion column, a 30-ml glass column, 1.9 cm in diameter and 10.5 cm long. Both columns have extra coarse, fritted glass disks at the bottom.

Five milliliters of distilled water are placed in the 30-ml column, and 15 ml Dowex 1 resin are poured into it. The cation column is filled by adding 170 ml Dowex 50W resin in the same way.

Millipore filtering apparatus (Pyrex Hydrosol Microanalysis Filter Holder)

Millipore Type OH membrane filter, 1.5- μ pore size, 2.5-cm diameter low-background beta counter.

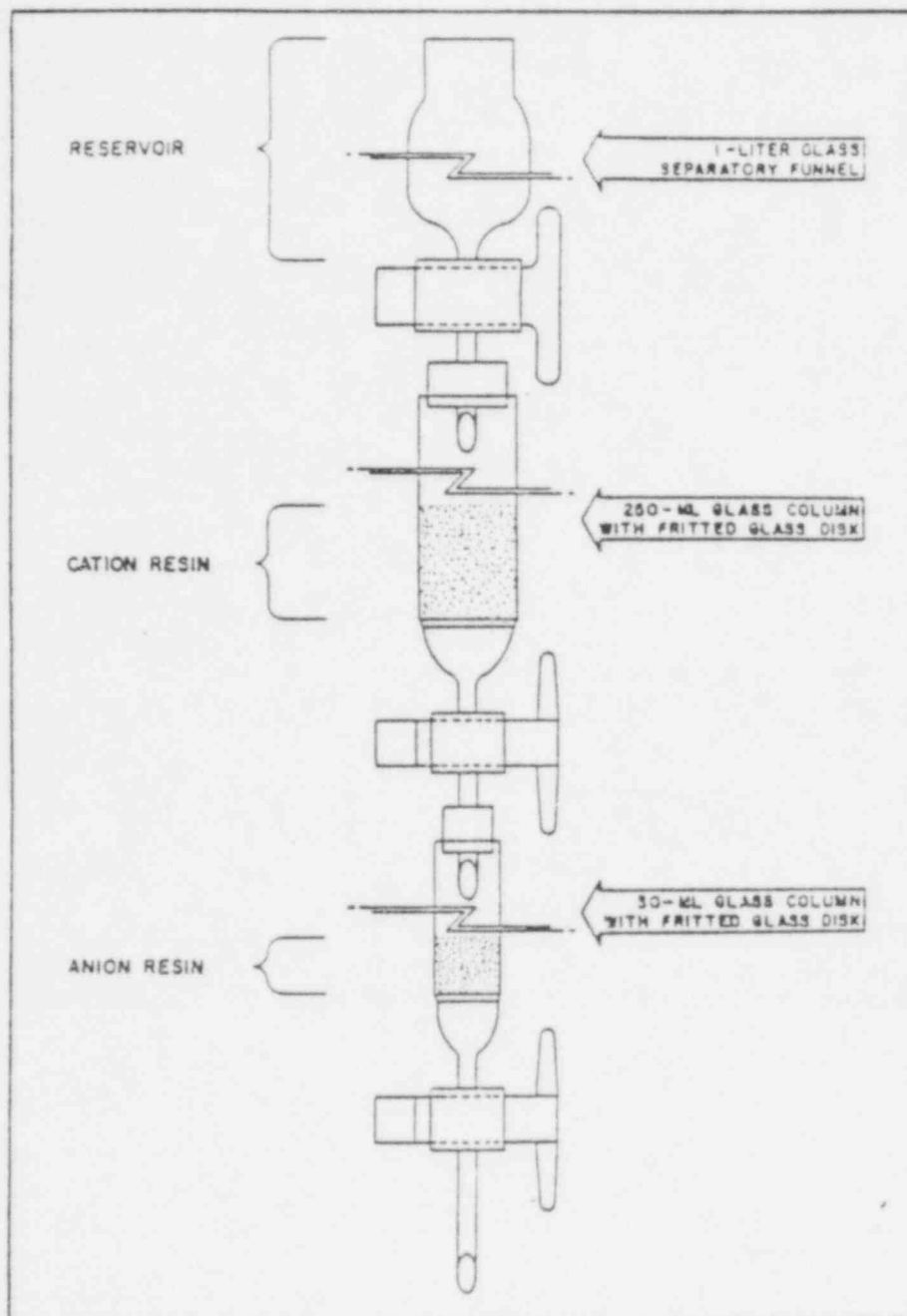


Figure 8-4 Ion-exchange system

Section 8.1 - Part APart A. Strontium-90Procedure

1. Place 1 liter of milk into the graduated reservoir. Pipette 1.0 ml each of yttrium, strontium, and barium carrier solutions into 10 ml of citrate solution: swirl to mix and dissolve the barium citrate which forms. Transfer this mixture quantitatively to the milk with 5 ml of distilled water, and mix well.
2. Open the stopcocks of the reservoir, anion column, and cation column, in that order. NOTE THE TIME. Control the flow rate at 10 milliliters per minute (ml/min) with the anion column stop-cock. Check occasionally by collecting effluent for 1 minute in a graduated cylinder. Stop flow when just enough milk remains in the columns to cover resin. NOTE THE TIME. Discard the effluent milk. RECORD THE MIDPOINT OF THE ELUTION PERIOD AS THE BEGINNING OF YTTRIUM 90 DECAY.
3. Replace the milk reservoir with a separatory funnel containing 300 ml of warm distilled water, and let the water flow through the columns at approximately 10 ml/min to displace the milk. Stop the flow when just enough water remains in the columns to cover the resin. Discard the effluent water.
4. Separate the columns.
In order to collect eluate for "total radio-strontium", barium, and calcium determinations, and to regenerate the cation column for subsequent use, follow Step 5, Part B.
5. Gradually add 75-100 ml of 2N HCl to the anion column. Control the effluent flow at 2 ml/min. Collect eluate in a 250-ml centrifuge bottle.
6. Add 5 ml of 2N oxalic acid to the eluate and adjust the pH to 1.5 with 6N NH_4OH using a pH meter.
7. Stir and heat to near boiling in a water bath (approx. 20 min).
8. Cool in an ice bath and centrifuge. Decant and discard the supernatant. Proceed as in (a) or (b) depending on whether Ba-La-140 is absent or present from the gamma analysis of the sample.

(a) If fresh fission products are known to be absent:

Dissolve the ppt in 10 ml of HNO_3 , filter solution through Whatman No. 541 paper into a 40 ml centrifuge tube. Wash paper, collecting the washing in tube and continue as in Step A-9.

Section 8.1 - Part A (Continued)(b) If fresh fission products are present:

Dissolve the ppt in 10 ml of HNO_3 , transfer the solution to a 60 ml separatory funnel, washing the tube with additional 10 ml of HNO_3 . Add 10 ml of equilibrated TBP, shake 2-3 min, and when separated drain and discard the lower acid phase. Add 15 ml of 14N HNO_3 to the separatory funnel, shake 2-3 min, drain and discard the lower acid phase. Repeat the 14N HNO_3 treatment to remove eight lanthanide elements and La-140 . Add 15 ml of H_2O to the separator and shake. Drain the lower phase into a 125-ml centrifuge tube. Repeat the wash, using 15 ml of 0.1N HNO_3 , adding it to the centrifuge tube.

9. Add 5 ml of 2N oxalic acid to the purified yttrium solution from (a) or (b). Adjust to a pH of 1.5 with NH_4OH , using a pH meter.
10. Digest the solution in a hot H_2O bath for 10 min. with occasional stirring. Cool in an ice bath (20 min).
11. Filter on a weighed Whatman No. 42 (2.1 cm) filter paper. Wash with H_2O , ethyl alcohol and ether and dry at room temperature and weigh.
12. Mount and count in a proportional counter.
13. If analysis for Sr-89 is not required, disregard Section 8.1-Part B. Use the computer program SR8990 to calculate (Sr-90) activity.

Section 8.1 - Part BPart BTotal Radiostrontium (Sr-89 separation)Procedure

Continue following columns separation (Step A-4).

5. Connect 1 l separator funnel containing 1 l of 4N NaCl to the cation column. Allow the solution to flow at 10 ml/min to elute the alkali metal and alkaline earth ions and to recharge the column. Collect 1 l of eluate into a 2 l beaker, but leave the resin covered with 2-3 ml of solution.
6. Wash the column with 500 ml of H₂O or more to remove excess NaCl. Discard the wash.
7. Remove 20 ml of the NaCl eluate into a small bottle for the determination of stable calcium. (See section 6.1).
8. Dilute the eluate to 1500 ml with distilled water.
9. Heat the solution to 85°-90° C (near boiling on a hot plate) and add, with constant stirring, 100 ml of 3N Na₂CO₃. Stir gently while on hot plate to prevent bumping. Let stand overnight.
10. Decant most of the supernate. Transfer the precipitate to a 250 ml centrifuge bottle.
11. Wash the precipitate twice with 50 ml portions of H₂O. Dry it in an oven at 110° C for 1-2 hours.
12. Dissolve the ppt slowly with vigorous stirring in 10 ml of 6N HNO₃ (with magnetic stirrer). Filter through Whatman No. 541 paper into a 40 ml centrifuge tube. Rinse the bottle with little 6N HNO₃ and pour the washings through the paper. To the filtrate, add slowly 30 ml of 21N HNO₃ (fuming). Stir well and cool in an ice bath. Centrifuge and discard supernatant.
13. Carefully add 30 ml of conc. HNO₃ to the precipitate. Heat in a H₂O bath with stirring for about 30 minutes. Cool the solution in an ice water bath for about 5 minutes. Centrifuge and discard supernatant.
14. Repeat step No. 13.

Section 8.1 - Part B (Continued)

15. Dissolve the ppt. in 10 ml. of H₂O and 5 ml. of NH₄AC buffer and heat in a water bath: Adjust pH to 5.5 using a pH meter and add immediately 1 ml. of 3N Na₂CrO₄ and mix well. Digest in a water bath for 5 min., centrifuge and decant the supernatant into another 40 ml. centrifuge tube.
16. Heat the supernate in a water bath. Adjust the pH to 8-8.5 with NH₄OH. With continuous stirring, cautiously add 5 ml of 3N Na₂CO₃ solution. Heat gently for 10 minutes. Centrifuge and decant the supernate. Wash the strontium carbonate precipitate with 0.1 N Na₂CO₃. Centrifuge again, and decant the supernate.
17. Dissolve the carbonate precipitate in 5 ml of 6N HNO₃. With continuous stirring, cautiously add 30 ml of fuming HNO₃ to the solution. (Stirring the solution longer helps in the precipitation of the strontium nitrate.) Cool in ice bath, centrifuge and decant the supernate.
18. Dissolve the strontium nitrate precipitate in 3 ml of H₂O and 5 ml of 6N HNO₃. Add cautiously, with continuous stirring, 20 ml of fuming HNO₃. Cool in an ice bath, centrifuge and discard supernatant. RECORD TIME AS BEGINNING OF Y-90 INGROWTH.
19. Dissolve the precipitate in 10 ml of H₂O. Heat in a water bath. Adjust the pH to 8-8.5. With continuous stirring, add 5 ml of 3 N Na₂CO₃ solution. Heat gently for 10 minutes.
20. Cool and filter on a weighed No. 42 Whatman (2.1 cm) filter paper. Wash thoroughly with water and alcohol.
21. Dry the precipitate in an oven at 105° C or under the lamp for 30 minutes. Cool and weigh.
22. Mount and count without delay in a proportional counter as total strontium.
23. Calculate Sr-89 and Sr-90 activity (pCi/l) using computer program SR8990.

Section 8.1 (Continued)

Calculations

Part A.

$$\text{Strontium 90 concentration (pCi/liter)} = \frac{A}{B \times C \times D \times E \times F}$$

Where:

- A = net beta count rate of yttrium 90 (cpm)
- B = recovery of yttrium carrier
- C = counter efficiency for counting yttrium-90 or yttrium oxalate mounted on a 2.1-cm diameter membrane filter (cpm/pCi)
- D = sample volume (liters)
- E = Correction factor $e^{-\lambda t}$ for yttrium-90 decay, where t is the time from midpoint of the elution time of milk (Step A-2) to the time of counting.
- F = Correction factor $1 - e^{-\lambda t}$ for the degree of equilibrium attained during the yttrium-90 ingrowth period, where t is the time from collection of the milk sample to the time of passage through the column (Step A-2)

Part B.

$$\text{Strontium 89 concentration (pCi/liter)} = \frac{1}{B \times C} \left(\frac{A}{D \times E} - F (G \times H + I \times J) \right)$$

Where:

- A = net beta count rate of "Total radiostrontium" (cpm)
- B = counter efficiency for counting strontium-89 as strontium oxalate mounted on a 2.1-cm diameter membrane filter (cpm/pCi)
- C = correction factor $e^{-\lambda t}$ for strontium-89 decay, where t is the time from sample collection to the time of counting
- D = recovery of strontium carrier
- E = volume of milk sample (liters)
- F = strontium 90 concentration (pCi/liter) from Part A
- G = self-absorption factor for strontium-90 as strontium oxalate mounted on a 2.1-cm diameter filter, obtained from a self-absorption curve prepared by plotting the fraction of a standard activity absorbed against density thickness of the sample (mg/cm^2)
- H = counter efficiency for counting strontium-90 as strontium oxalate mounted on a 2.1-cm diameter membrane filter (cpm/pCi)
- I = counter efficiency for counting yttrium-90 as yttrium oxalate mounted on a 2.1-cm diameter membrane filter (cpm/pCi)

Section 8.1 (Continued)

J = correction factor $1-e^{-\lambda t}$ for yttrium-90 ingrowth, where t is the time from the last decantation of the nitric acid (Step B-18).

Reference: Radioassay Procedures for Environmental Samples U.S. Department of Health, Education and Welfare. Environmental Health Series, January 1967.

Section 8.4

8.4 Strontium 89 and Strontium 90 in Water Samples

A. Principle of Method

The acidified sample of clear water with stable strontium, barium and calcium carriers is treated with oxalic acid at a pH of 3.0 to precipitate insoluble oxalates. The oxalates are dissolved in nitric acid and strontium nitrate is separated from calcium as a precipitate in 70% nitric acid. The residue is purified by adding iron and rare earth carriers and precipitating them as hydroxides. After a second strontium nitrate precipitation from 70% nitric acid, the nitrates are dissolved in water and with added yttrium carrier, are stored for ingrowth of yttrium-90. The strontium is again precipitated and separated from 70% nitric acid with the yttrium nitrate being in the supernate. Each fraction is precipitated separately as an oxalate and collected on No. 42 (2.1 cm) Whatman filter or planchet for counting either total radiostrontium or yttrium-90 or both.

Reagents

Acetic acid, CH₃COOH: 1.5N

Ammonium acetate, NH₄C₂H₃O₂: 3N

Ammonium acetate buffer: pH 5.0

Ammonium hydroxide, NH₄OH: concentrated (15 N), 6 N, 1 N

Ammonium oxalate, (NH₄)₂C₂O₄.H₂O: 0.5% w/v

Carrier solutions:

Ba²⁺ as barium nitrate, Ba(NO₃)₂: 20 mg Ba²⁺ per ml

Ca²⁺ as calcium nitrate, Ca(NO₃)₂.4H₂O: 40 mg Ca²⁺ per ml

Sr²⁺ as strontium nitrate, Sr(NO₃)₂: 20 mg Sr²⁺ per ml

Y³⁺ as yttrium nitrate, Y(NO₃)₃: 10 mg Y³⁺ per ml

Hydrochloric acid, HCl: concentrated (12 N), 0.5 N

Hydrogen peroxide, H₂O₂: 30% solution

Nitric acid, HNO₃: fuming (90%), concentrated (16 N), 6 N, 3 N

Oxalic acid, H₂C₂O₂. 2H₂O: Saturated at room temperature

Scavenger solutions: 20 mg Fe³⁺ per ml, 10 mg each Ce³⁺ and Zr⁴⁺ per ml

Fe³⁺ as ferric chloride, FeCl₃.6H₂O

Ce³⁺ as cerous nitrate, Ce(NO₃)₃.6H₂O

Zr⁴⁺ as zirconyl chloride, ZrOCl₂.8H₂O

Sodium Carbonate, Na₂CO₃: 3N, 0.1N

Sodium Chromate, Na₂CrO₄: 3N

Apparatus

Analytical balance

Low background beta counter

Medium - porosity filter stick

pH meter

Section 8.4 APart A. Strontium 89Procedure

1. Filter 1 liter of an acidified water sample using millipore filter paper.
2. Digest the filter paper with the residue with concentrated nitric acid (HNO_3) until all the organic matter is removed.
3. Evaporate to dryness and dissolve the residue with hot water and filter using No. 541 Whatman filter paper.
4. Combine the filtrates in a 2 liter beaker.
5. Add 1 ml of strontium carrier solution, 1 ml barium carrier solution, and if necessary, 1 ml of calcium carrier solution. (Improved precipitation may be obtained by adding calcium to soft waters.) Stir thoroughly and while stirring add 125 ml of saturated oxalic acid solution.
6. Using a pH meter, adjust the pH to 3.0 with 15 N NH_4OH , and allow the precipitate to settle for 5-6 hours.
7. Decant most of the supernate (liquid) and transfer the precipitate to a 250 ml centrifuge bottle. Wash the precipitate and the beaker wall with 0.5% ammonium oxalate and centrifuge. Discard the supernate.
8. Dissolve the precipitate with 10 ml of 6 N HNO_3 and transfer to a 250 ml beaker. Then use 20 ml of 16 N HNO_3 to rinse the centrifuge tube and combine it to the solution in the 250 ml beaker.
9. Evaporate the solution to dryness. Cool; then add 50 ml 16 N HNO_3 and repeat the acid addition and evaporation until the residue is colorless.
10. Transfer the residue to a 40-ml centrifuge tube, rinsing with a minimum volume of 16 N HNO_3 . Cool in a refrigerator overnight. Centrifuge at 1500-1800 rpm for 10 minutes, and discard the supernate.
11. Dissolve the precipitate in 5 ml of 6N HNO_3 and then add 30 ml of fuming nitric acid. Centrifuge, and discard the supernate.
12. Dissolve the nitrate precipitate in about 10 ml of distilled water. Add 1 ml of scavenger solution. Adjust the pH of the mixture to 7 with 6 N NH_4OH . Heat, stir, and filter through a Whatman No. 541 filter. Discard the mixed hydroxide precipitate.

Section 8.4 A (continued)Part A. Strontium 89Procedure (continued)

13. To the filtrate, add 5 ml of ammonium acetate buffer. Adjust the pH with 3N HNO₃ or NH₄OH to pH 5.5. (Note: the pH of the solution at this point is critical.) Add dropwise with stirring 1 ml of 3N Na₂CrO₄ solution. Heat in a water bath.
14. Cool and centrifuge. Decant the supernate into another centrifuge tube. Save the precipitate for Ba analysis if needed.
15. Heat the supernate in a water bath. Adjust the pH to 8-8.5 with NH₄OH. With continuous stirring, cautiously add 5 ml of 3N Na₂CO₃ solution. Heat gently for 10 minutes. Cool, centrifuge, and decant the supernate. Wash the precipitate with 0.1N Na₂CO₃. Centrifuge again and decant the supernate.
16. Dissolve the precipitate in no more than 4 ml of 3N HNO₃. Then add 20-30 ml of fuming HNO₃, cool in a water bath, and centrifuge. Decant and discard the supernate.
17. Repeat step 16. Then, RECORD THE TIME AND DATE AS THE BEGINNING OF YTTRIUM 90 INGROWTH. If no immediate count of total radiostrontium is desired add to the precipitate 1 ml of yttrium carrier solution and 4 ml of 6N HNO₃ and store 7-14 days to allow the yttrium 90 to grow in.
18. To determine total radiostrontium, dissolve the precipitate in 10 ml of water. Heat in water bath. Adjust the pH to 8-8.5. With continuous stirring add 5 ml of 3N Na₂CO₃ solution. Heat gently for 10 minutes.
19. Cool and filter on a weighed No. 42 (2.1 cm) Whatman filter paper. Wash thoroughly with water and alcohol.
20. Dry the precipitate under the lamp for 30 min. Cool and weigh.
21. Mount and count without delay its beta activity as "total radiostrontium" in a proportional counter.

Section 8.4Part B. Strontium 90Procedure

1. After counting total radiostrontium dissolve the precipitate on the filter in 6 N HNO_3 and transfer the solution to a 40 ml centrifuge tube. The total volume of dissolution and rinsing should be about 4 ml.
2. Add 1 ml of yttrium carrier solution and store until 7 to 14 days have elapsed since step 17 was completed.
3. Heat the equilibrated strontium-yttrium sample in a water bath at approximately 90°C . Adjust the pH to 8 with NH_4OH , stirring continuously.
4. Cool to room temperature in a cold water bath and centrifuge for 5 minutes. Record the hour and date of decantation as the end of the yttrium-90 ingrowth and the beginning of its decay in the yttrium fraction.
5. Dissolve by adding about 4 drops of HCl with stirring. Add 15-20 ml of water. Heat in a water bath and adjust the pH to 8 with NH_4OH , stirring continuously.
6. Cool to room temperature in a cold water bath and centrifuge for 5 minutes.
7. Repeat steps 5 and 6.
8. Add 3 drops of HCl to dissolve the precipitate, then add 20 ml of water. Filter using No. 541 filter paper. Heat in a water bath at approximately 90°C . Add 1 ml of saturated oxalic acid solution dropwise with vigorous stirring. Adjust to a pH of 2-3 with NH_4OH . Allow the precipitate to digest for about an hour.
9. Cool to room temperature in a cold water bath. Centrifuge for 10 minutes and decant most of the supernate. Filter by suction on a weighed filter paper. Wash the precipitate with water and alcohol.
10. Dry the precipitate under the lamp for 30 minutes. Cool and weigh. Mount and count without delay in a proportional counter.
11. Calculate Sr-89 and Sr-90 activity in pCi/l using the computer program for Sr-89,90.

Section 8.4 (continued)

Part B. Strontium 90

Calculations

For formulas used refer to Section 8.1.

Reference: Radioassay Procedures for Environmental Samples U.S. Department of Health, Education and Welfare. Environmental Health Series, January 1967.

Section 8.6

8.6 Strontium-89 and Strontium-90 in Milk (Ash), Vegetation, Fish, Wildlife, Soil and Bottom Sediment Samples - Sodium Carbonate Fusion.

Principle of Method

Strontium is separated from calcium, other fission products, and other natural radioactive elements. Fuming nitric acid separations remove the calcium and most of the other interfering ions. Radium, lead, and barium are removed with barium chromate. Traces of other fission products are scavenged with yttrium hydroxide. After the Sr-90 and Y-90 equilibrium has been attained, the Y-90 is precipitated as the hydroxide and converted to the oxalate for counting. Strontium is precipitated as the carbonate and counted for total activity. Strontium-89 activity is computed as the difference between the total radiostrontium and the strontium-90 (as yttrium-90) activity.

Reagents

Ammonium acetate buffer, (NH₄)₂ Ac:pH = 5.0, 6M

Ammonium hydroxide, NH₄OH:6N

Carrier Solutions:

Ba⁺², Ba(NO₃)₂:20 mg/ml of Ba⁺²

Fe⁺³, Fe(NO₃)₃, scavenger:5 mg/ml of Fe⁺³

Sr⁺², Sr(NO₃)₂:20 mg/ml of Sr⁺²

Y⁺³, Y(NO₃)₃:10 mg/ml of Y⁺³

Ethyl alcohol, C₂H₅OH:absolute

Hydrochloric acid, HCl:12N (conc.)

Nitric acid, HNO₃:16N (conc.), 6N, 3N, fuming

Oxalic acid, H₂C₂O₄:saturated

Potassium nitrate, KNO₃:powdered

Sodium carbonate, Na₂CO₃:powdered, 3N, 0.1N

Sodium chromate, Na₂CrO₄:3N

Sodium hydroxide, NaOH:pelTets

Apparatus

Teflon filter holder, or filter funnel and sample mount rings and discs

Magnetic stirrers with Teflon-Coated magnet bars

Mylar film

Glass fiber filters

Fisher filtrator

Brinkman dispenser - pipettor

Section 8.6 APart A. Sample Preparation - Sodium Carbonate FusionProcedure

1. Weigh out 3 g of ashed sample or silted soil and set aside.
2. Sift into a 250 ml nickel crucible enough Na_2CO_3 to very lightly cover the bottom.
3. Add 30 g of NaOH pellets and 5 g of KNO_3 .
4. Add the weighed ash sample and tap the crucible gently to shake the ash down among the pellets.
5. Sift from 10 to 20 grams of Na_2CO_3 over the ash so it is completely covered.
6. Place in a muffle furnace at 600°C for 20 to 30 minutes to melt and fuse the mixture.

NOTE: If carbon materials remain floating on the surface of the melt, cautiously add a few grains of KNO_3 and heat for another 5 to 10 minutes.

Decomposition of organic matter is complete when no further reaction is noticed on addition of KNO_3 .

7. Using a long-handled tongs, remove the crucible from the muffle furnace and immediately, but very cautiously, cool in an ice bath until the melt is completely solidified and cool enough to handle without gloves.

NOTE: It is very important that no moisture come in contact with the melt at this time. One drop of water in the crucible could render the melt very difficult, if not impossible, to remove.

8. Transfer the melt to a 250 ml centrifuge bottle using distilled water and stir until completely dispersed.

NOTE: Rotating the crucible in the palm of one's hand and very gently applying pressure should be sufficient to loosen the melt from the sides of the crucible.

9. Add 2 ml of strontium and 1 ml of barium carriers.
10. Bring to a gentle boil, cool, centrifuge and discard the supernatant.

Section 8.6 A (continued)Part A. Sample Preparation - Sodium Carbonate FusionProcedure (continued)

11. To the residue add 50 ml 3N Na_2CO_3 as a wash, swirl and disperse the residue, heat for 10 minutes in a hot water bath, centrifuge and discard the supernatant.
12. Repeat step (11) three times to put the precipitate in a suitable form for further analysis.
13. Dissolve the precipitate in 50 ml of concentrated HNO_3 , transfer to a 250 ml beaker, and take to dryness on a hot plate.

NOTE: Evaporation may be done rapidly at first, and then very slowly to prevent spattering.

A jelly-like substance may form at this point, due to hydrated silicic acid formed from the soluble silicates and will be removed in the following steps.

14. Bake the remaining residue for at least 1 hour at 120° to 130° C, cool, moisten the salts with 5 ml of HNO_3 and allow to stand at room temperature for 10 minutes. Then place on a hot plate, bring to a boil and add 45 ml of boiling water. DISPERSE ANY REMAINING RESIDUE WITH A GLASS STIRRING ROD AND FILTER IMMEDIATELY into a 250 ml beaker. Use Whatman No. 541 hardened filter paper.

NOTE: To separate the silicic acid the hydrated acid must be changed to a less hydrated and less soluble acid by baking at 100° to 130°C .

It is important at this point that evaporation be to complete dryness. (There should no longer be a smell of acid).

Addition of 5 ml of HNO_3 converts any metal oxides which may have been formed back to nitrates so they will be dissolved and not removed with the silicates.

Filtering must be done immediately as some of the silicates will tend to go back into solution. Also, due to this fact, removal of silicates by dehydration is not 100% efficient and the process must be repeated at least once and more often if necessary.

15. Evaporate and repeat step (14) at least once, a gain as often as necessary.

Section 8.6 A (continued)Part A. Sample Preparation - Sodium Carbonate FusionProcedure (continued)

16. Evaporate the solution in a beaker to dryness on a hot plate. Cool, then add 40 ml of concentrated HNO_3 and evaporate to 20-25 ml. Then add another 40 ml HNO_3 and repeat the procedure.

NOTE: The liquid portion of the sample at this point will be yellow. Should the color toward the end of the first evaporation be red-brown, or black, add more nitric acid and repeat the above procedure as often as necessary to obtain a clear yellow solution.

The dark samples described above have been known to explode if evaporated to dryness without adding additional portions of nitric acid. These samples should be handled in a hood with the window down as far as possible to prevent possible personal injury to the operator.

This step is to destroy any remaining organic materials. The darker colored solutions contain large amounts of organic matter.

17. Complete the analyses as described under Determination.

References: The basis for this procedure was presented by J.J. Bolan in the Public Health Service Manual, titled "Chemical Analysis of Environmental Radionuclides, Determination of radiostrontium in food" (1.11.3.A(8.65)). Modifications to this procedure were made by the North Dakota State Department of Health.

Section 8.6 BPart B. DeterminationI. Strontium - 89Procedure

1. Transfer the solution to a 40 ml conical, heavy-duty centrifuge tube using a minimum of conc. HNO_3 . Cool the centrifuge tube in an ice bath for about 10 minutes. Centrifuge and discard the supernatant.

NOTE: The precipitate consists of calcium, strontium and barium-radium nitrates. The supernatant contains part of the sample's calcium and phosphate content.

2. Add 30 ml of conc. HNO_3 to the precipitate. Heat in a hot water bath with stirring for about 10 minutes. Cool the solution in an ice bath with stirring for about 5 minutes. Centrifuge and discard the supernatant.

NOTE: Additional calcium is removed from the sample. Nitrate precipitations with 70% HNO_3 will afford a partial decontamination from soluble calcium while strontium, barium, and radium are completely precipitated.

The separation of calcium is best at 60% HNO_3 , however at 60% the precipitation of strontium is not complete. Therefore, it is common practice to precipitate $\text{Sr}(\text{NO}_3)_2$ with 70% HNO_3 which is the concentration of commercially available 16 N HNO_3 .

Most of the other fission products, induced activities and actinides are soluble in concentrated HNO_3 affording a good "gross" decontamination step from a wide spectrum of radionuclides. The precipitation is usually repeated several times.

3. Repeat step (2) two more times.
4. Dissolve the nitrate precipitate in about 10 ml distilled water. Add 1 ml of scavenger solution. Adjust the pH of the mixture to 7 with 6 N NH_4OH . Heat, stir, and filter through a Whatman No. 541 filter. Discard the mixed hydroxide precipitate.

Section 8.6Part B Determination1. Strontium-89Procedure (continued)

5. To the filtrate add 5 ml of ammonium acetate buffer (pH 5.0). Adjust the pH to 5.5 with 3N HNO₃ or 6N NH₄OH. (Note: The pH of the solution at this point is critical. Barium chromate will not precipitate completely in more acidic solution and strontium will partially precipitate in more basic solutions.) Add dropwise with stirring 1 ml of 3N Na₂CrO₄ solution. Heat in a water bath to about 90°C and centrifuge. Decant the supernate into another centrifuge tube. Save the precipitate for Ba analysis if needed.
6. Heat the supernate in a water bath. Adjust the pH to 8-8.5 with NH₄OH. With continuous stirring, cautiously add 5 ml of 3N Na₂CO₃ solution. Heat gently for 10 minutes. Centrifuge, and when completeness of precipitation has been verified by adding a few drops of Na₂CO₃, centrifuge and decant the supernate. Wash the strontium carbonate precipitate with 0.1N Na₂CO₃. Centrifuge again, and decant the supernate.
7. Dissolve the carbonate precipitate in 5 ml 6N HNO₃. With continuous stirring, cautiously add 20 ml fuming HNO₃ to the solution. (Stirring the solution longer helps in the precipitation of strontium nitrate). Cool in an ice bath, centrifuge and decant the supernate.
8. Dissolve the strontium nitrate precipitate in 3 ml H₂O and 5 ml 6N HNO₃. Add cautiously, with continuous stirring, 20 ml fuming HNO₃. Cool in ice bath, centrifuge and discard supernatant. RECORD TIME AS BEGINNING OF Y-90 INGROWTH.
9. Dissolve the precipitate in 10 ml of H₂O. Heat in a water bath. Adjust the pH to 8-8.5. With continuous stirring, add 5 ml of 3N Na₂CO₃ solution. Heat gently for 10 minutes.
10. Cool and filter on a weighed No. 42 (2.1 cm) Whatman filter paper. Wash thoroughly with water and alcohol.
11. Dry the precipitate under the lamp for 30 minutes. Cool and weigh.
12. Mount and count without delay in a proportional counter as total radiostrontium.

Section 8.6Part B DeterminationIi. Strontium-90Procedure

1. After counting total radiostrontium, dissolve the strontium carbonate precipitate on the filter in 6N HNO₃ and transfer the solution to a 40 ml centrifuge tube. The total volume of dissolution and rinsing should be about 4 ml.
2. Add 1 ml of yttrium carrier solution and store until 7 to 14 days have elapsed since Step B-I-8 was completed.
3. Heat the equilibrated strontium-yttrium sample in a water bath at approximately 90° C. Adjust the pH to 8 with NH₄OH, stirring continuously.
4. Cool to room temperature in a cold water bath and centrifuge for 5 minutes. Discard the supernate, record the time and date of the decantation as the end of the yttrium-90 ingrowth and the beginning of its decay in the yttrium fraction.
5. Dissolve precipitate by adding about 4 drops of HCl with stirring. Add 15-20 ml of water. Heat in a water bath and adjust the pH to 8 with NH₄OH, stirring continuously.
6. Cool to room temperature in a cold water bath and centrifuge for 5 minutes. Discard supernate.
7. Repeat steps 5 and 6.
8. Add 3 drops of HCl to dissolve the precipitate, then add 20 ml of water. Filter the solution using No. 541 Whatman hardened filter paper. Heat in a water bath at approximately 90° C. Add 1 ml of saturated oxalic acid solution dropwise with vigorous stirring. Adjust to a pH of 2-3 with NH₄OH. Allow the precipitate to digest for about an hour.
9. Cool to room temperature in a cold water bath. Centrifuge for 10 minutes and decant most of the supernate. Filter by suction on a weighed filter paper. Wash the precipitate with water and absolute ethyl alcohol.
10. Dry the precipitate under the lamp for 30 minutes. Cool and weigh. Mount and count without delay in a proportional counter as Y-90 (Sr-90).
11. Calculate Sr-89 and Sr-90 activity using the computer program for Sr-89,-90.

Section 8.6 B (continued)Part B DeterminationII. Strontium-90Calculations

$$a. \text{ Strontium-90 concentration (pCi/g)} = \frac{A}{B \times C \times D \times E \times F}$$

Where:

- A = net beta count rate of yttrium-90 (cpm)
- B = recovery of strontium carrier
- C = efficiency for counting yttrium-90 as yttrium oxalate (cpm/pCi)
- D = sample size (in grams)
- E = correction factor $e^{-\lambda t}$ for yttrium-90 decay, where t is the time from decantation of the strontium supernate (Step B-II-4) to the time of counting (Step B-II-10)
- F = correction factor $1 - e^{-\lambda t}$ for the degree of equilibrium attained during the yttrium-90 ingrowth period, where t is the time from strontium separation (Step B-I-8) to the time of strontium removal (Step B-II-4).

$$b. \text{ Strontium-89 concentration (pCi/g)} = \frac{1}{B \times C} \left(\frac{A}{D \times E} - F(G \times H + I \times J) \right)$$

Where:

- A = net beta count rate of "total radiostrontium": (cpm)
- B = counter efficiency for counting strontium-89 as strontium oxalate mounted on a 2.1 cm diameter membrane filter (cpm/pCi)
- C = correction factor $e^{-\lambda t}$ for strontium-89 decay, where t is the time from sample collection to the time of counting
- D = recovery of strontium carrier
- E = sample size (in grams)
- F = strontium-90 concentration (pCi/g)
- G = self-absorption factor for strontium-90 as strontium oxalate mounted on a 2.1 cm diameter membrane filter
- H = counter efficiency for counting strontium-90 as strontium oxalate mounted on a 2.1 cm diameter membrane filter (cpm/pCi)
- I = counter efficiency for counting yttrium-90 as yttrium oxalate mounted on a 2.1 cm diameter membrane filter (cpm/pCi).
- J = correction factor $1 - e^{-\lambda t}$ for yttrium-90 ingrowth, where t is the time from the last decantation of the nitric acid supernate from the strontium nitrate precipitate to the time of counting (Step B-I-8).

References: Radioassay Procedures for Environmental Samples. U.S. Department of Health, Education and Welfare Environmental Health Series, January 1967. HASL Procedure Manual edited by John H. Harley, 1972.

APPENDIX IV

ERRATA

Table 5.0-4

Environmental Radiological Monitoring Program Quarterly Summary

Name of facility Quad Cities Nuclear Power Station Docket No. 50-254, 50-265
 Location of facility Rock Island, Illinois Reporting Period 3rd Quarter 1984
 (County, State)

Sample Type (Units)	Type and Number of Analyses	LLD	Indicator Locations Mean ^a Range	Location with Highest Quarterly Mean		Control Locations Mean ^a Range	Number of Non-routine Results
				Location	Mean Range		
Air Particulates (pCi/m ³)	Gross Beta 77	0.01	0.028 (76/77) (0.017-0.051)	Q-05, Saddle Club Dairy 1.8 mi @ 160°	0.032 (12/12) (0.022-0.051)	None	0
Airborne Iodine (pCi/m ³)	I-131 35	0.10	<LLD	-	-	None	0
Gamma Background (TLDs) (mR/Qtr.)	Gamma Dose 16	3.0	11.2 (6/6) (8.5-12.6)	Q-11, Port Byron 8.0 mi @ 170°	13.1 (1/1) -	10.6 (10/10) (8.3-13.1)	0
Milk (pCi/l)	I-131 26	0.5	<LLD	-	-	None	0
Cooling Water (pCi/l)	Gross Beta 42	1.0	4.2 (13/13) (1.9-5.3)	Q-22A, Diffuser Pipe Blowdown at Station and Q-21, Inlet Canal at station	4.2 (13/13)	4.2 (13/13) (1.6-4.8)	0
	Tritium 1	200	<LLD	-	-	None	0
Public Water (pCi/l)	Gamma Spec. 6						
	Cs-134 10	10.0	<LLD	-	-	None	0
	Cs-137 10	10.0	<LLD	-	-	None	0
	Other Gammas 20.0		<LLD	-	-	None	0
Bottom Sediments	Gamma Spec. 1						
	Cs-134	0.1	<LLD	-	-	None	0
	Cs-137	0.1	0.68 (1/1)	Q-23, Lock & Dam #14 Mississippi River, 15.0 mi @ 220°	0.68 (1/1)	None	0
	Other Gammas	0.2	<LLD	-	-	None	0

^a Mean and range based on detectable measurements only. Fractions indicated in parentheses.

TABLE 5.0-1 (continued)

QUAD CITIES STANDARD RADIOLOGICAL MONITORING PROGRAM

<u>Sample Media</u>	<u>Collection Site</u>	<u>Type of Analysis</u>	<u>Frequency</u>	<u>Non-Routine Reporting Levels</u> ^b
5. Public Water	(a) East Moline Water Works (b) Davenport Water Works	1. Gamma Isotopic	1. Monthly Analysis of Weekly Composites	(See footnote e)
6. Cooling Water ^f	(a) Inlet (b) Discharge	1. Gross Beta	1. Weekly	
7. Sediment	(a) Lock and Dam No. 14	Gamma Isotopic	Annually	
8. Dairy Census	(a) Site Boundary to 2 miles (b) 2 miles to 5 miles (c) At dairies listed in item 4.	(a) Enumeration by a door-to-door or equivalent counting technique (b) Enumeration by using referenced information from county agricultural agents or other reliable sources. (c) Inquire as to feeding practices. (1) pasture only (2) Feed and chop only (3) Pasture and feed; if both, ask farmer to estimate fraction of food from pasture <25% 25-50% 50-75% >75%	Annually, during grazing season	

^a Additional information giving the distance and direction of individual sampling locations may be found in Appendix III of the 1978 Annual Report.

^b Average concentration over calendar quarter.

^c A gamma isotopic analysis shall be performed whenever the gross beta concentration in a sample exceeds by five times (5x) the average concentration of the preceding calendar quarter for the sample location.

^d Bi-weekly shall mean that the frequency is once every other week.

^e H-3 2×10^4 , Mn-54 1×10^3 , Fe-59 4×10^2 , Co-58 1×10^3 , Co-60 3×10^2 , Zn-65 3×10^2 , Zr-Nb-95 4×10^2 , I-131 2, Cs-134 30, Cs-137 50, Ba-La-140 2×10^2 pCi/l.

^f Provided by station personnel.