

COMPREHENSIVE RADIOLOGICAL SURVEY

OFF-SITE PROPERTY N/N' SOUTH NIAGARA FALLS STORAGE SITE LEWISTON, NEW YORK

Prepared for

U.S. Department of Energy as part of the Formerly Utilized Sites — Remedial Action Program

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FINAL REPORT

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COMPREHENSIVE RADIOLOGICAL SURVEY

OFF-SITE PROPERTY N/N' SOUTH NIAGARA FALLS STORAGE SITE LEWISTON, NEW YORK

INTRODUCTION

Beginning in 1944, the Manhattan Engineer District and its successor, the Atomic Energy Commission (AEC), used portions of the Lake Octario Ordnance Works, presently referred to as the Niagara Falls Storage Site (NFSS) and off-site properties, approximately 3 km northeast of Lewiston, NY, for storage of radioactive wastes. These were primarily residues from uranium processing operations; however, they also included: contaminated rubble and scrap from decommissioning activities, biological and miscellaneous wastes from the University οf Rochester, and low-level fission-product waste from contaminated-liquid evaporators at the Knolls Atomic Power Laboratory (KAPL). Receipt of radioactive waste was discontinued in 1954, and, following cleanup activities by Hooker Chemical Co., 525 hectares of the original 612 hectare site This property was eventually sold by the General were declared surplus. Services Administration to various private, commercial, and governmental agencies. 1

The Department of Labor is the current owner of a tract identified as off-site property N/N' South (see Figure 1). A radiological survey of that tract is the subject of this report.

SITE DESCRIPTION

General

Figure 2 is a plot plan of off-site property N/N' South. The property occupies approximately 31 nectares and is bounded on two sides by roads: Castle Garden to the west, and South Patrol Road on the southeast. Town of Lewiston property borders the site on the south and Modern Landfill, Inc., owns the property to the north. Unused railroad tracks cross the property in a southwest to northeast direction. There are major drainage ditches along Track Street and South Track Street, on both sides of the southwestern portion of the railroad

tracks and crossing the northwest corner of the site. The site is heavily wooded with the exception of the Track Street area and the northwest corner of the site.

Radiological History

Two small areas of property N/N' South were previously used for handling or storing contaminated material and low-level radioactive wastes. An incincrator in the northwest corner of the property was used by Hooker Electrochemical Company to burn low level radioactive waste containers. Aerial photographs indicate the incinerator was present as late as 1963. The Track Street area was a classification yard, used for temporary storage of metal scrap, building rubble, and other miscellaneous material; some of the material stored in this area was contaminated.

Both of these areas were included in the 1971-72 cleanup efforts; however, following decontamination elevated radiation levels were still present at both locations.² Scrap and building debris remain in the ditches adjacent to Track Street and small pieces of debris were noted in the incinerator area. The 1980 mobile gamma scan by Oak Ridge National Laboratory indicated elevated radiation levels in both of these areas.³

SURVEY PROCEDURES

The comprehensive survey of NFSS off-site property N/N' South was performed by the Radiological Site Assessment Program of Oak Ridge Associated Universities (ORAU). The survey was in accordance with a plan dated September 27, 1982, approved by the Department of Energy's Office of Occupational Safety. With the exception of shallow borehole drilling, the survey was conducted during the period of October 18-November 3, 1982; the borehole investigations were performed March 24-25, 1983. The objectives and procedures from that plan are presented in this section.

Objectives

The objective of the survey was to provide a comprehensive assessment of the radiological conditions on property N/N^{γ} South and associated potential

health effects, if any. Radiological information collected included:

- direct radiation exposure rates and surface beta-gamma dose rates,
- 2. locations of elevated surface residues,
- 3. concentrations of radionuclides in surface and subsurface soil, and
- 4. concentrations of radionuclides in ground water.

Procedures

1. Site Preparation

- Brosh and weeds were cleared as needed to provide access for gridding and surveying. This operation was performed under subcontract by Modern Disposal Co., Model City, NY.
- b. An 80 m grid system was established by McIntosh and McIntosh of Lockport, NY, under subcontract. This grid system is shown on Figure 2. The 80 m grid was subdivided into 10 m intervals in the area where the incinerator had been located and in the area bounded by Track Street and South Track Street (see Figures 4 and 5).
- Gamma exposure rate measurements were performed at the surface and at l m above the surface at each accessible grid line intersection. Measurements were performed using portable gamma NaI (TI) scintillation survey meters. Conversion of these measurements to exposure rates in microroentgens per hour ($\mu R/h$) was in accordance with cross calibration with a pressurized ionization chamber.
- 3. Beta-gamma dose rate measurements were performed 1 cm above the surface at each accessible grid line intersection. These measurements were conducted using thin-window (7 mg/cm²) G-M detectors and portable scaler/ratemeters. Measurements were also obtained with the detector shielded to evaluate contributions of non-penetrating beta and low-energy gamma radiations. Meter readings were converted to dose-rate in microrads per hour (prad/h) based on cross calibration

with a thin-window ionization chamber using soil samples containing elevated concentrations of Ra-226.

- 4. Surface (0-15 cm) soil samples of approximately 1 kg each were collected at each accessible grid line intersection.
- 5. Walkover surface scans were conducted over all accessible areas of the property. Scanning intervals were at 5-10 m in the heavily wooded regions and at 1-2 m along roads, railroads, and in the old incinerator and Track Street areas. Portable gamma scintillation survey meters were used for these scans. Locations of elevated contact radiation levels were noted and surface exposure rates were measured at these locations.
- 6. At locations of elevated surface radiation levels, beta-gamma dose rates and exposure rates at 1 m above the surface were measured. Surface soil samples were obtained from these locations, and, following sampling, the surface exposure levels were remeasured to evaluate the effectiveness of shallow sampling on the removal of the radiation source. The locations where these additional measurements and samples were obtained are indicated on Figures 4, 5, and 6.
- 7. Detection Sciences Group of Carlisle, MA, performed ground-penetrating radar surveys in the old incinerator and Track Street areas, where elevated direct radiation levels were noted, to identify subsurface objects or anomalies which might be indicative of waste burials on the site. Ground radar scans were also conducted at locations of proposed boreholes to identify the presence of underground piping or utilities which would preclude drilling.
- 8. Boreholes were drilled to provide a mechanism for logging subsurface direct radiation profiles and collecting subsurface soil and water samples. Fourteen boreholes to ground water depth (2-6 m) were drilled by Site Engineers, Inc., of Voorhees, NJ, using a truck-mounted 20 cm diameter hollow-stem auger. These holes were drilled at systematic locations throughout the property as shown on Figure 7.

Twelve shallower (0.5-1.5 m) borcholes were drilled by Earth Dimensions of East Aurora, NY, using a special augering rig able to negotiate swampy and muddy areas. The shallow boreholes were primarily at locations where direct radiation measurements and ground-penetrating radar had indicated possible residues (old incinerator and Track Street areas). The locations of these borcholes are shown on Figures 8 and 9.

Gamma scans of boreholes were performed to identify elevated radiation levels, which would indicate subsurface residues. Radiation profiles in the boreholes were then determined by measurements of gamma radiation at 30-50 cm intervals between the surface and ground water (deep holes) or the hole bottom (shallow holes). A collimated gamma scintillation detector and portable scaler were used for these measurements.

Ground water samples of approximately 3.5 liters each were collected from deep boreholes. Collection was performed using a hand bailer. Soil samples of approximately 1 kg each were collected from various depths in the holes by scraping the sides of the borehole with a specially constructed sampling tool.

9. Twenty soil samples and seven water samples were collected from the Lewiston area (but not the NFSS or associated off-site properties) to provide baseline concentrations of radionuclides for comparison purposes. Direct background radiation levels were measured at locations where baseline soil samples were collected. The locations of the baseline samples and background measurements are shown on Figure 10.

Sample Analysis and Interpretation of Results

Soil and sediment samples were analyzed by gamma spectrometry. Radium-226 was the major radionuclide of concern, although spectra were reviewed for Cs-137, U-235, U-238, and other gamma emitters. Radiochemical analysis for Sr-90 was conducted on several samples with elevated levels of Cs-137.

Water samples were analyzed for gross alpha and beta concentrations. Additional information concerning analytical equipment and procedures is contained in Appendix A.

Results of this survey were compared to applicable guidelines for formerly utilized radioactive materials handling sites as presented in Appendix B.

RESULTS

Background Levels and Baseline Concentrations

background exposure rates and baseline radionuclide concentrations in soil, determined for 20 locations (Figure 10) in the vicinity of the NFSS, are presented in Table 1-A. Exposure rates ranged from 6.8 to 8.8 μ R/h (typical levels for this area of New York). Concentrations of radionuclides in soil were: Ra-226, <0.09* to 1.22 pCi/g (picocuries per gram); U-235, <0.14 to 0.46 pCi/g; U-238, <2.20 to 6.26 pCi/g; Th-232, <0.32 to 1.18 pCi/g; and Cs-137, <0.02 to 1.05 pCi/g. These concentrations are typical of the radionuclide levels normally encountered in surface soils.

Radioactivity levels in baseline water samples are presented in Table 1-B. The gross alpha and gross beta concentrations ranged from 0.55 to 1.87 pCi/l (picocuries per liter) and <0.63 to 14.3 pCi/l, respectively. These are typical of concentrations normally occurring in surface water.

Direct Radiation Levels

Direct radiation levels, systematically measured at 80 m grid line intersections, are presented in Table 2. Radiation levels measured at 10 m intervals in the old incinerator and Track Street areas are presented in Tables 3 and 4 respectively. The gamma exposure rates at 1 m above the surface ranged from 9.5 to 15 pR/h at the 80 m intervals; contact gamma exposure rates and beta-gamma dose rates at these locations ranged from 8.7 to 16 μ R/h and 42 to 140 prad/h, respectively. In the incinerator area these ranges were exposure rate at 1 m, 10 to 39 μ R/h; exposure rate at contact, 10 to 93 μ R/h; and beta-gamma dose tate, 42-320 μ rad/h. Direct radiation levels in the Track

Street area were exposure rate at 1 m, 9.8 to 18 μRh ; exposure rate at contact, 8.7 to 20 $\mu R/h$; and beta-gamma dose rate, 34 to 160 $\mu rad/h$.

Beta-gamma measurements performed with the G-M detector shielded averaged approximately 20% less than those with the unshielded detector. This indicates only a small portion of the surface dose rate is due to nonpenetrating beta or low-energy photon radiations.

The walkover survey identified several areas with elevated surface radiation levels in the incinerator and Track Street areas. These locations are indicated on Figures 4, 5, and 6. Direct radiation levels at these locations are presented in Table 5. Gamma exposure rates at contact and at 1 m above the surface at these locations ranged from 20 to 410 μ R/h and from 15 to 39 μ R/h, respectively. Contact beta-gamma dose rates ranged from 140 to 3770 μ rad/h. The maximum contact radiation levels were at grid location 668E,30S - in the Track Street area. Sampling reduced the radiation level significantly at only one location (109E,210S). At other locations, direct radiation levels were not reduced by sampling; at some points levels actually increased following removal of soil samples. This suggests that the contamination at most locations extends greater than 15 cm below the surface and/or is diffused rather than in discrete particles.

Radionuclides Concentrations in the Surface Soil

Tables 6, 7, and 8 list the concentrations of radionuclides measured in surface soil from 80 m grid line intersections and from 10 m grid intersections in the incinerator and Track Street areas. The systematic samples from 80 m grid line intersections (see Table 6) contained Ra-226 concentrations ranging from <0.13 to 2.60 pCi/g. With the exception of the sample from location 740E,0S, the Ra-226 levels were not significantly different from baseline levels. Other radionuclide concentrations in these samples were also in the ranges of baseline levels.

In the old incinerator area, systematic Ra-226 and U-238 levels ranged from <0.13 to 12.8 pCi/g and <0.64 to 20.2 pCi/g, respectively (see Table 7). The highest levels of both Ra-226 and U-238 were in the sample from grid point

170E,30S. Concentrations in systematic soil samples from the Track Street area were: Ra+226, 0.16 to 9.74 pCi/g and U-238, 0.69 to 9.29 pCi/g (see Table 8). The locations with the highest levels of Ra-226 and U-238 were 660E,20S and 700E,80S respectively. Concentrations of U-235 and Cs-137 in systematic samples from both the incinerator and Track Street areas were not significantly different from those in baseline samples.

Radionuclide concentrations in surface soil samples, collected from locations of elevated radiation levels are presented in Table 9. Samples 81 through 89 were collected from the old incinerator area. All samples obtained from that area contained Ra-226 and U-238 concentrations above those in baseline samples. The highest Ra-226 concentration (141 pCi/g) was in sample 84 from grid location 163E,31S. Sample 83, at grid point 153E,36S, contained the highest U-238 levels (68.5 pCi/g). Several samples from this area also contained slightly elevated levels of U-235 and Cs-137.

Sample 810, collected near Castle Garden Road, consisted of a metallic-like rock. Due to its high activity and irregular geometry only qualitative gamma spectroscopy was performed; the major radionuclide was found to be Ra-226. A total Ra-226 activity of 3.73 μ Ci was determined by comparison of direct radiation levels with those from a standard encapsulated radium source.

Samples B11 through B19 were collected from the Track Street area. All except samples B11 and B21 had elevated Ra-226 levels. The highest Ra-226 level (430 pCi/g) was in sample B17 from grid location 668E,30S. Radionuclide concentrations in samples B11 and B12 consisted mainly of U-238; however, there were also elevated levels of U-235 and Th-232 in these samples. Other samples collected in the Track Street area also had elevated levels of U-235, U-238, and Cs-137.

Samples B6, B7, and B14 were also analyzed for Sr-90; levels were low, ranging from C.16 to 0.85 pCi/g.

Cround-Penetrating Radar Findings

The subcontractor report summarizing the ground-penetrating radar survey results for property N/N' South is provided as Appendix C. (This report also includes the findings on property Q, since the two properties were surveyed simultaneously.) At locations where surface contamination was noted in the old incinerator area, radar signatures indicative of non-ionic liquids were present. These anomalies are most likely due to the presence of petroleum solvents or oils that do not readily disperse in the ground. There is no evidence of buried residues in this area. The Track Street area survey indicated no evidence of subsurface objects. No pipes or other subsurface interferences were detected at proposed borehole locations.

Borehole Gamma-logging Measurements

Gamma scintillation measurements performed in borcholes indicated that contamination is confined to the upper 25-50 cm of soil. As evidenced by soil sample analysis, the gamma count rates determined by the borehole measurements were reliable indicators of elevated subsurface radionuclide levels. However, the gamma-logging data were not useful in quantifying radionuclide concentrations in the subsurface soil, because of the varying ratios of Ra-226, U-235, U-238, and Cs-137 occurring in soils from this site.

Radionuclide Concentrations in Subsurface Soil

Table 10 presents the radionuclide concentrations measured in soil samples from boreholes. The locations of the fourteen boreholes (H1-H14), selected to provide a representative coverage of the property, had subsurface radionuclide concentrations either in the range of baseline samples or less than the minimum detectable activity (MDA).

Borcholes H15-H21 were at locations where the walkover scan had identified possible surface contamination. Only one subsurface sample collected from the old incinerator area (H19 at 0.3 meters) had an elevated level of Ra-226 (11.6 pCi/g). The sample collected at 0.1 meter from borehole H-21 (also in the

the old incinerator area) had U-235 and U-238 concentrations of 2.36 and 28.6 pCi/g respectively. Other subsurface samples from the incinerator area and all subsurface samples obtained from the Track Street area had radionuclide concentrations either in the range of baseline samples or less than the minimum detectable activity (MDA).

Radionuclide Concentrations in Water

Water collected from the systematic boreholes had gross alpha and gross beta concentrations within the EPA drinking water criteria of 15 pCi/l and 50 pCi/l, respectively (see Table 11). High concentrations of dissolved solids in several samples resulted in residues, which adversely affected the detection sensitivities of the gross alpha and gross beta procedure.

COMPARISON OF SURVEY RESULTS WITH GUIDELINES

The guidelines applicable to cleanup of the Niagara Falls Storge Site off-site properties are presented in Appendix B. All exposure rates at 1 m above the ground surface on property N/N' South are well below the 60 μ R/h criteria established by the NRC for open land areas.

There are two surface areas where the Ra-226 soil concentrations, averaged over 100 m², exceed the level of 5 pC1/g above background. One of these is near grid point 160E,30S in the old incinerator area. The average concentration, based on biased and systematic camples, is 37.5 pCi/g. The contaminated area, shown on Figure 11, covers approximately 200 m² and is limited to the top 25 cm of soil. The second area is along a ditch in the vicinity of Track Street and covers approximately 800 m² to a depth of 25 cm (see Figure 12). The average concentration of Ra-226 in this area is about 122 pCi/g. Although many samples from both these areas had elevated levels of U-238, the criteria level of 40 pCi/g was not exceeded as averaged over 100 m². Other locations of Ra-226 and U-238 surface soil contamination, indicated on Figures 11 and 12, are small and isolated. At these locations the concentrations, averaged over 100 m², would be below the criteria level.

Ground penetrating radar identified anomalies in the incinerator area but did not indicate the presence of subsurface objects which would suggest buried materials. Borehole measurements and sampling indicate all subsurface soil concentrations are within the applicable guidelines of 15 pCi/g for Ra-226.

Radionuclide concentrations in surface and subsurface water are within the EPA Interim Drinking Water Standards.

An evaluation of the potential health effects associated with radiation levels and residual contamination on property N/N' South is presented in Appendix D. That section compares the radiation levels with background exposures in the Niagara, New York, area and the scientifically based guidelines established for the protection of radiation workers and the general public.

SUMMARY

A comprehensive survey of off-site property N/N' South at the Niagara Falls Storage site (former Lake Ontario Ordnance Works) was conducted by Oak Ridge Associated Universities. The survey included surface radiation scans, measurements of direct radiation levels, and analyses of radionuclide concentrations in surface and subsurface soil samples and in subsurface water samples. Ground penetrating radar was also used to identify subsurface anomalies, which might suggest buried radioactive residues or objects.

The results of the survey indicate general and isolated areas of surface soil contamination in the vicinity of the old incinerator and in the Track Street area of the property. The major contaminant is Ka-226, although elevated levels of U-238 were also noted in many of the samples. Smaller amounts of Cs-137, U-235, and Th-232 were also detected in several samples. Subsurface sampling and measurements indicate that this contamination is limited to the top 25 cm of soil. Soil concentrations exceed the criteria in a 800 m² area in the Track Street area and a 200 m² area in the vicinity of the old incinerator. These two areas could be brought into compliance with the removal of approximately 200 m³ and 50 m³ of soil, respectively. Other isolated areas of contamination could be eliminated by removal of much smaller amounts of surface soil.

Although there were small areas of contaminated residues on portions of this property, under present conditions of property use, these contaminants do not pose potential health risks. There is no evidence that migration of the radioactive materials is adversely affecting adjacent properties or the ground water.

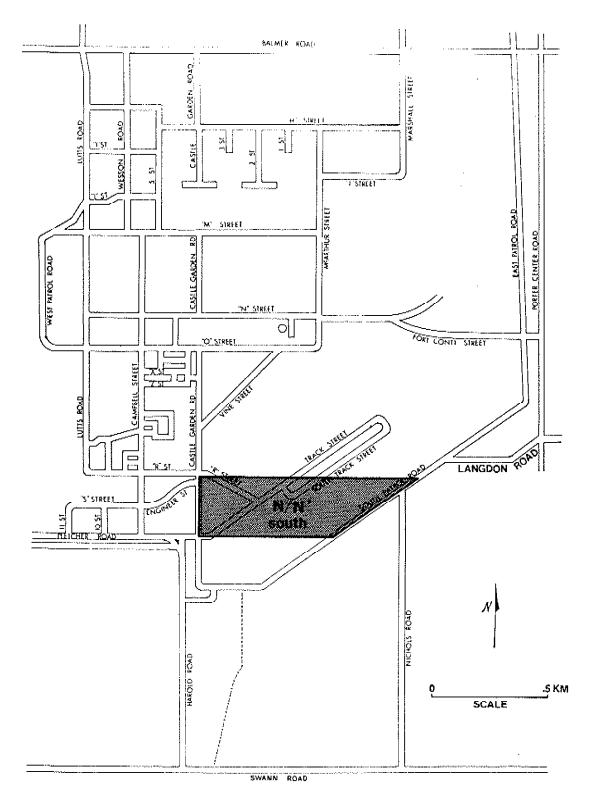
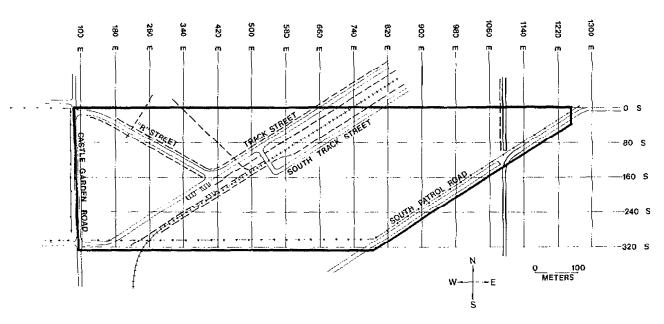


FIGURE 1. Map of Niagara Falls Storage Site and Off-Site Properties, Lewiston, New York, Indicating the Location of Property N/N' South.



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FIGURE 2. Plan View of Off-Site Property N/N' South, Indicating Prominent Surface Features and the Grid System Established for Survey Reference.

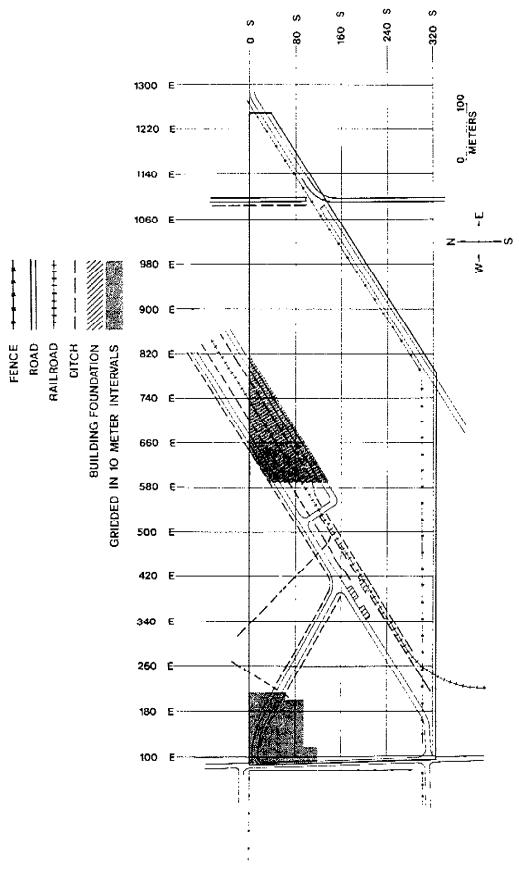


FIGURE 3. Map of Property N/N' South Indicating Areas Gridded in 10 Meter Intervals.

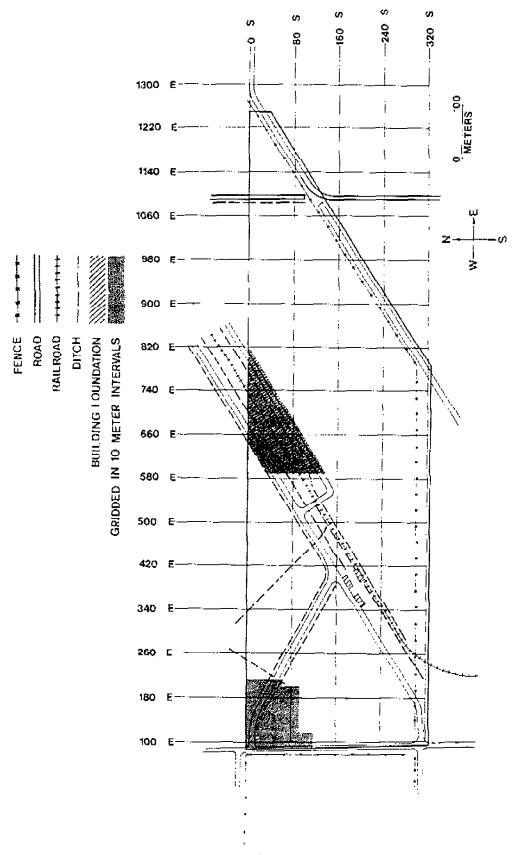


FIGURE 3. Map of Property N/N' South Indicating Areas Gridded in $10\ \text{Meter}$ Intervals.

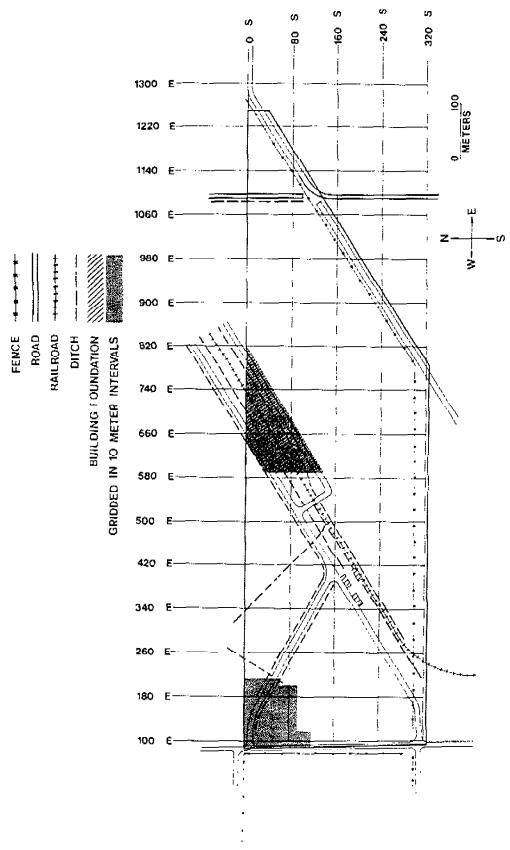


FIGURE 3. Map of Property N/N' South Indicating Areas Gridded in $10~{\rm Meter}$ Intervals.

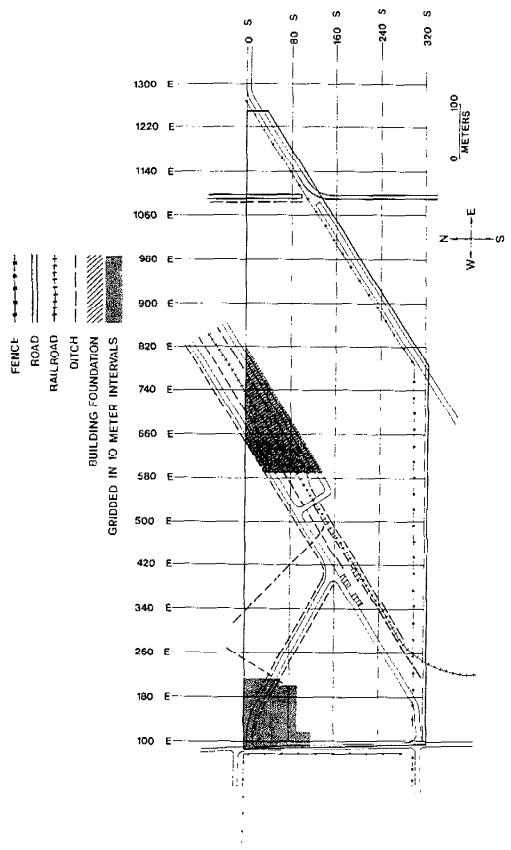


FIGURE 3. Map of Property N/N^{\star} South Indicating Areas Gridded in 10 Meter Intervals.

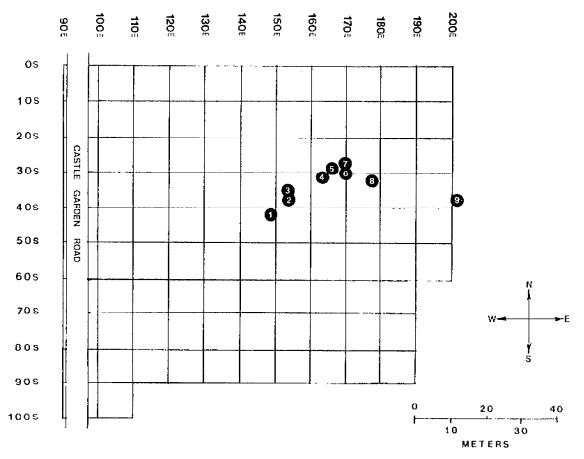
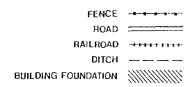


FIGURE 4. Locations of Elevated Direct Radiation Levels in the Old Incinerator Area.



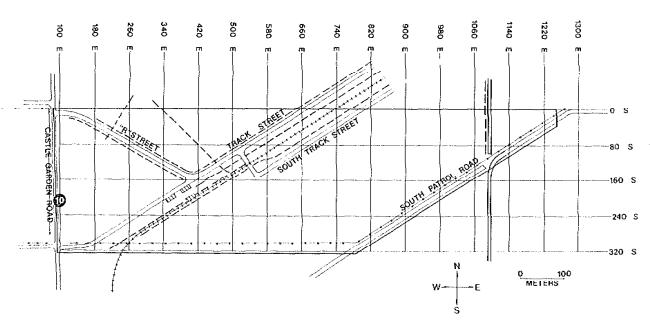


FIGURE 5. Location of an Elevated Direct Radiation Level Along Castle $\ensuremath{\mathsf{Garden}}$ Road.

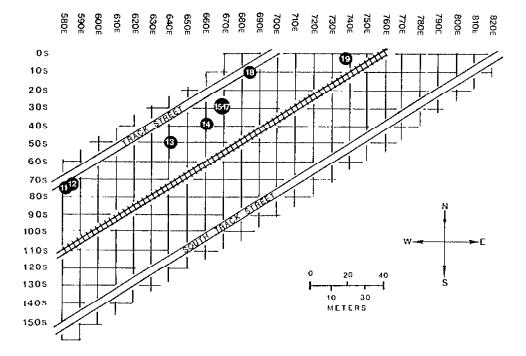


FIGURE 6. Locations of Elevated Direct Radiation Levels in the Track Street Area.

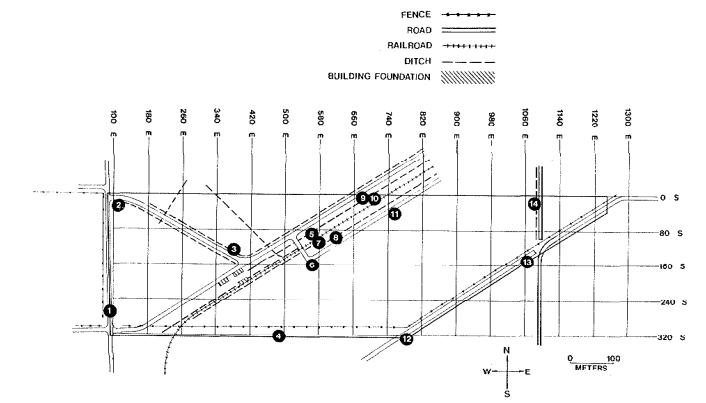


FIGURE 7. Locations of Deep Boreholes for Subsurface Investigations.

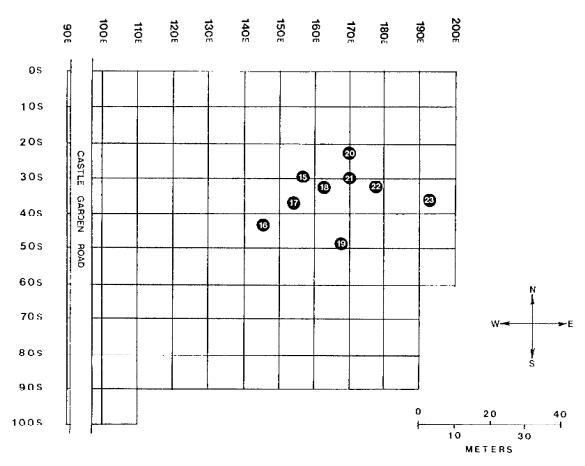


FIGURE 8. Locations of Shallow Boreholes in the Incinerator $$\operatorname{Areas}$.$

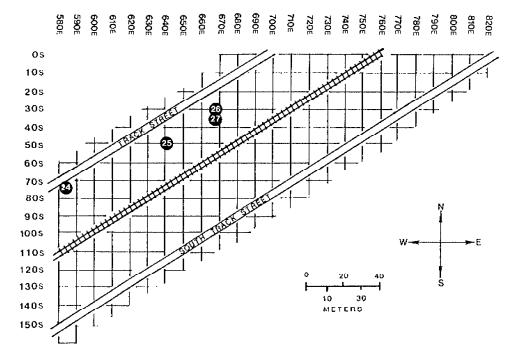


FIGURE 9. Locations of Shallow Boreholes in the Track Street Area.

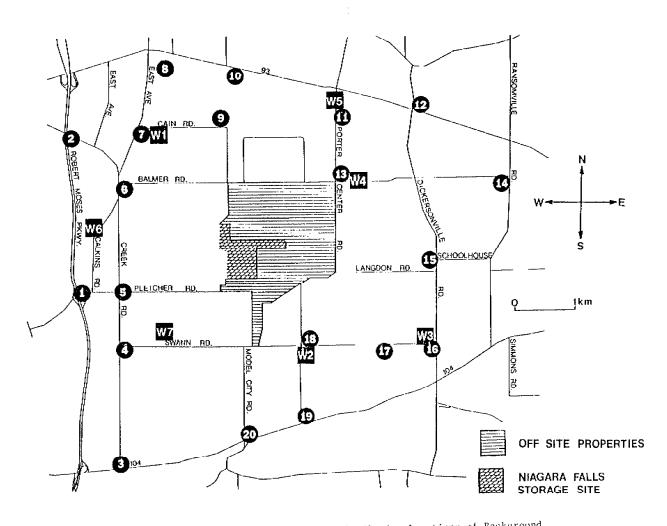


FIGURE 10. Map of Northern Niagara County, New York, Showing Locations of Background Measurements and Baseline Samples (#1-20: soil samples and direct measurements; W1-W7: water samples).

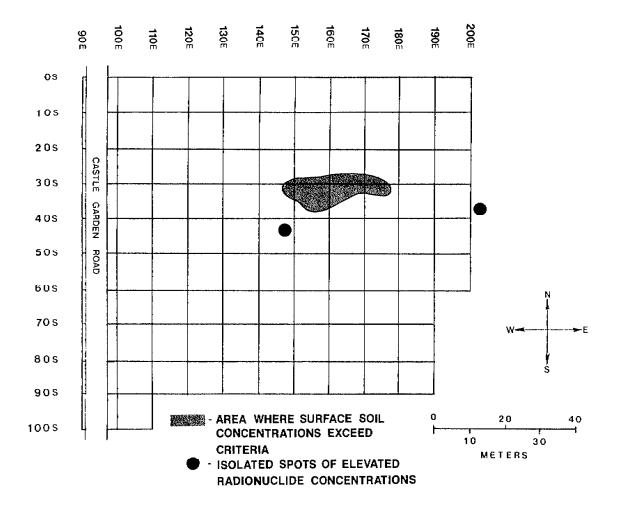


FIGURE 11. Map of the Incinerator Area, Indicating Locations Where Ra-226 Concentrations Exceed Criteria for Surface Soil.

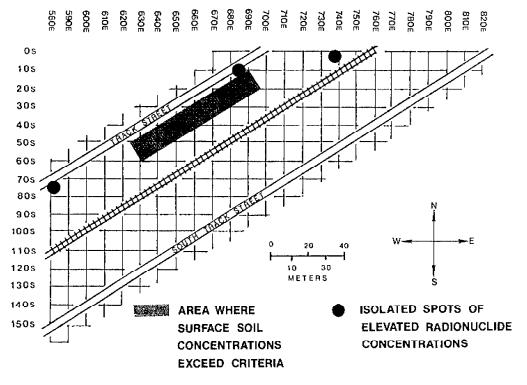


FIGURE 12. Map of the Track Street Area, Indicating Locations Where Ra-226 Concentrations Exceed Criteria for Surface Soil.

TABLE 1-A BACKGROUND EXPOSURE RATES AND
BASELINE RADIONUCLIDE CONCENTRATIONS IN SOIL

	Exposure Rateb		Radiomuclide Concentrations (pCi/g)						
Locationa	(yR/h)	Ra-226	U÷235	U-238	Th=232	Cs-137			
1	6.8	0.74 ± 0.16°	<0.19	<2.89	0.70 ± 0.46	0.29 ± 0.08			
2	6.8	0.75 <u>+</u> 0.19	<0.19	<3.35	<0.22	0.24 ± 0.08			
3	8.3	0.71 ± 0.18	0.46 <u>*</u> 0.41	<3.72	0.88 <u>+</u> 0.33	0.34 ± 0.09			
4	7.9	0.67 ± 0.18	<0.27	<4.10	1.18 ± 0.35	0.12 ± 0.07			
5	7.3	0.70 ± 0.16	<0.17	<3.34	0.68 ± 0.24	0.14 ± 0.07			
6	7.7	0.50 + 0.15	<0.16	<2.33	0.52 <u>+</u> 0.38	0.17 ± 0.09			
7	7.7	0.63 <u>*</u> 0.13	<0.17	<2.73	0.83 ± 0.24	0.35 👲 0.08			
8	7.6	0.59 <u>+</u> 0.12	<0.14	<2.20	0.54 + 0.23	<0.02			
9	7.1	0.63 • 0.20	<0.23	<4.16	0.83 👲 0.38	0.69 ± 0.11			
10	7,1	0.70 ± 0.16	<0.19	<2.98	<0.18	0.69 <u>+</u> 0.10			
11	6.7	<0.09	<0.19	<2.83	0.49 ± 0.31	0.48 ± 0.14			
12	7.1	0.48 <u>+</u> 0.13	<0.16	<2.84	0.65 + 0.26	0.68 ± 0.10			
13	6.7	0.57 ± 0.14	<0.17	<2.36	0.49 ± 0.26	0.41 + 0.08			
14	6.8	0.68 <u>+</u> 0.17	<0.19	<3.24	0.67 ± 0.25	0.70 ± 0.10			
15	8.2	0.65 ± 0.14	<0.17	<3.20	0.72 <u>+</u> 0.35	0.23 ± 0.08			
16	7.4	0.91 <u>*</u> U.17	<0.71	<3.58	0.83 🛨 0.28	0.61 ± 0.09			
17	7.0	0.48 + 0.14	<0.16	<2.73	0.32 ± 0.22	0.38 ± 0.08			
16	7.7	0.73 ± 0.16	<0.18	6.26 + 9.23	<0.23	0.32 ± 0.12			
19	8.8	1.22 + 0.22	<0.23	<3.79	1.08 ± 0.49	1.05 ± 0.13			
20	8.6	0.83 ± 0.17	<0.21	<3.59	0.84 <u>+</u> 0.29	0.08 ± 0.07			
ange	6,8 to 8,8	<0.09 to 1.22	<0.14 to 0.46	<2,20 to 6,76	<0.18 to 1.18	<0.02 to 1.05			

 $[^]a$ Refer to Figure 10. b Measured at 1 m above the surface. c Errors is 2_σ based on counting statistics only.

TABLE 1-B RADIONUCLIDE CONCENTRATIONS IN BASELINE WATER SAMPLES

ocationa	Radionuclide Conce	
ocation-	Gross Alpha	Gross Beta
Wl	0.95 ± 0.93 b	4.79 ± 1.15
W2	0.95 ± 0.94	9.17 ± 1.31
W3	0.55 ± 0.78	2.73 ± 1.05
W4	0.63 ± 0.89	5.37 ± 1.17
W5	0.73 ± 0.68	<0.64
W6	1.87 ± 1.84	14.3 <u>+</u> 2.4
W7	1.16 ± 0.66	<0.63
Range	0.55 to 1.87	<0.63 to 14.3

 $^{^{\}rm a}$ Refer to Figure 10. $^{\rm b}$ Errors are 2σ based on counting statistics.

TABLE 2

DIRECT RADIATION LEVELS SYSTEMATICALLY
MEASURED AT 80 M GRID LINE INTERSECTIONS

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface	Beta-Gamma Dose Rates at the Surface
Location	(υR/h)	(\pm R/h)	(hrad/h)
100E, OS	11	12	62
100E, 80S	11	11	68
100E,160s	11	11	69
100E,240S	11	11	75
100E,320s	9.8	8.7	48
180E, OS	11	11	62
180E, 80s	îî	11	83
180E,160S	11	12	94
180E,240S	11	12	68
180E,320S	11	11	68
260E, OS	11	11	83
260E, 80S	11	01	88
260E,160S	12	12	78
260E,240S	13	15	110
260E,320S	12	12	86
•	11	12	110
340E, OS 340E, 8OS	12	12	81
340E,160S	11	11	55
340E,240S	11	11	62
	13	11 12	65
340E,320S		12	81
420E, OS	11		
420E, 80S	11	11	78
420E,160s	11	12	55
420E,240S	11	11	55
420E,320S	11	12	68
500E, 0s	12	11	81
500E, 80S	12	12	91
500E,160S	13	13	78
500E,240s	12	11	73
500E,320S	11	12	57
580E, OS	11	12	78
580E, 80S	15	16	140
580E,160S	11	12	44
580E,240S	11	11	96
580E,320S	11	11	73
660E, OS	12	11	88
660E, 80S	14	13	68
660E,160S	10	11	49
660E,240S	11	11	68
660E,320S	11	11	68
740E, 0S	15	16	81
740E, 80S	11	11	42

TABLE 2, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 80 M GRID LINE INTERSECTIONS

Grid Location	Gamma Exposure Rates at 1 m Above the Surface (µR/h)	Gamma Exposure Rates at the Surtace $(\mu R/h)$	Beta-Gamma Dose Rates at the Surface (prad/h)
740E,160S	11	11	83
740E,240S	10	11	73
740E,320S	10	11	91
820E, OS	11	10	47
820E, 80S	11	11	81
820E,160S	11	11	52
820E,240S	11	11	78
820E,320S	11	11	78
900E, OS	12	12	7 8
900E, 80S	11	12	60
900E,160S	9.5	9.5	65
900E,240S	11	11	55
900E,320S	10	11	55
980E, OS	11	11	73
980E, 80S	11	11	85
980E,160S	12	11	62
980E,240S	11	11	79
1060E, OS	11	11	79
1060E, 80S	11	11	83
1060E,160S	11	11	59
1140E, OS	11	11	73
1140E, 80S	11	11	70
1220E, 0S	11	11	66

TABLE 3

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE AREA OF THE OLD INCINERATOR

Grid Location	Gamma Exposure Rates at 1 m Above the Surface (uR/h)	Gamma Exposure Rates at the Surface $\binom{1}{1}R/h$	Beta-Gamma Dose Rates at the Surface (urad/h)
		, j,	
1000 00		1.0	62
100E, OS	11 11	12 11	52 52
100E, 10S	13	15	86
100E, 20S		12	86
100E, 30S	11	11	61
100E, 40S	11		98
100E, 50S	11	11	68
100E, 60S	11	10	
100E, 70S	11	11	61
100E, 80S	11	11	68
100E, 90S	11	11	55 20
100E,100S	11	11	70
110E, OS	15	16	110
110E, 10S	15	17	100
110E, 20S	11	10	72
110E, 30S	10	10	64
110E, 40S	11	11	53
110E, 50S	11	11	62
110E, 60S	11	11	70
110E, 80S	11	11	7.5
110E, 90S	12	11	62
110E,100S	11	12	69
120E, OS	12	13	82
120E, 10S	12	12	68
120E, 20S	11	10	5 3
120E, 30S	11	11	62
120E, 40S	11	11	65
120E, 50S	11	12	81
120E, 505	11	12	64
120E, 70S	12	12	91
120E, 70S	12	12	81
	12	1.2	87
120E, 90S	11	11	78
130E, OS		11	7
130E, 10S	10	11	7 2 59
130E, 20S	10		78
130E, 30S	10	11	70 62
130E, 40S	11	11	
130E, 50S	12	12	64
130E, 60S	11	11	75

TABLE 3, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE AREA OF THE OLD INCINERATOR

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface	Beta-Gamma Dose Rates at the Surface (⊒rad/h)	
Location	(µ R/h)	(u R/h)		
130E, 70S	1.2	12	86	
130E, 80S	1 2	I 2	66	
130E, 90S	12	12	81	
140E, OS	12	12	75	
140E, 10S	11	11	65	
140E, 20S	10	11	72	
140E, 30S	11	11	61	
140E, 40S	12	12	83	
140E, 50S	12	12	62	
140E, 60S	12	11	68	
140E, 70S	12	12	78	
140E, 80S	11	11	66	
140E, 90S	12	12	79	
150E, OS	12	11	60	
150E, 10s	11	11	90	
150E, 20S	11	11	66	
150E, 30S	12	12	83	
150E, 40S	15	15	120	
150E, 50S	11	11	6.2	
150E, 60S	11	11	72	
150E, 70S	11	12	95	
150E, 80s	11	12	53	
150E, 90S	11	12	66	
160E, OS	11	11	73	
160E, 10S	11	11	42	
160E, 20S	11	12	74	
160E, 30S	13	13	120	
160E, 40S	12	12	74	
160E, 50S	11	12	73	
160E, 60S	12	11	65	
160E, 70S	$\overline{11}$	12	62	
160E, 80S	11	11	85	
160E, 90S	11	II	53	
170E, OS	11	11	62	
170E, 10S	11	11	48	
170E, 20S	12	îî	53	
170E, 30S	39	93	3 20	
170E, 40S	12	12	74	

TABLE 3, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE AREA OF THE OLD INCINERATOR

Grid Jocation	Gamma Exposure Rates at 1 m Above the Surface (µR/h)	Gamma Exposure Rates at the Surface ():R/h)	Beta-Gamma Dose Rates at the Surface (µrad/h)
COCACTON	(2 K/ II)	(): K/ II/	(μ140/π/
170E, 50S	12	11	88
170E, 60S	11	11	75
170E, 70s	11	$\frac{1}{1}$	88
170E, 80S	11	11	90
170E, 90S	11	11	73
180E, OS	11	11	62
180E, 10S	11	11	62
180E, 20S	12	12	42
180E, 30S	13	12	64
180E, 40S	12	12	83
180E, 50S	11	11	82
180E, 60S	11	11	69
180E, 70S	12	11	77
180E, 80S	11	11	83
180E, 90S	11	11	83
190E, OS	11	12	70
190E, 10S	11	12	72
190E, 20S	11	11	81
190E, 30S	11	11	$\overline{62}$
190E, 40S	12	12	99
190E, 50S	12	12	85
190E, 60S	12	11	87
190E, 70S	11	12	77
190E, 703	11	12	7,7 75
200E, 80S	11	12	62
200E, 10S	11	12	73
200E, 10S 200E, 20S	11	11	7 2
200E, 20S	11	11	68
200E, 40S	11	12	79
200E, 40S 200E, 50S	11	11	64
200E, 50S 200E, 60S	11	12	73

TABLE 4

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED
AT 10 M GRID LINE INTERSECTIONS IN THE TRACK STREET AREA

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface (\(\mu R/\h))	Beta-Gamma Dose Rates at the Surface (urad/h)
Location	(μ R/h)	(48/11)	(μιασ//
(000		10	81
680E, OS	12	12 10	61 57
690E, OS	11		78
700E, OS	12	13	60
710E, OS	13	12	78
720E, OS	14	15	
730E, OS	15	13	68
740E, OS	15	16	81
750E, OS	17	18	65
760E, OS	15	16	86
770E, OS	15	16	100
780E, OS	13	14	88
790E, OS	15	15	110
800E, OS	13	15	99
810E, OS	11	11	47
820E, OS	11	11	47
660E, 10S	12	12	78
670E, IOS	12	12	73
680E, 10S	14	15	120
690E, 10S	13	13	42
/00E, 10S	13	12	47
710E, 10S	14	15	65
720E, 10S	15	15	78
730E, 10S	17	15	81
740E, 10S	15	15	160
750E, 10S	15	15	100
760E, 10S	15	15	99
770E, 10S	13	15	91
780E, 10S	13	14	110
790£, 10S	13	13	78
800E, 10S	11	11	57
810E, 10S	10	10	49
820E, 10S	12	$\tilde{1}\tilde{2}$	60
650E, 20S	13	13	60
660E, 20S	13	13	81
670E, 20S	13	12	65
680E, 20S	13	15	83
690E, 20S	15	15	78
	14	15	88
700E, 20S 710E, 20S	14	15	65

TABLE 4, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE TRACK STREET AREA

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface	Beta-Gamma Dose Rates at the Surface
Location	(hR/h)	(uR/h)	(µrad/h)
720E, 20S	15	15	78
730E, 20S	16	18	94
740E, 20S	16	18	110
750E, 20S	15	15	91
760E, 20S	16	18	120
770E, 208	13	12	49
780E, 20S	11	12	73
790E, 20S	10	10	62
800E, 20S	11	12	47
640E, 30S	11	10	55
650E, 30S	12	12	60
660E, 30S	13	13	100
670E, 30S	18	20	86
680E, 30S	14	15	81
690E, 30S	14	14	70
700E, 30S	14	15	56
710E, 30S	15	16	91
720E, 30S	15	15	78
730E, 30S	1.5	15	110
740E, 30S	15	15	88
750E, 30S	15	15	140
760E, 30S	13	15	86
770E, 30S	11	12	62
780E, 30S	11	11	65
790E, 30S	12	12	83
620E, 40S	12	12	83
630E, 40S	11	12	44
640E, 40S	12	12	65
650E, 40S	11	11	81
660E, 40S	15	16	94
670E, 40S	12	13	52
680E, 40S	14	14	100
690E, 40S	13	13	83
700E, 40S	15	15	81
710E, 40S	15	15	83
720E, 40S	15	15	60
730E, 40S	15	15	94
740E, 40S	12	12	73
750E, 40S	12	12	73

TABLE 4, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE TRACK STREET AREA

Grid Location	Gamma Exposure Rates at 1 m Above the Surface ($\mu R/h$)	Gamma Exposure Rates at the Surface ($_{ m LR}/h$)	Beta-Gamma Dose Rates at the Surface (urad/h)
7605 400	10	0.0	
760E, 40S	10	9.8	60
770E, 40S	11	12	68
600E, 50S	12	12	62
610E, 50S	11	11	5 7
620E, 50S	12	12	73
630E, 50S	13	13	78
640E, 50S	16	16	120
650E, 50S	13	15	130
660E, 50S	. 14	14	96
670E, 50S	15	16	83
680E, 50S	15	18	150
690E, 50S	15	17	130
700E, 50S	15	17	83
710E, 50S	15	16	100
720E, 50S	12	12	70
730E, 50S	12	12	91
740E, 50S	10	9.8	68
750E, 50S	<u>:</u> 2	11	62
580E, 60S	12	12	75
590E, 60S	11	11	68
600E, 60S	13	13	7 8
610E, 60S	12	12	110
620E, 60S	15	15	83
630E, 60S	14	15	110
640E, 60S	14	15	100
650E, 60S	14	14	100
660E, 60S	15	16	120
670E, 60S	15	15	88
680E, 60S	17	18	150
690E, 60S	15	15	94
700E, 60S	15	17	110
	12	12	62
710E, 60S 720E, 60S	11	12	55
730E, 60S	11	11	83
740E, 60S	12	12	63 52
580E, 70S	12	12	68
590E, 70S 600E, 70S	13 12	12 12	83 62
OLD 18 / 11%	1.7	1.7	K 7

TABLE 4, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED AT 10 M GRID LINE INTERSECTIONS IN THE TRACK STREET AREA

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface	Beta-Gamma dose Rates at the Surface
Location	(kR/h)	(μR/h)	(urad/h)
620E, 70S	13	13	88
630E, 70S	14	13	110
640E, 70S		16	99
650E, 70S	15	16	110
660E, 70S	16	19	110
670E, 70S	17	18	100
680E, 70S	15	16	65
690E, 70S	12	13	65
700E, 70S	12	12	55
710E, 70S	10	9.8	52
580E, 80S	15	15	110
590E, 80S	14	15	73
600E, 80S	12	13	94
610E, 80S	13	13	65
620E, BOS	14	14	110
630E, 80S	15	15	160
640E, 80S	15	16	100
650E, 80S	15	15	110
660E, 80S	14	13	68
670E, 80S	13	12	49
680E, 80S	12	11	81
690E, 80S	10	8.7	52
580E, 90S	12	12	52
590E, 90S	12	12	110
600E, 90S	13	15	7 5
610E, 90S	15	15	100
620E, 90S	15	15	130
630E, 90S	15	16	96
640E, 90S	15	16	110
650E, 90S	13	13	83
660E, 90S	12	11	5 2
670E, 90S	12	12	86
680E, 90S	10	10	55
580E,100S	12	13	100
590E,100S	15	16	94
600E,100S	14	15	99
610E,100S	15	15	99
620E,100S	14	15	94
630E,100S	16	19	140

TABLE 4, cont.

DIRECT RADIATION LEVELS SYSTEMATICALLY MEASURED
AT 10 M GRID LINE INTERSECTIONS IN THE TRACK STREET AREA

Grid	Gamma Exposure Rates at 1 m Above the Surface	Gamma Exposure Rates at the Surface	Beta-Gamma Dose Rates at the Surface (prad/h)	
Location	(nR/h)	(u R/h)		
640E,100S	12	12	81	
650E,100S	12	12	78	
660E,100S	10	9.8	55	
670E,100S	12	12	78	
580E,110S	15	17	140	
590E,110S	15	18	110	
600E,110S	13	15	96	
610E,110S	12	12	75	
620E,110S	12	12	78	
630E,110S	11	11	55	
640E,110S	9.8	10	34	
650E,110S	11	11	52	
660E,110S	11	12	47	
580E,120S	12	12	91	
590E,120S	12	12	81	
600E,120S	11	11	62	
610E,120S	12	12	91	
620E,120S	13	13	55	
630E,120S	9.8	9.8	44	
580E,130S	12	12	65	
590E,130S	11	11	73	
600E,130S	12	12	60	
610E,130S	9.8	8.4	78	
620E,130S	11	11	88	
580E,140S	12	11	70	
590E,140S	11	11	52	
600E,140S	11	11	68	
580E,150S	9.8	8.7	52	
590E,150S	12	11	62	

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TABLE 5 DIRECT RADIATION LEVELS AT LOCATIONS IDENTIFIED BY THE WALKOVER SURFACE SCAN

		Expo	sure Rate (:R/h)	Sulface Dose Rate (grad/h)	Soil Sampleb	Contact Exposure Rate After Sample
location ^a	cation ^a Grid Location — C	Contact	1 to Above Surface	(trad/h)	SGII SEBPIE	Retoval (µR/h)
1	1481, 435	26	19	150	Dl	28
2	154E. 38S	35	22	200	B2	32
3	153E, 36S	51	24	250	В3	71
4	163E, 31S	7.5	24	160	B4	98
5	166E, 295	59	32	220	R5	43
6	170E, 30S	93	39	320	ь6	130
7	169E, 32S	26	21	140	ь7	26
6	1782, 328	71	32	3 50	в8	91
9	2077, 378	24	16	240	39	32
10	109E,210S	220	15	1230	310	13
11	552E, 76S	28	16	180	B11	32
12	584E, 74S	59	18	140	B1 2	63
13	641E, 50S	200	24	1470	B13	200
14	660E, 39S	22	16	160	B14	28
15	668E, 33S	97	26	530	B15	160
16	669E, 31S	51	20	530	B16	49
17	668E, 30S	410	38	3770	B17	550
18	684E, 10S	21	18	160	B18	21
19	738E, 45	20	18	160	519	20

a Refer to Figures 4, 5, and 6.
 b Soil concentrations presented in Table 9.

TABLE 6

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 80 M GRID LINE INTERSECTIONS

_			oncentrations (pCi/g)	
Location	Ra-226	U-235	v~23 B	CL-137
100E, 03	0.57 + 0.198	<0.24	<2.94	0.18 + 0.09
100E, 80S	0.71 + 0.18	<0.24	×4.09	0.08 - 0.09
100E,160s	1.36 ± 0.29	<0.24	≺3.94	0.66 ± 0.1
100E,240S	0.48 ± 0.21	<0.25	<4.33	0.47 ± 0.1
100E,320S	0.72 ± 0.28	<0.29	<5.15	0.29 ± 0.10
180E, QS	0.60 ± 0.23	<0.24	<3.27	0.47 ± 0.1
180E, 80S	0.42 🛨 0.23	<0.24	<5.67	0.59 <u>¥</u> 0.13
180E,160S	0.56 <u>+</u> 0.22	<0.24	<4.08	0.70 ± 0.13
180E,240S	0.53 <u>+</u> 0.24	<0.24	<3.99	0.34 ± 0.11
180E,320S	I.43 <u>+</u> 0.26	<0.24	<4.39	0.32 ± 0.13
260E, OS	0.66 + 0.21	1.23 ± 0.56	<4.06	0.40 + 0.1
260E, 80S	0.60 <u>+</u> 0.20	<0.24	<3.21	0.53 ± 0.16
260E,160S	0.67 ± 0.25	<0.24	<2.30	0.41 ± 0.1
260E,240S	0.48 ± 0.23	<0.24	<3.32	0.43 ± 0.13
2600,3203	0.55 <u>+</u> 0.25	<0.23	<4.53	0.49 <u>+</u> 0.1
340E. OS	0.78 👱 0.21	<0.24	<4.68	<0.04
340€, 803	0.52 <u>+</u> 0.23	<0.24	<4.16	<0.04
340E,160S	0.53 ± 0.20	0.52 <u>+</u> 0.46	<4.92	0.32 <u>+</u> 0.1;
340E,240S	0.53 <u>+</u> 0.19	<0.24	<3.76	0.26 ± 0.10
340E,320S	0.38 ± 0.30	<0.23	<3.79	0.26 ± 0.16
420E, OS	0.63 🛨 0.27	<0.23	<4.15	0.45 ± 0.13
420E, 80S	0.65 <u>+</u> 0.23	<0.24	<4.41	0.61 <u>+</u> 0.0
420E,160S	1.II <u>+</u> 0.29	<0.24	<4.47	0.54 <u>+</u> 0.1
420E,240S	0.36 ± 0.20	<0.24	<4.63	0.49 <u>+</u> 0.1
420E,320S	0.49 ± 0.20	<0.24	<4.81	0.79 + 0.1
500E, 0S	0.55 ± 0.26	<0.24	<3.81	<0.04
500E, 80S	0.43 ± 0.23	<0.24	<3.95	0.55 <u>+</u> 0.1
500E,160S	0.61 ± 0.26	<0.24	<4.68	0.30 <u>+</u> 0.1
500E,240S	0.56 ± 0.23	<0.25	<3.88	0.23 ± 0.1
500E,320S	0.47 ± 0.24	<0.24	8.54 ± 11.07	0.51 1 0.10
580E, OS	0.30 ± 0.20	<0.24	<3.99	<0.05
580E, 803	0.56 ± 0.20	<0.24	4.88 <u>+</u> 10.75	0.16 ± 0.0
580E,1GOS	0.64 ± 0.23	<0.24	<5.60	0.16 <u>+</u> 0.1.
580E,240S	0.70 ± 0.23	<0.25	<3.23	0.34 <u>+</u> 0.1:
580E,320S	1.28 ± 0.31	<0.25	<3.69	0.49 ± 0.1
660E, OS	0.43 ± 0.19	<0.24	<3.52	0.26 ± 0.11
660E, 80S	<0.16	<0.24	<3.76	0.55 <u>+</u> 0.1
660E,160S	0.47 ± 0.22	<0.21	<3.63	0.07 <u>+</u> 0.06
660E,240S	0.63 ± 0.40	<0.24	<2,04	0.32 ± 0.13
660E,320S	0.57 <u>+</u> 0.21	<0.22	<3.41	0.51 <u>+</u> 0.1
740E, OS	2.60 ± 0.22	<0.24	<5.60	0.82 <u>+</u> 0.1
740E, 80S	0.51 ± 0.20	<0.24	<4.13	0.25 <u>+</u> 0.1
740E,160S	<0.13	0.26 <u>+</u> 0.54	<3.45	0.63 ± 0.19
7408,2408	0.56 ± 0.23	<0.24	<1.94	0.38 ± 0.13
740E,320S	0.49 ± 0.21	0.65 <u>+</u> 0.56	<4.38	0.41 ± 0.13
829E, OS	0.77 <u>+</u> 0.21	<0.24	7.73 + 7.40	0.71 + 0.23

TABLE 6, cont.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES
FROM 80 M GRID LINE INTERSECTIONS

		Radionuclide Co	necentrations (pCi/	g)
Location	Ra-226	U-235	U-238	Cs-137
820E, 80S	0.64 + 0.28	<0.24	<3.70	0.33 + 0.11
820E,1605	0.49 🛨 0.21	<0.24	<3.80	0.42 🗓 0.15
820E,240S	0.45 <u>+</u> 0.19	<0.24	<5.14	0.37 ± 0.14
820E,320S	0.52 + 0.21	<0.24	<1.41	0.54 🛨 0.13
900E, OS	0.47 + 0.17	<0.24	<4.24	0.80 + 0.08
900E, 80S	0.56 + 0.33	<0.26	<5.31	0.59 + 0.18
900E,1603	C.55 ± 0.22	<0.24	<4.75	0.69 ± 0.14
900E,240S	0.56 ± 0.21	<0.24	<4.20	0.52 ± 0.12
9COE.323S	C.74 + 0.20	73.74	<4.91	0.67 ± C.12
980E, 0S	C.53 + O.19	<0.24	<4.85	0.50 ± C.13
950E, 805	C,42 - 0.19	<0.24	<4.51	0.25 🗓 0.11
980E,160S	0.50 ± 0.27	<0.24	<3.95	9.76 <u>+</u> 0.16
980E,240S	0.50 + 0.23	<0.24	<5.29	0.59 ± 0.15
1060E, OS	0.69 ± 0.20	<0.24	<4.18	0.34 ± 0.14
1060E, 80S	0.61 ± 0.23	<0.24	<6.21	0.31 ± 0.17
1060E,160S	0.76 🛖 0.20	<0.24	<3.49	0.58 ± 0.13
1140E, CS	0.61 + 0.21	<0.24	<4.52	0.43 4 0.13
1140E, 80S	0.68 <u>+</u> 0.25	<0.24	<3.22	0.25 <u>+</u> 0.13
1220E, 0S	0.45 + 0.19	<0.24	<4.72	C.78 <u>+</u> O.16

a Errors are 2c based on counting statistics.

TABLE 7

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE INCINERATOR AREA

100E, 05 100E, 105 100E, 205 100E, 305 100E, 305 100E, 405 100E, 505	0.57 ± 0.19° 0.54 ± 0.72 2.20 ± 0.28 0.52 ± 0.28 0.52 ± 0.70	<0.24 <0.24 <0.24	U-238 <2.94 <2.50	0.18 ± 0.0 <0.05
100E, 10S 100E, 20S 100E, 30S 100E, 40S 100E, 50S	0.54 ± 0.77 2.20 ± 0.28 0.52 ± 0.28	<0.24 <0.24	<2.50	
100E, 20S 100E, 30S 100E, 40S 100E, 50S	2.20 ± 0.28 0.52 ± 0.28	<0.24		<0.05
100E, 30S 100E, 40S 100E, 50S	n.52 ± 0.28			
100E, 40S 100E, 50S			<4.67	0.09 • 0.0
100E, 50S	0.52 <u>+</u> 0.20	<0.25	<4.07	0.45 ± 0.1
		<0.24	<1.89	0.30 ± 0.1
100E. 60S	<0.13	<0.24	5.69 <u>+</u> 8.91	0.71 + 0.0
	0.54 ± 0.02	<0.24	<3.80	0.12 ± 0.1
100E, 70S	0.72 ± 0.30	CO. 76	<4.15	0.17 ± 0.1
100E, 80S	0.71 ± 0.18	<0.24	<4.09	0.08 ± 0.0
10DE, 90S	0.84 <u>+</u> 0.22	<0.24	<4.05	0.09 ± 0.0
100E,100S	0.57 ± 0.25	<0.24	5.32 <u>+</u> 8.36	0.17 ± 0.1
110E, OS	0.67 ± 0.30	40.24	<4.19 <2.49	<0.04 0.15 ± 0.1
110E, 10S	0.61 ± 0.18	<0.24	<2.49	0.10 ± 0.1
110E, 20S	0.61 + 0.18	<0.24	<3.83	<0.03
110E, 30S	0.62 + 0.11	<0.24 <0.24	5.46 + 7.96	0.17 • 0.1
110E, 40S	0.60 ± 0.19		<3.28	<0.04
110E, 50S	0.47 <u>+</u> 0.18 0.71 + 0.18	<0.24	<2.73	0.10 + 0.1
110E, 60S		<0.24	<3.84	0.17 ± 0.0
110E, 70S	0.66 ± 0.21	<0.24 <0.24	<4.45	0.07 + 0.0
110E, 80S 110E, 90S	0.58 ± 0.19 0.82 ± 0.24	<0.24	<4.71	0.29 + 0.0
110E, 90S	0.75 ± 0.11	<0.24	<3.49	0.27 + 0.0
120E. OS	0.61 ± 0.11	<0.24	<3.83	<0.04
120E, 10S	0.85 ± 0.42	<0.24	<3.34	0.19 ± 0.1
120E, 20S	0.55 + 0.20	<0.24	<3.76	0.06 + 0.0
120E, 30S	<0.14	<0.24	<1.89	<0.03
120E, 40S	0.51 ± 0.07	<0.24	<4.06	<0.03
120E. 50S	0.59 + 0.30	<0.24	<4.64	<0.04
120E, 60S	0.42 . 0.21	<0.24	<4.11	0.50 4 0.1
120E, 70S	0.74 + 0.24	<0.28	<5.35	0.40 + 0.1
120E, 80S	0.42 + 0.20	<0.24	<5.19	0.08 + 0.0
120E, 90S	0.47 + 0.18	<0.25	<3.17	0.06 + 0.1
130E, OS	0.54 + 0.20	<0.94	<4.75	0_05 + 0_0
130E, 105	0.33 ± 0.21	<0.24	<3.95	0.51 + 0.1
130E, 20S	0.53 + 0.15	<0.24	<2.88	0.11 + 0.0
130E. 3US	0.74 + 0.18	<0.24	<3.06	0.12 ± 0.0
130E, 40S	0.41 + 0.15	<0.24	<3.02	<0.03
130E, 50S	0.63 ± 0.28	<0.24	<4.29	0.27 ± 0.1
130E, 60S	0.62 ± 0.20	0.18 ± 0.42 <0.24	<4.64 <3.62	$\begin{array}{c} 0.23 \pm 0.1 \\ 0.40 \pm 0.1 \end{array}$

TABLE 7, cont.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE INCINERATOR AREA

		Radionucliće (Concentrations (pCi/ _E))
Location	Fâ-776 ** *		U-236	Cb-131
130E, 80S	0.69 ± 0.22	<0.24	<5.05	0.39 + 0.1
130E, 90S	0.84 • 0.24	<0.24	<4.25	0.47 ± 0.1
140E, QS	0.57 + 0.19	<0.24	<3.78	<0.03
140E, 10S	0.46 + 0.22	<0.25	<2.91	0.29 + 0.1
140E, 20S	0.52 + 0.24	<0.24	7.86 + 8.36	0.05 ± 0.0
140F, 30S	0.65 🛨 0.24	₹0.24	6.04 - 12.29	<0.02
140E, 40S	0.55 <u>+</u> 0.11	<0.24	<3.60	<0.04
140E, 50S	0.96 + 0.37	< 0.24	<5.41	0.51 + 0.1
140E, 60S	0-59 ± 0.20	<0.24	<4.20	0.25 ± 0.1
140£, 70S	0.48 + 0.21	<0.24	<4.26	0.27 ± 0.1
140£, 80S	0.59 * 0.17	<0.24	<4.54	0.29 ± 0.1
140E, 90S	0.83 + 0.27	<0.24	<4.23	<0.03
150E, 0S	0.49 ± 0.16	<0.24	<3.57	0.41 <u>+</u> 0.1
150E, 10S	0.64 ± 0.23	<0.24	<3.92	0.61 ± 0.1
150E, 20S	0.56 ± 0.20	<0.24	<2.34	<0.03
150F, 30S	0.53 <u>+</u> 0.15	<0.24	<3.58	<0.03
150E, 40S	5-19 <u>+</u> 0.50	<0.24	<5.47	0.67 + 0.1
150E, 50S	0.59 <u>+</u> 0.19	<0.24	<3.63	0.25 + 0.1
1505, 603	0.54 ± 0.23	<0.24	<2.19	0-11 - 0-1
150E, 70S	0.51 + 0.18	<0.24	<3.92	0.09 ± 0.1
150E, 80S	0.63 ± 0.24	<0.24	<4.20	<0.03
150E, 90S	0.61 ± 0.21	<0.24	<3.80	0.28 ± 0.1
160E, OS	0.63 ± 0.24	<0.24	<3.86	<0.04
160E, 10S	U.46 <u>+</u> U.27	<0.24	<3,23	0.53 ± 0.1
160E, 20S	1.04 ± 0.29	0.20 ± 0.89	<5.32	0.51 ± 0.1
160E, 30S	6.61 + 0.60	1 67 + 1 06	19.9 👲 0.5	1.11 ± 0.1
160E. 40S	2.24 ± 0.34	<0.24 <0.74	<5.R2	0.89 ± 0.1
160E, 505	1.29 ± 0.29		<4.79	0.78 <u>+</u> 0.1
160E, 6DS	0.68 ± 0.21	<0.24	<2.45	0.11 ± 0.0
160E, 70S	0.56 ± 0.21	<0.24	<4.04	0.20 <u>+</u> 0.1
160E, 80S	<0.16	<0.24	<3.21	0.13 ± 0.1
160L, 90S	0.50 ± 0.16	<0.22	<3.15	0.33 + 0.1
170E, OS	0.82 + 0.21	<0.24	<3.14	0.45 + 0.1
170E, 10S	0.51 ± 0.18	<0.24	<3.39	0.57 <u>*</u> 0.1
170E, 20S	0.60 ± 0.39	<0.24	5.41 + 9.02	0.95 ± 0.1
170E, 30S		1.47 <u>+</u> 1.47	20.2 • 0.5	1.48 ± 0.2
170E, 40S	0.69 ± 0.24	<0.24	8.08 + 11.25	0.35 ± 0.1
170E, 50S	0.84 ± 0.22	<0.24	<3.82	0.32 ± 0.1
170F, 60S	0.53 ± 0.20	<0.24	<4.02	0.23 ± 0.1
170E, ⁷ 0S	0.67 + 0.22	<0.24	<4.51	0.06 ± 0.0

(1 (1)

TABLE 7, cont.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE INCINERATOR AREA

		Radionuclide C	oncentrations (pCi	ξ}
location	Rs-226	v-235	U-238	Cs-137
170E, &OS	0.49 ± 0.188	<0.24	<3.57	0.31 + 0.1
170E, 90S	0.64 ± 0.16	<0.24	<3.44	0.16 + 0.0
180E, OS	0.60 <u>+</u> 0.23	<0.24	<3.27	0.47 + 0.1
16GE, 10S	1.30 ± 0.24	<0.24	<4.48	0.70 + 0.1
160E, 20S	0.66 ± 0.21	<0.24	<4.21	0.34 ± 0.1
160E, 303	1.10 + 0.26	<0.24	54.67	0.40 ± 0.1
1801, 40s	0.54 <u>+</u> 0.27	<0.24	<5.05	0.40 + 0.1
160E, 50S	0.65 ± 0.19	<0.24	<4.46	0.19 + 0.1
1808, 605	0.66 + 0.23	<0.24	<3.58	0.19 ± 0.1
1807, 708	0.58 🗓 0.20	<0.24	<3.58	0.45 ± 0.1
180E, 80S	0.42 ± 0.23	<0.24	<5.67	0.59 + 0.1
180E, 30S	0.52 ± 0.19	<0.24	<4.37	0.21 + 0.0
190E, OS	0.45 + 0.23	<0.24	<2.89	<0.03
190E, 10S	0.72 + 0.20	<0.24	<4.65	0.14 + 0.0
190E, 20S	0.87 ± 0.26	<0.24	<4.01	0.55 • 0.1
190E, 30S	0.49 + 0.17	<0.24	<2.54	0.28 ± 0.0
190E, 40S	0.55 + 0.28	<0.25	<4.26	0.36 + 0.1
190E, 50S	0.62 + 0.20	<0.24	<1.36	0.37 + 0.1
190E, 60S	0.60 + 0.22	<0.24	<2.99	<0.04
190E, 70S	0.47 + 0.21	<0.24	<2.96	0.71 ± 0.1
190E, 80S	0.50 ± 0.24	<0.24	<2.35	0.17 + 0.0
200E, OS	0.55 + 0.20	<0.24	<3.96	0.06 + 0.0
2000, 103	0.46 - 0.22	NO.24	<4.30	U.20 + U.1
200E, 20S	6.61 <u>+</u> 0.50	<0.24	<6.04	0.46 ± 0.1
200E, 30S	0.48 + 0.19	<0.24	<2.46	0.52 + 0.1
200E, 40S	0.85 - 0.20	<0.24	<5.02	0.40 + 0.1
200F, 50S	<0.16	<0.24	<3.23	0.41 + 0.1
200E, 60S	0.53 + 0.22	<0.24	<0.64	0.21 + 0.0

a Errors are 20 based on counting statistics.

TABLE 8

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE TRACK STREET AREA

		Radionuclide Concentrations (gCu/g)					
Location	Ra-226	€-235	U-73B	C= -137			
670 Σ. 05	0.38 ± 0.13^{2}	<0.24	<5.64	0.77 <u>+</u> 0.17			
680E, 0S	0.56 ± 0.18	<0.24	<3.57	0.53 ± 0.13			
690E. 05	ъ	ь	ð	ь			
700E, 0S	1.23 ± 0.25	<0.24	<4.02	0.08 ± 0.11			
/10%, US	1.17 + 0.31	<0.24	<2.97	0.34 ± 0.11			
720E, 0S	1.59 ± 0.41	0.94 <u>*</u> 0.86	5.33 <u>+</u> 0.47	2.32 <u>+</u> 0.30			
730E, 08	1.56 + 0.38	<0.24	<5.60	0.84 ± 0.20			
740E, 05	2.60 + 0.22	<0.24	<4.57	0.82 + 0.18			
750E, 0S	4.41 ± 0.47	<0.24	3.44 <u>*</u> 0.54	0.50 ± 0.19			
7605, US	5.16 ± 0.02	<0.21	6.63 <u>+</u> 4.92	0.36 ± 0.08			
770%, OS	4.05 ± 0.51	<0.24	<7.71	1.10 ± 0.23			
780E, OS	0.91 <u>+</u> 0.28	<0.24	<5.13	0.17 ± 0.13			
79CE, 0S	2.44 <u>*</u> 0.33	<0.24	2.01 ± 0.46	0.59 ± 0.13			
600E, 05	1.51 <u>+</u> 0.33	<0.24	<6.39	1.21 ± 0.11			
810E, OS	0.59 ± 0.26	<0.24	<4.29	0.49 ± 0.19			
660E, 10S	0.70 ± 0.22	<0.24	0.97 ± 0.39	0.32 ± 0.11			
670F, 10S	2.13 ± 0.33	<0.24	<4.63	0.55 ± 0.14			
68CE, 108	1.30 ± 0.31	SQ.24	<5.69	0.66 ± 0.1			
690E, 105	1.10 ± 0.28	<0.24	<4.96	0.01 ± 0.1			
700E, 10S	0.91 ± 0.19	0.59 ± 0.45	<2.60	0.25 ± 0.0			
7108, 108	2.45 ± 0.38	<0.24	5.00 ± 0.52	0.71 + 0.1			
720E, 1GS	4.17 ± 0.46	<0.50	<7,81	0.51 ± 0.2			
790E, 105	2.25 <u>+</u> 0.36	<0.24	9.73 ± 0.14	1.01 ± 0.1			
740E, 10S	6.03 ± 0.17	0.67 ± 0.32	7.93 + 5.59	0.31 ± 0.0			
750E, 10S	3.37 <u>+</u> N.38	<0.24	<6.06	0.57 ± 0.1			
7601, 10S	2.41 ± 0.38	<0.24	<4.61	0.54 ± 0.1			
7701, 10S	2.58 ± 0.42	<0.24	<6.78	0.74 ± 0.2			
780E, 10S	1.64 + 0.40	<0.24	<4.83	0.58 ± 0.1			
790E, 10S	0.68 ± 0.21	<0.24	0.69 ± 0.35	0.69 ± 0.1			
800E, 10S	0.55 ± 0.20	<0.24	<4.04	0.77 ± 0.1			
610E, 10S	0.35 + 0.22	<0.24	<2.04	0.80 ± 0.1			
8202, 10 5	0.52 ± 0.24	<0.24	<5.15	0.33 ± 0.1			
650E, 20S	1.50 <u>+</u> 0.26	<0.24	<2.92	0.70 ± 0.1			
660E, 20S	9.74 ± 0.70	<0.60	<7.22	1.45 ± 0.2			
670E, 20S	0.86 ± 0.27	<0.24	<4.B3	<0.05			
680E, 20S	1.40 <u>+</u> 0.27	<0.24	<3.98	0.14 ± 0.1			
690E, 20S	6.49 ± 1.81	<2.27	2.68 ± 0.59	<0.29			
700E, 20S	1.45 ± 0.14	<0.24	<4.63	0.70 ± 0.1			
7101., 205	1.80 + 0.51	<0.24	<3.77	0.60 - 0.1			
770E, 20S	4.87 + 1.40	<1.68	1.49 ± 0.38	0.50 ± 0.9			
730E, 20S	3.37 + 0.43	<0.24	<5.77	0.66 + 0.1			

TABLE 8, CONT.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE TRACK STREET AREA

Radionuclide Concentrations (pCi/g)						
Location	Яя-276	U-235	U-23B	Cs-137		
740E, 20S	3.36 ± 0.44	<0.24	<5.65	0.73 ± 0.2		
750E, 20S	2.23 ± 0.38	<0.24	<5.71	0.68 ± 0.1		
760E, 20S	3.60 ± 0.41	<0.24	<5.61	0.47 ± 0.1		
770£, 20S	0.56 + 0.11	0.33 ± 0.59	<4.12	0.99 + 0.1		
780%. 20S	0.51 + 0.21	<0.24	<3.55	0.68 ± 0.1		
790F, 70S	0.59 + 0.21	<0.24	<4.48	0.22 ± 0.1		
800E, 70S	0.45 <u>+</u> 0.21	<0.24	<4.72	0.44 ± 0.1		
640E, 30S	3.71 ± 0.37	<0.74	<4.34	0.67 + 0.1		
650E, 30S	0.43 + 0.29	<0.24	<4.52	0.42 ± 0.1		
660E, 3QS	1.66 + 0.33	0.45 ± 0.78	<5.72	0.21 + 0.1		
670E, 30S	4.20 <u>+</u> 0.45	<0.24	<5.03	0.52 + 0.2		
680E, 3QS	3.19 ± 0.35	<0.24	<6.03	0.25 + 0.1		
690E, 30S	2.15 ± 0.44	<0.24	<6.20	0.78 + 0.2		
700E, 305	1.13 ± 0.33	0.55 ± 0.62	53.26	0,66 + 0.7		
710E, 30S	8.06 ± 0.20	<2.13	<7.48	<0.25		
720E, 30S	1.35 <u>+</u> 0.29	<0.24	1.36 ± 0.38	0.67 + 0.10		
730£, 30S	2.77 ± 0.26	<0.24	<5.81	0.79 ± 0.20		
740E, 305	1.34 ± 0.22	<0.24	<4.11	0.73 ± 0.16		
750E, 30S	1.95 ± 0.33	<0.24	<3.99	0.94 + 0.1		
760E, 3QS	1.30 ± 0.31	<0.24	<3.98	0.71 + 0.1		
770E, 30S	0.46 + 0.11	<0.24	<5.00	0.71 + 0.1		
760E, 305	0.57 ± 0.11	<0.23	<2.53	0.72 + 0.11		
790E, 30S	0.52 ± 0.19	<0.24	<3.74	0.55 ± 0.13		
620E, 40S	2.70 ± 0.35	<0.24	<4.55	0.55 + 0.13		
630E, 40S	1.82 <u>+</u> 0.29	<0.24	<3,42	0.53 ± 0.13		
640E, 405	0.56 + 0.21	<0.24	<2.55	0.83 ± 0.13		
630E, 405	ь	Ъ	b	Ь		
660E, 405	4.96 <u>+</u> 0.46	<0.24	<6.19	0.45 ± 0.16		
670E, 40S	0.88 <u>+</u> 0.30	<0.24	<3.51	0.88 ± 0.17		
680E, 405	1.54 ± 0.30	<0.24	<4.64	0.92 + 0.17		
690Z, 405	5.64 ± 1.83	<1.40	<11.18	<0.29		
700E, 40S	2.97 ± 0.22	<0.24	<5.68	0.76 ± 0.19		
710E, 4GS	3.36 <u>+</u> 0.37	<0.24	<5.06	0.38 ± 0.13		
720E, 40S	2.22 ± 0.34	0.65 <u>+</u> 0.84	<5.04	0.66 + 0.1		
730E, 40S	3.14 ± 0.42	<0.24	<4.74	1.35 2 0.2		
740£, 40S	0.54 ± 0.26	<0.24	<4.11	0.99 ± 0.10		
150€, 40S	0.70 <u>+</u> 0.56	<0.24	<4.50	0.72 + 0.11		
7602, 40s	0.22 ± 0.17	<0.17	<3.76	0.78 ± 0.11		
770E. 40S	0.58 ± 0.22	<0.24	<3.75	0.83 - 0.17		
610%, 50S	3.60 + 0.47	<0.24	1.03 + 0.30	0.82 + 0.22		

TABLE 8, cont.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE TRACK STREET AREA

Radionuclide Concentrations (pCi/g) location Ra-226 U-235 U-238 Cs-1					
location		U-235	0-236		
620E, 50S	0.60 + 0.29	<0.24	0.92 + 0.36	0.74 + 0.15	
63DE, 50S	0.72 ± 0.29		<3.63	1.28 + 0.21	
640E, 50S	ъ Ъ	ь	ъ	ъ	
650E, 50E		0.41 ± 0.63	₹6.59	0.56 ± 0.18	
660E, 50S	1.84 + 0.29	<0.24	6.51 ± 6.59	0.45 ± 0.14	
6708, 505	4.20 + 0.15	<0.16	2.90 ± 0.66	<0.25	
1801, 50s	0.64 + 0.19	<0.24	<3.49	<0.02	
h90%, 505	4.56 ± 0.46	<0.24	3.15 ± 0.50	0.77 + 0.16	
700E, 50S	2.95 ± 0.56	<0.24	<5.01	0.67 ± 0.17	
710£, 50S	1.61 + 0.41	<0.24	<7.38	0.51 ± 0.13	
720E 50S	0.61 + 0.24	<0.24	<4.93	0.72 * 0.16	
730E, 50S	0.49 + 0.16	<0.24	<3.05	0.55 ± 0.13	
740E, 50S	0.36 + 0.19	<0.24	<2.85	1.11 ± 0.10	
750E, 50S	0.47 + 0.18	<0.19	<3.59	0.31 <u>+</u> 0.10	
580E, 605	0.63 <u>+</u> 0.24	<0.26	<3.25	0.85 <u>+</u> 0.16	
590£, 60S	4.46 ± 0.38	<0.24	<4.87	0.70 ± 0.14	
6002, 605	0.99 <u>+</u> 0.26	<0.24	<3.76	0.69 ± 0.15	
610£, 60S	0.66 = 0.22	<0.24	6.20 ± 8.73	0.79 ± 0.16	
620E, 60S	5.56 ± 1.32	<1.69	1.94 <u>+</u> 0.59	<0.17	
630E, 60S	1.77 ± 0.31	0.42 <u>*</u> 0.74	<5.14	0.78 ± 0.17	
640E, 60S	0.77 ± 0.26	0.85 + 0.67	<3.68	1.25 ± Q.20	
650E, 60S	1.37 ± 0.38	<0.24	<4.46	0.77 <u>*</u> NO.1	
660E, 605	5.59 + 1.87	4.56 + 6.10	4.96 <u>+</u> 0.59	<0.28	
670E. 60S	2.90 ± 0.45	<0.24	<5.72	0.55 + 0.16	
680E, 60S	5.01 ± 0.56	<0.24	<6.72	0.72 ± 0.17	
690E, 60S	2.04 <u>+</u> 0.36	<0.24	<5.16	0.93 <u>+</u> 0.17	
700Σ, 60S	1.67 <u>+</u> 0.45	<0.24	<7.09	1.31 ± 0.23	
710E, 60S	0.75 ± 0.23	<0.24	<3.92	0.86 ± 0.17	
720E, 60S	0.60 ± 0.22	<0.23	<4.51	0.34 ± 0.13	
730E, 60S	0.57 ± 0.20	<0.24	<2.05	0.57 <u>+</u> 0.13	
740E, 60S	0.50 ± 0.19	<0.24	<3.19	0.71 + 0.0	
580E, 70S	2.17 <u>*</u> 0.37	<0.24	2.78 + 0.37	0.66 <u>+</u> 0.19	
590E, 70S	0.75 <u>+</u> 0.28	<0.24	<4.30	0.84 <u>+</u> 0.10	
600E, 70S	0.53 <u>+</u> 0.28	0.51 • 0.68	3.95 ± 0.36	0.86 ± 0.20	
610E, 70S	<0.19	<0.24	<4.00	0.71 ± 0.14	
620Z, 70S	$1.22 \pm 0.2I$		<4.26	0.78 <u>+</u> 0.11	
630E, 70S	2.09 + 0.36	<0.24	1.12 + 0.51	0.74 ± 0.13	
640E, 70S	5.31 ± 1.67	<1.61	<22.85	<0.26	
650E, 70S	2.56 <u>*</u> 0.37	<0.24	<7.53	0.66 • 0.16	
660E. 70S	1.21 + 0.27	<0.24	<4.48	0.36 • 0.1	

TABLE 8, cont.

RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE TRACK STREET AREA

	Radiomuclide Concentrations (pCi/g			
Loration	Ra-226	U-235	U-238	Cs-137
670%. 70S	0.72 + 0.33	<0.24	<4.47	
680E, 70S	0.75 + 0.25	<0.24	<4.14	0.82 ± 0.15
690E, 70S	0.83 ± 0.26	<0.29	<3.47	0.33 ± 0.16 0.73 ± 0.15
700E, 70S	0.69 ± 0.32	<0.24	(3.31	0.53 + 0.13
710E, 70S	0.35 + 0.17	<0.24	<2.55	0.69 ± 0.13
590E, 80S	1.07 + 0.36	<0.24	<5.23	1.20 + 0.21
600E, 80S	0.56 ± 0.25	<0.24	<6.43	0.82 ± 0.16
610E, 80S	0.64 + 0.25	<0.24	<4.92	<0.05
620E. 80S	1.89 + 0.33	<0.24	<5,32	1.06 + 0.20
630E, 80S	6.59 + 1.82	<2.38	2.05 + 0.56	
640E. 80S	3.84 + 0.43	<0.24	<4.59	0.60 ± 0.67
650E. 80S	5.11 ± 0.48	<0.24	<5.90	0.76 <u>+</u> 0.18 0.58 + 0.16
660E, 80S	<0.16	<0.24	<3.76	0.55 ± 0.14
670E, 80S	0.64 ± 0.30	0.37 + 0.56	<4.36	0.96 + 0.19
680E, 80S	1.44 ± 0.19	<0.24	<3.48	0.47 + 0.09
690E, 80S	<0.44	<1.06	0.36 + 0.27	1.37 + 0.72
700E, 80S	0.98 + 0.35	<0.25	9.29 ± 7.93	0.63 + 0.17
580E, 90S	0.63 + 0.29	<0.24	<4.71	0.19 ± 0.11
590E, 90S	0.96 + 0.28	<0.24	<5.70	0.63 + 0.18
600L 90S	0.80 ± 0.30	<0.24	<6.14	1.07 + 0.19
610E. 90S	<0.46	<0.22	4.70 ± 0.51	<0.24
620E, 90S	1.44 + 0.36	<0.24	1.20 + 0.37	0.85 + 0.16
630E, 90S	3.71 + 0.45	1.01 ± 0.75	<5.58	1.04 + 0.19
640E, 90S	2.83 • 0.42	1.29 ± 0.78	46,20	0.70 • 0.18
650E, 90S	0.55 ± 0.18	<0.24	<2.81	0.08 + 0.07
660E, 90S	0.69 + 0.25	<0.22	<2.52	0.75 + 0.18
670E. 90S	0.67 • 0.27	<0.24	<4,42	0.34 + 0.13
680E, 905	0.61 + 0.12	<0.24	<4.61	0.16 ± 0.10
580E, 100S	0.73 + 0.29	<0.24	<4.10	0.69 + 0.16
590E.100S	4.94 + 2.23	<2,22	3.78 + 0.66	<0.30
6COE, 100S	1.35 + 0.29	<0.24	<3.64	0.84 + 0.19
610E,100S	2.88 + 0.46	<0.24	<4.01	1.12 + 0.21
670E,100S	2.38 ± 0.32	<0.24	<4.72	0.80 + 0.18
630E,100S	1.86 ± 0.34	<0.24	<4.20	1.02 + 0.18
640E,100S	0.90 + 0.23	0.34 + 0.48	<4.36	0.54 + 0.10
650E.100S	0.70 + 0.21	<0.26	<4.80	0.55 + 0.16
660£,1003	0.50 ± 0.21	<0.20 <0.24	<4.80 <3.42	0.55 ± 0.16 0.63 ± 0.17
670E,100S	0.30 ± 0.19 0.31 ± 0.24	<0.24	<.78	0.52 ± 0.15

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TABLE 8, cont. RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM 10 M GRID INTERVALS IN THE TRACK STREET AREA

		Radicnuclide Concentrations (pCi/g)					
10	cation	Ra-226	υ-235	ν-238	Cs-137		
 580	E.110S	6.24 + 2.67	<1.79	4.62 + 0.54	0.68 <u>+</u> 0.69		
	11.1105	5.66 + 0.50	0.83 + 0.69	<6.76	0.74 ± 0.26		
	E,110S	1.92 + 0.41	<0.24	<5.17	1.21 ± 0.22		
	E,110S	0.68 ± 0.19	<0.24	<3.43	1.02 + 0.15		
	E,1108	0.36 ± 0.21	<0.22	<3.00	0.48 • 0.16		
	E 1105	0.64 + 0.22	<0.24	<3.96	0.67 ± 0.13		
	E.110S	0.31 + 0.29	<0.23	<2.20	0.86 + 0.12		
	E.1105	0.52 + 0.25	<0.24	<4.15	0.39 ± 0.13		
	E.110S	0.60 ± 0.21	0.22 ± 0.70	<1.57	<0.04		
580	E.120S	0.40 + 0.20	<0.24	<3.23	0.31 ± 0.13		
	DE.1208	0.65 + 0.25	<0.24	<4.26	0.71 ± 0.17		
600	E,120S	0.56 🛓 0.23	<0.24	<5.62	0.66 + 0.16		
610	E.120S	0.63 - 0.21	<0.24	<5.02	0.16 ± 0.00		
6.20)L,120S	0.87 + 0.27	<0.24	<4.89	0.22 ± 0.14		
630)E,120S	0.39 € 0.20	<0.24	<2.99	1.35 ± 0.19		
	E,1308	0.72 + 0.22	<0.24	<4.23	0.56 ± 0.13		
	E,130S	0.74 + 0.22	<0.24	4.16 <u>*</u> 8.76	0.49 ± 0.14		
	DE.130S	0.74 + 0.28	<0.24	<3.92	0.97 ± 0.22		
	DE.130S	0.28 + 0.15	<0.24	<2.31	0.87 ± 0.14		
629	DE.130S	0.74 + 0.23	<0.24	<3.58	0.75 ± 0.15		
	DE.140S	1.00 + 0.24	<0.24	<3.13	0.62 ± 0.16		
591	DE.1408	0.68 ± 0.21	<0.24	<3.30	0.69 ± 0.14		
	DE. 140S	0.45 + 0.21	<0.26	<4.50	0.45 ± 0.13		
	DE,1508	0.32 + 0.22	<0.24	<3.14	0.59 + 0.13		

 $^{^{\}rm a}$ Errors are 2σ based on counting statistics. $^{\rm b}$ Sample not obtained - grid point on paved surface.

TABLE 9 RADIONUCLIDE CONCENTRATIONS IN SURFACE SOIL SAMPLES FROM LOCATIONS IDENTIFIED BY THE WALKOVER SCAN

ample			Radionuclide Concentrations (pCi/g)b				
No.a	Location	Ra-226	บ−235	U-238	Cs-137		
в1	148E, 43S	3.99 + 1.43°	4.49 ± 2.33	51.6 <u>+</u> 0.7	0.51 <u>+</u> 0.17		
B2	154E, 38S	15.1 ± 0.9	1.73 ± 1.78	17-8 <u>+</u> 0-5	0.57 ± 0.23		
В3	153E, 36S	45.1 \pm 1.8	4.57 ± 2.48	68.5 ± 0.8	1.33 ± 0.39		
В4	163E, 31S	141 <u>+</u> 3	<0.25	23.3 ± 0.5	3.15 ± 0.59		
В5	166E, 29S	26.5 ± 1.2	0.93 ± 1.68	6.13 ± 0.39	6.52 ± 0.40		
В6	170E, 30S	88.3 ± 1.9	3.33 ± 2.88	39.3 ± 0.6	1.18 ± 0.22		
В7	169E, 32S	9.32 ± 0.72	1.18 ± 1.46	22-2 ± 0.5	5.93 <u>+</u> 0.43		
В8	178E, 32S	63.7 \pm 1.9	<0.24	22.3 <u>+</u> 0.5	1.09 ± 1.15		
В9	202E, 37S	23.3 ± 1.2	3.99 <u>+</u> 1.99	34.7 <u>+</u> 0.6	2.77 <u>+</u> 0.43		
B10	109E,210S	d					
B11e	582E, 76S	0.44 ± 0.56	9.08 ± 2.19	125 <u>+</u> 1	0.82 ± 0.27		
B12f	584E, 74S	0.98 ± 0.32	6.57 ± 2.11	119 <u>+</u> 1	0.68 ± 0.28		
B13	641E, 50S	246 <u>+</u> 3	6.64 ± 4.75	10.4 ± 0.5	<0.49		
B14	660E, 39S	1.39 ± 0.44	2.57 ± 1.42	32-2 + 0.5	3.40 <u>+</u> 0.74		
B15	668E, 33S	222 ± 3	6.87 ± 5.17	24.4 <u>+</u> 0.6	0.54 <u>+</u> 0.09		
B16	669E, 31S	55.3 + 1.7	3.79 ± 2.71	10.7 ± 0.5	2.16 ± 0.46		
B17	668E, 30S	431 <u>+</u> 4	<0.24	6.49 ± 0.39	<0.59		
B18	684E, 10S	6.05 + 0.54	<0.24	<6.31	2.94 ± 0.28		
B19	738E, 4S	6.95 ± 0.65	3.01 + 1.46	41.8 ± 0.7	0.55 ± 0.22		

a Refer to Figures 4, 5, and 6.
b Refer to Table 5 for direct radiation levels.
c Errors are 20 based on counting statistics.
d Ra-226 level 3.73 pCi.
e Sample B11 contains 13.9 ± 1.2 Th-232.
f Sample B12 contains 12.5 ± 1.2 Th-232.

TABLE 10 RADIONUCLIDE CONCENTRATIONS IN BOREHOLE SOIL SAMPLES

orehole	Grid	Deptb		Radionuclide Concentrations (pCi/g) Ra-226 U 235 U-238 Cs-137			
No.a	Location		Ra-226	u 235	U-238		
			0.33 + 0.24 ^b	<0.74	<4.59	<0.03	
	,	2.5	0.67 ± 0.23	<0.24	<2.64	0.57 <u>+</u> 1.49	
112	121E, 236	Surface				0.45 ± 0.07	
		0.5	0.75 ± 0.18 0.50 ± 0.14	<0.24 <0.24	<3.12	<0.02	
		1.0	0.50 <u>+</u> 0.14	<0.24	<3.44	<0.03	
		2.0	0.52 ± 0.15	0.27 ± 0.59	<2.79	<0.03	
		3.0	0.45 ± 0.16	<0.24	3.80 <u>+</u> 13.61	<0.02	
ыз	380E,140S	Surface	0.93 + 0.29	<0.24		0.39 ± 0.13	
	-	0.5	0.64 + 0.24	<0.24	<4.86	<0.03	
		1.0	0.64 + 0.24 0.54 + 0.22 0.75 + 0.20	<0.24	<5.12	<0.04	
		2.0	0.75 2 0.20	<0.24	54.86	0.05 <u>+</u> 0.05	
		3.0	0.38 - 0.19	<0.22	<4.01	<0.03	
E4	490E.321S	0.3	0.50 + 0.14	<0.24	<2.82	<0.02	
		3.0	$\begin{array}{c} 0.50 \pm 0.14 \\ 0.56 \pm 0.28 \end{array}$	<0.24	<4.09	<0.02	
н5	560E. 85S	Surface	2.32 ± 0.32	<0.24		0.90 • 0.14	
	•	0.5	0.49 + 0.16	<).24	,3.41	<0.03	
		1.0	0.49 ± 0.16 0.61 ± 0.17	<0.18	<3.44	<0.03	
			0.41 ± 0.16	<0.24	<4.34	0.47 + 0.12	
H6	560E,160S	Surface	$\begin{array}{c} 0.29 \pm 0.16 \\ 0.53 \pm 0.21 \\ 0.55 \pm 0.23 \\ 0.42 \pm 0.22 \end{array}$	<0.24	<1.98	0.15 ± 0.08	
	,	0.5	0.53 + 0.21	<0.24 <).24	<2.95	<0.03	
		1.0	0.55 ± 0.23	<0.24	<2.84	<0.04	
		2.0	0.42 ± 0.22	<0.19	<2.39	<0.03	
		3.0	1.39 ± 0.18	<0.24	<2.66	<0.03	
H?	580E.100S	Surface	0.37 + 0.36	<0.24		0.27 ± 0.10	
-		0.5	0.37 ± 0.36 0.44 ± 0.21	<0.24	<3.70	<0.03	
		1.0	0.51 + 0.18	<0.24 0.41 <u>+</u> 0.50	<4.65	<0.03	
		2.0	0.75 + 0.22	<0.24	<4.01	0.54 ± 0.09	
		3.0	$\begin{array}{c} 0.75 \pm 0.22 \\ 0.61 \pm 0.14 \end{array}$	<0.24	<4.34	<0.03	
18	625E.1158	Surface	3.14 ± 0.45		<5.63	0.19 ± 0.20	
	,	0.5	0.43 ± 0.20	<0.24	<4.33	<0.03	
		1_6	$\begin{array}{c} 0.43 \pm 0.20 \\ 0.62 \pm 0.25 \end{array}$	<0.24 <0.74	<5.35	<0.03	
		2.0	0.63 ± 0.19	<0.24	<4.23	0.06 + 0.06	
Н9	680E, 58	Surface	0.77 ± 0.25	<0.24	<2.97	0.47 ± 0.12	
	, >0		0.52 + 0.21	<0.24	<4.97	<0.03	

RADIONUCLIDE CONCENTRATIONS IN BOREHOLE SOIL SAMPLES

TABLE 10, cont.

orehole	Grid	Dept h	epth Radiomurbide Concentrations (pCi/g)					
No.	Location	(4)	Ru-226	U-235	C-238	Cs~137		
		1.0	0.60 ± 0.20	<0.24	<4.63	<0.03		
		2.0	0.66 ± 0.17	<0.24	<3.97	<0.03		
		3.0	0.59 ± 0.22	<0.24	<4.32	<0.03		
H10	702E, 20S			<0.24	<5.18	0.62 ± 0.14		
		0.5	0.68 + 0.24	<0.24	<3.14	<0.03		
		1.0	0.51 ± 0.22	<0.24	<3.54	<0,04		
		2.0	1.16 <u>+</u> 0.30	<0.24	<3.55	<0.03		
		3.0	0.50 ± 0.20	<0.24	<3.63	<0.04		
<u>u11</u>	745E, 37S	Surface		<0.24	₹4.14	0.64 ± 0.17		
		0.5	0.12 ± 0.25	<0.24	<4.99	<0.04		
		1.0	0.58 + 0.18	<0.24	<2.73	<0.04		
		2.0	0.59 ± 0.16	<0.24	<3.51	<0.03		
H1 2	7782,3195	0.3	0.50 ± 0.14	<0.24	<2.62	<0.02		
		3.0	0.56 ± 0.78	<0.24	<4.09	<0.02		
B13	1063E,155S		0.48 ± 0.17	<0.23	<3.77	<0.04		
		2.0	0.54 ± 0.17	<0.24	<3.50	<0.03		
		3.0	0.61 ± 0.18	<0.19	<3.47	<0.02		
H14	1080£, 20\$			<0.24	<3.10	0.65 ± 0.12		
		0.5	0.88 <u>+</u> 0.22	<0.24	<4.94	<0.04		
		1.0	1.17 ± 0.30	<0.24	<4.82	<0.04		
		2.0	1.16 + 0.28	<0.24	<4.37	<0.03		
		3.0	0.95 <u>+</u> 0.26	<).24	<4.43	<0.04		
H15	156E, 30S	Surface		<0.41	<8.03	0.16 ± 1.10		
		1.0	1.80 + 0.29	<0.36	2.73 * 0.38	<0.06		
H16	144E, 43S	Surfacec	1.66 • 0.32	0.68 + 0.61	10.6 ± 0.4	0.85 ± 0.37		
H17	154E, 38S	Surface	15.1 ± 0.9	1.73 ± 1.78	17.8 ± 0.5	0.57 ± 0.23		
H18	162E, 32S	0.3	0.66 <u>+</u> 1.16	<0.04	<5.87	<0.04		
H19	168E, 49S	0.3	11.6 ± 0.7	1.38 • 1.18	1.22 * 0.32	1.21 <u>*</u> 0.23		
E20	170E, 22S	Surface	0.63 ± 0.22	<0.30	<5.73	0.14 + 0.18		

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TABLE 10, cont. RADIONUCLIDE CONCENTRATIONS IN BOREHOLE SOIL SAMPLES

Burchole	Grid		Radiopurlide Concentrations (pCi/g)					
No.	Location		Ra-226	U-235	U-23B	Cs=137		
H21	170E, 30S	Surface 0.1	88.3 ± 1.9 3.36 ± 1.33	3.33 ± 2.86 2.36 ± 2.02	39.3 ± 0.6 78.6 ± 0.5	1.16 ± 0.22 <0.23		
Н22	178E, 32S	Surface¢	63.7 <u>*</u> 1.9	<0.24	22.3 <u>+</u> 0.5	1.09 ± 1.15		
H23	193E, 36S	1.0	0.89 + 0.24	<0.29	<6.05	<0.05		
1124	584E, 74S	Surface 0.5 1.0	$0.98 \pm 0.32 \\ \stackrel{< 0.16}{< 0.43} \pm 0.19$	6.57 <u>+</u> 2.11 <0.24 <0.24	119 <u>+</u> 1 <4.63 <4.41	0.68 ± 0.28 <0.04 <0.03		
H75	641E, 50S	Surface	246 • 3	6.64 <u>*</u> 4.75	10.4 ± 0.5	<0.49		
£26	666E, 30S	Surfacec	431 <u>+</u> 4	<0.24	6.49 ± 0.39	<0.59		
н27	668E, 33S	Surfacec	222 ± 3	6.67 ± 5.17	24.4 ± 0.6	0.54 ± 0.09		

a Refer to Figures 7, 8, and 9.
b Errors are 20 based on counting statistics.
c Subsurface sample not collected due to negative findings of borehole logging measurements.

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TABLE 11 RADIONUCLIDE CONCENTRATIONS IN SUBSURFACE WATER SAMPLES

entrations (pCi/1)	Radionuclide Conc	Grid	Borehole	Sample Boreh	
Gross Beta	Gross Alpha	Location	No.ª	No.	
8.53 ± 7.75	<5.21	92E,269	H1	Wl	
5.55 ± 2.13	9.06 ± 2.42b	121E, 23S	H2	W2	
2.34 ± 12.32	8.86 ± 9.21	380E,140S	н3	W3	
13.5 ± 5.9	<3.67	490E,321S	H4	W4	
21.9 + 2.7	3.23 + 1.96	560E, 85S	H5	₩5	
6.30 ± 2.23	1.47 ± 1.98	560E,160S	н6	W6	
28.5 ± 8.7	11.7 \pm 7.7	580E,100S	H7	W7	
12.6 + 8.3	1.85 + 7.04	625E,115S	Н8	W8	
24.5 + 5.9	4.47 + 4.25	680E, 5S	Н9	W9	
<1.36	<1.17	702E, 20S	H10	W10	
7.99 + 5.61	4,52 + 5,03	745E, 37S	н11	W11	
<5.42	8.53 ± 4.79	778E,319S	H12	W12	
3.60 + 8.27	13.9 + 6.8	1063E,155S	н13	W13	
44.0 ± 4.1	4.61 + 2.74	1080E, 20S	н14	W14	

 $[\]begin{array}{c} a \\ b \end{array} \ \, \begin{array}{c} \text{Refer to Figure 7.} \\ \text{Errors are 2}\sigma \ \, \text{based on counting statistics.} \end{array}$

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- 1. E.A. Vierzba and A. Wallo, A Background Report and Evaluation of Resurvey Requirements for the Former Atomic Energy Commission Portion of the Lake Ontario Ordnance Works, Aerospace Corp., November 1982.
- 2. Oak Ridge Operations, U.S. Atomic Energy Commission, <u>Radiation Survey and Decontamination Report of the Lake Ontario Ordnance Works Site</u>, Oak Ridge, TN, January 1973.
- 3. T.E. Myrick, et al., Preliminary Results of the Ground-Level Gamma-Ray Scanning Survey of the Former Lake Ontario Ordnance Works Site Draft Report, ORNL, Oak Ridge, TN, 1981.

APPENDIX A

INSTRUMENTATION AND ANALYTICAL PROCEDURES

APPENDIX A

Instrumentation and Analytical Procedures

Gamma Scintillation Measurement

Walkover surface scans and measurements of gamma exposure rates were performed using Eberline Model PRM-6 portable ratemeters with Victorean Model 489-55 gamma scintillation probes containing 3.2 cm x 3.8 cm NaI(Tl) scintillation crystals. Count rates were converted to exposure levels ($\mu R/h$) using factors determined by comparing the response of the scintillation detector with that of a Reuter Stokes model RSS-III pressurized ionization chamber at several locations on property N/N' South.

Beta-Gamma Dose Rate Measurements

Measurements were performed using Eberline "Rascal," Model PRS-1, portable scaler/ratemeters with Model HP-260 thin-window, pancake G-M, beta probes. Dose rates (μ rad/h) were determined by comparison of the response of a Victoreen Model 440 ionization chamber survey meter to that of the G-M probes for soil samples having elevated Ra-226 content.

Borehole Logging

Borehole gamma radiation measurements were performed using a Victoreen Model 489-55 gamma scintillation probe, connected to a Ludlum Model 2200 portable scaler. The scintillation probe was shielded by a 1.25 cm thick lead shield with four 2.5 cm x 7 mm holes evenly spaced around the region of the scintillation crystal. The probe was lowered into each hole using a tripod holder with a small winch. The length of the hole was scanned and measurements were performed at 30-50 cm intervals in all holes. The logging data were used to identify regions of possible residues and guide the selection of subsurface soil sampling locations. Due to the varying ratios of Ra-226, U-238, and Cs-137 there was no attempt to estimate soil radionuclide concentrations directly from the logging results.

Soil Sample Analysis

Soil samples were dried, mixed, and a portion placed in a 0.5 liter Marinelli beaker. The quantity placed in each beaker was chosen to reproduce the calibrated counting geometry and ranged from 400 to 600 g of soil. Net soil weights were determined and samples counted using a 23% Ge(Li) detector (Princeton Gamma Tech) coupled to a Nuclear Data model ND-680 pulse height analyzer system. Background and Compton stripping peak search, peak identification, and concentration calculations were performed using the computer capabilities inherent in the analyzer system. Energy peaks used for determination of radionuclides of concern were:

Ra-226 - 0.609 MeV from Bi-214 (secular equilibrium assumed)

U-235 - 0.143 MeV

U-238 - 1.001 MeV from Pa-234 (secular equilibrium assumed)

Th-232 - 0.911 MeV from Ac-228 (secular equilibrium assumed)

Cs-137 - 0.662 MeV

Several samples were also analyzed for U-238 by neutron activation. Approximately 12-20~g of soil was irradiated for 30 minutes in a neutron flux of $10^8~n/cm^2/sec$ from a 25 mg Cf-252 spontaneous fission source. After a two minute wait time, the U-239 peak (74.6 kev) was counted for 1000 seconds and the U-238 calculated.

Strontium-90 analysis was performed following standard procedures specified in "Radiochemical Analytical Procedures for Analysis of Environmental Samples," EMSL-LV-0539-17, March 1979.

Water samples were rough-filtered through Whatman No. 2 filter paper. Remaining suspended solids were removed by subsequent filtration through 0.45 µm membrane filters. The filtrate was acidified by addition of 20 ml of concentrated nitric acid. Fifty milliliters of each sample was evaporated to dryness and counted for gross alpha and gross beta using a Tennelec Model LB5100 low-background proportional counter.

Calibration and Quality Assurance

With the exception of the exposure and dose rate conversion factors for portable survey gamma and beta-gamma meters, all survey and laboratory instruments were calibrated with NBS-traceable standards. The calibration procedures for these portable instruments are described above.

Quality control procedures on all instruments included daily background and check-source measurements to confirm lack of malfunctions and nonstatistical deviations in equipment. The ORAU laboratory participates in the EPA Quality Assurance Program.

APPENDIX B

SUMMARY OF RADIATION GUIDELINES APPLICABLE TO OFF-SITE PROPERTIES AT THE NIAGARA FALLS STORAGE SITE

E,

SUMMARY OF RADIATION GUIDELINES APPLICABLE TO OFF-SITE PROPERTIES AT THE NIAGARA FALLS STORAGE SITE

	Made of Expasure	Exposure conditions	Guideline value	Guideline source
1.	External gamma radiation	Continuous exposure to individual in general population (whole body)	60 pR/be	Nuclear Regulatory Commission (NRC) Standards for Protection Against Radiation (10 CFR 20.105
		Indoor gamma radiation (above background)	20 uR/h	EPA Standards for Uranium Mill Tailings (40 CFR 192)
2.	Surface alpha contamination [®]	Ra-216 contamination fixed on surfaces	100 dpm/100 cm ²	KRC Guidelines for Facilities and Equipment Prior to Release for Uprestricted Use or
		Removable Ra-226	20 dpm/100 cm ²	Termination of Licenses for By-product, Source, or Special Nuclear Material (Adapted from NRC Reg. Guide 1.86)
٦.	Surface beta contamination®	Removable beta-gamma	1000 dpm/100 cm ²	Same as number 2
4.	Beta-gamma dose rate	Average dose rate on an area no greater than 1 u^2	0.2 mrad/b	Same as number 2
		Maximum dose rate in any $100~{ m cm}^2$ area	1.0 mrad/h	Same as number 2
5.	Exposure to radon	Maximum permissible concentration of Ru-220 in air in unrestricted areas	-3.0 pCi/1	NRC 10 CFR 20.103, Appendix B, Table 11
		Average annual radon daughter concentration (including background)	0.030 WL maximum 0.020 WL preferrable	EPA Standards for Fill Teilings

SUPPLIES APPLICABLE TO OFF-SITE PROFESTIES AT THE WINGARA FALLS SIGNAGE SITE, CODE.

Mode of Exposure	Exposure conditions	Cuiceline value	Guideline source
6. Radionuclides in water	Gross Alpha Gross Beta Naximum contaminant level fur combined Ra-226 and Ra-228 in drinking water	15 pci/1 50 pci/1 5 pci/1	EFA Interim Drinking Water Standards (40 GFR 141)
	Maximum permissible concen- tration of the following radionuclides in water for unrestricted area:		hau (10 CFR 20.103 Appendix E, Table II)
	R.a-276 U-238 Ta-230 Fb-210	30 pci/1 40,000 pci/1 2,000 pci/1 100 pci/1	
7. Radionuclides in soil	Concentration above background-averaged over an area of 100 m ²		
	Re-126	5 pCi/g (surface) 15 pCi/g (subsurface)	EPA Standards for Uranium Hill Tailings
	G-238 Th-232 Sr 90 G-137	40 pci/g 100 pci/g 100 pci/g EO pci/g	Interim Soil Limits for DSD Projects, LA-UR-79-1865-Rav, J.W. Healy et al.
	V-enriched in V-235	30 pCi/g	NRC Branch Technical Position Paper (Federal Register, October 23, 1981)

 $^{^{\}mbox{\scriptsize A}}$ Applicable to building and equipment surfaces only.

APPENDIX C

REPORT OF GROUND-PENETRATING RADAR SURVEY
OF OFF-SITE PROPERTY N/N' SOUTH
AT THE NIAGARA FALLS STORAGE SITE

FINAL REPORT

GROUND-PENETRATING RADAR SURVEY

OF AREAS Q AND N/N' AT THE

FORMER LAKE ONTARIO ORDNANCE WORKS

LEWISTON, NEW YORK

496 HEALD ROAD

CARLISLE, MASSACHUSETTS 01741

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FINAL REPORT

GROUND-PENETRATING RADAR SURVEY

OF AREAS Q AND N/N' AT THE

FORMER LAKE ONTARIO ORDNANCE WORKS

LEWISTON, NEW YORK

Prepared for
OAK RIDGE ASSOCIATED UNIVERSITIES, INC
Oak Ridge, Tennessee 37830

Purchase Order No. C-25303 Letter Release No. 3 Dated October 15, 1982

Report No. J147-82

December, 1982

DETECTION SCIENCES GROUP

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INTRODUCTION AND SUMMARY

Starting on October 25, and continuing through November 7, 1982, Detection Sciences Group conducted a ground-penetrating radar survey of designated locations in Area Q and Areas N/N' at the former Lake Ontario Ordnance Works, Lewiston, New York. The radar survey was performed in accordance with Oak Ridge Associated Universities, Inc. Purchase Order No. C-25303, Letter Release No. 3, dated October 15, 1982, and per field instructions received from O.R.A.U. personnel.

The first area to be surveyed was a site in Area N at the southeast corner of "R" Street and Castle Garden Road. The radar survey zone was from 20S to 50S, extending from 140E to 185E. The radar anomalies and the radar survey lines are shown in Figure 1. The next task to be performed was the inspection of 24 boring sites located in Area Q. The boring locations and the co-ordinates of the borings that were relocated are given in Table 1. The third task was to run a series of survey lines between 760S and 840S, extending from 160E to 180E in Area Q. The results of the survey in this location are shown in Figure 2. The next task was to run six survey lines parallel to Track Street and South Track Street in Area N'. The radar survey lines and the radar anomalies are shown in Figure 3. The last task was to inspect the locations of 3 borings to be placed in the corners of the southwest triangle of Area N, bounded by Castle Garden Road, "R" Street, and Track Street. These locations are tabulated in Table 2.

Out of the 24 boring locations surveyed in Area Q, 3 locations - Q1, Q4, and Q9 - were found to have underground pipes. These locations were moved 2 to 3 meters away from the pipes to avoid potential drilling problems. As a precautionary measure, 8 other boring locations in Area Q were moved 1 to 3 meters away from their original locations. In all cases, the relocated sites were chosen to be as near as possible to the original site chosen for the boring. Of the 3 boring sites in the southwest triangle of Area N, none had to be relocated.

Prior to the radar work, O.R.A.U. had placed pin flags at locations where monitoring instruments indicated activity. It was noted that the radar signatures obtained where there were clusters of pin flags were significantly different than the radar signatures observed over most of the surrounding area. Specifically, the radar signatures at the pin flag clusters tended to be much darker and more pronounced than the prevailing radar signatures of the overall site. The apparent correlation between the radar signatures and the pin flag clusters has been subdivided into two categories: those areas having very dark, strong signal returns, and those areas having moderately dark, strong returns. The difference between the two subdivisions is only a difference of degree. In general, this kind of radar signature is indicative of non-ionic liquids being present in the ground. Although water is itself a non-ionic liquid, the radar signatures are not typical for wet, saturated ground. More likely, the radar signatures are due to the presence of petroleum solvents or oils that do not readily disperse in the ground. Determination of the composition of the non-ionic liquids (including the possibility that it is perched water) would require physical samples to be taken from these locations, a task that is not part of the radar survey program.

DESCRIPTION OF THE SURVEY

The entire survey was conducted with the 120 MHz radar antenna. With the exception of the survey lines along Track Street and South Track Street, the survey was run with our antenna shield on the antenna to prevent any stray, above-ground reflections or electrical interference from obscuring the data.

The survey vehicle was used to tow the antenna on the long survey lines along Track Street and South Track Street. For the north-south survey lines between 760S and 840S in Area Q, we also utilized the vehicle to tow the antenna over the ground. All of the other survey work was performed by hand-pulling the 120 MHz antenna over the ground.

A significant gain was made in survey efficiency by using a labor-saving device consisting of a knotted rope 12 meters long. At the center of the rope was a loop through which a chaining pin or other pointed device could be inserted into the ground at the proposed location for a boring. The knots in the rope were spaced at 2 meter intervals, and were painted with high-visibility paint. Using this rope made it unnecessary to lay out a steel tape to measure and mark the 2 meter intervals used for ground locations in the vicinity of the proposed boring. Also, by leaving the center pin in place, the 12-meter rope was quickly rotated 90 degrees on the ground to mark off the orthogonal survey line crossing the proposed location of the boring. Using this technique, we were able to inspect more than twice as many boreholes in a single day as had previously been the case with manually laying out the 2-meter ground locations with a steel tape.

With the exception of using the antenna shield and the knotted marker rope, the survey methods were the same as have been described in previous reports.

: :
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:
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Area N'.

The area scanned by the radar between Track Street and South Track Street is shown in Figure 3. This area is surprisingly free of any radar evidence of past usage, and shows mostly undisturbed ground free from any type of contaminants. A few, highly localized concentrations of dark and medium dark radar reflectors are observed, but do not appear to have any significance (in contrast to the dark reflectors extending over significant portions of the site surveyed in Area N).

Borings - Area N.

The locations of the three boring locations in Area N that were inspected by radar are listed in Table II. The three sites were all satisfactory, and none of the borings were relocated.

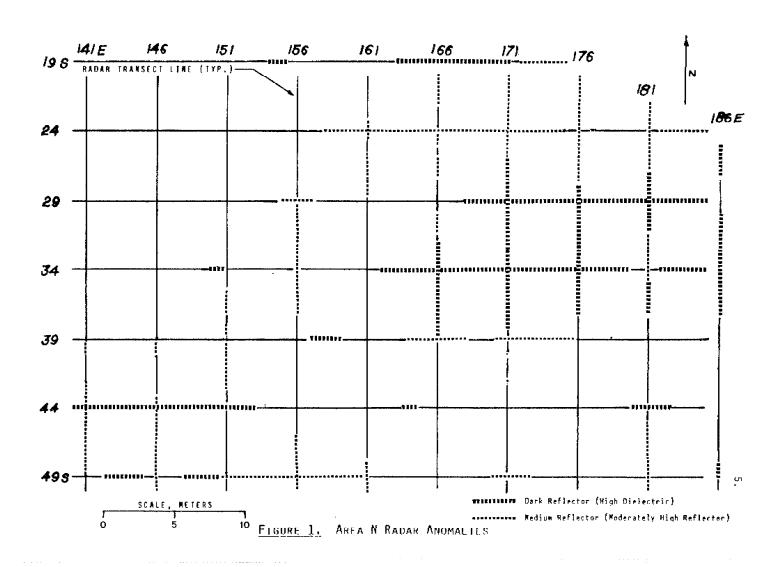


TABLE I
BORING LOCATIONS DETERMINED BY RADAR

AREA Q

Boring Number	Direction of Relocation	Proposed Location	Final Location	Notes
Q1	Move 2m North	1502S, 103E	1500S, 103E	N-S Pipe @ 107.5E
Q2	•	1504S, 112E	1504S, 112E	,
Q3	Move 2m West	1503S, 133E	1503\$, 131E	
Q4	Move 2m North	3501S, 22E	1499S, 22E	E-W Pipe @ 1502E
Q5	Move 3m North	1507S, 25E	1504\$, 25E	
Q6	-	1503S, 31E	1503S, 31E	
Q7	-	Storage Yard [†]	Storage Yard [†]	Gravel area
Q8	-	1340\$, 48E	7340S, 48E	
Q9	Move 3m East	1266S, 176E	7266S, 179E	N-S Pipe @ 1755
Q10	-	1222S, 155E	1222S, 155E	
QTI	Move 2m South	1167S, 159E	1169S, 159E	
Q12	Move lm West	1080S, 28E	1080\$, 27ā	
Q13	-	744S, 30E	744S, 30E	
Q14	Move 1m East	334S, 20E	334S, 21E	
Q15	-	569\$, 200E	569S, 200E	
216	-	544S, 440E	44S, 440E	
Q17	-	320S, 526E	320S, 526E	
819	-	330S, 760E	330S, 760E	
019	-	805\$, 175E	805S, 175E	Dark anomaly zone
020	Move 1m North	805S. 770F	804S, 170E	·
Q21	-	805S, 165E	805S, 165E	
Q22	-	820S, 175E	820S, 75E	
Q23	Move 2m South	820S, 170E	822S, 170E	
Q24	Move 2m North	820S, 165E	818S, 165E	

^{*}Town of Lewiston Public Works storage yard; 26.2m south, 8.8m west of wooden light pole.

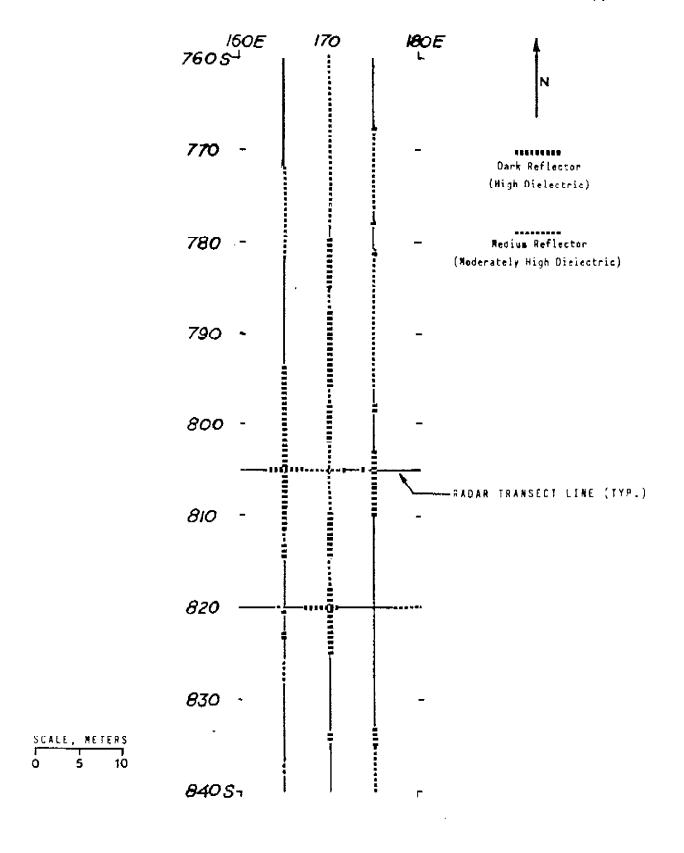


FIGURE 2. AREA Q RADAR ANOMALIES

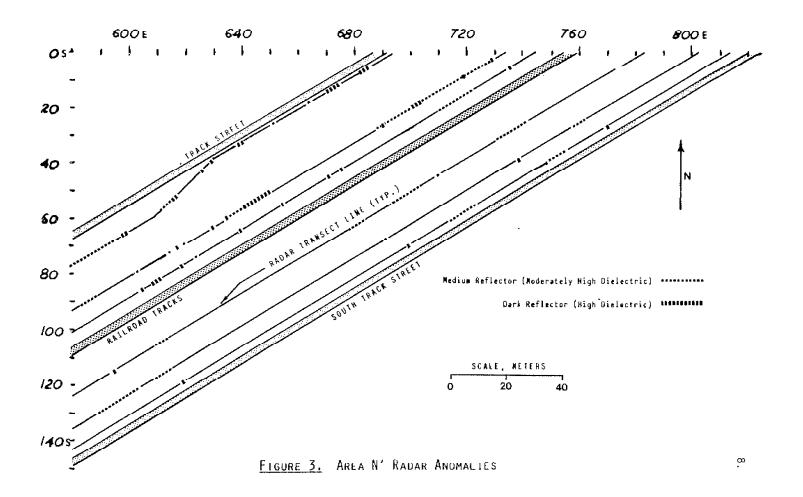


TABLE II

BORING LOCATIONS DETERMINED BY RADAR

AREA N

Boring Number	Direction of Relocation	Proposed. Location	Final Location	<u>Notes</u>
NI	-	27ØS, 59 Ø E	270S, 590E	
N2	-	40S, 102E	40S, 102E	
N3	-	15ØS, 4Ø4E	150S, 404E	

APPENDIX

Liquid Contaminants
Field Charts

LIQUID CONTAMINANTS

The presence of liquid contaminants in the ground creates distinct radar signatures according to the electrical properties of the liquid. Ionic liquids are electrically active due to the ions having electrical charge. Examples of ionic liquids are acids and salts. The presence of an ionic liquid in the ground lowers the electrical resistance of the ground, an effect that is observed by the radar system as a lighter-than-normal contrast.

Non-ionic liquids in the ground also modify the electrical characteristics of the ground, but in a different way: non-ionic liquids increase the dielectric constant of the ground, an effect that is observable by the radar system as a darker-than-normal contrast.

Fresh water is a special case of a non-ionic liquid. For reasons explained in the following paragraphs, water is usually distinguishable from other non-ionic liquids in the ground.

Ionic Liquids

As the radar signal travels into the ground, the energy is attenuated in proportion to the electrical conductivity of the ground. Normally, the ground has relatively high resistivity (the inverse of conductivity). The presence of electrically-active liquids will lower the electrical resistivity of the ground, resulting in more rapid loss of the radar signal. This snows in the radar record as an area of lighter-than-normal contrast. Experience with the radar system allows us to make close estimates of the resistivity values. In effect, the radar can be used as an imaging resistivity meter because of its extreme sensitivity to this parameter. Vertical and horizontal boundaries which only show as broad trend lines with conventional resistivity probes become sharply delineated on the radar record, thus allowing the boundaries of a contamination zone to be established with high accuracy.

Non-ionic Liquids

Non-ionic liquids such as petroleum-based solvents, oils, or other non-electrically active substances do not alter the electrical conductivity. Instead, non-ionic liquids tend to increase the dielectric constant of the ground. Higher dielectric values affect the coefficient of reflection according to the following equation:

$$o = \frac{\sqrt{\varepsilon_1} - \sqrt{\varepsilon_2}}{\sqrt{\varepsilon_1} + \sqrt{\varepsilon_2}}$$

where p - coefficient of reflection

 ϵ_1 = dielectric constant, layer l

ε₂ = dielectric constant, layer 2

Layer 1 is taken as the first layer through which the radar energy is passing, and layer 2 is the second layer. The reflection at the interface is dependent on the contrast, or relative difference, between the dielectric constants of the two layers.

When using this equation, it is convenient to use the relative dielectric constant normalized to air, which has a relative dielectric constant of unity. On this scale, various types of soils all have relative dielectric constants that are clustered in a narrow range between 3 and 5. (The numbers are dimensionless, being relative values.) These values are for dry conditions. The addition of moisture or other liquids greatly alters these values. For example, average moisture conditions in the ground result in a relative dielectric constant around 12. Wet, saturated ground can have relative dielectric values of 30 to 35. Peat, an extreme example of water-laden material, exhibits relative dielectric constants of 58 to 65. The addition of water can change the dielectric constant by more than an order of magnitude. The variability among soil types is less than a factor of two. Thus, the radar reflections, which depend on the contrast, or difference, between reflecting layers, is very sensitive to the presence of water.

A similar situation holds true for other liquids whose presence in the ground modify the dielectric constants. The difference between water and other liquids is the ease with which water disperses in the ground compared to oily or more viscous liquids. Also, the moisture content of the ground tends to reach steady-state, equilibrium conditions a foot or two below the surface. What this does is to create a general moisture "background" that tends to be constant over extended areas as well a exhibiting a constant vertical profile beginning near the surface.

Localized wet spots produce a greater contrast in the radar record, and are often observable at the surface as a low spot in the ground. What is important is that the wet spot retains its structural integrity on the radar record. Wet glacial till still looks like ordinary glacial till, soil horizons observed outside a wet area are still preserved in the wet area, etc. In the case of oil, sludge, and liquid industrial waste, the structural appearance of the ground is typically altered, because these materials do not disperse like water, and therefore tend to show localized alterations in the overall radar appearance of the ground. Another phenomenon is observed when liquids other than water are in the ground: the radar signal often resonates, or "rings", producing a series of identical waves on the radar record. This is most likely due to layering effects, where the depth of a liquid-laden layer produces a resonant cavity effect, resulting in multiple echoes from the same reflecting layer. Agair, because of the dispersive nature of water, such resonant cavities are almost never observed in wet, saturated ground.

These observations may be summed as follows: wet areas increase the radar contrast, but do not alter the structural appearance of the ground. Other liquids which do not have the dispersive properties of water also increase the contrast, but tend to alter the structural appearance of the ground. Finally, the observation of resonant signatures is most often associated with liquids other than water, and is seldom observed when water alone is present.

(617) 369 - 7999

RADARVISION

TAPE NO.: 8245

LOCATION: LEWISTON NY

CLIENT: ORAV

DATE: 10/25/82

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RADARVISION

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CLIENT: ORAU

DATE: 10/20/62

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(617) 369-7999

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APPENDIX D

EVALUATION OF RADIATION EXPOSURES ON OFF-SITE PROPERTY N/N' SOUTH NIAGARA FALLS STORAGE SITE LEWISTON, NEW YORK

C XICHAPPA

Evaluation of Radiation Exposures on Off-Site Property N/N' South Niagara Falls Storage Site Lewiston, New York

INTRODUCTION

The U.S. Department of Energy has completed a radiological survey and determined that portions of the Department of Labor property, Lewiston, New York, are contaminated with low-level radioactive residues resulting from previous uses This property was part of the former Lake Ontario Ordnance of this property. Works, now known as the Niagara Falls Storage Site (NFSS) and associated off-site properties, where radioactive wastes from Manhattan Engineer District and Atomic Energy Commission operations were handled and stored. These wastes were primarily residues from uranium processing operations. However, they also included contaminated rubble and scrap from decommissioned facilities, biological and miscellaneous wastes from the University of Rochester, and low-level fission product waste from contaminated-liquid evaporators at the Knolls Atomic Power Laboratory (KAPL) in Schenectady, New York. Receipt of additional wastes was discontinued at the site in 1954. Although some storage of radioactive materials on the Niagara Falls Storage Site continues under the control of the Department of Energy, work involving handling of radioactive waste has not been performed on the off-site properties for approximately 25 years.

In 1954 a preliminary cleanup of the site was performed by Hooker Chemical Company. Approximately 1298 acres of the original 1511 acre site were then declared excess and eventually sold by the General Services Administration to various private, commercial, and governmental agencies. The Department of Labor is the current owner of a 76.6 acre tract, identified as off-site property N/N' South. This property is not occupied or in use.

The property was surveyed by Oak Ridge Associated Universities, Oak Ridge, Tennessee, during October and November 1982, and March 1983, and found to contain radioactive contamination. The survey indicated radionuclides from the naturally occurring uranium and actinium decay series and small quantities of Cs-137.

Cosium-137 is a man-made radionuclide created through the fission process such as in a nuclear reactor. Cosium-137 has a half-life* of approximately 30 years and emits beta and gamma radiation. The naturally occurring decay series, known as the uranium and actinium series, are believed to have been created when the earth was formed, and they are still present today because of their very long half-lives. These series are presented in Tables D-1 and D-2.

As a radionuclide decays it changes into another substance. In the case of U-238, for example, the decay produces Th-234. Thorium-234 is called the "daughter" of U-238, U-238 is the "parent" of Th-234. In turn, Th-234 is the "parent" of Pa-234. Radioactive decay started by U-238, U-235, or Th-232 continues as shown in the tables until a stable nuclide is formed.

The radionuclides in these decay series are present in small quantities throughout the environment. Concentrations of them normally occur in soil, air, water, food, etc., and are referred to as background concentrations. Radiation exposures resulting from this environmental radioactivity are referred to as background exposures. These background exposures are not caused by any human activity, and to a large extent, can be controlled only through man's moving to areas with lower background exposures. Each and every human receives some background exposure daily.

The use of radioactive materials for scientific, industrial, or medical purposes may cause radiation exposures above the background level to be received by workers in the industry, and to a lesser extent, by members of the general public. Scientifically based guidelines have been developed to place an upper limit on these additional exposures. Limits established for exposures to the general public are much lower than the limits established for workers in the nuclear industry.

RADIATION LEVELS ON PROPERTY N/N' SOUTH

The survey identified elevated levels of cirect radiation and contamination of the soil above the normal background levels. The major radionuclides noted in these soils are Ra-226, U-238, and Cs-137. Increased levels of radioactivity

^{*} The half-life is the time required for half of the atome of a radioactive substance to disintegrate ("decay" or transform).

resulting from contaminated residues on this property can cause an increased radiation exposure to persons. The exposure potentially comes from one primary source or pathway: direct radiation emitted by the radionuclides in the residue or soil. Additional exposures may also be received through inhalation of suspended airborne particulates and radon gas and its daughter products* and ingestion of contaminated food or water. In Table D-3 the exposure levels associated with this property are summarized and compared with the guidelines and background levels.

External Radiation Exposure Levels

As Tables D-1 and D-2 indicate, several members of the naturally occurring decay series emit gamma radiation as does Cs-137. (Gamma rays are penetrating radiation like x-rays.) Contaminated areas can, therefore, be sources of external gamma radiation exposure.

The National Council on Radiation Protection and Measurements has recommended maximum annual whole-body exposure of 500,000 microroentgens** per year to an individual exposed in the general population. This is equivalent to a continuous level of approximately 57 microroentgens per hour. The maximum exposure level noted on this property was 550 microroentgens per hour. This level was present only at contact with one small isolated area of contaminated surface soil. average exposure rate of 12 microroentgens per hour at about 3 feet above the surface is a better estimate of the average exposure an individual might receive. For comparison, the average background level in the Lewiston area is about 8 microroentgens per hour, and continuous exposure at this level would produce an annual exposure of about 69,800 microroentgens. Also, a typical chest x-ray (according to data from the Department of Health and Human Services) might yield an exposure of about 27,000 microroentgens.

The soil is contaminated with radium and lesser levels of uranium and cesium which emit beta and gamma radiations. Nuclear Regulatory Commission (NRC) guidelines for decommissioning former nuclear facilities require that the dose

^{*} Radon-222 is a jas that results from the decay of radium-220, a member of the naturally occurring uranium series (see Table D-1).

A* The Roentgen is the unit of exposure to X- or gamma radiation. A microroentyen is one-millionth of a Roentgen.

rate (from beta and gamma radiation) measured at a distance of one centimeter above the surface does not exceed 1.0 millirad* per hour and 0.2 millirad per hour average. The maximum dose rate measured at this site was 3.8 millirad per hour and the average was 0.08 millirad per hour. Although the maximum level exceeds NRC guidelines, the primary concern of this guideline is exposure of skin surface. The thickness of ordinary shoe soles is adequate to protect the skin of feet from beta radiation. In most cases, exposures are negligible at a distance of 1 ft away from the surface and areas of body skin are adequately protected from these exposures if they remain away from these surfaces. Beta radiation from surface residues are therefore not a significant factor in evaluating the potential health effects at this site.

Exposure from Inhalation of Airborne Radioactive Particulates

A very small amount of the radioactive contamination on this property may become airborne by resuspension of particulates from the surface layer of soil. The actual fraction of material that becomes resuspended is dependent on a number of factors including surface conditions (e.g. damp, dry, covered by ground vegetation, etc.), particle size, activities on the site which disturb the surface soil, and micrometeorological conditions (e.g. surface wind speed and direction). Determining average levels of airborne radionuclides requires air sampling over an extended time period and was beyond the scope of the ORAU survey. However, an estimate of the potential airborne concentrations can be made based on the average concentration of radioactive material in the surface soil and using standard computation procedures of the Nuclear Regulatory Commission.

Areas of significantly higher surface contamination levels are isolated and small (usually less than 6 inches in diameter). The average surface soil concentration for property N/N' South is therefore best approximated by the samples collected at the grid line intersections. Radium-226 is the major radionuclide of concern on this site and the average concentration in surface soil is 1.34 picocuries** per gram or about 0.64 picocuries per gram above the level normally present in surface soils in the Lewiston area. The resulting

^{*} The rad is the unit of beta-gamma dose. A millirad is one-thousandth of a rad.

** The curie is the unit indicating the quantity of a radioactive substance. A picocurie is one-millionth-millionth of a curie.

concentration of resuspended Ra-226, based on a resuspension factor of 5 x 10^{-9} per meter, would be about 1 x 10^{-16} microcuries per cubic contineters of air. For comparison, the Nuclear Regulatory Commission's guideline level for continuous exposure of the general public is 2 x 10^{-12} microcuries per cubic centimeter. The estimated concentration of airborne Ra-226 is almost a factor of 20,000 less than the guidance level and would therefore not result in a significant increase in radiation exposure to individuals on this property.

Exposure From Inhalation of Radon in Air

The deposits of radium bearing residues in soil may be indirect sources of radiation exposure on site. As shown in Table D-1, Ra-226 changes to Rn-222 as a result of radioactive decay. Radon-222 is an inert gas which can emanate from the ground and, with its daughter products, result in lung exposures. Radon levels in the vicinity of the NFSS are continuously monitored by Department of Energy contractors. Sampling near property N/N' South indicted an average radon concentration of approximately 0.4 picocuries per liter of air from January 1981 through June 1981*. The guideline for continuous exposure of the general public is 3 picocuries per liter. For comparison the average area background level in the Lewiston area during 1979 and 1980 was 0.23 picocuries per liter. This source is not considered significant.

Other Exposure Considerations

Loose radioactive contamination can result in exposure through ingestion (eating or drinking) of contaminated foodstuffs. This site is not used for raising crops and average radionuclide concentrations in the ground water at this site are within the EFA drinking water limits. These pathways would not, therefore, result in significant exposures.

ESTIMATES OF HEALTH EFFECTS

The primary health effect associated with radiation exposure is an increased risk of cancer. In general, the risk is assumed to increase as the total dose of radiation increases. Total dose is dependent not only on exposure rate and

^{*} Data from National Lead Co. of Ohio, <u>Review of Radon Monitoring at the Niagara</u>
Falls Storage <u>Site</u>, Cincinnati, OH, March 31, 1980.

concentration levels on the property, but also on the nature and duration of the exposure. In addition, a given incividual's increased risk is dependent upon many factors including the individual's age at onset of exposure, variability in latency period (time between exposure and physical evidence of disease), the individual's personal habits and state of health, previous or concurrent exposure to other hazardous agents, and the individual's family medical history. Because of these variables, large uncertainties would exist in any estimates of the number of increased cancers in a relatively small exposed population such as might be the situation on this site. Estimates of the increased risks have been calculated and are given in Table D-4. Assumptions made in performing these calculations are:

- 1. The levels reported in Table D-3 are representative of the conditions and will not change during the year or from year to year.
- 2. Averge exposure levels in Table D-3 are representative of the averages to which an individual working on the property might be exposed.
- 3. An individual would spend a working lifetime, i.e. 40 hours per week, 50 weeks per year, for 45 years (age 20 to 65) on the site.
- 4. Background exposure rates to individuals while not on the property will be 8 microroentgens per hour from external gamma radiation.

The risk estimates are based on the 1980 National Academy of Sciences report, "The Effects on Populations of Exposure to Low Levels of Ionizing Radiation," and the 1977 report by the United States Scientific Committee on Effects of Atomic Radiation. The lifetime risk estimate used to calculate the values in Table D-4 is 100 cancer deaths per million persons exposed per rem of radiation exposure. It is believed by many radiation biologists that with low dose rates such as those encountered at this property, the actual risks of cancer are much less than 100 per million persons per rem, zero not being excluded.

The Ra-226 and radon air concentrations estimated or measured for this property are a small fraction of the guidance level and exposures and risk from these pathways would be negligible. In addition, exposures and risk from the pathways of ingestion of crops grown on contaminated soils and water containing

radionuclides from the soil are also considered negligible, based on the low-levels and use of this property. Exposures and risk are therefore limited to one pathway — direct exposure to gamma radiation.

The estimated increased risk due to cancer from exposure to the average radiation levels on property N/N' South, for a working lifetime is 0.036 per 1000 deaths. This can be compared with the average lifetime risks of cancer in Niagara County of 218 per 1000 deaths based on 1977 crude death rate statistics for this same year. The average lifetime risks of cancer in the State of New York and the United States are 216 per 1000 deaths and 203 per 1000 deaths, respectively. An individual working under the assumed conditions will therefore be subject to an increased risk of dying from cancer of 0.0036 percent or an increase in total risk from 21.8 to 21.8036 percent when compared to the average risk in Niagara County. This may also be expressed as a percent increase in overall risk of getting a fatal cancer of 0.0165 percent — a negligible increase.

SUMMARY

In summary, portions of Niagara Falls Storage Site off-site property N/N' South are contaminated with low-level residues containing Cs-137 and naturally occurring radionuclides. The level of Ra-226 contamination in the surface soil in some areas of the property exceeds the present criterion for release of this property for unrestricted use. Although under current conditions of property use this contamination is capable of producing slight radiation exposures to persons on the property, these exposures are well within the scientifically based guidelines and risks to such persons are negligible.

TABLE D-1
URANIUM DECAY SERIES

Parent	Half-Life	Major Decay Products	Daughter
Uranium-238	4.5 billion years	alpha	Thorium-234
Thorium-234	24 days	beta, gamma	Protactinium-234
Protactinium-234	1.2 minutes	beta, gamma	Uranium-234
Uranium−234	250,000 years	alpha	Thorium-230
Thorium-230	80,000 years	alpha	Radium-226
Radium-226	1,600 years	alpha	Radon-222
Radon-222	3.8 days	alpha	Polonium-218
Polonium-218	3 minutes	alpha	Lead-214
Lead-214	27 minutes	beta, gamma	Bismuth-214
Bismuth-214	20 minutes	beta, gamma	Polonium-214
Polonium-214	.0002 seconds	alpha	Lead-210
Lead-210	22 years	beta	Bismuth-210
Bismuth-210	5 days	beta	Polonium-210
Polonium-210	140 days	alpha	Lead-206
Lead-206	stable	none	none

TABLE D-2
ACTINIUM DECAY SERIES

Parent	Half-Life	Decay Products	Daughter
Jranium-235	710 million years	alpha	Thorium-231
Tnorium-231	25.5 hours	beta	Protactinium-231
Protactinium-231	32,000 years	alpha	Actinium-227
Actinium-227	21.6 years	beta, gamma	Thorium-227
Thorium-227	i8.2 days	alpha	Radium-223
Kadium-223	11.4 days	alpha	Radon-219
Radon-219	4.0 seconds	alpha	Polonium-215
Polonium-215	.0018 seconds	alpha	Lead-211
Lead-211	36.1 minutes	beta, gamma	Bismuth-211
Bismuth-211	2.15 minutes	alpha	Thallium-207
Thallium-207	4.79 minutes	beta	Lead-207

TABLE 0-3 SIMMARY OF EXPOSURE LEVELS ON SHOPERTY NAM' SOLIEN LEWISTON, NEW YORK

	Levels on Site				Guidelines for
Exposure Source	Average	Maximum	Background Levels	Guldelines for General Pubilc	Radlation Workers
Gamma Radiallon from Cesium-137 and uranium and actinium decay	12 μR/h ^a	550µR/ո	я _Џ R/ћ	0.5 rem ^b per year for Individual, equivalent to 250 kg/h above natural background for 40 h/wk and continuous exposure.	5 rems per year
Radionuclides (Radium÷226) in air	1 × 10 ⁻¹⁶ µCi/cc ^c	******	unknown	$2\times10^{-12}~\mu\text{GH/cc}$ for continuous (168 h/wk) exposure	$5 \times 10^{-11} \ \mu \text{Ci/cc}$ for 40 h/wk exposure
Radon in air	0.4 pCi/I	, ·-	U.23 pCi/i (1979 & 1980 avg.)	5 pCi/I	30 gCi∕l for 40 h/wk exposure
Radionuclides (gross alpha concentration) in Ground Water	5,9 pCi/(13.9 pCI/I	Appr. 0.8 pCi/l	15 pCi/i, EPA Standard for Public Drinking Water Systems	400 pC1/I
Radionuclides in Soft: ^d					
Radiuπ-126	1.3 pCi/q	431 pCi/g	Appr. 0.7 pC1/q	EPA Mill Tallines (Criteria is 5 pC1/g above background averaged over 100 m2 of surface soil.)	none
Cesiuπ−137	0.5 pSi/g	6.5 pCi∕y	Appr. 0.5 pCi/g	80 pCl/g above background (Criteria developed by Los Alamos sci. Lab. for cleanup at sites contaminated by fission product residues.	none

TABLE D-3, cont.

SUMMARY OF EXPOSURE LEVELS ON PROPERTY NAME SOUTH LEWISTON, NEW YORK

	Levels on Site				Guidellnes for
Exposure Source	Average	Maximum	Background Levels	Guldelines for General Public	Radiation Workers
Vranium−238	4.4 pCi/g	125 p01/g	Appr. 5.5 pCl/g	40 pCI/g Interim Soil timits for 0 A D Projects, LA-UR-79-1865-Rev, J.W. Healy, et al.	none
Uranluπ−235	0.3 pCi∕g	9 .1 pCI/g	Appr. 0.2 pCi/g	30 pCi/g (U enriched in U-235) NRC Branch Tochnical Paper (Federal Register, October 23, 1981)	none

^a The Roomigen (R) is a unit which was defined for radiation protection purposes for people exposed to penetrating

gamma radiation. A microroentgen (μ R) is one millionth of a Roentgen.

b The rem is the unit of lonizing radiation that produces the same biological damage in man as an absorbed dose of I rountgon of high voltage x-ray. A rountgen of gamma exposure to a non-is equivalent to one ϵ and

I rountgen of high voltage x-ray. A rountgen of gamma exposure to a non-is equivation, to one can.

The microcurie (μ Ci) and placeurie (pCi) are units which are defined for expressing the amount of radioactivity present in a substance. I μ Ci = 10^{-6} Ci, I pCi = 10^{-12} Ci,

d For soil concentrations, only the systematically collected soils were used in determining the average level.

TABLE D-4

SUMMARY OF WORKING LIFETIME RADIATION
EXPOSURES AND ESTIMATES OF ASSOCIATED CANCER RISK
FOR PROPERTY N/N' SOUTH, LEWISTON, NEW YORK

Exposure	Equivalent Corrected for Background	Increased Risk Due to All Cancers	
xternal gamma radiation	0.36 rems	0.036 per 1000ª	
nhalation of resuspende particulates	d negligible	0	
nhalation of radon	negligible	0	
ngestion of food and water contaminated by radioactive materials on-site	negligible	O	
OTAL	0.36 rems	0.036 per 1000 ^b	

 $^{^{\}rm a}$ Using the risk coefficient of 100 cancer deaths/10 $^{\rm 6}$ person rem. This is approximately a mean value from BEIR-III (1980) and UNSCEAR (1977).

b The average lifetime risk of death due to cancer in the United States is 203 per 1000 (20.3 percent); in Niagara County the average lifetime risk is 218 per 1000 (21.8 percent).