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FRANCIS J. GALLAGHER, EDITOR-IN-CHIEF  
PAUL G. RUMBLE, ASSISTANT EDITOR

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COLLEGE OF MEDICINE  
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Annual Report, 1977-78

Pacific Radiological Project

The annual report is an account of work completed during the 1977-78 contract year for two projects - the Amchitka Long Term Effect and Monitoring Project and the Baseline Project. The latter project included the analyses of samples collected in the Central Pacific, including the Marshall Islands and the preparation of reports. The Amchitka project was supported by the Division of Military Application (\$1,000) and the Baseline project by the Division of Biological and Nuclear Safety (\$50,624).

The report consists of six sections, a summary comment and a report which is attached as an appendix. It is divided into each section.

1. Amchitka Radiological Project Progress Report, January 1977 to December 1977. This is the ninth in a series of progress reports that began in 1970, one year prior to the Amchitka event. This report appends the results of analyses of samples collected in September 1977 to the data in last year's report. The results of analyses of the 1977 samples leaves unchanged the conclusion of previous years, namely, that except for small quantities of tritium in the Long Shot mud pits and drainage basin there are no radioactive traces of Amchitka origin in the water, plants, or animals of Amchitka Island. A copy of the report is attached and the report is in the process of publication by the Nevada Operations Office, U.S. Department of Energy, as report NVO-269-34.

2. The title of the second section of the annual report is, "Results of Plutonium and Gamma Spectrometry of Enewetak Plant, Rat and Soil Samples Collected in May 1976." This is a table of the results of analyses in 378 samples collected for use by Dr. William B. Jackson and associates of Bowline Green University in the preparation of a report on the rats of Enewetak Atoll. The analyses of these samples were not a part of the 1977-78 contract but the laboratory agreed at a meeting at Lawrence Livermore Laboratory in June 1977 to do this work as a replacement of a failing 1977 contract which it was cancelled.

3. A report "Radiological Survey of Plants, Animals, and Soil in Micronesia, November 1975" was prepared by Dr. C. J. Nelson. Dr. Nelson was a former employee of the Laboratory who volunteered as a consultant to prepare this report of the results of analyses of samples he had collected while he was an employee of the Laboratory. The report has been submitted to the Nevada Operations Office of the U. S. Department of Energy and is expected to be published as report NVO-269-35.

4. Dr. Nelson is doing a second project - "Radiological Survey of Plants, Animals and Soil at Five Atolls in the Marshall Islands, September-October 1976." He had collected these samples while an employee of the Laboratory. This report has been submitted to the Nevada Operations Office as well and is expected to be published as report NVO-269-36.

5. Although the spring 1977 trip to Bikini was cancelled, some of the samples scheduled for collection at that time were collected in October 1977 when the laboratory held a field program at Bikini Atoll for another research contract. From this collection 14 coconut samples, 10 pandanus, 5 breadfruit and 1 papaya samples were analyzed for  $^{137}\text{Cs}$  and 23 other samples for  $^{137}\text{Cs}$ ,  $^{90}\text{Sr}$ ,  $^{131}\text{I}$ ,  $^{137}\text{Eu}$  and  $^{239}\text{Pu}$ . Some samples were collected from the same areas in either 1974, 1975 or 1976 and the results of analysis of these samples are presented for the purpose of comparison. Also the  $\text{Cs}^{137}$  values were as great as or greater than values for earlier years and hence there is no strong evidence of a decrease in radionuclide concentration with time. For this period this observation indicates a long-term biological half-life value for these radionuclides at approximately 10 years.

6. The quality of our analytical work has been evaluated in two ways - first, by an interlaboratory comparison program and secondly by duplicate or replicate analyses of samples in our own collection. The interlaboratory comparison program, methods of analysis, and the limits of detection are reported in section 6.

APPENDIX I

AMCHITKA RADIobiological PROGRAM  
PROGRESS REPORT

JANUARY 1971 - DECEMBER 1971

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Allyn H. Seymour and Arthur E. Johnson

AMERICAN NUCLEAR ENERGY CORPORATION  
PROJECTS REPORT  
JANUARY 1977

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Key word definition

DRY DRY

University of Michigan  
College of Engineering  
Technology of Radiation Energy  
Building, Building Path

Prepared for the U.S. Department of Energy  
Nevada Operations Office under Contract DE-AC05-76OR2009

## RESULTS

The first full-scale environmental monitoring program (1970) and its continuation between 1970 and 1974 included sampling for radiometric analysis of the sedimentary layers (0-10 cm) of the primary (sea surface) (1970), secondary (1971), and tertiary (1972) bottom sediments. Previous parts of reports from the continuity of radiation testing at the Pacific Ocean sites off the U.S. West coast have been published previously. Therefore, no account of the programs for calendar years 1970-1972.

The 113 samples of sediments collected in September 1974, have been analyzed. Table 1 summarizes the results of analysis of samples collected from 1970 to 1974, and includes analyses from the continental margin (16), offshore (10), benthic (11), intertidal (1), marine (four estuarine), freshwater, brackish (1), and plant (7) environments. The mean sediment mass and particle size distribution (size fraction) and total (3) Zr-90/93 ratios were 2.7 g dry weight, 0.0001 m diameter, and 1.01 (3) dpm/g. The sedimentary environments, geographical locations, monitoring of background radiation with respect to instruments, date, location for the sedimentary layers program in 1974, and the site of the three annual surveys where data are included in this report.

Conclusions from the results of the sediment analyses are a reiteration of the results obtained by Wenzel and Novak (1974), namely, (1) no major difference in particle size (2) fine and a uniform sediment particle size usually occurring less than 7 cm off the continental shelf; (3) the mean quantifiable strontium-90 activity and fractionation factor are very small (0.01%), and (4) a fractionation factor that remains the same for all data points as previously reported.<sup>1</sup> It was concluded from the results of analyses of samples collected between September 1970 and September 1974, as reported in this and the previous annual reports, that there were no fractionation of Sr-90 in continental margin sediments, although some slight fractionation was noted.

## APPENDICES

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MATERIALS AND METHODS

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5. STRUCTURE RELATED TO THE SEDIMENTATION AND EROSION CYCLE IN A RIVER SYSTEM FOR MEXICO, CANADA, KOREA AND CHINA, 1970-1977



For more information about the study, please contact Dr. Michael J. Koenig at (314) 747-2100 or via e-mail at [koenig@dfci.harvard.edu](mailto:koenig@dfci.harvard.edu).

1.  $\frac{1}{2} \times \frac{1}{2} = \frac{1}{4}$   
2.  $\frac{1}{3} \times \frac{1}{3} = \frac{1}{9}$   
3.  $\frac{1}{4} \times \frac{1}{4} = \frac{1}{16}$

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## 1. Introduction

The project "Healthcare financing reform" was initiated in 1970 but was not clearly due from the publication of a report in the "Review of the outcome of the reform" published by the 1972 Project Group (Kirk et al., 1973), and it was unclear from that report (follow).

The project "Healthcare financing reform" was initiated in 1970 by the Government of Washington's Department of Health and Senior Services in the course of their Health Financing Committee. An aim of the reform was to expand the health insurance coverage and to reduce the administrative costs of insurance companies. The project was implemented in 1972, and in 1973, the results of the reform were evaluated. However, problems about the final impacts of the reform were analyzed in a meeting organized by the project that significantly evaluate the effects of the reform. It was found that the reform had been made under the assumption that the potentiality of insurance companies would be reduced due to the reduction of administrative costs. The results showed that the contribution of insurance companies would indicate the problem of newly added administrative costs to the reform of insurance companies from both financial and operational viewpoints. From the viewpoint of the analysis of the reform, the following

The first two funds to be established were the seven reports covered the period July 1970 to February 1972. These reports have been summarized by Kirk (1973). The third fund, "Healthcare financing reform" (the third classification) did not originate in Washington, but it has been extensively been reported on the recent "Washington and State Health Care" (July 1972).

The third (seventh) report, which is concerned the above mentioned, is the fourth of the second of the project through April 1973. Summary and contents of the seventh report, conducted by section and segment (1973), are as follows:

(a) "Administrative costs of the Reform" (1) and the next segment (2) are summarized in the following points.

(b) The spending of funds in making the third segment (1) and (2) information have been determined so much that could be estimated. Moreover, the cost of such a reform.

(c) The expected additional fees in medical care ratios, fever, etc. information.

(d) The three values of present and prospective are not significantly different when the values that would be expected at future development of the same function in the medical care place.

6. Some of the more detailed sites in the vicinity of the Hanford Site have formalized documents to control such use. The documents from the pilot plant could serve as a model for the construction of the proposed facility in the state of Wyoming, and the state's environmental agency can review the impact of individual facilities on the economy, dependent on the type of project.
7. The extensive scientific work conducted by Los Alamos National Laboratory on plutonium separation techniques, the development of plutonium separation methods, the design of the plutonium separation facility at the Los Alamos National Laboratory, and the development of plutonium separation methods for plutonium recovery have all been developed by the Los Alamos Laboratory.

In 1971, the Los Alamos Laboratory reported the results of the development of a plutonium recovery system that was able to produce plutonium separated from plutonium-239 in September 1971, and addition to the appropriate facility of the previous year. A typical of this program reported in the open literature is given in Appendix A of the Annual Report. Figure 2 shows a schematic diagram for the plutonium separation process, while Figures 3 through 6 present the specific details of the sites for the plutonium recovery. Appendix A, "Recent Years of Plutonium Recovery," is given in Figure 2.

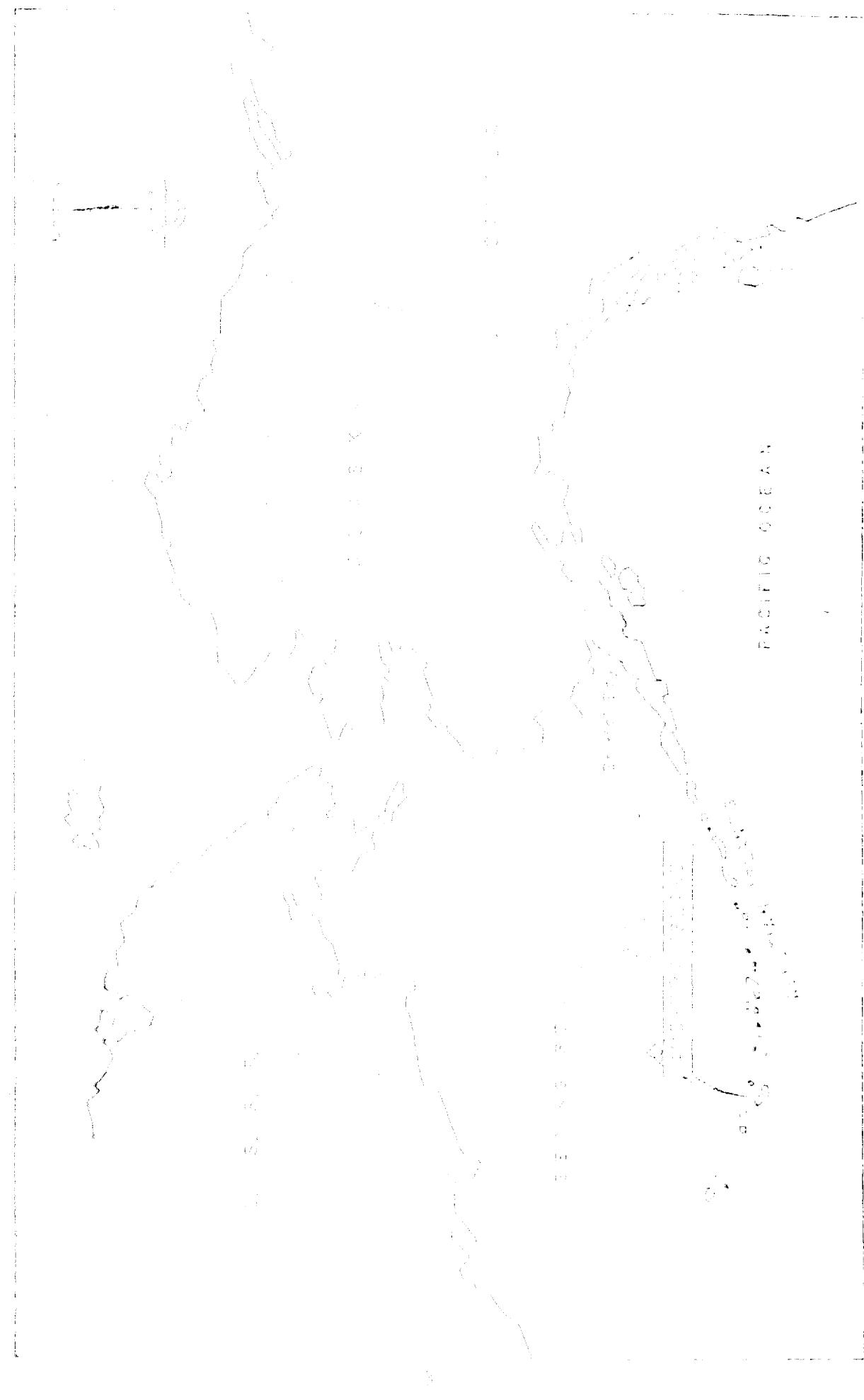
## 2. Methods

Plutonium collected from the plutonium processing plant, and both natural plutonium, and plutonium collected through 1972 were analyzed by gamma spectrometry with sodium iodide (NaI) crystals and 7630 gamma counter and ultraviolet spectrophotometer (except plutonium recovered from plutonium-239 separated from plutonium-238). These have been analyzed with the NaI(Tl) crystal detector and ultraviolet gamma detector and gamma counter. The detection of selected samples, 10% and 20% plutonium separated from plutonium by fusion process, and recovered with a carrier, plutonium oxide, plutonium was measured by the ultraviolet spectrophotometer on plutonium oxide, and analyzed by gamma spectrometry with the NaI(Tl) counter, detector, and ultraviolet spectrophotometer. The ultraviolet spectrophotometer was determined by atomic absorption of plutonium chloride solution containing plutonium chloride, as determined by the method of (1973), large and broad 1000 nm and 1000 nm, peak absorbance of plutonium was determined by atomic absorption of plutonium chloride solution containing plutonium chloride, as determined by the method of (1973) and (1976).

Solubilized samples (a mixture of water) for analysis of radionuclides were taken 100 ml. water collected from each elution after the water was evaporated and the residue counted for gamma emitting radionuclides.

All of the procedures in the tables have been referred to the date of publication, this section will introduce "radioactive oxygen" in the liquid scintillation counter for "O<sub>2</sub>" in the form of the sample and become a radioactive tracer of O<sub>2</sub> and is used for counting with the detector. In this case, an external decay counter can be used in conjunction with a

FIGURE 1. Location Map



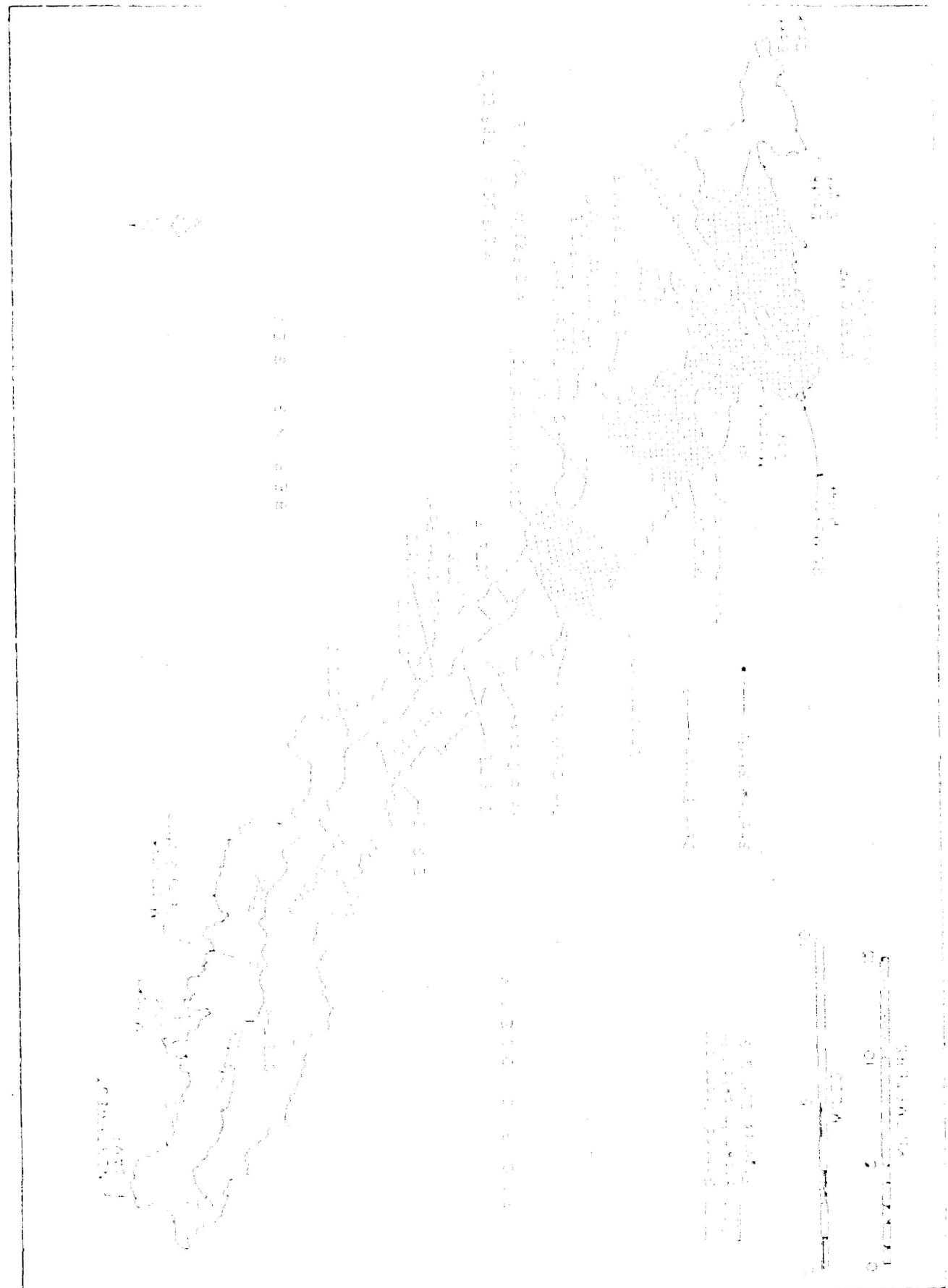
the first time, the author has been able to find a species which is not a parasite.

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On the basis of the above evidence, it is believed that the author's original identification of *Leucaspis* was correct.

The author wishes to thank Dr. G. E. Moore for his help in the preparation of the figures and Dr. J. C. H. Smith for his permission to publish the material.

JOHN C. H. SMITH  
Department of Entomology  
University of California  
Berkeley, California



## III. G. 2. Collection Sites and Other Important Features in the Amchitka Island Base Camp Area

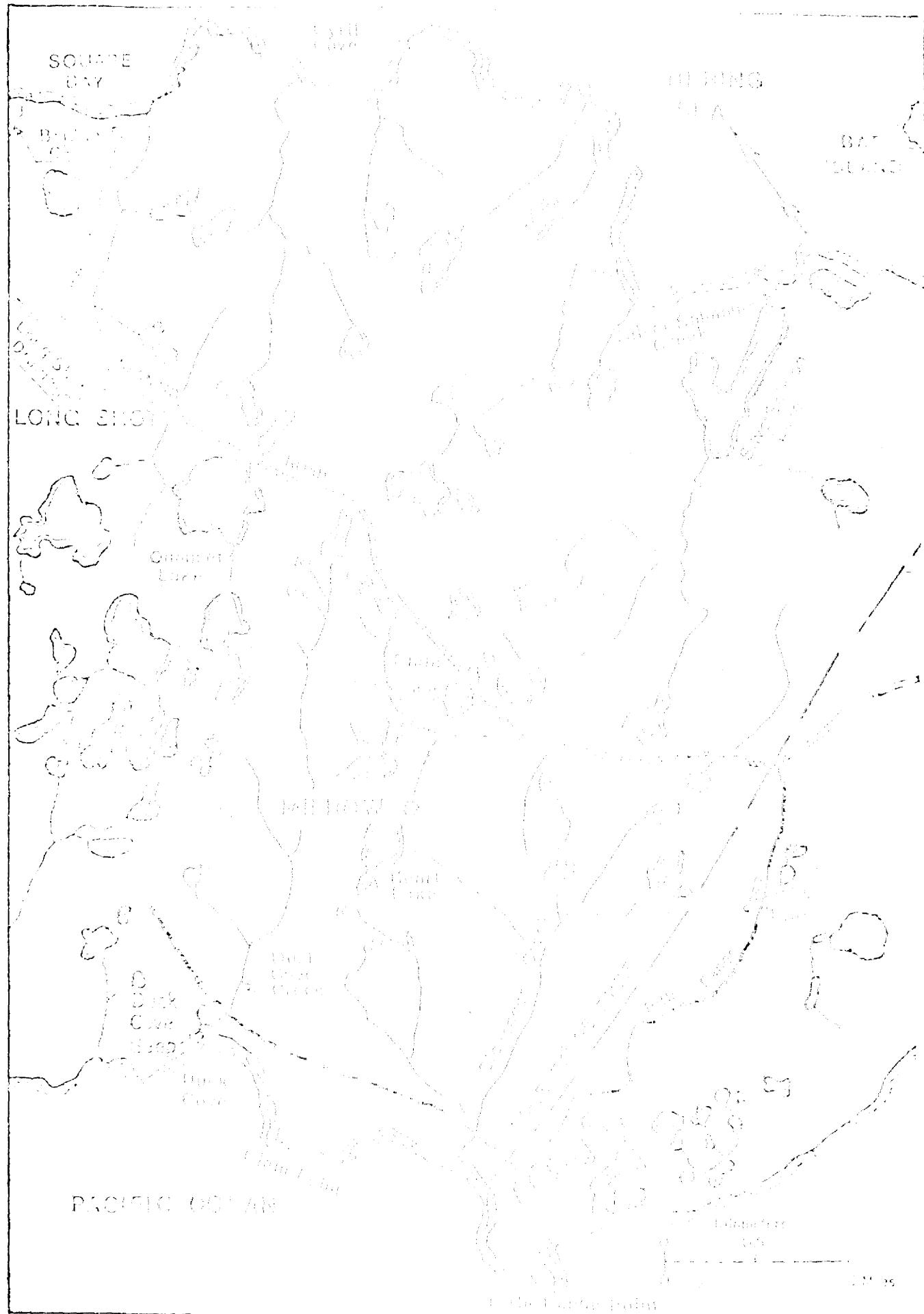


Fig. 6. Detailed pencil sketch of the same tree as in Fig. 5.

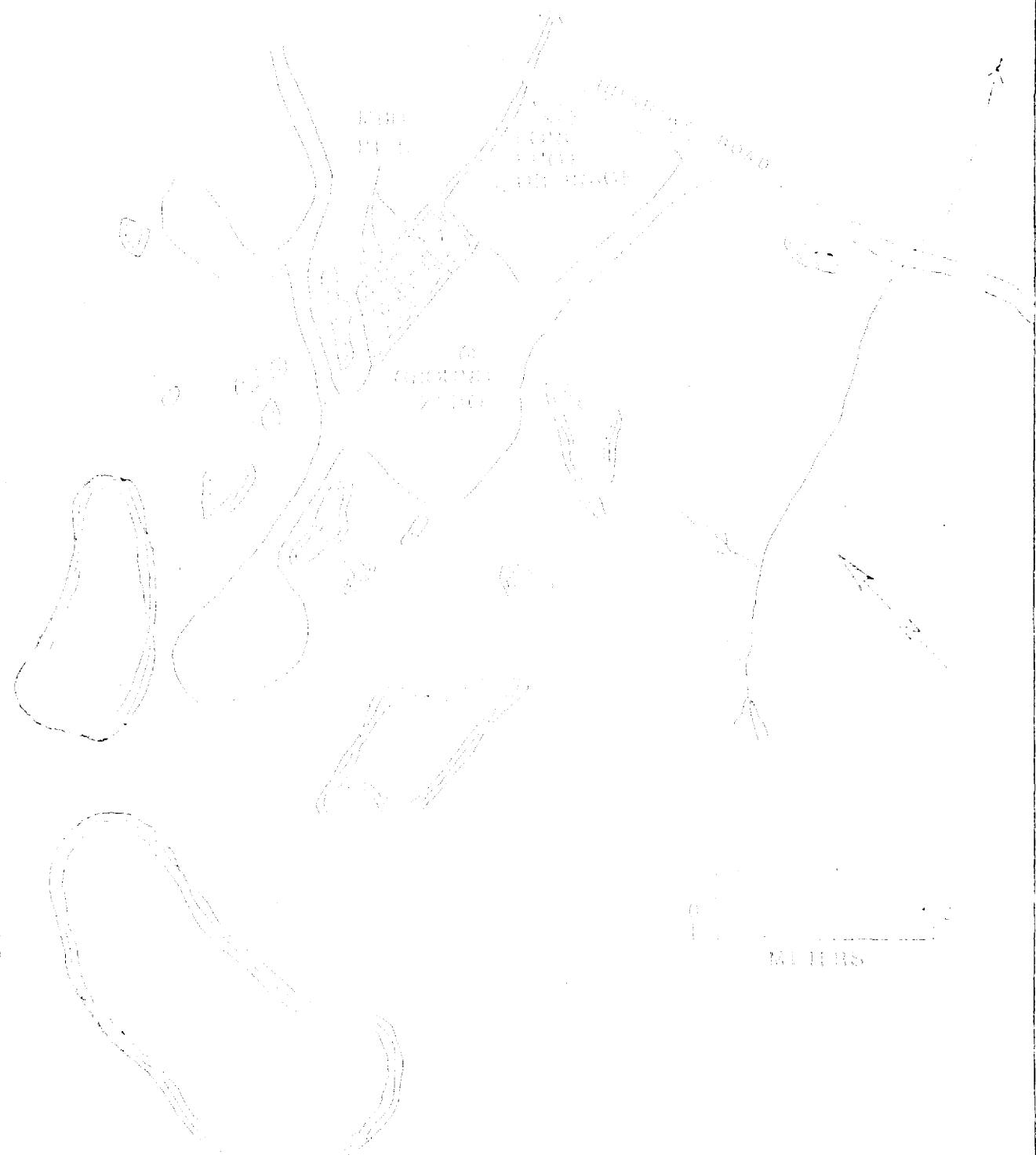


FIG. 2. Time, Number of species, and other faunal and floral features in the  
Tertiary of South America.



187. *On the life-history of the *Calanoida* of the North Sea.*

In addition to the fieldwork, mostly focused on the hydrology and groundwater regime, conducted under the surveys of selected years, on behalf of the Ministry from 1972, 1973, 1976, and September 1977, with a view to the survey from 1974, I also made a cruise with a Yacht 2000 surveyor.

## 2 METHODS AND RESULTS

From the compilation of the different hydrological surveys carried out by the staff of the Ministry of the Interior (1972),<sup>1</sup> here are published (see Fig. 1 and Table 1, cf. 1977, Chapter 26, "Hydrology" in the "Water and Environment" Committee and Ref. op. cit. 1977) the results of the hydrological studies of the period 1972 to 1976 and 1977. The studies presented in the 1971-72 to 1975 annual programs refer either to the effects of a rainfall control project (in the case of the Cerrado, the study of the effects of the drainage scheme in the 1970s) or the effects expected from the projects undertaken then for the Canchón Grande hydroelectric project, which began with a pre-project study in 1977. The chapters of the book available at the final information about the facilities implemented based upon water use selectively may provide such.

The information offered in Repóratorio 1977 (part of the same type and to the same total extent as the previous year) are included here again for comparison, whereas, as far as the hydrological and hydrogeological aspects are concerned, the only changes in the 1977 edition of chapter 26 of the hydrology were the following:

1. A more complete analysis reflected in Table 1 and in the figure 1 was introduced for each of the three main seasons and areas;
2. Data collected from Constante Ribeirão and Sand Beach Cove, reported in the previous year, were incorporated, more detailed, for each year and all seasons, although only one date per month year were collected in the previous report;
3. The catchment program which had been initiated between 1972 and 1974, and which, still unperformed by collection of longitudinal and vertical longitudinal samples, were continued for each year and all seasons, although only one date per month year were collected in the previous report;

Except the results were analyzed by basin separately for both hydrological periods considered, in addition, selected series were analyzed for the four different months (M, J, J, S) and for each year and each season, respectively, as presented in Tables 1 to 10.

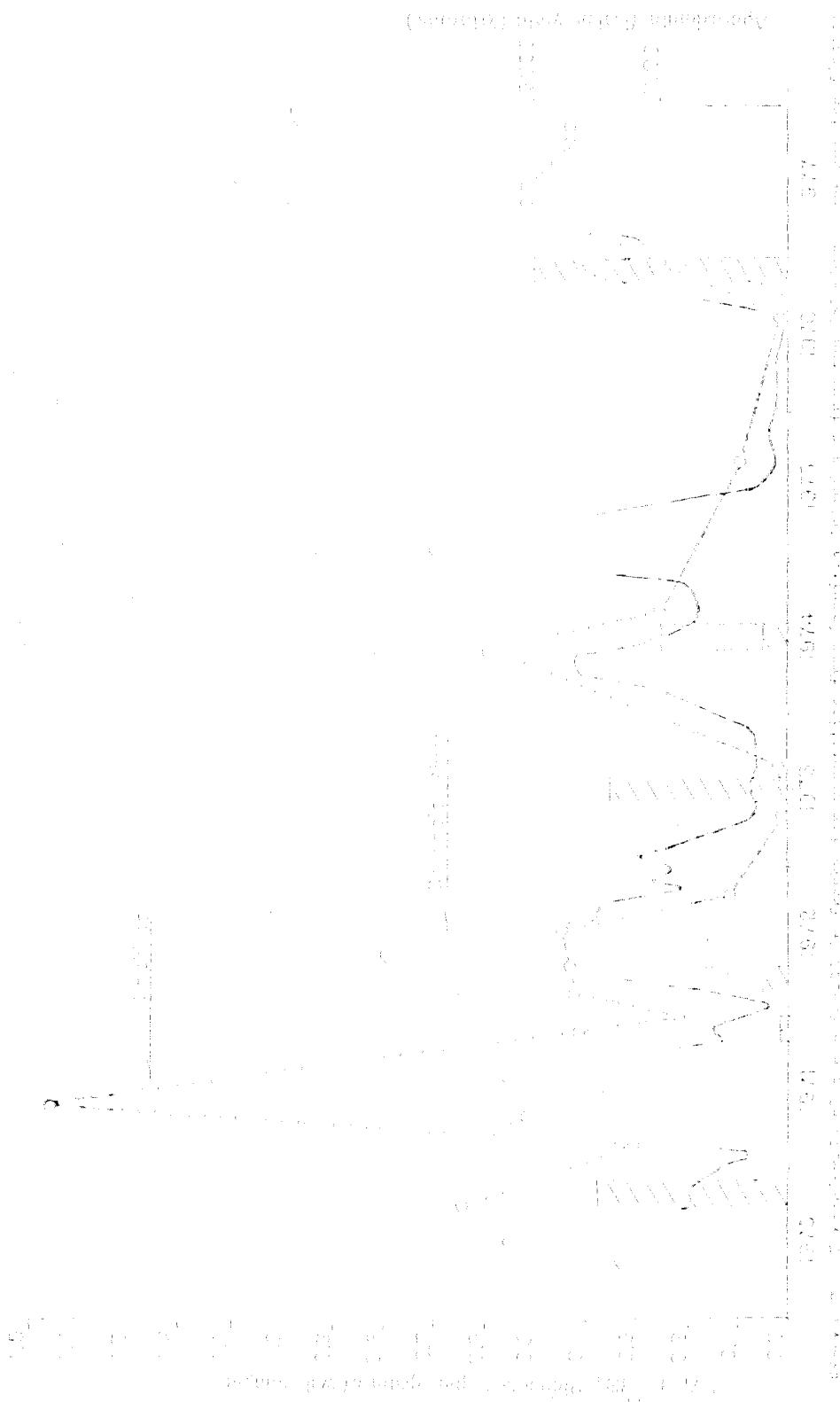
<sup>1</sup>The assistance of Álvaro Gómez, Glen Lofman, John Parker (Refugee and Exiles Foundation) (through) of the American Hydrological Federation program is gratefully acknowledged.

The results of date-specific analysis for three types of benthic macrofauna collected in freshwater are given in Table 6-10-6. The samples were from sand (a) area), fluvial (c) plain), and bottom (d) sediments (not specifically described) which represent algae, complex algae, soft-shelled clams, mussels, but only "sand" was present at all depths. Data in Table 6-10-6 are the values for the different years separately and the values for the year of collection are indicated "year of collection" after each year, i.e., 1970, 71, 72, 73, and 74.

To test for "post-hoc" the "year of collection" effect, the values for the amount of benthics in (% dry weight) (%db) in both the freshwater were compared with the amount of (% dry weight) (%db) from 1970, 71, and 72 samples from the Columbia River and with the value of date-specific denoted as "year" (denoted as "year" of collection) of 1973 data from the Columbia River. These data are presented in Figure 7.

The data in Figure 7 were selected for the following reasons: (1) 70 and 710 are samples of freshwater benthics; the water body is the Columbia River; the period is in the fall; the sediment texture was selected as an average value of index of specific surface of sediments from the Columbia River were selected as an input value; (2) the numbers samples from a total of approximately the 1000 samples in the river and the number of benthic samples denoted as "year" of collection with the 72% plain (%db) value because the 72% is the perfectly sample of fall benthic community in the samples of the Columbia. The results of one year of the Columbia River samples (fall) were to compare benthic samples (Columbia) were provided by Lamm (1970). The Columbia River samples were collected by the U.S. Army Corps of Engineers. The Columbia River samples were reported in terms of dry weight and for this reason the data are converted to dry weight. The data of wet weight of the samples in Figure 7 are converted to dry weight. The wet weight values are calculated from the wet weight values (Lamm, 1970) and determined from the measurements of 10 samples in 70, 71, 72, 73, and 74. The numbers of samples are denoted as "year" of collection in Figure 7. The nature of importance about the difference in the sediment texture (Lamm, 1970), the position of Figure 7 provide evidence for the following comments: (1) the benthic in the Columbia River samples are greater (2) the benthic in the river occur after Columbia became condition of the Columbia greater than yields (3) there are four "year" of collection" reflected in point in 1970, 71, 72, 73 and 74; (4) the collection of macrofauna in the Columbia River occurred in the autumn, fall, and fall-spring months in the Columbia.

From comments (3) and (4) above, it is our conclusion is that the date of collection of macrofauna with the Columbia River, in addition to the date of collection. Another method of determining the nature of the macrofauna is to determine the date of collection of the fauna (%db) and the proportion of short-term benthic samples collected in the Columbia. The date of collection of macrofauna can be determined from the date of collection, proportion of short-term benthic samples collected in the Columbia. The proportion of any kind of macrofauna in the benthic samples collected in the Columbia is denoted as "year" of collection. The data of the macrofauna measured in the Columbia are collected in Table 6-10-6. The data in Table 6-10-6 are converted to dry weight of the macrofauna (%db) and the proportion of short-term benthic samples collected in the Columbia (%db).



$^{137}\text{Cs}$  (6 and 30 days post-fallout) at the same sampling point, collected in the same way, then there is no difference in percent removal from the leaf surface of the fallout Cs-137 (Boatman 1974) and humic fulvic acid.

There has been no systematic collection of the radiocesium data for a specific area located in the set of sites, the availability of the samples related to the plant life occurring in the field. However, in Table 3, the September 1977 values from the marsh area near the Lake Ontario shore at Tonawanda show that the average Cs-137 concentration in the sediments was higher than the average radiocesium value in the vegetation, which may be due to the difference in the time of sampling. The average Cs-137 concentration in the sediments (Leavenworth Creek, Bay of Quinte) and the surface soils of the forested areas (Hedge Creek, Long Island, Chittenango, Seneca) are significantly lower than the average Cs-137 concentration in the vegetation. The availability of each sample was significantly different so that the mean of the availability for the vegetation and soil samples of Tonawanda, Bay of Quinte, Hedges Creek, Chittenango, and Seneca, the mean of the two, and one of each of the samples, for each location, is given from the Great Lakes Science and Resource Survey, 1977-1978. One sample, however, had to be omitted and a total of 10 samples were used to determine the significance of the mean differences. The 11th sample was omitted and the confidence limit was calculated using the Student's *t*-test. Hence, even though the differences were all on three-fold, the differences were not statistically significant because of the inaccuracy of the error bar and the number of samples.

Estimated Cs-137 values for Keweenaw are given in Table 4, for 1977, 1978, and in Table 5 for 1979, 1980, 1981. The results of a broader range of sediments, pollen, other microorganisms, and geochemical fractions, were not included in the present analysis because of the small number of samples, the variable nature of the samples, and the lack of precision. The values are calculated from the collection date, location of fallout and the date of collection, both indicated in Table 4 for the Keweenaw samples. The former values for 1977 and 1978 are greater than for the 1979 samples, except for otherwise noted.

Sample of the Keweenaw, 1977 (Table 7) had four fallout Cs-137 dates given and all other concentrations in 1977 and 1978 other years from the last 10 years were taken from Table 4 and Keweenaw. The first 117 Cs-137 dates recorded for Keweenaw probably demonstrate that in Table 2 the 29  $\mu\text{g}/\text{g}$  dry sample given in 1978 from the marsh gave an average of about 30% Cs-137 fallout. Thus, 30% Cs-137 were detected in the 1977 sample for the first time since 1976.

Estimated radiocesium values for the vegetation, remaining vegetation, in terms of  $\mu\text{g}/\text{g}$  of dry sample, are given in Table 6, were 6.2 for 1977 and ranged from 0.4 to 7.8 for the different sampling points.

Relatively occurring Cs-137 dates reported in Table 6 are the two earliest radiocesium originally detected in the Keweenaw, 1977, 1978, and analyzed in 1977. According to Dr. J. Norden, no greater than 2.7% Cs-137 detectable in five samples over the entire area was given 0.4  $\mu\text{g}/\text{g}$ .





100 and 500 m from the coastline, and for five of the four most heavily contaminated sites, the values for the bottom sediments from Government surveys at 100 m were about twice as great as the values for water samples taken at the same distance, and not matched by the 500 m data (Dovey et al., 1977). In the previous paper, the values for PCBs determined from the four 500 m surveys were converted into a mean of the four concentrations taken at the other depths, and the "mean" PCB concentration was calculated. It is clear from the data presented in the figure that each collected sample is in the fourth quartile. Because of samples taken at different distances from the coastline, it is difficult to interpret the previous paper.

Since 1976, and the last survey, many more open ocean surveys have been conducted, both for PCBs and the residue of chlorinated dioxins and furans (PCDD/Fs). The general conclusions from these are that the total PCB concentrations in the open ocean are higher than those in coastal waters, and that there are no obvious differences related to the location. The highest concentrations are probably those associated with the continental shelves off the Atlantic coast (Dovey et al., 1978), California (Hong et al., 1977), and Washington (Lindstrom and Lindberg, 1976) and were found to be higher, on average, than the levels in the waters immediately adjacent to the coast. In particular, the value of 1.5 ppm PCBs (Lindstrom and Lindberg, 1976), for this reason, the choice of 1.5 ppm PCBs as a reference is believed to be the same as for other areas of the world's oceans.

A long-term national survey programme, the National Marine Fisheries Survey (NMFS), was initiated in 1976 and the results of the survey for 1977 and the three previous years are presented in Fig. 1. The overall survey (Rideout et al., 1978) is based on a 100 km grid, and is similar to the 100 m grid used in the previous paper. The total greater than 0.01 ppm PCBs per square kilometer for the survey of 1977 is plotted in Fig. 1. The survey covers roughly the area from 10°N to 40°N and 100°W to 160°W. The survey never reaches far off-shore, and 17°N to the north and 14°S to the south are the extremes, the boundaries near the equator. Most of the data points represent five to thirty-five kilometers from the coast.

## 3. SURVEY AND CONCLUSION

The major objective of this paper is to determine the extent of pollution of the continental shelf of the Pacific Ocean by the chlorinated hydrocarbons, particularly of chlorinated and polychlorinated dibenz-p-dioxins and furans. The survey is a relatively extensive one which can be expected to be world-wide. The values of the concentrations in the surface sediments of the continental shelf would be somewhat higher than the values given in the survey for September 1977, and also higher than would be expected in the future, because of the evidence that in the lead-up to the coming year, i.e., 1978, the evidence shows the marine-life contamination by PCBs will be greater than would be expected from the factors except for a slight continuation of the long-term natural variation.

1. Mining areas are subject to many types of pollution which can damage the environment both land and marine, particularly at Ashfield Island and on the Bay Islands, with the resultant loss of economic value due to the loss of fish and other natural resources.

2. The effects of industrialisation on the environment have been studied by the Committee.

3. Radioactive products, radon and radon decay products have been detected in areas of the range from the plateau to the valley of the upper Río Grande.

4. Various forms of radioactive waste products from Ashfield Island and the Cuyamel River were gathered on the site and placed in containers.

5. Peaks of abundance of short-lived radionuclides occur at 50-1970-71, 1971 and 1977 and the long-lived radionuclides are also evident.

6. Two neutron products are shown in Table 1<sup>1</sup> to be the dominant fallout radionuclides in the samples and that the <sup>40</sup>K contribution which were present from the last Acidic plutonium extraction.

7. The radioactivity from fallout radionuclides generally increases with increasing distance from the sources.

8. There has been no response in the <sup>137</sup>Cs activity with a fall time from a nuclear emergency shorter than one year or less than underground water.

9. The background can always very easily exceed more than 10% of the total dose from the environment.

10. The repository detection and alarm system for the integrated analysis of the data is considered effective to detect changes in the standard operation conditions in the environs of the facility.

Supporting and Auxiliary Elements and their Weight to Dry Weight Ratio for Some Hardwood Cellulose Derivatives

Element	Type	Weight %	
		Cellulose	Residue
WATER SOLUBLE			
Acetyl cellulose	Acetyl	50.00	49.99
Acetyl cellulose	Methyl	50.00	49.99
Acetyl cellulose	Ethyl	50.00	49.99
Acetyl cellulose	Propyl	50.00	49.99
Acetyl cellulose	Butyl	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
ALKALI INSOLUBLE			
Acetyl cellulose	Acetyl	50.00	49.99
Acetyl cellulose	Methyl	50.00	49.99
Acetyl cellulose	Ethyl	50.00	49.99
Acetyl cellulose	Propyl	50.00	49.99
Acetyl cellulose	Butyl	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
ALKALI SOLUBLE			
Acetyl cellulose	Acetyl	50.00	49.99
Acetyl cellulose	Methyl	50.00	49.99
Acetyl cellulose	Ethyl	50.00	49.99
Acetyl cellulose	Propyl	50.00	49.99
Acetyl cellulose	Butyl	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
ALKALI INSOLUBLE			
Acetyl cellulose	Acetyl	50.00	49.99
Acetyl cellulose	Methyl	50.00	49.99
Acetyl cellulose	Ethyl	50.00	49.99
Acetyl cellulose	Propyl	50.00	49.99
Acetyl cellulose	Butyl	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
ALKALI SOLUBLE			
Acetyl cellulose	Acetyl	50.00	49.99
Acetyl cellulose	Methyl	50.00	49.99
Acetyl cellulose	Ethyl	50.00	49.99
Acetyl cellulose	Propyl	50.00	49.99
Acetyl cellulose	Butyl	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99
Acetyl cellulose	Ammonium	50.00	49.99
Acetyl cellulose	Chloroammonium	50.00	49.99
Acetyl cellulose	Sodium	50.00	49.99



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Table 3

Long-Run Efficient Production in the Resistor Industries, United States, 1900-1940

Industry	Period	Number of Firms	Number of Plants	Number of Workers	Output per Worker	Output per Plant	Output per Firm	Output per Firm per Worker	Output per Firm per Plant	Output per Worker per Plant	Output per Firm per Worker per Plant	Output per Firm per Worker per Plant per Firm
Aluminum	1900-1910	10	10	10,000	100	100	10	10	10	10	10	10
Aluminum	1910-1920	10	10	10,000	100	100	10	10	10	10	10	10
Aluminum	1920-1930	10	10	10,000	100	100	10	10	10	10	10	10
Aluminum	1930-1940	10	10	10,000	100	100	10	10	10	10	10	10
Automobiles	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Automobiles	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Automobiles	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Automobiles	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Bicycles	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Bicycles	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Bicycles	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Bicycles	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Clothing	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Clothing	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Clothing	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Clothing	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Electrical	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Electrical	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Electrical	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Electrical	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Furniture	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Furniture	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Furniture	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Furniture	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Leather Goods	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Leather Goods	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Leather Goods	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Leather Goods	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Machinery	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Machinery	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Machinery	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Machinery	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Metals	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Metals	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Metals	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Metals	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Plastics	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Plastics	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Plastics	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Plastics	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Textiles	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Textiles	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Textiles	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Textiles	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10
Wood Products	1900-1910	100	100	100,000	100	100	10	10	10	10	10	10
Wood Products	1910-1920	100	100	100,000	100	100	10	10	10	10	10	10
Wood Products	1920-1930	100	100	100,000	100	100	10	10	10	10	10	10
Wood Products	1930-1940	100	100	100,000	100	100	10	10	10	10	10	10

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Table 4













Table 10

Estimated Cost of the Standardized Collection Method

Number of sites	Number of samples per site	Number of samples per site per method	Number of samples per method	Cost per sample	Total cost
100	10	1	100	\$1.00	\$100.00
100	10	2	200	\$1.00	\$200.00
100	10	3	300	\$1.00	\$300.00
100	10	4	400	\$1.00	\$400.00
100	10	5	500	\$1.00	\$500.00
100	10	6	600	\$1.00	\$600.00
100	10	7	700	\$1.00	\$700.00
100	10	8	800	\$1.00	\$800.00
100	10	9	900	\$1.00	\$900.00
100	10	10	1000	\$1.00	\$1000.00
100	10	11	1100	\$1.00	\$1100.00
100	10	12	1200	\$1.00	\$1200.00
100	10	13	1300	\$1.00	\$1300.00
100	10	14	1400	\$1.00	\$1400.00
100	10	15	1500	\$1.00	\$1500.00
100	10	16	1600	\$1.00	\$1600.00
100	10	17	1700	\$1.00	\$1700.00
100	10	18	1800	\$1.00	\$1800.00
100	10	19	1900	\$1.00	\$1900.00
100	10	20	2000	\$1.00	\$2000.00
100	10	21	2100	\$1.00	\$2100.00
100	10	22	2200	\$1.00	\$2200.00
100	10	23	2300	\$1.00	\$2300.00
100	10	24	2400	\$1.00	\$2400.00
100	10	25	2500	\$1.00	\$2500.00
100	10	26	2600	\$1.00	\$2600.00
100	10	27	2700	\$1.00	\$2700.00
100	10	28	2800	\$1.00	\$2800.00
100	10	29	2900	\$1.00	\$2900.00
100	10	30	3000	\$1.00	\$3000.00
100	10	31	3100	\$1.00	\$3100.00
100	10	32	3200	\$1.00	\$3200.00
100	10	33	3300	\$1.00	\$3300.00
100	10	34	3400	\$1.00	\$3400.00
100	10	35	3500	\$1.00	\$3500.00
100	10	36	3600	\$1.00	\$3600.00
100	10	37	3700	\$1.00	\$3700.00
100	10	38	3800	\$1.00	\$3800.00
100	10	39	3900	\$1.00	\$3900.00
100	10	40	4000	\$1.00	\$4000.00
100	10	41	4100	\$1.00	\$4100.00
100	10	42	4200	\$1.00	\$4200.00
100	10	43	4300	\$1.00	\$4300.00
100	10	44	4400	\$1.00	\$4400.00
100	10	45	4500	\$1.00	\$4500.00
100	10	46	4600	\$1.00	\$4600.00
100	10	47	4700	\$1.00	\$4700.00
100	10	48	4800	\$1.00	\$4800.00
100	10	49	4900	\$1.00	\$4900.00
100	10	50	5000	\$1.00	\$5000.00
100	10	51	5100	\$1.00	\$5100.00
100	10	52	5200	\$1.00	\$5200.00
100	10	53	5300	\$1.00	\$5300.00
100	10	54	5400	\$1.00	\$5400.00
100	10	55	5500	\$1.00	\$5500.00
100	10	56	5600	\$1.00	\$5600.00
100	10	57	5700	\$1.00	\$5700.00
100	10	58	5800	\$1.00	\$5800.00
100	10	59	5900	\$1.00	\$5900.00
100	10	60	6000	\$1.00	\$6000.00
100	10	61	6100	\$1.00	\$6100.00
100	10	62	6200	\$1.00	\$6200.00
100	10	63	6300	\$1.00	\$6300.00
100	10	64	6400	\$1.00	\$6400.00
100	10	65	6500	\$1.00	\$6500.00
100	10	66	6600	\$1.00	\$6600.00
100	10	67	6700	\$1.00	\$6700.00
100	10	68	6800	\$1.00	\$6800.00
100	10	69	6900	\$1.00	\$6900.00
100	10	70	7000	\$1.00	\$7000.00
100	10	71	7100	\$1.00	\$7100.00
100	10	72	7200	\$1.00	\$7200.00
100	10	73	7300	\$1.00	\$7300.00
100	10	74	7400	\$1.00	\$7400.00
100	10	75	7500	\$1.00	\$7500.00
100	10	76	7600	\$1.00	\$7600.00
100	10	77	7700	\$1.00	\$7700.00
100	10	78	7800	\$1.00	\$7800.00
100	10	79	7900	\$1.00	\$7900.00
100	10	80	8000	\$1.00	\$8000.00
100	10	81	8100	\$1.00	\$8100.00
100	10	82	8200	\$1.00	\$8200.00
100	10	83	8300	\$1.00	\$8300.00
100	10	84	8400	\$1.00	\$8400.00
100	10	85	8500	\$1.00	\$8500.00
100	10	86	8600	\$1.00	\$8600.00
100	10	87	8700	\$1.00	\$8700.00
100	10	88	8800	\$1.00	\$8800.00
100	10	89	8900	\$1.00	\$8900.00
100	10	90	9000	\$1.00	\$9000.00
100	10	91	9100	\$1.00	\$9100.00
100	10	92	9200	\$1.00	\$9200.00
100	10	93	9300	\$1.00	\$9300.00
100	10	94	9400	\$1.00	\$9400.00
100	10	95	9500	\$1.00	\$9500.00
100	10	96	9600	\$1.00	\$9600.00
100	10	97	9700	\$1.00	\$9700.00
100	10	98	9800	\$1.00	\$9800.00
100	10	99	9900	\$1.00	\$9900.00
100	10	100	10000	\$1.00	\$10000.00

Table 10 illustrates the cost of the standardized collection method. The cost per sample is \$1.00. The cost per site is the number of samples per site times the cost per sample. The total cost is the number of sites times the cost per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 11 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 12 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 13 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

Table 14 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 15 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 16 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.

The following table shows the estimated cost of the standardized collection method for various numbers of sites and samples per site.

Table 17 illustrates the cost of the standardized collection method for various numbers of sites and samples per site.



TABLE III

Sedimentation and Conductivity in the Lake Ontario Bay Waters (in micrometers and milligrams per liter)

Lake Ontario	Conductivity micromhos/cm. <sup>a</sup>	Depth m.	Conductivity micromhos/cm. <sup>a</sup>	Depth m.	Conductivity micromhos/cm. <sup>a</sup>
South Bay	17.0	0.0	17.0	0.0	17.0
South Bay	17.0	10.0	16.9	10.0	16.7
South Bay	17.0	20.0	16.9	20.0	16.7
South Bay	17.0	30.0	16.9	30.0	16.7
South Bay	17.0	40.0	16.9	40.0	16.7
South Bay	17.0	50.0	17.0	50.0	17.0
South Bay	17.0	60.0	17.0	60.0	17.0
South Bay	17.0	70.0	17.0	70.0	17.0
South Bay	17.0	80.0	17.0	80.0	17.0
South Bay	17.0	90.0	17.0	90.0	17.0
South Bay	17.0	100.0	17.0	100.0	17.0
South Bay	17.0	110.0	17.0	110.0	17.0
South Bay	17.0	120.0	17.0	120.0	17.0
South Bay	17.0	130.0	17.0	130.0	17.0
South Bay	17.0	140.0	17.0	140.0	17.0
South Bay	17.0	150.0	17.0	150.0	17.0
South Bay	17.0	160.0	17.0	160.0	17.0
South Bay	17.0	170.0	17.0	170.0	17.0
South Bay	17.0	180.0	17.0	180.0	17.0
South Bay	17.0	190.0	17.0	190.0	17.0
South Bay	17.0	200.0	17.0	200.0	17.0
South Bay	17.0	210.0	17.0	210.0	17.0
South Bay	17.0	220.0	17.0	220.0	17.0
South Bay	17.0	230.0	17.0	230.0	17.0
South Bay	17.0	240.0	17.0	240.0	17.0
South Bay	17.0	250.0	17.0	250.0	17.0
South Bay	17.0	260.0	17.0	260.0	17.0
South Bay	17.0	270.0	17.0	270.0	17.0
South Bay	17.0	280.0	17.0	280.0	17.0
South Bay	17.0	290.0	17.0	290.0	17.0
South Bay	17.0	300.0	17.0	300.0	17.0
South Bay	17.0	310.0	17.0	310.0	17.0
South Bay	17.0	320.0	17.0	320.0	17.0
South Bay	17.0	330.0	17.0	330.0	17.0
South Bay	17.0	340.0	17.0	340.0	17.0
South Bay	17.0	350.0	17.0	350.0	17.0
South Bay	17.0	360.0	17.0	360.0	17.0
South Bay	17.0	370.0	17.0	370.0	17.0
South Bay	17.0	380.0	17.0	380.0	17.0
South Bay	17.0	390.0	17.0	390.0	17.0
South Bay	17.0	400.0	17.0	400.0	17.0
South Bay	17.0	410.0	17.0	410.0	17.0
South Bay	17.0	420.0	17.0	420.0	17.0
South Bay	17.0	430.0	17.0	430.0	17.0
South Bay	17.0	440.0	17.0	440.0	17.0
South Bay	17.0	450.0	17.0	450.0	17.0
South Bay	17.0	460.0	17.0	460.0	17.0
South Bay	17.0	470.0	17.0	470.0	17.0
South Bay	17.0	480.0	17.0	480.0	17.0
South Bay	17.0	490.0	17.0	490.0	17.0
South Bay	17.0	500.0	17.0	500.0	17.0
South Bay	17.0	510.0	17.0	510.0	17.0
South Bay	17.0	520.0	17.0	520.0	17.0
South Bay	17.0	530.0	17.0	530.0	17.0
South Bay	17.0	540.0	17.0	540.0	17.0
South Bay	17.0	550.0	17.0	550.0	17.0
South Bay	17.0	560.0	17.0	560.0	17.0
South Bay	17.0	570.0	17.0	570.0	17.0
South Bay	17.0	580.0	17.0	580.0	17.0
South Bay	17.0	590.0	17.0	590.0	17.0
South Bay	17.0	600.0	17.0	600.0	17.0
South Bay	17.0	610.0	17.0	610.0	17.0
South Bay	17.0	620.0	17.0	620.0	17.0
South Bay	17.0	630.0	17.0	630.0	17.0
South Bay	17.0	640.0	17.0	640.0	17.0
South Bay	17.0	650.0	17.0	650.0	17.0
South Bay	17.0	660.0	17.0	660.0	17.0
South Bay	17.0	670.0	17.0	670.0	17.0
South Bay	17.0	680.0	17.0	680.0	17.0
South Bay	17.0	690.0	17.0	690.0	17.0
South Bay	17.0	700.0	17.0	700.0	17.0
South Bay	17.0	710.0	17.0	710.0	17.0
South Bay	17.0	720.0	17.0	720.0	17.0
South Bay	17.0	730.0	17.0	730.0	17.0
South Bay	17.0	740.0	17.0	740.0	17.0
South Bay	17.0	750.0	17.0	750.0	17.0
South Bay	17.0	760.0	17.0	760.0	17.0
South Bay	17.0	770.0	17.0	770.0	17.0
South Bay	17.0	780.0	17.0	780.0	17.0
South Bay	17.0	790.0	17.0	790.0	17.0
South Bay	17.0	800.0	17.0	800.0	17.0
South Bay	17.0	810.0	17.0	810.0	17.0
South Bay	17.0	820.0	17.0	820.0	17.0
South Bay	17.0	830.0	17.0	830.0	17.0
South Bay	17.0	840.0	17.0	840.0	17.0
South Bay	17.0	850.0	17.0	850.0	17.0
South Bay	17.0	860.0	17.0	860.0	17.0
South Bay	17.0	870.0	17.0	870.0	17.0
South Bay	17.0	880.0	17.0	880.0	17.0
South Bay	17.0	890.0	17.0	890.0	17.0
South Bay	17.0	900.0	17.0	900.0	17.0
South Bay	17.0	910.0	17.0	910.0	17.0
South Bay	17.0	920.0	17.0	920.0	17.0
South Bay	17.0	930.0	17.0	930.0	17.0
South Bay	17.0	940.0	17.0	940.0	17.0
South Bay	17.0	950.0	17.0	950.0	17.0
South Bay	17.0	960.0	17.0	960.0	17.0
South Bay	17.0	970.0	17.0	970.0	17.0
South Bay	17.0	980.0	17.0	980.0	17.0
South Bay	17.0	990.0	17.0	990.0	17.0
South Bay	17.0	1000.0	17.0	1000.0	17.0

<sup>a</sup> Conductivity measured at 25°C. Salinity converted to micromhos/cm. by multiplying by 1.000.

and chloride values from which complete hydrographic conditions were determined. The temperature, conductivity, concentration of the sediment, and the water column depth at the time of sampling and of division in the lake were also determined.

#### Conductivity, Temperature, and Chloride

##### South Bay, Canfield Lake, Rock Cave,

##### South Bay, Canfield Lake, Kilding River, Rock Cave, Cleveland Creek





THE INFLUENCE OF SOIL ON PLANT GROWTH 131







Table 1.  $\langle \cos(\theta_{\text{obs}} - \theta_{\text{true}}) \rangle$

Table 16 (continued)

Location and Date	Condition	Radius of Search (m)	Intercept Probability	Probability of Detection
<b>1.1. Flooded Pathways (cont.)</b>				
1977	Pathway B	1	1100 ± 120	0.06 ± 0.01
1977	Pathway C	1	1000 ± 100	0.05 ± 0.01
1977	Pathway D	1	1100 ± 200	0.05 ± 0.01
1977	Pathway E	1	700 ± 100	0.04 ± 0.01
1977	Pathway F	1	1000 ± 100	0.04 ± 0.01
1977	Pathway G	1	2100 ± 200	0.04 ± 0.01
1977	Pathway H	1	1000 ± 100	0.04 ± 0.01
1977	Pathway I	1	1000 ± 100	0.04 ± 0.01
1977	Pathway J	1	1200 ± 100	0.04 ± 0.01
1977	Pathway K	1	1200 ± 100	0.04 ± 0.01
1977	Pathway L	1	1200 ± 100	0.04 ± 0.01
1977	Pathway M	1	1200 ± 100	0.04 ± 0.01
1977	Pathway N	1	1200 ± 100	0.04 ± 0.01
<b>1.2. Unflooded Pathways</b>				
August 1976	Distance below flood water level	1	800 ± 100	0.06 ± 0.01
C	Unflooded Road	1	600 ± 100	0.07 ± 0.01
A	100 meters below road	1	400 ± 100	0.07 ± 0.01
B	100 meters below road	1	300 ± 100	0.06 ± 0.01
D	200 meters above the bay	1	170 ± 10	0.04 ± 0.01
E	Road and creek	1	100 ± 10	0.04 ± 0.01
August 1976	Distance below flood water level	1	700 ± 100	0.07 ± 0.01
C	Unflooded Road	1	500 ± 100	0.07 ± 0.01
B	100 meters below road	1	300 ± 100	0.07 ± 0.01
F	200 meters below road <sup>1</sup>	1	150 ± 10	0.04 ± 0.01
G	400 meters below road <sup>1</sup>	1	100 ± 10	0.04 ± 0.01
H	500 meters below road <sup>1</sup>	1	50 ± 10	0.03 ± 0.01
I	700 meters above the bay	1	40 ± 10	0.03 ± 0.01
J	100 meters above the bay	1	20 ± 10	0.03 ± 0.01
September 1977	Unflooded Road	1	400 ± 100	0.06 ± 0.01
N	100 meters below road	1	100 ± 10	0.04 ± 0.01
M	200 meters below road <sup>1</sup>	1	60 ± 10	0.03 ± 0.01
O	400 meters below road <sup>1</sup>	1	10 ± 10	0.02 ± 0.01

10.2.10. *Geometric mean*



1. *What is the relationship between the two variables?*

<sup>1</sup> See the report of the Joint Committee on Taxation, Part I.





10. The following table gives the number of hours worked by each of the 100 workers.



U.S. Army, R., 1972, An Evaluation of Selected Methods, Aerial Survey, and Photogrammetry for Monitoring Water Quality in Rivers, in: *Final Report of the Inter-Agency Monitoring Project*, May 1972, Part 2.

Burkhardt, M. and P. A. Rappaport, 1972, *Aerial Survey and Photogrammetry for River Water Quality Monitoring*, Final Report, May 1972, Part 1, University of Colorado, Boulder, CO, U.S. Geological Survey, Denver, CO, 1972, 13 p.

DeGrazio, R. and M. J. DeGrazio, 1972, *Photogrammetric Methodology to Determine River Flow Rates*, Final Report, March 1972, The University of Arizona, Tucson, AZ, 10 p.

Tucker, R., 1972, *Archaeological Survey of the Columbia River in Oregon*, in: *Karlsruhe Conference Report, International Conference on Archaeology*, to be published.

United States Army Research and Development Command, 1972, *Guidelines for Report of Selected Topics*, Field Manual, Chapter 0134.

Upp, R. H., V. L. Ridder and T. K. Johnson, 1972, *Photogrammetry and Remote Sensing Handbook*, Volume 2, 7th edn.

7. Referring to one of previous Archaeological reports:

Year	Report No.	Author
1970/71	AMC 200-11	Peterson, E.
1972	AMC 200-12	Held, J. et al.
1972	AMC 200-13	Held, J. et al. et al.
1973	AMC 200-14	Peterson, M., H., and A. H., et al.
1974	AMC 200-15	Peterson, M., H., and A. H., et al.
1975	AMC 200-16	Peterson, M., H., and A. H., et al.
1976	AMC 200-17	Peterson, M., H., and A. H., et al.

(a) LOST AND FOUND (CONT.)

Item	Item No.	Item Description
1. Animal, Mammal		
a. Bear	1	Bear
b. Deer	2	Deer
c. Fox	3	Fox
d. Goat	4	Goat
e. Rabbit	5	Rabbit
f. Sheep	6	Sheep
g. Wolf	7	Wolf
2. Animal, Bird		
a. Duck	8	Duck
b. Hen	9	Hen
c. Pigeon	10	Pigeon
d. Sparrow	11	Sparrow
e. Swan	12	Swan
f. Turkey	13	Turkey
g. Vulture	14	Vulture
3. Animal, Fish		
a. Carp	15	Carp
b. Goldfish	16	Goldfish
c. Trout	17	Trout
4. Animal, Reptile		
a. Lizard	18	Lizard
b. Snake	19	Snake
5. Animal, Insect		
a. Bee	20	Bee
b. Fly	21	Fly
c. Grasshopper	22	Grasshopper
d. Moth	23	Moth
e. Spider	24	Spider
f. Wasp	25	Wasp
6. Animal, Fish		
a. Goldfish	26	Goldfish
b. Koi	27	Koi
7. Animal, Reptile		
a. Lizard	28	Lizard
b. Snake	29	Snake
8. Animal, Insect		
a. Bee	30	Bee
b. Fly	31	Fly
c. Grasshopper	32	Grasshopper
d. Moth	33	Moth
e. Spider	34	Spider
f. Wasp	35	Wasp
9. Animal, Fish		
a. Goldfish	36	Goldfish
b. Koi	37	Koi
10. Animal, Reptile		
a. Lizard	38	Lizard
b. Snake	39	Snake
11. Animal, Insect		
a. Bee	40	Bee
b. Fly	41	Fly
c. Grasshopper	42	Grasshopper
d. Moth	43	Moth
e. Spider	44	Spider
f. Wasp	45	Wasp
12. Animal, Fish		
a. Goldfish	46	Goldfish
b. Koi	47	Koi
13. Animal, Reptile		
a. Lizard	48	Lizard
b. Snake	49	Snake
14. Animal, Insect		
a. Bee	50	Bee
b. Fly	51	Fly
c. Grasshopper	52	Grasshopper
d. Moth	53	Moth
e. Spider	54	Spider
f. Wasp	55	Wasp
15. Animal, Fish		
a. Goldfish	56	Goldfish
b. Koi	57	Koi
16. Animal, Reptile		
a. Lizard	58	Lizard
b. Snake	59	Snake
17. Animal, Insect		
a. Bee	60	Bee
b. Fly	61	Fly
c. Grasshopper	62	Grasshopper
d. Moth	63	Moth
e. Spider	64	Spider
f. Wasp	65	Wasp
18. Animal, Fish		
a. Goldfish	66	Goldfish
b. Koi	67	Koi
19. Animal, Reptile		
a. Lizard	68	Lizard
b. Snake	69	Snake
20. Animal, Insect		
a. Bee	70	Bee
b. Fly	71	Fly
c. Grasshopper	72	Grasshopper
d. Moth	73	Moth
e. Spider	74	Spider
f. Wasp	75	Wasp
21. Animal, Fish		
a. Goldfish	76	Goldfish
b. Koi	77	Koi
22. Animal, Reptile		
a. Lizard	78	Lizard
b. Snake	79	Snake
23. Animal, Insect		
a. Bee	80	Bee
b. Fly	81	Fly
c. Grasshopper	82	Grasshopper
d. Moth	83	Moth
e. Spider	84	Spider
f. Wasp	85	Wasp
24. Animal, Fish		
a. Goldfish	86	Goldfish
b. Koi	87	Koi
25. Animal, Reptile		
a. Lizard	88	Lizard
b. Snake	89	Snake
26. Animal, Insect		
a. Bee	90	Bee
b. Fly	91	Fly
c. Grasshopper	92	Grasshopper
d. Moth	93	Moth
e. Spider	94	Spider
f. Wasp	95	Wasp
27. Animal, Fish		
a. Goldfish	96	Goldfish
b. Koi	97	Koi
28. Animal, Reptile		
a. Lizard	98	Lizard
b. Snake	99	Snake
29. Animal, Insect		
a. Bee	100	Bee
b. Fly	101	Fly
c. Grasshopper	102	Grasshopper
d. Moth	103	Moth
e. Spider	104	Spider
f. Wasp	105	Wasp

## (M) 1990 PMS INDEX (CONTINUED)

NAME	PAPERS	PAPERS
	$\{\Omega_1, \Omega_2\}$	$\{\Omega_1, \Omega_3\}$
2000-1990 PMS INDEX	0	0
2000-1990 PMS INDEX	2	2
2000-1990 PMS INDEX	6	13
2000-1990 PMS INDEX (continued) (continued)	3	10
2000-1990 PMS INDEX	7	17
2000-1990 PMS INDEX	8	16
2000-1990 PMS INDEX	8	17
2000-1990 PMS INDEX	8	17





APPENDIX 2

RESULTS OF PLUTONIUM AND GAMMA  
SPECTRUM ANALYSES OF ENEWETAK  
PLANT, RAT AND SOIL SAMPLES

COLLECTED IN MARCH 1976

UNIVERSITY OF CALIFORNIA  
LABORATORY REPORT

ATOMIC ENERGY RESEARCH

January 26, 1972

A. Nevinisi

Principle Investigator: Dr. William J. Stoeck  
Subject: Results of Plutonium Gamma Spectrum Analyses of  
Plutonium-treated and control samples collected in  
Korea, 1970

Rat, plant, and soil samples from Korea (*vide infra*) were sent to our laboratory by Dr. Jack O. Smith concerning the plutonium analysis. These samples had been directly dried and packed by Dr. Stoeck's group.

The soil samples were checked for gamma activity by use of standard sample holders and counted for 400 minutes on a 3730 detector system. The gamma-counting results for each sample are given in Table 1. The quantity of the individual radionuclides required was sufficient to fill the standard sample holder. The samples were added to the standard sample holder in batches of two or three for gamma counting. The results of analyses for gamma-emitting radionuclides found in these samples are given in Table 2.

After gamma-counting, the samples were prepared for plutonium analysis were spiked with  $^{238}\text{Pu}$  and chemically yielded plutonium. The laboratory standard methyl acetate was used for separation of the plutonium. Finally, plutonium was extracted onto cellulose filter and measured with a sodium bismuthate detector and scintillation analyzer. The results of plutonium yields are given in Table 3.

cc: Seymour  
Schell  
Nevinisi

Table 3. Significant differences between control and treated groups of *Glossy Ibis* and *Black-necked Stilt* at different stages of development.

“*It is a very good place to go to, but it is not a place to go to.*”



Table 1. Partitioning (%) and Distribution Coefficient of Treatedek in Marine Sediments Collected by Drift-Wall Collection at Shallow in March 1991. Values are  $\mu\text{g}/\text{dry weight}$  of sample.<sup>a</sup>

Sample	Depth (cm)	Mean Depth	SD
11	0-10 cm	0.107 $\pm$ 0.031	0.057 $\pm$ 0.123
17	0-10 cm	0.047 $\pm$ 0.016	0.043 $\pm$ 0.012
18	0-10 cm	0.034 $\pm$ 0.009	0.024 $\pm$ 0.005
24	0-10 cm	0.027 $\pm$ 0.006	0.031 $\pm$ 0.006
25	0-10 cm	0.014 $\pm$ 0.006	0.049 $\pm$ 0.022
26	0-10 cm	0.007 $\pm$ 0.007	0.007
27	0-10 cm	0.006 $\pm$ 0.001	0.006 $\pm$ 0.002
28	0-10 cm	0.011 $\pm$ 0.001	0.012 $\pm$ 0.004
29	0-10 cm	0.003 $\pm$ 0.001	0.01 $\pm$ 0.002
30	0-10 cm	0.007 $\pm$ 0.001	0.051 $\pm$ 0.010
31	0-10 cm	0.004 $\pm$ 0.002	0.062 $\pm$ 0.011
32	0-10 cm	0.007 $\pm$ 0.006	0.015 $\pm$ 0.003
33	0-10 cm	0.008 $\pm$ 0.009	0.75 $\pm$ 0.07
34	0-10 cm	0.005 $\pm$ 0.001	0.032 $\pm$ 0.009
35	0-10 cm	0.005 $\pm$ 0.001	0.014
36	0-10 cm	0.006 $\pm$ 0.001	-0.0028
37	0-10 cm	0.007 $\pm$ 0.003	0.001
38	0-10 cm	0.007 $\pm$ 0.001	0.06 $\pm$ 0.03
39	0-10 cm	0.004 $\pm$ 0.001	0.34 $\pm$ 0.32
40	0-10 cm	0.004 $\pm$ 0.001	-0.005
41	0-10 cm	0.003 $\pm$ 0.001	-0.004
42	0-10 cm	0.004 $\pm$ 0.001	-0.001
43	0-10 cm	0.003 $\pm$ 0.001	-0.003
44	0-10 cm	0.004 $\pm$ 0.001	0.009
45	0-10 cm	0.005 $\pm$ 0.001	0.005 $\pm$ 0.002
46	0-10 cm	0.003 $\pm$ 0.003	0.003 $\pm$ 0.002
47	0-10 cm	0.004 $\pm$ 0.001	0.02 $\pm$ 0.15
48	0-10 cm	0.004 $\pm$ 0.001	-0.002
49	0-10 cm	0.004 $\pm$ 0.001	0.03 $\pm$ 0.013
50-1	0-10 cm	0.005 $\pm$ 0.001	0.337 $\pm$ 0.34
50-2	n	0.11 $\pm$ 0.07	0.28 $\pm$ 0.1
50-3	n	0.11 $\pm$ 0.09	0.04 $\pm$ 0.11
50-4	n	0.063 $\pm$ 0.009	0.03 $\pm$ 0.01
50-5	n	0.056 $\pm$ 0.013	0.47 $\pm$ 0.31
50-6	n	0.067	0.21 $\pm$ 0.19

Table 3. Continued

Sample	Sample type	Mean	SD
DS-1	Soil (0-10 cm depth)	71.6 ± 77.7	154 ± 14.3
DS-2	Soil	136.1 ± 13.6	76.1 ± 3.6
DS-3	Soil	130.4 ± 1.7	71.4 ± 6.45
DS-4	Soil (0-10 cm depth)	9.71 ± 0.78	9.77 ± 1.17
DS-5	Soil	7.01 ± 0.37	7.89 ± 1.22
DS-6	Soil	7.66 ± 0.65	8.84 ± 1.22

± one sigma

\* All count rates from field samples obtained within the one-sigma, constant background error. The mean ( $\bar{x}$ ) values are blind values and one standard deviation divided by constant background.

APPENDIX 3

RADIOLOGICAL SURVEY OF PLANTS, ANIMALS AND SOIL  
IN MICRONESIA

NOVEMBER 1975

by

Vinton A. Nelson

MONITORING SURVEY OF HAZARD, BREAKS, AND ROT  
IN MICRONESIA, APRIL 1976

by

Victor A. Rehman

PR-Applied Physics

UNIVERSITY OF MICHIGAN  
COLLEGE OF MEDICINE  
Institute of Radiological Sciences  
Ann Arbor, Michigan 48106

Approved for the U.S. Department of Energy  
Review Operator, ORNL under Contract No. Y-16-S-08-0050

## REFERENCES

From 1976 to 1979 about 40 devices were monitored under water, over water, or land or in the atmosphere over the water of the central Pacific. Most of these truly took place at Rikulu and French Franks in the Marshall Islands, and some at Johnston, Oahu (Hawaii), and Midway (Northern Pacific). The distribution of neutrino fluxes predicted by these areas has been roughly, but relatively, accurately determined by the available calculations. The present report is part of a laboratory of Neutrino factory program begun in 1974 and described in a previous paper (Richter, 1977). The purpose of this study was to determine qualitatively and quantitatively, radioactive progeny from a common fission source released in Rikulu, other than those produced by the local fission during the test periods. This resulted while M. J. McAttee, Jr., the Marshall Islands team, brought up the French Islands, Rikulu, French Franks, Johnston, Oahu, and French Franks, in the early 1970's. Data was taken collected in the early 1970's prior to comparison with the current and kinds of neutrino fluxes found in mid-Pacific from Rikulu and French Franks at 73°.

## ACKNOWLEDGMENT

The areas mentioned above were visited in November 1975. The French Islands areas are shown specifically to the joint effort of Rikulu and French Franks. The white light camera and 3 Geiger counter sites within these areas are shown. The film was scanned directly with a scanner from Brookhaven National Laboratory (BNL) which had radiated previously according with sodium iodide (NaI) scintillation detector and a preamplifier in cascade. The results of the survey referred will be given in a separate briefer report. Personnel from

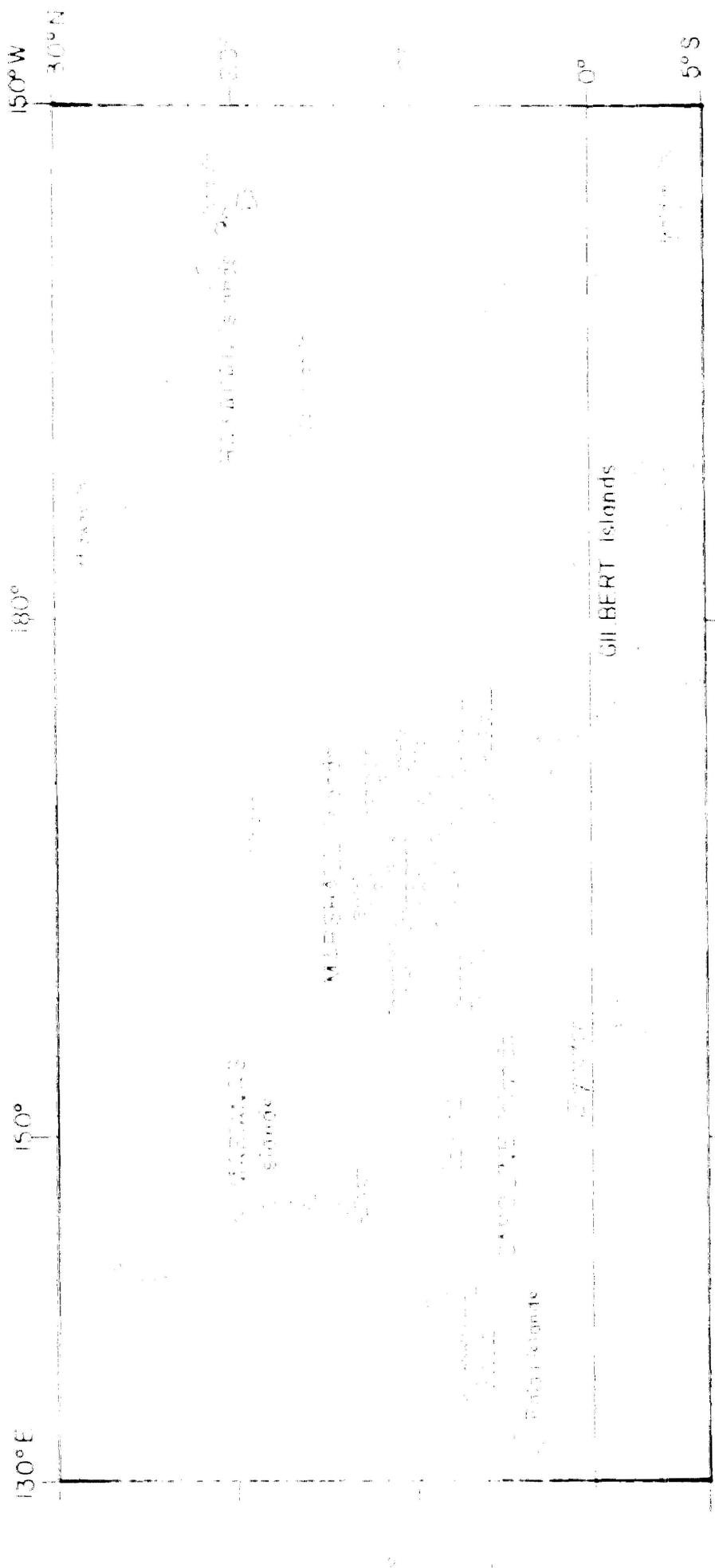
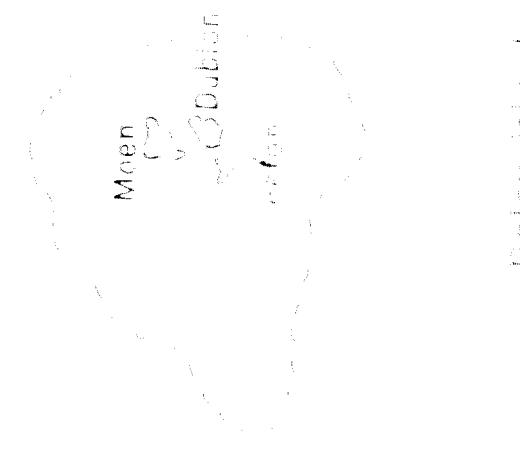


FIGURE 7. Salinity lines based on either frontal or off-frontal profiles for development of the AIL.

## Ponape District



## Truk Atoll



## Suam Island



## Koror Island

Hand-drawn map of Koror Island. The island is roughly triangular with a jagged coastline. Two districts are labeled: Shomizu (top right) and Arri (bottom right).

## Majuro Atoll

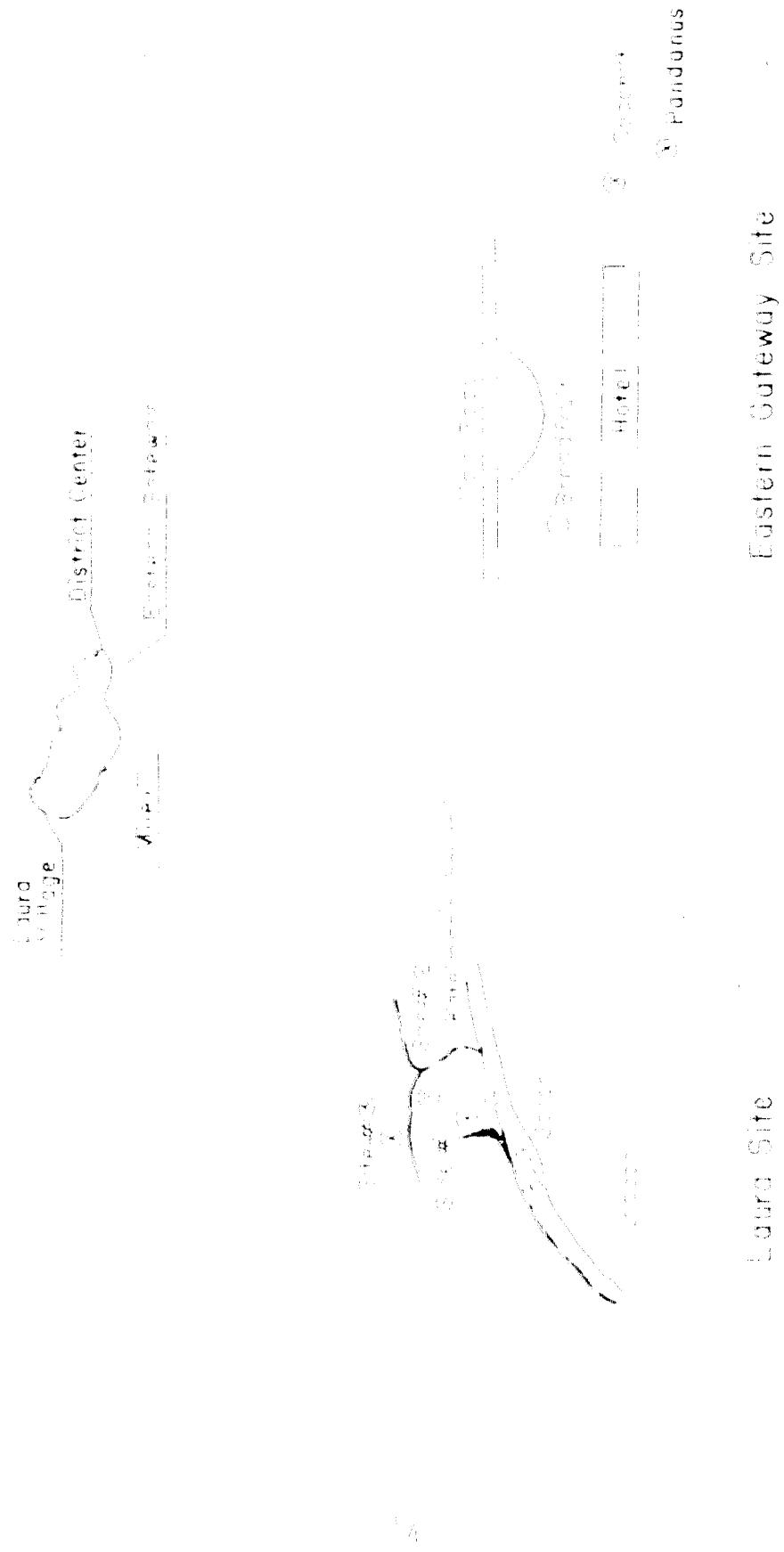


Figure 2. Locations camped at Majuro Atoll in November 1991.

248 energy-corrected gamma-ray biexponential and total samples with each sample forming the data set due to the effect of the beta-<sup>3</sup>-radioisotope (i.e., <sup>57</sup>Co, <sup>60</sup>Co, <sup>65</sup>Nickel, <sup>67</sup>Ruthenium, <sup>75</sup>Sodium, <sup>87</sup>Rubidium, <sup>90</sup>Samarium, etc.). A Monte Carlo multiple scattering of these elements was simulated and spectra data sets were subjected to provide data for estimation of future differentiation and quantities of radioactive isotopes in the raw gamma data.

The number of samples after division into counts per unit fractions, is shown in Table I. Twenty percent of the samples were biological specimens, and human subjects made thirty percent with counts on face (n=1,661) and acrylic (n=1,060). Approximately equal numbers of samples came from the first five major collection areas.

#### ANALYTICAL METHODS

##### Gamma-Ray Spectrometry

All of the samples were analyzed by gamma-ray spectrometry either with a 32-channel counter (thallium-doped) crystal and 700-channel pulse-height analyzer or with a germanium (lithium-doped) oxide detector and 4,096-channel, pulse-height analyzer. Both samples and raw data on the Ge(Li) system, and the digitized samples were analyzed on-line systems.

All samples were oven dried, ground and a portion counted in a sample holder of polyvinyl chloride (PVC) pipe 7 mm in diameter and either 1.2 cm or 0.6 cm for radionuclide measurement, using approximately tissue or 6 g grams of salt culture, respectively. The 1.2 cm holder, the sensitivities of the biexponential and total samples were 1.0 and 0.7, respectively. These samples were then analyzed for optimum counting conditions.

The gamma-shielding redondection to the samples counted on the NaI crystal were determined by a method of least squares. The radiometric values for samples counted on the Ge(Li) detector were calculated either manually or with

Table I. Description of Samples Collected on the November 1976 Trip to Keweenaw.

Sampling station	Sampled depth (m)	Sampled area (m <sup>2</sup> )				
1	29	10	6	67	26	16
2	73	13	16	86	39	17
3	31	14	1	66	14	12
4	31	13	1	70	16	7
5	25	15	3	63	12	13
Total	119	67	51	216	109	67

- a. The individual cores of the total area sampled have been divided into discrete increments of 1 m depth.
- b. Five cores were collected at each sampling station. Kerosene was added to each core, and a portion of each sample was extracted and analyzed for total methanol plus formaldehyde.

corrected by adding the counts from each energy of the channels, and applying the optimum thresholding to appropriate background counts, and applying a correction factor to convert counts to picocuries (pcu). First, the previously mentioned reference spectrum and the type of sample holder are used to calculate the total volume were converted to decay to the date of analysis.

### Counting and Quality Analysis

To measure  $^{238}\text{U}$  content,  $^{238}\text{U}$  was selectively separated from  $^{235}\text{U}$ , and the alpha signal was counted with a low-level beta counting system. The signal was corrected by the cascade effect, pile-up subtraction, and a low-energy alpha-suppression system. An additional buffer alpha detector at a right-angle provides the yield determined by sum of  $^{238}\text{U}$  counts.

### Error Budget

To account for all the errors of the measurement, first the two sigma of applied, counting time, the error for the sum of counts, then the sigma in the standard deviation are disregarded, count times, and the systematic error.

### Summary

Many factors influence the limit of detection, including the type of detector and its size, the detection order coincidence, the duration of the counting period, the size and density of the sample, and the geometric relationship of the sample and detector. Because the limits of detection varied considerably for various conditions and types of samples, it can be summarized by saying that detection effects were approximately as follows:

### By gamma detection

100%

95.4%

Concentration of  $\text{^{137}Cs}$  in dry weight of sample (ppm) = 0.12  $\mu\text{Ci}/\text{g} \times 10^3$

### By beta detection

100%

0.7  $\mu\text{Ci}/\text{g} \times 10^3$

### By alpha detection

100.0%

0.07  $\mu\text{Ci}/\text{g} \times 10^3$

## RESULTS AND DISCUSSION

Data are presented in Appendix Table I (Appendix II) for the results of the analysis of the samples collected by us in December 1975. All data are expressed in microcuries per gram of dry weight ( $\mu\text{Ci}/\text{g}$ , dry), except where expressly noted, while  $\mu\text{Ci}/\text{g}$  denotes the measured activity in dry weight material at the collection point. Thus the  $\mu\text{Ci}/\text{g}$ , dry values may be converted to  $\mu\text{Ci}/\text{g}$  wet by applying a factor of unity minus of the measured water content. Since there are greater differences in the availability value between different types of the organic matter than between the available and unavailable types, the results will be discussed by organic type.

### Fats

Seven species of fish were collected from the ocean in the six months. Gafftopsail, yellowtail, mackerel, were obtained from the three local ports and were collected in bottom trawl mesh. As shown by Appendix Table I, gafftopsail contained the only incorporate measureable tritium ( $\text{^{3H}}$ ) concentration greater than 0.6  $\mu\text{Ci}/\text{g}$ , dry and the average value was 4.8  $\mu\text{Ci}/\text{g}$ , dry. Tritium ( $\text{^{3H}}$ ) was detected in only 6 of 13 samples. The mean concentration of  $\text{^{3H}}$  measured was 0.9  $\mu\text{Ci}/\text{g}$ , dry. The two mackerel collected from Kurejima Harbor ( $\text{^{3H}}$ ) was detected in only 13 of 14 samples and the

Table 2. Mean Yield and Root Weight Ratio of Some  
Mycorrhizal Grains

Root Weight	Mean Yield	Root/ Yield	Deviation	
(kg/ha)				
Wheat	(6)	Uninoculated grain	2.36	+ 1.2
"	(3)	Vaccaria	4.65	+ 3.2
"	(1)	Luzerne	3.64	+ 1.1
Pearl millet	(2)	Sunflower	4.70	+ 1.7
"	(6)	Vaccaria	4.35	+ 0.7
"	(2)	Rainbow	3.45	+ 1.1
Soybean	(6)	Vaccaria + Manz	3.37	+ 1.5
"	(3)	Vaccaria	4.33	+ 0.8
Sorghum	(1)	Vaccaria	4.75	+ 1.2
"	(1)	Vaccaria + G. Hill	3.76	- 0.5
Flagstaff	(1)	Vaccaria	3.70	- 0.5
"	(1)	Vaccaria + Manz	3.09	- 0.5
Convolvulus	(1)	Vaccaria	3.56	- 0.5
"	(1)	Vaccaria + Manz	3.53	- 0.5
Jack	(1)	Luzerne	3.92	- 0.5
(kg/ha)				
Breadfruit	(17)	Coffee	6.17	+ 1.1
"	(16)	Guadua	6.24	+ 1.1
"	(15)	Luzerne	4.76	+ 1.64
Papaya	(17)	Coffee	7.23	+ 1.12
"	(13)	Theobroma	6.27	+ 0.7
"	(16)	Guadua	3.86	+ 0.7
Cocoyam	(11)	Guadua	2.60	+ 0.27
"	(3)	Guadua	2.27	+ 0.2
"	(1)	Coffee	3.72	+ 0.22
Taro	(8)	Guadua	2.71	+ 0.51
"	(7)	Coffee	7.00	+ 0.2
"	(7)	Guadua	16.70	+ 1.57
Papaya	(7)	Coffee	12.19	+ 3.12
"	(7)	Guadua	10.69	+ 2.11
"	(7)	Guadua	6.62	+ 1.3
Cassava	(1)	Root	2.10	- 0.5
Banana	(1)	Coffee	6.02	- 0.5

Table 7 (continued)

Specific	Knowles of Land	Mark	Learn Vest/Poly Ratio	Deviation
MANUFACTURERS				
Completion	{3}	Completion	{1, 3}	
Completion	{4}	Completion	{2, 3}	
Completion	{4}	Completion	{4, 5}	

Growth and Development

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*Journal of the American Statistical Association*, Vol. 33, No. 201, March, 1938.

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are presented in Appendix tables 7 (bottom) and 8 (from the soil profile), values generally correlated with  $(\text{cm}^3 \text{ g}^{-1})$  amounts of  $^{238}\text{U}$ ,  $^{226}\text{Ra}$ ,  $^{228}\text{Th}$ , and  $^{228}\text{Ra}$  measured from the volume (in cm<sup>3</sup>) of samples taken at 0–10, 10–20, and 20–30 cm depth from the surface (bottom of sample tube),  $R_{\text{eff}}$ , and  $R_{\text{soil}}$ . In addition to these values, figures were compared with values obtained by neutron activation analysis (NAA) (the well-known  $^{232}\text{Th}$  and  $^{226}\text{Ra}$  method) (Boggs et al., 1979; Boggs and Ritter, 1980; Boggs et al., 1981; Boggs and Ritter, 1982). The difference between the two methods was found to be within 10% (with a few exceptions) (Boggs et al., 1981). No major differences between the amount of  $^{226}\text{Ra}$  (in  $\text{Bq m}^{-2}$ ) found in the soil samples were observed except for three outliers.

Table 7 shows the variations of the measured higher values of  $^{226}\text{Ra}$  ( $^{226}\text{Ra}_{\text{soil}}$ ) and  $^{226}\text{Ra}_{\text{NAA}}$  did the wells differ from the other samples taken at 0–10 cm from the other distances. For example, the  $^{226}\text{Ra}$  values from the wells were, on average, one order of magnitude higher than  $^{226}\text{Ra}$  from other distances, while the  $^{226}\text{Ra}$  values from 0–10 cm order of magnitude higher. Both  $^{226}\text{Ra}$  and  $^{228}\text{Ra}$  were detected in soils from any distance other than 0 cm. The range of variation for the maximum value of  $^{226}\text{Ra}$  from all previous authors ( $^{226}\text{Ra}_{\text{soil}}$ ) is 0–1000 (Barro et al., 1972);  $^{226}\text{Ra}$  (Fritz et al., 1970);  $^{226}\text{Ra}$  (Liu et al., 1979) and  $^{226}\text{Ra}$  (Liu et al., 1980).

Results of the analyses of the soils confirm our initial hypothesis concerning the distribution of the radionuclides  $^{226}\text{Ra}$  and  $^{228}\text{Ra}$  concerned with distance, while the concentration of the relatively occurring radionuclides  $^{228}\text{Th}$  and  $^{228}\text{Ra}$  is relatively constant up to the depth of 10 cm (except outliers). The influence of  $^{226}\text{Ra}$  and  $^{228}\text{Ra}$  was present in the top layer of the soil profiles.

Considering only the bottom sediment (0–10 cm depth) the  $^{226}\text{Ra}_{\text{soil}}$  and  $^{226}\text{Ra}_{\text{NAA}}$  (in  $\text{Bq m}^{-2}$ ) from 0–10, 10–20, and 20–30 cm depth (bottom of sample tube) the values for these radionuclides increase from south to the north. This is to be expected in the Karpinsk and Kirovograd and are much less than the values from Bakh and Dnepropetrovsk (Boggs, 1977).

## SUMMARY AND CONCLUSIONS

This study of radionuclides in plants, soils, and maf. from Five Mile, 155 Km. N.W. of Fort Macleod, was one phase of the Pacific Geodiversity Program. This was a part of the larger part of the program to evaluate the kinds and amounts of radionuclides in biological and environmental samples from the Canadian prairies. The scientific purpose of this study was to measure the total and fractionated  $^{38}K$  found in vegetation and soil samples of  $^{38}K$  would which did not have an appreciable "potassium" from the beta of K-40 and K-40 decay.

Approximately 250 samples for this study were collected during November 1973, and 130 contained 309 strontium-90, and 67 contained 229-rhenium-186 respectively.

Results of the analyses indicate that naturally occurring  $^{38}K$  is the predominant radionuclide in the biological material. Consistently was there a fairly high radionuclide detected in most of the biological samples. A mean of 1.6% present in the maf. were usually less than 1 ppb/g of dry mass of the whole plants. No maf. had a detectable quantity of either the man-made plutonium or plutonium-239.

Soil samples from all the different areas generally contained 1 to 100 ppb/g strontium-90 and 10 to 20 ppb/g strontium-89. Below, and above, also contained less than 1 ppb/g naturally occurring fractions of radon-222, thorium-232, and uranium-238. Radon-222 and radon-226 were relatively low in the radon-free soils, and in addition contained 309 and 229-rhenium-186. Amounts of the naturally occurring radionuclides in the plants were much higher than amounts of either man-made radionuclides from the other sources.

Considering only the natural radionuclides, the values for 309, 229-rhenium-186, and 229-thorium-232 in the plants and soils are similar. Garry, Thistle, Brome, and Alfalfa were from the values for these three radionuclides. Total radionuclides for alfalfa, switchgrass, and kafir grass in the northern Barrell Hill area were from the values for these three radionuclides. Total radionuclides for alfalfa, switchgrass, and kafir grass in the southern Barrell Hill area were from the values for these three radionuclides.

and we find the following values of these additional first four angles from Table I  
and Table II above.

## REFERENCES

- Estimated incidence of land-dependent felonies in 1976. Anchorage City Sheriff's Office Report, KIA-188-AK-1-C-100, 1976.
- . 1976. Radioactivity survey and findings. RMC-RM-1, 1976, 10 pp.
- Nelson, M. R. 1977. Radioactive survey of plants, animals and the environment during and after fallout from the Marshall Islands. RMC-RM-1, University of Alaska Fairbanks, College of Fisheries, Department of Radiation Ecology, 60 pp.
- Nelson, M. R., and Thompson, A. H. 1976. Preliminary Environmental Survey Report. Juneau, Washington Public Power Supply Authority, Bureau of the US Army Corps of Engineers, USGS Service, University of Alaska, College of Fisheries, Anchorage, Alaska, 100 pp.
- . 1977. Radiological survey of Juneau, Alaska. Report January 1976 to December 1976. RMC-RM-1, University of Alaska, 1977, College of Fisheries, Laboratory of Radiation Ecology, 40 pp.

Appendix Table 1. Predominant radionuclides in fish Collected in Micronesia in November, 1975.

Table 1. (Continued)

Appendix Table 2 Some Radionuclides in Plants Collected on Majuro Atoll in November 1975.

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Appendix Table 3. Some Radionuclides in Plants Collected in Ponape District in November 1975.

THE SPECIFIC FEATURES OF THE POLYMERIZATION

b. *not* significant; the test sample count is less than the two-jinn counting error, that is not analyzed.

Appendix Table 4. Some Radionuclides in Plants Collected in Truk District in November 1975.

“*It is a good idea to have a few books on your shelf that you can read over again.*”

Appendix Table 5. Some Radionuclides in Plants Collected in the Palau Islands  
in November 1972.

The prior values are found in Fig. 1, normalized, according to Eq. (1) for a single sample.

Appendix Table 6. Some Radiociliates in Plants Collected on Guam in November 1975.

<sup>4</sup> See, e.g., *Opinion of the Justices to the Senate*, 18 U.S. 1, 10 (1803).

Table 7. Some Rachomuc Lides in Soil Collected on Majuro Atoll in November 1976.

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Appendix Table 8 . Some Radionuclides in Soil Collected in the Ponape District in November 1975.

Appendix Table 9. Some Radionuclides in Soil Collected at Truk in November 1975.

Appendix Table vi. Soils Radionuclides in Soil Collected in the Palau Islands in November 1975.

Figure 1 shows the results of the first sample collection. The figure shows the mean and standard deviation of the measured values.

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Appendix Table 11. Some Radionuclides in Soil Collected on Guam in November 1975.

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APPENDIX 4

RADIOLOGICAL SURVEY OF PLANTS, ANIMALS AND SOIL

AT FIVE ATOLLS IN THE MARSHALL ISLANDS

SEPTEMBER - OCTOBER 1970

By

John W. Nelson

WILLIAM J. NELSON

RADIOGRAPHIC IMAGE OF THE NUCLEAR EXPLOSION  
AT DAY 1000 IN THE MURKIN TEST SITE  
SOUTHERN CALIFORNIA

BY

VICTOR K. VADIM

P. E. NV

University of Washington  
College of Engineering  
Department of Radiology, School  
of Medicine, Seattle, WA

Prepared for the U. S. Department of Energy  
Nevada Operations Office under Contract No. DE-AC52-06NA2530

## INTRODUCTION

As stated in a previous publication report (Bellouin 1977),

"The Silvertonen (Specialized Laboratory) and (new Safety Standards and Compliance) portion of the International Radiation Control (IRC) and the Radiobiology Program (General Radiation Audit Program) began on July 1974 and is continuing. The objective of this program is to determine the kinds and amounts of radioactive materials distributed in the form, quantity, activity, and nature of the Central Pacific, especially the Marshall Islands, and to furnish the data in ASI/1974 and other appropriate areas. (However, if any of the laboratory, Nevada Operations Office (NO) staff, they may make an exception of the data of information received by the project giving the enclosed the Central Peacetime."

Here we report the results of the analysis of samples collected on a field trip conducted in September October 1974.

## MATERIALS AND METHODS

Areas visited in the Marshall Islands are shown in Figure 1. This trip was carried out by Mr. James L. Price, from the National Laboratory. Representative biological and soil samples were collected with emphasis on food items common to the diet of the Marshallese people (i.e., fish, coconut, papaya, breadfruit, coconut milk, etc.) in fairly mixed proportions. At these items, deer were collected and analyzed. Soils were collected to provide data for estimating future distribution and mobility of radionuclides in the environment and to do sampling of the data of Figure 1. Section 1.

The number of samples taken, given their size in soft fractions, is shown in Table 1. Slightly less than half the samples were sand-size, 5%, clayey, and loamy soils, and the rest were on the surface (0-25 cm) or on the (0-100 cm).

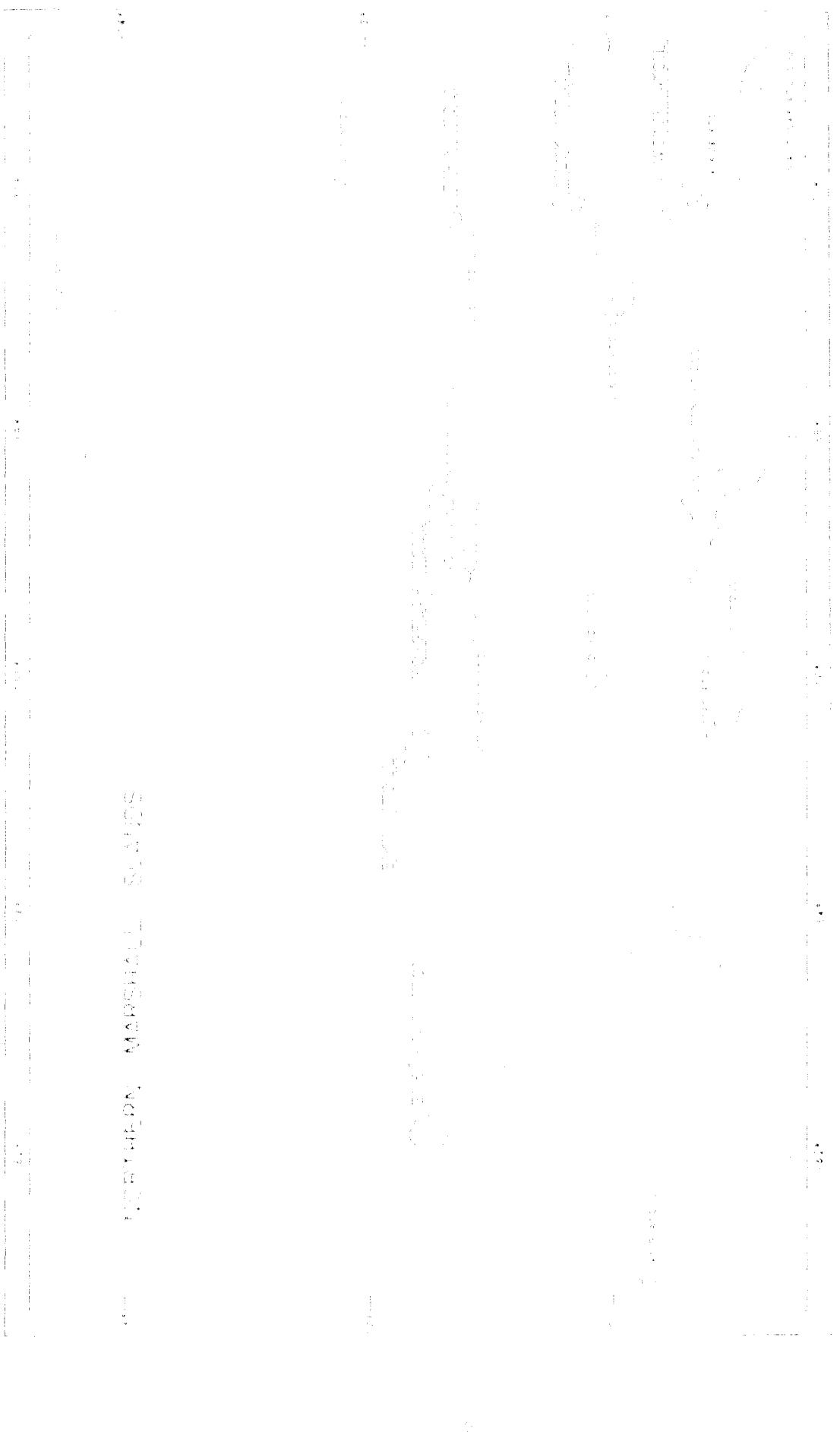
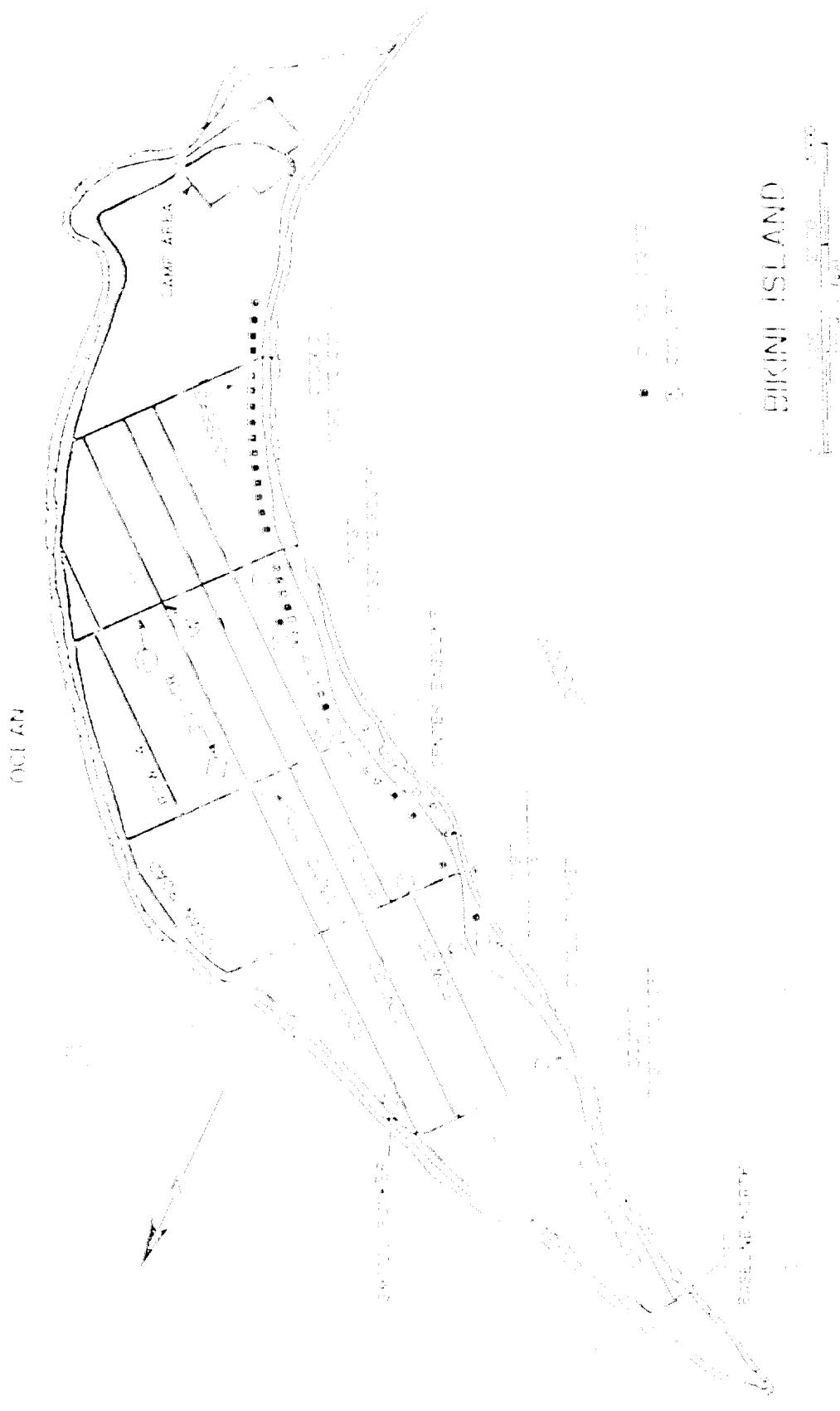


Figure 1. Five atolls (underlined) in the northern Marshall Islands, where samples were collected (solid black circles) during 1996.



Completion of Campfire sites on Västern Island, Riköni Atoll, October 1976.

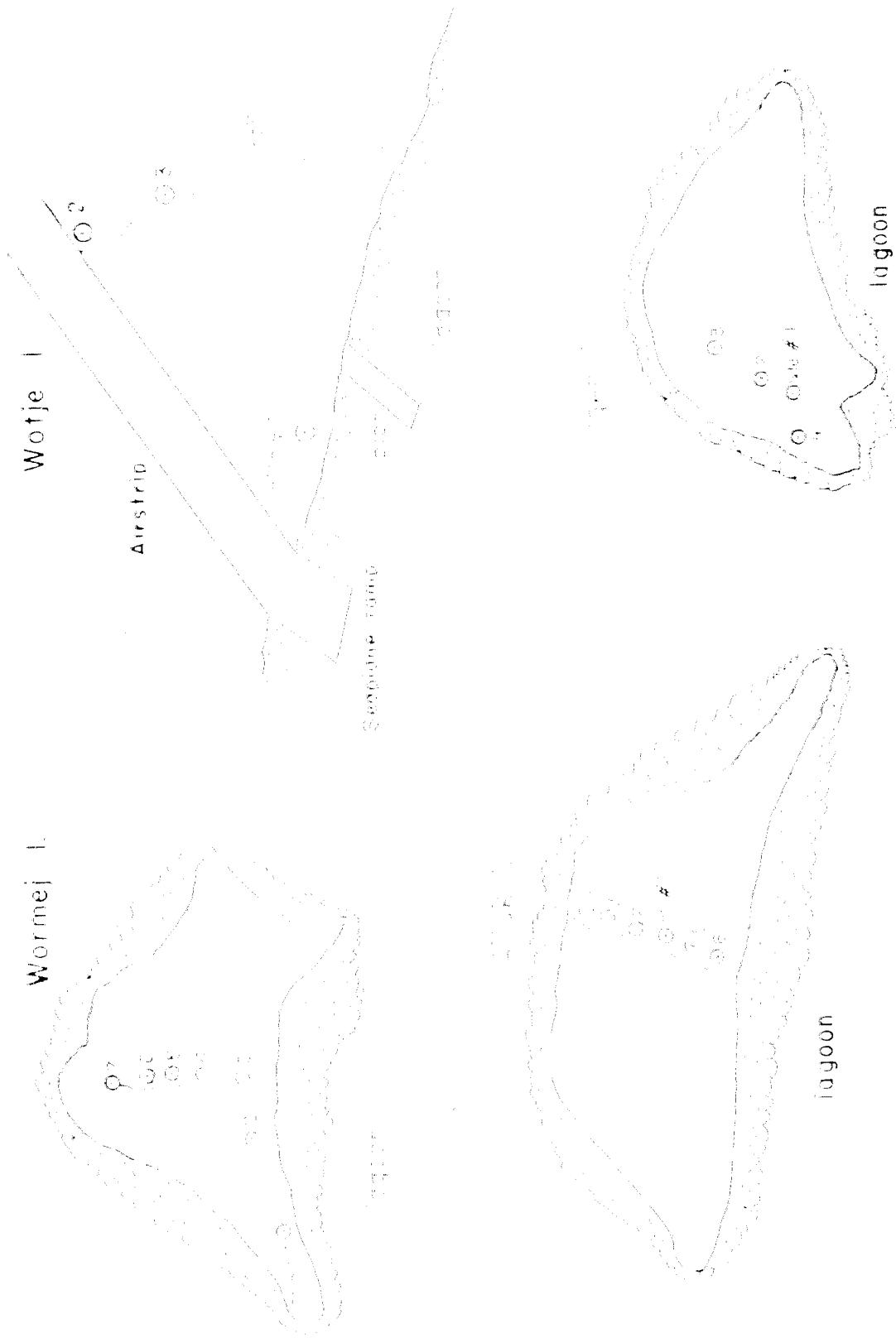
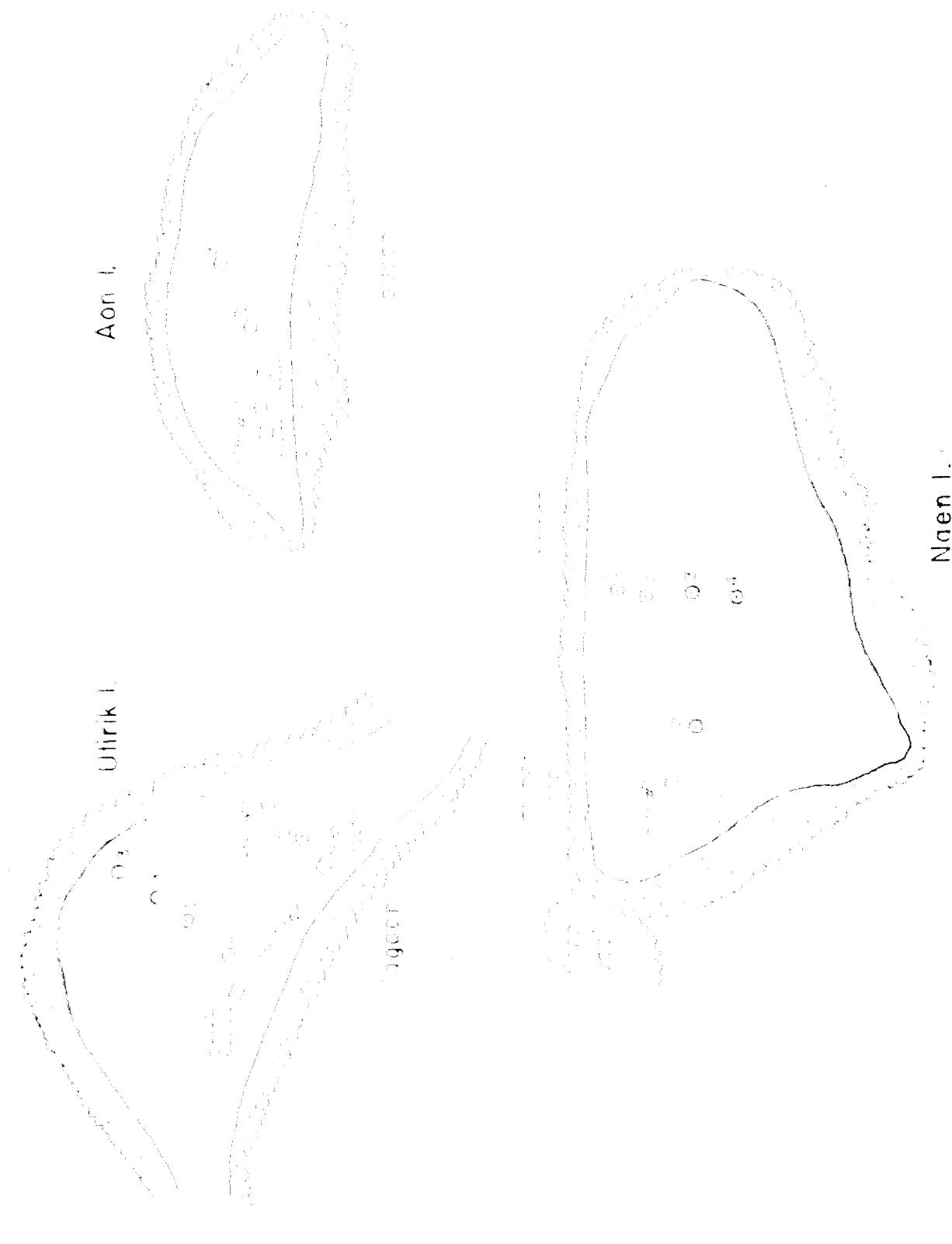


Figure 3. Sampling sites on Wotje and Wotje Islands, Wotje Atoll and Aitutak, Tuamotu Islands, French Polynesia.

**Figure 4.** Sampling Sites on Utirik and Aon Islands, Utirik Atoll and on Naen Island, Rongelap Atoll, September 1976.



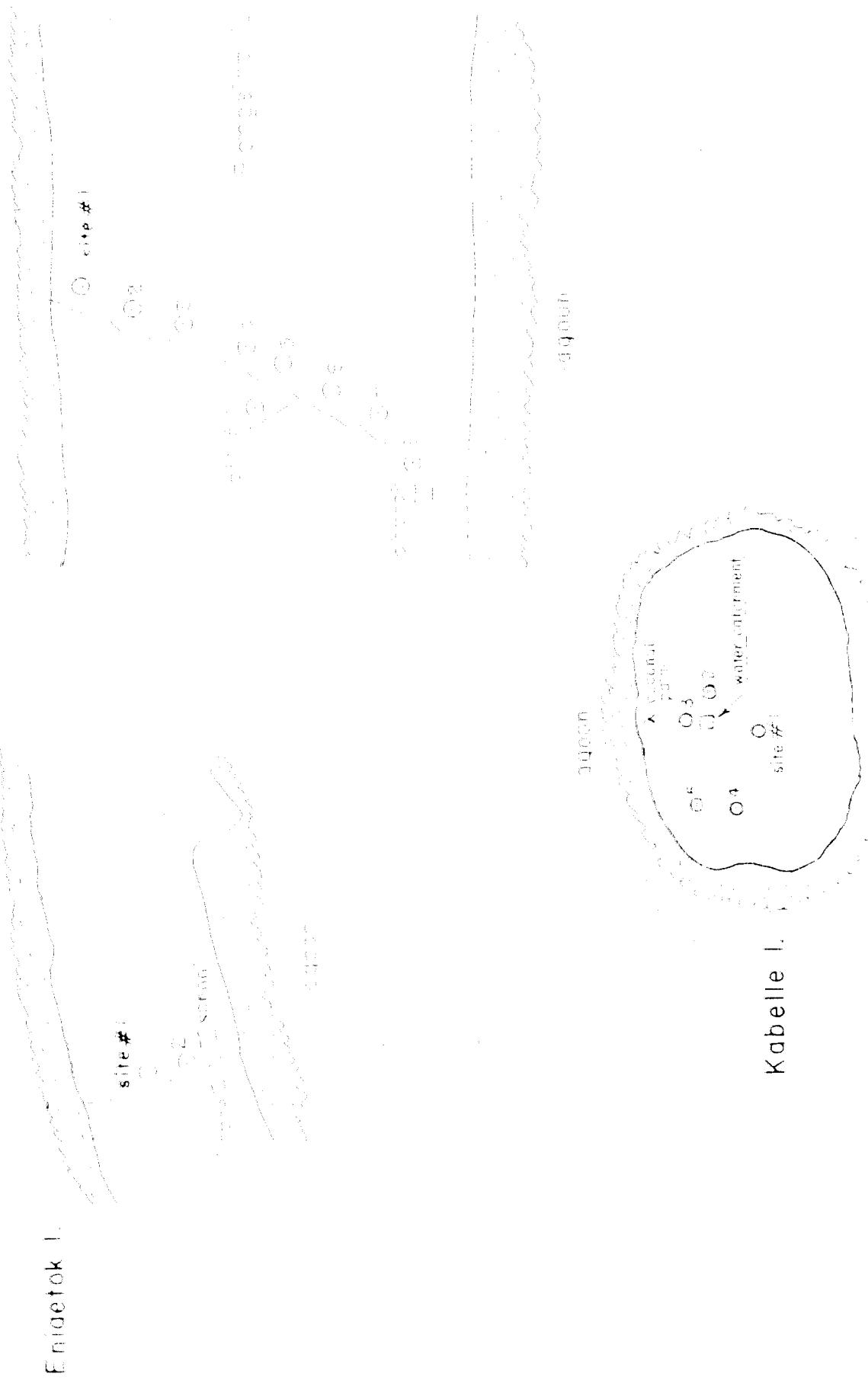


figure 5. Sampling sites on Eniaetok, Rongelap, and Kabelle Islands, Rongelap Atoll, September 1976.

Table 1. Number of samples processed and analyzed that were collected during the September 1976 field trip to the Marshfield Islands.

Sample Type	Number of Samples Collected		Number of Samples Processed		Number of Samples Analyzed	
	Number	Percent	Number	Percent	Number	Percent
Soil	100	100	100	100	100	100
Roots	100	100	100	100	100	100
Leaves	100	100	100	100	100	100
Flowers	100	100	100	100	100	100
Fruit	100	100	100	100	100	100
Stems	100	100	100	100	100	100
Whole plant	100	100	100	100	100	100
Total	600	100	600	100	600	100

In addition to the radiotracer counting, personnel from Deutsches Röntgenologisches Forschungsinstitut (DRFI) conducted an <sup>137</sup>Cs radiation survey (radioactive sodium iodide (NaI) scintillation detector) and a gamma line spectrometer. The results of the Bonnhaven analyses and each element may be compared with the 137Cs results for a series of joint reports to the local authorities.

## ANALYSIS METHODS

### Gamma Spectrometry

All of the samples were analyzed by gamma spectroscopy, either after a low background iodide (Thallium ortho-<sup>133</sup>Cs) crystal and 200-channel pulse-height analyzer or with a germanium (Germanium-doped) diode detector and 2048-channel pulse-height analyzer. Soil samples were analyzed on the GEMCO system and the plutonium samples were analyzed on path systems.

All samples were oven-dried, ground and a portion comminuted in polyethylene (PEO) film 2 inches in diameter and placed in one inch deep thin-walled sample holder for gamma-ray measurement. Fifty grams of the 100 gms of soil could be contained in the 2 x 2 inch container. Two aliquots of the biological and soil samples were 1 g and 1.5 g, respectively. These samples were then analyzed for concentrations of radionuclides.

The energy calibration and background subtraction was carried on the NaI detector with 6623 keV gamma line of Thallium-204. The calibrated dose values were then converted on the NaI(+) detector with 100% efficiency with a computer by scaling the counts down to a range of five channels and a sum in the spectrum, subtracting the appropriate background counts, and applying conversion factors to convert counts to picocuries (pCi). Sixty previously counted reference spectra for the different geometries and sample sizes were used. All values were corrected for decay to the date of collection.

## Structural-Function Relationship

To determine the structure, several methods can be separated from  $\text{^{13}C}$ , and  $\text{^{15}N}$  NMR, which filter paper and coupled with a flow system containing a flow cell. The  $\text{^{13}C}$  NMR extracted by the exchange, which is based on polarizing glass, and can be used to differentiate between which systems, which are suitable for alpha detection, and multiple carbon systems. Chemical shift difference by two or three atoms.

## Energy Inputs

For a sample complex, the energy given to an audience will be dictated by the efficiency of energy conversion in the process. The more time for more than one cycle, is considered cyclotron and decay mode, coupling energy.

## Effect of Detection

Many factors will affect the effect of detection, including the type of detection and analysis, the presence of other nuclear sides, the duration of the counting period, size, the and density of the sample, and the relationship between temperature and activity. Often, the form of detection is considerably different and uniform, but different types of samples, but can be measured by selecting the detection. And it can be approximately as follows:

By mass detection:

$$\begin{array}{ll} \text{^{13}C} & 7.1 \times 10^6 \text{ pCi/mg-Tera} \\ \text{^{15}N} & 0.41 \times 10^6 \text{ pCi/mg-Tera} \\ \text{^{36}Cl} & 0.01 \times 10^6 \text{ pCi/mg-Tera} \\ \text{^{37}Ar} & 0.07 \text{ pCi/mg-Tera} \end{array}$$

by beta detection:

$$\text{^{32}P} \quad 0.2 \text{ pCi/mg-Tera}$$

by alpha detection:

$$\text{^{232}Th} \quad 0.02 \text{ pCi/mg-Tera}$$

## RESULTS

Data are presented for the analysis of the samples of the complex collected by UIC from the Bernhard section in 1976. Appendix Table 1 contains all raw data for sample analysis. The data are first presented (Table 1) and then extracted by sampling location (Table 2), for plant material type. All data are given in picocuries per gram dry weight except where explicitly indicated.

### Plant Material

Plants from Pilgrim and Alluvium horizons in the soil and from wetland areas in the Alluvium and Alluvium-like (the Anterior Octopus) were analysed. Results of the analyses on these samples of flora, plants and soil are given and the radionuclides  $^{36}\text{Kr}$  and  $^{137}\text{Cs}^{134}\text{Cs}$  are given in Appendixes 1 (36Kr), 2 and 3 (plants) and 4 (soil).

In the 1976 sampling year, according to Kr was the most abundant radionuclide. Except for a small amount of  $^{36}\text{Kr}$  found in plants, no fallout radionuclides were identified above the other background. In plants,  $^{36}\text{Kr}$  was also the predominant radionuclide, however,  $^{36}\text{Kr}$  was above the limit of detection in all plant samples and  $^{137}\text{Cs}$  was measurable in about one third of the samples analysed. Of the plants sampled, scandium (Sc) was the most  $^{36}\text{Kr}$ -enriched measured element, followed by the metal  $^{238}\text{U}$ . Increasing the apparent timers for finding enrichment of the  $^{36}\text{Kr}$ .

Using 10% as the pre-enriched radionuclides in the soil samples from wetland and Alluvium soils, 90% for plants measured as  $^{36}\text{Kr}$  was found in all samples except four from River Alluvium, one from which sampling location on these two

studies. Concentrations of  $^{106}\text{Ru}$  in these three samples ranged from 0.1 to 1.6 pCi/g. All sulfur-30 and  $^{36}\text{Ar}^{38}\text{Kr}$  were incorporated in the samples analyzed. Concentrations were less than a picocurie in all samples except one, the sample with the highest  $^{36}\text{Ar}^{38}\text{Kr}$  found. Sulfur-34 was contained in all samples at 100% (the only sample where it was not 100% was determined by only a few of the sulfur analyses).

## RESULTS

The two previously identified sites in plant samples (Appendix Table I) were  $^{137}\text{Cs}$  and  $^{138}\text{Cs}$  (Appendix Table II). Of the samples analyzed, the radioactive portion of the three samples found could be attributed to the precipitation of  $^{137}\text{Cs}$  (average 14.0%). Values for  $^{137}\text{Cs}$  in the plants ranged up to 2.125 pCi/g. Plutonium-239 values were below the limits of detection in the five samples analyzed.

Sulfur concentrations in the plants contained  $^{34}\text{S}$ ,  $^{36}\text{S}$ , and  $^{38}\text{S}$  with the total of the samples analyzed for these radionuclides (Appendix Tables I and II, Appendix III) and the results collected in Appendix III. The sulfur concentration ranged up to 6.3 mol/g, averaging about 3 mol/g in the surface soils (samples 1 and 2) before the surface crusts of 10 to 100 mm thickness were removed. In the surface samples analyzed,  $^{34}\text{S}$  was a major component (up to 0.9 mol/g sulfur), whereas  $^{36}\text{S}$  was a minor component (0.1 to 0.2 mol/g sulfur) and  $^{38}\text{S}$  was present in amounts less than 0.1 mol/g.

## DISCUSSION

Most samples of marine organisms from Florida contained  $^{137}\text{Cs}$  and  $^{138}\text{Cs}$  with the exception of a conch shell (Appendix Table I and II). Collected samples ranged from 0.01 pCi/g (mullet) except for the 1979 red snapper which contained 0.07 pCi/g. Ranges of concentrations of the catched fish samples are consistent with those of the predominantly addressed finfish species (Appendix Table I and II). See in the column  $\delta^{34}\text{S}$  (Appendix Table I) the  $\delta^{34}\text{S}$  of the samples containing

to 100 ppb of  $^{238}\text{U}$  per gram of dry sediment, however, sediment in all samples contained up to 100 ppb of  $^{238}\text{U}$  per gram. The other filters contained 7 to 20 ppb of

the residue of the analysis of the soil samples are given in Appendix Tables 14 through 17. Concentrations of  $^{238}\text{U}$  and  $^{232}\text{Th}$  were also converted to and expressed in  $\mu\text{Ci/g}$ ,  $\text{Bq/g}$ ,  $\text{ppm}$ , and  $\text{ppb}$  and were also converted to and expressed in  $\text{Bq/g}$ . The sediments analyzed in these specific sediment sites (Antimony 125 to 127) were taken in the soils from New Bedford. Radon-222 values were found to be relatively independent of depth and ranged on Rm 1 (Soil 127) and 2 (Soil 126) from 400 to 1000  $\text{Bq/m}^2/\text{day}$  (soils 127 and 126 respectively). Concentrations of  $^{226}\text{Ra}$  taken from Rm 127 and 126 from New Bedford were less than a hundred ppb/g. Plutonium-238, -239, and -240 values in dredge 126 and 127 were commonly 1 to 5 ppb/g, while in Rm 126 values for these radionuclides ranged to 6 ppb/g.

#### Other Areas

Plants were the only sample type collected at Block Island, Rm 100. The analyses of these samples are in Appendix Table 18. Concentrations of  $^{238}\text{U}$  and  $^{232}\text{Th}$  are the same as previously described. The greatest U238 and Th232 values are in the leaves of the plants. Leaves from a plant near off-shore (Soil 128) contained 2000 ppb of  $^{238}\text{U}$  per gram while leaves from a plant near shore (Soil 129) contained 200 ppb of  $^{238}\text{U}$  per gram. Plutonium-238, -239, and -240 values were very low (concentrations of detection).

### DISCUSSION AND CONCLUSION

#### Concentration Trends

Having in mind the generally increasing  $\text{Ra}_{226}$  values were least in the samples from dredge 127, increased slightly on Rm 100 (soil) and increased significantly at Rm 126. The U238/U232 ratios of dry values increased to the west,

2.5 ft. thick. The sand is yellowish brown, fine-grained, and contains many small pebbles.

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collected by the Rennell Islands. A field report conducted for this project in September-October 1976 (Nestor 1978) which I have collected are about 21% and 19% *Leptosperma* 30% and 19% *Phormium* 22% 24% and 20% were present, 10% *Cordyline* and 5% *Acacia*. Other flora and 3% *C. tenuissima* are predominant in the northern island community and 1% *Acacia*, 10% *Leptosperma* and 7% *Phormium* are represented in the southern island community. *Leptosperma* 40% is the predominant species in the northern community while 10% is dominant in the majority of the southern sites.

Amounts of endemism vary between islands and between islands within Rennell Province vary with distance from the main island "Maine Atoll" and in relation to the elevation position from the March 1976 survey. Plants from Bikendrik Island had the highest amounts of endemism, with 50% *Leptosperma* plants from the island at Renell Province being endemic to the island. The southern islands off "Maine Atoll" are 10% *Maine Atoll* endemic to moderate amounts of endemism while Bikendrik Island plants had the lowest amount of endemism of 5% at the survey during the 1976 trip.

#### REFERENCES

- Bellows, M. A. 1977. A floristic survey of plants, animals, and ecology of three island and seven Atolls in the Rennell Islands, H. S. - 500 Report No. 1977. College of Fisheries, University of Washington, Seattle.

A P P E N D I X I A B I L I  
Some Radionuclides in Fish Collected at Rongelap,  
Atoll, and Etoe Atolls in September 1976

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A P P E N D I C E 2

Predominant Radionuclides in Plants Collected at Eloway [sic] in September 1947

the first time in the history of the world, the people of the United States have been compelled to make a choice between two political parties.

by the present author in Plants collected in Africa, also in self culture.

Kaufmann's Little Folio Encyclopedia, Vol. I, Part One

APPENDIX I TABLE A

Predominant Radionic Lichen, in Soil Collected on  
Mormo Island, Baitie Atoll, September 1976

Philosophical Contention in India, 1

**APPENDIX TABLE 5**  
Soil respiration rates in soil collected on  
active Islands, Native sites, and Control plots

Site	Depth (cm)	Mean $R_s$ (mg m <sup>-2</sup> h <sup>-1</sup> )		Mean $R_s$ (mg m <sup>-2</sup> h <sup>-1</sup> )	Mean $R_s$ (mg m <sup>-2</sup> h <sup>-1</sup> )
		Native	Control		
1	0-10	1.0	1.0	1.0	1.0
1	10-20	1.0	1.0	1.0	1.0
1	20-30	1.0	1.0	1.0	1.0
1	30-40	1.0	1.0	1.0	1.0
1	40-50	1.0	1.0	1.0	1.0
1	50-60	1.0	1.0	1.0	1.0
1	60-70	1.0	1.0	1.0	1.0
1	70-80	1.0	1.0	1.0	1.0
1	80-90	1.0	1.0	1.0	1.0
1	90-100	1.0	1.0	1.0	1.0
1	100-110	1.0	1.0	1.0	1.0
1	110-120	1.0	1.0	1.0	1.0
1	120-130	1.0	1.0	1.0	1.0
1	130-140	1.0	1.0	1.0	1.0
1	140-150	1.0	1.0	1.0	1.0
1	150-160	1.0	1.0	1.0	1.0
1	160-170	1.0	1.0	1.0	1.0
1	170-180	1.0	1.0	1.0	1.0
1	180-190	1.0	1.0	1.0	1.0
1	190-200	1.0	1.0	1.0	1.0
1	200-210	1.0	1.0	1.0	1.0
1	210-220	1.0	1.0	1.0	1.0
1	220-230	1.0	1.0	1.0	1.0
1	230-240	1.0	1.0	1.0	1.0
1	240-250	1.0	1.0	1.0	1.0
1	250-260	1.0	1.0	1.0	1.0
1	260-270	1.0	1.0	1.0	1.0
1	270-280	1.0	1.0	1.0	1.0
1	280-290	1.0	1.0	1.0	1.0
1	290-300	1.0	1.0	1.0	1.0
1	300-310	1.0	1.0	1.0	1.0
1	310-320	1.0	1.0	1.0	1.0
1	320-330	1.0	1.0	1.0	1.0
1	330-340	1.0	1.0	1.0	1.0
1	340-350	1.0	1.0	1.0	1.0
1	350-360	1.0	1.0	1.0	1.0
1	360-370	1.0	1.0	1.0	1.0
1	370-380	1.0	1.0	1.0	1.0
1	380-390	1.0	1.0	1.0	1.0
1	390-400	1.0	1.0	1.0	1.0
1	400-410	1.0	1.0	1.0	1.0
1	410-420	1.0	1.0	1.0	1.0
1	420-430	1.0	1.0	1.0	1.0
1	430-440	1.0	1.0	1.0	1.0
1	440-450	1.0	1.0	1.0	1.0
1	450-460	1.0	1.0	1.0	1.0
1	460-470	1.0	1.0	1.0	1.0
1	470-480	1.0	1.0	1.0	1.0
1	480-490	1.0	1.0	1.0	1.0
1	490-500	1.0	1.0	1.0	1.0
1	500-510	1.0	1.0	1.0	1.0
1	510-520	1.0	1.0	1.0	1.0
1	520-530	1.0	1.0	1.0	1.0
1	530-540	1.0	1.0	1.0	1.0
1	540-550	1.0	1.0	1.0	1.0
1	550-560	1.0	1.0	1.0	1.0
1	560-570	1.0	1.0	1.0	1.0
1	570-580	1.0	1.0	1.0	1.0
1	580-590	1.0	1.0	1.0	1.0
1	590-600	1.0	1.0	1.0	1.0
1	600-610	1.0	1.0	1.0	1.0
1	610-620	1.0	1.0	1.0	1.0
1	620-630	1.0	1.0	1.0	1.0
1	630-640	1.0	1.0	1.0	1.0
1	640-650	1.0	1.0	1.0	1.0
1	650-660	1.0	1.0	1.0	1.0
1	660-670	1.0	1.0	1.0	1.0
1	670-680	1.0	1.0	1.0	1.0
1	680-690	1.0	1.0	1.0	1.0
1	690-700	1.0	1.0	1.0	1.0
1	700-710	1.0	1.0	1.0	1.0
1	710-720	1.0	1.0	1.0	1.0
1	720-730	1.0	1.0	1.0	1.0
1	730-740	1.0	1.0	1.0	1.0
1	740-750	1.0	1.0	1.0	1.0
1	750-760	1.0	1.0	1.0	1.0
1	760-770	1.0	1.0	1.0	1.0
1	770-780	1.0	1.0	1.0	1.0
1	780-790	1.0	1.0	1.0	1.0
1	790-800	1.0	1.0	1.0	1.0
1	800-810	1.0	1.0	1.0	1.0
1	810-820	1.0	1.0	1.0	1.0
1	820-830	1.0	1.0	1.0	1.0
1	830-840	1.0	1.0	1.0	1.0
1	840-850	1.0	1.0	1.0	1.0
1	850-860	1.0	1.0	1.0	1.0
1	860-870	1.0	1.0	1.0	1.0
1	870-880	1.0	1.0	1.0	1.0
1	880-890	1.0	1.0	1.0	1.0
1	890-900	1.0	1.0	1.0	1.0
1	900-910	1.0	1.0	1.0	1.0
1	910-920	1.0	1.0	1.0	1.0
1	920-930	1.0	1.0	1.0	1.0
1	930-940	1.0	1.0	1.0	1.0
1	940-950	1.0	1.0	1.0	1.0
1	950-960	1.0	1.0	1.0	1.0
1	960-970	1.0	1.0	1.0	1.0
1	970-980	1.0	1.0	1.0	1.0
1	980-990	1.0	1.0	1.0	1.0
1	990-1000	1.0	1.0	1.0	1.0
2	0-10	1.0	1.0	1.0	1.0
2	10-20	1.0	1.0	1.0	1.0
2	20-30	1.0	1.0	1.0	1.0
2	30-40	1.0	1.0	1.0	1.0
2	40-50	1.0	1.0	1.0	1.0
2	50-60	1.0	1.0	1.0	1.0
2	60-70	1.0	1.0	1.0	1.0
2	70-80	1.0	1.0	1.0	1.0
2	80-90	1.0	1.0	1.0	1.0
2	90-100	1.0	1.0	1.0	1.0
2	100-110	1.0	1.0	1.0	1.0
2	110-120	1.0	1.0	1.0	1.0
2	120-130	1.0	1.0	1.0	1.0
2	130-140	1.0	1.0	1.0	1.0
2	140-150	1.0	1.0	1.0	1.0
2	150-160	1.0	1.0	1.0	1.0
2	160-170	1.0	1.0	1.0	1.0
2	170-180	1.0	1.0	1.0	1.0
2	180-190	1.0	1.0	1.0	1.0
2	190-200	1.0	1.0	1.0	1.0
2	200-210	1.0	1.0	1.0	1.0
2	210-220	1.0	1.0	1.0	1.0
2	220-230	1.0	1.0	1.0	1.0
2	230-240	1.0	1.0	1.0	1.0
2	240-250	1.0	1.0	1.0	1.0
2	250-260	1.0	1.0	1.0	1.0
2	260-270	1.0	1.0	1.0	1.0
2	270-280	1.0	1.0	1.0	1.0
2	280-290	1.0	1.0	1.0	1.0
2	290-300	1.0	1.0	1.0	1.0
2	300-310	1.0	1.0	1.0	1.0
2	310-320	1.0	1.0	1.0	1.0
2	320-330	1.0	1.0	1.0	1.0
2	330-340	1.0	1.0	1.0	1.0
2	340-350	1.0	1.0	1.0	1.0
2	350-360	1.0	1.0	1.0	1.0
2	360-370	1.0	1.0	1.0	1.0
2	370-380	1.0	1.0	1.0	1.0
2	380-390	1.0	1.0	1.0	1.0
2	390-400	1.0	1.0	1.0	1.0
2	400-410	1.0	1.0	1.0	1.0
2	410-420	1.0	1.0	1.0	1.0
2	420-430	1.0	1.0	1.0	1.0
2	430-440	1.0	1.0	1.0	1.0
2	440-450	1.0	1.0	1.0	1.0
2	450-460	1.0	1.0	1.0	1.0
2	460-470	1.0	1.0	1.0	1.0
2	470-480	1.0	1.0	1.0	1.0
2	480-490	1.0	1.0	1.0	1.0
2	490-500	1.0	1.0	1.0	1.0
2	500-510	1.0	1.0	1.0	1.0
2	510-520	1.0	1.0	1.0	1.0
2	520-530	1.0	1.0	1.0	1.0
2	530-540	1.0	1.0	1.0	1.0
2	540-550	1.0	1.0	1.0	1.0
2	550-560	1.0	1.0	1.0	1.0
2	560-570	1.0	1.0	1.0	1.0
2	570-580	1.0	1.0	1.0	1.0
2	580-590	1.0	1.0	1.0	1.0
2	590-600	1.0	1.0	1.0	1.0
2	600-610	1.0	1.0	1.0	1.0
2	610-620	1.0	1.0	1.0	1.0
2	620-630	1.0	1.0	1.0	1.0
2	630-640	1.0	1.0	1.0	1.0
2	640-650	1.0	1.0	1.0	1.0
2	650-660	1.0	1.0	1.0	1.0
2	660-670	1.0	1.0	1.0	1.0
2	670-680	1.0	1.0	1.0	1.0
2	680-690	1.0	1.0	1.0	1.0
2	690-700	1.0	1.0	1.0	1.0
2	700-710	1.0	1.0	1.0	1.0
2	710-720	1.0	1.0	1.0	1.0
2	720-730	1.0	1.0	1.0	1.0
2	730-740	1.0	1.0	1.0	1.0
2	740-750	1.0	1.0	1.0	1.0
2	750-760	1.0	1.0	1.0	1.0
2	760-770	1.0	1.0	1.0	1.0
2	770-780	1.0	1.0	1.0	1.0
2	780-790	1.0	1.0	1.0	1.0
2	790-800	1.0	1.0	1.0	1.0
2	800-810	1.0	1.0	1.0	1.0
2	810-820	1.0	1.0	1.0	1.0
2	820-830	1.0	1.0	1.0	1.0
2	830-840	1.0	1.0	1.0	1.0
2	840-850	1.0	1.0	1.0	1.0
2	850-860	1.0	1.0	1.0	1.0
2	860-870	1.0	1.0	1.0	1.0
2	870-880	1.0	1.0	1.0	1.0
2	880-890	1.0	1.0	1.0	1.0
2	890-900	1.0	1.0	1.0	1.0
2	900-910	1.0	1.0	1.0	1.0
2	910-920	1.0	1.0	1.0	1.0
2	920-930	1.0	1.0	1.0	1.0
2	930-940	1.0	1.0	1.0	1.0
2	940-950	1.0	1.0	1.0	1.0
2	950-960	1.0	1.0	1.0	1.0
2	960-970	1.0	1.0	1.0	1.0

A VARIATION IN THE  $\delta^{34}\text{S}$ 

From Redox-Precipitated Sulfide in Acidic Groundwater  
Arthur Klink, September 1976

Collection Site	Depth (m)	Redox-precipitate Concentration ( $\mu\text{g/g}$ )			
		$\text{Fe}^{2+}$	$\text{Mn}^{2+}$	$\text{Zn}^{2+}$	$\text{Pb}^{2+}$
Site #1	0-2.5	0.48±0.06	0.12±0.04	0.05±0.02	0.77±0.07
	2.5-5	0.28±0.03	0.08±0.02	0.03±0.01	
	5-10	0.28±0.03	0.08±0.02	0.03±0.01	
	10-15	0.16±0.02	0.03±0.01	0.01±0.005	
	15-20	0.07±0.01	0.02±0.01	0.01±0.005	
	20-25	0.05±0.01	0.02±0.01	0.01±0.005	
	25-30	0.03±0.01	0.01±0.005	0.01±0.005	
	30-35	0.02±0.01	0.01±0.005	0.01±0.005	
	35-40	0.01±0.005	0.01±0.005	0.01±0.005	
	40-45	0.01±0.005	0.01±0.005	0.01±0.005	
	45-50	0.01±0.005	0.01±0.005	0.01±0.005	
Site #2	0-2.5	0.13±0.03	0.04±0.01	0.01±0.005	
	2.5-5	0.07±0.02	0.02±0.01	0.01±0.005	
	5-10	0.05±0.01	0.02±0.01	0.01±0.005	
Site #3	0-2.5	0.22±0.04	0.05±0.02	0.02±0.01	
	2.5-5	0.09±0.02	0.03±0.01	0.01±0.005	
	5-10	0.07±0.02	0.03±0.01	0.01±0.005	
Site #4	0-2.5	0.09±0.02	0.03±0.01	0.01±0.005	
	2.5-5	0.05±0.01	0.02±0.01	0.01±0.005	
	5-10	0.04±0.01	0.02±0.01	0.01±0.005	
Site #5	0-2.5	0.09±0.02	0.03±0.01	0.01±0.005	
	2.5-5	0.05±0.01	0.02±0.01	0.01±0.005	
	5-10	0.04±0.01	0.02±0.01	0.01±0.005	
Site #6	0-2.5	0.11±0.03	0.04±0.02	0.02±0.01	
	2.5-5	0.06±0.02	0.03±0.01	0.01±0.005	
	5-10	0.05±0.02	0.03±0.01	0.01±0.005	
Site #7	0-2.5	0.12±0.03	0.04±0.02	0.02±0.01	
	2.5-5	0.07±0.02	0.03±0.01	0.01±0.005	
	5-10	0.06±0.02	0.03±0.01	0.01±0.005	
Site #8	0-2.5	0.13±0.03	0.05±0.02	0.03±0.01	
	2.5-5	0.08±0.02	0.04±0.01	0.02±0.01	
	5-10	0.07±0.02	0.04±0.01	0.02±0.01	
Site #9	0-2.5	0.14±0.03	0.06±0.02	0.03±0.01	
	2.5-5	0.09±0.02	0.05±0.01	0.02±0.01	
	5-10	0.08±0.02	0.05±0.01	0.02±0.01	
Site #10	0-2.5	0.15±0.03	0.07±0.02	0.04±0.01	
	2.5-5	0.10±0.02	0.06±0.01	0.03±0.01	
	5-10	0.09±0.02	0.06±0.01	0.03±0.01	
Site #11	0-2.5	0.16±0.03	0.08±0.02	0.05±0.01	
	2.5-5	0.11±0.02	0.07±0.01	0.04±0.01	
	5-10	0.10±0.02	0.07±0.01	0.04±0.01	
Site #12	0-2.5	0.17±0.03	0.09±0.02	0.06±0.01	
	2.5-5	0.12±0.02	0.08±0.01	0.05±0.01	
	5-10	0.11±0.02	0.08±0.01	0.05±0.01	
Site #13	0-2.5	0.18±0.03	0.10±0.02	0.07±0.01	
	2.5-5	0.13±0.02	0.09±0.01	0.06±0.01	
	5-10	0.12±0.02	0.09±0.01	0.06±0.01	
Site #14	0-2.5	0.19±0.03	0.11±0.02	0.08±0.01	
	2.5-5	0.14±0.02	0.10±0.01	0.07±0.01	
	5-10	0.13±0.02	0.10±0.01	0.07±0.01	
Site #15	0-2.5	0.20±0.03	0.12±0.02	0.09±0.01	
	2.5-5	0.15±0.02	0.11±0.01	0.08±0.01	
	5-10	0.14±0.02	0.11±0.01	0.08±0.01	
Site #16	0-2.5	0.21±0.03	0.13±0.02	0.10±0.01	
	2.5-5	0.16±0.02	0.12±0.01	0.09±0.01	
	5-10	0.15±0.02	0.12±0.01	0.09±0.01	
Site #17	0-2.5	0.22±0.03	0.14±0.02	0.11±0.01	
	2.5-5	0.17±0.02	0.13±0.01	0.10±0.01	
	5-10	0.16±0.02	0.13±0.01	0.10±0.01	
Site #18	0-2.5	0.23±0.03	0.15±0.02	0.12±0.01	
	2.5-5	0.18±0.02	0.14±0.01	0.11±0.01	
	5-10	0.17±0.02	0.14±0.01	0.11±0.01	
Site #19	0-2.5	0.24±0.03	0.16±0.02	0.13±0.01	
	2.5-5	0.19±0.02	0.15±0.01	0.12±0.01	
	5-10	0.18±0.02	0.15±0.01	0.12±0.01	
Site #20	0-2.5	0.25±0.03	0.17±0.02	0.14±0.01	
	2.5-5	0.20±0.02	0.16±0.01	0.13±0.01	
	5-10	0.19±0.02	0.16±0.01	0.13±0.01	
Site #21	0-2.5	0.26±0.03	0.18±0.02	0.15±0.01	
	2.5-5	0.21±0.02	0.17±0.01	0.14±0.01	
	5-10	0.20±0.02	0.17±0.01	0.14±0.01	
Site #22	0-2.5	0.27±0.03	0.19±0.02	0.16±0.01	
	2.5-5	0.22±0.02	0.18±0.01	0.15±0.01	
	5-10	0.21±0.02	0.18±0.01	0.15±0.01	
Site #23	0-2.5	0.28±0.03	0.20±0.02	0.17±0.01	
	2.5-5	0.23±0.02	0.19±0.01	0.16±0.01	
	5-10	0.22±0.02	0.19±0.01	0.16±0.01	
Site #24	0-2.5	0.29±0.03	0.21±0.02	0.18±0.01	
	2.5-5	0.24±0.02	0.20±0.01	0.17±0.01	
	5-10	0.23±0.02	0.20±0.01	0.17±0.01	
Site #25	0-2.5	0.30±0.03	0.22±0.02	0.19±0.01	
	2.5-5	0.25±0.02	0.21±0.01	0.18±0.01	
	5-10	0.24±0.02	0.21±0.01	0.18±0.01	
Site #26	0-2.5	0.31±0.03	0.23±0.02	0.20±0.01	
	2.5-5	0.26±0.02	0.22±0.01	0.19±0.01	
	5-10	0.25±0.02	0.22±0.01	0.19±0.01	
Site #27	0-2.5	0.32±0.03	0.24±0.02	0.21±0.01	
	2.5-5	0.27±0.02	0.23±0.01	0.20±0.01	
	5-10	0.26±0.02	0.23±0.01	0.20±0.01	
Site #28	0-2.5	0.33±0.03	0.25±0.02	0.22±0.01	
	2.5-5	0.28±0.02	0.24±0.01	0.21±0.01	
	5-10	0.27±0.02	0.24±0.01	0.21±0.01	
Site #29	0-2.5	0.34±0.03	0.26±0.02	0.23±0.01	
	2.5-5	0.29±0.02	0.25±0.01	0.22±0.01	
	5-10	0.28±0.02	0.25±0.01	0.22±0.01	
Site #30	0-2.5	0.35±0.03	0.27±0.02	0.24±0.01	
	2.5-5	0.30±0.02	0.26±0.01	0.23±0.01	
	5-10	0.29±0.02	0.26±0.01	0.23±0.01	
Site #31	0-2.5	0.36±0.03	0.28±0.02	0.25±0.01	
	2.5-5	0.31±0.02	0.27±0.01	0.24±0.01	
	5-10	0.30±0.02	0.27±0.01	0.24±0.01	
Site #32	0-2.5	0.37±0.03	0.29±0.02	0.26±0.01	
	2.5-5	0.32±0.02	0.28±0.01	0.25±0.01	
	5-10	0.31±0.02	0.28±0.01	0.25±0.01	
Site #33	0-2.5	0.38±0.03	0.30±0.02	0.27±0.01	
	2.5-5	0.33±0.02	0.29±0.01	0.26±0.01	
	5-10	0.32±0.02	0.29±0.01	0.26±0.01	
Site #34	0-2.5	0.39±0.03	0.31±0.02	0.28±0.01	
	2.5-5	0.34±0.02	0.30±0.01	0.27±0.01	
	5-10	0.33±0.02	0.30±0.01	0.27±0.01	
Site #35	0-2.5	0.40±0.03	0.32±0.02	0.29±0.01	
	2.5-5	0.35±0.02	0.31±0.01	0.28±0.01	
	5-10	0.34±0.02	0.31±0.01	0.28±0.01	
Site #36	0-2.5	0.41±0.03	0.33±0.02	0.30±0.01	
	2.5-5	0.36±0.02	0.32±0.01	0.29±0.01	
	5-10	0.35±0.02	0.32±0.01	0.29±0.01	
Site #37	0-2.5	0.42±0.03	0.34±0.02	0.31±0.01	
	2.5-5	0.37±0.02	0.33±0.01	0.30±0.01	
	5-10	0.36±0.02	0.33±0.01	0.30±0.01	
Site #38	0-2.5	0.43±0.03	0.35±0.02	0.32±0.01	
	2.5-5	0.38±0.02	0.34±0.01	0.31±0.01	
	5-10	0.37±0.02	0.34±0.01	0.31±0.01	
Site #39	0-2.5	0.44±0.03	0.36±0.02	0.33±0.01	
	2.5-5	0.39±0.02	0.35±0.01	0.32±0.01	
	5-10	0.38±0.02	0.35±0.01	0.32±0.01	
Site #40	0-2.5	0.45±0.03	0.37±0.02	0.34±0.01	
	2.5-5	0.40±0.02	0.36±0.01	0.33±0.01	
	5-10	0.39±0.02	0.36±0.01	0.33±0.01	
Site #41	0-2.5	0.46±0.03	0.38±0.02	0.35±0.01	
	2.5-5	0.41±0.02	0.37±0.01	0.34±0.01	
	5-10	0.40±0.02	0.37±0.01	0.34±0.01	
Site #42	0-2.5	0.47±0.03	0.39±0.02	0.36±0.01	
	2.5-5	0.42±0.02	0.38±0.01	0.35±0.01	
	5-10	0.41±0.02	0.38±0.01	0.35±0.01	
Site #43	0-2.5	0.48±0.03	0.40±0.02	0.37±0.01	
	2.5-5	0.43±0.02	0.39±0.01	0.36±0.01	
	5-10	0.42±0.02	0.39±0.01	0.36±0.01	
Site #44	0-2.5	0.49±0.03	0.41±0.02	0.38±0.01	
	2.5-5	0.44±0.02	0.40±0.01	0.37±0.01	
	5-10	0.43±0.02	0.40±0.01	0.37±0.01	
Site #45	0-2.5	0.50±0.03	0.42±0.02	0.39±0.01	
	2.5-5	0.45±0.02	0.41±0.01	0.38±0.01	
	5-10	0.44±0.02	0.41±0.01	0.38±0.01	
Site #46	0-2.5	0.51±0.03	0.43±0.02	0.40±0.01	
	2.5-5	0.46±0.02	0.42±0.01	0.39±0.01	
	5-10	0.45±0.02	0.42±0.01	0.39±0.01	
Site #47	0-2.5	0.52±0.03	0.44±0.02	0.41±0.01	
	2.5-5	0.47±0.02	0.43±0.01	0.40±0.01	
	5-10	0.46±0.02	0.43±0.01	0.40±0.01	
Site #48	0-2.5	0.53±0.03	0.45±0.02	0.42±0.01	
	2.5-5	0.48±0.02	0.44±0.01	0.41±0.01	
	5-10	0.47±0.02	0.44±0.01	0.41±0.01	
Site #49	0-2.5	0.54±0.03	0.46±0.02	0.43±0.01	
	2.5-5	0.49±0.02	0.45±0.01	0.42±0.01	
	5-10	0.48±0.02	0.45±0.01	0.42±0.01	
Site #50	0-2.5	0.55±0.03	0.47±0.02	0.44±0.01	
	2.5-5	0.50±0.02	0.46±0.01	0.43±0.01	
	5-10	0.49±0.02	0.46±0.01	0.43±0.01	
Site #51	0-2.5	0.56±0.03	0.48±0.02	0.45±0.01	
	2.5-5	0.51±0.02	0.47±0.01	0.44±0.01	

## 6.1.1. Results for $\lambda = 0.001$

Exact Radon transform with left-hand or right-hand  
Riesz kernels (Appendix A).

Condition	$\ f\ _2$ L <sup>2</sup> ( $\mathbb{R}^d$ )	Radon transform in $\mathbb{R}^d/\mathbb{R} \mathbf{e}_d$		
		$\ f\ _{L^2}$	$\ f\ _{L^1}$	$\ f\ _{L^\infty}$
Step 1	0.75	1.27(0.16)	0.75	0.36(0.02)
Step 2	0.75	0.70(0.01)	0.75	0.36(0.02)
Step 3	0.75	0.53(0.01)	0.53(0.01)	0.36(0.02)
Step 4	0.75	0.39(0.00)	0.39(0.00)	0.36(0.02)
Step 5	0.75	0.27(0.00)	0.27(0.00)	0.36(0.02)
Step 6	0.75	0.20(0.00)	0.20(0.00)	0.36(0.02)
Step 7	0.75	0.15(0.00)	0.15(0.00)	0.36(0.02)
Step 8	0.75	0.12(0.00)	0.12(0.00)	0.36(0.02)
Step 9	0.75	0.10(0.00)	0.10(0.00)	0.36(0.02)
Step 10	0.75	0.08(0.00)	0.08(0.00)	0.36(0.02)
Step 11	0.75	0.06(0.00)	0.06(0.00)	0.36(0.02)
Step 12	0.75	0.04(0.00)	0.04(0.00)	0.36(0.02)
Step 13	0.75	0.03(0.00)	0.03(0.00)	0.36(0.02)
Step 14	0.75	0.02(0.00)	0.02(0.00)	0.36(0.02)
Step 15	0.75	0.01(0.00)	0.01(0.00)	0.36(0.02)
Step 16	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 17	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 18	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 19	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 20	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 21	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 22	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 23	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 24	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 25	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 26	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 27	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 28	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 29	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 30	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 31	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 32	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 33	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 34	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 35	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 36	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 37	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 38	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 39	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 40	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 41	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 42	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 43	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 44	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 45	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 46	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 47	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 48	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 49	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 50	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 51	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 52	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 53	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 54	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 55	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 56	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 57	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 58	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 59	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 60	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 61	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 62	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 63	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 64	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 65	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 66	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 67	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 68	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 69	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 70	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 71	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 72	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 73	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 74	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 75	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 76	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 77	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 78	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 79	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 80	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 81	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 82	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 83	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 84	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 85	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 86	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 87	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 88	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 89	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 90	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 91	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 92	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 93	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 94	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 95	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 96	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 97	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 98	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 99	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)
Step 100	0.75	0.00(0.00)	0.00(0.00)	0.36(0.02)

i) Evolution values (mean, standard deviation and type I proportionality error) for a sample variance.

ii) Evolution (mean, standard deviation and type I proportionality error) for a sample average.

## APPENDIX I ABLE III Predominant Radionuclides in Plants, Estimated Activity Content (dpm/g)

to be obtained from the following tables:

<sup>1</sup> See, e.g., *U.S. v. Babbitt*, 100 F.3d 1250, 1254 (10th Cir. 1996) (“[T]he [Bald Eagle] Act is a strict statute that does not allow for any discretion in its enforcement.”).

## A Preliminary Analysis

Stage and Growth in the Infected and Uninfected (*U*) Avian Influenza Virus Strains

Category	Location	Initial Dose (EID <sub>50</sub> )	Relative decrease in titration at p.i./d.p.v.			
			0 d.p.v.	1 d.p.v.	2 d.p.v.	3 d.p.v.
Site A	0.25	0.1000, 0.05	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	2.50	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	10.00	0.1000, 0.05	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
Site B	0.25	0.1000, 0.05	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	2.50	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	10.00	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
Site C	0.25	0.1000, 0.05	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	2.50	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	10.00	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
Site D	0.25	0.1000, 0.05	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	2.50	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	10.00	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
Virus	0.25	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	2.50	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17
	10.00	100	0.500, 0.17	0.700, 0.17	0.700, 0.17	0.700, 0.17

Table 1

1. The relative titrations of the avirulent strains of the four types of avian influenza virus at different stages.

2. The relative titrations of the virulent strains of the four types of avian influenza virus at different stages.

Refractive index calibration in mm/m.

A P P E N D I X

Predominant Radionuclides in Tritium Clad, Tritium Isotopic Isolated, Tritium Isotopic Isolated, Tritium Isotopic

A P P E N D I X 1 A B C E 12

Predominant radioulnar ligamentous lesions in patients collected at Ronchi Lapi Atoll in September 1976

Kadison's like (concentrated) in the sense of

**Predominant Radionuclides in Coconut Crabs Collected  
at Kinglap Atoll in September 1976**

Tissue	Type of crab	Tissue part	Radionuclide concentration ( $\text{mCi} \text{kg}^{-1}$ , $\text{h}^{-1}$ )		
			$\text{Ra}$	$\text{U}$	$\text{Th}$
Carapace	Adult	Whole	1.0	0.0	0.0
Carapace	Adult	Exoskeleton	1.0	0.0	0.0
Carapace	Adult	Endoskeleton	1.0	0.0	0.0
Carapace	Immature	Whole	1.0	0.0	0.0
Carapace	Immature	Exoskeleton	1.0	0.0	0.0
Carapace	Immature	Endoskeleton	1.0	0.0	0.0
Foot	Adult	Whole	1.0	0.0	0.0
Foot	Adult	Exoskeleton	1.0	0.0	0.0
Foot	Adult	Endoskeleton	1.0	0.0	0.0
Foot	Immature	Whole	1.0	0.0	0.0
Foot	Immature	Exoskeleton	1.0	0.0	0.0
Foot	Immature	Endoskeleton	1.0	0.0	0.0
Muscle	Adult	Whole	1.0	0.0	0.0
Muscle	Adult	Exoskeleton	1.0	0.0	0.0
Muscle	Adult	Endoskeleton	1.0	0.0	0.0
Muscle	Immature	Whole	1.0	0.0	0.0
Muscle	Immature	Exoskeleton	1.0	0.0	0.0
Muscle	Immature	Endoskeleton	1.0	0.0	0.0
Respiratory tract	Adult	Whole	1.0	0.0	0.0
Respiratory tract	Adult	Exoskeleton	1.0	0.0	0.0
Respiratory tract	Adult	Endoskeleton	1.0	0.0	0.0
Respiratory tract	Immature	Whole	1.0	0.0	0.0
Respiratory tract	Immature	Exoskeleton	1.0	0.0	0.0
Respiratory tract	Immature	Endoskeleton	1.0	0.0	0.0
Whole body	Adult	Whole	1.0	0.0	0.0
Whole body	Adult	Exoskeleton	1.0	0.0	0.0
Whole body	Adult	Endoskeleton	1.0	0.0	0.0
Whole body	Immature	Whole	1.0	0.0	0.0
Whole body	Immature	Exoskeleton	1.0	0.0	0.0
Whole body	Immature	Endoskeleton	1.0	0.0	0.0

Estimated radionuclide concentrations in the tissues of coconut crabs collected at Kinglap Atoll in September 1976. The values represent the mean of three samples. The error in the mean is approximately  $\pm 10\%$ . The radionuclides measured were Ra-226, U-234, and Th-230. The radionuclides Ra-228, Ra-228, and U-238 were not detected.

The radionuclides Ra-226, U-234, and Th-230 were measured in the tissues of coconut crabs collected at Kinglap Atoll in September 1976. The estimated radionuclide concentrations in the tissues are given in Table I. The radionuclides Ra-228, Ra-228, and U-238 were not detected. The radionuclides Ra-226, U-234, and Th-230 were measured in the tissues of coconut crabs collected at Kinglap Atoll in September 1976. The estimated radionuclide concentrations in the tissues are given in Table I. The radionuclides Ra-228, Ra-228, and U-238 were not detected.

*Heavy metal(loid)s in soil collected on Rongelap Island,  
Rongelap Atoll, in September 1976.*

Site	Depth (cm)	Radiometric detection limit (ppm), dry wt.	
		Cr	As
1	0-10	1.0	0.05
1	10-20	1.0	0.05
1	20-30	1.0	0.05
1	30-40	1.0	0.05
1	40-50	1.0	0.05
1	50-60	1.0	0.05
1	60-70	1.0	0.05
1	70-80	1.0	0.05
1	80-90	1.0	0.05
1	90-100	1.0	0.05
1	100-110	1.0	0.05
1	110-120	1.0	0.05
1	120-130	1.0	0.05
1	130-140	1.0	0.05
1	140-150	1.0	0.05
1	150-160	1.0	0.05
1	160-170	1.0	0.05
1	170-180	1.0	0.05
1	180-190	1.0	0.05
1	190-200	1.0	0.05
1	200-210	1.0	0.05
1	210-220	1.0	0.05
1	220-230	1.0	0.05
1	230-240	1.0	0.05
1	240-250	1.0	0.05
1	250-260	1.0	0.05
1	260-270	1.0	0.05
1	270-280	1.0	0.05
1	280-290	1.0	0.05
1	290-300	1.0	0.05
1	300-310	1.0	0.05
1	310-320	1.0	0.05
1	320-330	1.0	0.05
1	330-340	1.0	0.05
1	340-350	1.0	0.05
1	350-360	1.0	0.05
1	360-370	1.0	0.05
1	370-380	1.0	0.05
1	380-390	1.0	0.05
1	390-400	1.0	0.05
1	400-410	1.0	0.05
1	410-420	1.0	0.05
1	420-430	1.0	0.05
1	430-440	1.0	0.05
1	440-450	1.0	0.05
1	450-460	1.0	0.05
1	460-470	1.0	0.05
1	470-480	1.0	0.05
1	480-490	1.0	0.05
1	490-500	1.0	0.05
1	500-510	1.0	0.05
1	510-520	1.0	0.05
1	520-530	1.0	0.05
1	530-540	1.0	0.05
1	540-550	1.0	0.05
1	550-560	1.0	0.05
1	560-570	1.0	0.05
1	570-580	1.0	0.05
1	580-590	1.0	0.05
1	590-600	1.0	0.05
1	600-610	1.0	0.05
1	610-620	1.0	0.05
1	620-630	1.0	0.05
1	630-640	1.0	0.05
1	640-650	1.0	0.05
1	650-660	1.0	0.05
1	660-670	1.0	0.05
1	670-680	1.0	0.05
1	680-690	1.0	0.05
1	690-700	1.0	0.05
1	700-710	1.0	0.05
1	710-720	1.0	0.05
1	720-730	1.0	0.05
1	730-740	1.0	0.05
1	740-750	1.0	0.05
1	750-760	1.0	0.05
1	760-770	1.0	0.05
1	770-780	1.0	0.05
1	780-790	1.0	0.05
1	790-800	1.0	0.05
1	800-810	1.0	0.05
1	810-820	1.0	0.05
1	820-830	1.0	0.05
1	830-840	1.0	0.05
1	840-850	1.0	0.05
1	850-860	1.0	0.05
1	860-870	1.0	0.05
1	870-880	1.0	0.05
1	880-890	1.0	0.05
1	890-900	1.0	0.05
1	900-910	1.0	0.05
1	910-920	1.0	0.05
1	920-930	1.0	0.05
1	930-940	1.0	0.05
1	940-950	1.0	0.05
1	950-960	1.0	0.05
1	960-970	1.0	0.05
1	970-980	1.0	0.05
1	980-990	1.0	0.05
1	990-1000	1.0	0.05
2	0-10	1.0	0.05
2	10-20	1.0	0.05
2	20-30	1.0	0.05
2	30-40	1.0	0.05
2	40-50	1.0	0.05
2	50-60	1.0	0.05
2	60-70	1.0	0.05
2	70-80	1.0	0.05
2	80-90	1.0	0.05
2	90-100	1.0	0.05
2	100-110	1.0	0.05
2	110-120	1.0	0.05
2	120-130	1.0	0.05
2	130-140	1.0	0.05
2	140-150	1.0	0.05
2	150-160	1.0	0.05
2	160-170	1.0	0.05
2	170-180	1.0	0.05
2	180-190	1.0	0.05
2	190-200	1.0	0.05
2	200-210	1.0	0.05
2	210-220	1.0	0.05
2	220-230	1.0	0.05
2	230-240	1.0	0.05
2	240-250	1.0	0.05
2	250-260	1.0	0.05
2	260-270	1.0	0.05
2	270-280	1.0	0.05
2	280-290	1.0	0.05
2	290-300	1.0	0.05
2	300-310	1.0	0.05
2	310-320	1.0	0.05
2	320-330	1.0	0.05
2	330-340	1.0	0.05
2	340-350	1.0	0.05
2	350-360	1.0	0.05
2	360-370	1.0	0.05
2	370-380	1.0	0.05
2	380-390	1.0	0.05
2	390-400	1.0	0.05
2	400-410	1.0	0.05
2	410-420	1.0	0.05
2	420-430	1.0	0.05
2	430-440	1.0	0.05
2	440-450	1.0	0.05
2	450-460	1.0	0.05
2	460-470	1.0	0.05
2	470-480	1.0	0.05
2	480-490	1.0	0.05
2	490-500	1.0	0.05
2	500-510	1.0	0.05
2	510-520	1.0	0.05
2	520-530	1.0	0.05
2	530-540	1.0	0.05
2	540-550	1.0	0.05
2	550-560	1.0	0.05
2	560-570	1.0	0.05
2	570-580	1.0	0.05
2	580-590	1.0	0.05
2	590-600	1.0	0.05
2	600-610	1.0	0.05
2	610-620	1.0	0.05
2	620-630	1.0	0.05
2	630-640	1.0	0.05
2	640-650	1.0	0.05
2	650-660	1.0	0.05
2	660-670	1.0	0.05
2	670-680	1.0	0.05
2	680-690	1.0	0.05
2	690-700	1.0	0.05
2	700-710	1.0	0.05
2	710-720	1.0	0.05
2	720-730	1.0	0.05
2	730-740	1.0	0.05
2	740-750	1.0	0.05
2	750-760	1.0	0.05
2	760-770	1.0	0.05
2	770-780	1.0	0.05
2	780-790	1.0	0.05
2	790-800	1.0	0.05
2	800-810	1.0	0.05
2	810-820	1.0	0.05
2	820-830	1.0	0.05
2	830-840	1.0	0.05
2	840-850	1.0	0.05
2	850-860	1.0	0.05
2	860-870	1.0	0.05
2	870-880	1.0	0.05
2	880-890	1.0	0.05
2	890-900	1.0	0.05
2	900-910	1.0	0.05
2	910-920	1.0	0.05
2	920-930	1.0	0.05
2	930-940	1.0	0.05
2	940-950	1.0	0.05
2	950-960	1.0	0.05
2	960-970	1.0	0.05
2	970-980	1.0	0.05
2	980-990	1.0	0.05
2	990-1000	1.0	0.05

Table A4 lists various detection limits for radiometric analysis for the two different methods.

The detection limits for the two methods are different because the two methods have different detection limits.

## APPENDIX TABLE I

## APPENDIX TABLE 16

Prevalent Radionuclides in Soil Collected on  
Fatile Island, Bemidji Atoll, in September 1960

Site No.	Sample No.	Radionuclide Concentration in Picocuries per gram		Net count	Net error <sup>a</sup>
		Counting error <sup>b</sup>	Interpolated error <sup>b</sup>		
1	1	0.0000	0.0000	0	0
2	2	0.0000	0.0000	0	0
3	3	0.0000	0.0000	0	0
4	4	0.0000	0.0000	0	0
5	5	0.0000	0.0000	0	0
6	6	0.0000	0.0000	0	0
7	7	0.0000	0.0000	0	0
8	8	0.0000	0.0000	0	0
9	9	0.0000	0.0000	0	0
10	10	0.0000	0.0000	0	0
11	11	0.0000	0.0000	0	0
12	12	0.0000	0.0000	0	0
13	13	0.0000	0.0000	0	0
14	14	0.0000	0.0000	0	0
15	15	0.0000	0.0000	0	0
16	16	0.0000	0.0000	0	0
17	17	0.0000	0.0000	0	0
18	18	0.0000	0.0000	0	0
19	19	0.0000	0.0000	0	0
20	20	0.0000	0.0000	0	0
21	21	0.0000	0.0000	0	0
22	22	0.0000	0.0000	0	0
23	23	0.0000	0.0000	0	0
24	24	0.0000	0.0000	0	0
25	25	0.0000	0.0000	0	0
26	26	0.0000	0.0000	0	0
27	27	0.0000	0.0000	0	0
28	28	0.0000	0.0000	0	0
29	29	0.0000	0.0000	0	0
30	30	0.0000	0.0000	0	0
31	31	0.0000	0.0000	0	0
32	32	0.0000	0.0000	0	0
33	33	0.0000	0.0000	0	0
34	34	0.0000	0.0000	0	0
35	35	0.0000	0.0000	0	0
36	36	0.0000	0.0000	0	0
37	37	0.0000	0.0000	0	0
38	38	0.0000	0.0000	0	0
39	39	0.0000	0.0000	0	0
40	40	0.0000	0.0000	0	0
41	41	0.0000	0.0000	0	0
42	42	0.0000	0.0000	0	0
43	43	0.0000	0.0000	0	0
44	44	0.0000	0.0000	0	0
45	45	0.0000	0.0000	0	0
46	46	0.0000	0.0000	0	0
47	47	0.0000	0.0000	0	0
48	48	0.0000	0.0000	0	0
49	49	0.0000	0.0000	0	0
50	50	0.0000	0.0000	0	0
51	51	0.0000	0.0000	0	0
52	52	0.0000	0.0000	0	0
53	53	0.0000	0.0000	0	0
54	54	0.0000	0.0000	0	0
55	55	0.0000	0.0000	0	0
56	56	0.0000	0.0000	0	0
57	57	0.0000	0.0000	0	0
58	58	0.0000	0.0000	0	0
59	59	0.0000	0.0000	0	0
60	60	0.0000	0.0000	0	0
61	61	0.0000	0.0000	0	0
62	62	0.0000	0.0000	0	0
63	63	0.0000	0.0000	0	0
64	64	0.0000	0.0000	0	0
65	65	0.0000	0.0000	0	0
66	66	0.0000	0.0000	0	0
67	67	0.0000	0.0000	0	0
68	68	0.0000	0.0000	0	0
69	69	0.0000	0.0000	0	0
70	70	0.0000	0.0000	0	0
71	71	0.0000	0.0000	0	0
72	72	0.0000	0.0000	0	0
73	73	0.0000	0.0000	0	0
74	74	0.0000	0.0000	0	0
75	75	0.0000	0.0000	0	0
76	76	0.0000	0.0000	0	0
77	77	0.0000	0.0000	0	0
78	78	0.0000	0.0000	0	0
79	79	0.0000	0.0000	0	0
80	80	0.0000	0.0000	0	0
81	81	0.0000	0.0000	0	0
82	82	0.0000	0.0000	0	0
83	83	0.0000	0.0000	0	0
84	84	0.0000	0.0000	0	0
85	85	0.0000	0.0000	0	0
86	86	0.0000	0.0000	0	0
87	87	0.0000	0.0000	0	0
88	88	0.0000	0.0000	0	0
89	89	0.0000	0.0000	0	0
90	90	0.0000	0.0000	0	0
91	91	0.0000	0.0000	0	0
92	92	0.0000	0.0000	0	0
93	93	0.0000	0.0000	0	0
94	94	0.0000	0.0000	0	0
95	95	0.0000	0.0000	0	0
96	96	0.0000	0.0000	0	0
97	97	0.0000	0.0000	0	0
98	98	0.0000	0.0000	0	0
99	99	0.0000	0.0000	0	0
100	100	0.0000	0.0000	0	0
101	101	0.0000	0.0000	0	0
102	102	0.0000	0.0000	0	0
103	103	0.0000	0.0000	0	0
104	104	0.0000	0.0000	0	0
105	105	0.0000	0.0000	0	0
106	106	0.0000	0.0000	0	0
107	107	0.0000	0.0000	0	0
108	108	0.0000	0.0000	0	0
109	109	0.0000	0.0000	0	0
110	110	0.0000	0.0000	0	0
111	111	0.0000	0.0000	0	0
112	112	0.0000	0.0000	0	0
113	113	0.0000	0.0000	0	0
114	114	0.0000	0.0000	0	0
115	115	0.0000	0.0000	0	0
116	116	0.0000	0.0000	0	0
117	117	0.0000	0.0000	0	0
118	118	0.0000	0.0000	0	0
119	119	0.0000	0.0000	0	0
120	120	0.0000	0.0000	0	0
121	121	0.0000	0.0000	0	0
122	122	0.0000	0.0000	0	0
123	123	0.0000	0.0000	0	0
124	124	0.0000	0.0000	0	0
125	125	0.0000	0.0000	0	0
126	126	0.0000	0.0000	0	0
127	127	0.0000	0.0000	0	0
128	128	0.0000	0.0000	0	0
129	129	0.0000	0.0000	0	0
130	130	0.0000	0.0000	0	0
131	131	0.0000	0.0000	0	0
132	132	0.0000	0.0000	0	0
133	133	0.0000	0.0000	0	0
134	134	0.0000	0.0000	0	0
135	135	0.0000	0.0000	0	0
136	136	0.0000	0.0000	0	0
137	137	0.0000	0.0000	0	0
138	138	0.0000	0.0000	0	0
139	139	0.0000	0.0000	0	0
140	140	0.0000	0.0000	0	0
141	141	0.0000	0.0000	0	0
142	142	0.0000	0.0000	0	0
143	143	0.0000	0.0000	0	0
144	144	0.0000	0.0000	0	0
145	145	0.0000	0.0000	0	0
146	146	0.0000	0.0000	0	0
147	147	0.0000	0.0000	0	0
148	148	0.0000	0.0000	0	0
149	149	0.0000	0.0000	0	0
150	150	0.0000	0.0000	0	0
151	151	0.0000	0.0000	0	0
152	152	0.0000	0.0000	0	0
153	153	0.0000	0.0000	0	0
154	154	0.0000	0.0000	0	0
155	155	0.0000	0.0000	0	0
156	156	0.0000	0.0000	0	0
157	157	0.0000	0.0000	0	0
158	158	0.0000	0.0000	0	0
159	159	0.0000	0.0000	0	0
160	160	0.0000	0.0000	0	0
161	161	0.0000	0.0000	0	0
162	162	0.0000	0.0000	0	0
163	163	0.0000	0.0000	0	0
164	164	0.0000	0.0000	0	0
165	165	0.0000	0.0000	0	0
166	166	0.0000	0.0000	0	0
167	167	0.0000	0.0000	0	0
168	168	0.0000	0.0000	0	0
169	169	0.0000	0.0000	0	0
170	170	0.0000	0.0000	0	0
171	171	0.0000	0.0000	0	0
172	172	0.0000	0.0000	0	0
173	173	0.0000	0.0000	0	0
174	174	0.0000	0.0000	0	0
175	175	0.0000	0.0000	0	0
176	176	0.0000	0.0000	0	0
177	177	0.0000	0.0000	0	0
178	178	0.0000	0.0000	0	0
179	179	0.0000	0.0000	0	0
180	180	0.0000	0.0000	0	0
181	181	0.0000	0.0000	0	0
182	182	0.0000	0.0000	0	0
183	183	0.0000	0.0000	0	0
184	184	0.0000	0.0000	0	0
185	185	0.0000	0.0000	0	0
186	186	0.0000	0.0000	0	0
187	187	0.0000	0.0000	0	0
188	188	0.0000	0.0000	0	0
189	189	0.0000	0.0000	0	0
190	190	0.0000	0.0000	0	0
191	191	0.0000	0.0000	0	0
192	192	0.0000	0.0000	0	0
193	193	0.0000	0.0000	0	0
194	194	0.0000	0.0000	0	0
195	195	0.0000	0.0000	0	0
196	196	0.0000	0.0000	0	0
197	197	0.0000	0.0000	0	0
198	198	0.0000	0.0000	0	0
199	199	0.0000</td			

APPENDIX I

periodic light. Radiotracers labeled in soil collected on flat land and hillsides were also used to determine how much water

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Table 17 (Cont.)

Radiolabelled Concentration in Pictures of  $\mu$ <sup>a</sup>

Collection location (cm)	Height (cm)	T <sub>3</sub>										T <sub>4</sub>										T <sub>5</sub>									
		100	150	200	250	300	350	400	450	500	550	600	650	700	750	800	850	900	1000	1100	1200	1300	1400	1500	1600	1700	1800	1900	2000		
1000	100	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	150	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	200	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	250	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	300	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	350	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	400	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	450	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	500	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	550	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	600	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	650	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	700	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	750	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	800	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	850	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	900	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1000	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1100	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1200	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1300	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1400	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1500	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1600	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1700	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1800	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	1900	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	
1000	2000	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	1.0	

1000 = 1000 nmol/L; 100 = 100 nmol/L; 10 = 10 nmol/L; 1 = 1 nmol/L. Values are expressed as mean  $\pm$  SD.

#### Effect of *l*-Glutamate on the Potency of *l*-DOPA and Tyrosine

The effect of *l*-glutamate on the potencies of *l*-DOPA and tyrosine was determined by the method of Cheng and Prusoff (1973).

Concentrations of *l*-DOPA and tyrosine were varied from 0.01 to 100  $\mu$ M. The concentration of *l*-glutamate was varied from 0 to 1000  $\mu$ M. The effect of *l*-glutamate on the potencies of *l*-DOPA and tyrosine was determined by the method of Cheng and Prusoff (1973).

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A P P L I D I X T A B L E 18  
Predominant Rationuclides in Plants Collected on  
Bikini Island in September and October 1946.

APPENDIX E

GAMMA EMITTING RADIONUCLIDES IN  
PLANTS AND SOIL SAMPLES COLLECTED  
AT BIKINI ISLAND IN OCTOBER 1977 AND APRIL 1976

J. K. Lutz et al.

## Gamma Radiation Factors in the Soil Surface

Collected at British Columbia in October 1977 and 1974-76

A field trip to British Columbia in the spring of 1977 was cancelled because of the unavailability of the required equipment. However, a collection of samples was made in October 1977 as part of another program, the Laboratory's "Biogeochronology of Paleorecords" project, by the Division of Biomedical and Environmental Research. This collection was limited to the most available basin and therefore was not as extensive as planned for the original field trip but some of the samples were collected at sites which had been collected in 1974, 1976 or 1976-77. Following is the analysis of the gamma emitting radionuclides in the October 1977 samples, detailed studies, and also, for the purpose of comparison, the results of analysis of similar samples collected in previous years.

The location of the site of British Columbia where samples were collected in October 1977 are shown in Figure 1. The measured gamma emitting radionuclide is  $^{137}\text{Cs}$  and the results of analysis for this radionuclide in 33 samples from ericaceous plants, coniferous, broadleaf, and alpine, are presented in Table 1 and in Figure 2. The values range from  $3.2 \times 10^3$  to  $4.7 \times 10^3$   $\mu\text{Ci per gram}$  of dry sample. Also found in Table 1 and Figure 2, the results of analysis for  $^{137}\text{Cs}$  in samples collected from the same collection site in either 1974, 1976 or 1976-77. A perusal of the data in Table 1 and Figure 2 indicates no strong seasonal trend for  $^{137}\text{Cs}$  in these samples for this 10-year period. In fact, samples in the 1977 values are greater than those for earlier years. However, the difference between values for the 1976-77 samples may be statistically significant because the values fall into two distinct groups and the sample may not have been collected at all areas from which samples in other years.

Soil profile samples were collected at British Columbia in 1974 and 1977, in October 1977 and the results of analysis of these samples for  $^{137}\text{Cs}$ ,

soil, topsoil, and bulk are given in Table 2. The measured radionuclide was  $^{137}\text{Cs}$  and the resulting  $\text{Bq/g}$  of soil profile abundance was  $3.6 \pm 0.5$ , also presented in Table 2 are the results of analysis of soil profile samples collected at PBI '76 in 1976. The 1977 values appear to be greater than the 1976 value for reasons that are not known but the difference may not be statistically significant.

The data for 33 soil and root profile samples from PBI '76 for 1977 and 1974 are shown graphically in Figure 3. The number above 30 on the 1977 values are apparently artifacts. But the 1976 values when summed, result in a further increase in the 1977 values. The reason is not known but it is likely that the  $^{137}\text{Cs}$  is being relatively exponentially exponential with the values of the surface and at 30, including 10<sup>3</sup> and 10<sup>4</sup>  $\mu\text{Ci per gram}$  of dry sample, respectively. Also in Figure 3 are the 1976 values for pumice, and lava-rock samples, all entered in the same file (PBI '76), in the soil profile samples. The maximum value was  $1.7 \times 10^3$   $\mu\text{Ci per gram}$  of dry sample and the minimum value of pumice root collected in October 1977.

The 1977 horizon (T16) was not supposed to be significantly greater at Site 9, M and C and at house 27 and 28 (Site 28) in recent years. For this reason, the persons collecting the samples were asked about the opportunity to collect samples in 1977 that could not be sampled in recent years. At Site 9 was positively identified but the trench could not be located which samples had been collected at. This could not be confirmed because of flooding. At Site 1 was dug in a depression in the ground but there was no positive identification that this was the location of T16. The trench at Site 1 which could not be located. As a result the previous tree ring dates sampled in 1977 could not be positively identified as there were several locations in the area. At house 21, the trench could not be found but the pieces of wood behind the house that was sampled in 1977 are probably the same material as originally. At Site 28, now 35, the trench could not be found and the tree sampled was one of three that was present. Therefore, because of the inability to obtain duplicate samples, it is preferred to the estimate of the difference in radiocarbon dates between years being used.



Table 1. Summary of 3% volume change of the coastal area around Boknafjord Island in October 1977 and ten similar samples collected between 1974 and 1976.

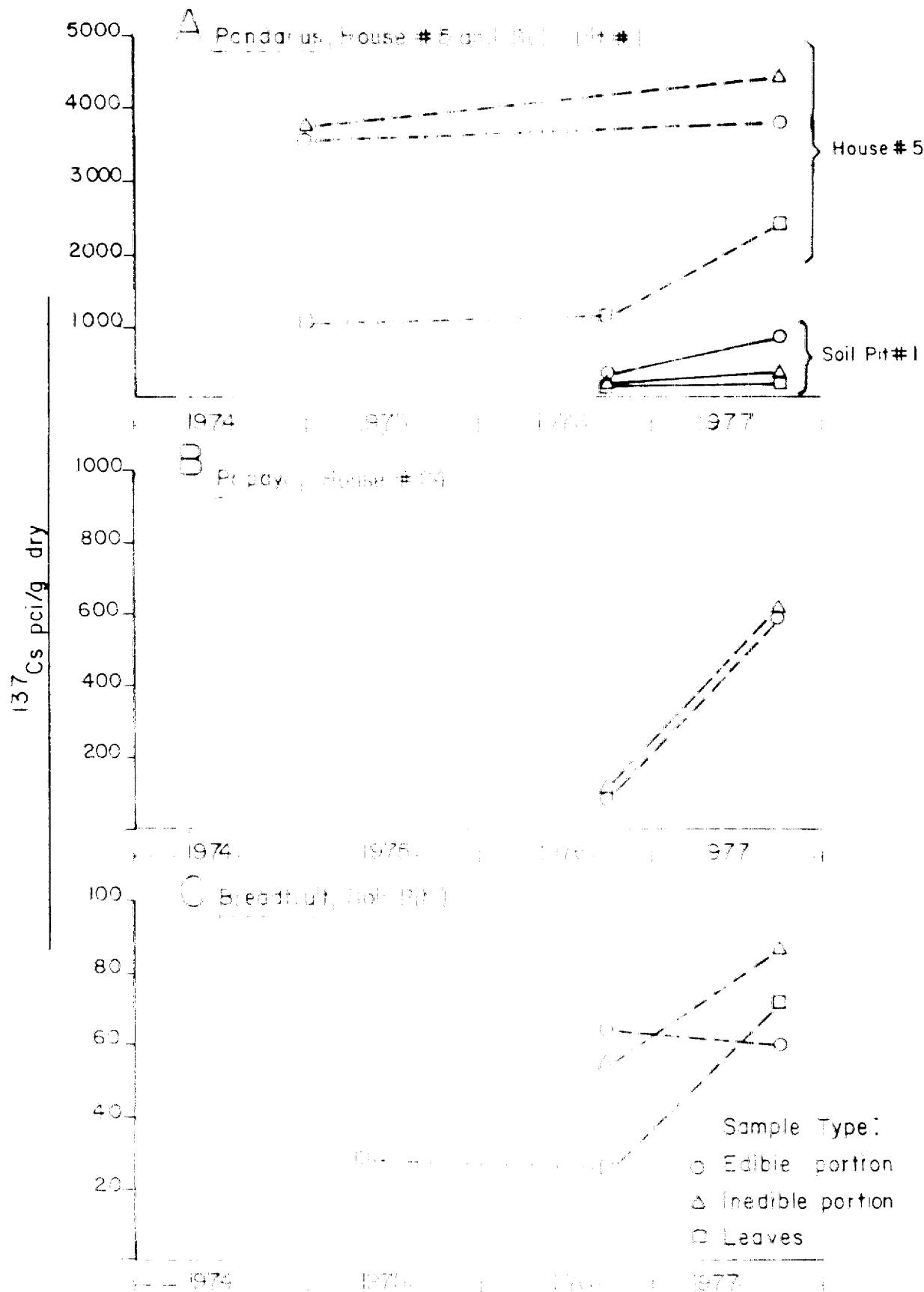


Figure 2. Continued activity for tree-plant samples collected at various locations in Guam during 1974-1977.

**Table 2.** Predominant radionuclides in soil samples collected at Bikini Island in October 1974 and April 1976

Collection Site	Soil Fraction	Collection Date	Radionuclide concentration in pCi/g, dry <sup>a</sup>				
			Ra	U	K	Co	Eu
Pit #9	0-2.5 cm	10 Oct. 1974	1.1 ± 0.1	1.2 ± 0.1	1.1 ± 0.1	1.8 ± 0.1	2.9 ± 0.3
	2.5-5	"	6.6 ± 0.1	10.1 ± 0.9	7.4 ± 0.1	7.4 ± 0.1	1.9 ± 0.3
	5-10	"	7.3 ± 0.1	10.1 ± 0.4	6.4 ± 0.1	6.4 ± 0.1	0.4 ± 0.1
	10-15	"	2.1 ± 0.1	13.0 ± 0.3	—	—	ns <sup>b</sup>
	15-20	"	—	3.7 ± 0.2	—	—	ns
	20-25	"	—	1.7 ± 0.1	—	—	ns
	25-35	"	0.4 ± 0.1	0.6 ± 0.1	—	—	ns
	35-47	"	0.7 ± 0.1	0.9 ± 0.1	0.7 ± 0.1	—	ns
Pit K	0-2.5 cm	10 Oct. 1977	2.6 ± 0.3	2.2 ± 0.5	0.6 ± 0.1	1.7 ± 0.2	—
	2.5-5	"	2.4 ± 0.1	2.2 ± 0.9	0.7 ± 0.1	0.7 ± 0.2	—
	5-10	"	2.6 ± 0.1	2.2 ± 0.2	0.6 ± 0.1	0.7 ± 0.2	—
	10-15	"	2.3 ± 0.1	2.6 ± 0.6	0.3 ± 0.1	0.4 ± 0.2	—
	15-25	"	1.5 ± 0.4	1.6 ± 0.6	0.3 ± 0.1	0.3 ± 0.2	—
	25-35	"	0.9 ± 0.2	1.6 ± 0.2	0.2 ± 0.03	0.2 ± 0.2	—
Pit #9	0-2.5 cm	10 Oct. 1974	3.0 ± 0.5	10.1 ± 0.6	1.2 ± 0.1	1.9 ± 0.1	—
	2.5-5	"	3.0 ± 0.2	22.0 ± 0.5	3.7 ± 0.1	1.7 ± 0.1	—
	5-10	"	2.7 ± 0.2	23.0 ± 0.4	0.6 ± 0.1	0.6 ± 0.1	—
	10-15	"	—	2.6 ± 0.1	—	—	ns
	15-25	"	—	1.7 ± 0.1	0.01 ± 0.02	—	ns
	25-35	"	—	0.6 ± 0.06	—	—	ns
	35-47	"	—	0.1 ± 0.03	—	—	ns
	75-100	"	—	0.3 ± 0.03	—	—	ns

a. values for single sample ± counting error (SEM).

b. ns = sample count not significant.

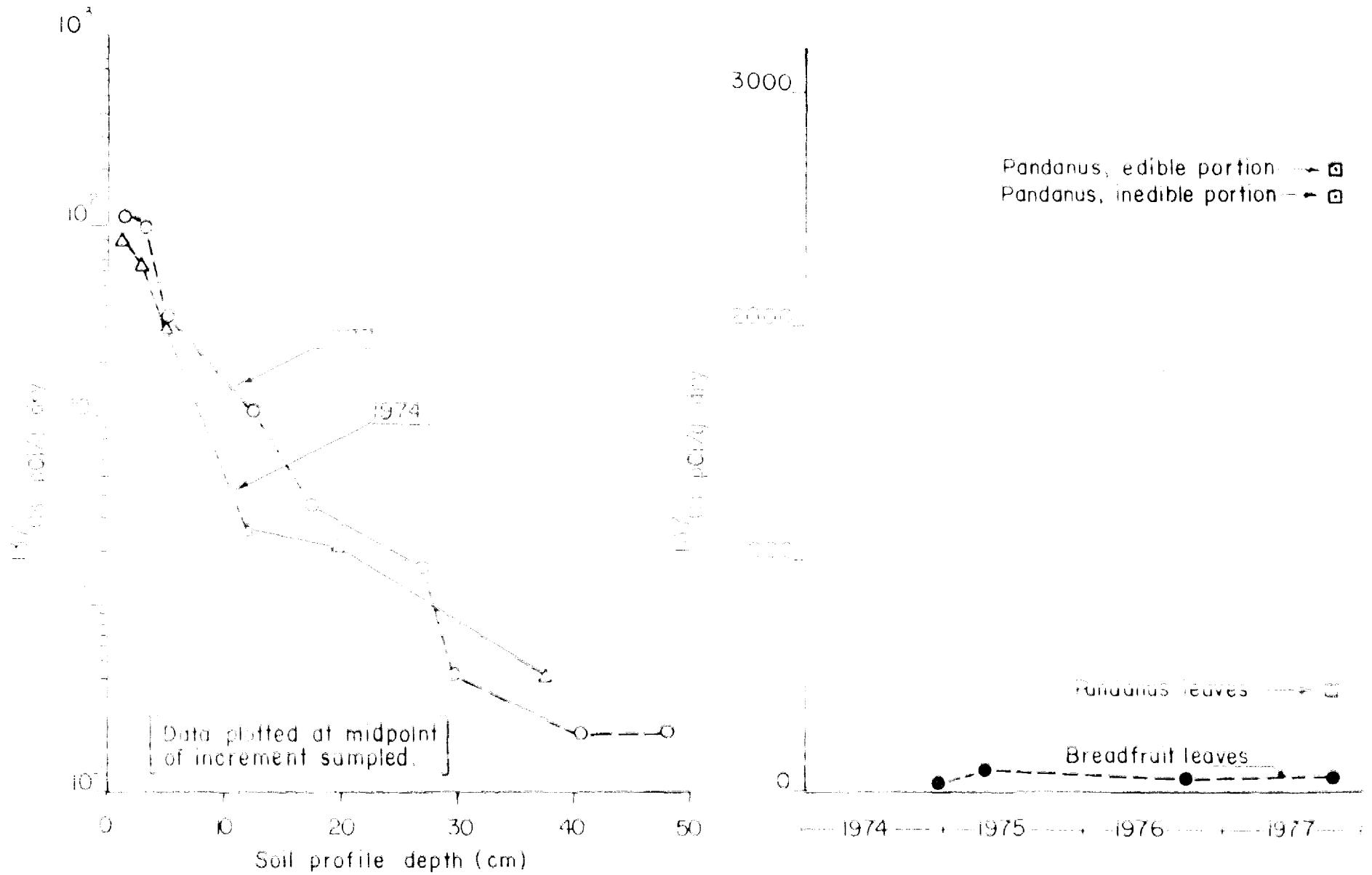


Figure 3. Cs-137 values in soil and plant samples from Ent. #9 on Bikini Island, 1974-1977.

## **APPENDIX 6**

### **INTERLABORATORY COMPARISON PROGRAM AND RADIONUCLIDE DETECTION LIMITS**

## Interlaboratory Comparison Programs and Radionuclide Detection Limits

Since 1971, the laboratory has participated extensively in interlaboratory comparison programs to evaluate the analytical merit of "unknown" radionuclides in both standard and environmental samples. From these results and from the results of routine analysis of duplicate radionuclide samples, we have corrected and improved our methods when needed and thereby continually established and maintained the quality of our analytical work.

For the interlaboratory programs we have analyzed about 170 samples including about 70 radionuclide species. The current programs are with the International Atomic Energy Agency (IAEA), the Environmental Protection Agency (EPA) and the Environmental Measurements Laboratory (EML) (formerly the Health and Safety Laboratory (HSL)). For IAEA, about 10 samples are analyzed yearly; for EPA, usually three water and sediment samples quarterly; and for EMIL, samples of fresh water, sea water, river sediments, marine sediments, fish meal, vegetation meal, and soils are analyzed quarterly.

The results of our自己's and of 17 other laboratories participating in these programs have been tabulated and a report prepared of the tabulation, "Summary of Quarterly Test of Results of Radionuclide Analysis". The report was prepared by Mr. W. A. Kendall (July 1970) and is available upon request. Generally, our results have compared favorably with known values in the standards and with the mean values for all laboratories for the "unknown" radionuclides in the environmental samples.

The methods of analysis used for the interlaboratory program and commonly used in our laboratory for the analysis of other radionuclides are as follows:

1. alpha emitting radionuclides - solution counting methods for  $^{210}\text{Po}$ ,  $^{239}\text{Pu}$ ,  $^{240}\text{Pu}$  and  $^{241}\text{Pu}$ ; zinc sulfide scintillation detector tube counting for gross alpha measurements.

2. beta emitting radionuclides - solution separation and counting for  $^{89}\text{Sr}$  and  $^{137}\text{Cs}$ ; liquid scintillation methods for background, gas counting methods for gross beta measurement.

3. gamma emitting radionuclides - (Li) glass or NaI(Tl) detection systems for radionuclides such as  $^{137}\text{Cs}$ ,  $^{60}\text{Co}$ ,  $^{131}\text{I}$ ,  $^{133}\text{Ba}$ ,  $^{137}\text{Cs}$ ,  $^{36}\text{Ar}$ , etc.; for  $\gamma$ -ray emitters such as  $^{137}\text{Cs}$ , chloride liquid scintillation counting.

The limits of detection for these methods are important since they govern the amount of a radionuclide that can be detected if it is present in a sample. Many factors influence the limit of detection, including the type of detector and analyzer, the presence of other radionuclides, the duration of the counting period, the size and density of the sample, and the geometry relationship of the sample and detector. Hence, the detectable limits of detection can vary considerably for various radionuclides and types of samples, but can be summarized by stating that the detection limits are approximately as follows:

By gamma detection:

$^{19}K$   $\gamma/\beta/\gamma/\alpha/\text{less}$

$^{7}Be$ ,  $^{103}Ru$ ,  $^{137}Cs$ ,  $^{138}Ba$ ,  $^{226}Ra$ ,  $^{230}Th$   $\gamma/\beta/\gamma/\alpha/\text{less}$

$^{95}Nb$ ,  $^{113}Cd$ ,  $^{137}Cs$ ,  $^{138}Ba$ ,  $^{226}Ra$   $\gamma/\beta/\gamma/\alpha/\text{less}$

By beta detection:

$^{19}K$   $\beta/\beta/\beta/\text{less}$

$^{85}Sr$   $\beta/\beta/\beta/\beta/\text{less}$

By X-ray detection:

$^{55}Fe$   $(\gamma, X) \rightarrow \text{less}$

By alpha detection:

$^{226}Ra$ ,  $^{232}Th$   $\alpha/\alpha/\alpha/\text{less}$