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Frequencies ARCENTES



Joseph Rivera suffered a heart attack and died on January 31, 1970. We are greatly saddened by the loss of our good friend and colleague. Soon after joining the Health and Safety Laboratory in 1960, Joe helped to implement the human diet and bone studies at HASL. He continued to pursue these and other investigations with the basic aim of estimating the exposure of man from radioactive fallout.

His strong interest in and involvement with the United Nations Scientific Committee on the Effects of Atomic Radiation, the National Committee on Radiation Protection and Measurements, and the Federal Radiation Council, exposed his scientific skills as well as his good humor to scientists from many areas of the world.

Joe was co-editor of the HASL "Quarterly". His advice and many contributions will be sorely missed. He was an independent thinker whose ideas and talents were sought in almost every endeavor at HASL.

HASL-224

UC-41, Health and Safety TID-4500, 54th Ed.

HEALTH AND SAFETY LABORATORY

FALLOUT PROGRAM QUARTERLY SUMMARY REPORT

(December 1, 1969 through March 1, 1970)

Prepared by

Edward P. Hardy, Jr.

Environmental Studies Division

Preceding reports in this series:

Year	HASL Report Nos.
1958	42, 51
1959	65
1960	77, 84, 88, 95
1961	105, 111, 113, 115
1962	117, 122, 127, 131
1963	132, 135, 138, 140
1964	142, 144, 146, 149
1965	155, 158, 161, 164
1966	165, 171, 172, 173
1967	174, 181, 182, 183
1968	184, 193, 197, 200
1969	204, 207, 210, 214
1970	2.1.7

April 1, 1970

UNITED STATES ATOMIC ENERGY COMMISSION New York Operations Office

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FALLOUT PROGRAM QUARTERLY SUMMARY REPORT

April 1, 1970

ABSTRACT

This report presents current data from the HASL Fallout Program, the National Radiation Laboratory in New Zealand, the Department of Scientific and Industrial Research in New Zealand, The EURATOM Joint Nuclear Research Centre and The Radiological Physics Division at Argonne National Laboratory. The initial section consists of interpretive reports and notes covering the following topics: radium-226 in diet, plutonium-239 anomaly in the troposphere, and the quality of radiochemical analyses in the HASL surface air sampling program. Subsequent sections include tabulations of radionuclide levels in stratospheric air, surface air, failout, milk, other diet components, and tap water. A bibliography of recent publications related to radionuclide studies is also presented.

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<u>Diets</u>

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INTRODUCTION

Every three months, the Health and Safety Laboratory issues a report summarizing current information obtained at HASL pertaining to fallout. This report, the latest in the series, contains information that became available during the period from December 1, 1969 to March 1, 1970. The next report is scheduled for publication July 1, 1970. Preceding reports in the series, starting with HASL-42, "Environmental Contamination from Weapons Tests", and continuing through HASL-224, (this report) may be purchased from the Clearinghouse for Federal Scientific and Technical Information, National Bureau of Standards, U. S. Department of Commerce, Springfield, Virginia 22151. A complete listing of these Fallout Program Quarterly Summary Reports is given on the title page of this report.

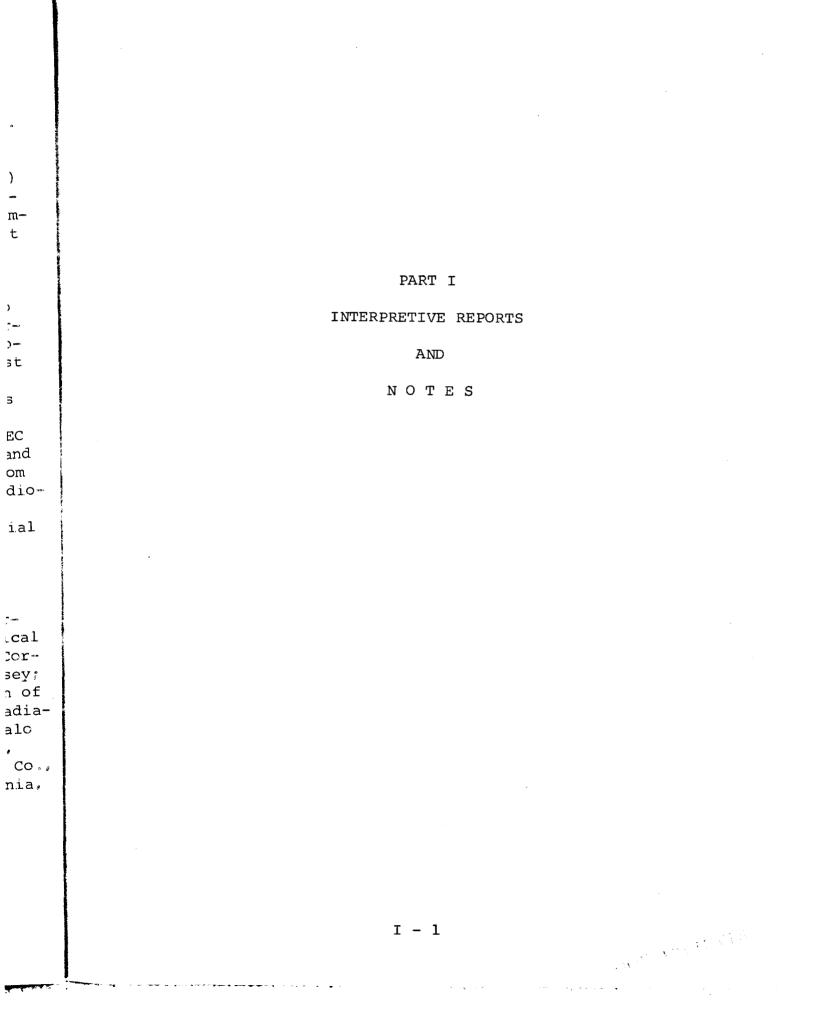
To give a more complete picture of the current fallout situation and to provide a medium for rapid publication of radionuclide data, these quarterly reports often contain information from other laboratories and programs, some of which are not part of the general AEC program. To assist in developing, as rapidly as possible, provisional interpretations of the data, special interpretive reports and notes prepared by scientists working in the field of fallout are also included from time to time. Many of these scientists are associated in some way with the general AEC program. Information developed outside of HASL is identified as such and is gratefully acknowledged by the Laboratory. In this report, data from the EURATOM Joint Nuclear Research Center at Ispra, the Division of Radiological Physics at Argonne National Laboratory, the National Radiation Laboratory in New Zealand and the Department of Scientific and Industrial Research in New Zealand, are given.

A portion of the radiochemical analyses either have been or are being carried out by commercial laboratories under contract to the HASL Environmental Studies Division. The results of these analyses are reported as part of HASL's regular fallout program. The contractor analytical laboratories which provided data are Nuclear Science and Engineering Corporation, Pittsburgh, Pennsylvania; Isotopes, Inc., Westwood, New Jersey; Radiochemistry Incorporated, Louisville, Kentucky; Tracerlab, Division of LFE, Richmond, California (now Trapelo Division/West); Controls for Radiation, Inc., Cambridge, Mass.; Hazleton-Nuclear Science Corporation, Palc Alto, Calif.; Food, Chemical and Research Laboratories, Inc., Seattle, Washington; Tracerlab, Division of LFE, Waltham, Mass; U. S. Testing Co., Inc., Richland, Washington, Custom Nuclear Co., Mountainview, California, and Ledoux and Co., Teaneck, New Jersey.

This report is divided into four main parts:

- Interpretive Reports and Notes 1.
- 2. HASL Fallout Program Data
- 3. Data from Sources Other than HASL
- 4. Recent Publications Related to Radionuclide Studies boy. Constraint

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RADIUM-226 IN THE DIET OF TWO U. S. CITIES

I. M. Fisenne and H. W. Keller

A revised estimate of the daily radium-226 dietary intake in three cities (New York City, Chicago and San Francisco) was reported in April 1969.⁽¹⁾ At that time, a comparison of the 1960 and 1966 diet estimates showed that the 1966 values were from 27% to 50% lower than the 1960 values. This difference was due to the change in the analytical procedure for radium-226. To obtain a more representative estimate of dietary intake, a second food sampling from the New York City and San Francisco was analyzed for radium-226. No Chicago diet samples were available for analyses, since food sampling from this city was terminated in 1968. Samples of New York City and San Francisco area tap water were also analyzed for radium-226.

The HASL food sampling program is described in HASL-200.⁽²⁾ The tap water samples from New York City represent an integrated one month collection. The San Francisco area tap water sample was collected on a single day.

The radium-226 diet level estimates are shown in Tables 1 and 2. Duplicate sample analyses are also listed. The yearly consumption figures are taken from a U. S. Department of Agriculture report.⁽³⁾ The average daily intake of radium-226 in the two cities may be computed from these data.

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New York CitySan Francisco1.7 pCi/day0.8 pCi/day

These values do not include the contribution of radium-226 from tap water. However, the radium-226 content of the water is low and would not materially increase the daily intake estimate.

<u>New York City</u>	San Francisco
0.011 pCi/liter	0.015 pCi/liter

Comparison of the two surveys in Tables 1 and 2 shows that although individual food types may vary by more than a factor of 2, the average daily intake for each city varies by less than 20% or within expected experimental error. It should also be noted that both the 1966 and 1968 food samplings were collected at mid-year so that possible seasonal variation was not studied. It seems unlikely that a significant seasonal variation in dietary radium-226 would exist for a given location.

The California State Department of Health has estimated the daily radium-226 intake of individuals receiving a hospital diet. The average daily radium-226 intake for Berkeley, California hospital diets from November 1967 to June 1968 was 0.4 pCi/day.⁽⁴⁾ Similar diets collected from April to December 1968 in San Francisco averaged 1.3 pCi/day. (5,6,7)

The radium-226 content of human bone from New York City and San Francisco is 0.036 pCi/g Ca and 0.031 pCi/g Ca, respectively, Since the dietary radium-226 levels for these cities are different by about a factor of two, it would be expected that the bone levels would reflect this difference. We suggest that a particular dietary component or a combination of a few components may control the radium-226 level in man. Little is known about the availability of radium-226 from foodstuffs, but there is indication that it is quite variable. We are initiating a few studies to attempt to correlate radium-226 bone levels with particular Part 16 Warry dietary components. I - 3

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Radium-226	in	New	York	City	Diet
Radium-220	111	New	TOTE	Orej.	

			19	66	19	68	Aver	
Diet Category	kg	g Ca yr	pCi kg	pCi yr	pCi kg	pCi yr	pCi kg	pCi yr
Dairy Products	200	216.0	0.25	50	0.30* 0.19*	60 38	0.25	50†
Fresh Vegetables	48	18.7	0.50	24	1.6 1.6	77 77	1.1	53
Canned Vegetables	22	4•4	0.65	14	0.68	15	0.67	15
Root Vegetables	10	3.8	1.4	14	1.2	12	1.3	13
Potatoes	38	3.8	2.8	106	1.7	65	2.3	87
Dry Beans	3	2.1	1.1	3.3	0.98	2.9	1.0	3
Fresh Fruit	59	9.4	0.43	25	0.20	12	0.32	19
Canned Fruit	11	0.6	0.17	1.9	0.15 0.16	1.7 1.8	0.16	1.8
Fruit Juice	28	2.5	0.42	12	0.90	25	0.66	18
Bakery Products	44	53.7	2.8	123	1.7	75	2.3	101
Flour	34	6.5	1.9	65	2.3	78	2.1	71
Whole Grain Products	11	10.3	2.2	24	2.7	30	2.5	28
Macaroni	3	0.6	2.1	6.3	1.4	4.2	1.8	5.4
Rice	3	1.1	0.76	2.3	3.3	9.9	2.0	60

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Diet Category	kg yr	g Ca yr	pCi kg	pCi yr	pCi kg	pCi yr	pCi kg	pCi yr
Meat	79	12.6	0.01	0.8	0.02	1.6	0.02	1.6
Poultry	20	6.0	0.76	15	0.10 0.11	2.0	0.44	8.8
Eggs	15	8.7	6.1	92	14 14	210 210	10	150
Fresh Fish	8	7.6	0.67	5.4	1.1	8.8	0.89	7.1
Shellfish	• . 1	1.6	0.80	0.8	0.90	0.9	0.85	0.9
Yearly Intake Daily Intake pCi/g Ca *Two different samples †Average of three sampl	les	370		584.8 1.6		680.2 1.8		639.6 1.7

н 1 5 Radium-226 in New York City Diet

TABLE 2

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Radium-226 in San Francisco Diet

			196	1966		1968		Average	
Diet Category	kg yr	g Ca yr	pCi kg	pCi yr	pCi kg	pCi yr	pCi kg	pCi yr	
Dairy Products	200	216.0	0.10	20	0.09 0.08	18 16	0.09	18	
Fresh Vegetables	48	18.7	0.48	23	0.80 0.88	38 42	0.66	32	
Canned Vegetables	22	4.4	0.35	7.7	0 .3 6	7.9	0.36	7.9	
Root Vegetables	10	3.8	1.2	12	1.5	15	1.4	14	
Potatoes	38	3.8	0.14	5.3	0.33	13	0.24	9.1	
Dry Beans	3	2.1	0.72	2.2	0.67	2.0	0.70	2.1	
Fresh Fruit	59	9.4	0.25	15	0.27	16	0.26	15	
Canned Fruit	11	0.6	0.70	7.7	0.16 0.19	1.8 2.1	0.44	4.8	
Fruit Juice	28	2.5	0.33	9.2	0.79	22	0.56	16	
Bakery Products	44	53.7	1.2	53	1.6	70	1.4	62	
Flour	34	6.5	1.4	48	1.4	48	1.4	48	
Whole Grain Products	11	10.3	2.1	23	2.2	24	2.2	24	
Macaroni	3	0.6	2.6	7.8	1.7	5.1	2.2	6.6	
Rice	3	1.1	0.24	0.7	0.33	1.0	0.29	0.9	
Meat	79	12.6	0.01	0.8	0.02	1.6	0.02	1.6	

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TABLE 2

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Diet Category	kg yr	g Ca yr	pCi kg	pCi yr	pCi kg	pCiyr	pCi kg	pCi yr
Poultry	20	6.0	0.43	8.6	0.38	7.6	0.41	8.2
Eggs	15	8.7	2.0	30	1.5	23	1.8	27
Fresh Fish	8	7.6	0.40	3.2	0.11	0.9	0.26	2.1
S hellfish	1	1.6	1.9	1.9	1.1	1.1	1.5	1.5
Yearly Intake Daily Intake pCi/g Ca		370		279.1 0.75			317.2 0.86	300.8 0.81

Radium-226 in San Francisco Diet

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All of the values reported by the contractor are corrected for a reagent blank, so that the blank values reported here reflect the contamination of the filter material both during manufacture and during handling at HASL and at the contractor laboratory.

Table 1 lists the results of the analyses of the blanks for each nuclide. In general these data indicate low levels of contamination for most of the nuclides, however it is significant to note that the average Pu-238 blank rose from 0.04 dpm in 1967 to 0.22 dpm in 1968. Because the amount of Pu-238 found in surface air samples is frequently in the range of 1 dpm this blank may represent a significant fraction of the total activity.

The results of analyses on standard samples are shown in Table 2. These data are indicative of the accuracy of the radiochemical analyses. The values shown are the average percent deviations between the added activities and the results reported by the contractor. Although most of the results are satisfactory, there appears to be a large positive bias in the Pu-238 values for much of 1968, which cannot be accounted for by the increase of the blank contamination. Samples submitted to the contractor during 1969 which were prepared with a new standard solution do not exhibit this bias, and it is therefore probable that the poor results reflect a degeneration of the Pu-238 standard solution used for the preparation of the 1968 guality control samples.

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Table l

Quality Control Results - Blanks - 1968

(Values in dpm/sample ± 1 std. Deviation)

	Fe-55	Sr-89	Sr-90	Zr-95	Cs-137	Ce-141	Ce-144	Pu-238	Pu-239	Stable Pb
January	42±67	7.5±4.6	0±.5	2.5±4.9	·	13±73	1.5±3.4	.34±.05	.04±.02	
Februar y	24±59		0.5±1.5	3.0±6.0			2.0±3.0	L	L	
March	62±137		23.8±12.4	0±6.4			6.7±4.1	.1 8±.01	.09±.03	
April	0±156		0±.03	1.4±13			5.4±2.5	0±.04	0±.03	
Мау	0±86		0±1.1	0±8.8			0±3.3	.29±.05	.03±.02	
June	96±72		1.6±4.3	3.1±6.1			0±1.9	.37±.05	0 ± .03	
July		7.1±20.1	0.5±1.3	0.9±7.2	0.2±0.4		2.1±2.8	.36±11	.12±.10	
August		0±2.3	0.5±0.4	9.3±4.4	0.7±4.0		2.0±1.6	.29±.07	.05±.05	
September		0 ±1.5	0 ± 0.4	2.8±3.0	1.5±0.5		1.6±1.6	.41±.06	0 ±.03	
October			0 ± 2.3		5.2±0.7		3 .0± 6.6	.11±.03	0 ±.02	.03
November			0.1±0.8	5.7±8.8	0.9±0.5		0.9±2.8	.06±.03	0 ±.03	
December			0.9±3.2	13.5±16.1	0.1±0.5		229±270	.11±.09	0 ±.07	0.0

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NOTE: L indicates sample lost

-- indicates no analysis

Table 2

Quality Control Results - Standards-1968 (Average values in % Deviation)

	<u>Fe-55</u>	Sr-89	<u>Sr-90</u>	Zr-95	<u>Cs-137</u>	<u>Ce-144</u>	<u>Pu-238</u>	Pu-239	Stable <u>Pb</u>
January	12.5	-12.5	-4.0	-13.0		-2.0	-8.5	-3,5	
February	18.0		-16.0	-19.5		-1.5	17.0	3.0	
March	21.5		-1.0	-21.0		-3.0	15.5	0.5	
April	5.5		-5.5	-13.5		0.5	16.0	-0.5	-1600 anna
Мау	-5.0		-22.0	-22.0		2.5	9.5	1.0	
June	13.0		-9.0	-16.0		3.5	10.0	1.0	
July			1.0	-6.0	11.0	10.0	10.0	1.0	
August		66.5	-13.5	-6.0	-5.0	-2.0	29.0	-3.5	
September		-11.5	- 6.5	-7.0	-8.0	2.0	9.0	0.5	
October			-3.5		-6.0	1.0	26.5	2.0	11.3
November		 	-1.5	-11,5	-	8.0	39.0	6.1	27.0
December	ca a		-5.0	-14.8	1.0	4.5	35.0	3.9	2.7

NOTE: -- indicates no analysis.

Plutonium-239 Anomaly in the Troposphere

by H. L. Volchok, (HASL) P. W. Krey, (HASL)

During the period mid-1968 through mid-1969, some unusual behavior of Pu-239 relative to Sr-90 has been observed in surface air. For many months and at many surface air sampling stations in the Northern Hemisphere, the ratio Pu-239/Sr-90 exceeded all values in the stratosphere at comparable times. Although a number of possibilities are discussed in this paper, this apparent enrichment of Pu-239 is not readily explainable.

Most of the Pu-239 existing in the world came about by the interaction of enormous neutron fluxes in atomic explosions with the U-238 present in such devices (1,2) or from unfissioned Pu-239 from the cores of the weapons. Much smaller amounts may have been released to the environment by a variety of accidents and incidents on the earth's surface such as airplane crashes at Thule, Greenland, and Palomares, Spain (3) or the recently publicized contamination in the neighborhood of Rocky Flats, Colorado (4). Sr-90 on the other hand has been released to the atmosphere mainly in nuclear explosions as a product of the fission reaction. Since the major production modes of these two radionuclides are very different, their distribution in the stratosphere and surface air have been studied for some time.

Stratospheric Data

The Pu-239/Sr-90 ratios at 19.2 km and just above the tropopause are illustrated in Figures 1 and 2 for the period August 1967 through April 1969. These figures were constructed from data obtained in Project Airstream (5,6,7,8) and represent the minimum and maximum altitudes routinely sampled in this program. Since the tropopause height varies with latitude, the altitudes in Figure 2 also vary as follows:

Latitude	<u>Altitude</u>
75 ⁰ n-65 ⁰ n	12.2 km
65 ⁰ n-36 ⁰ n	13.7 km
36 ⁰ n-34 ⁰ S	15.2 km
34 ⁰ s-51 ⁰ s	13.7 km

Figure 1 shows that at the higher altitude, the Pu-239/Sr-90 was uniformly less than 2% in both hemispheres, until the debris from the 6th Chinese test of June 1967 was intercepted in the last quarter of the year. At this point the ratio rose to greater than 2 but less than 3% and essentially remained within this range through all of 1968. The contours suggest that this debris at 19.2 km crossed the equator into the Southern Hemisphere in the spring of 1968 and influenced the ratio to as far south as the Airstream sampling network extends, 51°S.

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Figure 1 further shows that the French thermonuclear tests in mid-1968 abruptly ended the influence of the 6th Chinese test debris in the Southern Hemisphere, plunging the ratio to below 1%. The Chinese 8th test in December of 1968 apparently generated a ratio of Pu-239/Sr-90 of about 2% since the pattern in the Northern Hemisphere was not markedly affected in early 1969.

The Pu-239/Sr-90 in the lower stratosphere, just above the tropopause, is illustrated in Figure 2, and exhibited a similar pattern with one striking difference. At this altitude the high ratios generated by small French tests in the Southern Hemisphere in mid-1967 are clearly seen. Most important to note, however, is that in the Northern Hemisphere the highest ratios observed in the entire period shown were greater than 2.5% but less than 3%, and in early 1969 virtually all of the data were below 2.5%. This observation holds true not only for the lowest stratosphere but also for altitudes as high as 19.2 km (Fig. 1).

Surface Air Data

The Pu-239/Sr-90 ratios in surface air for the period 1966 through mid-1969 are shown in Figure 3. This figure was derived from the data obtained in the Health and Safety Laboratory (HASL) Surface Air Sampling Program (9).

Since the onset of our surface air analyses for Pu-239, in late 1965 through the beginning of 1968, rarely did any sample from the Northern Hemisphere indicate a ratio in excess of 2.5%, and the few that did were always analytically suspect. This relative constancy of the ratio had been noted by other investigators in earlier studies of stratospheric samples (1). In the Southern Hemisphere, high ratios in surface air were seen closely following the small French Tests in 1966 and 1967. The peak ratios appear to be displaced poleward relative to the reported latitude of the Southern Hemisphere tests. Recognizing that some debris from these tests did enter the lower stratosphere and that these peaks prevailed throughout the spring of each year, we feel that the geographical and temporal extent of the highs are not unexpected.

As a result of the 6th Chinese test (June 1967) most of which was deposited in the stratosphere, the gradual rise in the Pu-239/Sr-90 in the Northern Hemisphere surface air from less than 2% in mid-1967 to almost 3% by mid 1968 was not surprising. Starting with June 1968, however, and persisting well into 1969, numerous samples were measured with ratios over 3% and a few even exceeded 4%. From Figures 1 and 2 it seems clear that ratios such as these could not be derived from the debris in the stratosphere without some fractionation of these two nuclides between the tropopause and the earth's surface.

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Discussion

There are at least five possible explanations for the anomalously high Pu-239/Sr-90 ratios observed in surface air since mid-1968. These are as follows:

1. Systematic analytical bias. The possibility of a systematic error in the radiochemical analyses seems very remote because through the period of anomalous ratios, both the stratospheric and surface air samples were analyzed by a single contractor laboratory (Trapelo/West). The performance of the contractor during this period was monitoried by analysis of coded quality control samples and found to be excellent. Furthermore, the data are seen to be internally consistent, allowing for relatively smooth contouring.

2. Non-representative sampling. The representativeness of the samples in both the Airstream and Surface Air programs have always been considered adequate. The filter media, the particle size of the debris, the face velocities and the general conditions of sampling were all considered in design of the equipment in these programs. In this light, erratic bias in the samplers seems highly unlikely. In addition the continuing balance in the Sr-90 budget in

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all compartments of the earth's atmosphere and on its surface (10) suggests that the overall representativeness of the data is good.

3. Transport of debris from regions of higher ratio in either the troposphere or stratosphere. Figures 1, 2, and 3 clearly eliminate this possibility in that from mid-1968 there are no other data in either the stratosphere or surface air of either hemisphere with ratios as high as the anomalies under consideration. In view of our conclusion regarding the representativeness of the data, it appears that there are no hidden compartments or pockets in the atmosphere containing unexpected debris.

4. Tropospheric injections of Pu-239. A few major injections of Pu-239 in 1968 could theoretically have produced the high Pu-239/Sr-90 observed from mid year onward. The actual amount of "excess" Pu-239 in the surface air during the period of the anomaly, that is the amount not accountable as coming out of the stratosphere, was estimated by using average ratios and Sr-90 concentrations. It was found that 22% of the surface air ratio (on the average) was excess, which is equivalent to about 2 x 10^{-5} dpm/m³ of unaccountable Pu-239.

An upper and lower limit of the average monthly tropospheric
<u>D</u> inventory of the excess Pu-239 was then estimated from the above result, depending upon the assumed distribution with altitude. If the h excess assumed an altitude profile similar to that of stratospheric debris, an upper limit of 70 curies of excess Pu-239 in the troposphere was indicated. If the excess concentration is essentially constant to an elevation of 1000 meters, with nothing above, a lower limit of about 1 curie was obtained.

There were no reported atmospheric nuclear tests in the Northern Hemisphere from December 1967 to December 1968 (11); hence the mid-tolate-1968 anomalies cannot be explained as due to fresh debris. Other possible surface releases of Pu-239, such as airplane crashes with loss of nuclear material, leaks from processing plants or other similar sources cannot be evaluated with information presently available. However, since virtually all of the Pu-238 in surface air, in the period of the anomaly came out of the stratosphere (either from nuclear tests or from the burn-up of SNAP-9A) a sharp decrease in the ratio Pu-238/Pu-239 would have occurred in the event of significant tropospheric releases of Pu-239. As Figure 4 indicates, this sharp decline cannot be seen. Some decrease did occur in 1968, much of it reflecting fallout from

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China's 6th test in June 1967, but this did not closely match the Pu-239 increase either in magnitude or timing. Further, in the mid latitudes in 1968, and in all latitudes in 1969, increases in the Pu-238/Pu-239 occurred.

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On balance, we do not feel that the data support major tropospheric injections of Pu-239 to account for the anomalous Pu-239/Sr-90 ratios in the Northern Hemisphere. There is, however, a good deal of evidence of global, or at least hemispheric distribution of material emanating from limited sources at the earth's surface. As examples: lead, identifiable as originating in urban environments has been found in both the Arctic and Antarctic (12); dust, derived from the arid regions of West Africa, has been traced westward across the Atlantic Ocean to Barbados (13); and pesticide residues found in organs of Antarctic fauna (14), also must have been transported in the troposphere many thousands of miles. Thus there is not sufficient evidence at this time to Tule out this possibility.

5. Fractionation in the troposphere. This fifth possible explanation for the observed anomalies in the Pu-239/Sr-90 ratios offers much to the imagination, but there is little if any real positive evidence. Negatively, it can be demonstrated from Figure 3, that this had never happened to any observable extent in the preceeding two years. Also, it seems reasonable to assume that if the surface

air is enriched in Pu-239 in this particular period, somewhere in the atmosphere or on the ground we should observe data showing depletion in that isotope. Fallout data do not support this idea. Although sparse (only two deposition sites in the Northern Hemisphere had samples routinely analyzed for Pu-239), the fallout results indicate tha: Pu-239/Sr-90 ratios in precipitation are about the same as in the surface air (9, 15, 16, 17, 18, 19, 20, 21, 22, 23). On the other hard it certainly would not be unreasonable to expect fractionation between two such chemically different elements as Pu and Sr. Furthermore, since their origins differ markedly, they may very likely enter the tropospheron particles of different size. We do not propose to speculate further on the possible mechanisms of fractionation; however, it does seem clear: that an acceptable case for fractionation of these radionuclides. cc::ld be presented.

<u>Conclusions</u>

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The anomalous ratios of Pu-239/Sr-90 in Northern Hemisphere surface air in 1968-1969 remain unexplained. Neither of the most plausible explanations advanced in this report, tropospheric releases of Pu-239 and tropospheric fractionation of the radionuclides, have been experimentally substantiated. In fact both explanations seem to be contraindicated by other observations and data. Sampling and analysis for Pu-239 and Sr-90 will be continued in both the stratospheric and surface air programs, and other studies which may bear on this subject will be considered to help finally understand this paradox.

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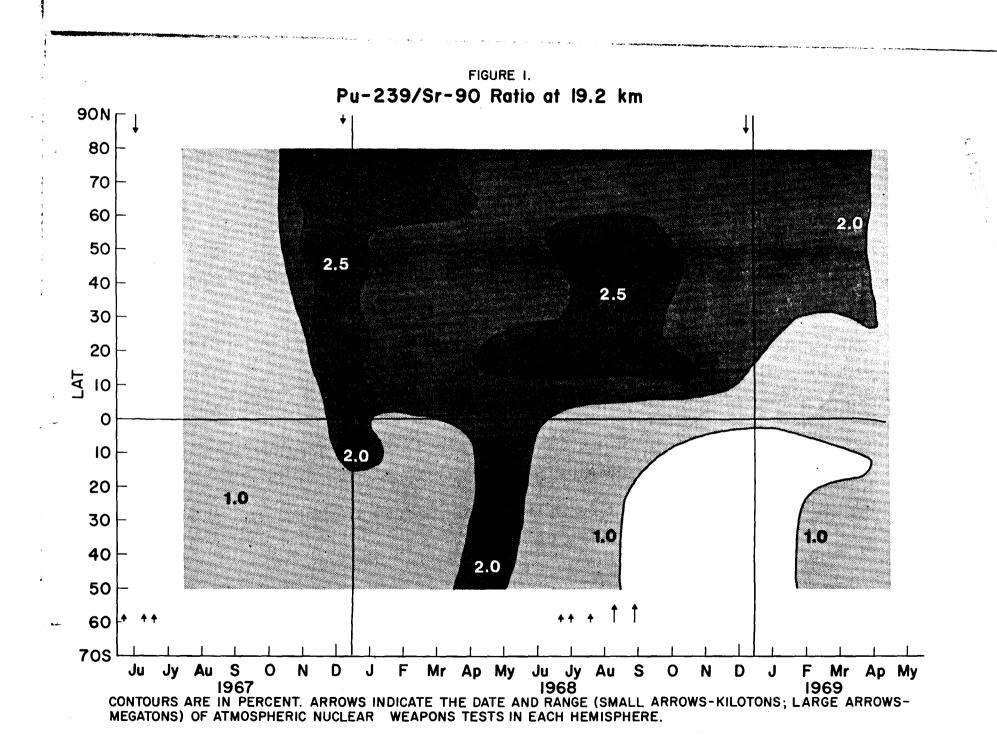
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USAEC Report HASL-217 (Appendix), January (1970), p. C-1



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FIGURE 2 Pu-239/Sr-90 Ratio Just Above the Tropopause 90N 80 70 2.5 60 2 2.5 50 40 30 20 OI LAT 0 11 10 1.02.5 20 3.0 30 1.0 40 50 ++ + 1 1 60 L+ - 🕈 🕈

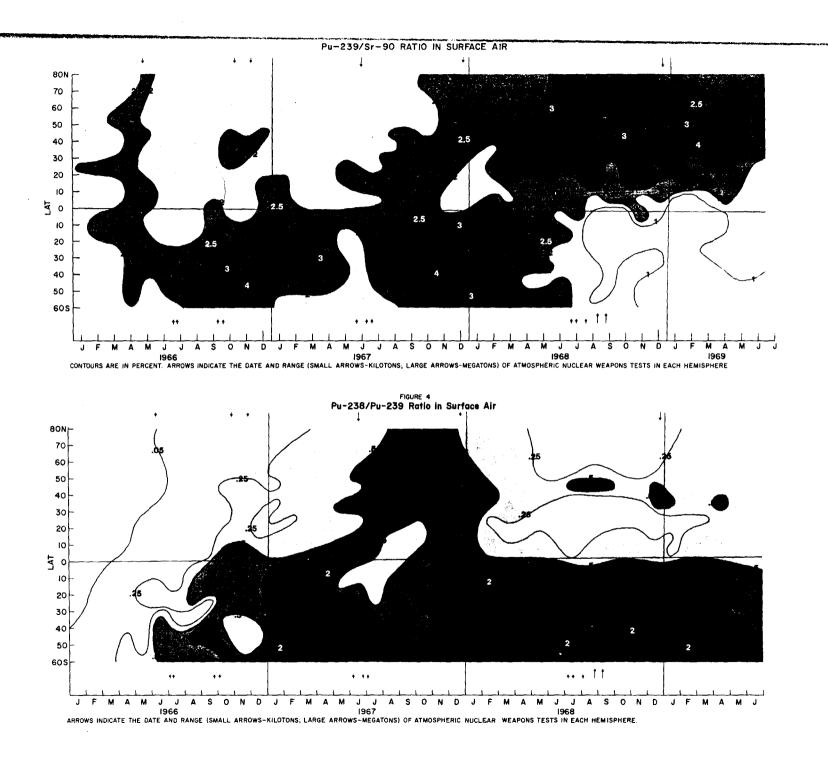
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WEAPONS TESTS IN EACH HEMISPHERE.



PART II

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HASL FALLOUT PROGRAM DATA



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1. Fallout Deposition

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- 1.1 Monthly Precipitation
 - 1.11 Sr^{90} and Sr^{89} in Monthly Deposition at World Land Sites

Precipitation and dry fallout are collected over monthly periods at stations in the United States and overseas. The samples are analyzed for Sr-90 and Sr-89 when it is expected to be present. A description of the sampling network and available data for each site are given in the <u>Appendix</u>, <u>Section A</u>.

1.12 Other Isotopes at Selected Sites

At a number of stations in the United States, monthly deposition collections were analyzed for radiostrontium and other nuclides of interest to the Atomic Energy Commission. Multinuclide analyses were discontinued as of July 1967 and the complete data reported in HASL-193, p. II-4 through II-25.

Plutonium analyses of monthly deposition are continuing at New York City, Melbourne, Seattle, Honolulu, Salisbury, Durban, and Rio de Janeiro. Available data are given in the <u>Appendix</u>, <u>Section C</u>.

1.2 Radiostrontium Deposition at Atlantic Ocean Weather Stations

Measurements of radicstrontium in precipitation and dry fallout collections at four U. S. Coast Guard Stations in the North Atlantic Ocean are carried out for comparison with land stations in the same latitude band. A description of the stations and available data are given in the <u>Appendix</u>, <u>Section B</u>.

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2. Surface Air Sampling Program

The Health and Safety Laboratory has been collecting surface air particulate samples at stations in the Western Hemisphere since January 1963. The filters are analyzed for a number of fission and activation product radionuclides. A description of the program and available data are given in the <u>Appendix</u>, <u>Section D</u>.

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3. Project Airstream

The Health and Safety Laboratory measures radioactivity in the lower stratosphere employing the RB-57F aircraft as a sampling platform. The aircraft are flown by the 58th Weather Reconnaissance Squadron under the direction of the 9th Weather Wing of the Air Weather Service. The missions are scheduled for early February, May, August and November and the coverage extends from $75^{\circ}N$ to $51^{\circ}S$ latitude in the Western Hemisphere. Air filter samples are collected from 12 to 19 km altitude and analyzed for ten radionuclides. A more complete description of the program and available data are given on pages II-9 to II-67.

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4. High Altitude Balloon Sampling Program

Balloon borne filtering devices are used to collect nuclear debris at altitudes from 24-41 km. Balloon launchings are conducted quarterly at Fairbanks, Alaska, 65[°]N; San Angelo, Texas, 31[°]N; Panama C.Z., 9[°]N; and Mildura, Australia, 34[°]S. Filters are analyzed for ten radionuclides. A more complete description of the program and available data were presented in HASL-217 cn pages II-148 to II-206. Corrected sampling dates for three samples are given in this report beginning on page II-68.

5. Radiostrontium in Milk and Tap Water

Strontium⁹⁰ levels in both powdered and fresh milk distributed in New York City and tap Water sampled at the Health and Safety Laboratory, have been measured on a monthly basis since 1954. These data are summarized in tabular and graphical form in the <u>Appendix</u>, <u>Section E</u>.

6. <u>Strontium⁹⁰ in Diets</u>

Quarterly estimates of the annual dietary intake of Sr⁹⁰ of New York City, Chicago, and San Francisco residents have been made based on analyses of foods purchased at these three cities every three months since 1960. Sampling in Chicago has been discontinued. The program is described and available data reported on p. II-4 to II-6 of this report.

7. <u>UNSCEAR - WHO Bone Program</u>

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Available Sr⁹⁰ data for human bone samples collected in 1969 from countries in Latin America and Africa are presented on pages II-7 and II-8 of this report.

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6. HASL Diet Studies: Fourth Quarter 1969

by J. Rivera, (HASL)

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Results of the measurements of the Sr⁹⁰ content of foods purchased in New York City and San Francisco in the fourth quarter of 1 are given in the following table. Estimates of the daily intake of based on these measurements and on the revised consumption statistic given in a recently available U. S. Department of Agriculture Report are also listed.

The estimates of daily Sr^{90} intake are a continuation of the HAS Tri-City diet studies which were started in March of 1960. Results c the earlier measurements along with those made during 1969 are shown graphically in the figure on page II-6. A complete description of th sampling methods and philosophy of the HASL diet studies was given in HASL-200⁽³⁾.

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STRONTIUM⁹⁰ IN NEW YORK CITY AND SAN FRANCISCO DIETS

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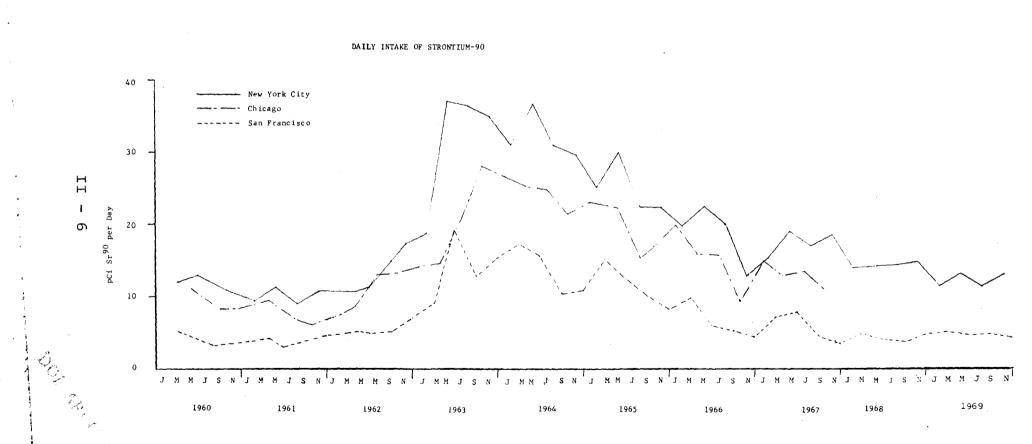
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	Diet Category	kq/yr.	gCa yr	% of yearly intake of_Ca	pCi Sr90	pCi Sr90	% of yearly intake of Sr-90	pCi Sr ⁹⁰	pCi Sr90 yr	% of yearly intake of Sr-90
	DAIRY PRODUCTS	200	216.0		9.2	1840		2.1	420	
				58			38			27
	FRESH VEGETABLES	48	18.7		18.1	869		4.0	192	
	CANNED VEGETABLES	22	4.4		8.2	180		3.4	75	
	ROOT VEGETABLES	10	3.8		7.8	78		4.5	45	
	POTATOES	38	3.8		6.5	247		1.1	42	
	DRY BEANS	3	2.1		36.8	110		11.1	33	
				9			31			24
	FRESH FRUIT	59	9.4		10.5	620		3.1	183	
	CANNED FRUIT	11	0.6		1.7	19		1.1	12	
н	FRUIT JUICES	28	2.5		3.7	104		1.8	50	
н				3			15			16
ł	DAVEDU DOODUGEG	44	F0 7		6 5	286		4.4	194	
ഗ	BAKERY PRODUCTS FLOUR	44 34	53.7		6.5 5.5	187		4.4 3.8	129	
•			6.5					5.8	75	
	WHOLE GRAIN PRODUCTS	11	10.3		16.2	178				
	MACARONI	3	0.6		4.0	12		3.1	9	
	RICE	3	1.1	20	1.7	5		2.5	8	26
				20			14			20
	MEAT	79	12.6		0.5	40		0.4	32	
	POULTRY	20	6.0		0.7	14		1.7	34	
	EGGS	15	8.7		1.7	26		2.5	38	
	FRESH FISH	8	7.6		0.3	2		0.3	2	
-	SHELL FISH	1	1.6		1.6	2		0.6	1	
-				1.0						7
				10			2			7
.#	YEARLY INTAKE		370			4819			1574	
	DAILY INTAKE = pCi/g Ca					13.0			4.2	



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John H. Harley

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In the work of the United Nations Scientific Committee on the Effects of Atomic Radiation, the major effort in evaluating fallout from weapons tests is directed towards estimating the dose commitment. A major contributor to bone dose is strontium-90 but there have not been adequate data for many regions in the world. The Committee decided that it would be useful to have even limited data for portions of the world not otherwise covered. It was agreed that some estimate could be made based on a small number of samples of adult bone and they requested the World Health Organization to assist in procuring samples.

Dr. E. I. Komorov of WHO has been coordinating this program. Samples for 1969 have been received from Senegal, Jamaica, Chile and Venezuela. Negotiations are under way to obtain additional material from other countries in Africa and from Asia. The samples are being analyzed by the Health and Safety Laboratory and the results are presented here as a tabulation of the available data.

Dr. Eduardo Penna-Franca of Brazil has also supplied samples, and they are included in the tabulation for comparison.

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			BONE FROM THE W	HU	
	_	pCi Sr ⁹⁰		-	pCi Sr ⁹⁰
<u>Aqe (yr.)</u>	Sex	per q Ca	Age (yr.)	Sex	per q Ca
	CHILE		VENE	ZUELA	
91	м	2.0	56	м	0.5, 0.6*
58	м	1.7	41	F	0.4
74	м	2.3, 2.0*	32	м	0.7
46	м	1.5	63	м	0.6
33	м	1.1	60	F	0.8
19	м	2.7	unknown	М	0.6
50	F	2.3	42	F	0.6
51	м	1.3	26	F	0.8
			67	м	0.7
			65	м	0.7
1	BRAZIL		50	F	0.7
			83	F	0.7
40	М	1.0	56	М	0.5
25	М	1.0	70	F	0.7
40	М	1.0	0.2	F	2.5
30	М	1.4, 1.4*	23	м	0.5
31	F	0.7, 0.7*	30	М	0.7
			26	F	0.7
			30	F	1.1
J	AMAICA		unknown	м	0.7
			24	F	0.8, 0.8*
62	F	1.2	25	М	0.7
42	F	1.4	•		
34	м	1.0	SENEO	AL	
60	F	0.6			
36	м	1.5	70	F	0.5
67	М	1.0	58	м	0.9
50	м	1.3	60	F	1.0
61	F	1.2	55	м	0.8, 0.8*
57	м	0.9	50	М	0.8
51	м	0.8, 0.8*	41	F	0.4
46	м	1.1	40	м	1.7
57	F	0.6	43	М	1.5
65	F	1.0	34	М	1.8
19	м	1.6	47	М	1.1
54	F	0.7	35	м	0.8
72	M	1.8	80	F	0.5
23	м	1.3			
28	F	2.2			
22	м	1.1			•
22	F	1.8			
21	M	1.6, 1.6*			
	м	0.8			
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3. PROJECT AIRSTREAM

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by Philip W. Krey (HASL) Michael Kleinman (HASL)

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Project Airstream is HASL's study of radioactivity in the lower tratosphere employing the RB-57F aircraft as a sampling platform. The ircraft are flown by the 58th Weather Reconnaissance Squadron under the direction of the 9th Weather Wing of the Air Weather Service. This project is a continuation of the Defense Atomic Support Agency's Project Stardust except that Airstream's sampling missions are limited to only one per season.

The data in this report cover the missions flown in April, July and October 1969. Previous reports containing results from this program are given in references 1 through 8.

FLIGHT SCHEDULE

Airstream missions are scheduled for February, May, August and November with a ± one month slippage. However, each mission must be completed within a nine day interval. The first Airstream mission was flown in August 1967. The flight trajectory and altitude coverage of an Airstream mission are shown in Figures 3a and 3b, respectively. Prior to February 1969, a large gap in the altitude coverage between 15.2 and 18.3 km existed in the sensitive polar regions of each hemisphere.

II - 9

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Beginning with the February 1969 mission, this gap was partially clo. by sampling at 16.3 km instead of 18.3 km at all latitudes poleward a about 35[°] in both hemispheres.

The coverage in Figure 3b extends almost continuously at the indicated altitudes from 75°N to 51°S latitude except for a slight di continuity between 10°S to 16°S. Each mission is accomplished by con ducting return flights northward and southward from each of the four Air Force Bases of operation:

Eielson AFB	64 ⁰ 40 ' N	147 ⁰ 06'W
Kirtland AFB	35 ⁰ 03 ' N	106 ⁰ 36'W
Albrook AFB	08 ⁰ 57'N	79034 W
Mendoza AFB	32 ⁰ 49'S	68 ⁰ 47'W

AIR FILTER SAMPLES

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Air filter samples are collected along the flight tract at latitude increments of approximately 3 to 4[°] at each of the prescribed altitudes using the U-1 foil system. This system permits the sequential insertion of up to 12 IPC No. 1478 filter papers (diameter 16-3/8 into the sampling duct near the bomb bay on the right side of the aircraft. The volume of air sampled by each filter is calculated by the methods developed under Project Stardust and updated by Krajewski (9), and are reported as standard cubic meters (SCM) under the ICAO standard atmosphere (760 mm Hg and 15°C).

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TOTAL GAMMA AND GAMMA SPECTRUM MEASUREMENTS

Ward of Upon arrival at HASL, the filters are coded, logged and quartered. The entire sample (or a representative fraction if the activity is too high) is folded and placed in a plastic box, 8 cm x 6.5 cm x 3.1 cm deep, for gamma spectrometric analysis on an 8" x 4" NaI (Tl) crystal. The total gamma activity is integrated between 100 Kev and 3.0 Mev, and the gamma concentration is reported as counts per minute (cpm) per 100 SCM on the counting date. The complex spectrum is then submitted for computer resolution by least squares fitting into its component members.

RADIOCHEMICAL ANALYSIS

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Based upon the gamma measurements, fractions of the filters are ticombined into appropriate composites which are sent to contractor laboratories for detailed radiochemical analyses including the following nuclides:

Fe-55Zr-95Pb-210Pu-238Sr-89Ce-141Po-210Pu-239,240Sr-90Ce-144Po-210Pu-239,240

At the present time, Trapelo Division/West formerly Tracerlab of Richmond, California is performing these analyses. Nuclide concentrations from radiochemical analyses are reported as picocuries per 100 standard cubic meters of air (pCi/100 SCM) at collection time. Following a previously established practice, Fe-55 is decay corrected to

II - 11

Mr. Martin

October 15, 1961 which is the average production date of this nucl in the 1961 test series. This is not to convey that all the Fe-55 currently in the stratosphere originated in the 1961 tests. To cor vert pCi/100 SCM to disintegrations per minute per 10^3 standard cut feet multiply by 0.629.

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One standard deviation of the counting error for all data in this report is less than $\pm 20\%$ and usually less than $\pm 10\%$ unless anno tated with the symbols:

- A: One standard deviation of the counting error is between $\pm 20 50\%$.
- B: One standard deviation of the counting error is between $\pm 51 100\%$.
- *: Activity is not detectable. This designation is applied to data when one standard deviation of the counting error is greater than ±100%.
- ?: The nuclide concentration of a specific sample is considered suspect because it is inconsistent with the concentration of the same nuclide in adjacent samples in space and time or because it is inconsistent with other nuclides in the same sample.

The nuclide activity for each sample is corrected for the normal radiochemical parameters such as chemical yield and detection efficiency, but it is important to note that a blank adjustment is also made. The value of the adjustment is determined for each nuclide by analyzing a number of blank samples, that is, samples containing no

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activity. Any activity detected in blank samples represents the contamination introduced into the sample by the laboratory reagents and equipment. The average blank value for each nuclide with its measure of uncertainty is then subtracted from the sample activity.

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The Po-210 activity has been decay corrected for ingrowth from its Pb-210 parent for the interval between its separation date and the day of collection. The reported error reflects the uncertainty of both the Pb-210 and Po-210 measurements.

RESULTS

An error in the volume computations of ten samples from the April 1969 mission was uncovered. The corrected radiochemical concentrations for these ten samples are given in Table 3a. The radiochemical analyses of the individual and composite samples from the July 1969 mission are reported in Table 3b.

The gross gamma and Zr-95 concentrations derived from NaI(T1) gamma spectral analyses of the October 1969 mission are given in Table 3c. Such spectra reflect the combined photopeaks of Zr-95 and its daughter Nb-95 at counting time. To calculate the quantity of Zr^{95} in this mixture, the production date of the fresh fission product debris must be known. The Zr-95 data in Table 3c were calculated on the assumption that most of the Zr-95 in each of three latitude regions of

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the stratosphere were produced by three separate events as follows:

Latitude Region	<u>Nuclear Test</u>
90N - 40N	Chinese test of September 29, 1969
40N - 10N	Chinese test of December 28, 1968
10N - >51S	French thermonuclear tests in mid- 1968

The samples in Tables 3b and 3c are grouped according to the altitude of collection beginning with 19.2 km. Within each altitude group, the samples are then listed with decreasing latitude. The fractions of each individual filter making up the composite are list immediately below the composite sample number. The collection parameters of the composite sample and the contractor laboratory performing the analyses are given prior to the nuclide concentrations.

QUALITY CONTROL

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To evaluate the contractor's performance in radiochemical analyses, HASL routinely submits blind duplicates, blanks and standards. The duplicates are identical composites submitted with different code numbers. The blanks are unexposed filters supplied by the Air Force. Standards are blank filters onto which calibrated solutions of various nuclides have been evaporated.

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Llows: These calibrated solutions are available from a number of sources (i.e., Radiochemistry Center, IAEA, Nuclear Chicago) and are recali-.969 brated at HASL. Generally, the agreement between HASL's measurement and the reported value is very good. HASL does not calibrate for Pb^{210} 68 iddirectly, and the supplier's value is accepted. HASL does calibrate for Po-210, and its evaluation of the Po-210 in a Pb-210 standard is now adopted rather than the equilibrium value from the Pb-210. :he itude The results of the quality control program for the October 1969 he mission are summarized in Table 3d. The standards indicate that the average accuracy of analysis is within ±10% or less. The blank anallisted yses indicate that the contamination introduced by normal handling araformand laboratory procedure is either unmeasurable or insignificant for all the nuclides studied. The duplicate samples show that the precision error of analyses is generally less than ±10% except when the counting error of the measurement becomes the major uncertainty. al-

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A serious exception to this general statement on analytical preds. cision is the Pu-238 and Pu-239 results from sample 2476 which differ widely from the results of its duplicate 2458. While plutonium concce. centrations and ratios from sample 2458 appear reasonable for the ious region of the stratosphere from which it was collected, they do not

II - 15

for sample 2476. The results of the other quality control sample in this and earlier reports attest to the overall reliability of the plutonium analyses. Therefore, it is likely that a sample s of the plutonium fraction of sample 2476 took place during analys although no positive evidence of such a switch was found. A thir duplicate of this sample will be submitted for Pu analyses to ver this conclusion.

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REFERENCES

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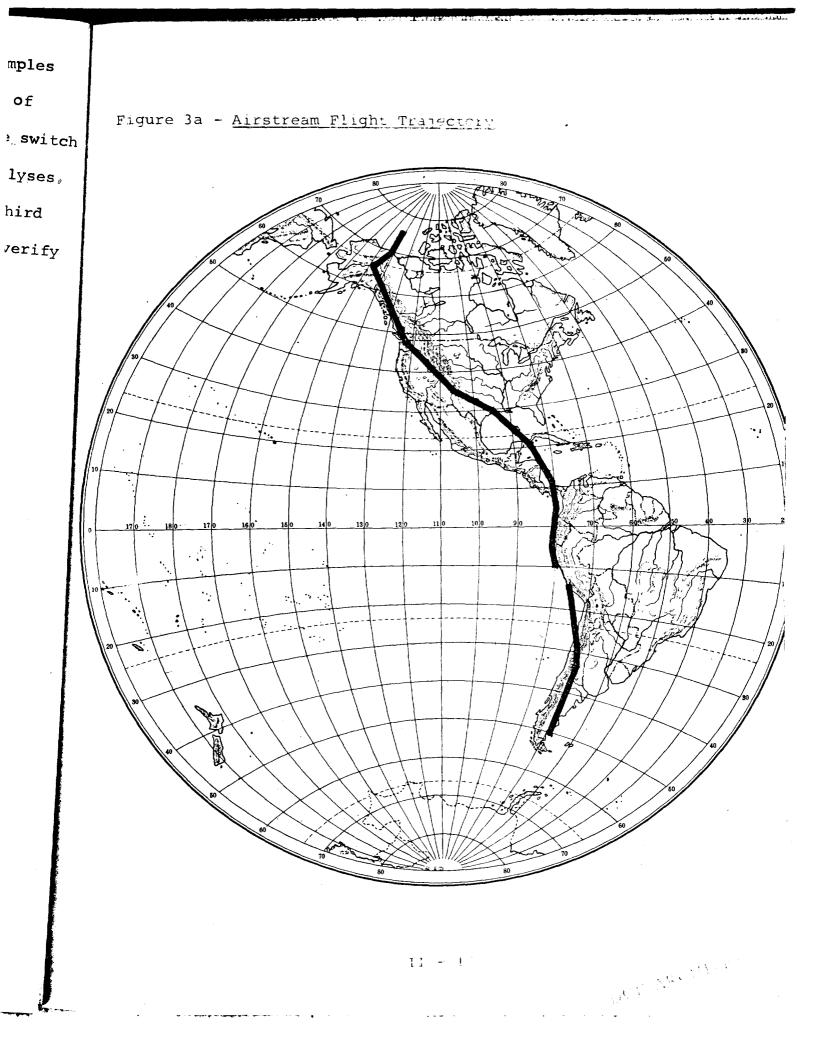
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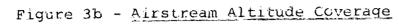
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- (6) Ibid, USAEC Report HASL-207, April (1969)
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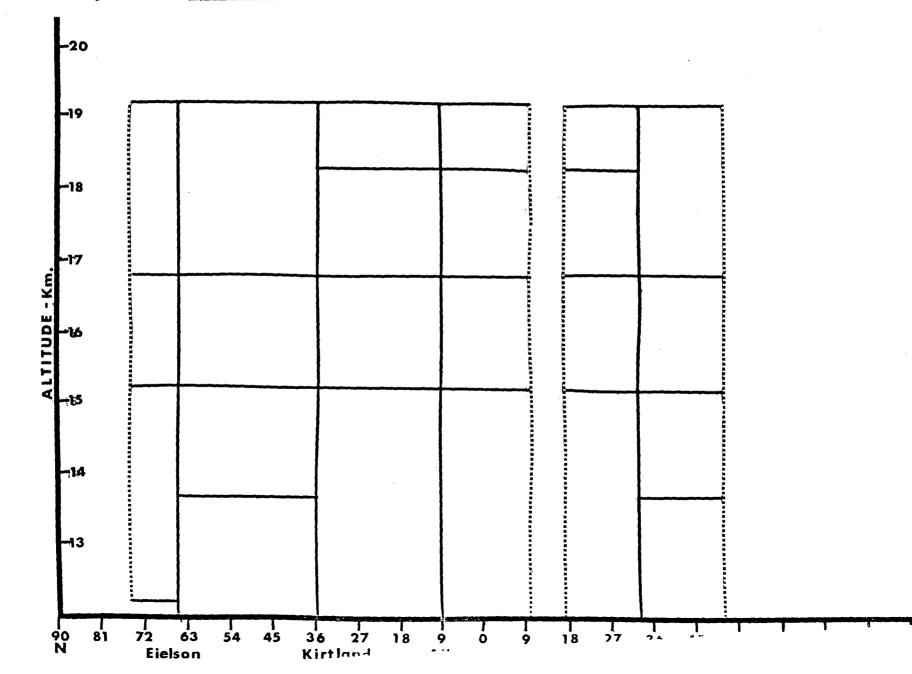
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			TAB	LE JA	
	F	ADIOCHEMICAL			POSITES
			19.2 KM		
SAMPLE NO. Composed of:1	2172 /4:2052 2094 2095 2096	2224 1/4:2052 2094 2095 2096	2173 1/2:2053 2055	2176 1/2:1943 2001	2177 1/2:1944 1945 1946 1947
MIDDOLUT OF	•				
MIDPOINT OF: COLLECTION	• •				
DATE	4/ 8/69	4/ 8/69	4/ 6/69	4/ 9/69	4/ 8/69
LAT. Long.	31N-21N 98W- 86W	31N-21N	21N-12N	75-205	205-305
VOL. OF AIR	3.080	98₩- 86₩ 3⊾080	86W- 81W	80W- 73W	73W- 68W
(100 SCM)	30000	3.000	2.680	4.320	5.720
LAB:	TLW	TLW	TLW	TEW	
FF				16.71	TLW
FE-55 SR-89	698.000	591.000	526.000		737.000
SR-90	3760.000 129.000	3660.000	3290.000	354.000	1150.000
ZR-95		126.000	117.000	64.200	184.000
CE-144	6410.000 3470.000	6530.000	5640.000	814.000	3220.000
PB-210		3360.000	2930.000	1160.000	4200.000
PO-210	0-561A	0.651A	000014	0.657A	0.496
PU-238	0.495	0.487	0.447	0.536	0.420
PU-239	0.443 2.494	0.507	0-468	0.462	1.288
0 237	26774	2.574	.2.214	0.641	2.409

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TABLE 3A

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RADIOCHEMICAL ANALYSIS OF APRIL 1969 COMPOSITES

16.8 KM

SAMPLE NO. 2190	2192
COMPOSED OF: 1/2:1973	1/2:2120
1974	2121
1975	2122
2081	
2082	

MIDPOINT OF:		
COLLECTION DATE	4/ 8/69	4/ 8/69
LAT.	59N-45N	35N-29N
LONG.	136W-117W	106W- 94W
VOL. OF AIR		7.440
(100 SCM)		1.0110
LAB:	TLW	TLW
FE-55	585.000	
SR-89	3500.000	2600.000
SR-90	106.000	81.900
ZR-95	6220.000	4460.000
CE-144	3320.000	2330.000
PB-210	0.516	0.622
PO-210	0.513	0.501
PU-238	0.482	0.229
PU-239	2.339	1.460
A:COUNTING	ERROR IS 20-5	0 PERCENT
	FRROR IS 51-1	

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT *:NOT DETECTABLE

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?:DATA SUSPECT *:

*:NOT DETECTABLE

TABLE 3A

RADIOCHEMICAL ANALYSIS OF APRIL 1969 COMPOSITES

13.7 KM

SAMPLE NO. COMPOSED OF:1/	2216 4:1961	2225 1/4:1961	2217 1/2:1964
	1962	1962	2107
	1963	1963	2101

MIDPOINT OF	•		
COLLECTION			
DATE	4/ 8/69	4/ 8/69	4/ 9/69
LAT.	61N-53N	61N-53N	53N-47N
LONG.	138W-129W	138W-129W	
VOL. OF AIR	5.300	5.300	129W-121W
(100 SCM)		000	9.330
			PC/100 SCM
LAB:	TLW	TLW	TLW
FE-55	779.000	928.000	457.000
SR-89	5980.000	5620.000	3260.000
SR-90	168.000	163.000	98.400
ZR-95	10100.000	9950.000	5510.000
CE-144	5050.000	4880.000	2800.000
PB-210	0.5794	0.431A	0.579
PO-210	0.398	0.397	0.469
PU-238	0.467	0.472	0.228
PU-239	3.289	3.230	1.848
A:COUNTING E	RROR IS 20-50	PERCENT	*:NOT DETECTAR

B:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT 7:DATA SUSPECT *:NOT DETECTABLE

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TABLE 3A

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RADIOCHEMICAL ANALYSIS DF APRIL 1969 COMPOSITES

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12.2 KM

SAMPLE NO. 2223 COMPOSED OF:1/2:1983 1984 1985

MIDPOINT OF: COLLECTION DATE 4/ 7/69 LAT. 75N-65N LONG. 146W-143W VOL. OF AIR 12.740 (100 SCM)

PC/100 SCM

LAB:	
SR-89	TLW (020,000
SR-90	4020.000
ZR-95	127.000
CE-144	7200.000 3590.000
PU-238	
PU-239	0.428 2.579
	24219

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT 7:DATA SUSPECT

*:NOT DETECTABLE

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SAMPLE NO. Composed of:1	2428	DIOCHEMICAL AND	ALYSIS OF JULY 1 19.2 KM	969 COMPOSITES		
		2429	19.2 KM			
		2420				
	2409 2410 2424 2425	2429 1/2:2308 2421 2422 2423	2430 1/2:2300 2309 2310 2311 2312	2431 1/2:2296 2297 2298 2299	2432 1/2:2269 2270 2295	2433 1/2:2271 2272 2273
MIDPOINT OF: COLLECTION DATE LAT. LONG. VOL. OF AIR (100 SCM)	7/23/69 75N-59N 146W-136W 6.160	7/24/69 59N-47N 136W-121W 5.290	7/23/69 47N-35N 121W-106W 6.010	7/23/69 35N-26N 106W- 91W 5.880	7/26/69 26N-18N 91W- 84W 4.090	7/27/69 18N- 9N 84W- 79W 4.110
			PC/100 SCM			
LAB:	TLW	TLW	TLW	TLW	TLW	TLW
SR-90 ZR-95 CE-144 PB-210 PO-210 PU-238 PU-239	67.200 688.000 1090.000 0.376A 0.166B 0.582 1.222	93.200 1250.000 1700.000 0.484A 0.361A 0.608 1.815	97.400 1180.000 1490.000 0.602A 0.300A 0.542 1.677	93.200 1390.000 1830.000 0.604A 0.344A 0.400 1.697	74.200 1100.000 1470.000 0.679A 0.472A 0.299 1.337	78.000 1220.000 1530.000 0.804A 0.675 0.356 1.492
B:COUNTING ER	ROR IS 51-100		*:NOT DETECTABLE	E		
	COLLECTION DATE LAT. LONG. VOL. OF AIR (100 SCM) -AB: SR-90 ZR-95 CE-144 PB-210 PO-210 PO-210 PU-238 PU-239 A:COUNTING ER B:COUNTING ER	MIDPOINT OF: COLLECTION DATE 7/23/69 LAT. 75N-59N LONG. 146W-136W VOL. OF AIR 6.160 (100 SCM) 67.200 AB: TLW SR-90 67.200 ZR-95 688.000 CE-144 1090.000 PB-210 0.376A PO-210 0.166B PU-238 0.582 PU-239 1.222	AIDPOINT OF: COLLECTION DATE 7/23/69 LAT. 75N-59N LONG. 146W-136W 136W-121W VOL. OF AIR 6.160 100 SCM) 5.290 AB: TLW SR-90 67.200 93.200 ZR-95 688.000 1250.000 CE-144 1090.000 1700.000 PB-210 0.376A 0.484A PO-210 0.166B 0.361A PU-238 0.582 0.608 PU-239 1.222 1.815	MIDPOINT OF: CDLLECTION DATE 7/23/69 LAT. 75N-59N LONG. 146W-136W JONG. 121W-106W VOL. OF AIR 6.160 SR-90 67.200 SR-90 67.200 SR-90 67.200 SR-90 67.200 SR-95 688.000 L250.000 1180.000 CE-144 1090.000 PO-210 0.376A O.376A 0.484A O.300A PU-238 0.582 O.4008 0.542 PU-239 1.222 1.815 1.677	AIDPOINT OF: COLLECTION DATE 7/23/69 7/23/69 7/23/69 LAT. 75N-59N 59N-47N 47N-35N 35N-26N LONG. 146W-136W 136W-121W 121W-106W 106W-91W VDL. OF AIR 6.160 5.290 6.010 5.880 (100 SCM) PC/100 SCM AB: TLW TLW TLW TLW SR-90 67.200 93.200 97.400 93.200 R-95 688.000 1250.000 1180.000 1390.000 ZE-144 1090.000 1700.000 1490.000 1830.000 2B-210 0.376A 0.484A 0.602A 0.604A 2O-210 0.166B 0.351A 0.300A 0.344A 2U-238 0.582 0.608 0.542 0.400 2U-239 1.222 1.815 1.677 1.697	AIDPOINT OF: COLLECTION DATE 7/23/69 7/23/69 7/23/69 DATE 7/23/69 7/23/69 7/26/69 LAT. 75N-59N 59N-47N 47N-35N 35N-26N 26N-18N LONG. 146W-136H 136W-121H 121W-106H 106H-91H 91H-84H VOL. OF AIR 6.160 5.290 6.010 5.880 4.090 (100 SCM) PC/100 SCM AB: TLW TLW TLW TLW TLW SR-90 67.200 93.200 97.400 93.200 74.200 2R-95 688.000 1250.000 1180.000 1390.000 1100.000 2E-144 1090.000 1700.000 1490.000 1830.000 1470.000 2B-210 0.376A 0.484A 0.602A 0.604A 0.679A 20-238 0.562 0.608 0.542 0.400 0.299 20-239 1.222 1.815 1.677 1.697 1.337 A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE #:NOT DETECTABLE

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SAMPLE NO. Composed of	2434 :1/2:2259 2260	2435 1/2:2256 2257 2258	TABLE ANALYSIS OF JULY 19.2 KM 2436 1/2:2346 2347 2348	3B 1969 COMPOSITES 2437 1/2:2349 2350 2366 2367	2438 1/2:2362 2363 2364 2365
MIDPOINT OF: COLLECTION DATE LAT. LONG. VOL. OF AIR (100 SCM)	7/25/69 9N- 1N 80W- 79W 3.380	7/25/69 1N-11S 81W- 78W 5.430	7/24/69 15S-26S 75W- 71W 4.310	7/24/69 265-405 71W- 68W 5°750	7/23/69 405-515 68W- 67W 4020
LAB: SR-90 ZR-95 CE-144 PB-210 PU-238 PU-239 A:COUNTING ERR B:COUNTING ERR ?:DATA SUSPECT	TLW 69.000 615.000 1090.000 0.676A 0.539A 0.335 1.070 OR IS 20-50 OR IS 51-100	TLW 55.600 511.000 922.000 0.711 0.433A 0.351 0.867 PERCENT	PC/100 SCM TLW 113.000 577.000 1720.000 0.655A 0.576A 0.728 1.334 *:NOT DETECTABLE	TLW 128.000 637.000 2020.000 0.762 0.230B 0.829 1.442	TLW 140.000 711.000 2170.000 0.404A 0.287A 1.002 1.745

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RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

18.3 KM

SAMPLE NO. 2439	2440	2441	2442	2443	2444
COMPOSED OF:1/2:2291	1/2:2267	1/2:2265	1/2:2252	1/2:2253	1/2:2343
2292	2294	2266	2277	2254	2344
2293		2276		2255	2345
2329					

MIDPOINT OF:						
COLLECTION DATE	7/24/69	7/25/69	7/27/69	7/26/69	7/25/69	7/24/69
LAT.	35N-26N	26N-18N	18N- 9N	9N- 1N	1N-11S	155-265
LONG.	106W- 91W	91W- 84W	84W- 79W	80W- 79W	81W- 78W	73W- 71W
VOL. OF AIR	7.690	2.800	4.060	3.790	6.570	5.820
(100 SCM)						
			PC/100 SCM			
LAB:	TLW	TLW	TLW	TLW	TLW	TLW
SR-90	86.400	65.500	61.200	44.900	42.200	70.500
ZR-95	1460.000	1080.000	951.000	521.000	466.000	463.000
CE-144	1800.000	1280.000	1220.000	777.000	699.000	1110.000
PB-210	0.698	1.200	1.021	0.948	0.865	1.161
PO-210	0.487	0.0 B	0.521A	0.306B	0.484A	*
PU-238	0.311	0.254	0.208	0.194	0.220	0.411
PU-239	1.559	1.203	1.058	0.722	0.649	0.910

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

18.3 KM

	SAMPLE NO.		
	COMPOSED OF:1	/2:2342 2354	
		2004	
	MIDPOINT OF:		
	COLLECTION		
	DATE	7/25/69	
	LAT.	265-335	
н	LONG.	71W- 68W	
н	VOL. OF AIR	3.930	
I	(100 SCM)		
26			
	LAB:	TLW	
	SR-90	106.000	
	ZR-95	600.000	
	CE-144	1850.000	
	PB-210	0.610A	
	PO-210	0.331A	
	PU-238 PU-239	0.739 1.348	
	PU-239	1.540	
	A:COUNTING ER	ROR IS 20-50 PERCENT	*:1
		ROR IS 51-100 PERCENT	
	?:DATA SUSPEC	. T	

NOT DETECTABLE

PC/100 SCM

RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

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16.8 KM

SAMPLE NO. COMPOSED OF:1	2446 1/4 : 2404	2475 1/4:2404	2447 1/2:2417	2448 1/2:2303	2449	2450
	2405 2406 2407	2405 2406 2407	2418 2419 2420	2304 2305 2306 2307	1/2:2320 2321 2322 2323 2328	1/2:2245 2246 2319
MIDPOINT OF: COLLECTION						
DATE	7/22/69	7/22/69	7/24/69	7/23/69	7 / 2 / / / 6	
LAT.	75N-61N	75N-61N	61N-50N	50N-39N	7/24/69 39N-26N	7/26/69
LONG.	150W-143W	150W-143W	138W-125W	125W-108W	108W- 91W	26N-18N 91W- 841
VOL. OF AIR (100 SCM)	3.820	3.820	7.530	10.090	13.850	6°950
LUU JUNI			PC/100 SCM			
LAB:	TLW	TLW	TLW	TLW	TLW	TLW
5R-90	97.700	103.000	106.000	40.400	47.900	
R-95	1390.000	1470.000	1540.000	721.000	803.000	46.600 714.000
E-144	1790.000	2000.000	1980.000	912.000	1030.000	951.000
B-210	0 . 569A	0.551A	0.790	1.188	0.975	1.30
0-210	0.295A	0.388B	0.396A	0°293B	0.443	0.434
	0.452	0.525	0.430	0.119	0.156	0.120
238 20-239	1.726	1.864	1.816			

RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES 16.8 KM SAMPLE NO. 2451 2452 2453 2454 2455 2340A 1/2:2237 COMPOSED 0F:1/2:2247 1/2:2234 1/2:2336 1/2:2338 1/2:2340 2238 2337 2248 2235 2339 2249 2236 MIDPOINT OF: COLLECTION DATE 7/27/69 7/23/69 7/23/69 7/24/69 7/24/69 7/24/69 LAT. 9N- 1N 1N-115 155-235 235-295 295-335-18N- 9N НН LONG. 84W- 79W 80W- 79W 81W- 78W 75W- 72W 72W- 69W 69W- 68W 7.190 5.800 8.180 5.580 3.920 2.780 VOL. OF AIR (100 SCM) N PC/100 SCM ä LAB: TLW TLW TLW TLW TLW TLW 37.300 30.100 SR-90 22.400 9.265 12.400 16.000 ZR-95 358.000 126.000 141.000 135.000 240.000 159.000 218.000 255.000 CE-144 489.000 167.000 588.000 476.000 1.280 1.448 0.947 PB-210 1.050 1.065 * 0.413A P0-210 0.329A 0.216B * 0.099 0.213 0.221 PU-238 0.056 0.052 0.059 0.392 0.172 0.182 0.207 0.429 0.399 PU-239 A:COUNTING ERROR IS 20-50 PERCENT ***:NOT DETECTABLE** B:COUNTING ERROR IS 51-100 PERCENT

TABLE 3B

?:DATA SUSPECT

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RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

16.8 KM

SAMPLE NO.	2 3 53A	2456	2360A	2361A
COMPOSED OF:	1/2:2353	1/2:2357	1/2:2360	1/2:2361
		2358		
		2359		

COLLECTION DATE	7/25/69	7/23/69	7/23/69	7/23/69
LAT.	335-375	375-465	465-495	49 S -51S
LONG.	69W- 68W	69W- 67W	67W- 67W	67W- 67W
VOL. OF AIR	3.030	5.420	1.680	1.210
(100 SCM)				
			PC/100 SCM	
LAB:	TLW	TLW	TLW	TLW
SR-90	126.000	66.100	141.000	99.100
ZR-95	548.000	361.000	670.000	514.000
CE-144	2060.000	1010.000	2270.000	1570.000
PB-210		0.797		
PD-210		0.481A		
PU-238	0.837	0.418	1.017	0.657
PU-239	1.470	0.743	1.638	1.214

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

II -

RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

15.2 KM

SAMPLE NO. Composed of:	2457 1/2:2388 2395 2396 2397	2458 1/4:2385 2386 2387	2476 1/4:2385 2386 2387	2459 1/2:2284 2285 2384	2460 1/2:2281 2282 2283	2461 1/2:2315 2316 2327 2330
MIDPOINT OF: COLLECTION H DATE LAT. LONG. WOL. OF AIR (100 SCM)	7/24/69 75N-61N 147W-138W 12.740	7/25/69 61N-53N 138W-129W 3.810	7/25/69 61N-53N 138W-129W 3∘810	7/24/69 53N-45N 129W-117W 9.110	7/24/69 45N-39N 117W-108W 8.110	7/25/69 39N-29N 106W- 94W 15.270
			PC/100 SCM			
LAB:	TLW	TLW	TLW	TLW	TLW	TLW
SR-90 ZR-95 CE-144 PB-210 PO-210 PU-238 PU-239 A:COUNTING EI B:COUNTING EI	98.300 1600.000 1880.000 0.897 0.498 0.378 1.926 RROR IS 20-50 RROR IS 51-100	82.800 1480.000 1700.000 0.920 0.333B 0.223 1.477 ₽ERCENT ≠	80.600 1350.000 1650.000 0.813A 0.380A 0.692 ? 0.718 ?	53.200 945.000 1210.000 1.007 0.405A 0.136 0.959	19.200 328.000 383.000 1.181 0.258A 0.042 0.314	14.700 251.000 328.000 1.254 0.3174 0.043 0.280

?:DATA SUSPECT

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RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

15.2 KM

SAMPLE NO. Composed of:1	2462 /2:2244 2317 2318	2463 1/2:2241 2242 2243	2464 1/2:2229 2230 2275 2278	2465 1/2:2231 2232 2233	2335A 1/2:2335	2466 1/2:2332 2333 2334
MIDPOINT OF: COLLECTION DATE LAT. LONG. VOL. OF AIR (100 SCM)	7/25/69 29N-21N 94W- 86W 10.730	7/27/69 21N-12N 86W- 81W 9.570	7/25/69 12N- 1N 81W- 79W 11.680	7/23/69 1N-11S 81W- 78W 11.280	7/24/69 15S-20S 75W- 73W 5.000	7/24/69 20S-29S 73W- 69W 10.000
			PC/100 SCM			
LAB:	TLW	TLW	TLW	TLW	TLW	TLW
SR-90 ZR-95 CE-144 PB-210 PO-210 PU-238 PU-239	9.891 159.000 200.000 1.162 0.337A 0.032 0.192	4.293 76.400 98.300 1.585 0.283A 0.014 0.078	2.362 34.700 41.800 0.924 0.157A 0.008A 0.025	1.206 14.500 19.700 0.720 * 0.003B 0.018	2.429 18.500 42.300 0.015A 0.038	9.514 61.600 144.000 1.058 0.436A 0.059 0.117
A:COUNTING ER B:COUNTING ER			*:NOT DETECTABL	C		

?:DATA SUSPECT

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RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

15.2 KM

SAMPLE NO. 2352A	2467	2468	2469
COMPOSED OF:1/2:2352	1/2:2378	1/2:2376	1/2:2374
	2379	2377	2375

MIDPOINT OF:				
COLLECTION				
DATE	7/25/69	7/23/69	7/23/69	7/23/69
LAT.	2.95-335	335-405	40S-46S	46S-51S
LONG.	69W- 68W	69W- 68W	68W- 67W	67W- 67W
VOL. OF AIR	3.390	7.920	6.060	4.420
(100 SCM)				
,			PC/100 SCM	
LAB:	TLW	TLW	TLW	TLW
SR-90	54.600	22.100	94.100	135.000
ZR-95	259.000	109.000	491.000	682.000
CE-144	876.000	322.000	1550.000	2000.000
PB-210		0.905	0.678	0.876
P0-210		0.4244	0°4364	0.2898
PU-238	0.356	0.136	0.625	0.921
PU-239	0.607	0.229	1.067	1.546

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:**DATA SUSPECT •

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			TABLE	38		
		RADIOCHEMICAL AN	ALYSIS OF JULY	1969 COMPOSITES	5	
			13.7 KM			
SAMPLE NO. COMPOSED OF:	2470 1/2:2380 2381 2382 2383	2286A 1/2:2286	2471 1/2:2287 2288 2289	2369A 1/2:2369	2472 1/2:2370 2371	2473 1/2:2372 2373
MIDPOINT OF: COLLECTION DATE LAT. LONG. VOL. OF AIR	7/25/69 61N-50N 138W-125W 13.380	7/24/69 50N-47N 125W-121W 4•250	7/24/69 47N-41N 121W-111W 10.770	7/23/69 375-40S 69W- 68W 3.600	7/23/69 40S-46S 68W- 67W 6.710	7/23/69 465-515 67W- 67W 5•440
ፚ (100 SCM)			PC/100 SCM			2.440
LAB:	TLW	TLW				
		1 L W	TLW	TLW	TLW	TLW
SR-90 ZR-95 CL-144 PB-210 PD-210	62.300 1030.000 1240.000 0.985 0.441	9.612 166.000 222.000	3.319 56.400 71.200 1.081 0.141B	9.979 - 53.700 151.000	18.700 101.000 307.000 0.900 0.378A	52.800 257.000 750.000 0.876 0.479A
₽U-238 ₽U-239	0.149	0.028A 0.193	0.012A 0.061	0.054 0.113	0.142	0.334 0.584

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RADIOCHEMICAL ANALYSIS OF JULY 1969 COMPOSITES

12.2 KM

SAMPLE NU. 2474	
COMPOSED OF:1/2:2392	
2393	
2394	

MIDPOINT OF: COLLECTION DATE LAT. LONG. VOL. OF AIR (100 SCM)	7/24/69 75N-65N 146W-143W 13.690
LAB:	TLW
SR-90	64.700
ZR-95	1190.000
CE-144	1410.000
PB-210	
PO-210	0.915
PU-238	0.493
	0.177
PU-239	1.304
A:COUNTING ERI B:COUNTING ERI	ROR IS 20-50 PERCENT ROR IS 51-100 PERCENT

?:DATA SUSPECT

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1 , 4 *:NOT DETECTABLE

PC/100 SCM

ALTITUDE 19.2 KM

SAMPLE NO.	2747	2748	2749	2764	2763	2762
FLIGHT NO.	288	288	288	296	296	296
DATE	10/13/69	10/13/69	10/13/69	10/14/69	10/14/69	10/14/69
TIME	0008-0044	0044-0117	0117-0145	0022-0110	0002-0022	2332-0002
LAT	75N-71N	71N-68N	68N-65N	65N-61N	61N-59N	59N-56N
LONG	143W-143W	144W-143W	146W-144W	147W-138W	138W-136W	136W-132W
VOL. OF AIR	2 . 90	2.70	2.53	4.23	1.76	2.66
(100 SCM)						2000
GROSS GAMMA/	466。	4 8500。	23600。	920000。	807000。	639000。
M/100 SCM		,				00,0000
H COUNT DATE	12/05/69	12/05/69	12/05/69	12/09/69	12/09/69	11/21/69
H						11/21/0/
1						
ω			PC/100 SCM			
ῶ ZR-95	163	18400	8160	337000	302000	257000
				55.000	502000	227000

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA** SUSPECT

				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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	SAMPLE ND. FLIGHT ND. DATE TIME LAT. LONG. VOL. OF AIR	2761 296 10/14/69 2304-2332 56N-53N 132W-129W 2°45	2760 296 10/14/69 2230~2304 53N~50N 129W-125W 2°80	2494 292 10/17/69 1830~1906 50N~47N 125₩-121W 3₀11	2495 292 10/17/69 1906-1932 47N-45N 121W-117W 2.27	2496 292 10/17/69 1932-1957 45N-43N 117W-114W 2°23	2497 292 10/17/69 1957-2022 43N-41N 114W-111W
	(100 SCM) GROSS GAMMA/ M/100 SCM	165000.	211000.	140000 o	367000。	260000.	2°22 131000°
	COUNT DATE	12/05/69	12/05/69	12/05/69	12/05/69	12/05/69	12/05/69
1	ZR-95	28100	73900	PC/100 SCM 49000	135000	96000	48300

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

*:NOT DETECTABLE

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR~95	CONCENTRATIONS	IN	OCTOBER	1969

SAMPLE NO. FLIGHT NO. DATE IIME LAT. LONG. VOL. OF AIR (100 SCM)	2498 292 10/17/69 2022-2045 41N-39N 111W-108W 2.05	2499 292 10/17/69 2045-2122 39N-35N 108W-106W 3.30	2525 298 10/13/69 2145-2221 39N-35N 108W-106W 3∘35	2514 293 10/13/69 2055-2134 35N-33N 106W-102W 3.55	2513 293 10/13/69 2022-2055 33N-31N 102W-98W 3.00	2512 293 10/13/69 1952-2022 31N-29N 98W- 94W 2.73
GROSS GAMMA/ M/100 SCM	48800.	70900.	2670.	848。	493 _°	897。
COUNT DATE	12/05/69	12/05/69	11/13/69	12/01/69	12/01/69	12/01/69
ZR-95	11400	16600	PC/100 SCM 435	216	134	248

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

***:NOT DETECTABLE**

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			TABLE 3C			
TOTAL GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM)	2511 293 10/13/69 1921-1952 29N-26N 94W- 91W 2.82	2510 293 10/13/69 1855-1921 26N-24N 91W- 89W 2.36	2637 291 10/16/69 1723-1759 24N-21N 89W- 86W 3.38	2638 291 10/16/69 1759-1832 21N-18N 86W- 84W 3.08	2639 291 10/16/69 1832-1901 18N-15N 84W- 82W 2.71	2640 291 10/16/69 1901-1929 15N-12N 82W- 81W 2°65
H GROSS GAMMA/	97 9。	1120.	1060.	990.	1210.	1020.
I COUNT DATE ω ω	12/01/69	11/12/69	11/19/69	11/19/69	11/19/69	11/19/69
ZR-95	267	244	PC/100 SCM 239	223	291	246

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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*:NOT DETECTABLE

				TABLE 3C			
TOTAL G	AMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

SAMPLE ND. Flight ND.	2641 291	2740 291	2739 291	2738 291	2737 291	2736
DATE	10/16/69	10/13/69	10/13/69	10/13/69	10/13/69	291 10/13/69
TIME LAT.	1929-2000 12N- 9N	1850-1922 9N- 5N	1815-1850 5N- 1N	1740-1815 1N- 3S	1703-1740 35- 75	1626-1703 7S-11S
LONG.	81W- 79W	80W- 79W	80W- 80W	81W- 80W	81W- 80W	80W- 78W
VOL. OF AIR (100 SCM)	2.93	2.90	3.24	3.24	3.38	3.51
H GROSS GAMMA/	809.	772。	639。	562.	648.	641.
COUNT DATE	11/19/69	12/04/69	11/23/69	11/22/69	11/22/69	11/22/69
			PC/100 SCM	1		
ZR-95	202	215	153	135	140	116

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

			ALITIOUE 19.2	Λ Π		
SAMPLE NO.	2615	2616	2617	2618	2619	2598
FLIGHT NO.	286	286	286	286	286	289
DATE	10/15/69	10/15/69	10/15/69	10/15/69	10/15/69	10/14/69
TIME	1552-1644	1644-1712	1712-1726	1726-1803	1803-1840	1852-1925
LAT。	155-205	205-235	235-265	265-295	295-335	335-375
LONG.	76W- 73W	73W- 72W	72W- 71W	71W- 69W	69W- 68W	69W- 68W
VOL. OF AIR	4.86	2.70	1.32	3.55	3.44	3.02
(100 SCM)						
H GROSS GAMMA/	844。	685.	1230。	625.	721。	821.
☐ M/100 SCM						
I COUNT DATE	11/18/69	11/18/69	11/18/69	11/18/69	11/18/69	11/17/69
40						
3						
			PC/100 SCM			
ZR-95	151	123	210	107	115	137

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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TOTAL GAMMA AND ZR-95 CONCENTRATIONS IN OCTOBER 1969

ALTITUDE 19-2 KM

TABLE 3C

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				TABLE 3C		
TOTAL	. GAMMA	AND	ZR-95	CONCENTRATIONS	IN OCTOBER	1969

SAMPLE NO.	2597	2596	2595	2594	2593
FLIGHT NO.	289	289	289	289	289
DATE	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69
TIME	1825-1852	1800-1825	1735-1800	1711-1735	1655-1711
LAT.	375-405	40S-43S	435-465	46 S -49S	495-515
LONG.	69W- 68W	68W- 67W	67W- 67W	67W- 67W	67W- 67W
VOL. OF AIR (100 SCM)	2.40	2.06	2.06	1.91	1.31
GRDSS GAMMA/ M/100 SCM	729.	1040。	966.	1030.	1110.
COUNT DATE	11/17/69	11/17/69	11/17/69	11/17/69	11/17/69

			PC/100 SCM			
ZR-95	115	177	157	172	182	

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

*:NOT DETECTABLE

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	.			TABLE 3C			
IUIAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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н	SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VDL. OF AIR (100 SCM) GROSS GAMMA/	2483 293 10/16/69 1926-2001 34N-33N 106W-102W 3.64 4070.	2505 293 10/13/69 1637-1642 33N-33N 102W-102W 0.54 2440.	2506 293 10/13/69 1642-1718 33N-31N 102W- 98W 3.90	2507 293 10/13/69 1718-1747 31N-29N 98W-94W 3.14	2508 293 10/13/69 1747-1826 29N-26N 94W-91W 4.23	2509 293 10/13/69 1826-1848 26N-24N 91W- 89W 2•38
I	M/100 SCM		24400	1030。	1060.	1030.	1230.
42	COUNT DATE	12/05/69	11/12/69	11/12/69	11/12/69	11/12/69	11/12/69
	ZR-95	966	424	PC/100 SCM 238	252	231	293

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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***:NOT DETECTABLE**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM)	2636 291 10/16/69 1645-1715 24N-21N 89W- 86W 3.53	2635 291 10/16/69 1613-1645 21N-18N 86W- 84W 3.77	2634 291 10/16/69 1545-1613 18N-15N 84W- 82W 3.21	2633 291 10/16/69 1515-1545 15N-12N 82W- 81W 3.45	2729 289 10/17/69 1631-1700 12N-9N 81W-79W 3.36	2728 289 10/17/69 1555-1631 9N- 5N 79W- 79W 4.14
GROSS GAMMA/	980。	833.	1160.	846.	714。	568.
COUNT DATE	11/19/69	11/19/69	11/19/69	11/19/69	11/22/69	11/22/69
ZR-95	249	196	PC/100 SCM 291	200	174	138

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

*:NOT DETECTABLE

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TABLE 3C							
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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SAMPLE NO.	2732	2733	2734	2603	2735	2604
FLIGHT NO.	291	291	291	289	291	289
DATE	10/13/69	10/13/69	10/13/69	10/12/69	10/13/69	10/12/69
TIME	1352-1426	1426-1503	1503-1542	1551-1631	1542-1619	1631-1708
LAT.	5N- 1N	1N- 35	35- 7S	7S-11S	7S-11S	11S-15S
LONG.	80W- 80W	81W- 80W	81W- 80W	80W- 78W	80W- 78W	78W- 76W
VOL. OF AIR	4.02	4.26	4.49	4.78	4.31	4.42
(100 SCM)						
¦ GROSS GAMMA/	463°	404.	419.	404.	457。	493.
M/100 SCM						
COUNT DATE	11/22/69	11/22/69	11/22/69	11/17/69	11/22/69	11/17/69
44						
			PC/100 SCM			
ZR-95	113	89	91	77	85	84

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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				TABLE 3C			
TOTAL G	AMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

SAMPLE NO. FLIGHT NO.	2605 289	2614 286	2606	2613	2607	2612
DATE	10/12/69	10/15/69	289 10/12/69	286 10/15/69	289 10/12∤69	286 10/15/69
TIME LAT.	1708-1756 15S-20S	1458-1541 14S-20S	1756-1827 205-235	1434-1458 205-235	1827-1853	1405-1434
LONG.	76W- 74W	75W- 73W	74W- 72W	73W- 72W	235-265 72W- 71W	235-265 72W- 71W
VOL. OF AIR (100 SCM)	5.62	5.17	3.55	2.88	2.93	3.48
GROSS GAMMA/ M/loo scm	826。	584.	738.	444.	717.	514.
COUNT DATE	11/17/69	11/18/69	11/17/69	11/18/69	11/18/69	11/18/69
30.05			PC/100 SCM			
ZR-95	151	102	127	80	128	82

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

*:NOT DETECTABLE

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			ALTIODE 10.5	KF1
SAMPLE NO.	2608	2611	2609	2563
FLIGHT NO.	289	286	289	289
DATE	10/12/69	10/15/69	10/12/69	10/15/69
TIME	1853-1919	1338-1405	1919-1941	1633-1706
LAT.	265-295	265-295	295-325	295-335
LONG.	71W- 69W	71W- 69W	69W- 68W	69W- 58W
VOL. OF AIR (100 SCM)	2.93	2.99	2.48	3-81
GROSS GAMMA/ M/100 SCM	570.	632.	5.	362.
COUNT DATE	11/18/69	11/18/69	11/18/69	12/01/69
			PC/100 SCM	4
ZR-95	112	107	*	63

ALTITUDE 18.3 KM

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

***:NOT DETECTABLE**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

ALTITUDE 16.8 KM

II - 47	SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM)	2746 288 10/13/69 2331-0004 75N-71N 143W-143W 4.81	2745 288 10/13/69 2304-2331 71N-68N 144W-143W 3.90	2744 288 10/13/69 2237-2304 68N-65N 146W-144W 4.09	2743 288 10/13/69 2147-2237 65N-61N 146W-138W 7.13	2756 296 10/14/69 2013-2033 61N-59N 138W-136W 2.80	2757 296 10/14/69 2033-2105 59N-56N 136W-132W 4.62
	GROSS GAMMA/ M/100 SCM COUNT DATE	1000.	3210.	4520°	127000.	525000°	46300.
		11/22/69	12/03/69	12/05/69	12/05/69	12/05/69	12/05/69
	ZR-95	432	445	PC/100 SCM 1600	46500	200000	18400
	A:COUNTING ER B:COUNTING ER	ROR IS 20-50 ROR IS 51-100	PERCENT	*:NOT DETECTABL	E		

?:DATA SUSPECT

				TABLE 3C			
TOTAL	GAMMA	AND	ZR95	CONCENTRATIONS	IN	OCTOBER	1969

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ALTITUDE 16.8 KM

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SAMPLE NO.	2758	2759	2493	2492	2491	2490
FLIGHT NO.	296	296	292	292	292	292
DATE	10/14/69	10/14/69	10/17/69	10/17/69	10/17/69	10/17/69
TIME	2105-2135	2135-2210	1743-1820	1714-1743	1645-1714	1619-1645
LAT.	56N-53N	53N-50N	50N-47N	47N-45N	45N-43N.	43N-41N
LONG.	132W-129W	129W-125W	125W-121W	121W-117W	117W-114W	114W-111W
H VOL. OF AIR	4.28	4.99	5.28	4.14	4.16	3.69
(100 SCM)						
GROSS GAMMA/	170000。	86200.	6420.	1270.	1250.	1170.
A M/100 SCM						
COUNT DATE	12/05/69	12/05/69	11/12/69	11/12/69	11/12/69	11/12/69
			PC/100 SCM	4		
ZR-95	57700	33100	1320	471	462	451
A - COUNTING C		OFREENT	+-NOT DETECTA	3. F		

A:COUNTING ERROR IS 20-50 PERCENT *: NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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ALTITUDE 16.8 KM

	SAMPLE NO.	2489	2488	2482 293	2549 298	2548 298	2547 298
	FLIGHT NO. DATE	292 10/17/69	292 10/17/69	10/16/69	10/14/69	10/14/69	10/14/69
	TIME LAT.	1555-1619 41N-39N	1534-1555 39N-36N	1847-1919 39N-35N	2051-2130 35N-33N	2016-2051 33N-31N	1945-2016 31N-29N
	LDNG.	111W-108W	108W-108W	108W-107W	106W-102W 6.00	102W- 98W 5.65	98W- 94W 5.03
ΤI	VOL. OF AIR (100 SCM)	3.49	3.27	4.62			
I	GRDSS GAMMA/ M/100 SCM	1280.	642.	939。	930.	1320.	2370.
49	COUNT DATE	11/12/69	11/12/69	11/10/69	11/14/69	11/14/69	11/14/69
	ZR-95	263	144	PC/100 SCM 183	170	205	323
	A:COUNTING EF	ROR IS 20-50	PERCENT	*:NOT DETECTAR	3LE		
		A COUNTING ENROR IS ED JO TEROCAT					

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

ALTITUDE 16.8 KM

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	SAMPLE NO.	2546	2545	2721	2722	2723	2724
	FLIGHT NO.	298	298	291	291	291	291
	DATE	10/14/69	10/14/69	10/15/69	10/15/69	10/15/69	10/15/69
	TIME	1909-1945	1845-1909	2017-2051	2051-2124	2124-2152	2152-2221
	LAT.	29N-26N	26N-24N	24N-21N	21N-18N	18N-15N	15N-12N
	LONG.	94W- 91W	91W- 89W	89W- 86W	86W- 84W	84W- 82W	82W- 81W
ПI	VOL. OF AIR (100 SCM)	5.86	3.90	5.47	5,38	4.66	4.85
1	GROSS GAMMA/	2580.	6130.	10700.	7810.	180.	245。
5 0		11/14/69	11/14/69	12/01/69	11/15/69	11/22/69	11/22/69
				PC/100 SCI	W		
	ZR-95	373	730	2030	1010	34	49
	A:COUNTING EF	ROR IS 20-50	PERCENT	*:NOT DETECTA	BLE		

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

ALTITUDE 16.8 KM

	SAMPLE NO.	2730	2652	2715	2651	2714	2650
	FLIGHT NO.	289	286	291	286	291	286
	DATE	10/17/69	10/16/69	10/13/69	10/16/69	10/13/69	10/16/69
	TIME	1704-1733	1933-1955	1907-1942	1859-1933	1830-1907	1824-1859
	LAT.	12N- 9N	8N- 5N	9N- 5N	5N- 1N	5N- 1N	1N- 3S
	LONG.	81W- 79W	80W- 79W	80W- 79W	80W- 80W	80W- 80W	81W- 80W
	VOL. OF AIR	4.54	3.50	5.74	5.41	6.26	5.57
	(100 SCM)						
	GROSS GAMMA/	405。	160.	103.	99.	79.	101.
н	M/100 SCM						
	COUNT DATE	11/21/69	11/21/69	11/22/69	11/21/69	11/21/69	11/21/69
51							
				PC/100 SCM	4		
	ZR-95	87	35	22	19	17	21

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

***:NOT DETECTABLE**

				ALILIUDE 10.8	К.М.		
	SAMPLE ND.	2713	2649	2712	2648	2711	2647
	FLIGHT NO.	291	286	291	286	291	286
	DATE	10/13/69	10/16/69	10/13/69	10/16/69	10/13/69	10/16/69
	TIME	1752-1830	1746-1824	1714-1752	1657-1746	1626-1714	1628-1657
	LAT.	1N- 3S	35- 75	35- 75	7S-12S	7S-11S	12S-15S
	LONG.	81W- 80W	81W- 80W	81W- 80W	80W- 78W	80W- 78W	78W- 77W
	VOL. OF AIR	6.43	6.01	6.55	7.93	7.99	4.66
	(100 SCM)						
НЦ	GROSS GAMMA/	92.	290.	86。	60.	125.	80.
	M/100 SCM						
1	COUNT DATE	11/22/69	11/21/69	11/22/69	11/21/69	11/22/69	11/21/69
52							
				PC/100 SCM			
	ZR-95	18	47	16	10	21	9
	A:COUNTING ER	ROR IS 20-50	PERCENT	*:NOT DETECTAB	LF		
	B:COUNTING ERROR IS 51-100 PERCENT						

ALTITUDE 16 8 KM

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?:DATA SUSPECT

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			TABLE 3C		
TOTAL GAMMA	AND	ZR-95	CONCENTRATIONS	IN OCTOBER	1969

ALTITUDE 16.8 KM

SAMPLE NO.	2582	2646	2583	2645	2584	2644
FLIGHT NO.	289	286	289	286	289	286
DATE	10/13/69	10/16/69	10/13/69	10/16/69	10/13/69	10/16/69
TIME	1636-1727	1538-1628	1727-1753	1509-1538	1753-1820	1439-1509
LAT.	155-205	15S-20S	205-235	205-235	235-265	235-265
LONG.	76W- 73W	77W- 74W	73W- 72W	74W- 73W	72W- 71W	73W- 71W
VOL. OF AIR	8.31	8.15	4.26	4.80	4.35	4 - 83
(100 SCM)	*					
H GRUSS GAMMA/	91.	15.	55.	42.	80.	88.
H M/100 SCM						
I COUNT DATE	11/15/69	11/21/69	11/15/69	11/21/69	11/15/69	11/21/69
л w						
ω.			PC/100 SCM			
ZR-95	15	67	9	4	13	10

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

				HEITIODE TO'O K	M		
	SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM) GRDSS GAMMA/	2585 289 10/13/69 1820-1848 26S-29S 71W- 70W 4.45	2643 286 10/16/69 1407-1439 265-295 71W- 69W 5.01	2586 289 10/13/69 1848-1923 29S-33S 70W- 69W 5.57	2562 289 10/15/69 1525-1601 335-375 69W-58W 5.48	2588 289 10/14/69 1431-1500 375-40S 69W- 68W 4.12	2589 289 10/14/69 1500-1525 40S-43S 68W- 67W 3.50
H H -	M/100 SCM	153.	147.	294。	651.	944.	991.
1 54	COUNT DATE	11/17/69	11/21/69	11/17/69	12/01/69	11/17/69	11/17/69
	ZR-95	26	25	PC/100 SCM 52	119	158	170
	A:COUNTING FRI	ROR IS 20-EO					

ALTITUDE 16.8 KM

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A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT **?:**DATA SUSPECT

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*:NOT DETECTABLE

				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

ALTITUDE 16.8 KM

SAMPLE NO.	2590	2591	2592
FLIGHT NO.	289	289	289
DATE	10/14/69	10/14/69	10/14/69
TIME	1525-1551	1551-1617	1618-1634
LAT.	43S-46S	46S-49S	495-515
LONG.	67W- 67W	67W- 67W	67W- 67W
VOL. OF AIR	3.54	3.54	2.18
(100 SCM)			
GROSS GAMMA/	1040.	1100.	1070.
M/100 SCM			
COUNT DATE	11/17/69	11/17/69	11/17/69
			PC/100 SCM
ZR-95	178	186	184

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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	SAMPLE NO. FLIGHT NO.	2772 296	2773 296	2774	2790	2789	2788
	DATE	10/13/69		296	288	288	288
L	TIME		10/13/69	10/13/69	10/14/69	10/14/69	10/14/69
		22242302	2302-2330	23302359	004 7 0125	0028-0047	23590028
	LATa	75N-71N	71N-68N	68N-65N	65N-61N	61N-59N	59N-56N
	LONG.	143W-143W	144W-143W	146W-144W	144W-139W	139W-136W	136W-132W
	/OL. OF AIR (100 SCM)	7.31	5.44	5.50	7.52	3.84	5.84
ψ.	GROSS GAMMA/	1420.	5990.	2220。	1640.	654。	1280 _°
	COUNT DATE	11/15/69	11/15/69	11/15/69	11/15/69	11/21/69	11/15/69
Z	<u>1</u> R-95	378	1400	PC/100 SCM 621	424	216	334 、
۵	COUNTING ER	ROR IS 20-50	PERCENT	*: ΝΩΤ <u>DETECTAB</u>	F		

B:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT 7:DATA SUSPECT

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***:NOT DETECTABLE**

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ALTITUDE 15.2 KM

SAMPLE NO.	2787	2786	2533	2532	2531	2530
FLIGHT ND.	288	288	293	293	293	293
DATE	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69
TIME	2328-2359	2255-2328	1859-1941	1827-1859	1759-1827	1731-1759
LAT.	56N-53N	53N-50N	50N-47N	47N-45N	45N-43N	43N-41N
LONG.	132W-129W	129W-125W	125W-121W	121W-117W	117W-114W	114W-111W
H VOL. OF AIR	6.16	6.33	8.13	5.98	5.15	5.43
[⊢] (100 SCM) ∣ GROSS GAMMA/	2820°	1190.	5410。	3080.	1820.	343.
∪ M/100 SCM V COUNT DATE	12/02/69	11/15/69	11/13/69	11/13/69	11/13/69	11/13/69
	•		PC/100 SCM			
ZR-95	991	472	1100	826	504	125
A .COUNTING ER	000 TC 20 E0	DEDCENT	*:NOT DETECTAB	1 6		

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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SAMPLE NO. FLIGHT NO. DATE IIME LAT. LONG. VOL. OF AIR (100 SCM)	2529 293 10/14/69 1705-1731 41N-39N 111W-109W 4.96	2481 293 10/16/69 1810-1843 39N-36N 108W-107W 6.90	2480 293 10/16/69 1702-1749 35N-33N 106W-102W 9.89	2486 292 10/15/69 1659-1742 35N-33N 106W-102W 9.39	2541 298 10/14/69 1635-1710 33N-31N 102W-98W 7.79	2542 298 10/14/69 1710-1740 31N-29N 98W- 94W 6.70
GROSS GAMMA/	381.	306。	138.	301.	146.	28.
COUNT DATE	11/13/69	11/10/69	11/10/69	11/12/69	11/14/69	11/14/69
ZR-95	6.9	65	PC/100 SCM 28	64	20	5

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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*:NOT DETECTABLE

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ALTITUDE 15.2 KM

SAMPLE NO.	2543	2544	2720	2719	2718	2717
FLIGHT NO.	298	298	291	291	291	291
DATE	10/14/69	10/14/69	10/15/69	10/15/69	10/15/69	10/15/69
TIME	1740-1818	1818-1841	1941-2012	1908-1941	1836-1908	1807-1836
LAT.	29N-26N	26N-24N	24N-21N	21N-18N	18N15N	15N-12N
LONG.	94W 91W	91W- 89W	89W- 86W	86W- 84W	84W- 82W	82W 81W
H VOL. OF AIR	8.46	5.18	6.77	7.20	7.10	6.44
(100 SCM)						
GROSS GAMMA/	29。	48.	226。	34.	155。	22。
ы м/100 SCM						
COUNT DATE	11/14/69	11/14/69	11/21/69	11/21/69	11/22/69	11/22/69
			PC/100 SCM			
ZR-95	4	8	34	8	25	3

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	ΙN	OCTOBER	1969

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SAMPLE NO. FLIGHT NO. DATE TIME LAT.	2716 291 10/15/69 1750-1807 12N-10N	2726 289 10/17/69 1440-1512 12N- 9N	2727 289 10/17/69 15121548 9N- 5N	2707 291 10/13/69 1350-1428 5N- 1N	2708 291 10/13/69 1428-1503 1N- 3S	2709 291 10/13/69 1503-1544 3S- 7S
LONG. VOL. OF AIR (100 SCM)	81W- 80W 3.77	81W- 79W 6.90	79W 79W 7.81	80W- 80W 8.78	81W- 80W 7.67	33-75 81W-80W 9°39
H GROSS GAMMA/ M/100 SCM	13.	30.	11.	10.	16.	12.
COUNT DATE	11/22/69	12/05/69	11/23/69	11/26/69	11/26/69	11/26/69
ZR-95	2	5	PC/100 SCM 2	2	2	2

A:COUNTING ERROR IS 20-50 PERCENT *:NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT **?:DATA SUSPECT**

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				TABLE 3C			
TOTAL	GAMMA	AND	ZR-95	CUNCENTRATIONS	IN	OCTOBER	1969

SAMPLE NO.	2710	2581	2580	2579	2578	2561
FLIGHT NO.	291	289	289	289	289	289
DATE	10/13/69	10/13/69	10/13/69	10/13/69	10/13/69	10/15/69
TIME	1544~1626	15401630	1512-1540	1445-1512	1414-1445	1412-1447
LAT.	7S-11S	15S-20S	205-235	235-265	265-295	295-335
LONG .	80W- 78W	76W- 73W	73W- 72W	72W- 71W	71W- 70W	69W- 58W
VOL. OF AIR	9.62	10.96	6.02	5.73	6.45	7.52
(100 SCM)						
🛏 GROSS GAMMA/	18.	18.	33 o	31.	42.	400.
H M/100 SCM						
I COUNT DATE	11/26/69	11/15/69	11/15/69	11/15/69	11/15/69	11/14/69
0						
H						
			PC/100 SCM			
ZR-95	3	3	5	5	6	66

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT *:NOT DETECTABLE

				TABLE 3C		
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN DCTOBER	1969

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SAMPLE NO.	2630	2629	2628	2627	2626	2625
FLIGHT NO.	286	286	286	286	286	286
DATE	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69
TIME	1817-1852	1747-1816	1722-1746	1656-1721	1632-1655	1614-1631
LAT.	33S37S	375-405	40S-43S	43S-46S	46S-49S	49S51S
LONG.	69W 68W	69W- 68W	68W- 67W	67W- 67W	67W- 67W	67W- 67W
VOL. OF AIR	6.75	5.59	4.43	4.49	4.13	3.05
(100 SCM)						
H GROSS GAMMA/	344.	660.	770.	768.	683。	620。
[™] M/100 SCM						
I COUNT DATE	11/19/69	11/19/69	11/19/69	11/18/69	11/18/69	11/18/69
6 N						
			PC/100 SCN	4		
ZR-95	63	115	133	128	121	118

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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***:NOT DETECTABLE**

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ALTITUDE 13.7 KM

	ZR-95	224	292	PC/100 SCM 660	422	279	249
ω 0	M/100 SCM Count date	11/21/69	11/21/69	11/15/69	11/15/69	11/13/69	11/13/69
1	(100 SCM) GROSS GAMMA/	800.	722。	1660.	1530.	725。	645。
ЧI	VOL. OF AIR	6.25	8.41	4.79	11.23	7.45	7.19
	LONG.	139W-136W	136W132W	132W-129W	129W-125W	125W-121W	121W-117W
	LAT.	61N-59N	59N56N	56N-53N	53N-50N	50N-47N	47N-45N
	TIME	2032-2058	2058-2133	2133-2209	2209-2255	1949-2020	2020-2052
	DATE	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69	10/14/69
	FLIGHT NO.	288	288	288	288	293	293
	SAMPLE NO.	2782	2783	2784	2785	2534	2535

A:COUNTING ERROR IS 20-50 PERCENT *: NOT DETECTABLE B:COUNTING ERROR IS 51-100 PERCENT

?:DATA SUSPECT

	- .			TABLE 3C		
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN OCTOBER	1969

ALTITUDE 13.7 KM

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SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM)	2536 293 10/14/69 2052-2119 45N-43N 117W-114W 6°26	293 293 10/14/69 10/14/69 2052-2119 2119-2148 45N-43N 43N-41N 117W-114W 114W-111W 6.26 6.89	2538 293 10/14/69 2148-2211 41N-39N 111W-109W 5.40	2620 286 10/14/69 1348-1421 37S-40S 69W- 68W 7.57	2621 286 10/14/69 1422-1451 40S-43S 68W 67W 6.51	2622 286 10/14/69 1452-1523 43S-46S 67W-67W 6.93
H GROSS GAMMA/ H M/100 SCM	677 _°	916。	741。	439。	521。	620.
າ COUNT DATE ອາ 4	11/13/69	11/14/69	11/14/69	11/18/69	11/18/69	11/18/69
ZR-95	271	225	PC/100 SCM 294	73	88	108
A:COUNTING ER	ROR IS 20-50	PERCENT	*:NOT DETECTABLE	E		

B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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TABLE 3C							
TOTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

ALTITUDE 13.7 KM

SAMPLE NO.	2623	2624
FLIGHT NO.	286	286
DATE	10/14/69	10/14/69
TIME	1524-1551	1552-1610
LAT.	465-495	495-515
LONG.	67W- 67W	67W- 67W
VOL. OF AIR	6.16	3.96
(100 SCM)		
GROSS GAMMA/	515.	530.
M/100 SCM		
COUNT DATE	11/18/69	11/18/69
ZR-95	83	89

PC/100 SCM

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT ***:NOT DETECTABLE**

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SAMPLE NO. FLIGHT NO. DATE TIME LAT. LONG. VOL. OF AIR (100 SCM)	2771 296 10/13/69 2139-2218 75N-71N 143W-143W 10.72	2770 296 10/13/69 2111-2139 71N-68N 144W-143W 7.70	2769 296 10/13/69 2044-2111 68N-65N 146W-144W 7.79
GRDSS GAMMA/ M/100 SCM	534。	210.	263。
COUNT DATE	11/15/69	11/22/69	11/22/69

				TABLE 3C			
TUTAL	GAMMA	AND	ZR-95	CONCENTRATIONS	IN	OCTOBER	1969

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ZR-95	100		PC/100 SCM
21 33	108	44	51

A:COUNTING ERROR IS 20-50 PERCENT B:COUNTING ERROR IS 51-100 PERCENT ?:DATA SUSPECT

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*:NOT DETECTABLE

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Table 3d

QUALITY CONTROL RESULTS

Report Date	Sample No.	Reference Date	_Sr-90_	<u>Zr-95</u>	<u>Ce-144</u>	Pb-210	<u>Po-210</u>	Pu-238	Pu-239
Standards	•				dpm	± % Standard	Deviation		
3/70	2477	7/25/69 added found % deviation	3.26x10 ³ 3.36x10 ³ ±1 +3.1	1020 891±4 -13	661 701±3 +6.0	6.78 7.77±31 +15	6.78 7.07±4 +4.3	2.96 2.96±3 0	4.10 4.4 6± 2 +8.8
44	2478	7/27/69 added found % deviation	3.91x10 ³ 3.42x10 ³ ±1 +12	891 835±4 ~6.3	560 607±2 +8.4	9.76 10. <u>4+</u> 19 +6.6	9.76 11.1±5 +14	2.96 2.56±5 -4.8	3.49 3.63<u>+</u>4 + 4. 0
		Avarage % deviation	+7.5	-9.6	+7.2	+11	+9.2	-2.4	+6.4

	$dpm \pm \%$ Standard Deviation									
Blanks										
2/70	2250A	7/27/69	*	*	2.99±69			*	*	
R	2301A	7/23/69	*	2.74±60	3.0 81 19			*	*	
						pCi/100 SCM				
Duplicates	2446		97.7±1	1390±1	1790±1	0.569±25	0.295±49	0.452±5	1.73±3	
	2475		103±3	1470±1	2000±1	0.55 1± 36	0.388±52	0.525±7	1.86±4	
	% deviat	cion about the average	±5 .3	±5 ,6	+11	±3.2	±27	±15	±7.2	
	2458		82 <i>.</i> 8±1	1480±2	1700±1	0.920±18	0.333±51	0.223±8	1.48±4	
	2476		80.6 ±2	1350±1	1650±1	0.813±22	0.380±48	0.692±4?	0,718±4?	
	% devia:	tion about the average	±2.7	±9.2	±3.0	±12	±13			
	i	Average % deviation	±4.0	±7.4	±7.0	±7.6	±20			

? Data suspect

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High Altitude Balloon Sampling Program

by P. W. Krey (HASL)

In HASL-217, the analyses of all high altitude balloon samples collected during 1968 were summarized. Unfortunately three samples collected in October 1968 at San Angelo, Texas were reported as having been flown in September. This error has been corrected in the accompanying tables which cover both the September and October flights made at San Angelo. None of the radiochemical data has been changed - only the collection dates.

Table 4

STRATOSPHERIC RADIONUCLIDE CONCENTRATIONS

BALLOON SAMPLES COLLECTED DURING SEPTEMBER 1968 LATITUDE, 31N SAN ANGELO, TEXAS

	ALTITUDE (KM)	21	24	27
	FLIGHT DAY	16	01	08
	HASL NUMBER	3011	2999	3001
	COLLECTION UNIT	D7-1	D7-1	D7-1
	ANALYTICAL LABORATORY	TLW	TLW	TLW
	GROSS GAMMA (CPM/KSCM)	1298.0#	1756.7#	608.2
	IRON-55	4200		
	STRONTIUM-89	285A	*	*
	STRONTIUM-90	543	282	97.2
	ZIRCONIUM-95	635	109A	*
-	CERIUM-144	4720	943	118
	POLONIUM-210	1.94A	1.76B	1.04B
	PLUTONIUM-238	8.37	7.03	6.82
	PLUTONIUM-239	13.0	5.80	2.37

A: One Standard Deviation of Counting Error is >20% to 50% of count.

B: One Standard Deviation of Counting Error is >50% to 100% of Count.

*: Standard Deviation Greater than Data Value.

4: Gross Gamma Count More than Two Weeks After Collection.

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Table 4

STRATOSPHERIC RADIONUCLIDE CONCENTRATIONS

BALLOON SAMPLES COLLECTED DURING SEPTEMBER 1968 LATITUDE, 31N SAN ANGELO, TEXAS

32	36	41
09	07	12
3002	3000	3009
AE-1	AE-1	HV3 K
TLW	TLW	WLT
706.2	776.0#	714.1
	PC/KSCM	
*	*	
32.7B	24.0B	61.1A
*	*	
65.0A	*	*
*	*	11.5A
3.88	2.33B	24.6
1.06A	*	.792B
	09 3002 AE-1 TLW 706.2 * 32.7B * 65.0A * 3.88	09 07 3002 3000 AE-1 AE-1 TLW TLW 706.2 776.0 ∰ PC/KSCM * * 32.7B 24.0B * * 65.0A * * 3.88 2.33B

A: One Standard Deviation of Counting Error is 20% to 50% of Count.

B: One Standard Deviation of Counting Error is 50% to 100% of Count.

*: Standard Deviation Greater than Data Value.

#: Gross Gamma Count more than Two Weeks After Collection.

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Table 4

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STRATOSPHERIC RADIONUCLIDE CONCENTRATIONS

BALLOON SAMPLES COLLECTED DURING OCTOBER 1968 LATITUDE, 31N SAN ANGELO, TEXAS

ALTITUDE (KM) FLIGHT DAY HASL NUMBER COLLECTION UNIT ANALYTICAL LABORATORY GROSS GAMMA (CPM/KSCM)	24 30 3045 D7-1 TLW 1187.1 拼	27 28 3035 D7-1 TLW 296.9#	30 29 3036 D7-1 TLW 198.5#
		0644664	
		PC/KSCM	
IRON-55	6460		
STRONTIUM-89	206A	*	
STRONTIUM-90	426	104	28.5
ZIRCONIUM-95	275A	*	
CERIUM-144	2560	165	*
POLONIUM-210	1.22B	2.48A	*
PLUTONIUM-238	7.61	6.77	3.09
PLUTONIUM-239	10.0	1.98	•883A

A: ONE STANDARD DEVIATION OF COUNTING ERROR IS >20% TO 50% OF COUNT. B: ONE STANDARD DEVIATION OF COUNTING ERROR IS >50% TO 100% OF COUNT. *: STANDARD DEVIATION GREATER THAN DATA VALUE #: GROSS GAMMA COUNT MORE THAN TWO WEEKS AFTER COLLECTION

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Part III

DATA FROM SOURCES OTHER THAN HASL

Numerous Fallout studies are conducted by other organizations in the United States and abroad. Some of these are sent to the editors for dissemination in these HASL Quarterly reports. Submitted data are reproduced essentially as received and no interpretation by HASL is attempted.

		Page
1.	National Radiation Laboratory, Department of Health Christchurch, New Zealand Environmental Radioactivity in New Zealand	
	Quarterly Report for April-June 1969: NRL-F35	III-2
	Quarterly Report for July-September 1969: NRL-F36	III-14
2.	Department of Scientific and Industrial Research The Institute of Nuclear Sciences Lower Hutt, New Zealand	III-24
	Radioisotopes in Rainwater: January - April 1968 May - August 1968 September - December 1968	
3.	Radiological Physics Division Argonne National Laboratory Cesium-137 in Various Chicago Foods (Coll. Month January 1970) by S. S. Brar and D. M. Nelson	III-28
4.	EURATOM Joint Nuclear Research Centre Ispra Establishment, Protection Service Site Survey and Meteorology Section, Quarterly	00

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Report

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REPORT No. NRL-F 35

DEPARTMENT OF HEALTH



QUARTERLY REPORT APRIL - JUNE 1969

ENVIRONMENTAL RADIOACTIVITY IN NEW ZEALAND

AND

MEASUREMENTS ON SAMPLES FROM FIJI AND RAROTONGA

NATIONAL RADIATION LABORATORY P.O. BOX 1456, CHRISTCHURCH, NEW ZEALAND

SYMBOLS UNITS AND EQUIVALENTS

UNITS OF RADIOACTIVITY

Ci ... Curie ... 3.7×10^{10} disintegrations per second mCi ... millicurie ... 10^{-3} Curies pci ... picocurie ... 10^{-12} Curies ... 2.22 disintegrations per minute

UNITS OF LENGTH, AREA, VOLUME AND MASS AND THEIR EQUIVALENTS IN THE IMPERIAL SYSTEM

cm ₂	centimetre 0.394 inches
km ²	square kilometre 0.386 square miles cubic metre 35.31 cubic feet
	litre 0.880 quart
g	gram 0.0353 ounce

NOTES

- 1. Unless otherwise noted, all times given in this report are New Zealand Standard time i.e. G.M.T. + 12 hours.
- 2. Radioactive fallout in rain is expressed as:
 - (a) Deposition millicuries per square kilometre (mCi/km²)
 - (b) Concentration picocuries per litre

(pCi/litre)

Concentration (pCi/litre) = $\frac{\text{deposition (mCi/km}^2)}{\text{rainfall (cm)}} \times 100$

Multiply mCi/km² by 2.59 to obtain mCi/sq. mile.

3. The levels of strontium-90 contamination in food and bone are given in "Strontium Units" i.e. picocuries strontium-90 per gram of calciumpCi Sr⁹⁰/g Ca.

Similarly caesium-137 results are given as picocuries of caesium-137 per gram of potassium.....pCi Cs^{137}/g K.

One litre of whole milk contains approximately:

1.2 g of calcium

1.4 g of potassium.

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POTENTIAL HEALTH HAZARD

The significance of the levels of radioactivity in environmental samples published in this Report may be understood more readily by comparing these levels with the following "permissible levels for the general population" which have been adopted for use in New Zealand.

These levels have been set as a guide to limit the controlled release of radioactive substances into the environment by licensed users in New Zealand.

They are levels which individually would not require remedial or preventive action and have been chosen to protect the most sensitive age group in the population. It is considered that any risk associated with these levels is exceedingly small and that levels many times as great would involve a hazard which is small compared to commonly accepted risks of life.

"Permissible levels" of Radioactivity

These levels were derived so as to ensure that the dose to any member of the public arising from the controlled use of radioactive materials does not exceed the Dose Limit recommended by the International Commission on Radiological Protection.

Strontium-90

In Milk: 270 Strontium Units - maintained indefinitely in the milk. In Bone: 67 Strontium Units.

Caesium-137

In Milk: 7,000 pCi/g K - maintained indefinitely in the milk.

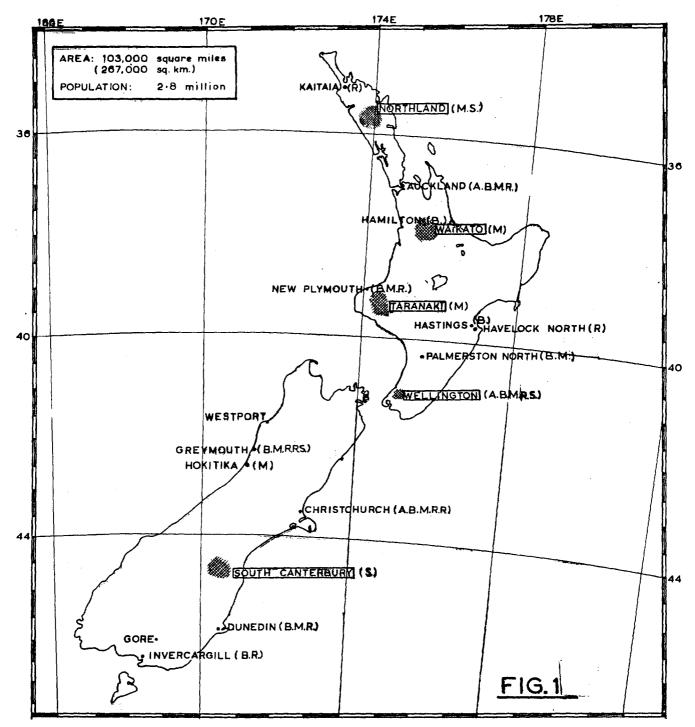
Iodine-131

In Milk: 200 pCi/litre - as an average intake over one year.

Total Beta Activity of Mixed Fission Products Between 10 and 80 Days Old

In Air: 300 pCi/m³ - for continuous breathing.

In Rainwater: 6,000 pCi/litre - for continuous consumption.



LOCATION OF COLLECTING STATIONS ESTABLISHED BY THE NATIONAL RADIATION LABORATORY FOR AIR(A), BONE(B), MILK(M), RAINWATER(R), AND SOIL (S) SAMPLES IN NEW ZEALAND one type of collection is performed (e.g. weekly and monthly rainwater collection) the appropriate symbol is shown twice. Collection areas not confined to a single location but extending over part of a province or district are shown thus WINAME

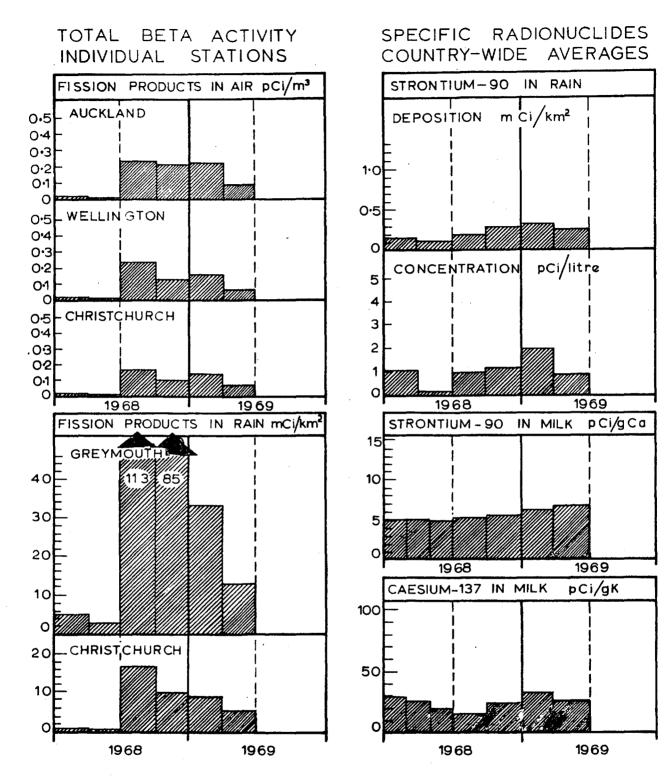


FIG. 2 SUMMARY OF ROUTINE MEASUREMENTS

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SUMMARY

GENERAL

Results of routine monitoring of fallout in air, rain and milk samples during the aecond quarter 1969 are tabulated in this Report. These results are summarized on a quarterly basis and are presented graphically in Fig. 2, together with results from previous Quarterly Reports during 1968 and 1969.

Additional information on sample collection and evaluation, comparison of levels of environmental radioactivity and health hazard assessment is given in the Annual Summary Report for 1968 "ENVIRONMENTAL RADIOACTIVITY IN NEW ZEALAND, Report No. NRL-F33" which also includes the results of extended monitoring of fallout from the French nuclear tests in the Pacific.

TOTAL BETA ACTIVITY - INDIVIDUAL STATIONS

During the first quarter 1969, and also during the two previous quarters, levels of fission products in air and rainwater increased. These higher levels resulted from the 1968 series of nuclear tests conducted by France in the Pacific area between 8 July and 9 September. This series included for the first time the detonation of two hydrogen bombs.

The expected decrease in levels, which had been evident during the months following the previous test series of 1966 and 1967 occurred more slowly following the 1968 nuclear tests. The total beta activity of air samples during the first quarter 1969 had not decreased significantly from the levels of the previous two quarterly periods. There has been, however, a significant decrease during the second quarter 1969. The total beta activity of rain samples has steadily decreased since the third quarter 1968. The greater delay in the reduction of fission product levels following the 1968 nuclear tests undoubtedly results from the injection of fission debris into higher altitudes when hydrogen bombs are detonated. The subsequent deposition of the debris is thus extended over a longer period.

The average levels in air during the second quarter 1969 were 0.09 pCi/m^3 at Auckland and 0.06 pCi/m^3 at Wellington and Christchurch. During the corresponding periods in 1967 and 1968 the highest levels were 0.03 pCi/m^3 (at Auckland and Christchurch) and 0.01 $pC1/m^3$ (at Auckland) respectively.

The total deposition in rain during the second quarter 1969 was 12.3 mCi/km² at Greymouth, and 5.4 mCi/km at Christchurch. The levels have steadily decreased since the third quarter 1968. During the second quarters of 1967 and 1968 the highest deposition was also at Greymouth: 4.7 and 2.3 mCi/km² respectively.

SPECIFIC RADIONUCLIDES - COUNTRY-WIDE AVERAGES

1. <u>STRONTIUM-90 IN RAIN.</u> The average deposition in rain decreased slightly from 0.34 mCi/km² during the first quarter 1969 to 0.27 mCi/km² during the second quarter 1969. This level is about one fifth of the highest level previously recorded i.e. during the first quarter 1965.

2. STRONTIUM-89 IN RAIN. The average deposition reached a maximum of 5.1 mCi/km² during October 1968, about one month after the conclusion of the 1968 nuclear tests. During₂the second quarter 1969 levels have been steadily falling: 0.5, 0.4 and 0.2 mCi/km² during April, May and June respectively. The previous highest level was 3.5 mCi/km² during November 1966.

<u>3.</u> STRONTIUM-90 IN MILK. The average level has increased slightly from 6.3 Strontium Units during the first quarter 1969 to 6.8 Strontium Units during the second quarter 1969. The highest level previously recorded was 15.9 Strontium Units during July-August 1964. The average level for the twelve months ending June 1969 (6.0 Strontium Units) is about 2% of the "permissible level" for the whole population⁽¹⁾.

4. CAESIUM-137 IN MILK. The average level has decreased slightly from 33 pCi/g K during the first quarter 1969 to 27 pCi/g K during the second quarter 1969. The highest level previously recorded was 81 pCi/g K during March-April 1965. The average level for the twelve months ending June 1969 (25 pCi/g K) is less than 0.4% of the "permissible level" for the whole population (1).

(1) SEE POTENTIAL HEALTH HAZARD - PAGE 3.

RESULTS OF ROUTINE MONITORING OF FALLOUT DURING SECOND QUARTER 1909

TABLE 1

TOTAL BETA ACTIVITY OF AIR SAMPLES

In Picocuries per Cubic Metre Four Days after Collection. Filters changed 3 times each week.

	Filters changed 3 times each week.								
AUC	KLAND	WELL	INGTON	CHRISTCHURCH					
Date	Total Beta	Date	Total Beta	Date	Total Beta				
Filter	Activity	Filter	Activity	Filter	Activity				
Removed	pCi/m ^{3°}	Removed	pCi/m ^{3*}	Removed	pCi/m ^{3°}				
2.4.69	0.11	4.4.69	0.07	2.4.69	0.08				
4.4.69	0.16	7.4.69	0.10	8.4.69	0.09				
7.4.69	0.12	9.4.69	N.S.	14.4.69	0.07				
9.4.69	0.12		0.06						
11.4.69		11.4.69	0.15	16.4.69 21.4.69	0.09				
14.4.69	0.13	14.4.69			0.05				
16.4.69	0.25	16.4.69	0.07	23.4.69	0.03				
18.4.69	0.15	18.4.69	0.12	24.4.69	0.05				
21.4.69	0.16	21.4.69	0.02	28.4.69	0.04				
	0.08	23.4.69	0.08	30.4.69	0.08				
23.4.69	0.07	25.4.69	0.06						
25.4.69	0.10	28.4.69	0.05						
28.4.69 30.4.69	0.08 0.14	30.4.69	0.11						
Average	0.14	Average	0.08	Average	0.06				
2.5.69	0.07	2.5.69	0.09	2.5.69	0.05				
5.5.69	0.07	5.5.69	N.S.	5.5.69	0.05				
7.5.69	0.03	7.5.69	0.07	7.5.69	0.07				
9.5.69	0.11	9.5.69	0.04	9.5.69	0.05				
12.5.69	0.08	12.5.69	0.06	12.5.69	0.03				
14.5.69	0.13	14.5.69	0.11	14.5.69	0.07				
16.5.69	0.09	16.5.69	0.13	16.5.69	0.06				
19.5.69	0.09	19.5.69	0.06	19.5.69	0.05				
21.5.69	0.05	21.5.69	0.10	21.5.69	0.10				
23.5.69	<0.01	23.5.69	0.08	23.5.69	0.04				
26.5.69	0.08	26.5.69	0.03	26.5.69	<0.01				
28.5.69	0.09	28.5.69	0.04	28.5.69	0.02				
30.5.69	0.04	30.5.69	0.05	30.5.69	0.03				
Average	0.07	Average	0.07	Average	0.05				
2.6.69	0.04	2.6.69	0.02	3.6.69	0.04				
4.6.69	0.02	4.6.69	0.04	6.6.69	0.06				
6.6.69	0.09	6.6.69	0.06	9.6.69	0.04				
9.6.69	0.10	9.6.69	0.04	11.6.69	0.03				
11.6.69	0.03	11.6.69	0.03	13.6.69	0.05				
12.6.69	0.18	13.6.69	0.04	16.6.69	0.02				
16.6.69	0.05	16.6.69	0.05	18.6.69	0.04				
18.6.69	0.07	18.6.69	0.06	20.6.69	0.03				
20.6.69	0.06	20.6.69	0.03	23.6.69	0.02				
23.6.69	0.04	23.6.69	0.02	25.6.69	0.05				
25.6.69	0.05	25.6.69	0.03	27.6.69	0.03				
27.6.69	0.08	27.6.69	0.05	30.6.69	0.03				
30.6.69	0.08	30.6.69	0.04		_				
Average	0.07	Average	0.04	Average	0.04				
Quarterly	0.09	Quarterly	0.06	Quarterly	0.06				
Average		Average		Average					
		l		U					

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TABLE 2 TOTAL BETA ACTIVITY OF WEEKLY RAINWATER COLLECTIONS FOUR DAYS AFTER COLLECTION							
АТ	DATE OF C FROM	OLLECTION TO	DEPOSITION mCi/km ²	RAINFALL cm	CONCENTRATION pCi/litre		
	29.3.69 5.4.69 12.4.69 19.4.69	5.4.69 12.4.69 19.4.69 26.4.69	1.5 2.2 1.4 2.0	6.2 5.5 7.8 7.8			
ĺ	29.3.69	26.4.69	7.1	27.3	. 26		
GREYMOUTH	26.4.69 3.5.69 10.5.69 17.5.69 24.5.69	3.5.69 10.5.69 17.5.69 24.5.69 31.5.69	0.2 1.3 0.4 1.6 0.2	1.0 2.8 1.0 8.9 0.9			
GR	26.4.59	31.5.69	. 3.7 .	14.6	25		
	31.5.69 7.6.69 14.6.69 21.6.69	7.6.69 14.6.69 21.6.69 28.6.69	0.2 <0.1 0.9 0.3	0.7 0.3 7.0 1.5	:		
	31.5.69	28.6.69	1.5	9.5	16		
	2nd QUART	ER 1969	12.3	51.4	24		
	28.3.69 3.4.69 11.4.69 18.4.69 24.4.69	3.4.69 11.4.69 18.4.69 24.4.69 2.5.69	0.2 1.5 0.3 1.2 0.3	0.2 0.8 <0.1 3.6 1.2			
	28.3.69	2.5.69	3.5	5.9	59		
CHRISTCHURCH	2.5.69 9.5.69 16.5.69 23.5.69	9.5.69 16.5.69 23.5.69 30.5.69	0.5 <0.1 0.1 0.3	1.7 <0.1 0.7 2.6			
RIS	2.5.69	30.5.69	1.0	5.1	20		
CH	30.5.69 6.6.69 13.6.69 20.6.69	6.6.69 13.6.69 20.6.69 27.6.69	0.1 <0.1 0.5 0.2	1.0 TRACE 0.8 1.5			
	30.5.69	27.6.69	0.9	3.3	27		
	2nd QUARTH	ER 1969	5.4	14.3	38		

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TABLE 3a STRONTIUM-90 IN RAIN SECOND QUARTER 1969									
COLLECT ING STATIONS]	DEPOSITIO mCi/km ²	N		RAINFALL cm		CONCENTRATION pCi/litre		
New Zealand	Apr.	May	Jun.	Apr.	May	Jun.	Apr.	May	Jun.
Kaitaia Auckland New Plymouth Havelock North Wellington Greymouth Christchurch Dunedin Invercargill	0.09 0.12 0.09 0.06 0.12 0.23 0.08 0.07 0.10	0.14 0.07 0.05 0.03 0.12 0.16 0.06 0.06 0.07	0.18 0.12 0.08 0.03 0.05 0.13 0.02 0.02 0.02	12.1 12.3 11.1 7.0 7.2 24.3 5.8 5.3 11.2	22.0 11.7 13.4 5.1 10.9 13.6 5.1 7.5 7.9	25.2 8.7 14.6 5.2 7.5 14.1 2.5 2.9 11.6	0.8 1.0 0.8 0.8 1.7 0.9 1.3 1.2 0.9	0.5 0.6 0.6 0.7 1.2 1.2 0.9 0.9	0.7 1.4 0.5 0.6 0.7 0.9 1.0 0.8 0.7
Country-Wide Averages Monthly Quarterly	0.11	0.08 0.27*	0.08	10.7	11.5	10.3	1.0	0.8	0.8
Pacific Islands					· · · · · · · · · · · · ·	<u></u>			
Suva, Fiji Rarotonga	0.10 0.07	N.S. 0.04	0.04 0.03	25.0 35.8	7.3 9.1	2.4 7.3	0.4 0.2	N.S. 0.5	1.7 0.4

* This value is the sum of the monthly depositions during the quarter.

N.S. No Sample.

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TABLE 3b	STRONTIUM-89 IN RAIN SECOND QUARTER 1969							
COLLECTING STATIONS		DEPOSITION RATIO mCi/km ² (at mid-month) Strontium-89/Strontium				ntium-90		
New Zealand	Apr.	May	Jun.	Apr.	May	Jun.		
Kaitaia Auckland New Plymouth Havelock North Wellington Greymouth Christchurch Dunedin Invercargill	0.4 0.5 0.4 0.2 0.4 1.1 0.4 0.4 0.5	03 04 07 02 02	0.1 0.3 0.1	4 4 3 5 6 5	5 3 6 8 3 5 3 3 3 3	2 4 3 2 2 4 3 4		
<u>Country-Wide Averages</u> Monthly Quarterly	0.5	04 1.1*	0.2	.5	4 .4	3		
Pacific Islands						·		
Suva, Fiji Rarotonga	0.•4 0.•4	N.S.	<01 <01	4 5	N. S. 4	2 3		

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* This value is the sum of the monthly depositions during the quarter.

N.S. No Sample.

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TABLE 4 STRONTIUM-90 AND CAESJUM-137 IN MILK 1969								
COLLECTING STATIONS	STRONTIUM-90 pCi/g Ca	CAESIUM-137 pCi/g K						
Northland Auckland Waikato Taranaki Palmerston North Wellington Westland * Christchurch Dunedin	April - June 5.1 4.9 5.9** 9.4 10.1 4.7 16.4 2.0 2.9	April 17 24 50 102 5 11 46 2 7	23 52					
Country-Wide Averages Monthly Quarterly	6 . 8	29	28 27	24				

- * The Westland Collecting Station was referred to as Greymouth or Hokitika in previous reports.
- ** April May only
- *** No Sample. Results in brackets are estimates used for calculating the country-wide average.

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REPORT No. NRL-F 36

DEPARTMENT OF HEALTH



QUARTERLY REPORT JULY-SEPTEMBER 1969

ENVIRONMENTAL RADIOACTIVITY IN NEW ZEALAND

AND

MEASUREMENTS ON SAMPLES FROM FIJI AND RAROTONGA

NATIONAL RADIATION LABORATORY P.O. BOX 1456, CHRISTCHURCH, NEW ZEALAND

SYMBOLS UNITS AND EQUIVALENTS

UNITS OF RADIOACTIVITY

Ci Curie Curie 10^{-3} Curies Ci	
pCi picocurie 10 ⁻¹² Curies 2.22 disintegrations per minute	
UNITS OF LENGTH, AREA, VOLUME AND MASS AND THEIR EQUIVALENTS IN THE IMPERIAL SYSTEM	i
cm ₂ centimetre 0.394 inches	
km ³ square kilometre 0.386 square miles	
m ³ cubic metre 35.31 cubic feet	
litre litre 0.880 quart	
g gram 0.0353 ounce	
NOTES	•
 Unless otherwise noted, all times given in this report are New Zealand Standard time i.e. G.M.T. + 12 hours. 	
2. Radioactive fallout in rain is expressed as:	
(a) Deposition - millicuries per square kilometre (mCi/km^2)	
(b) Concentration - picocuries per litre (pCi/litre)	
Concentration (pCi/litre) = $\frac{\text{deposition (mCi/km}^2)}{\text{rainfall (cm)}} \times 100$	
Multiply mCi/km ² by 2.59 to obtain mCi/sq. mile.	
3. The levels of strontium-90 contamination in food and bone are	
given in "Strontium Units" i.e. picocuries strontium-90 per gram	
of calciumpCi Sr ⁹⁰ /g Ca.	
Similarly caesium-137 results are given as picocuries of caesium-137 per gram of potassiumpCi Cs^{137}/g K.	
One litre of whole milk contains approximately:	
1.2 g of calcium	
1.4 g of potassium.	

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RESULTS OF ROUTINE MONITORING OF FALLOUT DURING FIRST QUARTER 1968

Total Beta A	stivity of	f Air Samp	les, Ta	ble 1.	••	• •	**	• •	7
Total Beta A	ctivity of	f Weekly R	ainwate	r Coll	ectio	ns,	Tabl	e 2.	8
Strontium-90	in Rain,	Table 3a.		•• ••	••	••	••	••	9
Strontium-89	in Rain,	Table 3b.	• •	•• ••	••	••	••	••	10
Strontium-90	and Caest	ium-137 in	Milk,	Table	4.		••	••	11

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POTENTIAL HEALTH HAZARD

The significance of the levels of radioactivity in environmental samples published in this Report may be understood more readily by comparing these levels with the following "permissible levels for the general population" which have been adopted for use in New Zealand.

These levels have been set as a guide to limit the controlled release of radioactive substances into the environment by licensed users in New Zealand.

They are levels which individually would not require remedial or preventive action and have been chosen to protect the most sensitive age group in the population. It is considered that any risk associated with these levels is exceedingly small and that levels many times as great would involve a hazard which is small compared to commonly accepted risks of life.

"Permissible levels" of Radioactivity

These levels were derived so as to ensure that the dose to any member of the public arising from the controlled use of radioactive materials does not exceed the Dose Limit recommended by the International Commission on Radiological Protection.

Strontium-90

In Milk:	270 Strontium Units - maintained indefinitely in the milk	•
In Bone:	67 Strontium Units.	

Caesium-137

In Milk: 7,000 pCi/g K - maintained indefinitely in the milk.

Iodine-131

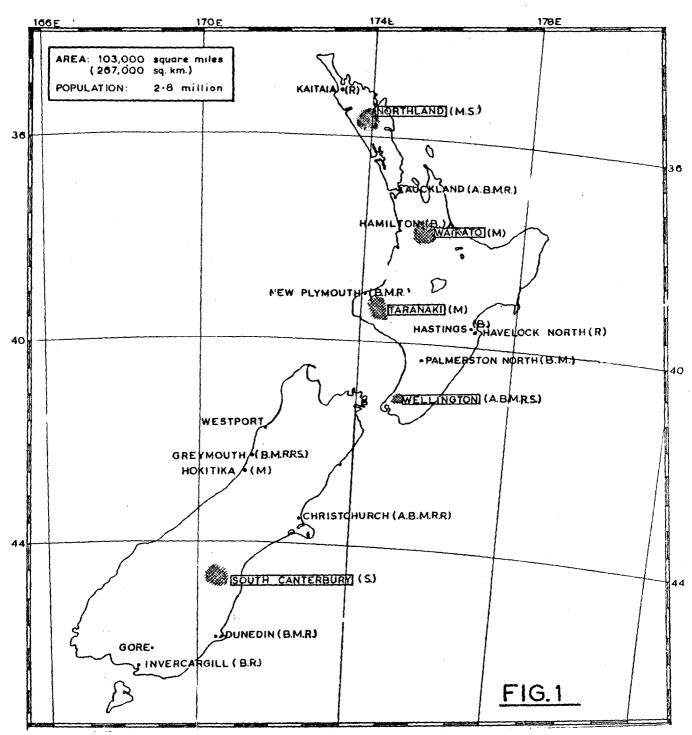
In Milk: 200 pCi/litre - as an average intake over one year.

Total Beta Activity of Mixed Fission Products Between 10 and 80 Days Old

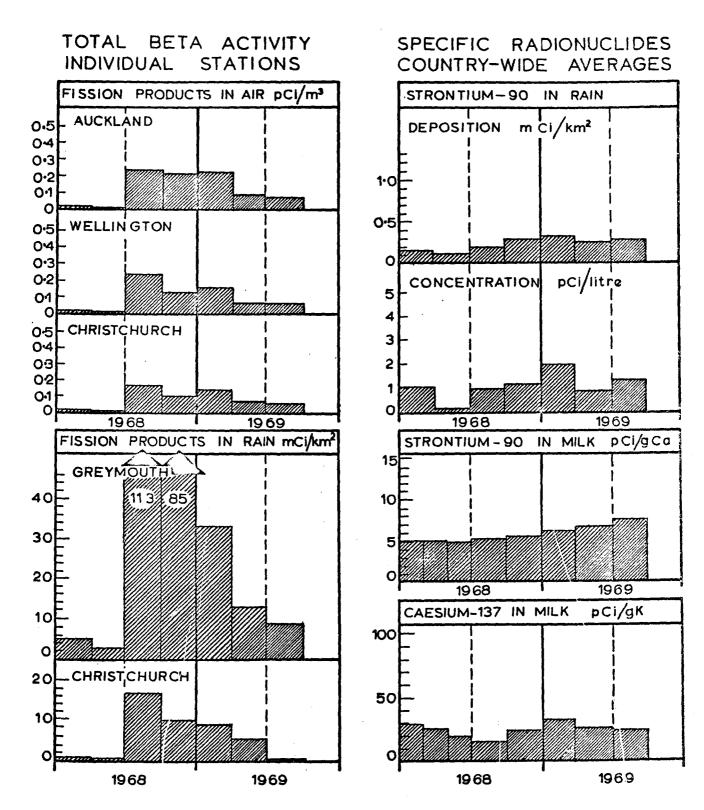
In Air: 300 pCi/m^3 - for continuous breathing.

In Rainwater: 6,000 pCi/litre - for continuous consumption,

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LOCATION OF COLLECTING STATIONS ESTABLISHED BY THE NATIONAL RADIATION LABORATORY FOR AIR(A), BONE(B), MILK(M), RAINWATER(R), AND SOIL (S), SAMPLES IN NEW ZEALAND one type of collection is performed (e.g. weekly and monthly rainwater collection) the appropriate symbol is shown twice. Collection areas not confined to a single location but extending over part of a province or district are shown thus (NAME)







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ROUTINE MEASUREMENTS

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SUMMARY

CENERAL

Results of routine monitoring of fallout in air, rain and milk samples during the third quarter 1969 are tabulated in this heport. These results are summarized on a quarterly basis and are presented graphically in Fig. 2, together with results from previous Quarterly Reports during 1968 and 1969.

Additional information on sumple collection and evaluation, comparison of levels of environmental radioactivity and health hazard assessment is given in the Annual Summary Report for 1968 "ENVIRONMENTAL RADIOACTIVITY IN NEW ZEALAND, Report No.NRL-F33" which also includes the results of extended monitoring of fallout from the French nuclear tests in the Pacific.

TOTAL BETA ACTIVITY - INDIVIDUAL STATIONS

As already noted in the reports NRL-F31 to NRL-P35, increased levels of fission products in air and rainwater were measured during the third quarter 1968. These levels resulted from the 1968 series of nuclear tests conducted by France in the Pacific Area between 8 July and 9 September. This series included for the first time the detonation of two hydrogen bombs.

The expected decrease in levels, which had been evident during the months following the previous test series of 1966 and 1967, occurred more slowly following the 1968 nuclear tests. The greater delay in the reduction of fission product levels following the 1968 nuclear tests undoubtedly resulted from the injection of fission debris into higher altitudes when the hydrogen bombs were tested. The subsequent deposition of the debris was thus extended over a longer period.

During the second and third quarters 1969, however, there were significant decreases in the levels of total beta activity of air and rainwater.

The average levels in air during the third quarter 1969 were 0.08 pCi/m³ at Auckland, 0.06 pCi/m³ at Vellington and 0.05 pCi/m³ at Christchurch. During the corresponding period in 1968 the levels were 0.23 pCi/m³ at Auckland and Wellington, and 0.16 pCi/m³ at Christchurch.

The total deposition in rain during the third quarter 1969 was 9.1 mCi/km² at Greymouth, and 0.4 mCi/km² at Christchurch. During the corresponding period in 1968 the deposition was 112.8 mCi/km² at Greymouth, and 16.4 mCi/km² at Christchurch.

SFECIFIC RADIONUCLIDES - COUNTRY-WIDE AVERAGES

1. STRONTIUM-90 IN RAIN The average deposition in rain increased slightly from 0.27 mCi/km² during the second quarter 1969 to 0.30 mCi/km² during the third quarter 1969. This level is less than one quarter of the highest level previously recorded, i.e., during the first quarter 1965.

2. STRONTIUM-89 IN RAIN The average deposition reached a maximum of 5.1 mC1/km² during October 1968, about one month after the conclusion of the 1968 nuclear tests. During the third quarter 1969 levels were about 0.1 mCi/km² per month.

3. STONTIUM-90 IN MILK The average level has increased from 6.8 Strontium Units during the second quarter 1969 to 7.9 Strontium Units during the third quarter 1969. The highest level previously recorded was 15.9 Strontium Units during July-August 1964. The average level for the twelve months ending September 1969 (6.6 Strontium Units) is about 2.5% of the "permissible level" for the whole population⁽¹⁾.

4. CAESIUM-137 IN MILK The average level has decreased slightly from 27 pCi/g K during the second quarter 1969 to 24 pCi/g K during the third quarter 1969. The highest level for the twelve nonths ending September 1969 (27 pCi/g K) is less than 0.4% of the "permissible level" for the whole population⁽¹⁾.

(1) See "POTENTIAL HEALTH HAZARD" - Page 3.

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TABLE 1	т	OTAL BETA ACTIVI	TY OF AIR SAMPL	ES	
	In Picocuries Fi	per Cubic Metre lters changed 3	Four Days afte times each week	r Collection.	
AUCK	LAND	WELLI	WELLINGTON CHRISTCHU		CHURCH
Date Filter Removed	Total Beta Activity pCi/m ³	Date Filter Removed	Total Beta Activity pCi/m3	Date Filter Removed	Total Beta Activity pCi/m ³
2.7.69 4.7.69	0.10 0.08	2.7.69 4.7.69	0.05 0.04	2.7.69 4.7.69	0.04 0.06
7.7.69 9.7.69 11.7.69	0.09 0.07	8.7.69 9.7.69	0.09 0.08	7.7.69 9.7.69	0.05 0.05
14.7.69	0.03 0.04 0.06	11.7.69 14.7.69 16.7.69	0.03 0.04 0.04	11.7.69 15.7.69 18.7.69	0.04 0.04 0.04
18.7.6 <u>9</u> 21.7.69	0.04 0.04	18.7.69 21.7.69	0.04 0.03	21.7.69 23.7.69	0.04 0.05 0.07
23.7.69 25.7.69 28.7.69	0.08 0.07 0.04	23.7.69 25.7.69 28.7.69	0.07 0.09 0.02	25.7.69 28.7.69 30.7.69	0.08 0.03 0.04
30.7.69 1.8.69	0.05 0.09	30.7.69 1.8.69	0.03 0.03	1.8.59	0.03
Average 4.8.69	0.06	Average 4.8.69	0.05	Average	0.05
6.8.69 8.8.69	0.13 0.04	6.8.69 8.8.69	0.05 0.05 0.17	4.8.69 6.8.69 8.8.69	0.03 0.05 0.04
11.8.69 13.8.69 15.8.69	0.07 0.05 0.08	11.8.69 13.8.69 15.8.69	0.05	11.8.59 14.8.59	0.05
18.8.69 20.8.69	0.13 0.10	18.8.69 20.8.69	0.08 0.02 0.07	15.8.69 18.8.69 20.8.69	0.03 0.04 0.05
22.8.69 25.8.69 27.8.69	0.07 0.03 0.08	22.8.69 25.8.69 27.8.69	0.07 0.05 0.09	25.8.69 27.8.69 29.8.69	0.09 0.12 0.04
29.8.69 1.9.69	0.10 0.08	29.8.69 1.9.69	0.05	1.9.69	0.04
Average 3.9.69	0.08	Average	0.07	Average	0.05
3.9.69 5.9.69 8.9.69 10.9.69	0.09 0.09 0.11 0.03	3.9.69 5.9.69 8.9.69 10.9.69	0.03 0.12 0.09 0.05	3.9.69 5.9.69 8.9.69 10.9.69	0.05 0.08 0.04 0.02
12.9.69 15.9.69 17.9.69	0.09 0.10 0.12	12.9.69 15.9.69 17.9.69	0.07 0.08 0.09	12.9.69 15.9.69 17.9.69	0.07 0.05 0.05
19.9.69 22.9.69	0.08 0.07	19.9.69 22.9.69	0.06 0.05	19.9.69 22.9.69	0.04 0.04
24.9.69 26.9.69 29.9.69	0.07 0.12 0.08	24.9.69 26.9.69 29.9.69	0.04 0.06 0.07	24.9.69 29.9.69 1.10.69	().06 0.08 0.08
1.10.69 Average	0.10	1.10.69 Average	0.08	Average	0.0.5
Quarterly Average	0.09	Quarterly Average	0.06	Quarterly Average	0.05

RESULTS OF ROUTINE MONITORING OF FALLOUT DURING THIRD QUARTER 1969

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TABLE 36 STRO	NTIUM-89	IN RAIN T	HIRD QUARTER	1969				
COLLECTING STATIONS	13	DEPOSITION mCi/km ² (at mid-month)			RATIO Strontium-89/Strontium-90			
New Zealand	July	Aug.	Sept.	July	Aug.	Sept.		
Kaitaia	0.2	<0.1	0.2	3 2	1	1		
Auckland		0.3	(0.2)**	2	2	(1)**		
New Plymouth		.0.3		2	2	(1)**		
Havelock North	<0.1		<0.1	3 2 2	2	<1		
Wellington	<u>ii</u>	0.1	0.1	2	1	2		
Greymouth		0.2	0.5	2	2	1		
Christchurch		0.2		2	1	<1		
Dunedin		<0.1		1 2	2 1	≤ 1		
Invercargil1	<0.1	<0.1	0.2	2	L	1		
Country-Wide Averages								
Monthly	0.1	0.1	0.2	2	2	1		
Quarterly		0.4*			2			
Pacific Islands	1							
Suva, Fiji	0.6	<0.1	***	4	<1	***		
• -		•	***		-	***		
Rarotonga	0.1	<0.1	***	3	<1	~ ~ ~		

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* This value is the sum of the monthly depositions during the quarter.

** Results in brackets are estimates used for calculating the country-wide average.

*** Sample transport delay. Results to be published in the fourth quarterly report.

TABLE 4 STRONTIUM-90 A	ND CAESIUM-137 IN MIL	K THIRD QUA	ARTER 1969				
COLLECTING STATIONS	STRONTIUM-90 pCi/g Ca		CAESIUM-137 pCi/g K				
	July - Sept.	July	Aug.	Sept.			
Northland	9.3	15	25	50			
Auckland	7.2	39	-25	27			
Waikato	6.9	27	29	35			
Taranaki	11.7	45	92	113			
Palmerston North	5.6		2	3			
Wellington	5.5	. 5 8	2	3 7			
Westland *	20.7	16	18	44			
Christchurch	1.8	2	くユ	<1			
Dunedin	2.7	3	(1)**	<1			
Country-Wide Averages			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,				
Monthly		18	22	31			
Quarterly	7.9		24				

- * The Westland Collecting Station was referred to as Greymouth or Hokitika in previous reports.
- ** No sample. Result in brackets is an estimate used for calculating the country-wide average.

2. Department of Scientific and Industrial Research The Institute of Nuclear Sciences Lower Hutt, New Zealand

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RADIOISOTOPES IN RAINWATER:

Results for

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January-April 1968	report No. 72	
May - August 1968	report No. 73	
September-December 1968	report No. 74	

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RADIOISOTOPES IN RAINWATER

Report No 72

Period: January - April 1968

Station: Gracefield, Lower Hutt, New Zealand, S 41° 14' E 174° 55'

Stainless steel collector 1 ft diameter

Sampling Period start finish	Activity: Sr ⁹⁰	microcuri Ba ¹⁴⁰	es/sq.mile Cs ¹³⁷	$\frac{\mathrm{Ba}^{140}}{\mathrm{Sr}^{90}}$	<u>Cs</u> 137 Sr90	Rain Inches	Remarks
1-2 2-1	248 <u>+</u> 20 (96)*		200 <u>+</u> 12 (77)*		0.81	4.69	monthly pot sample
Accumulated total or average for 1968	248 ± 20 (96)*		200 ± 12 (77)*		0.81	4.69	
2-1 3-1							recovery so low, no results recorded
Accumulated total or average for 1968	248 <u>+</u> 20 (96)*		200 ± 12 (77)*	-	0.81	7.33	
3-1 4-1	41.5±3 (16)*		22.3 ± 1.6 (8.6)*		0.54	0.66	monthly pot sample
accumulated total average For 1968	289 ± 20 (112)*		223 ± 12 (85.6) *		0.77	7.99	
4-1 5-1	234 <u>+</u> 16 (90)*		163 <u>+</u> 27 (63)*		0.70	14.36	monthly pot sample
accumulated total or average for 1968	523 ± 25 (202)*		386 <u>+</u> 30 (148.6)*	••	0.74	22.35	

* Microcuries per square kilometer

Note: Errors quoted are purely counting errors and do not include experimental errors or calibration uncertainties.

INSTITUTE OF NUCLEAR SCIENCES

Report 113/- WJM

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RADIOISOTOPES IN RAINWATER

Report No 73, Period: May 1968

Station - Gracefield, Lower Hutt, New Zealand S 41°14' E 174°55'

Stainless steel collector 1 fc diameter.

Sampling period		Activity: Microcuries/sq. mile			1/10, 00 137.	137, 90		
start	finish	Sr ⁹⁰	Ba ¹⁴⁰	Cs ¹³⁷	Ba ¹⁴⁰ /Sr ⁹⁰	Cs ¹³⁷ /Sr ⁹⁰	Rain, inches	Remarks
5 - 1	6 - 1	149 ± 11.0 (58.0)*		175 <u>+</u> 8 (58)*		· 1.17	8.89	Monthly pot sample
	ated cotal age for 968	672 <u>+</u> 27 (259)*		561 ± 31 (217)*		د0.8	31.24	
6 - 1	7 - 1	123 ± 8 (48)*		151 ± 2 (58)*		1.23	7.72	Monthly pot sample
	aced total age for 968	795 ± 28 (307)*		712 <u>+</u> 31 (275)*		0.90	38.96	
7 - 1	8 - 1	112 ± 7.3 (43)*	45,400 <u>±</u> 380 (17,900)*	155 ± 24 (55)*	404.0	1.39	<i>6.9</i> 0	Monthly pot sample
	ated total age for 968	907 ± 29 (350)*	45,400 <u>+</u> 380 (17,900)	987 ± 31 (330)*	404.0	1.09	45.86	
8 - 1	9 - 2	93.5 ± 3.2 (35)*	8,360 <u>+</u> 124 (3240)*	116 ± 1.35 (45)*	89.0	1.23	2.59	Monthly pot sample
	ated total age for 968	1000.5± 30 (385)*	53,760 ± 392 (21,140)*	110 <u>5</u> + 31.5 (37 5) *	246	1.10	48.45	

* microcuries per square kilometer

Note: Errors quoted are purely counting errors and do not include experimental errors or calibration uncertainities.

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RADIOISOTOPES IN RAINWATER.

Report No. 74 - September 1968 to December 1968

Station - Gracefield, Lower Hutt S41014' E174055'

Stainless Steel Collector lft. diameter

Samplin Start	g Period Finish	Acti Sr ⁹⁰	vity microcuri Ba ¹⁴⁰	e/Sq. mile Cs ¹³⁷	Ba ¹⁴⁰ /Sr ⁹⁰	Cs ¹³⁷ /9r ⁹⁰	Rain Inches	Remarks
9 - 2	10 - 1	153 <u>+</u> 6.5 59*	4530 <u>+</u> 67.1 1750*	186 <u>+</u> 1.9 71.5*	29_6	1.22	4,43	Monthly Pot Sample
	ted Total ge for 1968	1153,5 <u>+</u> 31 445*	58290 <u>+</u> 400 22890*	1289 <u>+</u> 32.0 446.5*	50,5	1.11	52,88	
10 - 1	11 - 1	262 <u>+</u> 6.4 105*	6360 <u>+</u> 81 2430*	555 <u>+</u> 3.8 214*	24	2.12	8.40	Monthly Pot Sample Sr ⁹⁰ probably low
	ted total ge for 1968	1415.5 <u>+</u> 31.5 550*	64,590 <u>+</u> 408 25,320*	1744 <u>+</u> 32.0 660.5*	46	1,21	61 .29	
11 - 1	12 - 1	190 <u>+</u> 6.4 73*	543 <u>+</u> 32.5 214*	214 <u>+</u> 6.7 82.5*	2.93	1,12	2.07	Monthly Pot Sample
	ted total ge for 1968	1605.5 <u>+</u> 32.0 623*	65,133 <u>+</u> 420 25,534*	1958 <u>+</u> 32.7 743*	40.7	1.20	63,35	
12 - 1	2 - 1 - 69	178 <u>+</u> 4.0 69*	-	296 <u>+</u> 3.2 114*		1.69	4.60	Northly Pot Sample Sr ⁹⁰ probebly low
	ted total ge for 1968	1783.5 <u>+</u> 32.0 692*	65,133 <u>+</u> 420 25,534*	2254 <u>+</u> 33 857*	36,6	1.24	67.95	

* Microcuries per square kilometer

Note errors quoted are purely counting errors and do not include experimental errors or calibrations uncertainties.

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Cesium-137 in Various Chicago Foods (Collection Month January, 1970)

S. S. Brar and D. M. Nelson Radiological Physics Division Argonne National Laboratory Argonne, Illinois 60439

Since April, 1961, the ¹³⁷Cs and potassium content of the Chicago portion of Tri-City Diet Sampling Program has been determined¹⁻⁴ in bulk food samples by gamma ray spectrometry using a 4" x 4" NaI (T1) crystal. The individual food components were counted for a minimum of 100 minutes, and from these measurements composite daily and yearly food intakes were obtained. Cesium-137 activity in food, now, is an order of magnitude lower than it was a few years ago; consequently, a new procedure for these measurements has been adopted in order to improve accuracy. The same variety of foods (all fresh vegetables; all fresh fruits, etc.) are composited before measurement, and the samples are counted a minimum of 400 minutes. The results of the January, 1970, quarter are tabulated in Tables I and II.

*Work performed under the auspices of the U. S. Atomic Energy Commission.

Table I

Cesium-137 in Chicago Diets (Adults)

January, 1970

		Potassium	137 _{Cs} pCi/kg	Potassium g/yr	137 _{Cs} pCi/yr
	kg/yr	g/kg	pur/kg	g/yr	po1/91
		1.0	18	37	666
White Bread	37	1.0	39	29	429
Whole Wheat Bread	11	2.6			429
Eggs	16	1.3	0.	21	0
Fresh Vegetables	43	3.5	· 0	151	-
Root Vegetables	17	2.9	0	49	0
Milk	221	1. 5	12	332	2652
Poultry 🔸	17	2.4	11	41	187
Fresh Fish	8	3.7	32	30	256
Flour	43	1.1	21	47	903
Macaroni	3	2.2	28	7	84
Meat	73	. 3.3	28	241	2044
Dried Beans	3	12.5	9	38	27
Fresh Fruit	68	2.3	0	156	0
Potatoes	45	4.2	8	189	360
Canned Fruits	26	1.0	14	2 6	364
Canned Fruit Juices	19	1.7	18	32	342
Canned Vegetables	20	.8	0	16	0
· · ·					
Tot al/yr				1442	8314
Total/day				4.0	23

* It is assumed in arriving at the average that nine times more ocean fish is consumed than fresh water fish.

Table II

Cesium-137 in Chicago Diets

(Infants)

January, 1970

	kg/yr	Pot assi um g/kg	137 _{Cs} pCi/kg	Potassium g/yr	137 _{Cs} pCi/yr
Evaporated Milk	137	3.2	34	438	4658
Formula Milk	. 37	1.7	30	63	1110
Cereals	8	6.9	26	55	208
Fruits	23	1.3	8	30	184
Meats	17	2,2	28	37	476
Vegetables	23	2.2	0	51	0
Total/yr				674	6636
Total/day				1.8	18

References

- S. S. Brar, et al., USAEC Report No. HASL-146, Cs-137 in Various Chicago Diets, pp. 225-232, July 1, 1964.
- 2. J. Rivera and J. J. Kelly, USAEC Report No. HASL-144, Cs-137 in Tri-City Diets, pp. 228, April 1, 1964
- 3. J. Rivera and J. H. Harley, USAEC Report No. HASL-147, Contributions to the Study of Fallout in Food Chains, pp. 31-35, July, 1964.
- S. S. Brar and D. M. Nelson, USAEC Report No. HASL-217, Cs-137 in Various Chicago Foods, pp. III-20 to III-23, January 1, 1970.

5.

EURATOM JOINT NUCLEAR RESEARCH CENTRE

ISPRA ESTABLISHMENT

Protection Service

Site Survey and Meteorology Section

QUARTERLY REPORT

The Euratom Ispra Establishment is located in Northern Italy 58 Km NW away from Milan and 14 Km W from Varese.

The activity levels shown in this report represent weaponstest fallout, and do not reflect any contamination from the site.

SAMPLE COLLECTION

a. Air

Air is drawn by pumps through paper filters at the rate of, at least, 500 m^3 / day, measured by gas meter.

The single daily filters are measured for gross beta radioactivity and then pooled to give monthly samples, for gamma spectrometry and radiochemical analyses.

b. Wet and dry deposition

These samples are collected monthly by means of four 1 m^2 stainless steel funnels, having the bottom always covered with deionized water. The collected water is evaporated and the dry residue analysed.

c. Milk

Milk is collected twice a week in four small local dairies to give 8 liters / month. About six liters dry matter are submitted to gamma spectrometry and two liters ashed for radiochemical determination of strontium-90.

CHEMICAL PROCEDURES AND COUNTING TECHNIQUES

- a. Strontium-90 is separated by the fuming nitric acid precipitation and then purified through hydroxides and chromates precipitations. The activity of the final strontium carbonate and yttrium oxalate precipitates is measured in low level anticoincidence beta counters.
- b. Cesium-137 is measured by direct gamma spectrometry on the unprocessed or dried samples and, whenever it is necessary, by gamma spectrometry after chemical separation. This is performed by filtration of the solution, obtained dissolving the sample, through a thin AMP (ammonium molybdophosphate) layer, by which cesium is retained. Details of this procedure may be found in the paper by E. Van der Stricht issued on "Radiochemical Acta" <u>3</u>, 193-199 (1964).
- c. Gamma emitting nuclides are measured by direct gamma spectrometry, using, also the spectrum stripping technique.
- d. Plutonium-239+240 is separated by anion exchange and electrodeposition; details of the procedure may be found in the paper by M.C. de Bortoli: "Radiochemical determination of plutonium in soil and other environmental samples", Anal. Chem. <u>39</u>, 375 (March 1967).

The activity is measured in a Frish grid ionisation chamber connected to a multichannel analyser.

EXTRAPOLATION OF THE DATA

Except when otherwise stated, the data presented in this report are extrapolated to the last day of the collecting period.

FALLOUT DEPOSITION

1969

SITE : ISPRA

III

ι ω 5 LAT. 45° 49' N

LONG, 8° 37' E

ALT. 250 m

Sale and the

" Gross bet	ta (1)	90 _{Sr}	89 _{Sr}	137 _{Cs}	239 _{Pu}	238 _{Pu}	Precipitation
mCi/Km ²	1	mCi/Km ²	mCi/Km ²	mCi/Km ²	Ci/Km ²	µCi/Km ²	mm
0.78	11-11	17 11 11 11 11 第二 11	t t t t ¥ t	E E E 30 E	t t t T		5.0
3.80	11-12	0.073	0.20	0.098	t t t ¥ t	t t t X t	123.4
0.95	13-1-70	0,012	0.008	0.013	0.22	0.06	22.4
	mCi/Km ² 0.78 3.80	0.78 11-11 3.80 11-12	mCi/Km ² Date(2) mCi/Km ² 0.78 11-11 x 3.80 11-12 0.073	mCi/Km ² Date (2) mCi/Km ² mCi/Km ² 0.78 11-11 # # 3.80 11-12 0.073 0.20	mCi/Km² Date(2) mCi/Km² mCi/Km² mCi/Km² 0.78 11-11 # # # 3.80 11-12 0.073 0.20 0.098	mCi/Km ² Date (2) mCi/Km ² mCi/Km ² mCi/Km ² mCi/Km ² mCi/Km ² 0.78 11-11 # # # # # # 3.80 11-12 0.073 0.20 0.098 #	mCi/Km^2 $Date(2)$ mCi/Km^2 mCi/Km^2 mCi/Km^2 $\mu Ci/Km^2$ $\mu Ci/Km^2$ 0.78 11-11 x x x x x 3.80 11-12 0.073 0.20 0.098 x x

(1) Potassium-40 equivalent (40 mg/cm²).

(2) Day and month of the gross beta measurement.

x Data not yet available.

AIR RADIOACTIVITY

1969

SITE: ISPRA

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A LAT. 45° 49' N

LONG. 8° 37' E

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ALT. 250 m

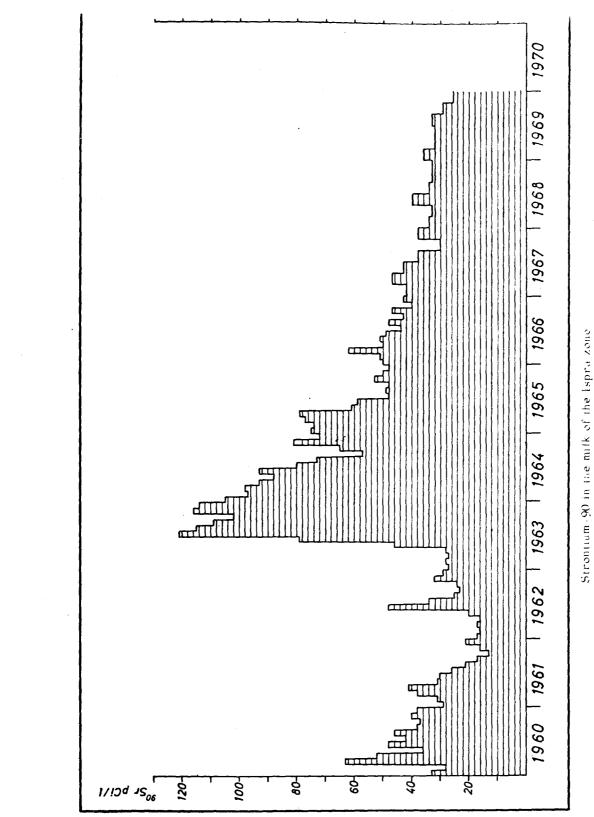
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Month	Gross beta pCi/m ³	⁹⁰ Sr 10 ⁻³ pCi/m ³	⁸⁹ Sr 10 ⁻³ pCi/m ³	¹³⁷ Cs 10 ⁻³ pCi/m ³	239 _{Pu} 10 ⁻⁵ pCi/m ³	238 _{Pu} 10 ⁻⁵ pCi/m ³
October	0.22	1.6	3.6	1.8	3.7	0.40
November	0.12	0.95	2.2	1.7	2.3	0.44
December	0.14	1.5	1.9	2.1	1.5	0.41
	1 1 1	f t t	t t t	t 4 1 8		

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PART IV

RECENT PUBLICATIONS RELATED TO RADIONUCLIDE STUDIES

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