

# DEPARTMENT OF HEALTH NEW ZEALAND

# ENVIRONMENTAL RADIOACTIVITY ANNUAL REPORT

1982 ISSN 0110 - 9944

P.O.BOX 25-099, CHRISTCHURCH NEW ZEALAND

**MARCH 1983** 

P ENV

#### **ACKNOWLEDGEMENT**

We gratefully acknowledge the assistance given by the staff of this and other Government Departments, especially the New Zealand Meteorological Service and the managers of milk processing plants. Without their co-operation the monitoring programme would not be possible. The Laboratory's Environmental Radioactivity Section organised the monitoring and analysed the samples. The Officer in Charge, Mr L.P. Gregory, was responsible for reporting and interpreting results. He was assisted professionally by Dr K.M. Matthews, and technically by Mrs M. Marsh and Miss M-J Okey.

for H.R. Atkinson (Director)

Published with the authority of the Director-General of Health

#### UNITS AND CONVERSION FACTORS

The International System of Units (SI) is used in this report. Conversion factors are given below for the relationship between SI units and those used prior to 1980. The SI unit of radioactivity, the becquerel (Bq), is 1 disintegration per second. The multiple of this unit used here is the megabecquerel (MBq) =  $10^6$  Bq, and the submultiple used is the millibecquerel (mBq) =  $10^{-3}$  Bq.

Deposition is given in megabecquerels per square kilometre (MBq/km²).

To convert to mCi/km² multiply by 0.027.

#### Concentration

in air: is given in millibecquerels per cubic metre  $(mBq/m^3)$ .

To convert to pCi/m³ multiply by 0.027.

in rain: is given in becquerels per litre (Bq/l). It is calculated by dividing the deposition by the rainfall in millimetres.

To convert to pCi/l multiply by 27.

in milk: strontium-90 is given in becquerels per gram of calcium (Bq/gCa).

caesium-137 is given in becquerels per gram of potassium (Bq/gK).

To convert to pCi/gCa or pCi/gK multiply by 27.

(One litre of milk contains about 1.2 g of calcium and about 1.4 g of potassium.)

#### LIMIT OF DETECTION AND PRECISION OF MEASUREMENT

In recent years some sample activities have been below the limit of detection. The detection limit ( $L_D$  counts) used here is calculated from the formula:  $L_D = 4.65 \, \sqrt{B}$ , where B is the background counts during the counting interval (EML Procedures Manual, HASL-300, Section D-08. U.S. Dept of Energy). The limit of detection is given in appropriate units in a footnote to the tables where relevant.

The precision of measurement is based on a counting error of 2 Poisson standard deviations (95% confidence level).

#### REFERENCE LEVELS

The following levels, against which measured levels reported here may be compared, have been adopted for New Zealand:

Mixed fission products between 10 and 80 days old (Total Beta Activity)

in air: 11 000 mBg/m<sup>3</sup> in rain: 220 Bg/1

strontium-90 in milk: 10 Bq/gCa caesium-137 in milk: 260 Bq/gK

#### SUMMARY

During 1982 the average deposition of strontium-90 at New Zealand monitoring stations was 1.2 megabecquerels per square kilometre ( $MBq/km^2$ ). This is the lowest annual deposition since monitoring commenced in 1960.

Large-scale nuclear weapons tests were conducted by the USSR and USA in 1961 and 1962. Subsequently average strontium-90 depositions at New Zealand stations increased to maximum values of 130 MBq/km² in 1964 and 110 MBq/km² in 1965. French atmospheric nuclear weapons tests were conducted in the South Pacific from 1966 to 1974. During this time average depositions ranged from 9.4 to 51 MBq/km². In 1975 these tests were replaced by underground tests and since then strontium-90 depositions have decreased significantly. Monitoring of total beta activity in air and rain at 5 Pacific Island stations continues. Fresh fission products from possible venting of underground tests have not been detected.

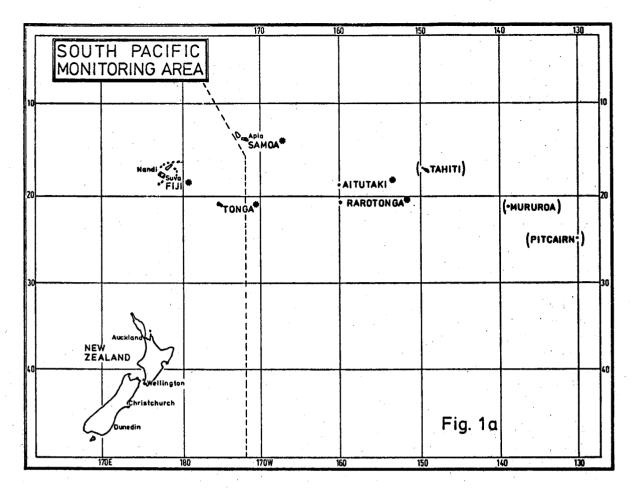
The concentrations of strontium-90 and caesium-137 in New Zealand milk reflect the changes in fallout deposition. Average concentrations during recent years are the lowest since measurements started.

The levels recorded during 1982 are very small fractions of the reference levels and thus do not constitute a public health hazard. Moreover, the radiation dose to the general population resulting from the long-term average levels, summarised herein, is small compared not only with the dose from the natural background but also with that from common variations in the natural background.

#### CONTENTS

1
1
3
5
5
8

page



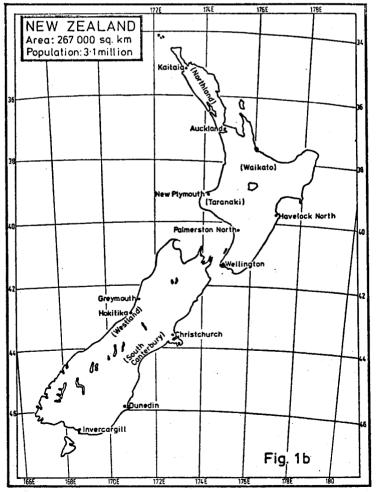


FIG. 1 MONITORING AND COLLECTING STATIONS ON PACIFIC ISLANDS AND IN NEW ZEALAND.

#### INTRODUCTION

This report continues the series of reports on environmental radioactivity monitoring in New Zealand, and in the South Pacific area, which have been published since 1961.

In September 1974 France terminated atmospheric nuclear testing which had been conducted in the Tuamotu Archipelago in the South Pacific since 1966, and in June 1975 commenced underground testing in the same area. The Laboratory's programme was changed to monitor any venting to the atmosphere of fission products from underground tests. The current programme consists of continuous air and rainwater monitoring at 5 Pacific island and 4 New Zealand stations. Samples are sent to the Laboratory for measurement of total beta activity.

The routine programme, monitoring long-term radioactive fallout from earlier atmospheric tests, was also continued during 1982. Here emphasis is given to the measurement of the two most potentially hazardous long-lived radionuclides, strontium-90 and caesium-137. Depositions of strontium-90 in rain are measured at 9 New Zealand and 2 Pacific island stations. (Naturally occurring lead-210 is also evaluated concurrently.) Strontium-90 and caesium-137 concentrations are measured in milk from 9 New Zealand stations.

Monitoring and sample collecting stations are shown in Fig. 1 a and b.

Fallout levels in recent years and particularly since 1976 have been very low and reports covering this period have been abbreviated. The reader is referred to the earlier annual reports (1), and special reports on French atmospheric nuclear tests (2). These give additional information on terms of reference. potential health hazard, adoption of reference levels, and technical information on procedures. They also include graphical presentations of results allowing historical and geographical comparisons.

#### TOTAL BETA ACTIVITY IN AIR AND RAIN

Normally the short-lived decay products of naturally occurring radon account for most of the beta activity in air. Ground level air over continents has a beta activity commonly ranging between 2 000 and 20 000 millibecquerels per cubic metre (mBq/m3), but under certain conditions the beta activity may be up to ten times the upper value of this range.

#### (1) "Environmental Radioactivity":

Annual Report 1971, Report No. NRL-F/48, June 1972 (summarising previous results)

Annual Report 1972, Report No. NRL-F/50, April 1973

Annual Report 1973, Report No. NRL-F/52, June

Annual Report 1974, Report No. NRL-F/54, June

Annual Report 1975, Report No. NRL-F/55, June 1976

Annual Report 1976, Report No. NRL-F/56, April 1977 (abridged)

Annual Report 1977, Report No. NRL-F/57, April 1978 (abridged)

Annual Report 1978, Report No. NRL-F/58, April 1979 (abridged)

Annual Report 1979, Report No. NRL-F/59, April 1980 (abridged)

Annual Report 1980, Report No. NRL-F/60, April 1981 (abridged) Annual Report 1981, Report No. NRL-F/61, March 1982 (abridged)

"Environmental Radioactivity. Fallout from Nuclear Weapons Tests Conducted by France in the South Pacific . . . and comparisons with previous test series." Report Nos: NRL-F/47, March 1972 (summarising all previous monitoring results since 1966); NRL-F/49, October 1972; NRL-F/51 November 1973, and NRL-F/53, November 1974.

Air filter and rainwater samples are measured 4 days after collection when naturally occurring beta activity has decayed and any residual beta activity from radioactive fallout can be assessed. Hereafter the term "total beta activity" refers to this residual fission product radioactivity in the sample and excludes naturally occurring radioactivity.

#### 1. Fission Products in Air

During 1982 air was monitored continuously at the New Zealand and Pacific island stations listed in Table 1. The air filters were changed 3 times each week and were despatched to the Laboratory for measurement of total beta activity. Average concentrations each month during 1982 did not exceed the limit of detection at any station. Because the levels were so low, individual results are not tabulated in the Appendix. However, average concentrations during the year are included in Table 1 for comparison with those of previous years.

TABLE 1 - Total Beta Activity in Air: Average Concentrations (mBq/m3)

	Nev	v Zealand	Station	ıs*	1	Pacifíc I	sland S	tations*	
•	AK	WN	HK	CH	FJ	SM	TO	AI	RA
1966 1 <b>9</b> 67	5.2 3.0	(3.7) 1.9		4.1 2.2		19	966 <b>-</b> 197	' <b>4</b>	· 3 72
1968	4.4	3.7		2.6	(Fren	ch atmosp	heric n	uclear t	ests)
1969 1970	4.4 5.9	3.3 4.4	(4.4)	2.6 3.7		nge: 1.5 Pacific	- 230	mBq/m³	+he
1971	7.8	4.4	5.9	5.6		ecial mor		and the second second second	
1972	2.2	1.9	1.9	1.9	Sp	eciai moi.	II COL III 9	program	unes
1973	0.7	0.4	0.7	0.7				14	tradición de
1974	3.0	1.9	2.6	1.9				100	10g - C - 1
1975	1.1	1.1	1.1	0.7	(<0.3)	(<0.3)	(0.4)	(<0.3)	· (0.4)
1976	0.4	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3
1977	<0.3	<1.1	<0.3	<0.3	<0.3	<0.3	<1.1	<1.1	<0.3
1978	<0.3	<0.3	<0.3	<0.3	<1.1	<1.1	<1.1	<1.1	<1.1
1979	<0.3	<0.3	<0.3	<0.3	<1.1	<1.1	<1.1	<1.1	<1.1
1980	<0.3	<0.3	<0.3	<0.3	<1.1	<1.1	<1.1	<1.1%	√<1. l
1981	<0.3	<0.3	<0.3	<0.3	<1.1	<1.1	<1.1	<1.1	<1.1
1982	<0.3	<0.3	<0.3	<0.3_	<1.1	<1.1	<1.1	<1.1	<1.1

<sup>\*</sup> The stations are: Auckland, Wellington, Hokitika, and Christchurch; Fiji, Samoa, Tonga, Aitutaki, and Rarotonga.

The limit of detection is 0.3 or 1.1 mBq/m3 depending on air samplers used.

The concentration of fission products in air at New Zealand stations was latitude dependent during the French South Pacific atmospheric tests (1966-1974). For example, concentrations were usually higher at Auckland than at Christchurch. Moreover, at the Pacific island stations concentrations were significantly higher, e.g., during the special monitoring programmes, lasting from 3 to 6 months, average concentrations ranged from 1.5 - 230 mBq/m³ depending on the locality, the meteorological conditions, and the extent of nuclear testing (2).

Since mid-1975, about 9 months after the termination of the French atmospheric nuclear tests, concentrations have been near or below the limit of detection at all stations. No fresh fission products have been detected since underground testing started.

Concentrations of fission products in air tabulated here, particularly those during recent years, are very small fractions of the reference levels.

<sup>( )</sup> Incomplete year.

#### 2. Fission Products in Rain

During 1982 weekly funnel and bottle collections of rainwater were made at the same stations providing air filter samples. The rainwater samples were despatched to the Laboratory for processing and measurement of total beta activity. Results are given in deposition units - megabecquerels per square kilometre (MBq/km²). Quarterly cumulative results during 1982 are given in Table 7, Appendix. Annual totals, rounded to the nearest 10 MBq/km², are listed for each station in Table 2.

TABLE 2 - Total Beta Activity in Rain: Total Deposition (MBq/km²)

	Ne	w Zealan	d Station	s*		Pacific	Island S	Stations	k
•	AK	WN	HK	СН	FJ	SM	TO	ΑĪ	RA
1963				1040					
1964				560					
1965				630					
1966			(3920)†	1180				_	
1967			2850	520			1966 – 197		
1968	*		7590	1040	(Frei	nch atmo	spheric i	nuclear	tests)
1969			2260	670	ra	ange: l	00 - 20	000 MBa/	km²
1970	(3740)	(2780)	4920	960	at		c Island.		
1971	3630	2960	3660	1180			onitorin		
1972	930	8 10	1220	560	-			J <u>F</u> J	
1973	190	260	300	150					
1974	2180	2220	1550	810					
1975	340	470	710	460	(90)	(140)	(70)	(110)	(140)
1976	90	110	160	60	100	110	150	140	110
1977	70	100	120	60	110	170	130	140	150
1978	120	100	180	70	160	230	210	230	220
1979	100	110	190	60	120	120	250	220	190
1980	70	100	140	60	100	180	70	200	160
1981	. 100	110	220	80	170	210	180	180	190
1982	80	100	180	60	130	160	130	170	140

<sup>\*</sup> See Table 1 (footnote) for station names.

Levels of fisson products in rainwater samples have been very low at all stations during recent years. During the period 1976 to 1982 annual depositions ranged from  $60 - 250 \text{ MBq/km}^2$  and average concentrations ranged from 0.03 - 0.15 becquerels per litre (Bq/1). These concentrations are very small fractions of the reference level.

#### STRONTIUM-90 DEPOSITION

#### 1. Routine Measurement

Strontium-90 deposition measurements started at 6 stations in New Zealand in 1960, and at Suva in 1961. Since 1963 measurements have been made at 9 New Zealand stations, and since 1967 at Rarotonga also. Collections are made continuously in high-walled stainless steel pots which are changed each month. The rainwater sample is passed through a column of cation exchange resin on site. The resin is then mailed to the Laboratory for measurement of strontium-90. Because the deposition rates have been so low in recent years, samples collected since 1976 have been aggregated quarterly to obtain increased measurement sensitivity.

Quarterly results for deposition, concentration, and rainfall at each station during 1982 are given in Table 8, Appendix. Annual depositions at each station since measurements commenced are listed in Table 3.

<sup>()</sup> Incomplete year.

<sup>†</sup> Commenced Greymouth Jul '66, transferred to Hokitika Jan '76.

TABLE 3 - Annual Deposition of Strontium-90 (MBq/km<sup>2</sup>)

			Ne	w Zeal	and St	ations	*				Is	ific land ions*
	KA	AK	NP	HN	WN	HK	CH	DN	IN	NZ Av	SU	ŘA
1960		46		26	28	56	18		17	32		
1961		40		28	42	81	24		43	43	38	
1962		65		35	68	100	26		43	56	60	
1963	67	72	7.3	37	75	140	46	38	63	68	90	
1964	150	150	200	59	130	290	48	66	110	130	91	
1965	110	110	150	62	140	220	63	74	100	110	73	
1966	58	47	71	31	59	81	26	27	41	49	44	
1967	38	33	46	18	35	63	14	24	32	34	28	(33)
1968	33	27	35	21	32	53	16	15	20	28	38	24
1969	57	46	54	25	39	83	26	25	45	44	46	27
1970	37	32	45	22	45	79	19	20	25	36	33	35
1971	75	50	68	38	44	92	24	30	40	51	(56)	(12)
1972	33	27	32	18	30	65	16	21	32	30	35	28
1973	16	9.6	13	7.4	14	2.1	5.9	6.7	11	12	14	(21)
1974	13	8.5	11	5.6	10	17	6.3	5.9	7.0	9.4	11	11
1975	9.5	8.9	11	5.6	10	21	7.0	6.3	9.6	9.9	8.5	5.2
1976	4.1	3.7	4.8	2.6	5.6	7.0	3.3	3.0	3.3	4.2	4.4	4.4
1977	2.6	3.0	2.6	1.5	3.7	5.2	1.1	1.5	1.9	2.6	3.0	1.1
1978	3.0	1.9	2.2	1.9	2.2	5.6	1.1	1.9	1.5	2.4	3.0	1.5
1979	3.3	2.6	3.0	2.2	3.3	5.9	1.1	1.5	1.5	2.7	2.6	0.7
1980	2.5	1.9	2.3	1.7	2.6	4.6	1.2	1.2	1.7	2.2	1.7	2.0
1981	2.0	1.1	1.8	1.0	1.5	4.2	0.7	1.4	1.2	1.7	1.4	0.9
1982	1.3	0.9	1.5	0.9	1.1	2.8	0.7	0.6	0.9	1.2	0.9	0.6
Annua	l Rain	fall (1	mm)†									

1410 1070 1430 810 1410 2600 640 690 1060 3210 1830

\* The New Zealand stations are: Kaitaia, Auckland, New Plymouth, Havelock

North, Wellington, Hokitika (Greymouth before 1976), Christchurch, Dunedin

It is evident that the deposition of strontium-90 is rainfall dependent and high rainfall areas such as Hokitika show elevated values compared to low rainfall areas such as Christchurch.

The large-scale Northern Hemisphere (USSR) and Pacific area (USA) nuclear tests, conducted in 1961 and 1962 before the signing of the Partial Test Ban Treaty, resulted in a delayed stratospheric fallout over New Zealand. Maximum annual depositions occurred in 1964 and 1965, exceeding a station average of 100 MBq/km<sup>2</sup>. Annual depositions then decreased markedly.

From 1966 to 1974 smaller-scale atmospheric nuclear tests were conducted by France in the South Pacific each year except 1969. Each series, lasting from 1 to 3 months and comprising from 3 to 8 nuclear explosions, took place during the Southern Hemisphere winter. Forty-one nuclear devices were reported to have been exploded, most of them being in the low to medium power (kiloton) range. However, megaton explosions were reported to have taken place on 5 occasions (2). The annual deposition of strontium-90 in New Zealand increased again during the period 1969 to 1971 reaching a second smaller maximum in 1971, less than one-half of the 1964 maximum. Since then the annual depositions have decreased progressively. During 1982 the average deposition at New Zealand stations was 1.2 MBq/km², less than 1% of the 1964 maximum.

and Invercargill. The Pacific island stations are: Suva (Fiji) and Rarotonga. † Mean annual rainfall, since 1968, rounded to the nearest 10 mm. Values in parenthesis are estimates.

Interpretation of the long-term results summarised in Table 3 and commentary on the origin of the strontium-90, fallout mechanism, latitudinal effect and rainfall dependence, were made in earlier reports (1).

#### LEAD-210 DEPOSITION

Lead-210 is a naturally occurring radionuclide produced in the atmosphere by decay of gaseous radon exhaled from land surfaces. Lead-210 deposition is measured along with strontium-90 in the same monthly rainwater collections.

Individual monthly depositions during 1982 are given in Table 9, Appendix. Annual depositions at each station since 1968 are listed in Table 4.

TABLE 4 - Annual Deposition of Lead-210 (MBq/km<sup>2</sup>)

											Is1	ific and
			Net			Station					Stati	
	KA	AK	NP	HN	WN	HK	CH	DN	IN	NZ $Av$	SU	RA
1967†	23	43	64	27	38	88	13	21	37	39	46	22
1968	65	61	77	31	69	118	24	28	39	57	91	23
1969	68	49	57	33	44	146	21	34	48	<i>56</i>	71	36
1970	53	37	67	23	56	114	24	27	34	48	68	31
1971	77	38	36	24	47	87	19	27	40	44	68	
1972	84	60	74	33	63	126	26	45	48	62	98	
1973	71	53	85	30	67	122	18	25	30	56	78	_
1974	46	40	65	28	60	105	26	24	21	46	69	148
1975	60	56	67	41	73	146	34	34	41	61	108	52
1976	52	49	53	34	54	101	26	26	28	47	75	_
1977	49	41	62	28	57	97	23	27	40	47	62	
1978	60	42	60	28	52	107	27	33	37	50	83	-
1979	70	56	70	38	76	167	27	34	38	64	79	_
1980	74	49	66	34	68	144	26	41	48	61	74	_
1981	61	40	61	28	61	146	18	27	34	<b>53</b>	-	_
1982	55	39	62	20	56	114	18	25	47	48	_	_

<sup>\*</sup> See Table 3 for station names.

The deposition of lead-210 is rainfall dependent like strontium-90. However, unlike strontium-90 there is little significant change in the New Zealand station average annual deposition which is consistently about 50 MBq/km $^2$ . Because of the current low strontium-90 deposition, lead-210 deposition is now about 40 times higher.

### STRONTIUM-90 AND CAESIUM-137 IN MILK

Monitoring of strontium-90 in New Zealand milk started in 1961, and caesium-137 in 1964. Since 1965 measurements have been made on milk samples from 9 collecting stations. Caesium-137 and potassium are determined monthly by gamma spectroscopy. Samples are then aggregated quarterly for strontium-90 and calcium determinations.

#### 1. Strontium-90

Quarterly average concentrations of strontium-90 in milk at each station during 1982 are given in Table 10, Appendix. Annual average concentrations at each station since 1961 are listed in Table 5.

<sup>†</sup> May-December only.

TABLE 5 - Strontium-90 in Milk: Annual Averages (Bq/gCa)

			(	Collect	ting Sta	ations*				
	ND	AK	WK	TA	PN	WN	WD ·	СН	DN	Average
1961	0.17		0.15	0.26			0.47	0.059		0.22
1962	0.23	0.20	0.18	0.35	0.16		0.50	0.078	0.11	0.23
1963	0.28	0.20	0.21	0.37	0.18		0.64	0.10	0.14	0.26
1964	0.41	0.34	0.35	0.63	0.26		0.96	0.096	0.15	0.40
1965	0.39	0.35	0.36	0.62	0.31	0.33	1.07	0.16	0.27	0.43
1966	0.24	0.23	0.23	0.46	0.18	0.23	0.84	0.089	0.15	0.29
1967	0.19	0.19	0.19	0.38	0.14	0.20	0.66	0.070	0.11	0.24
1968	0.15	0.14	0.15	0.30	0.13	0.18	0.52	0.059	0.089	0.19
1969	0.23	0.22	0.20	0.35	0.21	0.19	0.66	0.063	0.11	0.25
1970	0.19	0.19	0.19	0.36	0.13	0.17	0.78	0.081	0.093	0.24
1971	0.27	0.21	0.22	0.38	0.19	0.18	0.68	0.074	0.11	0.26
1972	0.18	0.17	0.16	0.30	0.19	0.15	0.54	0.070	0.11	0.21
1973	0.14	0.13	0.13	0.21	0.10	0.13	0.40	0.044	0.070	0.15
1974	0.12	0.11	0.10	0.20	0.093	0.11	0.33	0.048	0.070	0.13
1975	0.11	0.10	0.11	0.19	0.089	0.13	0.32	0.044	0.059	0.13
1976	0.096	0.089	0.093	0.13	0.059	0.089	0.23	0.041	0.041	0.096
1977	0.078	0.078	0.089	0.14	0.052	0.078	0.19	0.037	0.044	0.087
1978	0.063	0.078	0.081	0.11	0.056	0.078	0.14	0.037	0.041	0.076
1979	0.059	0.067	0.093	0.16	0.056	0.070	0.14	0.037	0.033	0.079
1980	0.059	0.060	0.088	0.14	0.049	0.066	0.13	0.032	0.035	0.073
1981	0.053	0.054	0.074	0.12	0.043	0.054	0.12	0.027	0.027	0.063
1982	0.050	0.051	0.069	0.10	0.030	0.047	0.093	0.023	0.024	0.054
Average	0.17	0.16	0.16	0.28	0.13	0.14	0.47	0.062	0.090	0.19

<sup>\*</sup> The stations are: Northland, Auckland, Waikato, Taranaki, Palmerston North, Wellington, Westland, Christchurch, and Dunedin.

Average concentrations of strontium-90 in New Zealand milk reached maximum values of 0.40 and 0.43 Bq/gCa during 1964 and 1965 when the rate of strontium-90 deposition was also a maximum. Levels then decreased, reaching a minimum of 0.19 Bq/gCa in 1968, indicating that levels in milk are dependent on the rate of fallout. However, during the period 1965-68 strontium-90 in milk decreased at a slower rate than strontium-90 deposition, indicating some uptake by grass of the cumulative deposit in the soil. After the start of French Pacific nuclear tests in 1966, concentrations increased slightly during the period 1969-71. However, with decreasing fallout depositions since 1973, concentrations in milk decreased again. The average concentration during 1982, 0.054 Bq/gCa, was the lowest recorded since measurements commenced.

Milk samples from the lowest and highest rainfall stations, i.e., Christchurch and Westland, give the range of strontium-90 contamination in New Zealand milk. Generally the extent of this range is from about one-half to about twice the country-wide average.

#### 2. Caesium-137

Monthly average concentrations of caesium-137 in milk at each station during 1982 are given in Table 11, Appendix. Annual average concentrations at each station since 1964 are listed in Table 6.

TABLE 6 - Caesium-137 in Milk: Annual Averages (Bq/gK)

		. ´		Collect	ting Sta	ations*				
	ND	AK	WK	TA	PN	WN	WD	СН	DN	Average
1964	1.8	1.9	2.6	6.2	0.7		2.8	0.2	0.4	2.1
1965	2.0	2.0	3.1	6.9	1.0	1.1	2.9	0.4	0.7	2.2
1966	1.4	1.2	2.2	5.2	0.4	0.7	1.6	0.1	0.3	1.5
1967	1.0	1.0	1.8	4.6	0.3	0.5	1.2	0.1	0.2	1.2
1968	0.6	0.7	1.3	3.8	0.1	0.3	0.8	<0.1	0.1	0.9
1969	1.0	1.0	1.5	3.7	0.2	0.3	1.4	0.1	0.1	1.0
1970	0.8	0.7	1.3	3.3	0.2	0.4	1.4	0.1	0.2	0.9
1971	0.9	0.7	1.3	3.0	0.3	0.3	1.1	0.1	0.2	0.9
1972	0.8	0.6	1.0	2.7	0.1	0.3	0.8	0.1	0.1	0.7
1973	0.5	0.3	0.8	1.8	0.1	0.1	0.5	<0.1	0.1	0.5
1974	0.3	0.3	0.6	1.5	0.1	0.1	0.3	<0.1	<0.1	0.4
1975	0.3	0.3	0.5	1.3	<0.1	0.1	0.3	<0.1	<0.1	0.3
1976	0.2	0.2	0.4	0.9	0.1	0.1	0.1	<0.1	0.1	0.2
1977	0.2	0.1	0.4	1.1	<0.1	0.1	0.2	<0.1	<0.1	0.3
1978	0.1	0.1	0.3	0.7	<0.1	<0.1	0.1	<0.1	<0.1	0.2
1979	0.1	0.1	0.4	1.2	<0.1	<0.1	0.1	<0.1	<0.1	0,2
1980	0.2	0.2	0.4	1.0	<0.1	<0.1	0.2	<0.1	<0.1	0.2
1981	<0.1	<0.1	0.2	0.9	<0.1	<0.1	<0.1	<0.1	<0.1	0.1
1982	<0.1	<0.1	0.3	0.9	<0.1	<0.1	<0.1	<0.1	<0.1	0.1
Average	0.6	0.6	1.1	2.7	0.2	0.3	0.8	0.1	0.2	0.7

<sup>\*</sup> See Table 5 for station names.

The highest concentrations of caesium-137 in milk were recorded in 1964 and 1965 when strontium-90 deposition and concentration in milk were also at their highest. Concentrations have decreased steadily since then. During the last 2 years average concentrations were 0.1 Bq/l, the lowest recorded since measurements commenced.

The "soil effect" leading to high caesium-137 concentrations in milk at Taranaki, and to a lesser extent at Waikato, has been the subject of a special survey (Annual Report 1977 (1)).

## 3. Comparison of Measured Levels with the Reference Levels

When measured levels in milk are compared with the reference levels, long-term averages are more meaningful. Since measurements commenced, the country-wide average levels of strontium-90 (0.19 Bq/gCa) and caesium-137 (0.7 Bq/gK) have been 2% and 0.3% of the reference levels respectively. The stations with the highest levels of contamination have corresponding percentages about 2.5 times and 4 times higher respectively.

Thus the long-term average levels, even at the stations with highest concentrations, are very small fractions of the reference levels and do not constitute a public health hazard.

### APPENDIX

TABLE 7 - Total Beta Activity in Rain 1982 (Weekly Collections):

Cumulative Rainfall (mm)
Cumulative Deposition (MBq/km²)
Average Concentration (Bq/1)

New Zealand Stations	lst Quarter	2nd Quarter	3rd Quarter	4th Quarter	1982
Auckland mm MBq/km <sup>2</sup> Bq/l	168 21 ± 7 0.13 ± 0.04	304 23 ± 7 0.08 ± 0.02	171 22 ± 7 0.13 ± 0.04	175 15 ± 7 0.09 ± 0.04	818 81 ± 14 0.10 ± 0.02
Wellington mm MBq/km <sup>2</sup> Bq/l	98 27 ± 7 0.28 ± 0.07	338 29 ± 7 0.09 ± 0.02	256 25 ± 7 0.10 ± 0.03	238 17 ± 6 0.07 ± 0.03	930 98 ± 14 0.11 ± 0.02
Hokitika mm MBq/km <sup>2</sup> Bq/l	518 49 ± 8 0.09 ± 0.02	483 38 ± 7 0.08 ± 0.01	623 49 ± 7 0.08 ± 0.01	815 47 ± 8 0.06 ± 0.01	2439 183 ± 15 0.08 ± 0.01
Christchurch mm MBq/km <sup>2</sup> Bq/1	49 18 ± 7 0.37 ± 0.14	102 15 ± 7 0.15 ± 0.07	92 18 ± 7 0.20 ± 0.08	207 13 ± 7 0.06 ± 0.03	450 64 ± 14 0.14 ± 0.03
Average MBq/km² Bq/l	29 0.22	26 0.10	29 0.13	23 0.07	107 0.11
Pacific Island Stations					
	1071 65 ± 19 0.06 ± 0.02	174 32 ± 17 0.18 ± 0.10	235 13 ± 18 0.06 ± 0.08	260 19 ± 14 0.07 ± 0.05	1740 129 ± 34 0.07 ± 0.02
Stations Nandi, Fiji mm MBq/km²	65 ± 19	$32 \pm 17$	13 ± 18	$19 \pm 14$	129 ± 34
Stations Nandi, Fiji mm MBq/km² Bq/1 Samoa mm MBq/km²	$65 \pm 19 \\ 0.06 \pm 0.02$ $1345 \\ 92 \pm 27$	32 ± 17 0.18 ± 0.10 244 20 ± 19 0.08 ± 0.08 286 31 ± 22	13 ± 18 0.06 ± 0.08 440 36 ± 19 0.08 ± 0.04 346 21 ± 20	19 ± 14 0.07 ± 0.05 314 13 ± 19 0.04 ± 0.06	129 ± 34 0.07 ± 0.02 2343 161 ± 43 0.07 ± 0.02 2018 130 ± 43
Stations Nandi, Fiji mm MBq/km² Bq/1 Samoa mm MBq/km² Bq/1 Tonga mm MBq/km²	65 ± 19 0.06 ± 0.02 1345 92 ± 27 0.07 ± 0.02 1254 35 ± 24	32 ± 17 0.18 ± 0.10 244 20 ± 19 0.08 ± 0.08 286 31 ± 22 0.11 ± 0.08 312 67 ± 21	13 ± 18 0.06 ± 0.08 440 36 ± 19 0.08 ± 0.04 346 21 ± 20 0.06 ± 0.06 270 40 ± 20	19 ± 14 0.07 ± 0.05 314 13 ± 19 0.04 ± 0.06 132 43 ± 20 0.33 ± 0.15 48 38 ± 19	129 ± 34 0.07 ± 0.02 2343 161 ± 43 0.07 ± 0.02 2018 130 ± 43 0.06 ± 0.02

The error term is  $\pm 2$  standard deviations

TABLE 8 - Strontium-90 in Rain 1982:

Rainfall (mm)
Deposition (MBq/km²)
Concentration (Bq/1)

New Zealand 'Stations		lst Quarter	2nd Quarter	3rd Quarter	4th Quarter	<u>Total</u>	Av
Kaitaia	mm MBq/km² Bq/1	193 0.4 0.002	303 0.3 0.001	309 0.4 0.001	176 0.2 0.001	981 1.3	0.001
Auckland	mm MBq/km² Bq/1	168 0.3 0.002	305 0.2 <0.001	172 0.3 0.001	176 0.1 <0.001	821 0.9	0.001
New Plymouth	mm MBq/km² Bq/1	260 0.5 0.002	326 0.4 0.001	343 0.2 <0.001	316 0.4 0.001	12 <b>45</b> 1.5	0.001
Havelock North	mm MBq/km² Bq/1	85 0.3 0.003	246 0.5 0.002	127 0.1 0.001	64 <0.1 <0.001	522 0.9	0.002
Wellington	mm MBq/km² Bq/1	230 0.4 0.002	363 0.3 <0.001	250 0.1 <0.001	207* 0.3* <0.001	1050 1.1	<0.001
Hokitika	mm MBq/km² Bq/1	558 1.0 0.002	595 0.4 <0.001	681 0.5 <0.001	789 0.9 0.001	2623 2.8	0.001
Christchurch	mm MBq/km² Bq/1	45 0.2 0.005	104 0.1 0.001	92 0.2 0.002	207 0.2 <0.001	<b>448</b> 0.7	0.002
Dunedin	mm MBq/km² Bq/1	190 0.3 0.002	163 0.2 <0.001	90 <0.1 <0.001	339 0.1 <0.001	782 0.6	<0.001
Invercargill	mm MBq/km <sup>2</sup> Bq/1	381 0.4 0.001	287 0.2 <0.001	252 <0.1 <0.001	395 0.3 <0.001	1315 0.9	<0.001
Average	mm MBq/km <sup>2</sup> Bq/l	234 0.4 0.002	299 0.3 <0.001	257 0.2 <0.001	297 0.3 <0.001	1087 1.2	0.001
Pacific Island Stations							
Suva, Fiji	mm MBq/km² Bq/1	1060 0.3 <0.001	725 <0.1 <0.001	573 0.4 <0.001	377 0.2 <0.001	2735 0.9	<0.001
Rarotonga	mm MBq/km² Bq/1	377 0.2 <0.001	429 0.1 <0.001	241 <0.1 <0.001	56 0.3 0.005	1103 0.6	<0.001

<sup>\*</sup> Results for 2 months only

The counting error for deposition results is about  $\pm \ 0.08 \ \mathrm{MBg/km^2}$ 

- Lead-210 in Rain 1982: Deposition  $(MBq/km^2)$ TABLE 9

New Zealand											•		
Stations	Jan	Feb	Mar	Apr	May	Jun	Jul	Aug	Sep	Oct	Nov	Dec	Tota1
Kaitaia	2.4	4.0	3.6	3.6	2.1	4.1	10.8	4.3	6.5	5.9	4.9	2.7	55
Auckland	1.9	2.0	4.5	3.9	4.4	2.5	2.8	2.8	3.1	5.5	(2.0)	3.1	39
New Plymouth	(1.9)	5.1	6.3	5.6	5.2	2.5	3.8	7.3	4.6	5.4	6.7	7.1	62
Havelock North	1.2	2.1	1.9	2.5	0.8	3.6	1.5	1.3	1.7	1.3	(1.2)	(0.5)	20
Wellington	1.4	4.1	3.7	2.3	7.6	5.7	0.9	2.9	6.7	3.7	(5.0)	(1.0)	. 56
Hokitika	14.4	8.4	9.9	5.8	15.8	6.2	8.9	1.4	3.2	12.8	14.9	17.5	114
Christchurch	1.3	1.2	6.0	8.1	1.3	1.7	1.5	0.1	.8	2.1	(1.0)	(2.0)	18
Dunedin	3.8	1.7	1.6	0.7	2.0	1.7	0.5	1.2	0.7	3.0	3.8	4.1	25
Invercargill	4.6	3.1	1.7	1.2	1.5	9.4	2.2	8.7	1.3	4.2	8.9	7.3	47
Average	3.7	3.5	3.4	3.0	4.5	3.6	4.0	3.4	3.3	4.9	5.1	5.7	48
Pacific Island													
Stations													
Suva, Fiji	NS	3.1	9.4	5.5	4.8	7.1	9.0	7.0	5.8	2.0	7.5	2.2	ļ
Rarotonga	1.6	NS	1.3	3.1	3.7	1.2	2.4	5.1	2.1	NS	NS	NS	ı

The counting error is less than 0.3 MBq/km $^2$  (less than 0.2 for most results). NS No result available. ( ) Estimate.

(Bq/gCa)	)
1982:	
-90 in Milk l	
in	
TABLE 10 - Strontium-90 in Milk 1982: (Bq/gCa)	
i	
0	
TABLE	

Ĺ

	lst Quarter	2nd Quarter	3rd Quarter	4th Quarter	AV
Northland	0.050	0.049	0.052	0.049	0.050
Auckland	0.056	0.047	0.050	0.050	0.051
Waikato	0.106	0.059	0.059	0.052	0.069
Taranaki	0.102	0.089	0.103	0.110	0.101
Palmerston North	0.012	0.038	0.044	0.027	0.030
Wellington	0.055	1	0.041	970.0	0.047
Westland	0.107	0.088	0.088	0.089	0.093
Christchurch	0.024	0.020	0.025	0.022	0.023
Dunedin	0.022	0.021	0.029	0.025	0.024
Average	0.059	0.051	0.055	0.052	0.054

The counting error is about ± 0.002 Bg/gCa

TABLE 11 - Caesium-137 in Milk 1982: (Bq/gK)

AV	<0.1	<0.1	0.28	0.88	<0.1	<0.1	<0.1	<0.1	<0.1
Dec	0.14	ND	0.35	1.11	0.12	QN	NON	N N	QN
Nov	0.14	ON	0.34	0.84	ON	ON	0.10	ON	ı
0ct	R	0.13	0.33	0.87	Q.	QN	2	<u>R</u>	QN
Sep	QN	ON	0.14	0.65	ON	ON	ND	ND	ND
Aug	Ø	QN	0.18	0.15	R	N ON	ON	N	QN
Jul	0.12	QN	0.10	0.51	N N	ON	QN	QN ON	NO
Jun	Q.	ON	0.13	0.68	ND	QN	CIN	Ø	QN
May	0.14	1	0.23	0.74	R	Ø	ı	CN CN	N N
Apr	ON	0.18	0.16	0.73	R	0.10	R	R	QN Q
Mar	N	R	0.53	1.59	R	R	0.12	QN N	QN <sup>*</sup>
Feb	0.11	N ON	0.46	1.39	ON	ON	0.13	R	GN.
Jan	0.11	S	0.35	1.32	ON	ON ON	S	<u>R</u>	<u>R</u>
	Northland	Auckland	Waikato	Taranaki	Palmerston North	Wellington	Westland	Christchurch	Dunedin

The counting error is about  $\pm$  0.07 Bq/gk. ND Not detected. The limit of detection is about 0.1 Bq/gK. - Not determined.

0.1

Average