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OCCUPATIONAL HEALTH DIVISION

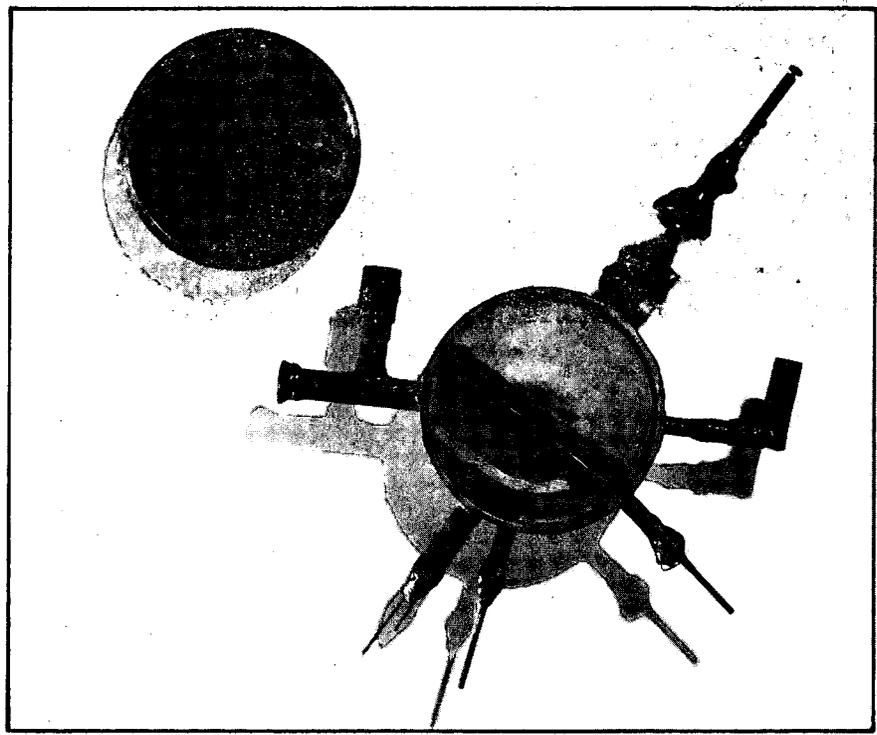
Annual Environmental Monitoring Report of the Lawrence Berkeley Laboratory

1987

Prepared by the Staff of the
Occupational Health Division

April 1988

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LBL-25110

ANNUAL ENVIRONMENTAL MONITORING REPORT
OF THE
LAWRENCE BERKELEY LABORATORY

1987

Prepared by the Staff of the
Occupational Health Division
Lawrence Berkeley Laboratory
University of California
Berkeley, California 94720

Gary E. Schleimer
Editor

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PREFACE

In 1976 R.H. Thomas published the *LBL Annual Environmental Monitoring Report* in two parts. Part I (LBL-4678) discussed in detail the modeling used to determine the population dose equivalent due to Laboratory radiological operations. That volume also described natural radiation background, geological features, climate and meteorology, and the environmental surveillance program of the Lawrence Berkeley Laboratory (LBL). Part II (LBL-4827) included only the results of the sampling and measuring programs and other data necessary to determine the environmental impact of the Laboratory's radiological operations for 1975. A format similar to LBL-4827 was used in the 1976, 1977, 1978, and 1979 Annual Monitoring Reports (LBLs 6405, 7530, 9080, and 11192, respectively).

While the 1980 Annual Report, LBL-12604, was kept brief, abstracted sections from LBL-4678 were included so that the document might stand alone. The same format has been used in this report, along with updates to LBL-4678 where appropriate, and a greatly expanded description of LBL's nonradiological environmental activities.

Readers wishing a more comprehensive discussion of LBL site characteristics and population dose modeling may obtain a copy of LBL-4678 from

Gary E. Schleimer
Environmental Health and Safety Department
Building 75, Room 112
Lawrence Berkeley Laboratory
Berkeley, CA 94720

R.O. Pauer and G.E. Schleimer of the Environmental Health and Safety Department of the Engineering Division contributed to the preparation of this report.

The bulk of the sample preparation and lab work was done by V.J. Montoya. Sample assays and computer data entry were done by W.B. Cornica. Special assays of air samples were performed by A.R. Smith.

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CONTENTS

Preface	iii
List of Tables	v
List of Figures	vii
Abstract	1
Introduction	1
1987 Environmental Monitoring Summary	3
1987 Environmental Activities and Permits Issued	4
Environmental Monitoring Results	5
Radiological Results	5
Penetrating Radiation	5
Airborne Radionuclides	9
Waterborne Radionuclides	11
Ground Water	19
Nonradioactive Pollutants	19
Waterborne Pollutants	19
Site Wastewater Discharges	25
Population Dose Resulting from LBL Operations	25
Accelerator-Produced Radiation	27
Airborne Radionuclides	28
Trends—LBL Environmental Impact	30
Accelerator-Produced Penetrating Radiation	30
Airborne and Waterborne Radionuclides	30
Quality Assurance	30
References	37

LIST OF TABLES

No.		Page
1.	Location of LBL monitoring stations.....	8
2.	Effective dose equivalent at LBL boundary due to accelerator operation, 1987	8
3.	Total quantities of radionuclides discharged into the atmosphere, 1987	11
4.	Summary of air samples, 1987.....	12
5.	Annual gross radioactivity found in LBL perimeter air samples, 1978–1987	13
6.	Summary of radioiodine in perimeter air samples, 1987	14
7.	Summary of airborne environmental HTO and $^{14}\text{CO}_2$ sampling, 1987	14
8.	Summary of perimeter airborne environmental HTO and $^{14}\text{CO}_2$ sampling, 1978–1987	15
9.	Summary of atmospheric deposition samples, 1987.....	16
10.	LBL perimeter-station deposition trends, 1978–1987	17
11.	Summary of surface- and drinking-water samples, 1987	20
12.	Summary of surface- and drinking-water samples, 1978–1987	21
13.	Summary of sewage sampling data, 1987.....	22
14.	Sanitary-sewer discharge trends, 1978–1987	23
15.	Summary of ground water samples, 1987.....	24
16.	Summary of Building 25 wastewater sample results, 1987	25
17.	Summary of Building 77 wastewater sampling results, 1987.....	26
18a.	Summary of Strawberry Sanitary Sewer sampling results, 1987	26
18b.	Summary of Hearst Sanitary Sewer sampling results, 1987.....	27
19.	Collective effective dose equivalent resulting from LBL airborne radionuclide releases, 1987	29

No.		Page
20.	Population dose equivalent, resulting from LBL operations, 1987	29
21.	LBL QAP sample results, 1987	36

LIST OF FIGURES

No.		Page
1.	Lawrence Berkeley Laboratory Buildings.....	6
2.	Environmental monitoring, Lawrence Berkeley Laboratory	18
3.	Annual accelerator-produced dose equivalent at the Olympus Gate Environmental Monitoring Station, 1959–1987	31
4.	Annual accelerator-produced dose equivalent at Building 90 Environmental Monitoring Station, 1962–1987	32
5.	Annual accelerator-produced dose equivalent at the 88-Inch Cyclotron Environmental Monitoring Station, 1963–1987.....	33
6.	Annual accelerator-produced dose equivalent at the Panoramic Way Environmental Monitoring Station, 1963–1987	34
7.	Annual releases of tritium (HTO) from the Building 75 Tritium Facility, 1969–1987	35

ANNUAL ENVIRONMENTAL MONITORING REPORT
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LAWRENCE BERKELEY LABORATORY, 1987

ABSTRACT

The Environmental Monitoring Program of the Lawrence Berkeley Laboratory is described. Data for 1987 are presented and general trends are discussed.

INTRODUCTION

Laboratory Operations

The Lawrence Berkeley Laboratory (LBL) is a multiprogram national laboratory managed by the University of California (UC) for the U.S. Department of Energy (DOE). LBL's major role is to conduct basic and applied science research that is appropriate for an energy research laboratory. LBL, birthplace of the cyclotron, was founded by the late Nobel Laureate Ernest Orlando Lawrence 56 years ago.

The Laboratory also supports nationwide university-based research by providing national facilities, including the National Center for Electron Microscopy, four large accelerators, several small accelerators, a number of radiochemical laboratories, several large gamma irradiators, and a tritium (^3H) labeling laboratory. The Bevatron (Building 51 in Fig. 1) is the most massive of LBL's accelerators. Originally designed as a 6-GeV proton synchrotron, it is presently capable of accelerating ions up to ^{40}Ca , from 20 MeV/nucleon to 2.1 GeV/nucleon, and ions up to uranium to 1 GeV/nucleon. For certain beams the SuperHILAC is used as an injector. (This combination is called the Bevalac.) The SuperHILAC (Building 71), a heavy-ion accelerator, is a multiprogrammable research accelerator in its own right and produces ion beams up to 8.5 MeV/nucleon. The 88-Inch Variable Energy Sector-Focused Cyclotron (Building 88) routinely produces intense beams of protons to about 60 MeV, alpha particles to 140 MeV, and heavy ions to mass 40 to energies of 350 MeV. The 184-Inch Synchrocyclotron (Building 6) provides alpha particle beams with energies up to approximately 1 GeV. Aside from shutdown periods, the first two of these accelerators provide beams around the clock. The 88-Inch Cyclotron provides beams ~120 hr/wk. The 184-Inch Cyclotron ceased operations in late 1987, is being dismantled, and an intense light source is to be built in its place.

The tritium facility located in Building 75 was designed to handle kilocurie quantities of tritium (a radioactive isotope of hydrogen- ^3H) used as a labeling agent for a variety of molecules subsequently employed in chemical and biomedical research. The facility was expanded during 1983 and is now funded by the National Institutes of Health.

Radiochemical and radiobiological studies performed in many laboratories at LBL typically use millicurie quantities of a great variety of radionuclides. The workplace and effluent release points are continuously sampled at all installations where significant quantities of radionuclides are handled.

The Site

LBL is situated upon a hillside above the main campus of UC. The 130-acre site is located on the west-facing slope of the Berkeley Hills, at elevations ranging from 150 to 350 meters above sea level. Most of the site is within the City of Berkeley, but about one-quarter of the eastern part is within the City of Oakland. It is located three miles east of San Francisco Bay and about fifteen miles east of San Francisco.

LBL is located in an urban environment on land owned by the University. The LBL site is bordered on the north by predominately single-family homes and on the west by multiunit dwellings, student residence halls, and commercial districts. The area to the south, which is part of the University lands, is maintained in a largely natural state and includes recreational facilities and the University Botanical

Garden. The population within an 80-km (50-mi) radius of the Laboratory is approximately 5.1 million (1980 census).¹

The Laboratory's activities are located both on site and off site. There are 67 buildings on the LBL hillside site, plus additional facilities located on the University campus, notably the Donner Laboratory of Biology and Medicine and the Melvin Calvin Laboratory. The on-site space consists of 1,350,000 gross square feet (gsf) in about 60 buildings: 1,307,000 in DOE buildings and trailers and 43,000 in University-owned buildings. Off-site space utilized by LBL consists of 260,000 gsf in various University buildings on the UCB campus and 130,000 gsf in leased facilities in Emeryville and Berkeley.

The Laboratory's population is approximately 3,850, including about 600 visiting scientists and engineers. About 3,100 are located on site, 700 are located in campus buildings, and about 50 are in off-site leased space.

The Climate

The climate of the LBL site is greatly influenced by its nearness to the Pacific Ocean and its exposure to the maritime air that flows in from the San Francisco Bay. Seasonal temperature variations are small, with a mean temperature difference between the summer 63°F and winter 48°F of only 15 degrees. Relative humidity ranges from 85–90% in the early morning to 65–75% in the afternoon. The average annual rainfall is 25 inches. About 95% of the rainfall occurs from October through April, and intensities are seldom greater than 0.5 inch per hour. Thunderstorms, hail, and snow are extremely rare. Winds are usually light, but summer sea breezes range up to 20–30 mph. Winter storm winds from the south or southwest have somewhat lesser velocities.

Geology

Most of the LBL site is underlain by complex sedimentary and volcanic rock. In general, the bedrock is relatively weak and weathers deeply. Consequently, a colluvial cover has been produced that is a few feet thick. The major geologic unit consists of poorly consolidated sandstones, siltstones, claystones, and conglomerates of relatively low strength and hardness. These rocks are blanketed by clay soils having high shrink-swell characteristics. The western and southern portions of the site are underlain by moderately well consolidated shales, siltstones, sandstones, and conglomerates. Throughout most of the upper elevations a volcanic unit overlays and is interbedded with the upper layers of the major geologic unit.

The Hydrogeology

Highly complex ground-water conditions are present at LBL. Year-round springs, annual surface seeps and variable water levels in observation wells indicate discontinuous and localized aquifers. These conditions are due to a combination of factors: open fracture volcanic flow rock, impervious claystone interbeds, permeable sandstone lenses, and irregular fracture patterns associated with past folding and faulting. During the rainy season, ground-water levels increase and cause a decrease in slope stability. Consequently, the Laboratory has installed an elaborate ground-water detection and drainage system. The drainage system uses both pumped vertical and free-flowing horizontal wells (hydraugers). Although ground-water wells are not used as a source of Laboratory or local community drinking water, two hydraugers (shown as dotted lines on Fig. 1) are sampled for gross radioactivity and tritium. Ground-water drainage feeds into Blackberry Creek on the north portion and into Strawberry Creek on the south portion of the Laboratory. Both creeks eventually flow through the Berkeley campus and then into the City of Berkeley storm drainage system, which empties into San Francisco Bay.

Water Supply

The Laboratory's primary water supply is the East Bay Municipal Utility District (EBMUD) Shasta Reservoir, which holds approximately two million gallons. The Laboratory's high pressure fire and domestic systems are supplied from this reservoir. A secondary source is the EBMUD's Berkeley View Tank, which holds approximately one million gallons. Water mains have automatic shutoff valves for protection in case of a main breakage. The LBL water distribution system operates entirely by gravity flow, requiring no pumps or energy consumption. The Laboratory has recently installed two 200,000-gallon water storage tanks at separate locations for fire protection. Diesel-powered pumps provide the necessary

flow and pressure for maintaining a reliable fire protection system during emergencies.

Sanitary Sewer Systems

The west-side LBL sanitary system connects to the City of Berkeley sewer main at Hearst Avenue. On the south side of the Laboratory, a second connection is also made to the City of Berkeley system. The Berkeley system flows to the EBMUD Sewage Treatment Facility, where the wastewater undergoes primary and secondary treatment before its discharge to San Francisco Bay. To ensure that its wastewater complies with the EBMUD discharge limits, the Laboratory monitors its wastewater for pH, toxic metals, and radioactivity. In addition, wastewater from both plating shops and a chemistry building is monitored and treated appropriately before discharge.

Storm Drainage System

Because of its hillside location and moderate annual rainfall, surface run-off is a prevalent feature at LBL. Consequently, an inclusive storm system, designed and installed in the 1960s, discharges into the Blackberry Creek watershed on the north side of LBL and the Strawberry Creek watershed on the south side. This system provides for runoff intensities expected in a 25-year maximum-intensity storm.

1987 ENVIRONMENTAL MONITORING SUMMARY

In order to establish whether LBL research activities produced any impact on the population surrounding the Laboratory, a program of environmental air and water sampling and continuous radiation monitoring was carried on throughout the year.² For 1987, as in the previous several years, dose equivalents attributable to LBL radiological operations were a small fraction of both the relevant radiation protection guidelines (RPG)³ and of the natural radiation background. [The reader should note that throughout this report the phrase "population dose" should be taken to mean collective effective dose equivalent (CEDE) and "dose" or "dose equivalent" to mean effective dose equivalent.]

The maximum effective dose equivalent delivered to a hypothetical member of the community is defined as the maximum perimeter dose equivalent. That value [the 1987 dose equivalent at the Olympus Gate Environmental Monitoring Station (MS) B-13D] was ≤ 4.1 mrem (3.5 mrem from direct radiation and 0.6 mrem from radionuclide releases), about 4% of the RPG. The hypothetical maximum exposure to an individual from airborne radionuclides would be to a person residing just outside the western LBL perimeter. The 1987 effective dose equivalent to such a person would have been ≤ 1.6 mrem—less than 2% of the RPG. The total population dose equivalent attributable to LBL operations during 1987 was ≤ 13 man-rem, an average of about 0.003% of the RPG of 100 mrem maximum effective dose equivalent to individual members of the surrounding population. CEDE is defined as the sum of the "doses" delivered to all individuals within an 80-km (50-mi) radius of the Laboratory.

Small amounts of ^{14}C , ^{35}S , ^{125}Xe , ^{125}I , ^{131}I , and unidentified alpha and beta-gamma emitters were released from LBL laboratory stacks. The collective effective population dose equivalent attributable to the foregoing releases is ~ 0.1 man-rem. The majority of the impact of LBL radionuclide operations is from the airborne release of 310 Ci of tritium (as HTO), which is responsible for a CEDE of approximately ≤ 10 man-rem, and a hypothetical maximum off-site individual exposure (from airborne radionuclides) of ≤ 1.6 mrem.

To put the Laboratory's impact into perspective, an approximate value for absorbed dose from external and internal natural sources (e.g., cosmic rays, radiation from continental rocks, naturally occurring radioactive potassium-40 in the muscles and bones) to each person within 80 km (50 mi) of LBL is roughly 0.1 rem/yr, which produces a natural annual population dose of $\sim 510,000$ man-rem.

Gross data for radioactivity in air and water for the period 1978–1987 are presented for comparison with the 1987 data. These gross data show that, except for periods following atmospheric nuclear weapons tests (China, 1980) and the Chernobyl fire (1986), gross radioactivity concentrations in air and water in the vicinity of LBL show only small fluctuations about background levels.

1987 ENVIRONMENTAL ACTIVITIES AND PERMITS ISSUED

Pursuant to LBL's long-term development plan, two environmental assessments (EAs) were completed in 1987 by the Laboratory's Plant Engineering Department in consultation with Ira Fink and Associates of Berkeley, California. These assessments have the following titles:

1. Environmental Assessment: 1-2 GeV Synchrotron Radiation Source
2. Lab-Wide Site Development Plan Final Report

Copies of these assessments were presented to the San Francisco Operations Office of DOE and to the UC. The assessments were prepared in compliance with the National Environmental Policy Act (NEPA).

For further information about these assessments contact

Donald G. Eagling
 Plant Manager
 Building 90G
 Lawrence Berkeley Laboratory
 Berkeley, CA 94720

In order to carry on its research, LBL designs and builds much of its required apparatus. These activities require substantial technical support, including the operation of fabrication, assembly, testing, and waste-handling facilities. The Laboratory operates these facilities under a series of environmental permits issued by state and local agencies. A list of these permits by type and issuing agency, with expiration date, is given below.

Environmental Permits

- 1) Source Operating Permits, Bay Area Air Quality Management District, Expire July 1, 1988.
 - Vapor Degreaser, Building 25A
 - Cold Cleaner, Building 46
 - Vapor Degreaser, Building 53
 - Ultrasonic Degreaser, Building 53
 - Machine Shop Tools, Building 53
 - Machine Shop Tools, Building 58
 - Vapor Degreaser, Building 64
 - Machine Shop Tools, Building 70A
 - Sawdust Collector, Building 74
 - Cold Cleaners (2), Building 76
 - Machine Shop Tools, Building 76
 - Sawdust Collector, Building 76
 - Paint Spray Booth, Building 76
 - Gasoline Storage Tank, Building 76
 - Vapor/Spray Degreaser, Building 77
 - Solder/Grinding Hood, Building 77
 - Ultrasonic Degreaser, Building 77
 - Machine Shop Tools (2), Building 77
 - Paint Spray Booth, Building 77
 - Sandblast Exhaust, Building 77
 - Ceramic Machine Shop Tools, Building 77
 - Paint Drying Oven, Building 77
 - Solvent Cleaning, Building 77
 - Metal Rack Saw, Building 79
 - Machine Shop Tools (2), Building 88

- Solder Hood, Building 88
 - Solvent Cleaning, Building 934
- 2) Wastewater Discharge Permit, East Bay Municipal Utility District, Expires June 8, 1988.
 - Plating Shop, Building 25
 - Plating Shop, Building 77
 - 3) Hazardous Waste Facility Permit, California Department of Health Services, Expires November 7, 1988.
 - 4) Storage Tank Registration. California Department of Health Services. Eight underground storage tanks, seven for petroleum fuel, one for waste oil.

Environmental Activities

1) An underground diesel storage tank, which had released an unknown amount of diesel into the soil, was removed. In addition, contaminated soil around and beneath the tank was also removed. Soil samples, taken during the soil removal operation, indicated that the contaminants had been removed to an acceptable level. In order to confirm that any remaining contaminants are not infiltrating into the groundwater, a well was installed below the former location of the tank and groundwater samples will be taken semiannually for at least the next year.

2) The LBL Environmental Impact Report had been accepted by the UC in order to extend its contract with the DOE to manage the Laboratory. This report assessed the potential impacts on the environment if the elements contained in the *Lawrence Berkeley Laboratory Site Development Plan* were to be implemented by continuation of the existing UC/DOE management arrangement. The current contract expired on September 30, 1987. The extension covers the period from October 1, 1987 through September 30, 1992.

3) A grease/oil interceptor had been installed at a steam cleaning pad. The effluent from steam cleaning operations is collected in an underground catch basin and pumped successively through an oil separator and then through a grease separator prior to discharge to the sanitary sewer system. The oil and grease is removed from the separator and properly disposed.

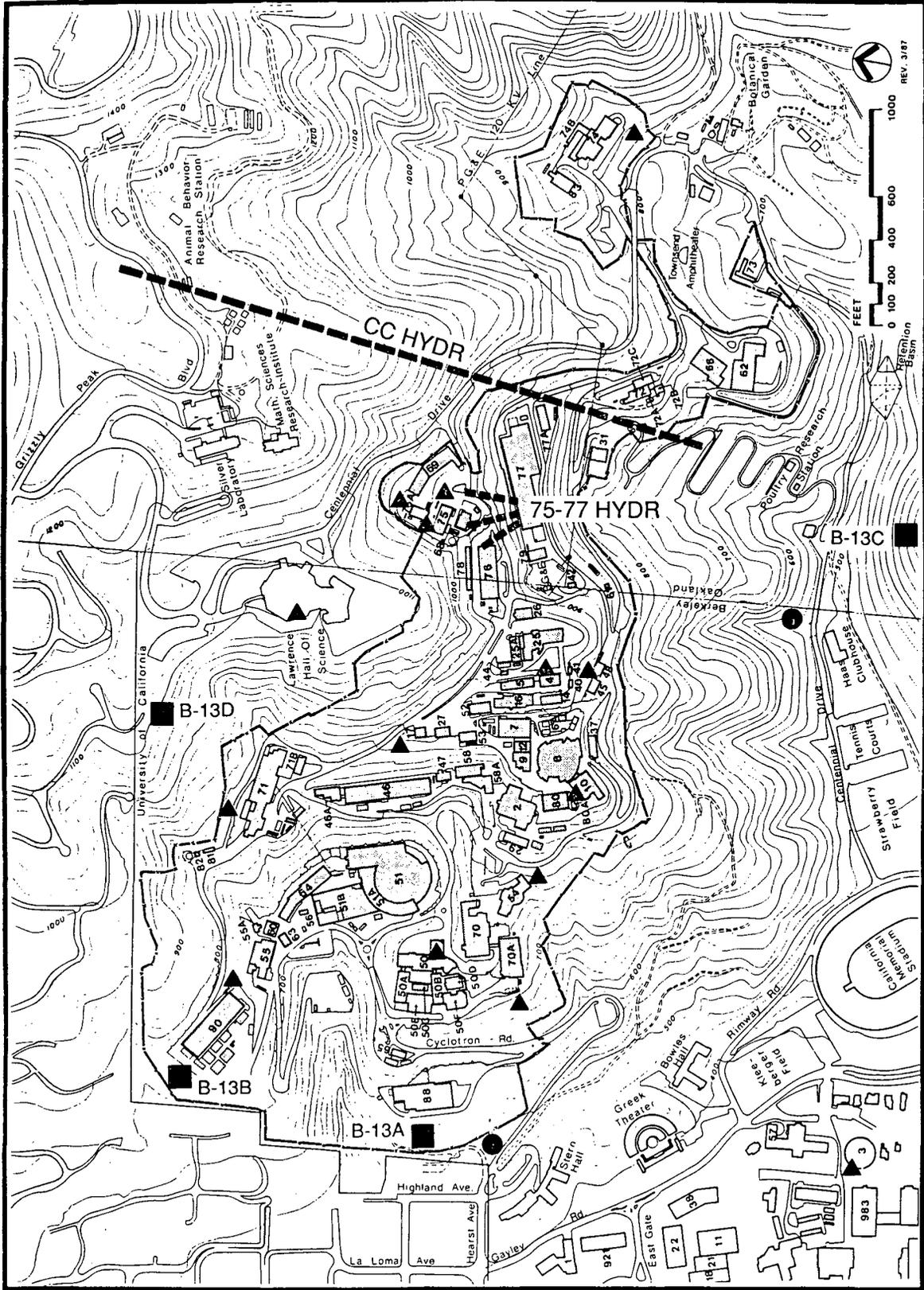
ENVIRONMENTAL MONITORING RESULTS

Radiological Results

Penetrating Radiation

To determine the radiological impact of LBL accelerator operations, we maintain permanent monitoring stations at four points about LBL's perimeter (see Fig. 1 and Table 1).

Each station contains sensitive neutron and gamma pulse counters. The neutron detectors are ~500-cm³ cylindrical BF₃ gas-proportional counters housed in 2.5-inch-thick cylindrical paraffin moderators. The gamma detectors are energy-compensated Geiger-Muller chambers. The output pulses from each of the eight detectors (one of each type is installed at each monitoring station) are prescaled and telemetered to registers in Building 75.⁴ Each LBL accelerator building contains at least one somewhat smaller moderated BF₃ neutron detector, whose output pulses are also prescaled and telemetered to Building 75. By comparing the accelerator neutron monitor output with the output of the perimeter-station neutron monitors, one may assign the perimeter dose equivalent to the accelerator responsible for it. Operational checks of the system are performed daily, and detectors are calibrated semiannually. A typical dose equivalent value for a perimeter-monitoring-station neutron detector corresponds to 0.43 μ rem/pulse. A gamma register-pulse corresponds to about 1.3 μ rem.



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- Perimeter Stations
- Sewer Samplers
- ▲ Environmental Sampling Site

Figure 1. Lawrence Berkeley Laboratory buildings.

KEY TO LBL BUILDINGS SHOWN IN FIGURE 1

Bldg. No.	Description		
HILL-SITE BUILDINGS			
		66	Surface Science & Catalysis Lab
		68	Upper Pump House
		69	Materiel Management & Purchasing
		70	Nuclear Science, Applied Science & Earth Sciences
4	Magnetic Fusion Energy (MFE)	70A	Nuclear Science, Materials & Chemical Sciences & Earth Sciences
5	Magnetic Fusion Energy (MFE)	71	Heavy Ion Linear Accelerator (HILAC)
6	184-Inch Cyclotron	71A	HILAC Rectifier
6A	Utilities Service	71B	HILAC Annex
7	Central Stores & Electronics Shops	72	National Center for Electron Microscopy
9	Magnetic Fusion Energy	72A	High Voltage Electron Microscope (HVEM)
10	Biomedical Research and Photography	72B	Atomic Resolution Microscope (ARM)
12	Central Stores Annex	72C	ARM Support Laboratory
14	Accelerator & Fusion Research & Earth Science	73	Atmospheric Aerosol Research
16	Magnetic Fusion Energy Laboratory	74	Biomedical Laboratory
17	Solar Refrigeration Process & Salvage	74B	Biomedical Laboratory Annex
25	Mechanical Technology	75	Radioisotope Service & National Tritium Facility (NTF)
25A	Electronics Shops	75A	Compactor Processing & Storage Facility
26	Medical Services	76	Craft & Maintenance Shops
27	Cable Shop & High Voltage Test	77	Mechanical Shops
29	Detector & Instrumentation Projects & Biomedical Research & Radiation Effects	77A	UHV Phase I Assembly Facility*
31	Chicken Creek Barn	78	Craft Stores
37	Utilities Service	79	Metal Stores
40	Electronics Warehouse	80	General Research Laboratory
41	Computer Aided Drafting	80A	Telephone Services
42	Earth Science Field Service	81	Liquid Gas Storage
44	Indoor Air Pollution Studies	82	Lower Pump House
45	Fire Apparatus	83	Cell Culture Laboratory
46	Accelerator Development, Electronics Projects & Real Time Systems Group (RTSG)	88	88-Inch Cyclotron
46A	Real Time Systems Group (RTSG)	90	Accounting & Financial Mgmt., Applied Science, Employment, Engineering, Personnel, Protective Services, Superconducting Super Colliding Group (SSC) & TID
47	Advanced Accelerator Study		
48	Fire Station		
50	Physics, Accelerator & Fusion Research & Nuclear Science		
50A	Physics, Director's Office & Earth Science		
50B	Physics & Computer Center		
50C	CAM Division Office & Physics		
50D	MCSD & Nuclear Science		
50E	Earth Sciences	B-4A	Safety Equipment Storage
50F	Information & Computing Sciences Division, Public Information & Patents	B-6B	Deionizer Building
51	Bevalac/Bevatron	B-7A	Radio Shop
51A	Bevatron Experimental Area	B-7B	Office Trailer
51B	External Particle Beam (EPB) Hall	B-7C	Office Trailer
52	Magnetic Fusion Energy Laboratory	B-7E	Office Trailer
53	SuperHILAC Development	B-13A	Environmental Monitoring West of 88
54	Cafeteria	B-13B	Environmental Monitoring West of 90
55	Research Medicine	B-13C	Environmental Monitoring South of UC Recreation Area
55A	Nuclear Magnetic Resonance (NMR)	B-13D	Environmental Monitoring North of 71
56	Cryogenic Facility	B-13E	Sewer Monitoring Station, Hearst Avenue
58	Accelerator Research & Development	B-13F	Sewer Monitoring Station, Strawberry Canyon
58A	Accelerator Research & Development Addition	B-13G	Waste Monitoring Station, West of 70
60	High Bay Laboratory	B-16A	Power Supply House
61	Standby Propane Plant	B-29A	Office Trailer
62	Materials & Chemical Sciences	B-29B	Office Trailer
63	Accelerator & Fusion Research	B-29C	Office Trailer
64	Accelerator & Fusion Research	B-75B	Office Trailer
65	Administrative Data Processing		
			SMALL BUILDINGS AND TRAILERS

Table 1. Location of LBL monitoring stations (MS).

Building No.	Name
B-13A	Building 88 Environmental MS
B-13B	Building 90 Environmental MS
B-13C	Panoramic Environmental MS
B-13D	Olympus Gate Environmental MS

The neutron background attributable to cosmic rays measured at LBL exhibits small fluctuations about a mean value of 3.3 mrem/year.⁵ Table 2 lists the accelerator-produced fence-post dose equivalents measured at each environmental monitoring station during 1987. The fence-post neutron dose equivalent and gamma-ray dose equivalent attributable to LBL accelerator operations in 1987 (see Table 2) are characterized as follows.

1. The 184-Inch Cyclotron produced no dose discernible above background as measured at the Panoramic Environmental Monitoring Station.
2. The SuperHILAC and Bevatron contributed approximately 60% and 40%, respectively, of the fence-post dose equivalent measured at the Olympus Gate Environmental Monitoring Station. The 3.5 mrem was delivered fairly uniformly during the operating year.
3. The 88-Inch Cyclotron fence-post dose equivalent of 2.1 ± 0.4 mrem is primarily attributable to stray neutrons and scattered photons produced during 13 light-ion (helium-3, p^+ , D^+ , helium-4) runs during

Table 2. Fence-post annual effective dose equivalent at the LBL boundary due to accelerator operation, 1987.

Station	1987 total above background		
	gamma (mrem)	n (mrem)	Total ^a (mrem)
Olympus Gate MS	0	3.5 ± 0.7	3.5 ± 0.7
Building 90 MS	0	0.7 ± 0.3	0.7 ± 0.3
Building 88 MS	0.7 ± 0.2	1.4 ± 0.3	2.1 ± 0.4
Panoramic MS	0	≤ 0.7	≤ 0.7
Standard for comparison (Dose to individuals at maximum point of exposure)			100 ^b

^aThe errors shown are those associated with the actual counts and calibration-source uncertainties. Neutron flux-to-dose equivalent conversion factors are not known to this accuracy.

^bSource: Reference 3.

1987. (The fence-post dose equivalent from radionuclide releases at this station was calculated to be 0.6 mrem for 1987.)

The DOE Orders, which provide detailed requirements for radiation protection, under which DOE contractors (LBL, for example) operate, include a table (see Ref. 3) that assigns dose equivalent rate vs. neutron flux density values for neutrons of various energies. In the interest of more accurately reporting the impact of the 88-Inch Cyclotron on LBL's neighbors, measurements of the average energies of the stray neutrons that were produced during the 88-Inch Cyclotron light-ion runs were made at the 88-Inch Environmental Monitoring Station (EMS) in 1985. The measurements⁶ indicated that previously reported values of fence-post dose equivalent were conservatively reported by a factor of more than five. The value of 1.4 ± 0.3 mrem attributable to neutron fluence reported for 1987 reflects less conservative but more realistic neutron energy vs. dose equivalent conversion factors. A neutron spectrometer was installed in the 88 EMS in early 1988 and neutron spectral measurements are ongoing.

4. With the exception of two short periods at the 88-Inch Cyclotron EMS, the continuous gamma measurements telemetered from the four monitoring stations showed no significant correlation with LBL accelerator operation during 1987 and were thus interpreted as constituting the natural gamma background for 1987. The mean value of gamma background inside the monitoring stations was 84 ± 5 mrem for 1987.

LBL's Environmental Health and Safety (EH&S) Department operates a radiological and chemical waste storage yard north and an instrument calibration facility south of Building 75. (The small trailer "complex" on Fig. 1 south of Building 75 is Building 75B, which houses EH&S administrative and operational personnel.)

A recording Geiger-Muller instrument in the southeast corner of Building 75B continuously monitored impact from calibration activities. The instrument recorded a total exposure of 113 ± 7 mrem during 1987 for a net annual effective dose equivalent attributable to calibration activity of 29 ± 7 mrem. A similar instrument located in Building 75A recorded a total exposure of 99 ± 6 mrem for a net exposure of 15 ± 8 mrem.

The 75B instrument is located roughly 10 m from sources of radiation, 70 m from the perimeter fence, 270 m from the nearest commercial (40 hour/wk) occupancy [the Lawrence Hall of Science (LHS)], and 500 m from the nearest home.

The ~30 mrem net exposure at 75B predicts an impact of ~0.7 mrem/yr at the perimeter; < 0.01 mrem/yr (40 hours/wk occupancy) at LHS; and < 0.01 mrem/yr at the nearest home. The sources of radiation monitored by the Building 75A instrument (mainly packaged low-level radioactive waste) are approximately midway between the LBL perimeter fence and the monitor, implying a fence-post dose at that location of ~15 mrem/yr. However, the perimeter fence adjacent to 75A is on UC land and the distances to the nearest occupancies (LHS and private homes) are approximately the same as from the calibration facility discussed above. Thus the impact from this radiation source to the public would be approximately one-half of that from the calibration facility.

LBL has several multicurie gamma irradiators used in radiobiological and radiochemical research. The largest of these units is a ⁶⁰Co unit housed in an interlocked, massive, reinforced concrete-covered labyrinth built as part of LBL's Building 74. (This unit is also the irradiator closest to the LBL perimeter.) Surveys taken when the irradiator was upgraded and reloaded found no area where the stray radiation field exceeded 1 mrem/hr, 1 meter from the outside walls or ceiling. This irradiator is ~80 m from the LBL perimeter fence, 150 m from the nearest "commercial" occupancy (a UCB Botanical Garden building), and more than 700 m from the nearest house. The projected annual dose equivalents to members of the public would be: at the perimeter fence <1.4 mrem/yr; at the Botanical Garden house (40-hr/wk occupancy) <0.1 mrem/yr; and at the nearest house <0.02 mrem/yr (168-hr/wk occupancy).

Airborne Radionuclides

Gross atmospheric particulate beta and alpha activities are measured by air sampling at 14 points: Four perimeter environmental monitoring stations and 10 of the 14 "environmental sampling sites" identified in Fig. 1. (The sites on the north side of Building 75 and the roof of Building 4 are rain collectors. The sites at LHS and west of Building 69 are tritium samplers.) The Building 3 site contains samplers for HTO (tritiated water) and ¹⁴CO₂. Atmospheric air is also sampled for radioiodines at the four perimeter monitoring stations.

The gross beta and alpha sampling media are 10 cm × 23 cm (4 × 9 inch) fiberglass-polyester filters through which air is pumped at 113 l/min (4 ft³/min) at the on-site locations, and 75 l/min (2.7 ft³/min) at the perimeter stations. TEDA-doped activated carbon cartridges are used to sample air for radioiodine at the four perimeter stations. Samples are removed weekly. Before they are counted, they are set aside for five days to enable short-lived radon and thorium daughters (naturally occurring airborne radionuclides) to decay. The filters are loaded into an automatic counter that determines their gross alpha activity by means of a large-area 0.25-mil Mylar window gas proportional counter. Gross beta activity is counted with Geiger-Muller detectors with 30 mg/cm² windows. The detection limit for alpha emitters is 3×10^{-15} μCi/ml. The detection limit for beta emitters is 120×10^{-15} μCi/ml. To ensure accuracy of all counting results, each group of samples counted includes at least one radiation standard sample and a number of background samples. Radioiodines in air, specifically ¹²⁵I and ¹³¹I are assayed by analyzing the activated carbon cartridges with a sodium iodide detector connected to a multichannel analyzer. The detection limits for ¹²⁵I and ¹³¹I are 4×10^{-15} μCi/ml and 12×10^{-15} μCi/ml, respectively.

Alan R. Smith of LBL's low background counting facility (LBCF), located in Bldg. 72, aggregated the 14 weekly environmental particulate air samples into sets and analyzed the sets for airborne particulate gamma-emitting nuclides. The sets were allowed to decay for at least two weeks and then analyzed with a large high-purity germanium detector (HPGE). Each set represented particulates collected from ~14,500 M³ of air, and was counted for a minimum of 1,000 minutes. Aside from very low concentrations of ¹³⁷Cs attributable to atmospheric nuclear weapons testing and the 1986 Chernobyl fire (¹³⁷Cs was found in concentrations of roughly 2×10^{-17} μCi/ml, about 0.000005% of the RPG), the only other gamma-emitters found in the samples were ⁷Be and ²¹⁰Pb. The ⁷Be is produced by cosmic-ray interactions with atmospheric nitrogen (and can also be produced by accelerators). It was found in concentrations ranging from 1.2×10^{-14} to 1.5×10^{-13} μCi/ml and averaged 4.8×10^{-14} μCi/ml, which is 0.001% of the RPG. The detection limit for ⁷Be is 2×10^{-16} μCi/ml for a 1,000 minute count. The concentrations of ²¹⁰Pb, a natural air contaminant, were not computed.

Inasmuch as the DOE Orders³ make no provision for unidentified radionuclides, throughout this report unidentified radionuclides will be conservatively labeled thorium-232 if they are alpha-emitting material or strontium-90 if beta-emitting material. The assertion of conservatism is made because, while ⁹⁰Sr and ²³²Th are found at LBL, they are only in a few LBL laboratories and, for isotopes used at LBL, represent the most restrictive beta and alpha emitters, respectively, listed in Reference 3. Although ²²⁷Ac, which is 4500 times more restrictive a beta emitter than ⁹⁰Sr, is also found at LBL, its most likely state is in equilibrium with its alpha emitting daughters, 18-day ²²⁷Th and 14-day ²²³Ra, and it would thus be detected as an alpha emitter.

Tritium, as HTO, is sampled by passing atmospheric air through a column containing silica gel. Adsorbed water is "exchanged" into distilled water, and an aliquot (5 ml) is placed in a vial and counted in a liquid scintillation counter. The detection limit for HTO in air is 500×10^{-12} μCi/ml.

As with gross alpha and beta samples, silica gel HTO samples are changed weekly. Each of the four perimeter environmental monitoring stations contains a tritium sampler, as does the Building 3 site. The stack from the tritium labeling facility is also monitored for tritium as described above. An additional site, is located at the northeast corner of Building 69A.

The concentration of ¹⁴CO₂ in air is determined by air sampling with NaOH. Samples are changed weekly. Air is bubbled through a jar containing 30 ml of 0.2 M NaOH and thymol blue as a pH indicator. If acid fumes in the sampled air drop the pH of the sample to about 6, a color change results, and the sample is assumed to be invalid (an infrequent occurrence). An aliquot (5 ml) of the NaOH is added to a scintillation cocktail and counted in a liquid scintillation counter. The detection limit for ¹⁴CO₂ is 200×10^{-12} μCi/ml.

The total quantities of radionuclides discharged into the atmosphere are summarized in Table 3. Aside from the tritium release that is four times the 1986 value, the figures are similar to those of last year, and the releases resulted in a small collective effective dose equivalent (see Table 19). One may note that a number of the average values listed in several of the tables in this report (notably Tables 4, 6, 7, 9, 11, and 13) are less than the minimum values listed for individual samples. The foregoing occurs whenever the

actual average value of a substance measured is less than the detection limit for that substance in an individual sample, and the average represents the arithmetic sum of all measurements divided by the number of measurements taken (as in this report). The uncertainties listed with tabular quantities represent 95% confidence limits of the assay values (or sum of assay values).

Although small quantities of radionuclides (Table 3) were discharged into the atmosphere during 1987, the data from the general environmental air sampling were within the range of normal background. The Table 4 data for 1987 may be compared with data from Table 5, which lists LBL perimeter air sample data maxima and averages for the period 1978–1987.

The radioiodine sampling program (Table 6) detected no significant ^{125}I or ^{131}I in perimeter air during 1987. The environmental air sampling program for ^{14}C and ^3H found detectable concentrations of these nuclides (Tables 7 and 8). Essentially, 100% of the tritium released from LBL was discharged from the Building 75 stacks.

All measurements of atmospheric deposition at outlying perimeter stations lie within the range of normal background; however, tritium was detected in rainfall collected within the Laboratory boundary near the stack from the Building 75 Tritium Facility (Tables 9 and 10). The deposition values, adjusted for rainfall, are compared with drinking-water standards (Ref. 3) assuming that all beta activity is ^{90}Sr and all alpha activity is ^{232}Th (conservative assumptions for both). The drinking-water tritium standard³ is used for tritium in Table 9.

Local drinking water is supplied by the East Bay Municipal Utility District (EBMUD) from sources located > 150 km east of LBL. EBMUD uses no well water or local surface water as drinking water.

Waterborne Radionuclides

Rainwater, creek water, and sewage from LBL's two sewer outfalls are analyzed for gross beta and alpha emitters (see Fig. 1; the Strawberry Sanitary Sewer is the southern site, Hearst is the western sewer).

Table 3. Total quantities of radionuclides discharged into the atmosphere, 1987.

Nuclide	Quantity discharged (Ci)
Tritium (as HTO)	310
Xenon-122,125	4×10^{-2}
Carbon-14 (as $^{14}\text{CO}_2$)	1×10^{-2}
Iodine-125	3×10^{-3}
Sulfur-35	6×10^{-5}
Iodine-131	5×10^{-5}
Unidentified beta-gamma emitters ^a	$< 3 \times 10^{-5}$
Unidentified alpha emitters ^b	$< 10^{-6}$

^aConservatively assumed to be ^{90}Sr .

^bConservatively assumed to be ^{232}Th .

Table 4. Summary of air samples, 1987.

	No. of samples	Concentration (10^{-15} $\mu\text{Ci/ml}$)						Average as % of standard	
		Alpha			Beta			Alpha	Beta
		Avg.	Min.	Max. ^a	Avg.	Min.	Max. ^a		
On-site average of 10 locations	497	0.3 ± 0.1	< 2	4 ± 2	11 ± 4	< 110	140 ± 110	4	0.1
<i>Perimeter Stations</i>									
Bldg. 88	48	≤ 0.5	< 3	4 ± 3	≤ 16	< 160	< 160	≤ 7	≤ 0.2
Bldg. 90	48	≤ 0.5	< 3	5 ± 3	21 ± 17	≤ 16	200 ± 160	≤ 7	0.2
Panoramic Way	49	≤ 0.5	< 3	5 ± 3	≤ 16	< 160	≤ 160	≤ 7	≤ 0.2
Olympus Gate	46	≤ 0.5	< 3	4 ± 3	≤ 16	< 160	< 160	≤ 7	≤ 0.2
Standard for Comparison ^b		7			$9,000$				

^aHighest single weekly sample.

^bReference 3: alpha conservatively assumed to be ²³²Th; beta assumed to be ⁹⁰Sr.

Table 5. Annual gross radioactivity found in LBL perimeter air samples, 1978–1987.

Year	No. of Samples	Concentration (10^{-15} $\mu\text{Ci/ml}$)			
		Alpha		Beta	
		Avg.	Max.	Avg.	Max.
1978	198	0.8 ± 0.2	5	60 ± 10	210
1979	202	1.5 ± 0.3	7	28 ± 12	230
1980	204	1.0 ± 0.3	6	28 ± 12	240
1981	195	1.1 ± 0.2	5	120 ± 40	500 ^a
1982	197	0.9 ± 0.2	4 ± 2	14 ± 10	140 ± 100
1983	201	0.49 ± 0.1	2	< 6	110 ± 80
1984	187	0.46 ± 0.1	3 ± 2	< 6	120 ± 100
1985	198	0.54 ± 0.2	4 ± 3	12 ± 6	120 ± 80
1986	195	0.5 ± 0.2	9 ± 3	40 ± 10	700 ± 100 ^b
1987	191	≤ 0.5	5 ± 3	≤ 16	200 ± 160
Standard for comparison ^c		7		9000	

^aThe Peoples Republic of China conducted an atmospheric nuclear test on October 15, 1980. Radionuclides from the test were not detected in LBL air samples until early 1981.

^bChernobyl fire, April 26, 1986.

^cReference 3: alpha conservatively assumed to be ^{232}Th ; beta conservatively assumed to be ^{90}Sr .

Table 6. Summary of radioiodine in perimeter air samples, 1987.

Perimeter Station	No. of samples	Concentration (10^{-15} $\mu\text{Ci/ml}$)						Average as % of standard	
		^{125}I			^{131}I			^{125}I	^{131}I
		Avg.	Min.	Max.	Avg.	Min.	Max.		
Bldg. 88	41	< 1	< 4	≤ 7	< 2	< 12	≤ 20	< 0.0002	< 0.0005
Bldg. 90	42	< 1	< 4	≤ 6	< 2	< 12	≤ 20	< 0.0002	< 0.0005
Panoramic Way	39	≤ 1	< 4	≤ 7	< 2	< 12	≤ 20	< 0.0002	< 0.0005
Olympus Gate	42	≤ 1	< 4	≤ 6	< 2	< 12	≤ 20	≤ 0.0002	< 0.0005
Standard ^a for comparison		5×10^5			4×10^5				

^aReference 3.Table 7. Summary of airborne environmental HTO and $^{14}\text{CO}_2$ sampling, 1987.

	No. of samples	Concentration (10^{-9} $\mu\text{Ci/ml}$)			Average as % of standard ^a
		Avg.	Min.	Max.	
Samples for Tritium as HTO					
<i>On-Site</i>					
ENV 69A	51	1.1 ± 0.5	< 0.5	8 ± 2	1.1
Bldg. 3 roof	51	< 0.2	< 0.5	2.8 ± 0.5	< 0.2
<i>Perimeter</i>					
LHS	50	< 0.2	< 0.5	2.6 ± 0.5	< 0.2
B-13D (Olympus)	47	< 0.2	< 0.5	5 ± 1	< 0.2
Standard for Comparison ^a		100			
Samples for Carbon-14 (as $^{14}\text{CO}_2$)					
<i>On-Site</i>					
Bldg. 3 roof	51	< 0.1	< 0.3	0.4 ± 0.1	< 0.02
Standard for Comparison ^a		500			

^aReference 3.

Table 8. Summary of perimeter airborne environmental HTO and $^{14}\text{CO}_2$ sampling, 1978–1987.

Year	No. of Samples	Concentration (10^{-9} $\mu\text{Ci/ml}$)				
		HTO		No. of Samples	$^{14}\text{CO}_2$	
		Avg.	Max.		Avg.	Max.
1978	101	2.2	9	50	0.12	0.45
1979	101	1	3.4	49	0.026	0.37
1980	103	< 0.2	0.4	52	< 0.07	0.35
1981	100	< 0.2	1.1	50	< 0.06	0.2
1982	102	0.3 ± 0.1	3 ± 1	51	< 0.04	0.3 ± 0.2
1983	101	0.4 ± 0.1	3 ± 1	49	< 0.01	0.3 ± 0.2
1984	97	0.5	7 ± 3	51	0.6	30 ± 10
1985	102	≤ 0.3	5 ± 1	50	≤ 0.1	1.1
1986	100	0.5 ± 0.1	12 ± 3	51	0.07 ± 0.02	0.4 ± 0.1
1987	97	< 0.5	5 ± 1	51	< 0.05	0.4 ± 0.1
Standard for comparison ^a		100			500	

^aReference 3.

Table 9. Summary of atmospheric deposition, 1987.

	Total deposition ($10^{-3} \mu\text{Ci}/\text{m}^2$)						Tritium in rainfall as HTO ^a ($\mu\text{Ci}/\text{m}^2$)		
	No. of samples	Alpha		Beta			No. of samples	Avg.	Max. ^{b,d}
		Avg.	Max. ^b	Avg.	Min.	Max. ^b			
On-Site (9 locations)	108	0.015 ± 0.006	0.18 ± 0.04	0.56 ± 0.06	0.17 ± 0.09	1.2 ± 0.1	122	5.4 ± 1	44 ± 9
Perimeter (4 locations)	48	≤ 0.04	0.06 ± 0.04	0.8 ± 0.5	0.4 ± 0.1	1.6 ± 0.1	24	0.1 ± 0.05	0.2 ± 0.1
Perimeter Averages as a % of Standard		≤ 0.2		≤ 0.2				0.009	
Drinking-water standard $\times 534^c$		27		534				1068	

^aThe on-site tritium-in-rainfall data are computed from samples taken at 11 locations.

^bHighest total for any one site.

^cThe standards used for comparison are derived from Reference 3 for ²³²Th (alpha values) and ⁹⁰Sr (beta values). The deposition represents that quantity of activity found in 534 liters of water (the average quantity of rainfall/ m^2 during 1987). Thus, the values used are 534 times the Reference 3 values. [No standards for comparison have been established, so drinking-water standards (radionuclide concentration/l) are used.]

^dThe location of this deposition collector is on the north side of Bldg. 75. The average HTO concentration in samples taken from the 75 collector was $8.3 \times 10^{-5} \mu\text{Ci}/\text{ml}$ or about 3% of the HTO drinking water standard.

Table 10. LBL perimeter station deposition trends, 1978–1987.

Year	No. of Samples	Rainfall (cm)	Concentration ($10^{-3} \mu\text{Ci}/\text{m}^2$)				$(\mu\text{Ci}/\text{m}^2)$		
			Alpha		Beta		No. of Samples	HTO	
			Avg.	Max.	Avg.	Max.		Avg.	Max.
1978	46	73.6	0.18	< 0.8	11	16	24	1.4	2
1979	47	79.9	0.04	0.1	2.6	5 ± 2	38	0.2	0.4
1980	47	57.3	0.04	0.06	2.5	6	32	< 0.2	< 0.6
1981	48	83.1	< 0.01	0.09	6.9	9.7	36	< 0.1	< 0.2
1982	48	109.0	< 0.01	0.017	1.9	5.2	36	< 0.2	0.3
1983	48	119.4	0.02	0.07	1.6	3.5	36	< 0.2	0.4
1984	48	45.5	0.05	0.08	< 1	3	36	< 0.2	0.2
1985	48	44.5	0.02	0.4	0.7	2	27	< 0.2	0.2
1986	48	81.4	0.03	0.04	0.8 ± 0.2	2	29	0.1	0.3
1987	48	53.4	≤ 0.04	0.06	0.8 ± 0.5	2	24	0.1	0.2

Additionally, sewer effluent is analyzed for gross halogen (radioiodine) content and for tritium. Rainwater is also analyzed for tritium as is the ground water, which flows from the horizontal wells (hydraugers), whose bores are represented by the heavy dashed lines in Fig. 1.

Sewer outfalls are sampled continuously, sample-to-flow ratios are designed to be between 10 and 20 ppm, and composite samples are taken weekly. The five creek sample points indicated in Fig. 2 are sampled weekly. A one-quart grab sample is taken from each site and analyzed for gross alpha and beta emitters.

The four perimeter environmental monitoring stations have 46-cm-diameter (18-in.) cylindrical rainfall collectors on their roofs. During rainy months (generally October through May) rainwater is picked up monthly and analyzed for gross alpha and beta activities and for tritium. During the dry California summer, each collector is rinsed with a quart of tap water, and the rinse is analyzed for "dry deposition." The 10 other atmospheric sampling sites alluded to in the air sampling section of this report also contain 46-cm-diameter (18-in.) combination rain/dry deposition collectors, which are sampled on a monthly basis in the same manner as the four perimeter environmental monitoring stations.

Rain that falls into the collector on the north side of Building 75 is analyzed on a storm-by-storm basis for tritium and gross alpha and beta activities. Tritium analysis of water samples is accomplished by liquid scintillation counting. Water samples are prepared for gross alpha and beta analysis by acidification (HNO_3) and evaporation into 2-inch stainless steel planchettes. Organic residues not wet-ashed by the nitric acid treatment are oxidized by flaming the planchettes.

Since radioiodine is driven out of the water samples when they are acidified, aliquots of the sewer effluent samples are preserved for radioiodine analysis. The iodine contained in the samples is precipitated with silver using stable KI as a carrier. The iodine aliquots are filtered, and the filtrate is processed in the same manner as the acid (HNO_3) samples described earlier. After the filtrate planchette has been flamed, the filter containing any precipitated radioiodine is placed in the planchette and is counted.

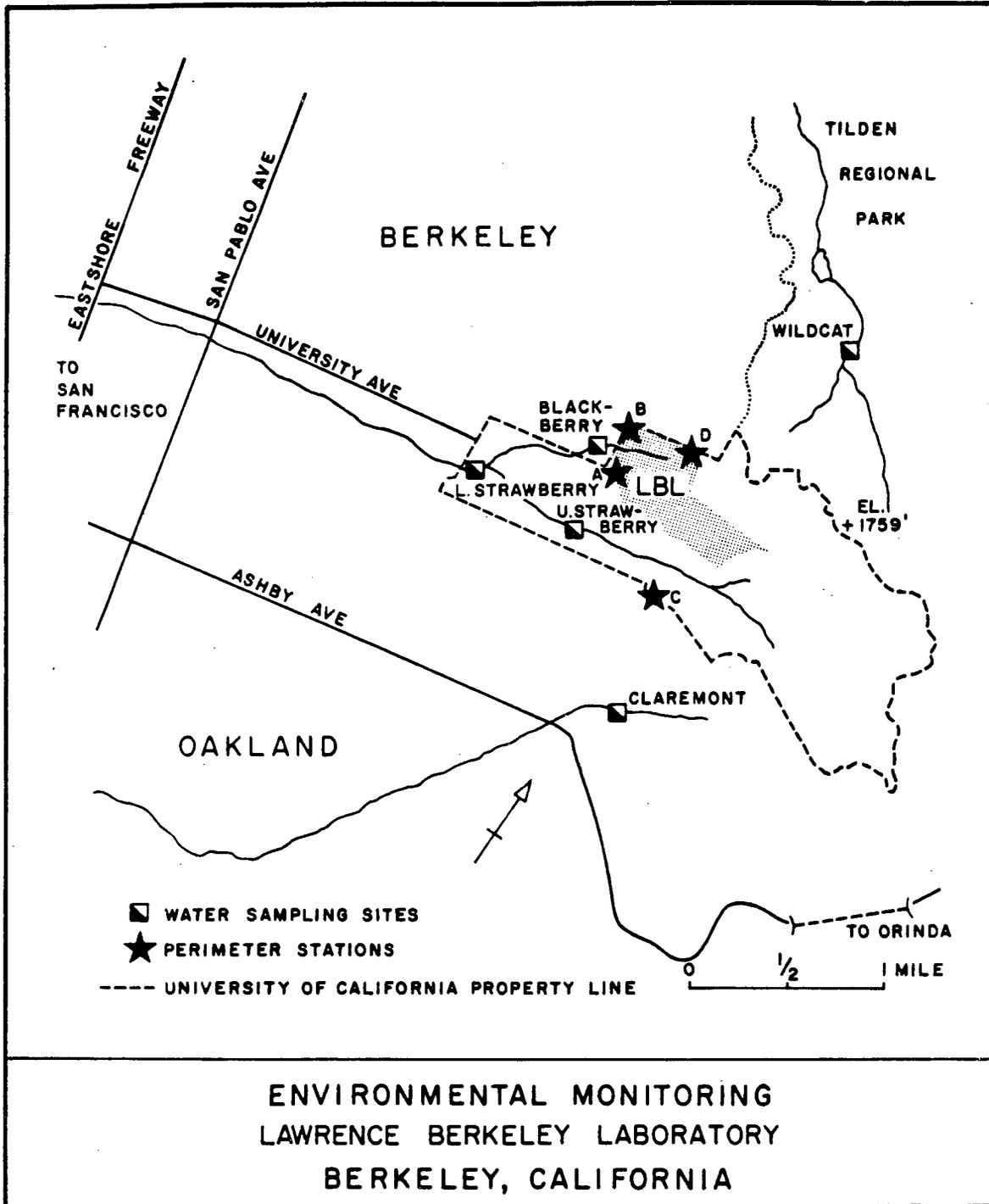


Figure 2. Environmental Monitoring, Lawrence Berkeley Laboratory.

The prepared planchettes are weighed (the tare weight of each planchette is first determined) and counted in a thin-window, low-background gas proportional counter for both gross alpha and beta activities. Since the samples are thick, self-absorption is computed based on areal sample density, which is the sample weight divided by the area of the planchette (20.26 cm²), assuming an alpha energy of 5.2 MeV and a beta energy of 1 MeV.

Table 11 summarizes the 1987 data from the surface-water and tap-water sampling programs. These results are similar to those obtained in past years and all lie within the normal range of background activity. There is no reason to suspect that any of the observed radioactivity originated from LBL. Table 12 summarizes the surface- and drinking-water samples for 1978–1987.

Table 13 summarizes the sewage sampling data for 1987. The average and maximum values listed for sewer beta concentrations reflect the weekly activity found in the hotter of the acid or radioiodine planchettes. Table 14 summarizes the sewage data for the years 1978–1987.

Ground Water

During 1987 the sampling of two of LBL's many hydraugers was started. Samples taken were counted for gross alpha and beta activity and tritium. One hydrauger, designated 75-77 hydr, describes a group of 3 bores, which were drilled horizontally ~200 ft into the earth fill where Buildings 75, 75A, 75B, and 69 were built. The bores are manifolded together and drain north of Bldg. 77 (see Fig. 1 for the approximate "fan out" of the hydraugers' bores). The second hydrauger, which was designated CC hydr, is a ~2500-ft-long horizontal bore from the Chicken Creek access road into Little Grizzly Peak (see Fig. 1). Both hydraugers continued to flow throughout 1987. The "75-77" hydrauger was chosen to be sampled since it drains water from the earth fill that is rained upon by the highest measured tritium-in-rainfall concentration (see Table 9). The "CC" hydrauger is the deepest hydrauger at LBL and is sampled so that the deepest available ground water can be assayed.

Measurable tritium was found in samples taken from 75-77 hydr. Table 15 summarizes the hydrauger (ground-water) sampling data for 1987.

Nonradioactive Pollutants

Waterborne Pollutants

Plating Shop Wastewater Discharges. There are two plating shops at LBL: Building 25 and Building 77. Both shops are subject to the EPA Metal Finishing Pretreatment Standard (40 CFR 433). In general, this standard establishes wastewater discharge limits for cyanide and certain toxic metals. The Categorical Pretreatment Standards have been adopted by EBMUD in Ordinance No. 296.

Wastewater samples are taken from both plating shops to verify compliance with the discharge limits. The samples represent a 24-hour average discharge and are taken before the wastewater combines with wastewater from nonelectroplating operations.

Periodically, EBMUD also obtains samples and reports their results to LBL.

Building 25 Plating Shop

As required by the EBMUD wastewater discharge permit, wastewater samples were taken quarterly from the discharge of the Building 25 treatment unit. The parameters to be monitored were chromium, copper, and lead. In addition, EBMUD collected three samples throughout the year and reported their results to LBL.

One of the seven samples taken indicated a discharge violation for lead. The subsequent investigation determined that a malfunctioning pH meter on the treatment unit probably led to this violation. After this meter was cleaned and recalibrated, discharge levels returned to within allowable limits. There were no other discharge violations detected.

Table 11. Summary of surface- and drinking-water samples, 1987.

	No. of samples	Concentration (10^{-9} $\mu\text{Ci/ml}$)						Average as % of standard	
		Alpha			Beta			Alpha	Beta
		Avg.	Min.	Max.	Avg.	Min.	Max.		
<i>On-site streams</i>									
Blackberry	51	≤ 0.2	≤ 0.6	≤ 2	1.9 ± 0.12	0.9 ± 0.7	3.5 ± 0.9	≤ 0.4	0.2
Lower Strawberry	50	0.2 ± 0.1	≤ 0.7	5 ± 3	1.7 ± 0.12	0.8 ± 0.7	13 ± 2	0.4	0.2
Upper Strawberry	51	0.3 ± 0.2	≤ 0.8	7 ± 4	1.7 ± 0.22	≤ 0.8	3.5 ± 1	0.6	0.2
Average		≤ 0.2			1.8 ± 0.1			≤ 0.4	0.2
<i>Off-site streams</i>									
Claremont	51	0.5 ± 0.2	≤ 0.3	< 3	1.9 ± 0.1	≤ 0.8	5 ± 1	1	0.2
Wildcat	51	≤ 0.2	≤ 0.4	< 2	1.1 ± 0.1	≤ 0.8	3 ± 1	≤ 0.4	0.1
Tap Water	51	≤ 0.03	≤ 0.1	≤ 0.4	0.7 ± 0.1	≤ 0.5	1.5 ± 0.7	≤ 0.06	0.07
Standard of Comparison ^a		50			1000				

^aReference 3: alpha assumed to be ^{232}Th ; beta assumed to be ^{90}Sr .

Table 12. Summary of surface- and drinking-water samples, 1978–1987.

Year	Concentration (10^{-9} $\mu\text{Ci/ml}$)											
	Three On-site Streams				Two Off-site Streams				Drinking Water			
	Alpha		Beta		Alpha		Beta		Alpha		Beta	
	Avg.	Max.	Avg.	Max.	Avg.	Max.	Avg.	Max.	Avg.	Max.	Avg.	Max.
1978	<0.1	6	3.5 ± 0.1	17	<0.3	4	1.8 ± 0.1	9	<0.1	0.3	1.3 ± 0.1	3
1979	<0.2	14	3 ± 0.1	27	<0.08	5	1.4 ± 0.1	3	<0.1	---	0.8 ± 0.1	---
1980	<0.2	4	2 ± 0.1	9	<0.3	3	1.2 ± 0.1	4	<0.1	0.5	0.8 ± 0.1	3
1981	<0.2	3	3.1 ± 0.1	45	<0.2	3	1.6 ± 0.1	22	<0.1	0.4	1.0 ± 0.1	---
1982	<0.3	3 ± 2	1.7 ± 0.1	5 ± 1	<0.3	5 ± 3	1.4 ± 0.1	6 ± 1	<0.1	1.1 ± 0.5	0.9 ± 0.1	2.2 ± 1
1983	<0.1	4 ± 2	1.5 ± 0.1	4 ± 1	<0.3	<2	1.2 ± 0.1	4 ± 2	<0.04	1.2 ± 0.5	0.9 ± 0.1	2.3 ± 0.7
1984	<0.13	<2	1.6 ± 0.3	3 ± 1	0.6 ± 0.3	3 ± 2	1	8 ± 1	0.03	0.3	0.9 ± 0.1	7 ± 1
1985	<0.2	<2	2 ± 0.5	25 ± 2	≤ 0.3	≤ 3	1 ± 0.1	5 ± 1	0.06 ± 0.05	≤ 2	0.9 ± 0.1	2 ± 1
1986	<0.2	8 ± 5	2.3 ± 0.1	27 ± 2	0.4 ± 0.3	4 ± 3	1.6 ± 0.1	10 ± 2	0.06 ± 0.04	<0.4	1.1 ± 0.1	6 ± 2
1987	≤ 0.2	7 ± 4	1.7 ± 0.1	13 ± 2	0.4 ± 0.2	≤ 3	1.5 ± 0.2	5 ± 1	<0.03	<0.4	0.7 ± 0.1	1.5 ± 0.7

Table 13a. Summary of sewage sampling data, 1987.

Total quantities discharged	Total volume (10 ⁶ liters)	Alpha (μ Ci)	Beta (mCi)	Tritium (Ci)
Hearst Sewer	140	≤ 14	2 ± 0.4	< 0.04
Strawberry Sewer	120	≤ 15	22 ± 5	0.7 ± 0.2

Table 13b. Summary of sewage sampling data, 1987 (continued).

Net concentrations	No. of samples	Concentration (10 ⁻⁹ μ Ci/ml)						No. of samples	Concentration (10 ⁻⁶ μ Ci/ml)			Average as % of drinking-water standard		
		Alpha			Beta				Tritium			Alpha	Beta	Tritium
		Avg.	Min.	Max.	Avg.	Min.	Max.		Avg.	Min.	Max.	%	%	%
Hearst	44	≤ 0.11	≤ 0.4	< 1.4	11 ± 2	≤ 2.4	80 ± 20	49	< 0.3	≤ 0.5	6 ± 2	≤ 0.2	1	< 0.02
Strawberry	48	≤ 0.12	≤ 0.4	1.2 ± 1.1	180 ± 40	5 ± 3	2200 ± 500	48	5 ± 1	≤ 0.5	20 ± 4	≤ 0.2	20	0.2
Overall	92	≤ 0.08			90 ± 40				3 ± 1			≤ 0.2	9	0.2
Standard for comparison ^a		50			1000				2000					

^aSource: Reference 3.

^bConservatively assumed to be ²³²Th.

^cConservatively assumed to be ⁹⁰Sr.

Note: The standards cited here are for specific radionuclides in *drinking water*, not sewage, and are provided for comparison purposes only.

Table 14. Sanitary-sewer discharge trends, 1978–1987.

Year	No. of Samples	Concentration (10^{-9} $\mu\text{Ci/ml}$)										
		Hearst					Strawberry					
		Total Flow (10^6 l)	Gross alpha		Gross beta		Total Flow (10^6 l)	Gross alpha		Gross beta		
	Avg.	Max.	Avg.	Max.	Avg.	Max.	Avg.	Max.	Avg.	Max.		
1978	50	229	0.9	34	33	824	48	233	0.6	9	14000	92000
1979	43	247	0.2	5	15	25	45	302	0.5	5	2600	14000
1980	48	288	0.4	3	22	220	46	135	0.3	6	180	1000
1981	49	281	< 0.2	1	21	150	43	89	0.5	14	240	2500
1982	42	300	0.05	1.1	20	460 \pm 20	29	180	0.5	17 \pm 12	60	640 \pm 40
1983	49	190	0.06	< 5	9	80 \pm 7	38	140	< 0.4	< 20	60	800 \pm 40
1984	51	170	0.02	< 5	80	1100 \pm 50	39	74	0.02	< 2	70	250 \pm 10
1985	50	160	< 0.2	< 3	15	90 \pm 10	49	120	< 0.2	< 2	140	1600 \pm 30
1986	47	200	< 0.1	1 \pm 0.3	10 \pm 1	50 \pm 10	47	110	< 0.1	1.1 \pm 0.3	400 \pm 10	4200 \pm 700
1987	44	140	\leq 0.1	\leq 1.4	11 \pm 2	80 \pm 20	48	120	\leq 0.1	1.2 \pm 1.1	180 \pm 40	2200 \pm 500

Table 15. Summary of ground-water samples, 1987.

Hydrauger Designation	No. of Samples	Concentration (10^{-9} $\mu\text{Ci/ml}$)						Concentration (10^{-6} $\mu\text{Ci/ml}$)			Average as % of drinking-water standard		
		Alpha			Beta			HTO			Alpha	Beta	Tritium
		Avg.	Min.	Max.	Avg.	Min.	Max.	Avg.	Min.	Max.	%	%	%
75-77 Hydr.	11	≤ 1	< 0.5	≤ 6	2 ± 0.4	< 0.9	4 ± 2	6 ± 1	< 0.9	10 ± 1	2	0.3	0.2
CC Hydr.	5	≤ 0.6	< 0.6	≤ 0.9	≤ 0.5	< 0.9	≤ 0.9	≤ 0.6	< 0.4	< 1	1	≤ 0.05	< 0.03
Drinking water standard ^a		50			1000			2000					

^aReference 3 alpha assumed to be ^{232}Th , beta assumed to be ^{90}Sr .

Table 16. Summary of Building 25 wastewater sample results.^a

	Chromium (ppm)	Copper (ppm)	Lead (ppm)
Minimum:	0.020	1.20	0.180
Maximum:	0.130	2.90	0.810
2 X SD:	0.096	1.45	0.585
Average:	0.063	1.90	0.397
% of Limit:	2.29	56.21	57.5
# > Limit:	0	0	1
Limit:	2.77	3.38	0.69

^aSummary of results from 7 samples.

Table 16 summarizes the results from the samples taken by LBL and EBMUD.

Building 77 Plating Shop

As required by EBMUD, wastewater samples were taken bimonthly from the discharge of the Building 77 treatment unit. Samples were analyzed for cadmium, chromium, copper, lead, nickel, zinc, and cyanide. In addition, EBMUD collected three samples throughout the year and reported their results to LBL. There were no discharge violations detected by LBL or EBMUD.

Table 17 summarizes the results of the samples taken by LBL and EBMUD.

Site Wastewater Discharges

There are two sanitary sewer systems serving LBL: Strawberry Sanitary Sewer and Hearst Sanitary Sewer. Effluent from each sewer system is monitored at the LBL boundary. Sampling is performed to assure compliance with the site discharge limits mandated by the EBMUD Ordinance No. 270. In this case the EBMUD does not require a compliance report from the Laboratory.

At both sites, a series of flow proportioned grab samples were collected and analyzed for a set of regulated heavy metals and oil and grease. No discharge violations were found at either site.

Tables 18a and 18b summarize the analytical results from the Strawberry and Hearst Sanitary Sewer samples, respectively.

POPULATION DOSE RESULTING FROM LBL OPERATIONS

The development of LBL's model used to assess the population dose equivalent attributable to penetrating radiation and airborne radionuclides, respectively, is detailed in Ref. 5. The model used population figures from the 1970 U.S. census.

Table 17. Summary of Building 77 wastewater sample results, 1987.^a

	Cadmium (ppm)	Chromium (ppm)	Copper (ppm)	Lead (ppm)	Nickel (ppm)	Zinc (ppm)	Cyanide Total (ppm)
Minimum:	0.003	0.050	0.170	0.010	0.230	0.052	0.010
Maximum:	0.160	0.330	0.760	0.100	1.100	0.11	0.020
2 X SD:	0.121	0.246	0.426	0.061	0.640	0.039	0.010
Average:	0.052	0.194	0.340	0.042	0.610	0.076	0.014
% of Limit:	7.594	7.004	10.059	6.087	15.327	2.912	1.167
# > Limit:	0	0	0	0	0	0	0
Limit:	0.69	2.77	3.38	0.69	3.98	2.61	1.20

^aSummary of results from 9 samples.Table 18a. Summary of Strawberry Sanitary Sewer sampling results, 1987.^a

	Cadmium (ppm)	Chromium (ppm)	Copper (ppm)	Iron (ppm)	Lead (ppm)	Nickel (ppm)	Silver (ppm)	Zinc (ppm)	Oil & Grease (ppm)
Minimum:	0.003	0.010	0.034	0.310	0.010	0.020	0.003	0.120	2
Maximum:	0.050	0.190	2.60	15.00	0.380	1.000	0.340	1.90	63
2 X SD:	0.023	0.085	1.22	7.676	0.170	0.354	0.133	0.745	26.0
Average:	0.009	0.050	0.57	5.745	0.059	0.129	0.031	0.562	14.7
% of Limit:	0.900	2.51	11.3	5.74	2.97	2.59	3.09	11.2	5.89
# > Limit:	0	0	0	0	0	0	0	0	0
Limit:	1	2	5	100	2	5	1	5	250

^aSummary of results from 36 samples.

Table 18b. Summary of Hearst Sanitary Sewer sampling results, 1987.^a

	Cadmium (ppm)	Chromium (ppm)	Copper (ppm)	Iron (ppm)	Lead (ppm)	Nickel (ppm)	Silver (ppm)	Zinc (ppm)	Oil & Grease (ppm)
Minimum:	0.003	0.010	0.091	0.280	0.010	0.010	0.005	0.130	3
Maximum:	0.040	0.250	3.00	4.90	0.630	0.110	0.320	2.95	40
2 X SD:	0.021	0.118	1.38	2.38	0.27	0.052	0.150	1.47	20.2
Average:	0.007	0.070	0.75	1.48	0.095	0.020	0.063	0.73	14.9
% of Limit:	0.682	3.500	14.972	1.482	4.757	0.403	6.309	14.554	5.974
# > Limit:	0	0	0	0	0	0	0	0	0
Limit:	1	2	5	100	2	5	1	5	250

^aSummary of results from 35 samples.

While the population within 80 km (50 mi) of LBL increased by 13% during the 1970s^{1,7,8} from 4.6 to 5.1 million people, the populations of Berkeley and Oakland, the two cities immediately adjacent to LBL, declined. Recomputing the population dose model with population statistics from the 1980 census produced no significant difference in its impact/insult value.

Accelerator-Produced Radiation

The LBL model developed by Thomas⁵ for determining population dose equivalent from the maximum measured value of perimeter (fence-post) dose assumes that the fence-post rate changes are uncorrelated with fluctuations in population. During 1987 the maximum fence-post dose was measured at the Olympus Gate Monitoring Station and was 3.5 mrem for the year (Table 2). An examination of the time sequence of the telemetered neutron fluence from the Olympus detector indicated that the neutron fluence peaks correlated well with the fluence peaks from the neutron detector located in the Bevatron approximately 40% of the time and with the peaks from the HILAC detector 60% of the time. The Bevatron and the SuperHILAC operated continuously seven days a week during 1987 except for maintenance, a "summer" shutdown from June 1 through October 1, and a year-end shutdown December 23, 1987 to January 6, 1988.

Aside from shutdown periods the modest fence-post dose equivalent was produced with reasonable uniformity throughout the year and does not seriously compromise the Thomas model's assumptions (student populations were low during the summer shutdown). The model's expression relating population dose equivalent M (in man-rem) to maximum measured fence-post dose H_0 (in rem) is

$$M < 10^3 \times H_0 (1.0 - 0.56f), \quad (1)$$

where f = the fraction of the fence-post dose contributed by the 88-Inch Cyclotron and/or the SuperHILAC. Since 40% of the fence-post dose has been assigned to the Bevatron, $f = 0.6$ [in Eq. (1)].

Thus the expression becomes

$$M < 10^3 (1 - 0.34) H_0. \quad (2)$$

Since H_0 was 3.5 mrem (or 0.0035 Rem), the population dose equivalent attributable to LBL accelerator operation during 1987 was

< 2.3 man-rem.

Airborne Radionuclides

The CEDE resulting from airborne releases of radionuclides is listed in Table 19. The US Environmental Protection Agency (EPA) regulations in 40 CFR 61 require that facilities releasing airborne radionuclides compute the impact of such releases using AIRDOSE-EPA or an approved code. In this report, MICROAIRDOSE, a microcomputer version of the AIRDOSE-EPA radionuclide dispersion and dose assessment code, was used (see Ref. 9). This code was used to compute both collective-effective-dose equivalent and the effective dose equivalent to a maximally exposed individual (radionuclides only).

MICROAIRDOSE computes contributions to the doses from inhalation, ingestion, and exposures from surface contamination and immersion. The code requires:

- a) radionuclide release data;
- b) committed dose-equivalent factors for released radionuclides;
- c) site-specific meteorological data;
- d) agricultural parameters;
- e) site-specific food and water source parameters;
- f) radionuclide-independent parameters; and
- g) distribution of the population within 80 km (50 mi) of LBL.

The data were obtained from the following sources:

- a) Table 3 of this report is used.
- b) values are from Ref. 10.
- c) 1960-1964 Oakland Airport five-year average data were used. While it is most desirable to use on-site meteorology data for the "release year" (1987), the US EPA Region IX regional meteorologist (Ref. 11) indicated that the use of the Oakland Airport five-year average data is, for this application, an acceptable second choice.
- d) Default parameters provided with the MICROAIRDOSE code were used from Ref. 12.
- e) Food and water source parameters were compiled by Victor J. Montoya of LBL EH&S Department's Environmental Surveillance Group from data provided by the water boards and agricultural commissioners of the 11 San Francisco Bay Area counties. The average values for foodstuffs and water not collected or grown within 80 km (50 mi) of LBL were found to be as follows: 35% of the drinking water is imported; 95% of the produce and leafy vegetables are imported; 25% of the milk is imported; and 90% of the meat is imported. (Imported food and water are assumed to be uncontaminated.)
- f) Values are from Ref. 13.
- g) The population distribution about LBL was compiled into 16 compass directions of 10 radial sectors each by Winifred B. Cornica of LBL EH&S Environmental Surveillance Group using data in Ref. 8.

The same values for a-d and f and g were used by MICROAIRDOSE to compute the maximally exposed individual, but it was assumed that 100% of all food and water were grown or collected locally. (More than 90% of the exposure from tritium is assigned to intakes of tritium in food and water. Since there are no known drinking water wells in the communities immediately adjacent to LBL, the value of ≤ 1.6 mrem maximum individual dose-equivalent-from-radionuclides should be quite conservative.) Table 20 summarizes the total CEDE due to LBL operations.

Table 19. Collective effective dose equivalent resulting from LBL airborne nuclide releases, 1987.

Nuclide	(man-rem)
H-3	9.9
Xe-125	< 0.001
C-14	0.01
I-125	0.02
S-35	< 0.001
I-131	< 0.001
Unidentified alpha emitters	0.07
Unidentified beta emitters	0.02
Total	10.0

Table 20. Population effective dose equivalent, resulting from LBL operations, 1987.^a

Contributing factor	Population effective dose equivalent (man-rem)
Penetrating radiation from accelerator operations	2.3
Radionuclide release (from Table 19)	10
LBL-produced effective population dose equivalent	< 13

^aFor 1987, the population dose attributable to natural background sources for the population within 80 km (50 mi) of LBL was approximately 5.1×10^6 persons \times 0.1 rem/person-yr = 5.1×10^5 man-rem.

TRENDS—LBL ENVIRONMENTAL IMPACT

Accelerator-Produced Penetrating Radiation

Figures 3–6 show the annual accelerator-produced dose equivalent reported by the four perimeter environmental monitoring stations from the year they were established to date. During the past several years, the LBL accelerators have run heavy ions during a significant fraction of their operating schedules. Successful work in beam development had served to increase beam currents in recent years and had increased the dose equivalent at the Building 88 EMS somewhat. That upward trend was reversed in 1983. The maximum perimeter dose equivalent (Fig. 3) remains a diminishing fraction of the radiation protection guidelines³ reflecting improvements in accelerator beam optics, local shielding, and cave selection.

Airborne and Waterborne Radionuclides

Figure 7 shows the annual releases of tritium (as HTO) from the Building 75 Tritium Facility from 1974 through 1987.

The 310 curies released during routine operations in 1987 is approximately four times of the 1986 releases and is responsible for approximately 80% of the LBL-produced population-dose equivalent from all sources for 1987. The operational personnel of the tritium facility are continuing to investigate all sources of release so that future releases may be minimized. The releases occur during molecular tagging and tritium waste processing. The increased releases during 1987 reflect a very active program compared to the previous year.

With the exception of occasional known releases, the atmospheric sampling program has yielded data over the past few years that are within the range of normal background.

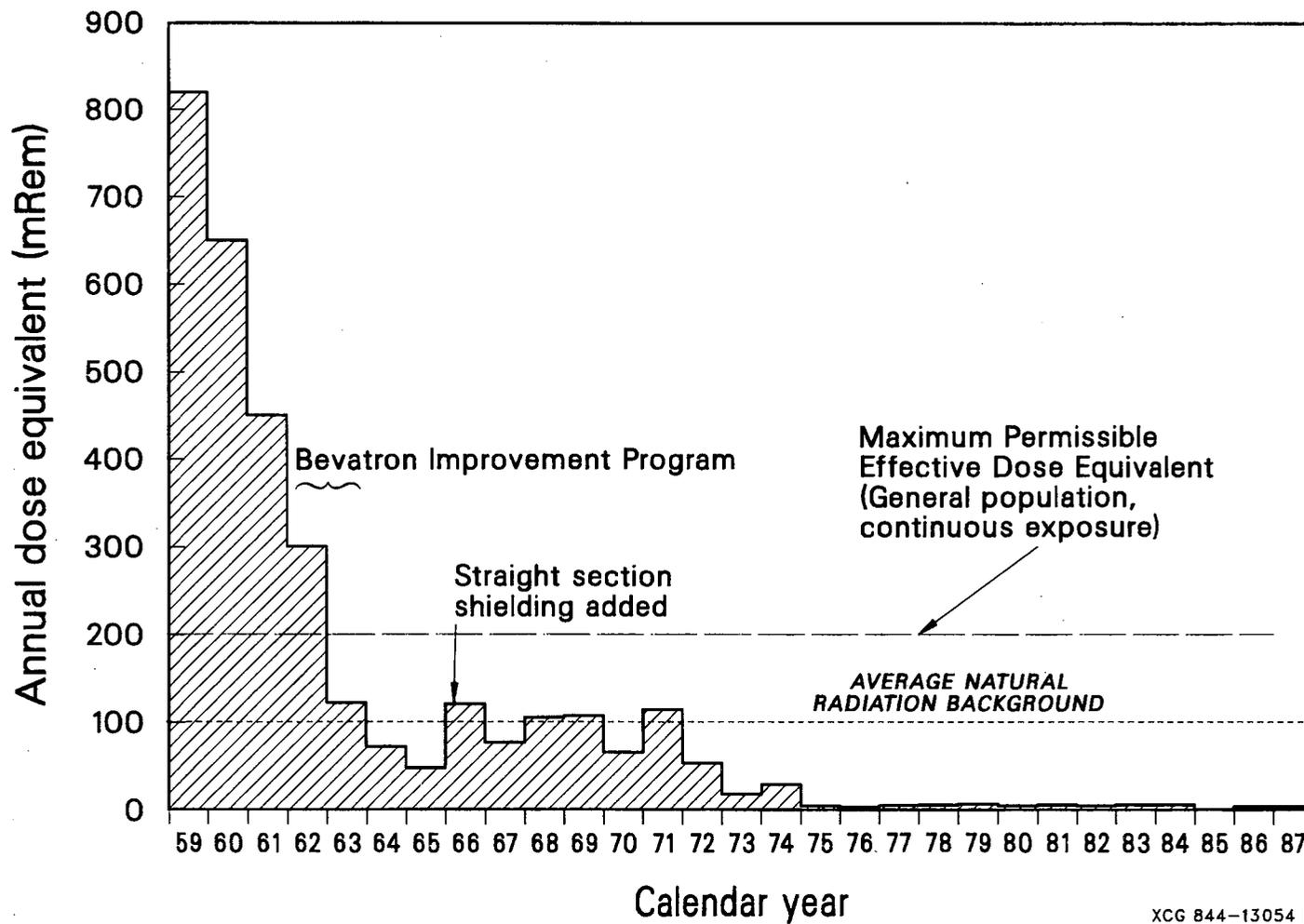
The surface-water program has always yielded results within the range of normal background. Because no substantial changes in the quantities of radionuclides used are anticipated, no changes are expected in these observations.

Under the terms of its license, the UC campus has discharged radionuclides into the Strawberry sewer, complicating the analysis of LBL sewer-sampling data. After 1979 the University discharges were sharply curtailed and are expected to remain so in the future.

QUALITY ASSURANCE

During 1987, in addition to the quality control procedures described in the body of this report, samples that were blind-spiked with tritium were worked up along with each group of environmental samples assayed for HTO.

The LBL Environmental Surveillance Group analyzed DOE's Environmental Measurements Laboratory (EML) QAPXXVI and QAPXXVII Water Samples for tritium and air samples for several gamma emitting nuclides (as reported in References 14 and 15). The results are tabulated in Table 21. The improvement in the gamma emitters-in-air results between May and September reflects increased experience with our group's first high resolution (HPGE) detector.



XCG 844-13054 D
4/15/88

Figure 3. Annual accelerator-produced dose equivalent reported by the Olympus Gate Environmental Monitoring Station, 1959-1987. Maximum Permissible Dose (General Population) is the maximum permissible dose equivalent to any single individual in the general non-Laboratory population. The maximum permissible continuous average effective dose equivalent to the general population is 100 mrem/year (excluding natural radiation background).

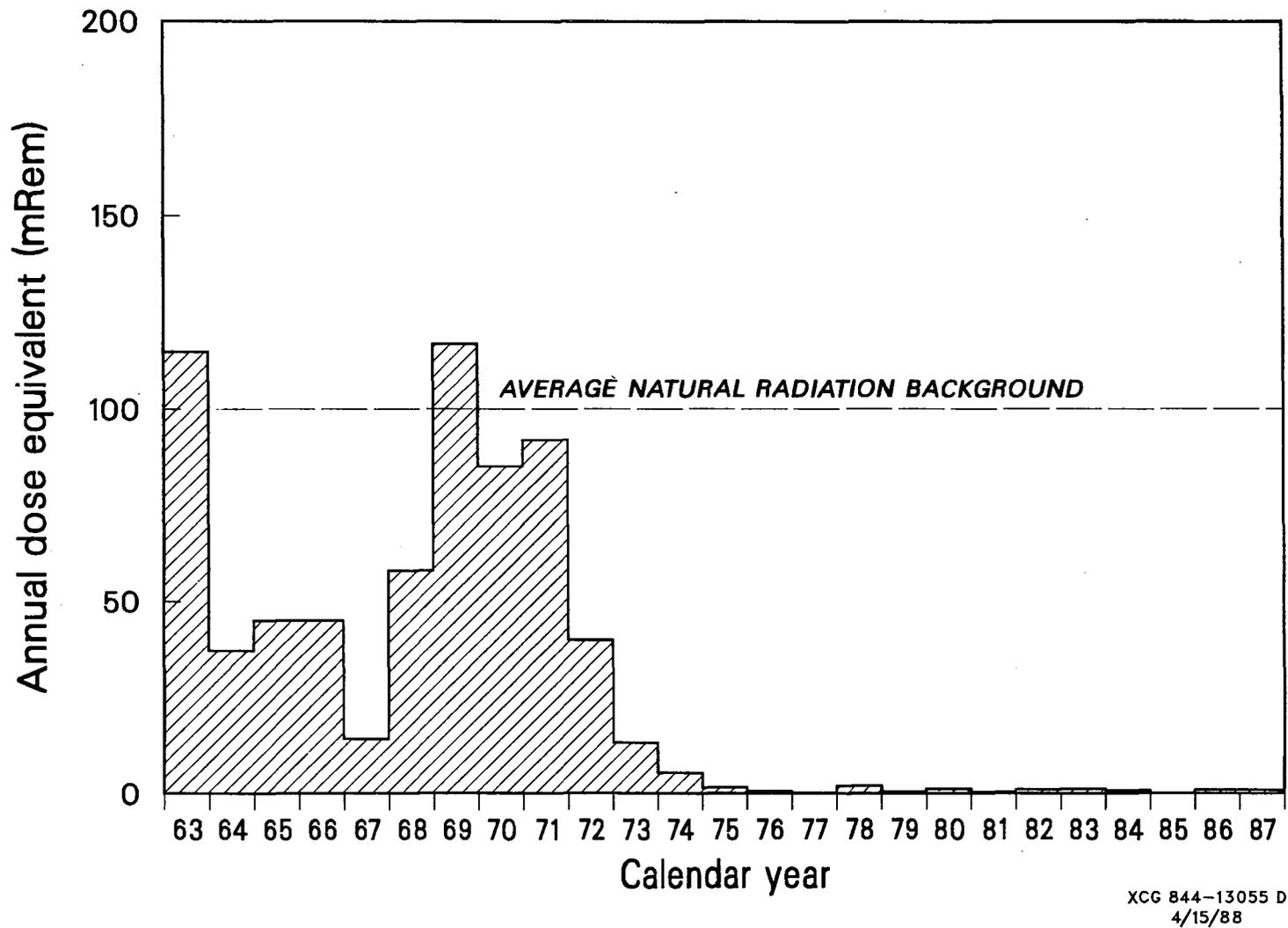


Figure 4. Annual accelerator-produced dose equivalent reported by the Building 90 Environmental Monitoring Station, 1962-1987.

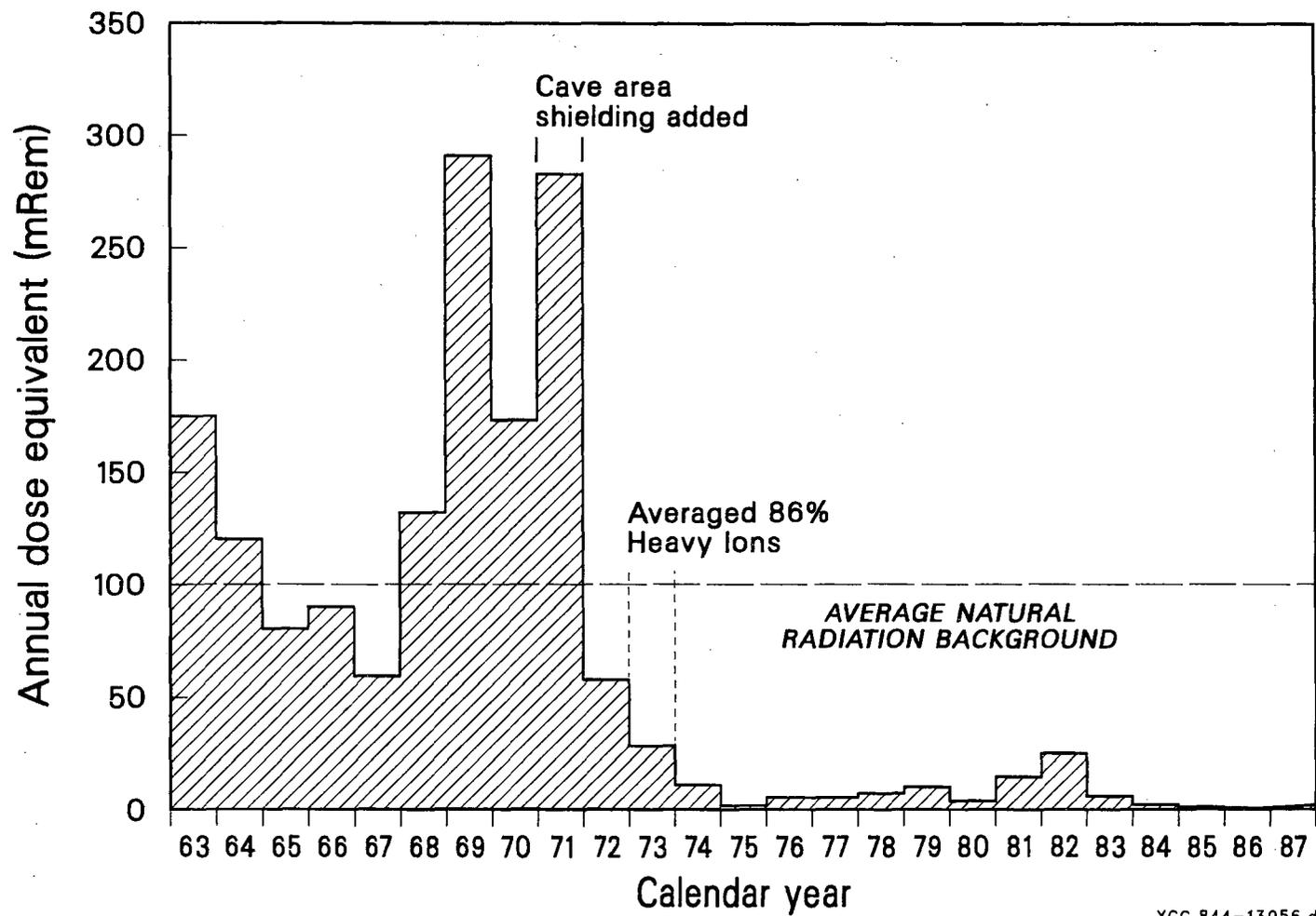


Figure 5. Annual accelerator-produced dose equivalent reported by the 88-Inch Cyclotron Environmental Monitoring Station, 1963-1987.

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4/15/88

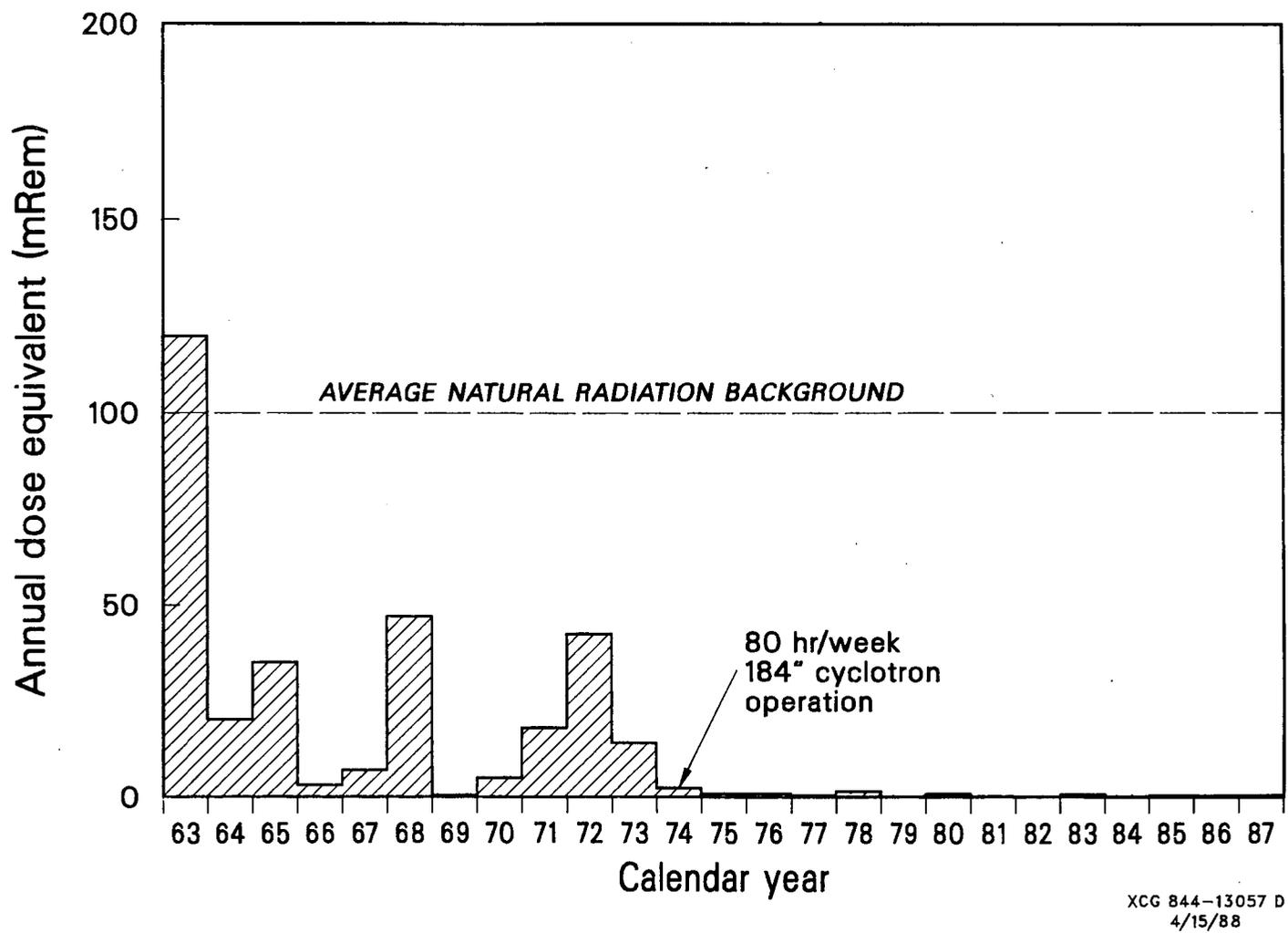
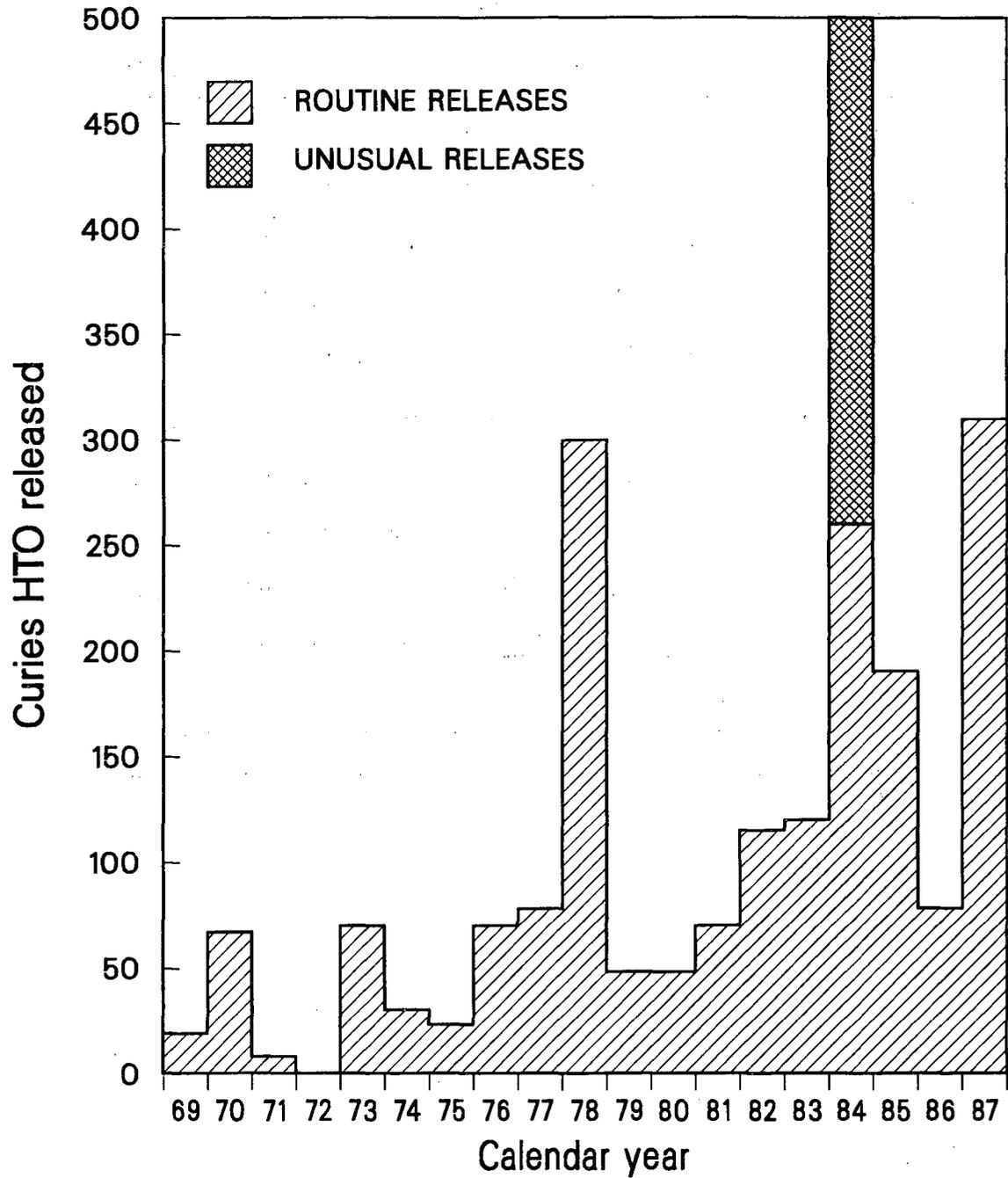


Figure 6. Annual accelerator-produced dose equivalent reported by the Panoramic Way Environmental Monitoring Station, 1963-1987.

Annual HTO Released



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4/15/88

Figure 7. Annual releases of tritium (HTO) from the Building 75 Tritium Facility, 1969-1987.

Table 21. LBL QAP sample results, 1987.

QAP Sample #	Date	Media	Nuclide	Reported LBL Results ^a (± percent)	EML Value	Ratio LBL/EML
XXVI	5/87	Air	BE-7	$6 \times 10^3 \pm 16$	4.64×10^3	1.29
		Air	MN-54	$8 \times 10^2 \pm 20$	4.55×10^2	1.76
		Air	CO-60	$7 \times 10^2 \pm 14$	4.44×10^2	1.58
		Air	CS-137	$5.7 \times 10^2 \pm 19$	4.70×10^2	1.21
		Water	H-3	30 ± 10	33.7	0.89
XXVII	9/87	Air	BE-7	$8 \times 10^2 \pm 50$	8.96×10^2	0.89
		Air	RU-106	$3 \times 10^2 \pm 50$	2.51×10^2	1.20
		Air	SB-125	$1 \times 10^3 \pm 10$	9.63×10^2	1.04
		Air	CS-137	$3 \times 10^2 \pm 13$	2.90×10^2	1.03
		Air	CE-144	$3.2 \times 10^2 \pm 25$	4.06×10^2	0.79
Water	H-3	19 ± 10	19.1	0.99		

^aResults for water are in Pci/ml; results for air are in Pci/sample. Reported errors are in percent.

REFERENCES

1. U.S. Department of Commerce Bureau of the Census, Characteristics of the Population: Number of Inhabitants--California 1980, PC 80 1 AC (March 1982).
2. U.S. Department of Energy, Effluent and Environmental Monitoring Program Requirements, DOE 5484.1, Chapter III (1981) and (1985).
3. U.S. Department of Energy, Requirements for Radiation Protection, DOE 5480.XX, Attachment 1, Table 1 [Concentration Guides for the Protection of the Public] (1987 in draft).
4. Dakin, H.S. and Stephens, L.D., Environmental Radiation Telemetry System, Lawrence Radiation Laboratory report UCRL-16482 (1967).
5. Thomas, R.H. (ed.), The Environmental Surveillance Program of the Lawrence Berkeley Laboratory, Lawrence Berkeley Laboratory report LBL-4678 (1976).
6. Greenhouse, N.A., private communication.
7. University of California Systemwide News, UC Headcount Environments (October 31, 1983).
8. U.S. Department of Commerce Bureau of the Census, Census Tracts--San Francisco-Oakland, California (et al.) Standard Metropolitan Statistical Area (SMSA), PHC 80 2 321 (July 1983).
9. "MICROAIRDOSE" Radiological Assessments Corporation, Neeses, SC, copyright 1987.
10. Corley, J.P. (ed.) Committed Dose Equivalent Tables for US Department of Energy Population Dose Calculations, prepared for the US Department of Energy, Office of Operational Safety by Pacific Northwest Laboratory, Richland, WA, DOE/EH, 1985.
11. Vimont, John, private communication (March 1988).
12. Hoffman, F.O. and Baes, C.F., II (eds), A Statistical Analysis of Selected Parameters for Prediction Food Chain Transport and Internal Dose of Radionuclides. Final Report. ORNL/NUREG/TM-282, 1979.
13. Ng, Y.C., et al., Prediction of the Maximum Dosage to Man from the Fallout of Nuclear Devices, UCRL-50163 (1968).
14. Sanderson, C.G. and Feiner, M.S., Semi-Annual Department of Energy Quality Assessment Program Report, Environmental Measurements Laboratory EML-498, September 1, 1987.
15. Sanderson, C.G. and Feiner, M.S., Semi-Annual Department of Energy Quality Assessment Program Report, Environmental Measurements Laboratory EML-503, January 4, 1988.

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