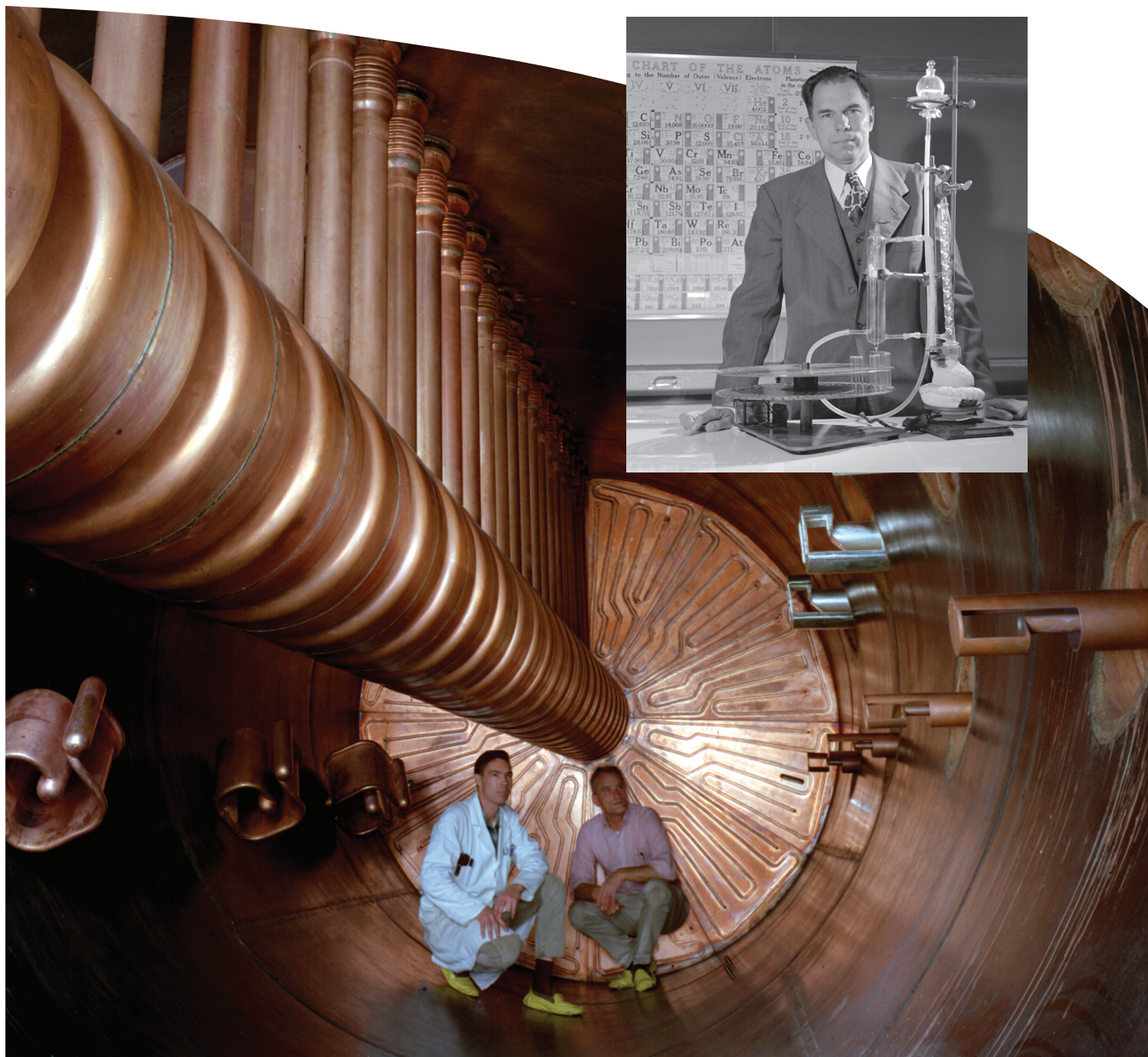




ACS
Chemistry for Life®

National Historic Chemical Landmarks

Chemists and Chemistry that Transformed Our Lives



Discovery of Transuranium Elements at Berkeley Lab

LAWRENCE BERKELEY NATIONAL LABORATORY

American Chemical Society

“A century ago, the periodic table looked much different than it does today. It had empty spots for elements that had not yet been found, and ended at uranium (element 92), the heaviest known element until 1940. But scientists were dreaming about artificially creating even heavier elements.”

*— From “16 Elements: Berkeley Lab’s Contributions to the Periodic Table,”
by Julie Chao and Glenn Roberts Jr., Lawrence Berkeley National Laboratory*



The number of protons in a nucleus determines an element’s atomic number. For instance, uranium has 92 protons and therefore its atomic number is 92. In addition, an element may have different isotopes, which contain different numbers of neutrons. The isotope uranium-238, for example, contains 92 protons and 146 neutrons.

The quest to understand what comprises the world around us dates back to ancient times. As early as the fourth century BCE, the Greek philosopher Aristotle proposed that the physical universe consisted of varying combinations of four “elements”—earth, water, air, and fire. Over the next few hundred years, practitioners isolated and used elements that meet our modern definition—they were fundamental substances consisting of one type of atom that singly or in combination constitute all matter.

Some of these elements, like gold, silver, and tin, were found in nature in relatively pure form; others, such as lead, mercury, and sulfur, had to be isolated from their ores. The 18th-century development of experimental science allowed rapid discovery of more new elements. But uranium, identified in 1789, remained the heaviest known chemical element for more than 150 years.

NEW ELEMENTS AT BERKELEY

In the mid-1930s, a new breed of nuclear scientists, made up of chemists and physicists, became intrigued with the possibility of synthesizing new elements not found in nature. Their dream was finally realized in 1937 when Italian mineralogist Carlo Perrier and physicist Emilio Segrè discovered technetium. Since then, several other elements were created or discovered for the first time at Lawrence Berkeley National Laboratory (Berkeley Lab) in Berkeley, California. This body of work has contributed to a better understanding of the structure of the atom’s nucleus and the nature of matter.

Forming new elements involves changing the nuclei of known

atoms by fusing them with other nuclei or with neutrons. Since nuclei contain positively charged protons as well as charge-free neutrons, fusing one nucleus with another requires overcoming the tremendous repulsion between the two positively charged nuclei. The forces required are millions of times greater than those involved in, say, the explosion of TNT.

Devices called particle accelerators have been used to provide energetic beams of various charged particles to produce the desired nuclear reactions with suitable targets. Accelerators can be linear, in which the beam of particles is accelerated in a straight line, or circular, as in the cyclotron invented by the American physicist Ernest O. Lawrence (1901–1958). Both accelerator types have been used in the discovery of elements at Berkeley Lab.

Synthesis of new elements at the lab began with the creation of neptunium (atomic number 93), the first element beyond uranium in the periodic table, by Edwin McMillan (1901–1991) and Philip Abelson (1913–2004) in 1940. Their work involved irradiating a uranium target with neutrons and was conducted at the Radiation Laboratory at the University of California, Berkeley (predecessor to Berkeley Lab).

GLENN SEABORG AND THE HEAVY ELEMENTS

Later, protons or deuterons (nuclei of hydrogen or deuterium atoms), alpha particles (nuclei of helium atoms), and heavier particles were used as projectiles. One outcome of this effort was plutonium (94), which was created in 1940 by bombarding uranium with deuterons—work conducted by a team led by Glenn Seaborg (1912–1999).

Seaborg was a promising nuclear chemist whose creativity in studying radioactive isotopes caught the attention of leaders of the Manhattan Project, an effort for nuclear weapons development during World War II. Seaborg moved temporarily to work with the Metallurgical Laboratory at the University of Chicago for this work in the early 1940s. While at Chicago, he continued his work to discover new elements with collaborators from Berkeley Lab, resulting in the discovery of americium (95) and curium (96) in 1944. Seaborg’s “actinide hypothesis,” one of his major contributions to chemistry, proposed the organization of the actinide series (atomic numbers 89–103) under the lanthanides (atomic numbers 57–71) and resulted in the configuration that the periodic table shows today.

When Seaborg and his research group returned to Berkeley Lab after the war, they soon developed new methods to form and detect radioactive elements and used them in the discoveries of berkelium (97) in late 1949 and californium (98) in early 1950. Isolation and identification of these elements required chemical separations, a particularly difficult problem because their chemistry was completely unknown.

In November 1952, the first thermonuclear device, known as the hydrogen bomb or H-bomb, was detonated in the South Pacific by Los Alamos Scientific Laboratory. Much to everyone’s surprise, analyses of the debris conducted by Berkeley Lab showed that two new elements, later named einsteinium (99) and fermium (100), had been produced. The huge, 10-megaton blast had



The element 104 discovery team in 1969: Matti Nurmia (from left), James Harris, Kari Eskola, Pirkko Eskola, and Albert Ghiorso.

created an enormous and nearly instantaneous neutron flux that resulted in the capture of at least 17 neutrons by uranium-238.

Within a couple of years, einsteinium and fermium were also produced in high-flux neutron reactors, but it soon became disappointingly clear that the neutron-capture path could not create elements beyond atomic number 100. Attention turned to using light-ion bombardments to add the necessary numbers of protons. Even with new technologies coming to the rescue, it took a few years to design and build linear accelerators and cyclotrons to accelerate the heavier projectiles.

HEAVIER ELEMENTS, ONE ATOM AT A TIME

In 1955, mendelevium (101) was formed by bombardment of einsteinium-253 with a beam of helium-4 ions (alpha particles). The successful identification of mendelevium was performed using separation by a recoil method proposed by Berkeley Lab's Albert Ghiorso (1915–2010). This method took advantage of the feeble recoil imparted in the fusion reaction of helium with the highly radioactive einsteinium target. Recoil kicked the mendelevium atoms out of the thin target onto a gold foil catcher. Chemical processing then proved that a new element had indeed been produced. Seventeen atoms in all were detected. This new separation technique was a powerful tool that would be used for subsequent new element experiments. Mendelevium

was the first element identified on an "atom-at-a-time" basis and the heaviest element to be first identified by chemical separation.

With the completion of the heavy ion linear accelerator (HILAC) at Berkeley Lab in 1957, a double-recoil method was put to work to identify nobelium (102). Nobelium atoms, recoiling from a curium target bombarded by carbon-12 ions, were stopped in helium gas and deposited onto a moving conveyor belt that carried them underneath a negatively charged collector. When the nobelium atoms alpha-decayed on the belt, the resulting fermium "daughter" atoms were kicked off the surface by the recoil from the alpha particles. These atoms were picked up by the collector and shown to behave chemically like fermium. This was the first use of the mother-daughter relationship to prove the atomic number of a new element. Although it was successful, the double-recoil method only worked if the isotopes' half-lives were suitably long. Faster methods were needed to measure the activities of less stable isotopes.

The first important improvement came with the invention of solid-state detectors to measure the energies of the various alpha emitters. In the case of lawrencium (103), first produced and identified at the HILAC in 1961, the recoiling atoms were deposited into a metallized Mylar tape, which was then moved past a series of solid-state detectors for measurement

of the short-lived alpha activity of the lawrencium-258 nuclei. Another equally important development was a gas jet system that transported the activities outside the target chamber where they could be viewed by the new detectors. Many variants of these quick and efficient methods were developed over time. With these new tools it became possible to produce and identify still heavier and shorter-lived elements.

The Berkeley Lab group gradually developed a new apparatus called the vertical wheel. It was used at the HILAC in 1969 to perform the first positive identification of rutherfordium (104) by measuring the decay of its isotopes 257 and 259. Dubnium (105) was first positively identified in 1970 using the vertical wheel to measure decay of dubnium daughters.

The vertical wheel reached its ultimate capability in 1974 in the element 106 discovery experiment by a Berkeley-Livermore group. In the experiment, the relationships—mother, daughter, and granddaughter—of isotope 263 of the new element 106 and its known descendants, isotope 259 of element 104 and isotope 255 of element 102, were demonstrated. The proposal of the name seaborgium for this element produced a dramatic worldwide discussion prior to its ultimate acceptance.

In 1964–65, scientists at Berkeley Lab reported calculations predicting an "island of nuclear stability" on the unknown far reaches of the periodic table, where nuclei with half-lives as long as a billion years could exist. This prediction has guided subsequent work in the field of nuclear science and the search for new elements and isotopes of known elements. This area of research continues, with scientists around the world continuing to search for ever-heavier elements, hoping to someday land at the island of nuclear stability.

Discovery of New Chemical Elements

A National Historic Chemical Landmark

The American Chemical Society (ACS) initially honored Lawrence Berkeley National Laboratory's discovery of new chemical elements with a National Historic Chemical Landmark (NHCL) in a ceremony at the lab in Berkeley, California, on March 11, 2000. The achievement was honored again at a rededication ceremony at Lawrence Hall of Science in Berkeley on August 11, 2019, during the International Year of the Periodic Table. The updated commemorative plaque reads:

Between 1940 and 1974, teams of scientists working at the site now known as Lawrence Berkeley National Laboratory discovered more than a dozen new elements beyond element 92 (uranium). These researchers developed theories and techniques to expand the periodic table by creating and identifying heavy elements with ever higher atomic numbers, including plutonium, berkelium, californium, and lawrencium. This series of achievements culminated in 1974 with the creation of element 106, which was named seaborgium to honor Nobel Laureate Glenn Seaborg (1912-99), who played a key role in many of these discoveries at the laboratory.

About the National Historic Chemical Landmarks Program

ACS established the NHCL program in 1992 to enhance public appreciation for the contributions of the chemical sciences to modern life in the U.S. and to encourage a sense of pride in their practitioners. The program recognizes seminal achievements in the chemical sciences, records their histories and provides information and resources about NHCL achievements. Prospective subjects are nominated by ACS local sections, divisions, or committees, reviewed by the ACS NHCL Subcommittee, and approved by the ACS Board Committee on Public Affairs and Public Relations.

ACS, the world's largest scientific society, is a not-for-profit organization chartered by the U.S. Congress. ACS is a global leader in providing access to chemistry-related information and research through its multiple databases, peer-reviewed journals, and scientific conferences. Its main offices are in Washington, D.C., and Columbus, Ohio.

Acknowledgments:

ACS External Affairs & Communications produced this booklet and wishes to thank contributors and reviewers, especially those at Lawrence Berkeley National Laboratory; the University of California, Berkeley; Lawrence Hall of Science; the ACS California Section; and the ACS NHCL Subcommittee.

Cover: Researchers pose inside the post-stripper tank of the heavy ion linear accelerator known as HILAC. Inset: Glenn Seaborg with an ion-exchange column in 1950. Photos courtesy of Lawrence Berkeley National Laboratory.

Designed by Barb Swartz, Design One

Printed by CAS, a division of the American Chemical Society

© 2019 American Chemical Society

2019 American Chemical Society

Bonnie Charpentier, President
Luis Echegoyen, President-Elect
Peter Dorhout, Immediate Past
President

John Adams, Chair, Board of Directors

2000 ACS Division of the History of Chemistry

Stephen Weininger, Chair

2000 ACS California Section

Alexander Mihailovski, Chair

2019 ACS California Section

Patrick Lee, Chair

2000 Lawrence Berkeley National Laboratory (LBNL)

Charles Shank, Director

2019 LBNL

Michael Witherell, Director

2000 Historical Site Committee

Albert Ghiorso, LBNL
Darleane Hoffman, LBNL and the
University of California
Mary Singleton, California ACS Section

2019 Lawrence Hall of Science

Rena Dorph, Director

2000 ACS National Historic Chemical Landmarks (NHCL) Subcommittee

Paul Anderson, Chair

2019 ACS NHCL Subcommittee

Vera Mainz, Chair

American Chemical Society

National Historic Chemical Landmarks Program
External Affairs & Communications
Office of the Secretary and General Counsel
1155 Sixteenth Street, NW
Washington, D.C. 20036
landmarks@acs.org
www.acs.org/landmarks