cross-section had been seen. This effect was so strong and clear that the group immediately sat down and wrote the draft of their paper announcing the discovery of the new particle. The following day Richter and Ting met at a committee meeting and to their mutual astonishment started to tell each other about the interesting physics results they had found.

Burton Richter was born in 1931 in New York. He decided to become a scientist as a child in high school, a physicist as an undergraduate, and a particle physicist after obtaining his PhD from MIT in 1956. Richter, always fascinated by electrons, sought a job at SLAC where he performed experiments demonstrating the range of validity of quantum electrodynamics

down to very small distances. He has long been a proponent of the virtues of electron-electron scattering and was instrumental in the commissioning of the e^+e^- colliding beam machine at SLAC. During this time he designed an e^+e^- storage ring and worked for ten years to create the complex known as SPEAR. His team working on SPEAR began experiments in 1973 culminating in the discovery of the ψ family of particles and the subsequent measurement of their properties.

Samuel Ting was born in 1936 in Michigan but returned to China with his parents when he was two months old. At the age of 20 he returned to America with \$100, very little knowledge of English and a determination to go to university. Within a short time

he had obtained a degree and a PhD from the University of Michigan and decided to become an experimental physicist. After a year at CERN he returned to teach at Columbia University in 1965. At Columbia he proposed an experiment at DESY to check a recent experimental result which appeared to show a violation of quantum electrodynamics. This was accepted, and in 1966 he began a long association with DESY in which he performed experiments extending the range of validity of quantum electrodynamics and determining the properties of vector mesons. His work over the last decade on lepton production culminated in the discovery of the J/ψ by its decay into electron pairs. □

The Nobel prizes (2): Chemistry

Boranes surgery

A Special Correspondent looks at the work of William Lipscomb

IT is no surprise to chemists that Bill Lipscomb should have been awarded the Nobel Prize for his beautiful work on the boron hydrides. The layman may, however, be baffled, wondering what these oddly-named entities might be, and why they should have merited years of study by a leading crystallographer who is no less well known for his more recent work on the structure of enzymes.

The story goes back more than half a century, to the time when the great inorganic chemist Alfred Stock reported that the simplest compound of boron and hydrogen is not BH_3 , as one would expect by analogy with BF_3 , but its dimer B_2H_6 . The trouble with B_2H_6 , and with all the other boron hydrides which he described, was its "electron deficiency". If the molecule had an ethane-like geometry (as the evidence of electron diffraction was held to indicate) it would need 7 electron-pair bonds to hold its atoms together, but only 12 valency electrons were available for the purpose. This ominous crack in orthodox valency theory was papered over by Linus Pauling, who advanced special reasons why the B-H bond should be able to make do with less than 2 electrons, and proposed structures for the higher boron hydrides in line with this idea.

In 1943 Longuet-Higgins and Bell showed that the chemical and spectroscopic evidence on B_2H_6 harmonised less well with an ethane-like structure than with a bridge structure, in which

two of the H atoms are situated in the middle of the molecule. It appeared that B_2H_6 manages to hold together, not by starving all its bonds of electrons but by sharing two electron pairs between the two sides of the central bridge. The bridge structure was confirmed soon afterwards by W. C. Price, and in 1949 it was proposed that one should regard B_2H_6 as containing two "liaisons triatomiques", or "3-centre bonds", neatly describable in terms of localised molecular orbitals. But the higher boron hydrides remained a puzzle.

At this point the crystallographers grasped the initiative, and Kasper, Lucht and Harker led off with a definitive determination of the molecular structure of crystalline $B_{10}H_{12}$. Their results if anything deepened the mystery: the bonding in this molecule clearly involved principles more general than those which had been advanced for B_2H_6 , and it became clear that a full understanding of these principles must await the collection of structural evidence on the other boron hydrides.

It was Lipscomb who took up this challenge. Undaunted by the pathological instability of most boron hydrides and the acute difficulty of "seeing" hydrogen atoms by X-ray diffraction, he and his colleagues completed a sequence of brilliant experimental studies of the hydrides described by Stock, and of others isolated in his own laboratory. A pattern soon

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William Lipscomb

began to emerge which could be interpreted in terms of "3-centre bonds" of various novel kinds, together with "many-centre bonds", for whose possible existence there were solid grounds in molecular orbital theory. Simultaneous theoretical studies in England and by Lipscomb's group predicted the existence of a stable ion $B_{12}H_{12}^{--}$ with icosahedral symmetry.

The preparation of the potassium salt, and the subsequent determination of its structure by Wunderlich and Lipscomb in 1960, set the final seal on the molecular orbital theory of the bonding in these remarkable substances. The fascinating work of Lipscomb and his colleagues is a story of the fruitful interplay between imaginative theory and skilled experiment, leading to a genuine "breakthrough" in our understanding of the forces which bind atoms together. No longer is there a dichotomy between the chemist's "covalent" bond and the "metallic" bond of the physicist: Lipscomb's work showed us that at least one element succeeds, in its hydrides, in forming both kinds of bond within the same molecular species. □