intice constant) about In_{0.91}Te. The In₃Te₄ ippears to form peritectically; when it is melted under a pressure of ~30 kbar, then cooled and pressure released, a mixture of at least the defect N₄Cl-type and In₃Te₄ phases is obtained. It is probable that the In₃Te₄ phase, in this case, contains excess Te.

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The pressure-induced In₃Te₄ phase is a superconductor. The most nearly stoichiometric one has transition at 1·25-1·15°K. The superconductivity of this phase has been discussed in the section on the NaCl-type phases and in Ref. 2.

50ME RESULTS OF PRESSURE TEMPERATURE EXPERIMENTS ON In₂Te₃

The pressure-temperature diagram of In₂Te₃ determined by differential thermal analysis will be the subject of a subsequent paper. In this section we discuss the nature of some of the specimens obtained after being subjected to various pressures and temperatures, then cooled and pressure released. The results on the In₂Te₃ are the most difficult to unravel.

It appears that there exists at high pressures an In2Tc3 phase which is isostructural with the wellknown Bi2Te3, (21) but it cannot be retained metastably as a single phase. It is not a trivial matter to discern the Bi₂Te₃-type phase in the powder patterns, because whenever it is present, the In₃Te₄-type phase is present also and the structures of the two are closely related. Both structures belong to the same space group; both have all atoms on the threefold axis. As shown earlier, In3Te4 has the sequence -In-Te-Te-In-In-Te-Te-In-, dividing the threefold axis body diagonal in approximately sevenths, while the In2Te3 must have the sequence -Te-Te-In-In-Te-Tedividing the threefold axis body diagonal in approximately fifths. The hexagonal a-axes of the two phases are very nearly equal in length and as one would expect the ratio of the hexagonal c-axes (threefold axis body diagonals) of the In2Te3 to In Tea is very nearly 5:7 (see below for lattice constants of the former).

The deduction of the existence of a pressure-induced Bi₂Te₃-type compound was based mainly on the results of 'single crystal' X-ray diffraction photography. In each of two runs, a sizeable crystalline piece of normal form In₂Te₃ was put

into a Teflon cell containing low viscosity silicone oil. One was pressurized at about 32 kbar at 400°C. the other at about 29 kbar and 340°C. What appeared by microscopic examination to be a single crystal was isolated from each. In both cases however, three phases were intimately cocrystallized. One of these phases has not been identified but accounts for 'extra' lines found in the powder photographs of most of the specimens for which 'extra' lines occur. The other two phases were the In₃Te₄ and In₂Te₃ types with their hexagonal a- and c-axes in apparently exact alignment. The In₂Te₃ type phase had hexagonal axes a = 4.28 + 0.01 and $c = 29.5 \pm 0.1$ Å in both cases as measured on Buerger precession camera photographs. The In₃Te₄ phase from the higher pressure and temperature experiment had a = 4.28 ± 0.01 , $c = 40.2 \pm 0.1$ Å; the other had a = 4.28 ± 0.1 , $c = 40.4 \pm 0.1$ Å.

The c-axis lengths of the In₃Te₄ type phases were substantially smaller than found for stoichiometric or near-stoichiometric In₃Te₄. Further, the 00·9 and 00·21 reflections were no longer observable on the precession camera photographs. It is probable that changes in stoichiometry produce vacancies and shifts in atomic positions.

Unfortunately, we have not been able to obtain metastably a single phase nonstoichiometric In₃Te₄ type. However, superconductivity tests also indicate that some of the In₃Te₄ types phases obtained in the various runs on In2Te3 are nonstoichiometric. This conclusion is based on the ideas set forth above on the nonstoichiometric NaCl-type phases. On this basis, excess Te in the In₃Te₄ type phase should result in a decrease of the superconducting transition temperature. For the specimen obtained by pressurizing the In₂Te₃ in the Teflon cell at 32 kbar and 340°C, about 60 per cent was found to be superconducting at 0.96-0.77°K. A specimen that was subjected to 60 kbar and 730°C in a differential thermal analysis experiment was found to contain the NaCl, In3Te4 and In₂Te₃ types. This specimen had two transitions, about 20 per cent of it at 1.25-1.00°K (the NaCltype phase) and about 20 per cent at 0.5-0.36°K, most likely, the In₃Te₄-type phase. On the basis of our ideas, it is considered very unlikely that the In₂Te₃ phase is superconducting.

A diffractometer pattern was taken of In₂Te₃ which had been pressurized at about 29 kbar and