CdAs₂I, phase C and As coexist. Finally at still higher pressures only phase C and As are found.

Resistance measurements along the 415 °C isotherm yielded a break in the curve of resistance versus pressure at 33 kbar. This is the boundary between the region where CdAs₂I is predominant with traces of phase C present, but no As, and the region where CdAs₂I, phase C and As coexist. This point on both isotherms is in good agreement with the previously proposed⁵ CdAs₂I/II phase boundary. As in the case of ZnAs₂ it is certain that what was previously thought to be the CdAs₂I/II boundary, is the present decomposition boundary. An experiment on the same CdAs₂ as was previously used⁵, was made, where the CdAs₂ was contained in a niobium capsule. The results were in perfect agreement with the present results.

Efforts to determine the exact composition of phase C were made, with less success. Mixtures of Cd_2As_2 and As and mixtures of Cd and As did not react to yield pure phase C at all, when the mixtures had the expected CdAs composition. However, when the mixtures had a CdAs₂ composition phase C was obtained, but with excess As present. Raising the temperature above the previously determined melting temperature produced a new quenchable phase D, with a diffraction pattern very rich in diffraction peaks. Table II presents a summary of these efforts from the various starting compositions tried.

X-ray quantitative analysis⁹ techniques proved less consistent than previously, and the results, although indicating a composition close to CdAs, are not considered reliable enough.

The products from certain high pressure experiments were tempered in a similar fashion to those of ZnAs₂. The tempering time was ~ 17 hours and 300 °C was found to be sufficient to produce the atmospheric pressure phases. A schematic representation of the relationships between the phases is presented in Fig. 6.

The diffraction pattern obtained for phase C (CdAs), was indexed using the method of DE WoLFF¹⁰, and yielded an orthorhombic cell with $a_o = 5.993$ Å, $b_o = 7.819$ Å, and $c_o = 8.010$ Å. Systematic absences showed the space group to be Pbca. Details of the crystallographic work will be published⁷.

In an attempt to understand the relative stabilities phases C and D and therefore also phases A and B, the following experiments were made. CdAs₂ was held at 10 kbar, 900 °C for $\sim 1/2$ hour, cooled slowly to ~ 600 °C, and was found to yield a glassy substance upon quenching to ambient. This is in agreement with earlier work at atmospheric pressure¹³. This suggests that phases D and B are the



Fig. 6. Stability relationships between phases obtained from high pressures and high temperatures in the Cd_3As_2 -As system.

products of the high pressure, high temperature region only. In the previous study⁵, an extra set of signals obtained in the high pressure region, were ascribed to the CdAs₂II-As eutectic. From the present results it is known that CdAs₂II does not exist. It is therefore tempting to assume that these signals belong to the CdAs-As eutectic, or that they are related to the melting of phase D, which is created when the melting temperature of phase C is exceeded for any length of time. This would tie in very well with the heating mechanism used to produce the double set of D.T.A. signals⁵. In the previous work it was noted that the strength of the lower signal (related to the amount of material present causing this signal), could be increased by spending more time above the melting temperature of phase C. The lower D.T.A. signal could be removed by lowering the temperature to ~25 °C. To relate this to the present work three results are presented in Table III. These results are in good agreement with the previous work, where only short periods were spent above the melting temperature. However, from the present results it appears that once phase D has been formed for longer periods of time, perhaps thereby destroying any seeding mechanisms present, the reverse transformation from phase D to phase C is not obtained. It is also possible that further change in composition occurs. A similar relationship between phases A and B is thought to exist.

Discussion

From the present work it appears that what was previously⁵ thought to be the melting curves of CdAs₂II and ZnAs₂II are probably the melting curves of CdAs and ZnAs. The phases B and D are thought to be metastable, and probably involve some process in the liquids of CdAs and ZnAs.

The phases ZnAs and CdAs are isostructural⁷ with ZnSb and CdSb¹⁴. A recent paper on the phase diagram of CdSb¹⁵ indicated a falling melting curve for CdSb, followed by a region above ~35 kbar, where CdSb decomposes into Cd and Sb. Certain questions arise from this paper¹⁶. Above ~35 kbar, the rising portion of the melting curve, is attributed to the Cd-Sb eutectic. An extrapolation of this curve to 1 bar, yields $\sim 30-50$ °C for the metastable eutectic temperature. However, an estimate of the metastable eutectic temperature from the composition-temperature phase diagram, yields a value of ~150 °C. Although these values come from long extrapolations, the lack of agreement is noticeable. Furthermore, the SbII/III transformation causes a sharp break in the melting curve of Sb. Correspondingly there should be a slope change in the Cd-Sb eutectic temperature at ~65 kbar where the Cd-SbII to Cd-SbIII transformation occurs. No such slope change is evident. Is this curve related to the Cd-Sb eutectic temperature, or does it belong to some other phase, which although stable at 50 kbar, 350 °C, decomposes at 50 kbar and lower temperatures to a mixture of Cd and Sb? BELASH and PONYATOVSKII¹⁵ noted that four weak diffraction lines could not be indexed as either pure Cd or pure Sb, and they attributed them to minor contaminations. It is possible that these four diffraction lines have a greater significance. In situ X-ray diffraction experiments at 60 kbar, 300 °C are needed.

Does this decomposition of CdSb have a parallel for the present CdAs and ZnAs phases ? If what was previously thought to be the melting curve of CdAs₂II is the melting curve of CdAs, then the parallel is clear. Above ~45 kbar the melting curve of CdAs is already falling, as for CdSb at 1 bar. Will CdAs decompose further at still higher pressures,

Table III. Experiments made on pure CdAs₂ to check the relationship between CdAs and phase D.

| Starting substance | Treatment | Product |
|------------------------|--|---------------------|
| Pure CdAs ₂ | 40 kbar, 650 °C for $\frac{1}{2}$ hour – cool slowly at 40 kbar to ~ 20 °C – hold for $\frac{1}{2}$ hour – heat slowly to 500 °C at 40 kbar – hold for 1 hour – quench to ambient. | Phase D + As |
| Pure CdAs ₂ | 40 kbar, 500 °C for $1/4$ hour – heat to 650 °C at 40 kbar, and hold for 1 minute – quench to ambient. | Phase $D + As$ |
| Pure CdAs ₂ | 40 kbar, 500 °C for $1/4$ hour – heat to 650 °C at 40 kbar, and hold for 1 minute – slow cool at 40 kbar to ~20 °C – hold for $1/2$ hour – heat slowly to 500 °C at 40 kbar, and hold for $3/4$ hour – quench to ambient. | CdAs (phase C) + As |