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Pressure-induced reconstructive phase transition in Cd_3As_2 Monika Gamża¹, Paolo Abrami², Lawrence V. D. Gammond², Jake Ayres², Israel Osmond², Takaki Muramatsu², Robert Armstrong², Hugh Perryman², Dominik Daisenberger³, Sitikantha Das^{2,4}, and Sven Friedemann^{2,*}¹*Jeremiah Horrocks Institute for Mathematics, Physics and Astrophysics, University of Central Lancashire, Preston PR1 2HE, United Kingdom*²*HH Wills Laboratory, University of Bristol, Bristol, BS8 1TL, United Kingdom*³*Diamond Light Source, Harwell Science and Innovation Campus, Didcot, OX11 0DE, United Kingdom*⁴*Department of Physics, Indian Institute of Technology, Kharagpur, 721 302, India*

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Cadmium arsenide (Cd_3As_2) hosts massless Dirac electrons in its ambient-condition tetragonal phase. We report x-ray diffraction and electrical resistivity measurements of Cd_3As_2 upon cycling pressure beyond the critical pressure of the tetragonal phase and back to ambient conditions. We find that, at room temperature, the transition between the low- and high-pressure phases results in large microstrain and reduced crystallite size, both on rising and falling pressure. This leads to nonreversible electronic properties, including self-doping associated with defects and a reduction of the electron mobility by an order of magnitude due to increased scattering. This paper indicates that the structural transformation is sluggish and shows a sizable hysteresis of over 1 GPa. Therefore, we conclude that the transition is first-order reconstructive, with chemical bonds being broken and rearranged in the high-pressure phase. Using the diffraction measurements, we demonstrate that annealing at $\sim 200^\circ\text{C}$ greatly improves the crystallinity of the high-pressure phase. We show that its Bragg peaks can be indexed as a primitive orthorhombic lattice with $a_{\text{HP}} \approx 8.68 \text{ \AA}$, $b_{\text{HP}} \approx 17.15 \text{ \AA}$, and $c_{\text{HP}} \approx 18.58 \text{ \AA}$. The diffraction study indicates that, during the structural transformation, a new phase with another primitive orthorhombic structure may also be stabilized by deviatoric stress, providing an additional venue for tuning the unconventional electronic states in Cd_3As_2 .

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I. INTRODUCTION

Electronic states with topological properties have been observed in several materials in recent years [1]. Among these, Cd_3As_2 is particularly important, as it is stable in air and is easily synthesised via standard methods. The ambient-condition tetragonal phase of Cd_3As_2 hosts gapless electronic states with a linear dispersion relation caused by symmetry-protected band crossings along Γ -Z in the Brillouin zone, as predicted by Wang *et al.* [2] and subsequently observed by angle-resolved photoemission spectroscopy and transport studies [3–6]. The topological protection leads to Fermi arcs on the surface of Cd_3As_2 crystals [7] and quantum Hall states along the edges [8,9]. Recently, topologically protected states have also been observed in thin films of Cd_3As_2 [10–12]. Thus, Cd_3As_2 is currently being explored for technological applications based on the topological properties, e.g., for optical switches, electronic devices operating at frequencies in the terahertz regime, and high-sensitivity photodetectors [13–15]. In addition, Cd_3As_2 is a prime candidate for applications based on the large magnetoresistance, room-temperature chiral charge pumping, thermoelectric properties, and possible topological superconductivity [16–20]. Assessing the stability of the ambient-condition phase of Cd_3As_2 and understanding

conditions from which this phase can be recovered with ultrahigh carrier mobilities after applying strain is of paramount importance for the development of devices based on this topological semimetal.

The ambient-condition phase of Cd_3As_2 (phase I) hosting the topologically protected electronic excitations has a distorted superstructure of the CaF_2 type [21]. Since Cd_3As_2 is Cd deficient compared with the ideal antifluorite stoichiometry, missing two out of eight Cd atoms needed to form a simple cube around the As, the superstructure can be envisioned as consisting of distorted $\text{Cd}_6\Box_2$ cubes (where \Box denotes an empty vertex) that are ordered in a spiraling corkscrew fashion along an axis parallel to the c direction of the tetragonal crystal lattice. The latest single-crystal x-ray diffraction (XRD) study revealed that each corkscrew is surrounded by corkscrews of the opposite chirality, resulting in the centrosymmetric crystal structure with the space group $I4_1/acd$ [21]. As a consequence of the centrosymmetric symmetry, no spin splitting is allowed at the symmetry-protected band crossings, and a bulk Dirac point at the Fermi energy along Γ -Z results, making the ambient-condition phase of Cd_3As_2 a three-dimensional analog of graphene [22]. Breaking various symmetries using tuning parameters such as temperature or pressure, P , may drive this system into other nontrivial quantum states [3,23,24].

With increasing temperature, Cd_3As_2 undergoes three consecutive structural phase transitions. It transforms to a

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Zn_3As_2 -type structure ($P4_2/nbc$; phase II) at $\sim 220^\circ\text{C}$ and to another primitive tetragonal phase of the Zn_3P_2 -type ($P4_2/nmc$; phase III) at $\sim 470^\circ\text{C}$ before entering the CaF_2 -type face-centred cubic (fcc) structure ($Fm\bar{3}m$; phase IV) at $\sim 600^\circ\text{C}$, which remains stable to the melting temperature [25–27]. Crystal structures of the high-temperature phases can be viewed as antifluorite superstructures containing 128, 32, and 8 distorted $\text{Cd}_6\Box_2$ cubes per conventional unit cell, respectively, and the transitions between them are related to changes in distribution of Cd atoms among the tetrahedral voids [21]. In the high-temperature fcc phase, the Cd ions are disordered [26], whereas in all the lower-temperature variants, the Cd atoms form ordered patterns and shift from the ideal antifluorite positions toward the empty vertices of $\text{Cd}_6\Box_2$ cubes. The transition between the Zn_3As_2 - and Zn_3P_2 -type phases at 470°C has all the signatures of a typical first-order structural change; it proceeds quickly and with a notable thermal hysteresis. In contrast, the transition at 220°C from the Zn_3As_2 -type structure to the ambient-temperature phase I progresses slowly and shows no distinct hysteresis [26]. Evidence for an intermediate phase with disordered crystal structure was found at the isomorphic transition in Zn_3As_2 [26].

At high pressures (P), additional phases of Cd_3As_2 have been identified [25,27]. At room temperature, a high-pressure semiconducting phase V was reported with a trigonal, orthorhombic, or monoclinic symmetry [25,28–30] above $P_c \approx 2.3$ GPa; this phase was mapped out to be stable up to 550°C . Above 550°C , three more high-pressure phases were detected but have not yet been characterized. Raman and optical reflectivity measurements indicated the presence of a consecutive phase at pressures above 9.5 and 8 GPa, respectively [31,32] where superconductivity was observed [30].

Although the presence of the Cd_3As_2 phase V at high pressures has been established, the correlation between the structural and electronic characteristics is still under debate. In particular, some studies show that the metal-to-semiconductor transition coincides with the structural phase transition from the ambient-condition phase I to the high-pressure phase V [29], whereas other reports indicate that the onset of semiconducting behavior occurs at much lower pressures (~ 1.1 GPa) compared with the structural phase transition [30]. Furthermore, recent magnetotransport measurements revealed a sudden shrinkage of the Fermi surface and an anomaly in the phase factor coincident with an anisotropic compression of the crystal lattice at $P \approx 1.3$ GPa, interpreted as an indication for opening of a band gap [33]. Those inconsistent findings motivated us to investigate in detail the pressure-driven evolution of the crystal structure and transport properties of Cd_3As_2 between the ambient-condition phase I and the high-pressure phase V by means of high-resolution powder XRD, electrical resistivity, and Hall effect measurements.

II. METHODS

A. Experimental details

Needle-shaped single crystals were grown from self-flux in Cd excess, as described in Ref. [21]. High-purity Cd and As were loaded into an alumina crucible and sealed inside

a quartz ampoule. The materials were heated to 825°C and slowly cooled to 425°C before being centrifuged to remove excess Cd. The single-crystal nature of selected samples was confirmed with XRD. Stoichiometric composition of representative single crystals was confirmed with energy dispersive x-ray spectroscopy. Our samples show a residual resistivity ratio of seven, slightly larger than samples in previous high-pressure studies [29,30,33]. As-grown single crystals displaying clear facets were used for our electrical transport study. For powder XRD measurements, crystals were ground with a slight amount of glycerol to avoid dust.

Electrical resistivity and Hall effect measurements at room temperature were conducted in piston-cylinder cells with glycerol as a pressure transmitting medium. The resistance of a manganin wire was used as a continuous pressure gauge with *in situ* calibration on the resistive transitions of bismuth I-II and II-III [34]. The load on the piston-cylinder cell was regulated with a home-build hydraulic setup enabling slow pressure sweeps with virtually constant sweep rate dP/dt (see inset of Fig. 2). The resistance and Hall effect were measured using a standard 4-point technique with a Stanford-Research SIM921 ac resistance bridge. The Hall effect was measured at fixed pressures in a ± 0.2 T electromagnet, giving rise to a transverse resistance R_t linear in magnetic field B . The Hall coefficient R_H was extracted from the slope of linear fits to $R_t(B)$. Sample dimensions were measured with an optical microscope; the uncertainty of the thickness measurement propagates to a systematic uncertainty of $\pm 15\%$ in R_H .

High-pressure powder XRD measurements were performed in membrane-driven diamond anvil cells (DACs) at the I15 beamline at Diamond Light Source. X-ray radiation with wavelength $\lambda = 0.4246(5)$ Å was used throughout as calibrated on a LaB_6 standard. Pressure was measured using the ruby fluorescence method [35]. Diffraction peaks of residual cadmium flux mixed with Cd_3As_2 were used as a second pressure marker, thus allowing us to assess errors in the measured pressure values, as shown in Fig. 1. Three sample loadings were prepared and are labeled as sample 1, sample 2, and sample 3. For the first two loadings, glycerol was used as a pressure medium, whereas sample 3 was immersed in silicon oil. Temperatures up to 250°C have been accessed with an external heater for samples 2 and 3. Diffraction patterns were acquired on a MAR345 image plate detector and integrated with the DAWN software [36] using the calibration parameters obtained from the LaB_6 standard.

B. Analysis of XRD patterns

Refinements of the diffraction patterns were carried out with Jana2006 [38]. Profile refinements using the Le Bail method were used for evaluating the structural parameters and for performing strain and size analyses. Contributions to the diffraction peak broadening originating in grain size and microstrain were separated based on their different dependencies on the scattering angle θ . To this end, reflections visible in diffraction patterns in the 2θ range from 6° to $\sim 16^\circ$ were fitted using Lorentzian and pseudo-Voigt shape functions [39]. The variations of the Lorentzian and Gaussian half widths with

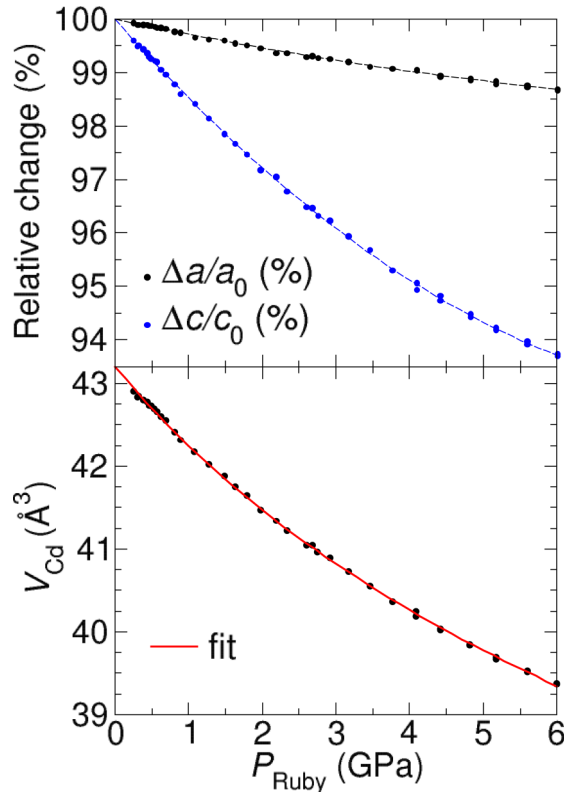


FIG. 1. Unit cell volume of Cd (bottom panel) and relative changes in its lattice parameters (top panel) obtained from Le Bail refinements of powder XRD patterns recorded on sample 1 at room temperature and plotted as a function of pressure measured using ruby fluorescence. Dashed lines are guides to the eye. Solid red line is the fit to the experimental data points using the Murnaghan equation of state with $K_0 = 43$ GPa, $K'_0 = 8$, and $V_0 = 43.2 \text{ \AA}^3$, in agreement with Ref. [37]. Error bars in the pressure values evaluated based on the deviations of the fit from the data points do not exceed 0.05 GPa.

Bragg angle θ are described by the equations

$$H_L = \frac{X}{\cos\theta} + Y \tan\theta + \Gamma_L(\theta), \quad (1)$$

and

$$H_G^2 = 8 \ln 2 \left[\frac{P}{\cos^2\theta} + U \tan^2\theta + \Gamma_G^2(\theta) \right]. \quad (2)$$

In each of Eqs. (1) and (2), the first term accounts for isotropic Scherrer particle broadening, the second term describes microstrain broadening related to lattice defects, while the last term stands for the Lorentzian (Γ_L) and Gaussian (Γ_G) parts of instrumental broadening. The latter was assessed from powder XRD pattern of virtually strainfree LaB_6 powder with controlled grain size in the range of tens of micrometers. In the refinements, only one term describing grain size effect and one term describing microstrain broadening were used throughout.

According to the Scherrer formula [40,41], the volume weighted crystallite size D_V is given by

$$D_V = \frac{K\lambda}{\beta \cos\theta}, \quad (3)$$

where K denotes the Scherrer constant and is assumed to be 1, λ is the wavelength of the radiation used for the XRD study, while β represents integral breadth of a reflection located at 2θ expected, assuming that the broadening originates solely in the grain size effect [42]. Therefore, the volume averaged size of crystallites can be calculated from refined X or P values in degrees using the formulas

$$D_{V,L} = \frac{360K\lambda}{\pi^2 X} \quad \text{and} \quad D_{V,G} = \frac{180K\lambda}{\sqrt{2\pi^3 P}}, \quad (4)$$

for Lorentzian and Gaussian-type broadening attributed to the grain size effect, respectively. The microstrain is given by

$$\epsilon = \frac{\beta}{\tan\theta}. \quad (5)$$

Therefore, the respective Lorentzian and Gaussian contributions are determined as

$$\epsilon_L = \frac{3\pi^2 Y}{1440} \quad \text{and} \quad \epsilon_G = \frac{\sqrt{2\pi^3 U}}{720}. \quad (6)$$

XRD data for the high-pressure phase of Cd_3As_2 was also analyzed by combining a Williamson-Hall method [43] with line profile analysis. To this end, a profile fitting procedure was adopted in which the measured XRD patterns were decomposed into individual peaks described using a pseudo-Voigt or Lorentzian shape function. According to the Williamson-Hall equation:

$$\beta \cos\theta = \frac{K\lambda}{D_V} + 4\epsilon \sin\theta. \quad (7)$$

Therefore, the microstrain ϵ and the volume-weighted crystallite size D_V are calculated from the slope and the intercept of the plot of $\beta \cos\theta$ against $\sin\theta$ obtained after correcting integral breadths of the diffraction peaks for the instrumental broadening, assuming $K = 1$.

Indexing of powder XRD patterns was performed using McMaille [44] and DICVOL06 [45] computer codes. Profile decomposition of the measured XRD patterns gave angular positions and intensities of individual peaks which were used as an input. Searching for candidate crystal lattices with volumes of up to 6000 \AA^3 was carried out in all crystal systems by means of Monte Carlo algorithms [44] or an exhaustive trial-and-error method with variation of parameters by successive dichotomy and partitioning of the unit cell volume [45]. Space groups for the resulting sets of lattice parameters were identified using JANA2006 [38].

III. EXPERIMENTAL RESULTS

We start by presenting continuous resistivity as well as discrete Hall effect measurements in Sec. IIIA, showing the hysteresis and nonreversible behavior of the electronic properties upon pressure cycling of Cd_3As_2 into phase V and back to low pressure. Subsequently, we introduce the XRD results in Sec. IIIB, which demonstrate the reversion to the ambient-pressure tetragonal phase I and show poor crystallinity after the pressure cycling. Together, these measurements provide evidence for a first-order reconstructive phase transition. We evaluate the compressibility of the low- and high-pressure phases of Cd_3As_2 at room temperature. We show that

annealing of the high-pressure phase at $\sim 200^\circ\text{C}$ greatly improves its crystallinity and thus allows us to reveal more details about its crystal lattice. Finally, we indicate the formation of a new primitive orthorhombic phase prompted presumably by deviatoric stress upon releasing pressure through the structural phase transition in Cd_3As_2 .

A. Electrical transport

As a crosscheck, we first measured the resistance of Cd_3As_2 on compression and decompression while staying within phase I at room temperature. Using a pristine single crystal, the pressure was first increased to 2 GPa and subsequently released back down to 0.5 GPa. As shown by gray lines in Fig. 2(a), $\rho(P)$ is fully reversible within phase I and increases gradually with pressure.

Once cycling the pressure through the phase transition into the Cd_3As_2 phase V at room temperature, the resistance of Cd_3As_2 shows large hysteresis and nonreversible behavior. Subsequent to the cycling at low pressures, the pressure has been ramped to 3 GPa and released back down to 0.5 GPa, as shown by the red data in Fig. 2(a). The transition from phase I to phase V is clearly seen as a pronounced steplike increase in the resistance with a sharp onset at $P_{c\uparrow} = 2.3$ GPa. Above 2.6 GPa, the resistance follows again a linear increase with further increasing pressure. The small linear increase below $P_{c\uparrow}$, the jump at $P_{c\uparrow}$, and the steeper linear increase above $P_{c\uparrow}$ are in very good agreement with earlier studies [29,30]. Upon releasing the pressure, the resistance follows this linear regime to well below $P_{c\downarrow}$, indicating a large hysteresis of the phase transition. Only at $P_{c\downarrow} = 1$ GPa, a large rise and sharp cusp is observed, indicating a phase transition. Most notably though, the resistance does not return to the low-pressure value. We note that the linear behavior observed below $P_{c\downarrow}$ after pressure cycling extrapolates to a value more than double the starting value, yet we show below in our analysis of the x-ray measurements that this second transition at $P_{c\downarrow}$ is associated with the reverse transition from phase V to phase I. All characteristics of the pressure-dependent resistance have been reproduced in three samples from the same batch.

Our Hall effect measurements at room temperature indicate a large increase in charge carrier concentration in the high-pressure phase of Cd_3As_2 . The results shown in Fig. 2(b) were obtained upon stepwise increasing the pressure to 3 GPa and subsequently releasing it down to 0.5 GPa at room temperature. While raising the pressure, the Hall coefficient remains roughly constant at $R_H = 4.5 \times 10^{-6} \text{ m}^3 \text{ C}^{-1}$, in qualitative agreement with previous measurements at 2 K [33]. We note that the systematic difference by a factor of 3 and 10 with earlier studies by He *et al.* [6] and Zhang *et al.* [33], respectively, is probably due to uncertainties in the sample thickness and/or different levels of doping while the difference between our room temperature and the previous low-temperature measurements is small [6]. The qualitative agreement continues at high pressures where, above $P_{c\uparrow}$, the Hall coefficient is reduced to $R_H = 1.6 \text{ m}^3 \text{ C}^{-1}$. In a simple single-band model, the drop of R_H at $P_{c\uparrow}$ corresponds to a threefold increase of the charge carrier concentration n in the high-pressure phase. This is similar to the sixfold increase of n observed at 2 K by Zhang *et al.* [33]. Together with the increase in resistance above $P_{c\uparrow}$, this suggests an order of magnitude reduction of the

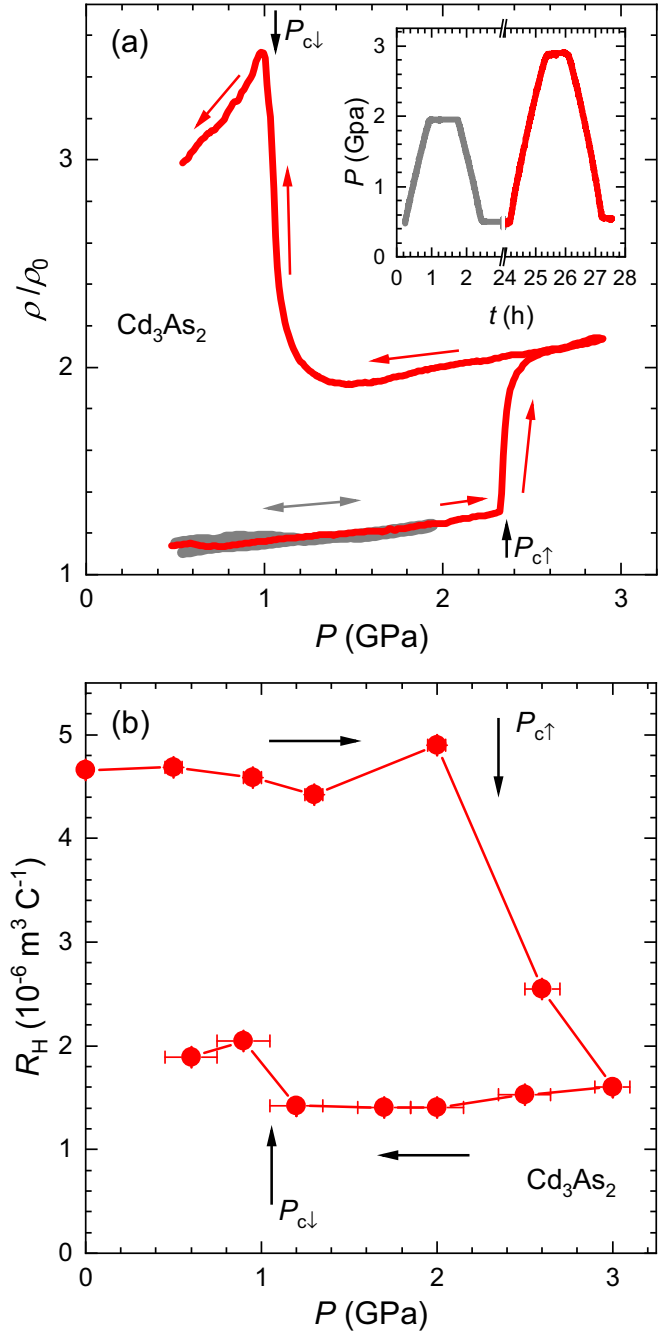


FIG. 2. (a) Nonreversible room-temperature resistivity ρ/ρ_0 where $\rho_0 = \rho(P = 0)$ and (b) Hall effect under pressure in Cd_3As_2 . The initial pressure cycle from ambient pressure to 2 GPa carried out on a pristine sample is shown in gray. It was followed by the second pressure cycle to 2.9 GPa and back down to 0.5 GPa, shown in red. Horizontal arrows indicate the evolution of pressure, vertical arrows highlight the position of the critical pressure on increasing pressure ($P_{c\uparrow}$) and decreasing pressure ($P_{c\downarrow}$). Inset illustrates the pressure protocol for the two cycles.

carrier mobility in the high-pressure phase, as also observed by Zhang *et al.* [33].

The initial value of R_H is not recovered after the pressure cycling. On decreasing pressure, the Hall coefficient remains roughly constant down to $P_{c\downarrow}$, with only a small increase

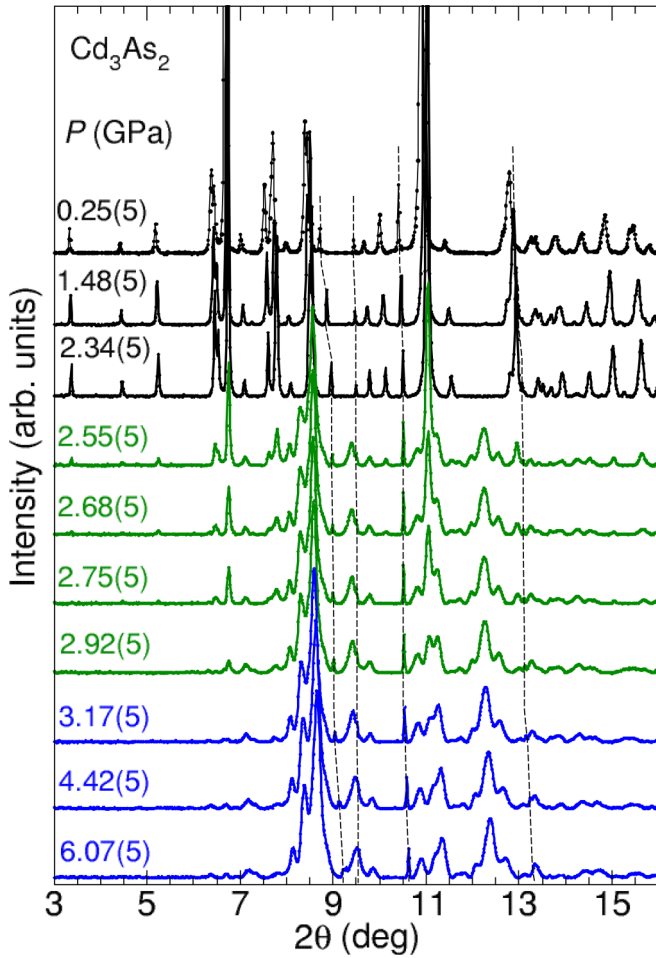


FIG. 3. Evolution of the XRD pattern recorded for Cd_3As_2 at room temperature in sequence proceeding from top to bottom, i.e., pressure raised from 0.25(5) GPa to 6.07(5) GPa. Diffraction patterns consistent with the ambient-condition phase I are shown in black, while blue color is used for the XRD data corresponding to the high-pressure phase V. Diffraction patterns in green contain Bragg peaks originating in both the low- and high-pressure phases of Cd_3As_2 , thus implying coexistence of phases I and V at pressures between 2.45(10) and ~ 3 GPa. Dashed vertical lines indicate positions of peaks originating in Cd.

below $P_{c\downarrow}$ to $R_H = 2\text{ m}^3\text{ C}^{-1}$, a value close to early measurements of polycrystalline samples [46]. The irreversible behavior in $\rho(P)$ and $R_H(P)$ suggests either the transformation into a phase different than the original ambient-condition phase I or fundamentally modified electronic behavior like increased scattering and/or charge carrier doping. To explore the origin of the nonreversibility in the resistivity and Hall coefficient, we performed detailed powder XRD measurements.

B. X-ray diffraction

Our XRD study on a pristine sample of the ambient-condition phase of Cd_3As_2 after loading it into the pressure cell indicate that the powder was well crystalline and single phased, except for a small contribution of cadmium flux detected as characteristic peaks at $2\theta \simeq 8.8^\circ, 9.5^\circ, 10.4^\circ$, and 16.2° (see Fig. 3), and used as a second pressure marker

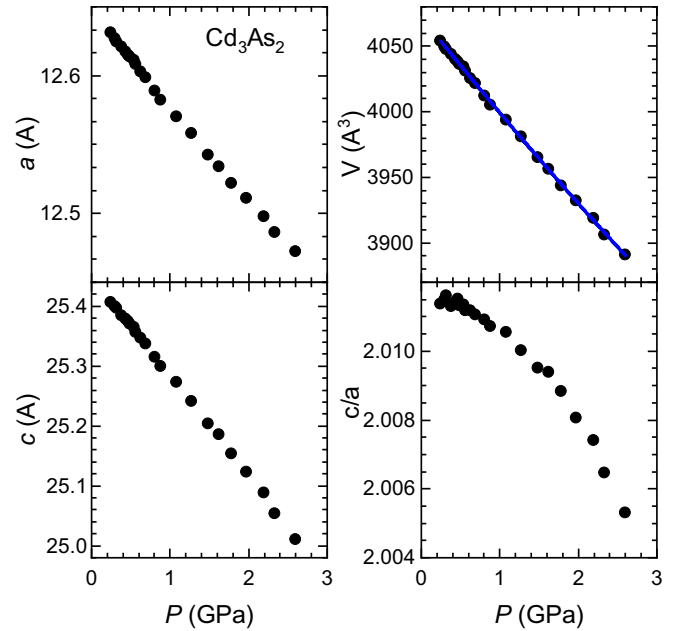


FIG. 4. Lattice parameters, their ratio, and the conventional unit cell volume of the ambient-condition phase of Cd_3As_2 at room temperature plotted as a function of pressure. Error bars are smaller than the size of the circles. Solid blue line is the fit to the experimental data points using the Murnaghan equation of state.

(see Fig. 1). To evaluate microstrain and sizes of Cd_3As_2 crystallites, Le Bail refinements of powder XRD patterns collected from several different sample spots were carried out. Excellent fits to the measured XRD data were achieved using pseudo-Voigt peak profiles with Gaussian contribution describing broadening due to the average grain size $D_V = 170(30)$ nm, and with Lorentzian distribution accounting for microstrain $\epsilon = 0.24(2)\%$. The rather significant positive microstrain indicates the presence of local compressive strain fields. They were likely created when dissipating mechanical energy during grinding single crystals of Cd_3As_2 into a fine powder suitable for the powder XRD study. The refined lattice parameters of the Cd_3As_2 phase I [$a = 12.6303(2)$ Å, $c = 25.409(1)$ Å, see Fig. 4] are very close to those reported in previous studies [21,33,47].

Figure 3 shows the evolution of XRD patterns collected under pressure at room temperature. With increasing pressure, the diffraction peaks are gradually shifted to larger diffraction angles, indicating a pressure-induced shrinkage of the unit cell. For pressures up to $P_{c\uparrow}$, the reflection pattern is consistent with the ambient-condition tetragonal phase I of Cd_3As_2 . The widths and shapes of all the peaks remain virtually the same compared with immediately after closing the pressure cell, indicating an elastic response of the crystal lattice to the applied stress field and giving no evidence for stress heterogeneity which would otherwise result in broadening of diffraction peaks. Lattice parameters derived from Le Bail refinements of the XRD patterns are shown in Fig. 4. The pressure-induced changes in the unit cell volume are well described by the Murnaghan equation of state fit, resulting in a bulk modulus $K_0 = 54(1)$ GPa, the pressure derivative of the bulk modulus $K'_0 = 2.3(8)$, and the ambient pressure

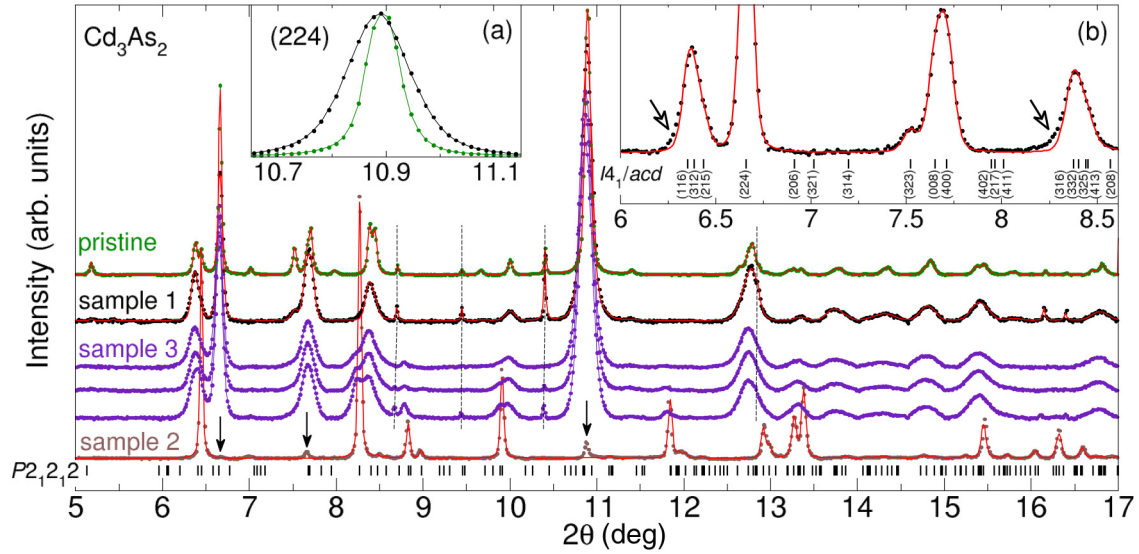


FIG. 5. Powder XRD patterns recorded at room temperature on Cd_3As_2 before [green dots, $P = 0.25(5)$ GPa] and after the pressure cycling. During the pressure cycling, the high-pressure phase was either kept at room temperature [sample 1: black dots, $P = 0.18(5)$ GPa] or was annealed at temperatures up to 240°C and cooled down to room temperature before pressure was released [sample 3: purple dots, $P = 0$; sample 2: beige dots, $P = 0.6(1)$ GPa]. Solid red line on top of the XRD data for sample 2 represents Le Bail refinement performed assuming pseudo-Voigt profiles for diffraction peaks consistent with an orthorhombic lattice with the space group $P2_12_12$ and the lattice parameters $a_{\text{IP}} = 8.026(2)$ Å, $b_{\text{IP}} = 4.1145(5)$ Å, and $c_{\text{IP}} = 18.988(1)$ Å. Angular positions of diffraction peaks expected for this orthorhombic lattice are indicated by short vertical black lines. Solid red lines on top of the XRD patterns for samples 1 and pristine correspond to Le Bail refinements performed assuming the ambient-condition phase I crystal lattice. Vertical arrows mark peaks attributed to a small amount of the ambient-condition phase in sample 2. Dashed vertical lines indicate positions of diffraction peaks originating in Cd. Inset (a) shows the (224) reflection of the tetragonal $I4_1/acd$ phase I of Cd_3As_2 before and after the pressure cycling performed at room temperature, normalized to the maximum intensity. Inset (b) shows enlarged angular range in which a pronounced asymmetric broadening is seen for some Bragg peaks of the ambient-condition phase of Cd_3As_2 recovered after the room temperature pressure cycling. Short black lines in (b) indicate positions of reflections compatible with the crystal structure of the ambient-condition phase.

volume $V_0 = 4072(1)$ Å³. The bulk modulus of the ambient-condition phase of Cd_3As_2 is smaller than that of copper and comparable with elemental Cd.

At pressures above ~ 2.5 GPa, a clear alteration of the XRD pattern is observed (see Fig. 3). This includes the loss of many peaks, e.g., for $2\theta \leq 6^\circ$, and the emergence of new peaks incompatible with the space group $I4_1/acd$ of the phase I, e.g., at $2\theta \simeq 12.5^\circ$, implying a change in the symmetry of the crystal lattice. The transition progresses slowly, with intensities of diffraction peaks originating in the low- and high-pressure phases changing gradually in the pressure range from ~ 2.5 to ~ 2.9 GPa and thus signifying a distinct phase coexistence regime.

The diffraction pattern of the high-pressure phase shown in Fig. 3 is similar to those reported in [29,30,33], where it was interpreted as a monoclinic structure. All the diffraction peaks of the high-pressure phase appear very broad. At the same time, the small width of reflections originating from residual Cd flux highlights the high-quality pressure conditions with negligible pressure gradients. Thus, we conclude that the broadening of the Cd_3As_2 reflections is intrinsic to the sample. Since at low 2θ the effect of microstrain on widths of peaks is insignificant, such a huge broadening even at lowest diffraction angles indicates that, during the pressure-induced structural phase transition, the hundreds of nanometer-sized grains observed in the pristine phase I form much smaller domains in the high-pressure Cd_3As_2 phase V, presumably

due to local strains arising from enhanced crystalline defects. Estimates for the volume-weighted domain size performed using Eq. (3) based on integral breaths of the first four reflections of the high-pressure phase after correcting for the instrumental broadening gave $D_V \approx 16(6)$ nm, an order of magnitude smaller than the average size of crystallites of the low-pressure phase before the transition.

Upon release of pressure to below 1.1 GPa, the high-pressure phase V of Cd_3As_2 converts back into the ambient-condition tetragonal structure. The phase transformation is complete at $P \approx 0.7$ GPa. While the peak positions of the tetragonal phase I have been recovered, the peak widths and shapes recorded after the pressure cycling are remarkably different, as shown in Fig. 5. A strong perturbation of the relative peak intensities is evident especially in the range of $3^\circ \leq 2\theta \leq 7^\circ$. Furthermore, the diffraction peaks of the ambient-pressure phase recovered after releasing pressure are much broader compared with the pristine sample. For instance, the (224) reflection shown in the inset of Fig. 5(a) is broadened to nearly twice its original full width at half maximum. Size and strain analyses based on Le Bail refinements of powder XRD patterns recorded after reducing pressure to 0.2–0.7 GPa indicated that the broadening of diffraction peaks from the low-pressure phase of Cd_3As_2 is dominated by microstrain. The best fits to the experimental data were achieved assuming Gaussian distribution describing the microstrain $\epsilon = 0.57(2)\%$ and a small Lorentzian or Gaussian contribution

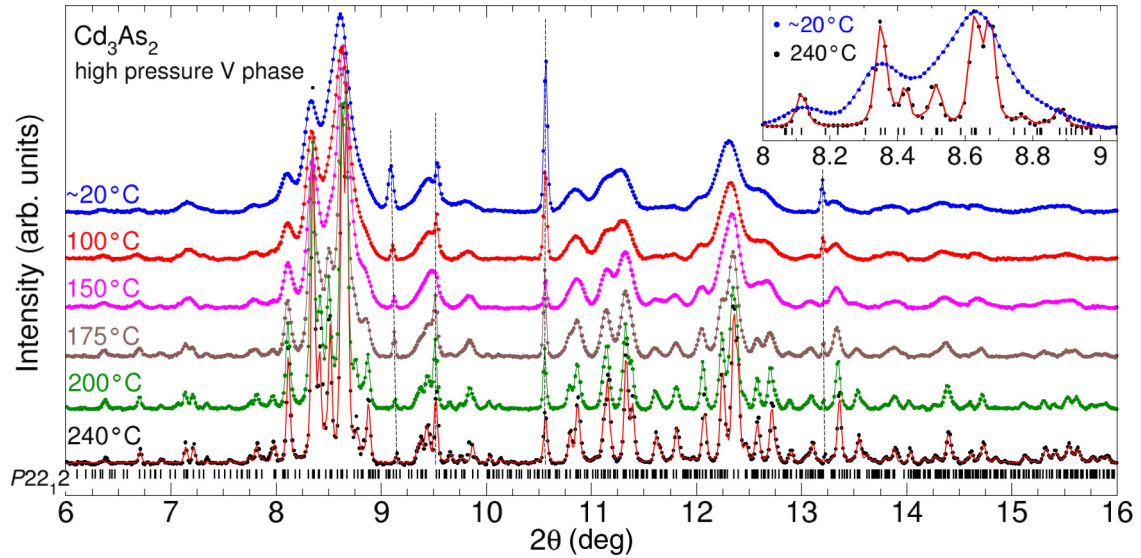


FIG. 6. XRD patterns for the high-pressure phase V of Cd_3As_2 measured on heating from room temperature to 240°C during which the load was kept constant on the DAC but pressure has increased from 4.18(5) GPa at room temperature to ~ 6 GPa at 240°C . Thick solid red lines represent Le Bail refinement performed assuming pseudo-Voigt profiles for diffraction peaks consistent with an orthorhombic lattice with the space group $P2_12_12_1$ and the lattice parameters $a_{\text{HP}} = 8.68 \text{ \AA}$, $b_{\text{HP}} = 17.15 \text{ \AA}$, and $c_{\text{HP}} = 18.58 \text{ \AA}$. Angular positions of diffraction peaks expected for this orthorhombic lattice are indicated by short vertical black lines. Dashed vertical lines show positions of diffraction peaks originating in Cd. Inset shows enlarged angular range with the most intense peak manifold at room temperature (blue dots) and after annealing at 240°C (black dots) normalized to the maximum intensity. The XRD pattern of the annealed sample revealed five intense Bragg peaks for 2θ between 8.2° and 8.7° , whereas for the $P2_1/c$ structure proposed by Zhang *et al.* [29], only two Bragg reflections are expected in this angular range.

included to account for broadening due to the average coherent crystallite size $D_V = 140(60) \text{ nm}$ similar to that before the pressure cycling. Noteworthy, a detailed inspection of the diffraction patterns measured after releasing pressure revealed that peak broadening is anisotropic and shows a distinct asymmetry, e.g., (116) and (316) peaks have pronounced tails on the low angle sides, indicated by arrows in Fig. 5(b). Both the asymmetry and hkl dependence of peak profiles are typical for systems with a large degree of microstrain arising from high concentrations of dislocations and dislocation-type lattice defects [48–51]. In summary, the analysis of the peak widths provides evidence for large nonuniform strain and reduced crystallite domain size in the ambient-condition phase I of Cd_3As_2 recovered after the pressure cycling.

The high-pressure phase of Cd_3As_2 can be annealed at temperatures between ~ 150 and 240°C . The evolution of the XRD patterns collected a few minutes apart during heating is shown in Fig. 6. Upon increasing the temperature, the positions of diffraction peaks remain virtually unchanged due to a cancelation of thermal expansion and further compression as a result of the increasing pressure in the cell with friction locking the membrane. Most notably, raising the temperature above $\sim 150^\circ\text{C}$ leads to a systematic narrowing of the diffraction peaks within minutes between measurements. The process of decreasing the peak widths accelerates with increasing temperature, as expected for a recrystallization anneal and grain growth [52]. It ends shortly after the temperature of 200°C is reached. The Williamson-Hall analyses (not shown) performed on several diffraction patterns recorded at temperatures between 230 and 240°C

on samples 2 and 3 indicated that, after the annealing, for both samples the microstrain was not larger than $\sim 0.06(\%)$, whereas the volume-weighted domain size increased from $\sim 16(6) \text{ nm}$ before the heat treatment to $\sim 80(15) \text{ nm}$ after the annealing.

The crystal structure of the high-pressure phase of Cd_3As_2 remains elusive, but XRD patterns recorded on the annealed samples reveal more details about the symmetry of the crystal lattice. The previously suggested monoclinic lattice ($P2_1/c$ [29]) is incompatible with the observed XRD patterns for the high-pressure phase V of Cd_3As_2 after the annealing. In particular, for the $P2_1/c$ structure proposed by Zhang *et al.* [29], only two reflections are expected to contribute to the most intense peak manifold (see Fig. 1(d) in Ref. [29]), whereas the diffraction patterns of the annealed samples revealed five intense Bragg peaks in the angular range of 8.2° to 8.7° , as shown in Fig. 6. Further, two Bragg peaks were detected at $2\theta \approx 8.76^\circ$ and 8.88° , while no reflections are expected for the $P2_1/c$ structure in this angular range. The XRD patterns measured on samples 2 and 3 after the annealing at temperatures up to 240°C at $P = 5(1) \text{ GPa}$ were virtually identical. Bragg reflections detected consistently in those diffraction patterns were used for the purpose of indexing. Searching for candidate crystal lattices did not give satisfactory results for cubic, tetragonal, hexagonal, and trigonal crystal systems with unit cell volumes of up to 6000 \AA^3 . Among orthorhombic structures, the best indexing solution with the smallest unit cell volume that accounts for the observed diffraction peaks has a primitive lattice with $a_{\text{HP}} \approx 8.68 \text{ \AA}$, $b_{\text{HP}} \approx 17.15 \text{ \AA}$, and $c_{\text{HP}} \approx 18.58 \text{ \AA}$. We note that such a crystal

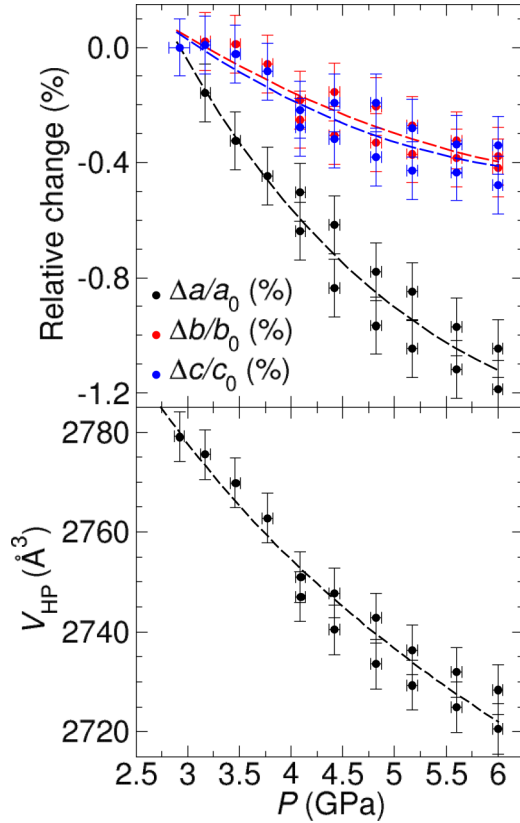


FIG. 7. The pressure dependence of the unit cell volume for the high-pressure phase V of Cd_3As_2 (bottom panel) and the pressure-induced relative changes in its lattice parameters (top panel) recorded at room temperature. Dashed lines are guides to the eye.

lattice may be considered as a superstructure of the anti-fluorite structure type made of 96 distorted $\text{Cd}_6\Box_2$ cubes, similarly to the ambient-pressure polymorphs. Searching for monoclinic cells gave candidate lattices with comparable or only slightly smaller unit cell volumes ($\Delta V/V \lesssim 12\%$). Notably, none of the lattices with smaller unit cell volumes could be interpreted in terms of the anti-fluorite-type building blocks. Therefore, we conclude that the structure of the high-pressure phase of Cd_3As_2 is likely primitive orthorhombic. Space group tests indicated the chiral $P22_12$ or one of the lattices without systematic extinctions, i.e., centrosymmetric $Pmmm$, noncentrosymmetric chiral $P222$ or achiral $P2mm$, $Pm2m$, or $Pmm2$. Indexing of the high-pressure Cd_3As_2 phase V assuming the space group $P22_12$ is shown in Fig. 6.

To evaluate the compressibility of the high-pressure phase of Cd_3As_2 , sequential Le Bail refinements of powder XRD patterns collected at room temperature were performed assuming the orthorhombic crystal lattice. The resulting pressure dependencies of the lattice parameters and the unit cell volume are shown in Fig. 7. The isothermal compressibility of the high-pressure phase immediately after the structural transformation is found to be $\kappa_T = 0.9(1) \times 10^{-11}$ Pa, 45 % smaller than the compressibility of phase I [$\kappa_T = 1.65(2) \times 10^{-11}$ Pa, see Fig. 4], and it decreases further with pressure. The notable reduction in compressibility can be rationalized by an increase in the density of Cd_3As_2 at the structural

transition by $\sim 4.8(2)\%$ estimated from the lattice parameters of the low- and high-pressure phases, assuming that the latter is orthorhombic and consists of 96 distorted $\text{Cd}_6\Box_2$ cubes.

Upon lowering pressure below ~ 1 GPa, the diffraction peaks of the annealed high-pressure phase in sample 3 start to diminish, and finally, the reflection pattern characteristic for the low-pressure phase of Cd_3As_2 emerges. As shown in Fig. 5, the broadening of the diffraction peaks is very similar to that observed for sample 1 after the pressure cycling at room temperature. However, next to the peaks expected for the ambient-condition phase of Cd_3As_2 , there are also distinct additional contributions to the diffraction patterns at $2\theta \approx 8.25^\circ$, 8.78° , 9.89° , and 11.80° that may indicate the formation of a new phase. While intensities of those extra peaks vary between different sample spots, their angular positions remain the same. We note that the additional features correspond well to the diffraction peaks observed for sample 2 after annealing at temperatures up to 240°C at pressures of 6 GPa and releasing the pressure. Since for sample 2 glycerol was used as a pressure-transmitting medium causing significant deviations from hydrostatic conditions in the accessed T - P regime [53,54], we surmise that deviatoric stress induced by both the nonhydrostaticity of the pressure medium and larger size of the high-pressure phase crystallites can stabilize an intermediate metastable phase of Cd_3As_2 forming during the pressure-driven structural phase transition at room temperature.

The diffraction peaks of the intermediate phase of Cd_3As_2 observed at ambient conditions can be indexed as a primitive orthorhombic structure with the lattice parameters $a_{IP} = 8.026(2)$ \AA , $b_{IP} = 4.1145(5)$ \AA , and $c_{IP} = 18.988(1)$ \AA and the space group $P2_12_12$. We note that those lattice parameters are related to the lattice parameters of the high-pressure phase: $a_{IP} \approx a_{HP}$, $b_{IP} \approx b_{HP}$, and $c_{IP} \approx c_{HP}/4$, with the differences of $+7.7\%$, -2.2% , and $+3.9\%$, respectively. Therefore, we suspect that the intermediate phase of Cd_3As_2 can be considered as another superstructure of the anti-fluorite structure type. Le Bail refinements performed assuming pseudo-Voigt profiles for Bragg peaks compatible with the orthorhombic lattice indicated a sizable degree of nonuniform strain $\epsilon = 0.16(2)\%$. The average crystallite size $D_V = 108(10)$ nm estimated from the refinements is comparable with the mean size of coherent domains of the high-pressure phase after annealing at temperatures up to 240°C .

IV. DISCUSSION

Our high-pressure studies show that, for bulk Cd_3As_2 , the phase I hosting topologically protected electronic states can be maintained up to 2.4 GPa but cannot be recovered with similar high mobilities once this level of compression is exceeded. Recently, Goyal *et al.* [24] found that the quantum Hall effect associated with the surface states persists in thin films of Cd_3As_2 under strain up to 0.5%. While Goyal *et al.* use biaxial strain in the (112) plane, the magnitude of the strain is equivalent to the modification of the crystal parameters at 1 GPa in our work using hydrostatic pressure. Hence, we expect that thin films with strain above $\approx 1\%$ will fall outside

the stability range of Cd_3As_2 phase I and will no longer host quantum Hall effect states.

Our high-resolution powder XRD measurements allow us to evaluate in detail the changes in the lattice parameters of the ambient-condition tetragonal phase I of Cd_3As_2 induced by the applied pressure. While the observed pressure dependence of the unit cell volume shown in Fig. 4 is similar to those reported in Refs. [29,33], our XRD data does not give any indication for a sudden jump in c/a at pressures below 1.5 GPa which Zhang *et al.* attributed to an additional topological phase transition [33]. Instead, our XRD study reveals a smooth decrease of the c/a ratio with applied pressure. Remarkably, the observed changes in the c/a ratio accord well with the pressure dependence of the Fermi wave vector derived from Shubnikov-de Haas oscillation measurements [33]. Both the axial compression along the c direction and the unexpected shrinking of the Fermi surface accelerate at $P \gtrsim 1.7$ GPa. First-principles electronic band structure calculations performed by Zhang *et al.* [33] indicated that the shortening of the lattice parameter c should shift the Dirac nodes toward the center of the Brillouin zone, thus reducing the Fermi wave vector and providing an explanation for the gradual increase of resistivity under pressure within the ambient-condition phase of Cd_3As_2 (see Fig. 2(a) and Ref. [29]).

Our high-pressure studies provide a confirmation that the I–V transition in Cd_3As_2 is of first-order type. The electrical resistivity and powder XRD measurements both detect the phase transition with an onset at ~ 2.4 and ~ 1.3 GPa on raising and releasing pressure, respectively. The sizable hysteresis evidences the first-order nature of the structural transformation, which shows a significant width of ~ 0.2 and ~ 0.5 GPa at $P_{c\uparrow}$ and ~ 0.4 and ~ 0.5 GPa at $P_{c\downarrow}$ based on the electrical resistivity and XRD data, respectively. We note that some discrepancies between the two methods are not unusual because the former potentially probes a percolative path, whereas the latter reflects changes in the entire sample volume. Importantly, the distinct pressure ranges with coexistence of the low- and high-pressure phases indicate sluggishness and thus an energy barrier between phases I and V of Cd_3As_2 . Therefore, we deduce that the transition process involves drastic changes in the system of chemical bonds and leads to a major reorganization of the crystal structure.

The observed increase in width of diffraction peaks at the transition from phase I to phase V indicates that the transformation causes fragmentation of crystallites into much smaller domains. Therefore, we attribute the tenfold reduction in mobilities of charge carriers inferred from the electrical transport study to the combined effects of an increased scattering resulting from the strongly reduced coherent domain size and the loss of Dirac electrons with high Fermi velocity caused by changes in the symmetry of the crystal lattice overcompensating the increase in charge carrier concentration.

He *et al.* have interpreted the superconductivity observed above 8 GPa to retain topological character [30]. This identification of topological superconductivity was based on presence of Dirac electrons with topological superconductivity possibly stabilized by a reduction of the crystal symmetry in the high-pressure phase [55]. However, our identification of the

reconstructive I–V phase transition suggests a fundamental modification of the structure. While we show that the high-pressure phase may contain the same $\text{Cd}_6\Box_2$ building blocks, they are very likely distorted and arranged in a different configuration. Hence, the Dirac electrons which are sensitive to the orbitals at the voids [22] are likely absent in the high-pressure phase V, and superconductivity in this phase is not of topological character. At the same time, our high-temperature annealing experiments highlight a route to study the superconductivity in this phase in more detail for samples with high crystallinity.

Recovering the ambient-condition phase I after cycling pressure at room temperature beyond $P_{c\uparrow}$ resulted in a large degree of microstrain. Estimates from profile refinements using the Le Bail method gave $\epsilon \approx 0.57(2)\%$, which is three orders of magnitude larger than ϵ values $\sim 10^{-6}$ expected for nearly strainfree crystals [56] and nearly three times larger than the microstrain induced by grinding crystals into a fine powder with tens of nanometer sized grains. Although results of the size and strain analyses need to be considered with caution because of significant peak overlap, the large positive ϵ indicates the presence of local compressive stress fields arising from lattice disorder such as dislocations, stacking faults, intercalations of atoms, or grain boundaries. Such defects may give rise to unintentional doping [57,58], providing a plausible explanation for the observed decrease in the Hall coefficient for the ambient-condition phase I of Cd_3As_2 after the pressure cycling [see Fig. 2(b)]. Thus, we infer that the unexpected increase in resistivity after returning to the low-pressure regime below $P_{c\downarrow}$ shown in Fig. 2(a) originates in a largely enhanced scattering that outweighs the expected increase in carrier mobilities due to the formation of bulk Dirac points and the gain in the charge carriers from self-doping. We note that studies on Cd_3As_2 epilayers revealed a substantial effect of extended defects on electron mobilities [59]. Moreover, for epitaxial (112) Cd_3As_2 films, a systematic decrease in Hall mobilities with the compressive in-plane strain ($\epsilon_s \gtrsim 0.5\%$) was observed [24]. Further studies, including high-resolution transmission electron microscopy imaging, are needed to address the origin of the increased scattering in the ambient-condition phase of Cd_3As_2 retrieved after the room-temperature pressure cycling.

V. CONCLUSIONS

Our high-pressure studies highlight the disruptive nature of the pressure-driven structural phase transition in Cd_3As_2 . We find large changes in both the electronic and microstructural characteristics at the room-temperature structural transformation. The reduction of the carrier mobility is shown to result from the formation of microstrain and fragmentation of crystallites into smaller domains caused by the phase transition. Our transport measurements indicate that the increased resistance is dominated by the enhanced scattering outweighing the gain in charge carriers in the high-pressure phase V. We show that annealing of the high-pressure phase at temperatures of $\sim 200^\circ\text{C}$ results in a largely improved crystallinity and thus should enable future studies aiming at solving the crystal structure and exploring

the electronic properties. Finally, we observe the formation of a new primitive orthorhombic phase prompted presumably by deviatoric stress upon releasing pressure through the structural phase transition. Our results highlight the importance of considering strain and crystallite size when tailoring electronic properties of Cd_3As_2 for applications of bulk and thin films.

Data are available at the University of Bristol data repository [data.bris](https://data.bris.ac.uk/) [60].

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- [39] The pseudo-Voigt function is an approximation of the Voigt function defined as
- $$pV(x) = \eta L(x) + (1 - \eta)G(x), \quad (8)$$
- with $L(x)$ and $G(x)$ denoting Lorentzian and Gaussian parts, respectively, that are of the same full width at half maximum (FWHM) H , and are weighted by η adopting values between 0 and 1 and thus shifting the profile more toward pure Gaussian or pure Lorentzian. The H and η parameters are related to the FWHM values of the deconvoluted Lorentzian (H_L) and Gaussian (H_G) functions [61]:
- $$H = H_G^5 + 2.69269H_G^4H_L + 2.42843H_G^3H_L^2 + 4.47163H_G^2H_L^3 + 0.07842H_GH_L^4 + H_L^5,$$
- $$\eta = 1.36603 \frac{H_L}{H} - 0.47719 \left(\frac{H_L}{H} \right)^2 + 0.11116 \left(\frac{H_L}{H} \right)^3.$$
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- $$\beta_L = \frac{\pi H_L}{2}, \quad (9)$$
- $$\beta_G = \frac{H_G}{2} \sqrt{\frac{\pi}{\ln 2}}, \quad (10)$$
- $$\beta_{pV} = \frac{0.5\pi H}{\eta + (1 - \eta)\sqrt{\pi \ln 2}}, \quad (11)$$
- respectively.
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