

Anomalous anisotropic compression behavior of superconducting CrAs under high pressure

Zhenhai Yu^{a,1}, Wei Wu^{b,1}, Qingyang Hu^a, Jinggeng Zhao^c, Chunyu Li^a, Ke Yang^d, Jinguang Cheng^b, Jianlin Luo^{b,e}, Lin Wang^{a,f,2}, and Ho-kwang Mao^{a,2}

^aCenter for High Pressure Science and Technology Advanced Research, Shanghai 201203, People's Republic of China; ^bBeijing National Lab for Condensed Matter Physics, Institute of Physics, Chinese Academy of Sciences, Beijing 100190, People's Republic of China; ^cNatural Science Research Center, Academy of Fundamental and Interdisciplinary Sciences, Harbin Institute of Technology, Harbin 150080, People's Republic of China; ^dShanghai Institute of Applied Physics, Chinese Academy of Sciences, Shanghai 201203, People's Republic of China; ^eCollaborative Innovation Center of Quantum Matter, Beijing 100190, People's Republic of China; and ^fState Key Laboratory of Superhard Materials, Jilin University, Changchun 130012, People's Republic of China

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CrAs was observed to possess the bulk superconductivity under high-pressure conditions. To understand the superconducting mechanism and explore the correlation between the structure and superconductivity, the high-pressure structural evolution of CrAs was investigated using the angle-dispersive X-ray diffraction (XRD) method. The structure of CrAs remains stable up to 1.8 GPa, whereas the lattice parameters exhibit anomalous compression behaviors. With increasing pressure, the lattice parameters *a* and *c* both demonstrate a nonmonotonic change, and the lattice parameter *b* undergoes a rapid contraction at ~0.18–0.35 GPa, which suggests that a pressure-induced isostructural phase transition occurs in CrAs. Above the phase transition pressure, the axial compressibilities of CrAs present remarkable anisotropy. A schematic band model was used to address the anomalous compression behavior of CrAs. The present results shed light on the structural and related electronic responses to high pressure, which play a key role toward understanding the superconductivity of CrAs.

CrAs | anisotropic compression behavior | high pressure | phase transformation

The discovery of superconductivity ($T_c = 26$ K) in ZrCuSiAs-type LaFeAs(O_{1-x}F_x) (1) inspired extensively experimental and theoretical research on the quaternary “1111” compounds RFeAsO, where R represents a lanthanide (La, Ce, Nd, etc.) (2–5). From the point of view of crystallography, the newly discovered iron-based superconductor exhibits a quasi-2D structural character at ambient conditions. Like the CuO₂ plane in copper oxide high-temperature superconductors, the Fe₂As₂ layers serve as the conduction planes for charge carriers, and the other building blocks are the charge reservoir layers, dominating the carrier density or the chemical potential. The fundamental structural factors of these compounds are of scientific interest as a meaning of achieving a higher superconducting transition temperature. Previous investigation suggests that the symmetry of the tetrahedral FeAs₄ (6), the FeAs interlayer spacing (7), and the anion (such as As) heights from the iron layer (8, 9) are all crucial aspects of the structure. The FeAs planes in binary transition metal monoarsenides (TAs) may serve an analogous role to that served by the CuO plane as a structural proxy for the layered cuprate perovskite family of compounds. These considerations may shed light on the exploration of novel superconductors (10, 11). Moreover, CrAs, in which low-lying states near the Fermi level are dominated by 3d electrons, was found to exhibit superconductivity on the verge of antiferromagnetic order via the application of an external high pressure (10).

Transition-metal-based monoarsenides such as CrAs and MnAs have been extensively studied during the past decades mainly due to their complex magnetism and magnetocaloric effects (12, 13). Furthermore, antiferromagnetic order in FeAs was observed below 70 K (14), whereas the discovery of iron-arsenide-based superconductors has renewed interest in FeAs. From a crystallographic viewpoint, CrAs is isostructural with MnAs, FeAs, and VAs, which implies that novel physical properties may

originate from the above structural characteristics. The lattice parameter *b* (space group with *Pnma* setting) of CrAs is the largest among transition-metal-based monoarsenides at ambient conditions, as shown in Fig. S1. CrAs crystallizes into a MnP-type structure ($a = 5.6411$ Å, $b = 3.4797$ Å, and $c = 6.1959$ Å), as shown in Fig. S24 at ambient conditions. Neutron diffraction studies of CrAs showed that it is a double-helical-type antiferromagnet with a Néel temperature (T_N) of ~260 K (15). The first-order magnetic transition of CrAs at T_N is accompanied by discontinuous changes of the lattice parameters. The most dramatic change is a sudden expansion of the lattice parameter *b* by ~4% below T_N (16). In our previous study (17), high-quality single-crystal CrAs was grown successfully using the Sn flux method. A clear magnetic transition was observed along with a sharp drop of the susceptibility and the resistivity as well at about 270 K. Whereas investigation of the temperature-dependent resistivity, $\rho(T)$, of CrAs showed that superconductivity did not appear even down to 350 mK at ambient pressure, it is interesting that high pressure has been found to drive the resistivity drop progressively toward higher temperatures with increasing pressure inside a diamond anvil cell, and, ultimately, the occurrence of superconductivity was observed near 1 K at a pressure of ~0.3 GPa (10).

As mentioned above, the application of high pressure has been verified as an effective approach to induce unconventional superconductivity. The pressure dependence of T_N for CrAs has

Significance

We performed high-pressure structural studies of CrAs to understand the physical mechanism of its pressure-induced superconductivity. It was found that the lattice parameters *a* and *c* both demonstrate a nonmonotonic change, and the lattice parameter *b* undergoes a rapid contraction at ~0.18–0.35 GPa, which suggests that a pressure-induced isostructural phase transition occurs in CrAs. Above the phase transition pressure, the axial compressibilities of CrAs present remarkable anisotropy. The pressure-related structural changes occurred concurrently with the appearance of bulk superconductivity, shedding light on the structural and related electronic responses to high pressure, which play a key role toward understanding the superconductivity of CrAs.

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¹Z.Y. and W.W. contributed equally to this work.

²To whom correspondence may be addressed. Email: wanglin@hpstar.ac.cn or hmao@carnegiescience.edu.

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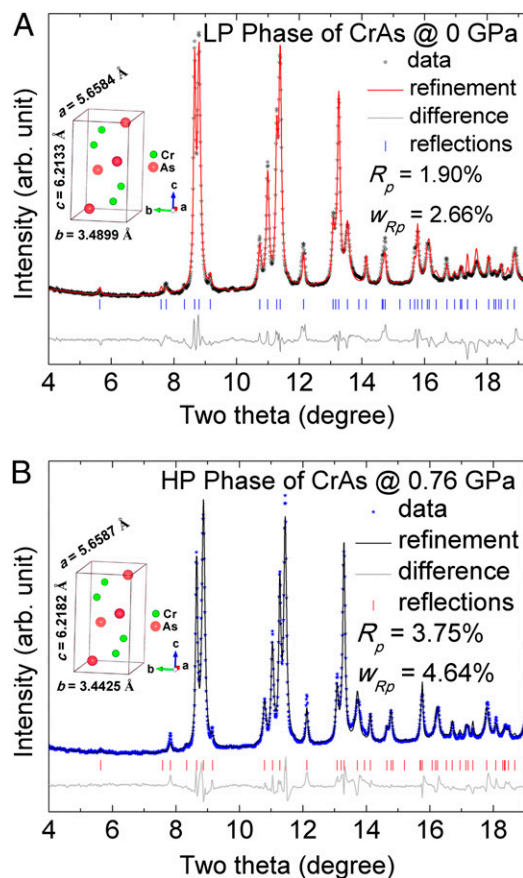


Fig. 1. Refinement of AD-XRD data of CrAs at ambient conditions (A) and 0.76 GPa (B). Measured (dots) and calculated (solid lines) patterns are shown, together with the difference curve (gray lines) and calculated positions of Bragg reflections (tick marks).

also been investigated by Keller et al. (18), where T_N was observed to decrease with increasing pressure (18, 19). Measurement of the electrical conductivity of CrAs at 4.2–350 K under high pressure up to 0.9 GPa was performed by Zavadskii and Sibarova (20). It was shown that CrAs exhibits metallic conductivity over

the entire range of pressure and temperature considered. Recently, Shen et al. investigated the structural and magnetic phase diagram of CrAs as a function of the temperature and pressure (21). The experimental results indicated that a spin reorientation was induced above the critical pressure (~ 0.6 GPa). To our knowledge, most of the studies reported focused on investigation of the chemical doping, temperature, or pressure effect on the structural phase transitions or on the magnetic and electrical properties of CrAs. The crystal structure of CrAs under high pressure has not been well studied, and therefore the correlation between the physical properties and crystal structure of CrAs has not yet been well understood. For example, the mechanism whereby the lattice parameters of CrAs show a juncture at T_N is still not fully understood. Therefore, it is essential to understand the relation between the structure and the properties of CrAs under compression. For this purpose, an in situ high-pressure X-ray diffraction (XRD) investigation of CrAs is undertaken in the present study using the angle-dispersive XRD technique with very fine pressure steps. It was found that the lattice parameters present anomalous compression behaviors under high pressure, which was interpreted using a schematic band model. The structural evolution was consistent with superconducting behavior under high pressure, providing crystallographic evidence for an enhanced appreciation of the physical mechanism.

Results and Discussion

Fig. 1A shows the XRD pattern of CrAs at ambient conditions with refinement based on a previously reported model (14). The tick marks below the diffraction pattern indicate the locations of the calculated diffraction peaks in the plot. The residual difference between the calculated and the experimental patterns is shown at the bottom of the plot. The good refinement confirmed that the crystal structure of CrAs at ambient conditions is $Pnma$, as shown in Fig. 1A, *Inset*. The detailed refinement results were shown in Table S1.

Fig. 2A presents selected patterns of the angle-dispersive XRD results of CrAs under various pressures. The sample was pressurized in small steps of ~ 0.1 GPa. At first glance, it can be clearly observed in the figure that the Bragg peaks shift toward higher angles owing to lattice contraction without any modification to the overall peak patterns, and no new diffraction peaks were observed in the XRD patterns up to a pressure of 1.81 GPa, except for an apparent separation of the diffraction peaks. To obtain further insight into the effect of the pressure on the

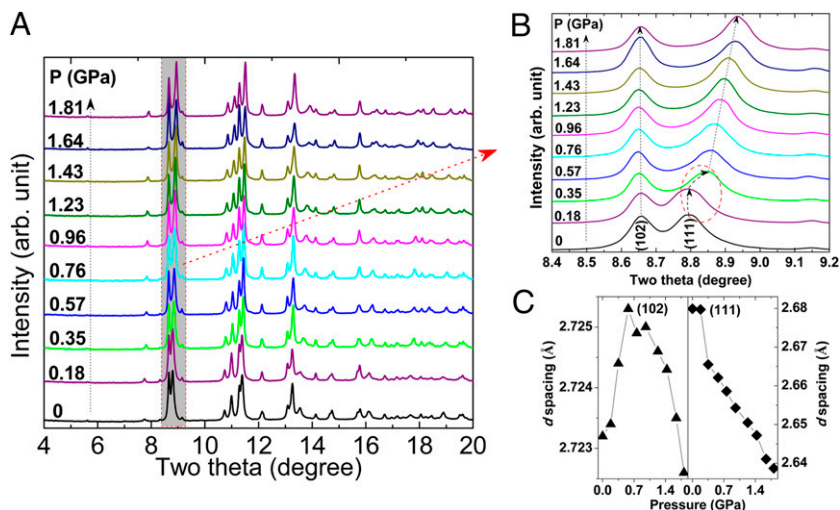


Fig. 2. (A) Selected patterns of the AD-XRD results of CrAs under various pressures. (B) Local diffraction peaks magnified to highlight the pressure dependence. (C) Selected d spacing as a function of pressure for CrAs.

The axial compressibility of CrAs is remarkably anisotropic after the isostructural phase transition point, and the relative axial compressibilities along the a -, b -, and c axis are $7.98 \times 10^{-3} \text{ \AA/GPa}$, $3.84 \times 10^{-2} \text{ \AA/GPa}$, and $3.71 \times 10^{-3} \text{ \AA/GPa}$, respectively. The compressibilities along the a - and c axis are more than an order of magnitude smaller than that along the b axis.

CrAs could be considered as a 3D network comprising CrAs₆ octahedra, in contrast with the quasi-layers in the iron-based superconductor. With regard to the anisotropic axial compressibility of CrAs, it could be explained by the different stacking of the CrAs₆ octahedra along the different crystallographic axes. The structure of the Cr-centered CrAs₆ octahedra viewed from different directions is depicted in Fig. 4 A–C. The CrAs₆ octahedra are face-connected along the a axis and edge-shared along the b axis. Neighboring CrAs₆ layers along the c axis are one-half edge-shared and one-half vertex-shared connections. The average numbers of the CrAs octahedral layers along the a -, b -, and c axis per unit cell are 2, 1, and 2, respectively, and the thicknesses of every CrAs₆ octahedral layer along the a -, b -, and c axis is 2.8206 (5.6411/2), 3.4797 (3.4797/1), and 3.0980 (6.1959/2) Å, respectively. Thus, the CrAs₆ octahedral layers along the b axis are sparser than those along the a - and c axis.

The calculated bonding lengths and bonding angles for the environment surrounding the Cr atoms indicate that the CrAs₆ octahedron is not perfect at ambient conditions. The pressure dependence of the As–Cr–As bonding angle ($\angle\text{As–Cr–As}$) of CrAs is shown in Fig. 4D. It can be seen from Fig. 4D that the drastic contraction of lattice parameter b resulted in two As–Cr–As angles exhibiting a contrary pressure dependence. Furthermore, greater distortion of the CrAs₆ octahedra and the shortening of the Cr–As bonding lengths make the CrAs₆ octahedra greatly compressed along the b axis under high pressure. The pressure dependence of the As–Cr–As bonding angle is shown in Fig. 4D, which also gives rise to an increase of the crystal-field splitting energy. Selected bonding lengths ($d_{\text{Cr–As}}$) of CrAs as a function of pressure are shown in Fig. S5. It can be seen from Fig. S5 that the bonding length correlated to the lattice parameters a and c shows subtle variation under high pressure, whereas the bonding length correlated to the lattice parameter b exhibits significant contraction as the pressure increases.

The band model of TAs was first proposed by Goodenough (23), Menyuk et al. (24), and Goodenough and Kafalas (25), and later developed by Boller and Kallel (26). Here, we interpret the anomalous compression behaviors of CrAs at ~ 0.18 – 0.35 GPa in terms of the above-mentioned band model. The electronic configuration of the Cr³⁺ ions in CrAs is $3d^3$. In light of crystal ligand field theory, CrAs crystallizes into the MnP-type structure, and the central Cr atoms are octahedrally coordinated with six As atoms. The d states of the Cr atom split into t_{2g} (d_{xy} , d_{yz} , and d_{zx}) and e_g (d_{z^2} and $d_{x^2-y^2}$) manifolds. The t_{2g} states of Cr hybridize with the p state of As atoms, forming the bonding and the antibonding states. Therefore, the e_g states of Cr are nonbonding and form the narrow states. The present calculated band structure of CrAs at ambient pressure is shown in Fig. S6A, where the Fermi level lies inside the energy band, indicating its metallic characteristics, and is consistent with the experimental results (10). The band scheme of CrAs under ambient and high pressures is shown in Fig. 5. The ions being arranged along the b axis space so far apart (~ 3.45 Å) that their wave functions of the $3d$ electrons possess little overlap, resulting in weak electron repulsion along this axis. Therefore, the b axis is much more compressible at the onset of pressure application compared with the a - and c axis. As the pressure increases, the widths of all energy bands in CrAs including the energy band along the $\gamma \rightarrow Y$ direction related to the electronic structure along the b axis in real space (hereafter EB _{b}) are broadened. When the applied pressure is increased to the critical point (~ 0.18 – 0.35 GPa), the EB _{b} just crosses into the Fermi level with partially filled electrons.

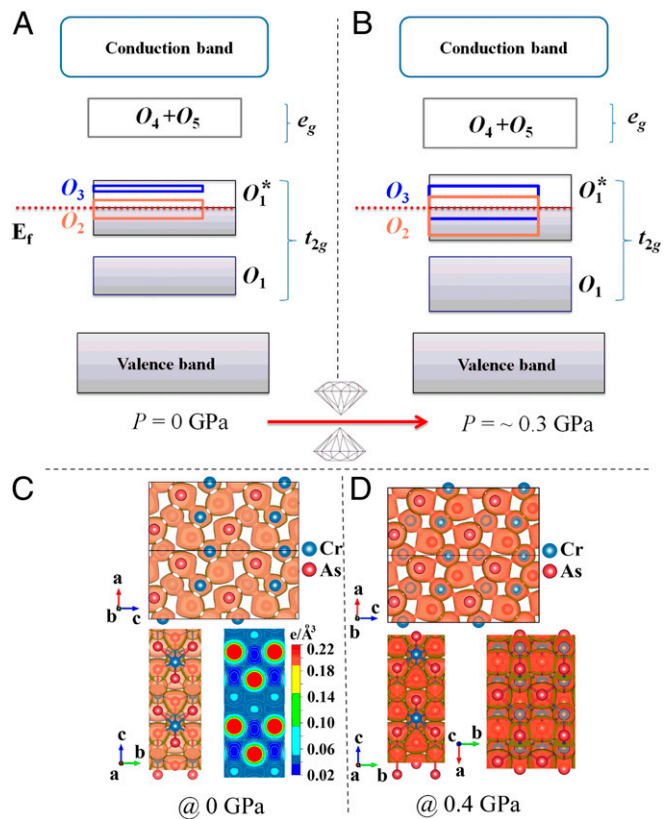


Fig. 5. Band scheme of CrAs at (A) ambient and (B) high pressures. (C) Charge density projected along the a – c plane. The visualized isosurfaces equal to a charge density of 0.05 e/\AA^3 . Atoms in the front layer (bright colors) are inserted into the interstitials of the back layer (atoms in gloomy colors) upon compression. Charge density projected along the b – c plane. A slice of charge density at the plane of $c = 0$ Å is shown in contours. The interatomic forces are derived from the gradient of energy, also known as the charge density. Because the charge density is less steep along the b axis, it undoubtedly achieves better compressibility. (D) Charge density projected along the a – c , b – c , and a – b planes of CrAs under 0.4 GPa.

The electronic repulsion also causes the b axis to become more incompressible. Therefore, the energy band model satisfactorily addresses the pressure-dependency curve of the lattice parameter b given in Fig. 3B and the electronic responses to high pressure, which is important in understanding the pressure-induced superconductivity of CrAs. Charge density projected along the a – c and b – c planes of CrAs under ambient pressure are shown in Fig. 5C. A slice of charge density at $c = 0$ Å is shown in contours (Fig. 5C, Right). The charge density is less dense (deeper color) along the b axis, resulting in a better compressibility. The results drawn here are consistent with the foregoing energy band model analysis. With increasing pressure the gradient of charge density decreases as shown in Fig. 5D. These findings provide consistent evidence with pressure-induced variation of crystal structure (i.e., volume contraction and distortion of CrAs₆ octahedral units in CrAs), in which the change of electronic structure was mainly affected by the shrinking of the b axis.

At ambient conditions, CrAs possesses the paramagnetic metallic characteristic and the electrons are in itinerant status. The itinerant state approaches low temperature as the pressure increases. This tendency continues until the T_N is suppressed to very low temperature. The superconductivity appears when the itinerant state occurs around ~ 2 K.

As seen from Fig. 3, the lattice parameters a and c of CrAs exhibited elongation while lattice parameter b experienced rapid

contraction ~ 0.18 – 0.35 GPa. Based on the experimental results of ref. 21, this suggests that the magnetism of CrAs may exhibit changes such as an abrupt drop of the magnetic propagation vector at a critical pressure (~ 0.6 GPa) (21). In addition, the large decrease of b axis observed in the present work has not been reported by Shen et al. (21).

Conclusions

In this paper, we presented crystallographic structural studies of the superconductor CrAs under high pressure with small pressure steps using synchrotron X-ray powder diffraction, which we believe to be heretofore unreported. The lattice parameters of CrAs exhibited anomalous variations as a function of pressure. We suggested that this was a pressure-induced isostructural phase transition. In the presently investigated pressure range, CrAs was remarkably anisotropic. Note that the transition pressure (~ 0.18 – 0.35 GPa) obtained from this work coincides with that of the emergence of pressure-induced bulk superconductivity in CrAs. The currently observed pressure-induced isostructural phase transition in CrAs deviates markedly from previously documented classifications.

Materials and Methods

The high-pressure synchrotron XRD experiment was carried out using a symmetric diamond anvil cell. T301 stainless steel served as the gasket. The

150- μm -diameter sample chamber was filled with a mixture of CrAs powder, a ruby chip, and silicone oil as the pressure-transmitting medium. Angle-dispersive XRD (AD-XRD) patterns were obtained with a MarCCD detector using monochromatic radiation with a wavelength of 0.4112 \AA at the X17C beamline of the NSLS, Brookhaven National Laboratory (BNL), and at the BL15U1 beamline of the SSRF using a wavelength of 0.6199 \AA . The Fit2D software package was used to process the data (27). The pressures were determined according to the fluorescence shift of ruby (28). The XRD patterns of CrAs were analyzed with Rietveld refinement using the GSAS program package (29) with a user interface EXPGUI (30). The AD-XRD experiments were carried out at room temperature.

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