

Superconductivity up to 243 K in the yttrium-hydrogen system under high pressure

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The discovery of superconducting H₃S with a critical temperature $T_c \sim 200$ K opened a door to room temperature superconductivity and stimulated further extensive studies of hydrogen-rich compounds stabilized by high pressure. Here, we report a comprehensive study of the yttrium-hydrogen system with the highest predicted T_c s among binary compounds and discuss the contradictions between different theoretical calculations and experimental data. We synthesized yttrium hydrides with the compositions of YH₃, YH₄, YH₆ and YH₉ in a diamond anvil cell and studied their crystal structures, electrical and magnetic transport properties, and isotopic effects. We found superconductivity in the *Im-3m* YH₆ and *P6₃/mmc* YH₉ phases with maximal T_c s of ~ 220 K at 183 GPa and ~ 243 K at 201 GPa, respectively. *Fm-3m* YH₁₀ with the highest predicted $T_c > 300$ K was not observed in our experiments, and instead, YH₉ was found to be the hydrogen-richest yttrium hydride in the studied pressure and temperature range up to record 410 GPa and 2250 K.

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High-temperature superconductivity (HTSC) is of great interest and importance because superconductors operating under ambient or technologically accessible pressure and temperature conditions can vastly improve many areas of technology. The conventional theories of Bardeen–Cooper–Schrieffer¹ and Migdal–Eliashberg^{2,3} indicate that superconductivity, even at room temperature, cannot be excluded. However, for a long time, the critical temperature of conventional superconductors has been limited to $T_c = 39$ K in MgB_2 ⁴. The discovery of unconventional superconductivity in the cuprates family has provided a strong motivation to further search for superconducting materials at even higher temperatures. Despite great efforts, superconductivity soon reached its limit in T_c at ~ 133 K⁵ (~ 164 K under pressure⁶). A microscopic theory of high- T_c unconventional superconductors is still lacking, which hampers further advances in this field.

The breakthrough and tremendous progress for reaching HTSC came with the discovery of the “Earth temperature superconductivity” at 203 K (-70 °C) in H_3S at ~ 150 GPa⁷. This achievement was based on the general assumption by Ashcroft, who suggested that hydrogen-dominant materials are promising candidates for HTSC⁸, which has then been greatly supported by modern ab initio computational methods for predicting the crystal structure and properties of novel materials^{9–11}. This work showed a clear route towards room-temperature superconductivity and initiated extensive high-pressure studies of hydrides, which are conventional phonon-mediated superconductors. Soon T_c reached 252 K in LaH_{10} at 170 GPa¹² ($T_c = 260$ K was claimed in Somayazulu et al.¹³), and very recently, $T_c = 287$ K was reported for the ternary carbon–sulfur–hydrogen system¹⁴. In addition, many other superconductors with high T_c s were found (see Review¹⁵ and the many refs. within).

While the claim of room-temperature superconductivity¹⁴ is waiting for its verification and clarification of the composition and crystal structure of the phase responsible for such high T_c , the superconductivity in H_3S and LaH_{10} compounds has been independently reproduced by several groups. For instance, the superconducting transitions in H_3S were confirmed with reported T_c s between ~ 183 and 200 K^{16–19}, and superconductivity in LaH_{10} was verified in close agreement at ~ 250 K^{13,20,21}.

Binary hydrides remain the most promising system for fruitful synergy between theory and experiment. Nevertheless, there is a discrepancy between the measured and predicted T_c s and phase diagrams, and new experimental data are crucial for further improvement of the computational methods. Recent theoretical reports provide convincing predictions for HTSC in the yttrium–hydrogen system. According to the calculations, T_c should be as high as 303 K at 400 GPa²² or 305–326 K at 250 GPa²³ in face-centred cubic (*fcc*) YH_{10} . In addition to YH_{10} , there are other phases with high calculated T_c s, which are predicted to be stable at lower pressures, e.g., hexagonal close packed (*hcp*) YH_9 with a T_c of 253–276 K stable at 200 GPa²² and body-centred cubic (*bcc*) YH_6 with a T_c of 251–264 K stable at 110 GPa²⁴.

In the present work, we report an experimental study of the yttrium–hydrogen system at high pressures and two superconducting *bcc*- YH_6 and *hcp*- YH_9 phases. Both phases have the same crystal structures as those predicted by the calculations^{22–24}, however, the observed T_c s are significantly lower than the calculated values by ~ 30 K. We could not synthesise *fcc*- YH_{10} , which has the highest computed T_c among the predicted yttrium hydrides. Instead, we revealed that *hcp*- YH_9 is the persistent hydrogen-richest yttrium hydride in the wide pressure-temperature domain up to 410 GPa and 2250 K – under the same conditions where the thermodynamic stability of the *fcc*- YH_{10} phase was predicted²². Here, we also discuss the contradictions between different experimental data for the yttrium–hydrogen system. Our preliminary report²⁵ and the present more comprehensive data are in good agreement with the data from

Troyan et al.²⁶ but evidently contradict the claim of $T_c = 262$ K at 182 GPa reported by Snider et al.²⁷.

Results and discussion

Synthesis of samples. We prepared various yttrium hydrides in a diamond anvil cell (DAC) by compressing either yttrium metal in H_2/D_2 (9 samples), yttrium trihydride YH_3/YD_3 in H_2/D_2 (9 samples), or YH_3 with NH_3BH_3 (7 samples). The details for the 31 different samples synthesised and analysed in the present study are summarised in Supplementary Table 1.

In our experiments, yttrium metal reacted with the surrounding hydrogen fluid at 17 GPa at room temperature, which was the lowest pressure at which X-ray diffraction patterns were collected (samples 26 and 27; Supplementary Fig. 1). This chemical reaction occurs even at lower pressures of ~ 1 GPa according to refs. 28,29.

Yttrium hydrides with a higher hydrogen content require much higher pressures for their formation. We synthesised body-centred tetragonal (*bct*) YH_4/YD_4 and *bcc*- YH_6/YD_6 within a pressure range of 160–175 GPa (Figs. 1 and 2) after heating YH_3/YD_3 in H_2/D_2 at ~ 1500 K with the aid of a pulsed laser (samples 5, 7 and 24). Both phases can also be formed by exposing the reactants to higher pressures of ~ 200 –244 GPa at room temperature for several weeks (samples 1, 2, 6, 9 and 10).

Both *bct*- YH_4/YD_4 and *bcc*- YH_6/YD_6 were observed within a wide pressure range of ~ 160 –395 GPa (Fig. 2). Laser heating of these phases with excess H_2/D_2 at pressures above 185 GPa results in the formation of the *hcp*- YH_9/YD_9 phase (samples 1, 2, 3, 6, 7, 10, 17–19 and 21–23).

X-ray diffraction studies. The crystal structures of novel yttrium hydrides were determined by X-ray powder diffraction. The Rietveld refinements for the typical powder diffraction patterns of the *I4/mmm* YH_4 , *Im-3m* YH_6 and *P6₃/mmc* YH_9 phases are shown in Fig. 1. The lattice parameters of these phases were determined at 135–410 GPa for a series of different samples and are listed in Supplementary Tables 1–5. The volumes of the crystal lattices of the synthesised yttrium hydrides are summarised in Fig. 2a as a function of pressure and are well approximated by the Vinet³⁰ equation of state (fitting parameters are given in Supplementary Table 6). The lattice volumes of *bct*- YH_4 , *bcc*- YH_6 and *hcp*- YH_9 (Fig. 2a) and the *c/a* ratios of *bct*- YH_4 and *hcp*- YH_9 (Fig. 2b) agree well with the theoretical predictions^{22–24,26,31,32}, which justifies the assignment of compositions for the hydrides with $\text{H}/\text{Y} > 3$.

We also independently estimated the compositions of new yttrium hydrides from the X-ray diffraction data by analysing the hydrogen-induced volume expansion V_{H} . In contrast to the lanthanum–hydrogen system¹², in which the V_{H} of ~ 1.8 Å³/H(D) atom is nearly identical in all lanthanum hydrides at approximately 150 GPa, the corresponding volume expansion is anomalously large in YH_3 and differs from that in the yttrium hydrides with higher hydrogen content. As a result, *bct*- YH_4/YD_4 and *fcc*- YH_3/YD_3 have almost the same lattice volumes within a pressure range of ~ 150 –250 GPa. Using the extrapolated equation of state for pure yttrium³³, the value of V_{H} in *bct*- $\text{YH}_4(\text{YD}_4)$ is ~ 1.7 – 1.8 Å³/H(D) atom at 180–200 GPa and is comparable with the values measured in other hydrogen-rich metal hydrides^{12,34–37}. The derived values provide the estimation for compositions of the new *bcc* and *hcp* phases as $\text{YH}_{5.7(3)}$ and $\text{YH}_{8.5(5)}$.

The synthesis of YH_{10} , which, according to theory, is the most promising candidate for the highest T_c of 305–326 K²³, was of particular interest for the present study. The calculations on the stability of *fcc*- YH_{10} provide contradictory results. Some

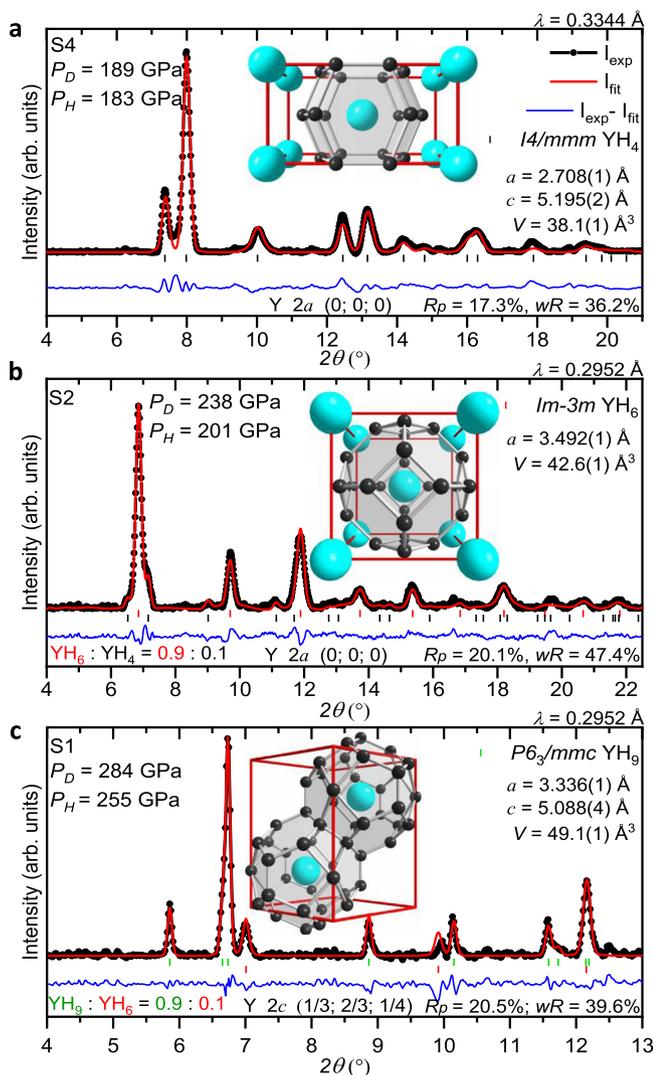


Fig. 1 X-ray powder diffraction patterns of the synthesised yttrium hydrides. **a** $I4/mmm$ YH_4 phase in sample 4 (S4) after pulsed laser heating at $P_H = 183$ GPa; **b** $Im-3m$ YH_6 phase in unheated sample 2 (S2) at $P_H = 201$ GPa with a T_c of ~ 211 K; **c** $P6_3/mmc$ YH_9 phase in sample 1 (S1) after pulsed laser heating at $P_H = 255$ GPa with a $T_c = 235$ K. See Supplementary Table 1 for details. P_H and P_D correspond to the pressures estimated by H_2 (D_2) vibron scale⁵⁴ and diamond scale⁵⁵, respectively. The black circles and red and blue curves correspond to the experimental data, Rietveld refinement fits and residues, respectively. The black, red and green ticks indicate the calculated peak positions for the $I4/mmm$ YH_4 , $Im-3m$ YH_6 and $P6_3/mmc$ YH_9 phases, respectively. The weight fractions for the phases, refined lattice parameters and coordinates for Y atoms are shown for each refinement. The fragments of the crystal structure with the characteristic YH_{18} , YH_{24} and YH_{29} coordination polyhedra (cages) are shown as insets. The large cyan and small black spheres show the positions of the Y and H atoms in the crystallographic unit cells according to Peng et al.²².

predictions show that this phase becomes stable at 250 GPa²³, which is an apparent contradiction with our observations. Other calculations suggest that at $T = 0$ K, YH_{10} is thermodynamically unstable at any pressure²². However, the difference in the Gibbs free energies between fcc - YH_{10} and hcp - YH_9 should decrease at higher temperatures, and fcc - YH_{10} should become more favourable at temperatures above 1500 K at 375 GPa or above 1100 K at 400 GPa²².

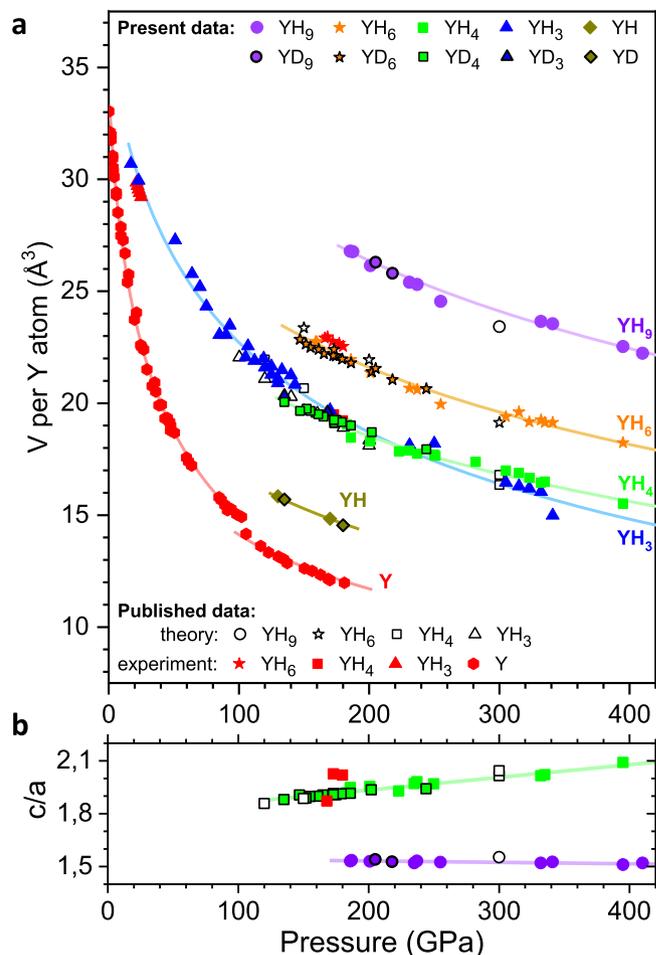


Fig. 2 Volume of various synthesised yttrium hydrides as a function of pressure. **a** The experimental data for the $P6_3/mmc$ YH_9 , $Im-3m$ YH_6 , $I4/mmm$ YH_4 , $Fm-3m$ YH_3 and $Fm-3m$ YH phases are shown as filled violet circles, orange stars, green squares, blue triangles and dark yellow rhombuses, respectively. The data for the corresponding yttrium deuterides are outlined in black. Open black circles, stars, squares and triangles correspond to the theoretically predicted structures for YH_9 ²², YH_6 ^{22-24,26}, YH_4 ^{22-24,26} and YH_3 ^{31,32}, respectively. Experimental data for pure Y ^{33,63}, YH_3 ⁵⁹, YH_4 ²⁶ and YH_6 ²⁶ taken from the literature are depicted by the red symbols. Solid curves correspond to the Vinet³⁰ equation of state fitting. The pressure was estimated from the frequency of the H_2 (D_2) vibron⁵⁴ for the samples with excess H_2 (D_2) and from the high-frequency edge of the Raman line from the stressed diamond anvils⁵⁵ for the remainder of our samples. **b** The pressure dependence for the lattice parameter ratio c/a in the $P6_3/mmc$ and $I4/mmm$ phases with linear fits.

Guided by these calculations, we prepared samples of YH_3 with NH_3BH_3 to study the yttrium-hydrogen system at very high pressures of ~ 325 – 410 GPa (see Supplementary Table 1). We did not observe fcc - YH_{10} in the final quenched products after pulsed laser heating of the samples up to 1600–2250 K. There was no hint of the fcc - YH_{10} phase immediately during pulsed laser heating; only the temperature-induced thermal expansion of the crystal structure of the hcp - YH_3 phase at 410 GPa and 2250(10) K was detected (Fig. 3). Notably, excess H_2 is hard to control in experiments with NH_3BH_3 as the source of hydrogen. Nonetheless, it is evident that excess H_2 was realised in sample 17 because the initial YH_3 completely transformed into single-phase $P6_3/mmc$ YH_9 after laser heating (Fig. 3). It is possible that the predicted YH_{10} exists at even higher pressures and temperatures,

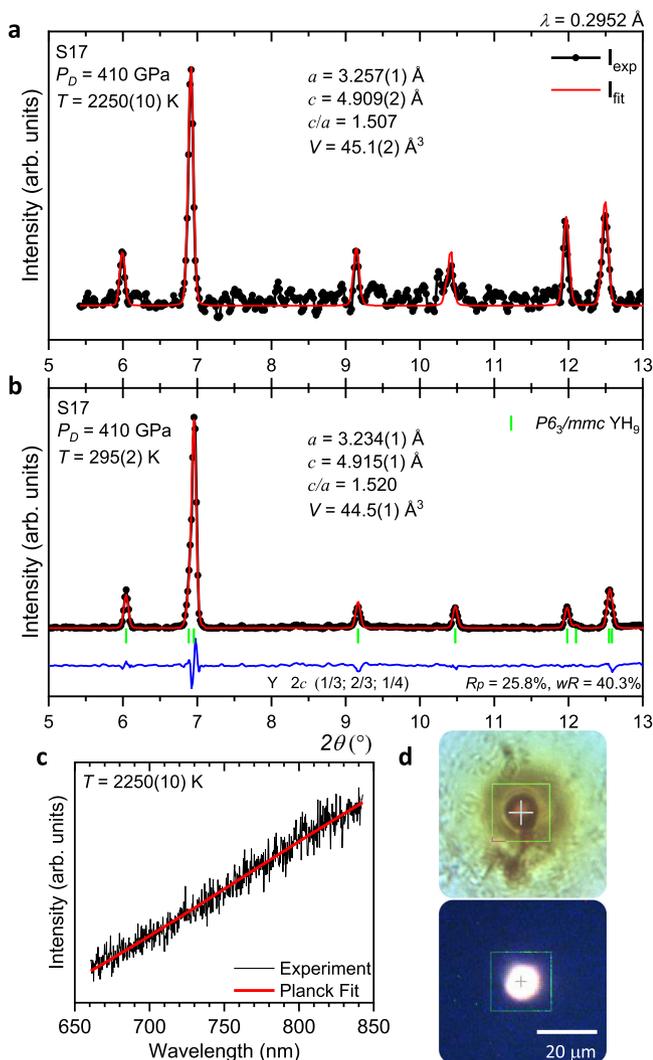


Fig. 3 X-ray diffraction study of the yttrium-hydrogen system at extreme pressure and temperature conditions. **a** X-ray powder diffraction pattern collected from sample 17 (S17) ($\text{YH}_3 + \text{NH}_3\text{BH}_3$) at 410 GPa directly at the moment of pulsed laser heating at 2250(10) K (black circles) and Le Bail refinement (red curve). **b** X-ray powder diffraction pattern of sample 17 at 410 GPa after subsequent quenching to ambient temperature, corresponding to the pure $P6_3/mmc \text{ YH}_9$ phase. Black circles, red and blue curves correspond to the experimental data, Rietveld refinement fits and residues, respectively. Green ticks indicate the calculated peak positions for the $P6_3/mmc \text{ YH}_9$ phase. **c** The thermal radiation spectrum measured during pulsed laser heating (black curve) and the fit to Planck's radiation law (red curve). **d** Photos of the sample at ambient temperature (top) and during pulsed laser heating (bottom).

but an experiment at such extreme conditions is currently challenging.

Superconductivity. The electrical resistance measurements of new yttrium hydrides revealed superconductivity with high T_c s (Fig. 4). The observed superconducting transitions were unambiguously assigned to either $hcp\text{-YH}_9/\text{YD}_9$ or $bcc\text{-YH}_6/\text{YD}_6$ by analysing the phase content in several of the prepared samples (see details in Supplementary Table 1). According to the X-ray diffraction data, some samples contained variable amounts of lower hydrides, namely, $fcc\text{-YH}_3$ and $bct\text{-YH}_4$, originating from the areas near the electrical leads. These areas were relatively poorly heated on purpose by the pulsed laser to prevent the

failure of the electrical leads on the samples. The presence of these impurities does not alter the observed HTSC in our samples (see below).

The T_c of $hcp\text{-YH}_9$ is higher than that of the $bcc\text{-YH}_6$ phase, as follows from the electrical measurements for sample 2 (the black and red curves in Fig. 4a). Prior to laser heating, the sample contained $bcc\text{-YH}_6$ and exhibited a superconducting transition with a T_c of ~ 211 K at 201 GPa. After heating at 2000(10) K, most of the $bcc\text{-YH}_6$ phase transformed into $hcp\text{-YH}_9$, and the superconducting transition shifted to a higher temperature of ~ 243 K (the X-ray diffraction patterns of sample 2 are shown in Supplementary Fig. 2d, e). Identical behaviour was observed for the deuterides in sample 6 at 202 GPa (the black and red curves in Fig. 4b); i.e., the initial $bcc\text{-YD}_6$ phase had a T_c of 165 K, and $hcp\text{-YD}_9$, which formed after laser heating, exhibited a higher T_c of 172 K (the X-ray diffraction patterns of sample 6 are presented in Supplementary Fig. 3).

The pressure dependencies of T_c for YH_9 and YH_6 have a “dome-like” shape with the highest measured T_c of 243 K at 201 GPa and 224 K at 166 GPa, respectively (Fig. 4e). Similar maxima at the $T_c(P)$ dependence were previously observed in H_3S^7 and LaH_{10}^{12} at ~ 150 GPa. The decrease in T_c with increasing pressure in the $bcc\text{-YH}_6$ and $hcp\text{-YH}_9$ phases is likely due to either the pressure-induced stiffening of the phonon frequencies similar to that in $bcc\text{-H}_3\text{S}^{38,39}$ or the presence of a flat region on the Fermi surface and the appearance of a two-gap structure similar to that in $fcc\text{-LaH}_{10}^{40,41}$. The data for $bcc\text{-YH}_6$ and YD_6 independently measured by Troyan et al.²⁶ agree with this trend. The sharp drop in T_c for YH_9 and YH_6 at pressures below ~ 195 and ~ 165 GPa, respectively, should be associated with the structural distortions and phase transformations between the high-pressure high-symmetry and low-pressure low-symmetry phases, as was recently demonstrated for $\text{H}_3\text{S}^{18,42}$ and $\text{LaH}_{10}^{21,43}$.

It should be noted that some samples synthesised by keeping reactants at room temperature for several weeks without laser heating (samples 1, 2, 6 and 10) exhibit T_c s that are lower by $\sim 5\text{--}10$ K in comparison to the samples prepared via laser heating-assisted synthesis (the corresponding symbols are outlined by the red circles in Fig. 4e, f). This effect should be attributed to the poorer crystallinity and homogeneity of the superconducting phase in the non-annealed samples, which manifested in the broadening of the Bragg reflections in the X-ray diffraction powder patterns (Supplementary Figs. 2 and 3). Similar behaviour was previously shown in $\text{H}_3\text{S}^{7,18}$.

Isotope effect. The substitution of hydrogen by deuterium in the samples resulted in a pronounced shift in T_c to lower temperatures. The transition temperature shifted to ~ 168 K for YD_6 in sample 7 at 173 GPa and ~ 172 K for YD_9 in sample 6 at 205 GPa (Fig. 4c, d). This isotope effect supports the conventional phonon-assisted mechanism of superconductivity. Using the T_c values measured for $hcp\text{-YH}_9/\text{YD}_9$ and $bcc\text{-YH}_6/\text{YD}_6$ within the pressure range of 183–205 GPa (samples 2, 4, 6 and 7), we calculated the isotope effect coefficient, $\alpha = -\frac{d\ln T_c}{d\ln M}$, where M is the atomic mass, to be 0.39 for the $bcc\text{-YH}_6/\text{YD}_6$ phase and 0.50 for the $hcp\text{-YH}_9/\text{YD}_9$ phase. The isotope coefficient value for $bcc\text{-YH}_6/\text{YD}_6$ is smaller than the maximal expected BCS value of 0.5 for a harmonic case. This result likely stems from the anharmonic effects and the contribution of acoustic phonons to electron-phonon coupling.

Resistance measurements under high magnetic fields. In addition to the observed drops in the resistance to a zero value and the isotope effect, the onset of superconducting order was

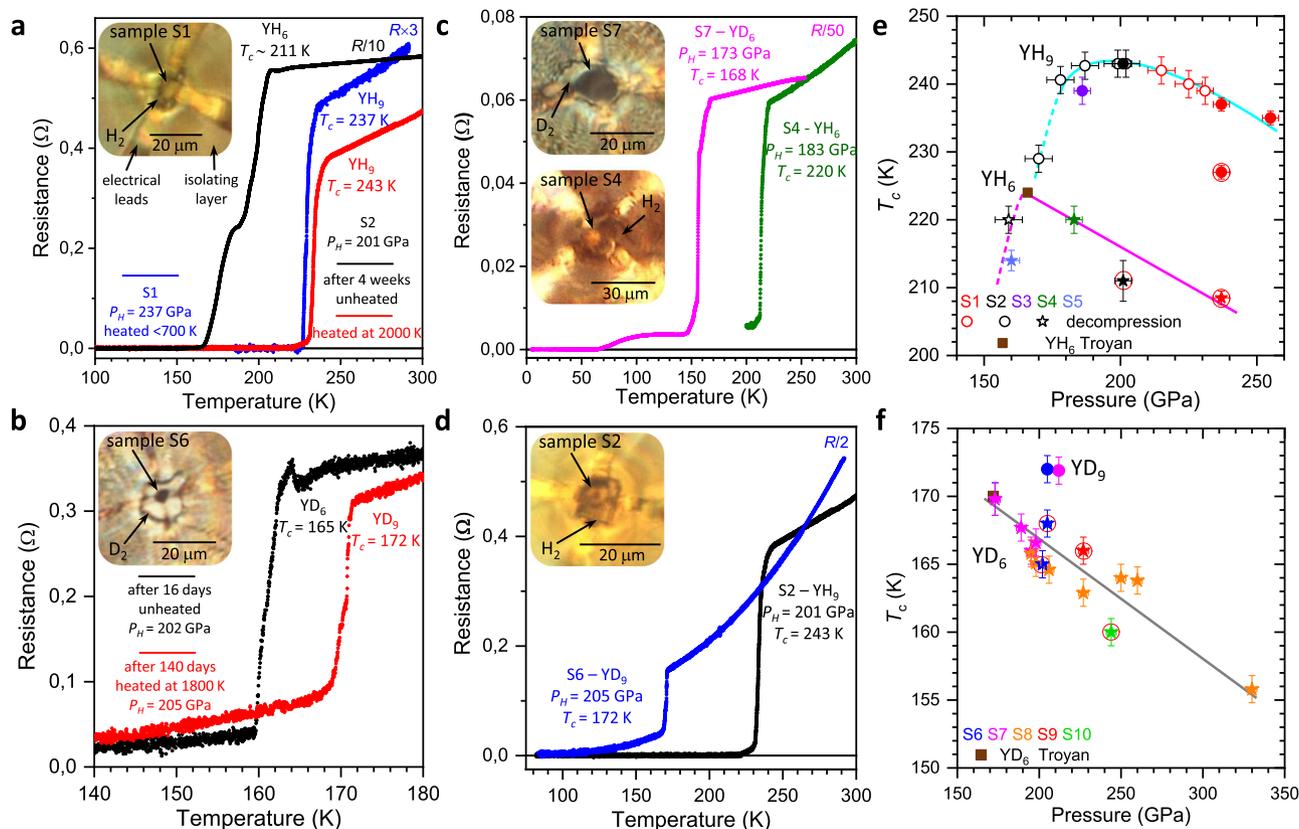


Fig. 4 Superconducting transitions in the synthesised yttrium hydrides and deuterides. **a, b** The temperature dependencies of the resistance of the yttrium-hydrogen and yttrium-deuterium samples measured with a four-probe technique in a van der Pauw geometry, demonstrating the shift in T_c to higher temperatures after pulsed laser heating of the samples. In **a**, Black and red curves correspond to the temperature dependence of the resistance of the $Im-3m$ YH_6 phase, which was formed after exposing YH_3 to H_2 at $P_H = 201$ GPa for 3 weeks, and the $P6_3/mmc$ YH_9 phase, which was synthesised after subsequent heating at 2000(10) K, in sample 2. The blue curve corresponds to the $P6_3/mmc$ YH_9 phase synthesised at $P_H = 237$ GPa in sample 1 (S1). In **b**, similarly, T_c increases from 165 K for $Im-3m$ YD_6 to 172 K for $P6_3/mmc$ YD_9 in sample 6 (S6) at $P_H \sim 205$ GPa. **c, d** The temperature dependencies of the resistance of $Im-3m$ YH_6 (YD_6) and $P6_3/mmc$ YH_9 (YD_9) synthesised in samples 2, 4, 6 and 7 (S2, S4, S6, S7) demonstrate the shift in the superconducting transitions with isotopic substitution. The absolute resistance values for some samples were multiplied by the specified constant factors for better presentation. The insets show photos of the samples and arrangements for electric transport measurements. **e, f** The pressure dependence of T_c for the superconducting transitions in $Im-3m$ YH_6 (stars) and $P6_3/mmc$ YH_9 (circles) phases and the corresponding deuterides, respectively. Different colours represent different samples. Open symbols are the data obtained on subsequent decompression. Symbols marked by red circles are the data for unheated samples. Cyan and grey curves are the guides for the eye. Brown squares depict the data from Troyan et al.²⁶. Error bars are defined the same as in Drozdov et al.¹². The horizontal and vertical error bars correspond to the uncertainty in the precise value of the pressure (inherent error bars of the method used) and in the determination of the correct value of T_c (criteria-dependent), respectively.

independently verified by the magneto-transport measurements under magnetic fields up to 9T. While the magnetic field has a negligible effect on the resistance of the normal metal, T_c is strongly reduced as the magnetic field increases, and superconductivity is completely suppressed at fields above the upper critical field H_{c2} . H_{c2} is the most direct probe of the coherence length of the superconducting order parameter $\xi = \sqrt{\frac{\phi_0}{2\pi H_{c2}}}$, where ϕ_0 is the magnetic flux quantum. Figure 5a, b show the dependence of the superconducting transition in samples 5 and 6 as a function of external magnetic field. To estimate the H_{c2} and ξ at zero temperature, we plotted the dependence of T_c on the applied external magnetic field, following the criterion of 90% of the resistance in the metallic state (Fig. 5c). The temperature dependence of H_{c2} can be approximated by the Ginzburg–Landau (GL) equation⁴⁴ for $\frac{T_c - T}{T_c} \ll 1$ and more accurately by the Werthamer–Helfand–Hohenberg (WHH)⁴⁵ model for all temperatures. The light and dark curves in Fig. 5c show the results of the fits for the experimental values of $H_{c2}(T)$ to the GL and WHH relations. These fits yield $H_{c2}^{WHH}(0\text{ K}) = 60\text{ T}$ ($H_{c2}^{GL}(0\text{ K}) = 46\text{ T}$)

for hcp - YD_9 and $H_{c2}^{WHH}(0\text{ K}) = 157\text{ T}$ ($H_{c2}^{GL}(0\text{ K}) = 107\text{ T}$) for bcc - YH_6 . The latter values are in good agreement with the estimate of $H_{c2}(0\text{ K}) = 116\text{--}158\text{ T}$ by Troyan et al.²⁶. The corresponding coherence length ξ (0 K) in bcc - YH_6 and hcp - YD_9 are 1.45–1.75 nm and 2.3–2.7 nm, respectively.

We estimated $H_{c2}^{WHH}(0\text{ K}) = 120\text{ T}$ ($H_{c2}^{GL}(0\text{ K}) = 92\text{ T}$) for hcp - YH_9 using the relation of $H_{c2} \sim (\frac{T_c}{v_F})^2$ and assuming the same Fermi velocities v_F in YH_9 and YD_9 counterparts. Generally, higher T_c values correlate with higher $H_{c2}(0\text{ K})$ values in the studied hydride superconductors, e.g. $H_{c2} = 144\text{ T}$ and $T_c = 250\text{ K}$ were observed in LaH_{10} ^{12,21}, $H_{c2} = 88\text{ T}$ and $T_c = 197\text{ K}$ in H_3S ¹⁶, $H_{c2} = 45\text{ T}$ and $T_c = 153\text{ K}$ in ThH_{10} and $H_{c2} = 38\text{ T}$ and $T_c = 145\text{ K}$ in ThH_9 ³⁶, $H_{c2} = 29\text{ T}$ and $T_c = 98\text{ K}$ in CeH_{10} ⁴⁶, and $H_{c2} = 11\text{ T}$ and $T_c = 68\text{ K}$ in SnH_x ⁴⁷. Conversely, the hcp - YH_9 phase has higher $T_c = 243\text{ K}$ but lower H_{c2} comparing with bcc - YH_6 phase. This is likely caused by the difference in electronic band structure in these phases, and further theoretical calculations are required to explain this anomaly.

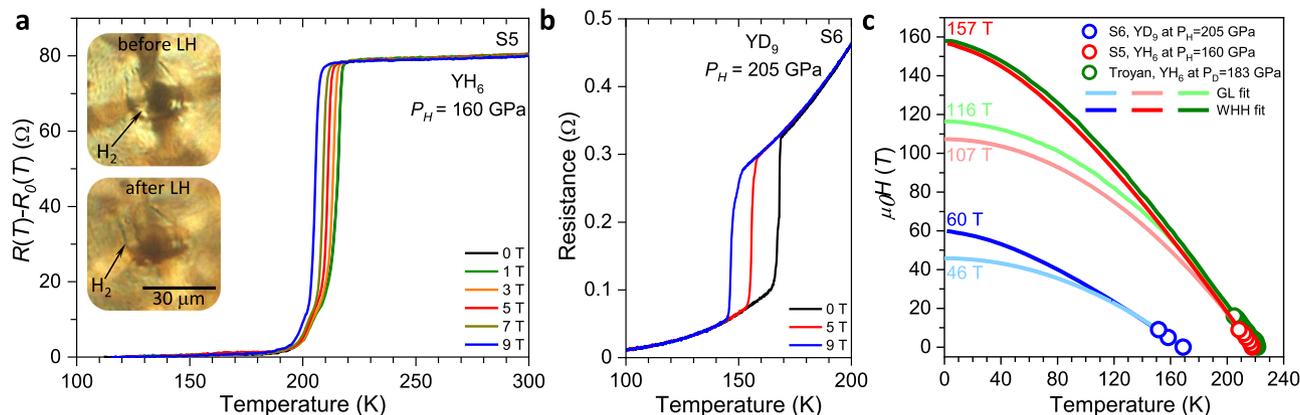


Fig. 5 Temperature dependence of the resistance for the *Im-3m* YH_6 and *P6₃/mmc* YD_9 phases under external magnetic field. **a, b** DC field measurements for the *Im-3m* YH_6 phase at $P_H = 160$ GPa in sample 5 (S5) and the *P6₃/mmc* YD_9 phase at $P_H = 205$ GPa in sample 6 (S6), respectively. **c** Fits of the superconducting upper critical field H_{c2} to the Werthamer-Helfand-Hohenberg (WHH) and Ginzburg-Landau (GL) formalisms. Red and blue circles denote the H_{c2} s measured for *Im-3m* YH_6 at $P_H = 160$ GPa and *P6₃/mmc* YD_9 phase at $P_H = 205$ GPa, respectively. The dark and light curves are the WHH and GL fits to the experimental data. Green circles and green curves are the data for *Im-3m* YH_6 phase at $P_D = 183$ GPa from Troyan et al.²⁶.

We found that despite a substantial difference in the $H_{c2}(0\text{ K})$ values for YD_9 and YH_6 samples, the Fermi velocities v_F estimated via the BCS relation: $\xi = 0.18 \frac{h v_F}{k_B T_c}$ are quite similar, i.e. 2.85×10^5 and 2.3×10^5 m/s, respectively. Similar v_F values were reported for other superconductors of so-called “superhydride” family including SnH_x ⁴⁷ and LaH_{10} ²¹. This indicates that the dispersion of the charge carriers contributing to the superconductivity does not significantly change between different superconducting hydrides. Interestingly, nearly constant values of v_F were also revealed for various unconventional high-temperature superconductors of the cuprates family⁴⁸.

Stability range of yttrium hydrides. The predicted crystal structures of the YH_3 , YH_4 , YH_6 and YH_9 compositions are in excellent agreement with the experimental results. To assess the accuracy of the calculations, we compared the predicted formation pressures for these novel hydrides with our experimental observations. The problem of determining the equilibrium pressure in experiments is often exacerbated by the presence of large baric hysteresis between the formation and decomposition pressures. It is generally accepted that the equilibrium pressure in most metal-hydrogen systems is much closer to the decomposition pressure of the high-pressure phase^{49,50}. Decreasing pressure in sample 2 resulted in the decomposition of YH_9 into *bcc*- YH_6 at 159 GPa (Supplementary Fig. 2f), which is considerably higher than the predicted YH_9 equilibrium formation pressure of 100 GPa²². The sharp drop in T_c for *hcp*- YH_9 at ~ 185 GPa (open black star in Fig. 4e) indicates that this phase is dynamically unstable below this pressure. Decreasing pressure in sample 24 resulted in a decomposition of *bcc*- YD_6 at ~ 135 GPa, whereas *bct*- YD_4 was stable down to at least ~ 135 GPa (Supplementary Tables 1 and 3). This result is in reasonable agreement with the predicted equilibrium formation pressure of 110 GPa for both phases²⁴.

The predicted T_c s of 251–264 K for *bcc*- YH_6 ²⁴ and 253–276 K²² for the *hcp*- YH_9 phase are ~ 30 K higher than the present experimental values. Recent calculations²⁶ for the *bcc*- YH_6 phase, which accounted for the anharmonicity, demonstrate a T_c of 236–247 K, which is still significantly higher than the experimental value. In addition, we did not observe superconductivity in *fcc*- YH_3 upon cooling down to 5 K in the pressure range of 15–180 GPa or in *bct*- YH_4 upon cooling down to 78 K at 250 GPa (Supplementary Fig. 4), while both phases were predicted to be

superconductors with T_c s of 40 K⁵¹ and 84–95 K^{24,32}, respectively.

Comparison with other works. In the recent report, Snider et al.²⁷ claimed a significantly higher maximum $T_c = 262$ K at 182(8) GPa in the yttrium-hydrogen system. Their values of T_c s measured in the pressure range of ~ 134 –187 GPa and the pressure dependence of T_c contradict the results of both the present work and Troyan et al.²⁶ (Supplementary Fig. 5). This strong disagreement raises a question about the material studied in Snider et al.²⁷. Unfortunately, the lack of any X-ray structural characterisation of the samples in Snider et al.²⁷ makes the direct comparison impossible. The superconductivity was putatively assigned to *P6₃/mmc* YH_9 based on a comparison of the measured and computed Raman spectra²⁷. However, Raman spectroscopy is not a reliable method for the identification of a crystal structure. Moreover, *bct*- YH_4 , *bcc*- YH_6 and *hcp*- YH_9 are good metals and could not account for the observed Raman spectra. Presently, we observed Raman spectra only for mixtures with a hydrogen-depleted *fcc* phase with a composition close to YH_1/YD_1 , which was formed during the compression of the initial $\text{YH}_{2.92(5)}$ and $\text{YD}_{2.87(5)}$ above ~ 100 GPa (see Methods section for details and Supplementary Fig. 6). Furthermore, the superconducting transitions measured in Snider et al.²⁷ below ~ 185 GPa cannot be assigned either to *hcp*- YH_9/YD_9 because it is unstable in this pressure range or to *bcc*- YH_6 because this phase has a significantly lower T_c .

Summary. We report on the superconductivity in *hcp*- YH_9 with a maximum T_c of 243 K at 201 GPa, which is the second highest T_c measured for the family of transition element superhydrides, and *bcc*- YH_6 with a T_c of 220 K at 183 GPa. At higher pressure, both phases demonstrate a decrease in T_c . The decrease in T_c under external magnetic fields additionally confirms the superconductivity in novel yttrium superhydrides, and the isotopic shift in the superconducting transition in deuterides to lower temperatures supports the conventional phonon-assisted mechanism of superconductivity. We found good agreement between the predicted and experimental crystal structures and the $V(P)$ dependencies of the synthesised hydrides. However, the measured T_c s for *bcc*- YH_6 and *hcp*- YH_9 are markedly lower than the computed values of 251–264 K for YH_6 ²⁴ and 253–276 K for YH_9 ²². We did not find the *fcc*- YH_{10} phase despite extensive trials at pressures up to 410 GPa and temperatures up to 2250 K.

Methods

Diamond anvil cell. Typically, DACs have diamonds bevelled at 8° to a diameter of ~250 µm with a culet size of ~15–35 µm. The diamond anvils had a toroidal profile for the samples pressurised over ~200 GPa, which was machined by a focused beam of xenon ions (FERA3, Tescan). Four tantalum or tungsten leads covered by gold were deposited onto the surface of one diamond anvil in a van der Pauw geometry. A metallic gasket (T301 stainless steel) was thoroughly isolated from the sputtered leads by a non-conducting layer prepared from a mixture of low-viscosity epoxy resin and a fine powder of either CaF₂, CaO, MgO, CaSO₄, cBN or Al₂O₃. The insulating gasket was pressed to a thickness of 3–5 µm, and a hole with a diameter of ~2/3 of the culet size was drilled by a pulsed ultraviolet laser.

Preparation of samples. Yttrium hydrides were synthesised in situ in a DAC via a direct reaction between either metallic yttrium (99.9%, Sigma Aldrich) and H₂ (99.999%) (D2, 99.75%, Spectra Gases) or, alternatively, YH₃ (YD₃) and H₂ (D₂) at pressures up to ~250 GPa. As an alternative source of hydrogen, NH₃BH₃ (97%, Sigma Aldrich) was used at pressures of 250–410 GPa. The Y or YH₃ (YD₃) pieces were typically 5–15 µm in diameter and 1–2 µm thick. The samples were handled in an inert Ar or N₂ atmosphere with residual O₂ and H₂O contents of <0.1 ppm to prevent oxidation. The procedure of hydrogen gas clamping and laser heating-assisted synthesis was the same as that for lanthanum hydrides¹². One-side heating of the sample was performed with the aid of a pulsed YAG laser. Elevated temperatures accelerate the diffusion of hydrogen into the metal; however, hot hydrogen can easily break diamond anvils by penetrating deep into microcracks at the surface of diamond. We avoided this by the rigorous polishing and etching of diamonds.

All samples synthesised and studied in the present work are summarised in Supplementary Table 1.

The YH₃ and YD₃ samples used as the initial materials in the DAC experiments were synthesised using bulk yttrium metal that was preliminarily annealed in a vacuum of ~10⁻³ Torr at 400 °C and then exposed to H₂(D₂) gas at a pressure of ~100 bars at 400 °C for 4 h and then at 200 °C for 24 h in a high-pressure Sievert-type apparatus⁵². According to a weighting method, the products were YH_{2.92(5)} and YD_{2.87(5)}. The samples were powdered and analysed with an Empyrean X-ray diffractometer in an inert atmosphere under ambient conditions. These materials consisted of pure single-phase *hcp*-YH₃ and YD₃ (Supplementary Fig. 7). The lattice parameters of both products were in agreement with published data⁵³. For brevity, these materials are referred to as YH₃ and YD₃ throughout the paper.

Electrical transport measurements. DC electrical measurements were performed on cooling and warming cycles with an electrical current of 10⁻⁵–10⁻³ A (Keithley 6220 and 2000). The present data were taken upon warming as it yields a more accurate temperature reading; that is, the cell was warmed up slowly (0.2 K min⁻¹) in a quasi-isothermal environment without coolant flow. The temperature was measured with an accuracy of ~1 K by a Si diode thermometer (Lakeshore DT-470) attached to the DAC body. The T_c was determined from the onset of superconductivity at the point of the apparent deviation of the temperature dependence of the resistance from the normal metallic behaviour.

Alongside the standard stainless steel DACs, special types of DACs with external diameters of 20 mm and 8.8 mm made of non-magnetic materials were used for measurements under external magnetic fields using a 9T Quantum Design Physical Property Measurement System (PPMS).

Estimation of pressure. The pressure in the DACs was estimated using the H₂ (D₂) vibron scale⁵⁴ if the corresponding vibron could be observed in the Raman spectra or diamond scale⁵⁵ based on the shift of the Raman line edge of stressed diamond and marked throughout the text as P_H and P_D, respectively. Typically, the second scale provides overestimated pressure values by ~5–40 GPa, which is a result of a large pressure gradient between the soft H₂/D₂ medium and the surrounding harder gasket. Unless otherwise stated, the pressure values displayed in the figures were estimated using the H₂ (D₂) vibron scale. Additionally, the pressure in samples 22, 26 and 27 was estimated using the equation of state of MgO⁵⁶, which served as a gasket material.

X-ray diffraction measurements. X-ray diffraction data were collected at beamline 13-IDB at GSECARS, Advanced Photon Source, using λ₁ = 0.2952 Å and λ₂ = 0.3344 Å, a beam spot size of ~2.5 × 2.5 µm², and a Pilatus 1 M CdTe detector. A typical exposure time varied between 10 and 300 s. To examine the formation of new yttrium hydrides at 325–410 GPa at high temperatures, we collected X-ray powder diffraction patterns in situ at high temperature. Each pattern was collected by accumulating 5 × 10⁵ shots with a duration of 1 µs, which were synchronised with laser heating pulses. The temperature was determined by the thermal emission from the sample measured with a PI-MAX3 detector. Primary processing and integration of the powder patterns were performed using Dioptas software⁵⁷. The indexing of powder patterns and refinement of the crystal structures were done with the GSAS and EXPGUI packages⁵⁸.

YH₃ and other phases formed under hydrogen deficiency. In separate experiments, we characterised YH₃ and YD₃, which were the starting materials for the synthesis of higher hydrides in our study, during compression up to 180 GPa. *Hcp*-

YH₃ is the yttrium hydride with the highest hydrogen content under ambient conditions. It is a black narrow-bandgap semiconductor with a metallic lustre. At increasing pressure, *hcp*-YH₃ undergoes a continuous phase transition at ~10–25 GPa into *fcc*-YH₃^{52,59,60}. A further pressure increase results in continuous metallisation, which is accompanied by a disappearance of the Raman spectrum at ~80 GPa and a significant decrease in electrical resistance from ~50 Ω at 16 GPa to ~0.12 Ω at 81 GPa (Supplementary Fig. 8). This behaviour agrees well with previous measurements⁶¹. YH₃ and YD₃ retain the *fcc* metal sublattice upon compression up to ~150 GPa (samples 12–13 and 26–30), in agreement with theoretical calculations³¹. However, we observed the formation of another *fcc* phase in addition to *fcc*-YH₃(YD₃) at pressures above ~100 GPa during the compression of the initial YH_{2.92(5)} and YD_{2.87(5)} samples (Supplementary Fig. 6). The formation of this new phase is accompanied by the appearance of a strong Raman spectrum. Its lattice volume is smaller by ~5 Å³ per Y atom than that of YH₃/YD₃, likely indicating a reduced hydrogen content close to YH₁/YD₁. We did not observe this hydrogen-depleted phase in samples compressed in H₂ or D₂ medium (samples 25, 27–29). Thus, the formation of the *fcc*-YH₁/YD₁ phase is likely driven by the non-stoichiometric composition of the initial materials. A similar phenomenon was previously observed in substoichiometric LaH_{2.3} upon compression in an inert medium, and it was attributed to a disproportionation reaction into the hydrogen-enriched stoichiometric LaH₃ and hydrogen-depleted solid solution⁶².

In addition to the *I4/mmm* YH₄, *Im-3m* YH₆ and *P6₃/mmc* YH₉ phases discussed in detail in the main text, we observed some unidentified impurity phases with complex X-ray powder diffraction patterns. Typical X-ray diffraction powder patterns of unidentified impurities are plotted in Supplementary Fig. 9a. Since such phases were found in samples with an evident deficiency of H₂ (D₂) or in the poorly heated areas of samples, their H/Y ratio is likely <9. Troyan et al.²⁶ also found some new phases at pressures of 165–180 GPa and assigned them to the YH₇ and Y₂H₁₅ hydrides. None of these phases fit the reflections from the unidentified phases observed in the present study.

In sample 28 at a lower pressure of 105 GPa, we found a new *bcc* phase with a composition close to YH₄ (Supplementary Fig. 9b, c).

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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Author contributions

M.I.E. guided the work. P.K., V.S.M., M.A.K., A.P.D. and S.P.B. prepared the samples and measured the superconducting transition. V.S.M., M.A.K. and V.B.P. performed X-ray diffraction studies with the assistance of S.C. and E.G.; the data were processed by V.S.M. and M.A.K.; S.M., L.B. and F.F.B. performed studies under external magnetic fields. D.A.K. assisted in the preparation of diamond anvils. V.S.M., M.I.E. and M.A.K. wrote the manuscript with input from all co-authors.

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