

# Anisotropic Dependence of Superconductivity on Uniaxial Pressure in CeIrIn<sub>5</sub>

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We measure the effect of uniaxial pressure on the superconducting transition temperature  $T_c$  in CeIrIn<sub>5</sub>. We find a linear change in  $T_c$  with both  $a$ -axis and  $c$ -axis pressure, with slopes of 56 mK/kbar and -66 mK/kbar, respectively. By comparing results from doping studies and different types of pressure measurements, we separate the influences of hybridization and dimensionality on  $T_c$ . We find the true geometric influence, for constant hybridization, is  $\partial T_c / \partial (c/a) = 44$  K.

The low-temperature phases of heavy fermion materials have attracted much attention in recent years. They exhibit a range of correlated phases, including several types of magnetism. Superconducting regimes emerge near zero-temperature magnetic phase transitions. Non-Fermi liquid behavior also appears near these quantum critical points, and can persist to significant temperatures. Tuning through alloying, pressure, or applied field allows exploration of the exact balance among the phases.

One of the most-studied heavy fermions recently has been CeMIn<sub>5</sub> (M = Ir, Rh, Co). The proximity of these Ce-based 115 compounds to an antiferromagnetic quantum critical point leads to a rich phase diagram [1]. CeRhIn<sub>5</sub> at ambient pressure has a superconducting transition near 0.1 K, deep within an antiferromagnetic phase [2]. Hydrostatic pressure destroys the magnetism and raises  $T_c$  to a maximum of 2.1 K at 16 kbar [3, 4]. On the other hand, CeIrIn<sub>5</sub> and CeCoIn<sub>5</sub> superconduct at ambient pressure [5, 6], with several similarities to the high-temperature cuprate superconductors. The 115 materials have a tetragonal, HoCoGa<sub>5</sub> crystal structure which can be viewed as alternating layers of CeIn<sub>3</sub> and MIn<sub>2</sub> stacked along the (0 0 1) direction [3, 5, 6], reminiscent of the copper-oxygen planes in the cuprates. Power-law temperature dependences in heat capacity, thermal conductivity and spin-lattice relaxation rate in the superconducting state suggest  $d$ -wave symmetry of the superconducting order parameter [7]. Furthermore, emergence of superconductivity near  $T_N$  and coexistence of homogeneous antiferromagnetism and superconductivity provide evidence that magnetic fluctuations mediate Cooper-pairing in these systems [8, 10, 11].

One major influence on the superconducting transition temperature is the hybridization between the Ce  $f$ -electrons and In  $p$ -electrons [12, 13, 14], which controls the spin-fluctuation temperature,  $T_{sf}$ .  $T_{sf}$  is proportional to  $T_c$  and roughly inversely proportional to  $\gamma$ , the Sommerfeld coefficient of the normal-phase specific heat. Smaller  $\gamma$ 's in the isostructural PuMGa<sub>5</sub> compounds (M = Rh, Co) partially account for their much higher superconducting transition temperatures. However, even within the CeMIn<sub>5</sub> series,  $\gamma$  alone cannot describe all the variation in  $T_c$ ; for example, while  $\gamma$  changes by just over a factor of 2 between M = Ir and M = Co,  $T_c$  changes

by nearly a factor of 6. Thus hybridization cannot be the sole influence on  $T_c$ .

Mean-field theoretical models of magnetically mediated superconductivity indicate a strong dependence of  $T_c$  on dimensionality [15, 16, 17]. Measurements on CeM<sub>1-x</sub>M'<sub>x</sub>In<sub>5</sub> show a linear relationship between  $T_c$  and the ratio of the tetragonal lattice constants  $c/a$  [8]. Interestingly, for the Pu-based 115 materials  $T_c$  is also linear in  $c/a$ , with the same relative change in  $T_c$  with dimensionality,  $\frac{1}{T_c} \frac{dT_c}{d(c/a)}$  [12]. The agreement in slopes makes sense if the difference in hybridization sets the overall temperature scale for each family but dimensionality governs the behavior within each family. Another way to control dimensionality and hybridization is by applying pressure. Under hydrostatic pressure,  $c/a$  is not even monotonic for CeRhIn<sub>5</sub> and CeCoIn<sub>5</sub>, and  $T_c$  is not linear in  $c/a$ . However, all the CeMIn<sub>5</sub> compounds have similar hybridization at the pressure  $P_{max}$  which maximizes the  $T_c$ . Considering  $T_c$  and  $c/a$  at  $P_{max}$ , where the effects of hybridization differences are reduced, does give a linear relationship [13, 18]. Kumar *et al.* argue that hydrostatic pressure mainly alters the hybridization. The similar hybridizations at  $P_{max}$  suggest that hybridization determines the pressure  $P_{max}$  while dimensionality governs the value of  $T_c$  at  $P_{max}$  [13].

Uniaxial pressure is a natural technique for further exploring the effects of dimensionality, since the  $c/a$  ratio of an individual sample can be increased or decreased depending on the pressure axis. Uniaxial pressure leads to fairly small changes in hybridization, since lattice constants decrease along the direction of applied force but increase in the perpendicular directions. On the other hand, the effects on  $c/a$  are much larger than for a similar hydrostatic pressure. To date, the effects of uniaxial pressure on the 115 materials have not been measured directly, although thermal expansion measurements combined with Ehrenfest relations predict the change in  $T_c$  with uniaxial pressure in the zero pressure limit [19]. Here we explore the dependence of  $T_c$  on pressure and hence on  $c/a$ , and combine our results with those on hydrostatic pressure and alloying to extract the dependence of  $T_c$  on dimensionality.

Samples were grown in alumina crucibles from a molten metal flux containing stoichiometric amounts of Ce and Ir,

and excess indium. We confirmed through transmission Laue X-ray diffraction that the lattice constants of our samples matched those reported elsewhere [20].

Our uniaxial pressure apparatus, shown in Figure 1, is a helium-activated bellows mounted on a dilution refrigerator [21], and permits changes in pressure without thermally cycling the sample. We monitor the pressure through a piezo-electric crystal in the pressure column. The maximum achievable pressure depends on the size of the sample, but is typically about 10 kbar. We measure the superconducting transition with adiabatic heat capacity. Superconducting NbTi spacers between the sample and the pressure cell serve as the thermal link.

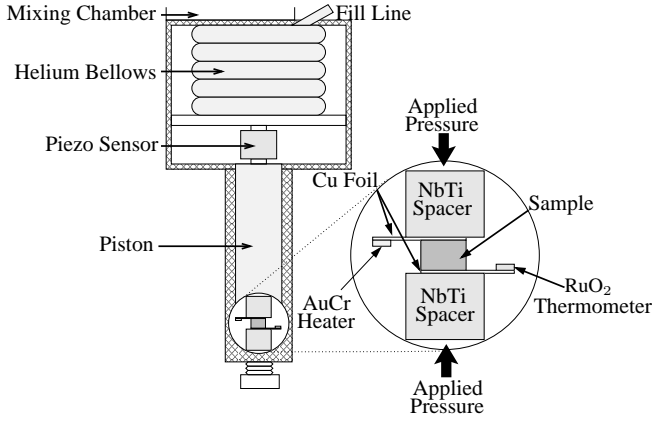


FIG. 1: Helium bellows setup for measuring heat capacity under uniaxial pressure.

As can be seen from Figure 1, the sample is unconstrained in directions perpendicular to applied pressure. To apply pressure along a specific axis, we first orient our sample using Laue X-ray diffraction. We then polish the sample according to the desired orientation. We preserve as much bulk as possible during the polishing, since the observed time constant for thermal decay is proportional to the sample's mass. Our largest samples for pressure along the  $c$  and  $a$  axes were approximately 30 mg and 80 mg, respectively.

Figure 2 shows representative heat capacity data. Pressure applied along the  $c$  axis shifts the transition to lower temperature, while  $a$ -axis pressure has the opposite effect. The data are scaled, with a single scaling constant used for each sample, regardless of pressure. The average  $C/T$  over all pressures at 800 mK is set to 700 mJ/mol K<sup>2</sup>.

For  $a$ -axis pressure, we fit a function  $\gamma_s + a_1 T^n + a_2 T^{-3}$  to the heat capacity in the superconducting phase [5]. The form fits well at all pressures, with the exponent  $n$  always close to 1, as expected for line nodes in the energy gap. For  $c$ -axis pressure, the reduced  $T_c$  leaves too small a temperature range for reliable fits. If we fix  $n = 1$  and refit the data, we find a slight decrease in both  $\gamma_s$  and  $a_1$  with pressure, as opposed to the sharp change in  $\gamma_s$  observed with uniaxial [21] and hydrostatic [22, 23] in other heavy fermion superconductors.

We define  $T_c$  as the temperature where  $C/T$  equals the average of its maximum value in the superconducting region

and its value in the normal phase at the onset of the transition. We also model the transition as an abrupt discontinuity with the restriction that the normal-phase entropy is the same as for the actual data. The two methods give consistent values of  $T_c$ . However, under  $c$ -axis pressure  $T_c$  decreases until eventually there is not enough data in the superconducting region to extrapolate the value of  $C/T$  in an equal-entropy calculation. For consistency between  $c$  and  $a$ -axis pressure, all values we report here use the average- $C/T$  method for  $T_c$  in both pressure directions. The transition temperatures for different pressures, obtained using this average  $C/T$  method, are shown in Figure 3. In our pressure range, the change in  $T_c$  is linear. It equals 56 mK/kbar for  $a$ -axis pressure, -66 mK/kbar for  $c$ -axis pressure. These values agree fairly well with those derived for the zero-pressure limit from thermal expansion data: 54 mK/kbar and -89 mK/kbar, respectively [19]. For the tetragonal crystal structure, we can also compare our results to the effect of hydrostatic pressure through  $\partial T_c / \partial p_V = 2\partial T_c / \partial p_a + \partial T_c / \partial p_c$ . Our data yield a value of  $\partial T_c / \partial p_V = 46$  mK/kbar, somewhat larger than the 25 mK/kbar obtained through direct hydrostatic pressure measurements [24].

We also comment on the normal state heat capacity. We find no significant change with  $a$ -axis pressure but an increase  $0.03 \pm 0.01$  J/kbar mol K<sup>2</sup> with  $c$ -axis pressure. Previous hydrostatic pressure measurements [24] found a decrease of -0.02 J/kbar mol K<sup>2</sup>. Although the sign differences here and in  $\delta T_c / \delta P$  may be related, the magnitudes of the heat capacity changes do not correspond to the overall Ce-In hybridization for the different types of pressure. Instead they suggest that hybridization in certain directions affects the heat capacity more than in others. The superconducting jump averages  $\Delta C / \gamma T_c = 0.73 \pm 0.05$  J/mol K<sup>2</sup> and has no apparent trend with pressure, consistent with previous experiments [5, 20, 24].

We take as the width of the transition the span between

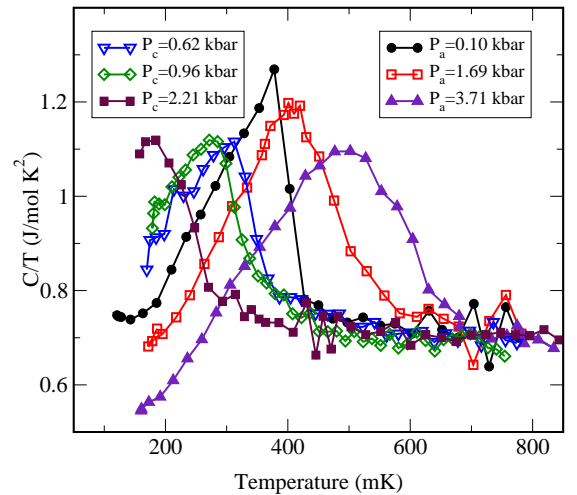


FIG. 2: Heat capacity at various pressures. The data have been scaled by the normal phase heat capacity; see text.

the temperatures corresponding to 20% and 80% of the  $C/T$  range during the transition, as shown in the inset of Figure 3. Under pressure the transition becomes broader and more rounded. Although an inhomogenous pressure distribution across the face of the sample or a variation in the cross-sectional area over the height of the sample would broaden the transition, reasonable values for these effects would produce less than 15% of the observed broadening.

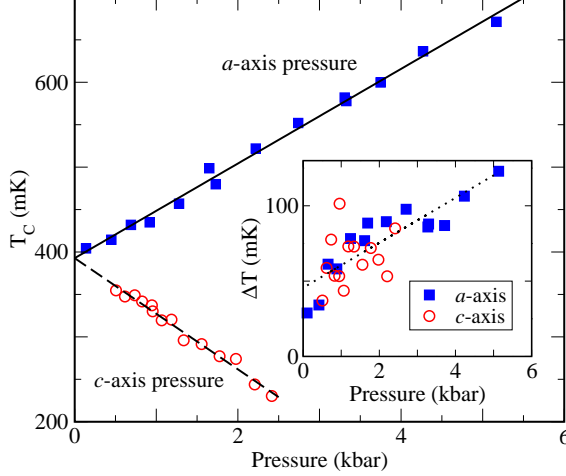


FIG. 3: Transition temperature as a function of uniaxial pressure.  $T_c$  was determined using the temperature corresponding to the average of the  $C/T$  values at the peak and onset of the transition. Lines are linear fits, with slope 56 mK/kbar and -66 mK/kbar. Inset: 20%-80% transition width vs. pressure. The dotted line is a guide to the eye.

We apply a small pressure during cooldown to keep the sample in place. This initial pressure differs for each cooldown but is typically on the order of 0.3 kbar. To determine the initial pressure, we extrapolate  $T_c$  versus  $p$  to where the transition temperature reaches that measured on samples outside of the pressure cell. The pressure labels for the curves in Figure 2 are adjusted for the initial pressure, so that the values listed are the correct pressures.

Figure 4 redisplayes the data of Figure 3 to  $T_c$  as a function of  $c/a$ . We use room temperature lattice constants [20] and the elastic constants of CeRhIn<sub>5</sub> [18] to compute  $c/a$  at different pressures [20]. We assume that the CeIrIn<sub>5</sub> elastic constants are similar [19]. Also, at our low to moderate pressures, the stress/strain relations are still linear.

Just as for measurements across the CeMIn<sub>5</sub> family, we find that  $T_c$  increases with increasing  $c/a$ . The dotted line of Figure 4 is at the ambient-pressure value for  $c/a$ . Applying  $c$ -axis pressure decreases  $c$  while increasing  $a$  through the Poisson ratio, so the data from  $c$ -axis pressures appear to the left of the dotted line. On the other hand,  $a$ -axis pressure decreases the  $a$  lattice constant along the pressure direction and increases  $c$ , but also increases the  $a$ -axis lattice constant perpendicular to the pressure direction. Since the largest change in lattice constant is along the pressure direction, the overall effect is an increase in  $c/a$ , and the  $a$ -axis data appear to the right of the dotted line. However, for  $a$ -axis pressure the ratio  $c/a$  is not a single well-defined quantity. The value of  $a$  is minimum

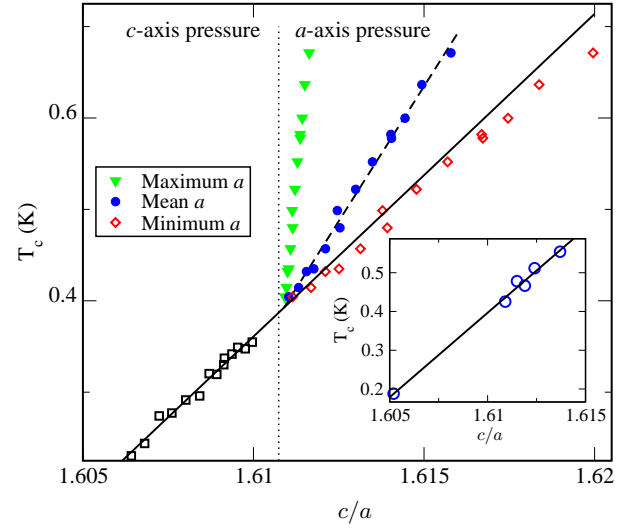


FIG. 4:  $T_c$  vs.  $c/a$ . For  $a$ -axis pressure, we calculate  $c/a$  using the minimum, geometric mean, and maximum  $a$  values; see text for details. The solid line is a least-squares fit to the  $c$ -axis data, with slope 35 K. The dashed line is a least-squares fit to the  $a$ -axis data using the mean values for  $a$ . Inset: putative  $T_c$  vs.  $c/a$  from substitution and pressure data, with hybridization potential  $V_{pf}$  equal for all points.

in the direction parallel to the applied pressure and maximum perpendicular to the pressure. Both of these extreme values are used to calculate  $c/a$ , with the results plotted in Figure 4. The remaining data set in Figure 4 uses the geometric mean of the maximum and minimum  $a$  values, which is the natural way to compare in-plane areas with the perpendicular lattice constant. As seen in Figure 4, using the mean value gives a kink in  $dT_c/d(c/a)$  at the crossover from  $c$ -axis to  $a$ -axis pressure. The larger slope of the  $a$ -axis data may indicate the influence of hybridization. Since the hybridization depends on the spacing between atoms, it increases for both directions of uniaxial pressure. However, the increase is small because the decrease in atomic spacing along the direction of the applied pressure is partly compensated by increased spacings in the perpendicular directions.

By considering several different methods of shifting  $T_c$ , we can separate the true influence of geometry on the transition temperature from the effect of hybridization. To do this, in Table I we consider the effects of hydrostatic pressure, chemical substitution, and our  $a$ -axis and  $c$ -axis pressure measurements. We use Harrison's calculation [25] of  $V_{pf}$  in a tight binding approximation,  $\eta_{pf} \frac{\hbar^2}{m} \frac{\sqrt{r_p r_f^5}}{d^5}$ . Here  $\eta_{pf} = 10\sqrt{21}/\pi$  is a dimensionless constant,  $m$  is the electron mass, and  $r_f = 0.445\text{\AA}$  and  $r_p = 19.1\text{\AA}$  are wave function radii [13, 26]. For  $r_p$  we extrapolate values from [26], p. 644. For hydrostatic pressure data, we take the percentage changes in  $V_{pf}$  and  $c/a$  from Table II and Figure 2 of [13].

Each experiment in Table I—substitution, applied pressure, or a combination of the two—increases  $V_{pf}$  over its value in CeIrIn<sub>5</sub> at ambient pressure, with a percent change given by  $\Delta V_{pf}$ . For each type of measurement we simultaneously scale

TABLE I: Values used in adjusting for effect of hybridization changes on  $T_c$ . Zero-pressure structural parameters from [20, 27]; hydrostatic pressure data from [13]. See discussion in text.

	$T_c$ (K)	$c/a$	$V_{pf}$	$\Delta V_{pf}(\%)$	$T_c$ (K) at $\Delta V_{pf} = 0.2\%$	$c/a$ at $\Delta V_{pf} = 0.2\%$
CeIrIn <sub>5</sub> (P=0)	0.40	1.6109	2.0240	–	–	–
CeIrIn <sub>5</sub> (29 kbar)	1.05	1.6099	2.1277	5.1	0.4254	1.6109
CeCoIn <sub>5</sub> (P=0)	2.30	1.6368	2.0930	3.4	0.5116	1.6124
CeCoIn <sub>5</sub> (14 kbar)	2.60	1.6435	2.1578	6.6	0.4666	1.6119
CeRhIn <sub>5</sub> (24 kbar)	2.50	1.6270	2.1326	5.4	0.4783	1.6115
CeIrIn <sub>5</sub> (5.17 kbar, $a$ -axis)	0.67	1.6158	2.0312	0.35	0.5543	1.6137
CeIrIn <sub>5</sub> (2.42 kbar, $c$ -axis)	0.23	1.6063	2.0273	0.16	0.1880	1.6052

the changes in  $T_c$ ,  $c/a$ , and  $V_{pf}$  by a single factor to reach  $\Delta V_{pf} = 0.2\%$ . Our scaled values show how  $T_c$  would vary as a function of  $c/a$  with  $V_{pf}$  held constant. The results from the final two columns of Table I also appear in the inset of Figure 4. The solid line is a best fit to all six points, with slope 44 K. Significantly, the slope remains 44 K if the fit omits the  $c$ -axis pressure data, which has particularly small hybridization change and lies far from the other points.

The broken symmetry perpendicular to the  $c$ -axis when pressure is applied along the  $a$  axis has little effect on the superconductivity, judging from the linearity of the fit in the Figure 4 inset. We note that using the minimum value of the lattice constant  $a$  preserves the linearity of  $T_c$  vs.  $c/a$ . Although probably coincidental, this may indicate how the destruction of tetragonal symmetry affects the superconductivity.

Our measurements show that up to a few kbar  $T_c$  changes linearly with uniaxial pressure, at 56 mK/kbar for  $a$ -axis pressure and -66 mK/kbar for  $c$ -axis pressure. This confirms the important role of  $c/a$  in controlling the onset of superconductivity. Changes in  $fp$  hybridization between the Ce and neighboring In atoms also have a large effect. By comparing several techniques that alter the dimensionality and the hybridization to different degrees, we control for hybridization changes and find a pure geometric influence of  $\partial T_c / \partial (c/a) = 44$  K.

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