Research Article

Pressure Induced Suppression to the Valence Change Transition in EuPdAs

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By applying a hydrostatic pressure, we have successfully suppressed the valence change transition in EuPdAs. The studied compound EuPdAs crystallizes in a P63/mmc space group. Through resistivity and magnetic susceptibility measurements, we find that EuPdAs shows a phase transition at 180 K and another transition below 10 K at ambient pressure, as was reported before. The overall transport and magnetic behavior is to some extent similar to that of the parent phase of iron based superconductors. With application of a hydrostatic pressure, the transition at 180 K is sensitively suppressed with a pressure as low as 0.48 GPa. However, superconductivity has not been induced with pressure up to 1.90 GPa.

1. Introduction

The EuPdAs alloy adopts the hexagonal CaIn₂-type structure. The transport and magnetic measurements of this material reveal a similarity to the parent phase of iron based superconductors [1, 2]. Particularly, there is a clear transition at around 180 K, which is of first order and accompanied with a large volume collapse caused by a strong contraction of the c-axis [3–5]. The transition at 180 K was reported to be induced by the fluctuation of Eu valence from 2.15 to 2.40 between 300 K and 4 K as revealed by experiments of Moessbauer measurement and μSR method [3, 6, 7]. However, whether this transition is related to some kind of magnetic transition is still unclear. And as it is known in the iron based superconductors, there is a close relationship between the high temperature phase transition and the emergence of superconductivity [8, 9]. Inspired by this similarity, we intend to check whether the transition at 180 K can be suppressed under hydrostatic pressure, as many iron-based superconductors do, and, if the transition is really related to a certain magnetic transition, whether superconductivity can be induced with the suppression of the transition.

2. Experimental Details

The polycrystalline sample of EuPdAs was synthesized by solid state reaction method, using EuAs as the precursor made by reaction of Eu and As (purity 99.9%, Alfa Aesar). High purity EuAs and Pd powder (99.99% purity, Cuibolin) were weighed in stoichiometric ratio, mixed well, and pressed into a pellet in an argon filled glove box. The pellet was loaded into an alumina crucible, which was sealed in an evacuated quartz ampule and then kept inside a PID controlled furnace. The temperature of the furnace was raised to 1000°C with a heating rate of 1.2°C/min and maintained for 24 hours. The sintered sample was cooled down naturally to room temperature with the furnace power turned off from 1000°C.

X-ray diffraction (XRD) measurements were performed on a Bruker D8 advanced diffractometer with the Cu-Kα radiation. Dc magnetization measurements were carried out with a SQUID-VSM-7T (Quantum Design). Measurements of resistivity under pressure were performed up to 2.3 GPa on a physical property measurement system (PPMS-16T, Quantum Design) by using HPC-33 piston type pressure cell with the Quantum Design dc resistivity and ac transport options. For the resistive measurements, silver leads with a diameter of 50 μm were glued to the EuPdAs sample in a standard four-probe method by using silver epoxy, and the sample was immersed in the pressure transmitting medium (Daphne 7373) in a Teflon cap with a diameter of 4 mm. Hydrostatic pressure was generated by a BeCu/NiCrAl clamped piston-cylinder cell. The pressure upon the sample was calibrated.
with the shift in $T_c$ of a high purity Sn sample by measuring the temperature dependence of resistivity.

3. Results and Discussion

Figure 1 shows the powder XRD patterns of the EuPdAs sample at room temperature, along with the result of the Rietveld structural refinement using the TOPAS program. On the whole, it is clear that the main diffraction peaks can be indexed well by the $P6_3/mmc$ space group (hexagonal structure) with $a = 4.262 \, \text{Å}$ and $c = 8.535 \, \text{Å}$ with Eu$_2$As$_2$ and Pd$_2$As as the impurity phases. The ratio between EuPdAs and the impurities is found to be 90:10.

![Figure 1: Powder X-ray diffraction patterns and the Rietveld refinement profile of the EuPdAs sample at room temperature.](image)

The resistivity of the system is generally about 100–300 $\mu\Omega\text{cm}$, which manifests that the sample can be categorized as a bad metal. Since Eu has usually anionic state of Eu$^{2+}$ and/or Eu$^{3+}$, while the As can stay at several anionic states such as As$^{3-}$ and As$^{5-}$, in this case the ionic state of Pd can be

which can be easily observed in the inset of Figure 2(b). And above 180 K, the magnetic susceptibility of EuPdAs follows Curie-Weiss law behavior very well.

The temperature dependence of electrical resistivity for EuPdAs with temperature ranging from 2.3 K to 300 K is shown in Figure 3. The 8 K and 180 K kinks also clearly appear in these resistivity curves, which is consistent with the magnetic measurements (while the 5.4 K anomaly is absent in the resistive curve). The overall transport behavior, especially the 180 K anomaly, reminds us of some parent phases of iron-based materials, like BaFe$_2$As$_2$, which exhibits antiferromagnetic order originated from Fe ions around 140 K. When we apply pressure or do some chemical substitutions on BaFe$_2$As$_2$, the 140 K transition will be gradually suppressed to lower temperature, eventually, the antiferromagnetic transition will totally disappear and superconducting state will be induced [2, 10].

Inspired by the knowledge in the parent phase of iron-based superconductors, we suspect that the 180 K anomaly for EuPdAs may be related to some kind of magnetic order of the Pd ions. If we can find some way to suppress this transition, we may expect some interesting properties, like superconductivity. To realize this purpose, we try to apply a hydrostatic pressure to EuPdAs.

The temperature dependence of resistivity for EuPdAs at various pressures with temperature ranging from 2.3 K to 250 K is illustrated in Figure 4. We find that the transition at 180 K is totally suppressed with a pressure of 0.48 GPa, and the resistivity value is depressed monotonously upon increasing pressure. However, no superconducting transition has been observed with pressure up to 1.90 GPa, while the 8 K anomaly is always preserved. Since the 180 K transition is so sensitive to pressure, we deem that it is merely a valence change transition and may not relate to the formation of any long range magnetic order. In addition, the 180 K anomaly can completely recover when pressure is removed (see Figure 3, the red curve); this reminds us of some pressure-induced iron-based superconductors, like CaFe$_2$As$_2$, in which the crystal structure will completely recover with pressure removed [11].

It is reported that the transition at 180 K is accompanied with a giant volume collapse of 1.9% caused by a strong contraction of the c-axis (2.3%) between 180 K and 150 K [3–5]; on the other hand, the lattice parameters will experience a strong but continuous decrease with increasing pressure [12]. Under the above considerations, it is reasonable to assume that the pressure we applied can result in the volume collapse at room temperature, which may induce the valence shift to 2.4 to occur at much higher temperature. Thus when applying pressure, the behaviour of the resistivity curve is well preserved below 180 K, while the 180 K transition disappears and the curve above 180 K extends smoothly from that at lower temperature.

The resistivity of the system is generally about 100–300 $\mu\Omega\text{cm}$, which manifests that the sample can be categorized as a bad metal. Since Eu has usually anionic state of Eu$^{2+}$ and/or Eu$^{3+}$, while the As can stay at several anionic states such as As$^{3-}$ and As$^{5-}$, in this case the ionic state of Pd can be
Figure 2: (a) Temperature dependence of dc magnetic susceptibility for EuPdAs as measured at an applied magnetic field of 20 Oe in the temperature range of 4.5 K to 50 K. Both magnetic susceptibilities measured in zero-field-cooled (ZFC) and field-cooled (FC) modes are shown. (b) Temperature dependence of dc magnetic susceptibility for EuPdAs under 1T in the temperature range of 4.5 K to 300 K. The inset shows 1/\chi versus T to clarify the 180 K anomaly.

Figure 3: Temperature dependence of electrical resistivity for EuPdAs in the temperature range 2.3 K to 300 K (the black curve). The red curve is the result measured after pressure is removed (sample still immersed in the pressure transmitting medium).

Figure 4: Temperature dependence of electrical resistivity for EuPdAs at various pressures in the temperature range 2.3 K to 250 K.

widely tuned. This provides a good way to tune the electronic states of the materials. This may provide an explanation to the transition at around 180 K, which is most probably induced by a valence change effect. From our recent study, one can see that the magnetic moment detected in the material is quite weak, either from the Pd or Eu ions. Nor has the long range magnetic order formation been found on the Pd ions. It would be interesting to use chemical doping to tune the electronic state 4d orbital in order to achieve a good balance between the itinerancy and local magnetism.

4. Conclusion

In summary, we have synthesized polycrystalline samples of EuPdAs by solid state reaction method. Through resistivity and magnetic susceptibility measurements, we find that EuPdAs shows a phase transition at 180 K, as well as a second phase transition below 10 K at ambient pressure. With application of hydrostatic pressure, the transition at 180 K is totally suppressed under a pressure of 0.48 GPa, and the resistivity curve is depressed monotonously upon increasing pressure. However, superconductivity has not been induced at low temperature with pressure up to 1.90 GPa. In order to achieve
some novel electronic properties, such as superconductivity, we suggest that tuning the occupation of the 4d orbitals of Pd with electrons would be essential.

**Conflict of Interests**

The authors declare that there is no conflict of interests regarding the publication of this paper.

**References**


