UC Irvine

UC Irvine Previously Published Works

Title

Discovery of novel magnetic features in the heavy-electron compound U₂Zn₁₇

Permalink

<https://escholarship.org/uc/item/9277c8f6>

Journal

Journal of Magnetism and Magnetic Materials, 108(1-3)

ISSN

0304-8853

Authors

Schenck, A
Amato, A
Birrner, P
[et al.](#)

Publication Date

1992-02-03

DOI

10.1016/0304-8853(92)91365-Z

Copyright Information

This work is made available under the terms of a Creative Commons Attribution License, available at <https://creativecommons.org/licenses/by/4.0/>

Peer reviewed

Discovery of novel magnetic features in the heavy-electron compound U_2Zn_{17}

A. Schenck ^a, A. Amato ^a, P. Birrer ^a, F.N. Gygax ^a, B. Hitti ^a, E. Lippelt ^a, S. Barth ^b, H.R. Ott ^b and Z. Fisk ^c

^a Institut für Mittelenergiephysik der ETH Zürich, CH-5232 Villigen PSI, Switzerland

^b Laboratorium für Festkörperphysik der ETH-Zürich, CH-8093 Zürich, Switzerland

^c Los Alamos National Laboratory, Los Alamos, NM 87545, USA

Transverse field μ^+ SR-measurements on a single crystal sample of U_2Zn_{17} reveal a strikingly non uniform magnetic behavior. While part ($\sim 30\%$) of the U-5f moments remain paramagnetic below $T_N = 9.7$ K (at least for external fields ≥ 0.2 T) another part ($\sim 20\%$) enters into a complex order below 9.7 K which is associated with huge internal field spreads at the μ^+ sites. Above T_N this part causes an anisotropic μ^+ Knight shift involving direction cosines up to 8th order. Only the remaining fraction is compatible with the simple AF order suggested by neutron diffraction studies.

Measurements of the magnetic susceptibility, specific heat and resistivity have revealed that U_2Zn_{17} undergoes a magnetic phase transition at 9.7 K [1], which was subsequently shown by neutron diffraction studies to lead to a simple Antiferromagnetic (AF) structure [2].

However, μ SR measurements on both polycrystalline [3] and monocrystalline samples [4] have indicated that the magnetic properties are not as simple as implied by the neutron diffraction results. We have refined and extended the analysis of the single-crystal data and have found additional striking features which we wish to report in this contribution.

For details on the sample and the measurements see ref. [4]. The measurements to be discussed here were performed in external fields of $H_{ext} = 0.4$ and 0.03 T in the temperature range 4–295 K.

The observed transverse field μ SR signal consists of four components which are distinguished by their frequency or Knight-shifts which are different both respect to their temperature and orientational dependences. In the following we list some but not all of the striking features of the four components.

Component no. 1: Accounts for $\sim 67\%$ of the total signal (this number and the corresponding ones below seem to be temperature independent). It includes some background signal from μ^+ stopping outside the single crystal specimen. Its frequency ν_1 is, within the present accuracy, solely determined by H_{ext} (no detectable Knight-shift). It does not reflect the phase transition at 9.7 K.

Component no. 2: Accounts for $\sim 20\%$ of the total signal. It shows a pronounced angular dependence of ν_2 , which is shown in fig. 1 for temperatures $\geq T_N$ and $H_{ext} = 0.4$ T. The angular dependence is well described by $\nu_2 = \nu_2^{(0)} + \nu_2^{(2)}P_2(\cos \theta) + \nu_2^{(4)}P_4(\cos \theta)$, where θ is the angle between H_{ext} and the hexagonal c-axis. This angular dependence is also observed below T_N for

$H_{ext} \geq 0.2$ T. However, for $H_{ext} = 0.03$ T $\nu_2(\theta)$ assumes a different form as the temperature is lowered from 9.5 to 4 K (see fig. 2).

Fig. 1. Orientational dependence of ν_2 at 0.4 T for various temperatures. θ_L is the angle between the c-axis and H_{ext} , shifted by an offset angle of 22° ($\theta_L = \theta + \theta_{off}$). The solid line represents a fit (see text).

Fig. 2. Orientational dependence of ν_2 at 0.03 T. Note the change in the angular dependence at the lowest temperatures as compared to fig. 1.

Component no. 3 and 4: Both appear with the same amplitude and account together for the remaining 13% of the total signal. Again a very strong angular dependence is observed above T_N which is very unusual in that 6th order and 8th order Legendre polynomials have to be included in order to describe $\nu_3(\theta)$ and $\nu_4(\theta)$ (the $T = 9.8$ K data are shown in fig. 3). In addition a strong dependence on temperature is manifest. Below T_N these two components are not detectable leading to a missing μ^+ polarization. This loss in polarization has been observed before also in a polycrystalline U_2Zn_{17} sample [3] suggesting strongly that it reflects some intrinsic property of U_2Zn_{17} .

In the following we discuss some of the implications arising from the features of the four components. Component no. 2: Above T_N the general features of this component can be understood consistently in terms of a particular μ^+ -site, e.g. $(\frac{1}{3}, \frac{1}{3}, \frac{1}{6})$, and dipole fields at this site originating from the neighboring effective U-5f-moments induced by H_{ext} in accordance with the magnetic susceptibility [5]. The fact that the form of the angular dependence persists unchanged to $T < T_N$, at least for $H_{ext} > 0.2$ T, implies that the neighboring U-moments are still able to align freely along H_{ext} (i.e. remain paramagnetic) and hence cannot take part in the AF order below 9.7 K. However, as fig. 2 reveals, the apparent paramagnetism seems to become suppressed for smaller H_{ext} and temperatures well below T_N signaling the presence of some residual interactions among the U-ions which is in competition with the Zeeman interaction.

Components no. 3 and 4: The disappearance of these components below T_N has been shown to be the result of a huge static field spread which appears in part of the sample volume below T_N [3]. The present analysis shows that this fraction of the volume is also associated with distinctly different properties above T_N as manifested in the unusual angular dependence of $\nu_3(\theta)$ and $\nu_4(\theta)$. This angular dependence cannot origi-

nate from the angular dependence of dipole fields even when taking into account the tensorial character of the magnetic susceptibility, but must involve a complex dependence of moments at the U-sites on the direction and probably strength of H_{ext} . Such a complex behavior may again be a result of competing interactions, perhaps involving quadrupole-quadrupole interactions as seen in e.g. CeB_6 [6]. The presence of competing interactions may also lead to a complex AF order in this fraction of the sample volume producing the observed huge field spread below T_N .

Fig. 3. Orientational dependence of ν_3 and ν_4 at 0.4 T and 9.8 K. The solid lines represent fits (see text). The errors, except where explicitly shown, are smaller than the dot size.

Component no. 1: The independence of ν_1 on angular orientation (and also its temperature independence) can be explained if some fraction of the μ^+ are located at another interstitial site (e.g. (000)), where the net dipole field from the induced U-moments are reduced by a factor of 30 in comparison to the site ($\frac{1}{3}, \frac{1}{3}, \frac{1}{6}$). Below T_N the dipole fields from the ordered moments will cancel to zero in case of the simple AF structure suggested in [2]. This would then leave component no. 1 unaffected.

U_2Zn_{17} shows a very peculiar magnetic behavior which suggests that several magnetically inequivalent regions or domains may coexist in a single crystal

specimen. Moreover, the observations imply strongly that at least in part of the sample volume the behavior of the U-5f moments is governed by competing interactions. It is suggested that quadrupole-quadrupole interactions may be involved.

References

- [1] H.R. Ott et al., Phys. Rev. Lett. 52 (1985) 1551.
- [2] D.E. Cox et al., Phys. Rev. B 33 (1986) 3614.
- [3] S. Barth et al., Hyperfine Interactions 31 (1986) 397.
- [4] S. Barth et al., Hyperfine Interactions 50 (1989) 711.
- [5] See e.g. J.M. Effantin et al., J. Magn. Magn. Mater. 47 & 48 (1985) 145.