

# Transport and Magnetic Properties of Dy<sub>2</sub>CoGa<sub>8</sub>

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The magnetic susceptibility, the specific heat and the electrical resistivity of the single crystal Dy<sub>2</sub>CoGa<sub>8</sub> have been measured. That compound crystallizes in a tetragonal structure with space group P4/mmm. It is found at ambient pressure that Dy<sub>2</sub>CoGa<sub>8</sub> presents an antiferromagnetic order below the Néel temperature,  $T_N = 16$  K. The electronic specific heat coefficient is about 180 mJ/K<sup>2</sup>. A strong uniaxial anisotropy is observed of the magnetic susceptibility,  $\chi$ . At higher temperatures above 150 K both  $\chi_a$  and  $\chi_c$  follows the Curie-law with an effective magnetic moment of 10.94  $\mu_B$ /Dy that is close to the expected value of for a Dy<sup>3+</sup> free ion. Crystalline electric field levels were derived from the magnetization and specific heat data.

*Keywords:* Dy<sub>2</sub>CoGa<sub>8</sub>; intermetallic compounds; antiferromagnetism.

La susceptibilidad magnética, el calor específico y la resistencia eléctrica de un mono cristal de Dy<sub>2</sub>CoGa<sub>8</sub> fue medida. El compuesto cristaliza en una estructura tetragonal con grupo espacial P4/mmm. Se encuentra a la presión ambiente que Dy<sub>2</sub>CoGa<sub>8</sub> presenta un orden antiferromagnético por debajo de la temperatura de Néel,  $T_N = 16$  K. El coeficiente electrónico del calor específico es de unos 180 mJ/K<sup>2</sup>. Un fuerte anisotropía uniaxial se observa de la susceptibilidad magnética,  $\chi$ . A temperaturas altas por encima de 150 K tanto  $\chi_a$  y  $\chi_c$  siguen la ley de Curie con un momento magnético efectivo de 10.94  $\mu_B$ /Dy que está cerca del valor esperado del ion libre Dy<sup>3+</sup>. Los niveles del campo eléctrico cristalino se derivaron de los datos de la magnetización y calor específico.

*Descriptores:* Dy<sub>2</sub>CoGa<sub>8</sub>; compuestos intermetálicos; antiferromagnetismo.

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## 1. Introduction

In the past few years, considerable interest has developed in the heavy fermions family systems CeTIn<sub>5</sub> and Ce<sub>2</sub>TIn<sub>8</sub>. [1,2] These compounds belong to the more general family R<sub>n</sub>TGa<sub>3n+2</sub> compounds with the Ho<sub>2</sub>CoGa<sub>8</sub> type structure are abundant due to their robust structure and the interesting physical properties that result from the ability of this structure to adopt different elements with various atomic sizes. The physical properties of this family of materials are also quite interesting and often display strongly correlated electron behavior especially for Ce. In particular, Ce<sub>2</sub>TIn<sub>8</sub> ( $n = 2$ ) can be described as bilayer variants of the single-layer CeTIn<sub>5</sub> with a tetragonal structure in which 2 layers of CeIn<sub>3</sub> units are stacked sequentially along the  $c$  axis with intervening monolayer of TIn<sub>2</sub> [3]. The  $n = 2$  materials possess similar Sommerfeld coefficients to that of their  $n = 1$  counterparts. At atmospheric pressure the bilayer T = Rh material is shown to order antiferromagnetically below Néel temperature ( $T_N$ )  $\approx 2.8$  K while the T=Ir material remains a heavy-fermion paramagnet to 50 mK, with no evidence for a phase transition [4,5].

This paper, concerning the magnetic properties and transport study of Dy<sub>2</sub>CoGa<sub>8</sub> single crystal. For this compound the exchange interactions and crystalline electric field (CEF) are two factors that influence the stability of the magnetic ordering of the rare-earth magnetic moments. Competition between these two interactions leads to a large vari-

ety of magnetic structures being observed in ternary rare-earth intermetallic compounds. Because of the large interatomic distances, the exchange interactions between the localized 4f electrons are indirect and are probably mediated via the conduction electrons (Ruderman Kittel Kasuya Yosida (RKKY) model).

## 2. Experimental details

X-ray powder diffraction (XRD) patterns were obtained in a Rigaku diffractometer at room temperature. Magnetization measurements have been taken in a Quantum Design superconducting quantum interference device (SQUID) MPMS-7T dc-magnetometer. Specific heat and electrical resistivity measurements were performed in a PPMS quantum design system.

Single crystalline samples of Dy<sub>2</sub>CoGa<sub>8</sub> were grown by gallium self-flux technique. The starting materials used for the preparation of Dy<sub>2</sub>CoGa<sub>8</sub> single crystals were high purity metals of Dy (99.95%), Co (99.9%), and Ga (99.999%) The crystals grew as plates (6×5×3 mm) with the  $c$  axis perpendicular to the large face. The compound Dy<sub>2</sub>CoGa<sub>8</sub> crystallizes in a tetragonal unit cell, space group I23, P4/mmm. The unit cell parameters determined by X ray powder diffraction are  $a = 4.219(5)$  Å and  $c = 10.991(2)$  Å and are in good agreement with the results found by Joshi *et al.* [6].

### 3. Results

The temperature dependence of the magnetic susceptibility of Dy<sub>2</sub>CoGa<sub>8</sub> is displayed in Fig. 1. An antiferromagnetic transition occurs at  $T_N = 18$  K, as can be seen in Fig. 1. In the paramagnetic state, the inverse susceptibility was fitted to the Curie–Weiss law with effective moment  $\mu_{\text{eff}} = 10.4 \mu_B/\text{Dy}$  and Curie temperature  $\theta = -45$  K and  $-6$  K in plane and along the c-axis, respectively. The effective moments are close to the theoretical value  $\mu_{\text{eff}} = 10.63 \mu_B/\text{Dy}$ .

In Dy<sub>2</sub>CoGa<sub>8</sub> intermetallic, the easy axis of magnetization lies along the [001] direction. The anisotropic behavior of the magnetization in both the paramagnetic and antiferromagnetically ordered states arising due to the influence of crystal electric fields (CEF). The ratio and  $\chi^c/\chi^a$  taken at  $T_N$  is mainly determined by the tetragonal CEF.

For Dy<sup>3+</sup> in a tetragonal environment, the CEF Hamiltonian can be written as

$$H_{\text{CEF}} = B_2^0 O_2^0 + B_4^0 O_4^0 + B_4^4 O_4^4 + B_6^0 O_6^0 + B_6^4 O_6^4, \quad (1)$$

where the  $B_l^m$  are the CEF parameters and the  $O_l^m$  are the Stevens operator equivalents.

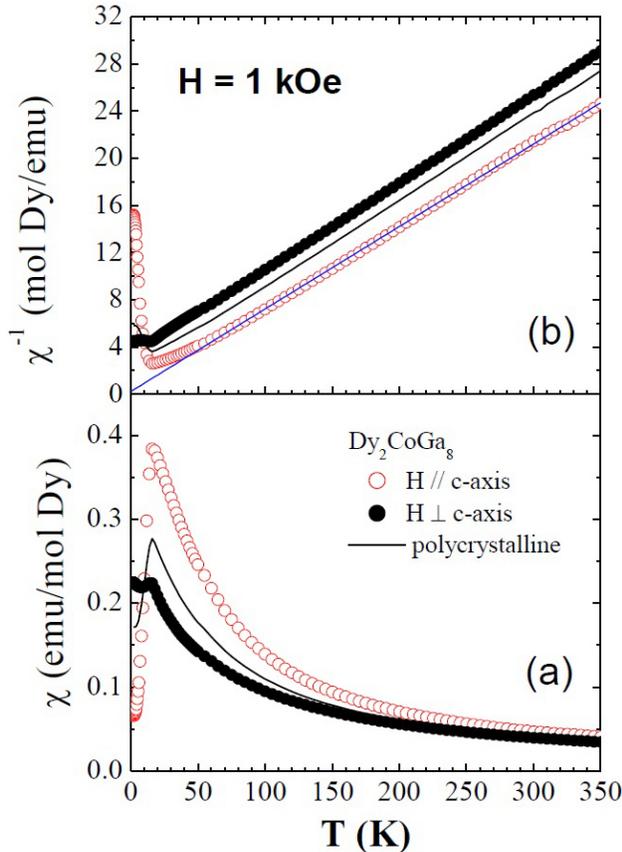


FIGURE 1. (a) The temperature dependence of magnetic susceptibility,  $\chi$  and (b)  $\chi^{-1}$  vs.  $T$  for Dy<sub>2</sub>CoGa<sub>8</sub> under a magnetic field of 1 kOe. The solid line through the data point indicates the CEF fit.

The inverse magnetic susceptibility including the molecular field contribution  $\lambda$  is given by

$$\chi^{-1} = \chi_{\text{CEF}}^{-1} - \lambda,$$

The inverse susceptibility of the Dy<sub>2</sub>CoGa<sub>8</sub> as shown in Fig. 1b, was fitted to the above discussed CEF model. The values of the CEF parameters obtained of the fit are presented in Table I

The temperature dependence of the electrical resistivity for Dy<sub>2</sub>CoGa<sub>8</sub> is plotted in Fig. 2. The inset shows the low-temperature resistivity data. The room-temperature value of the electrical resistivity is about  $50 \mu\Omega\text{cm}$  and the high temperature data show a weak metallic behavior for this compound. The resistivity for the current  $J$  along the [100] direction shows peaks at 9 and 6 K, both below  $T_N$  estimated of susceptibility peak

The temperature dependence of the electrical resistivity clearly reveals that the electrical resistivity has two peaks around 9 K and 6 K, and starts to decrease rapidly below to lower temperatures. Such a feature of the resistivity agrees with the notion of the presence of the CEF effect in this sample.

The temperature dependence of the specific-heat divided by temperature is plotted in Fig. 3. To calculate the magnetic entropy, the phonon contribution was estimated from the specific-heat data of Lu<sub>2</sub>CoGa<sub>8</sub> and subtracted from the total specific heat of the former. The magnetic contribution is plotted in Fig. 3 (square solid). The peak not sharp marks the Néel temperature in agreement with the magnetic measurements. The magnetic entropy recovered at the antiferromagnetic transition is  $8R\ln 2$ .

TABLE I. CEF parameters for Dy<sub>2</sub>CoGa<sub>8</sub> obtained from the inverse susceptibility fit.

$T_N$ (K)	$B_2^0$ (K)	$B_4^0$ (K)	$B_4^4$ (K)	$B_6^0$ (K)	$B_6^4$ (K)	$\lambda$
18	-0.6	0.004	0.001	$8.0 \times 10^{-8}$	$2.0 \times 10^{-4}$	-2,4

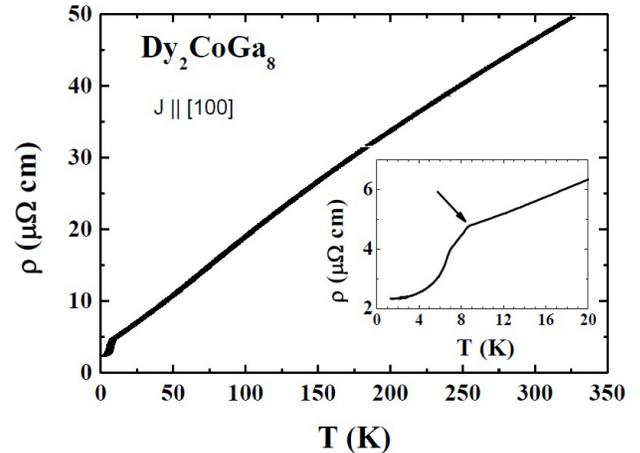


FIGURE 2. Temperature dependence of the electrical resistivity for Dy<sub>2</sub>CoGa<sub>8</sub>. The current has been applied parallel in the  $ab$  plane. The inset shows the low- $T$  range.

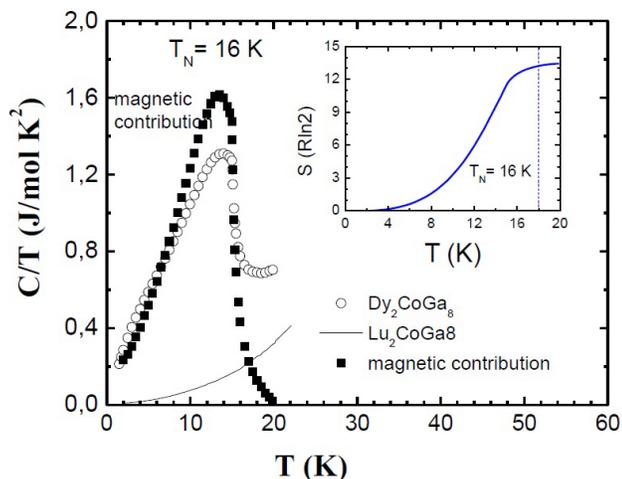


FIGURE 3. The specific heat data as a function of temperature. Inset is shown the magnetic entropy.

#### 4. Conclusions

Measurements of magnetic susceptibility, heat capacity, and electrical resistivity in single crystal  $\text{Dy}_2\text{CoGa}_8$  are re-

ported. This compound orders antiferromagnetically with a  $T_N = 16$  K, and the strong anisotropic magnetic susceptibility is higher for the field applied along the  $c$  axis. The anisotropic behavior of the susceptibility below the Néel temperature shows the  $c$ -axis as the easy axis of magnetization. Unfortunately we cannot synthesize the relative  $\text{DyCoGa}_5$  to compare its  $T_N$  with of  $\text{Dy118}$ . It is observed that density of states at the Fermi level decreases on replacing In by Ga in these  $\text{R}_2\text{CoX}_8$  ( $\text{R} = \text{rare earth}$ ;  $\text{X} = \text{In}$  and  $\text{Ga}$ ) series of compounds, it should result in a weaker conduction electron mediated RKKY magnetic interactions between the rare-earth ions in  $\text{R}_2\text{CoGa}_8$  compounds.

A detailed CEF study including fits to the existing data and additional experiments and inelastic neutron scattering experiments for a more direct determination of the CEF scheme of level for this compound is desirable.

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