

Specific-heat study on successive magnetic transitions in α -Dy2S3 single crystals under magnetic fields

メタデータ	言語: eng
	出版者: Springer
	公開日: 2020-01-30
	キーワード (Ja):
	キーワード (En): rare-earth sulfides, specific heat,
	successive magnetc transitions, magnetic entropy
	作成者: 戎, 修二, 牛木, 雄史, 高橋, 信
	メールアドレス:
	所属:
URL	http://hdl.handle.net/10258/00010128

Specific heat study on successive magnetic transitions in

α-Dy₂S₃ single crystal under magnetic fields

Shuji Ebisu, Yuji Ushiki and Shin Takahashi

Division of Applied Sciences, Muroran Institute of Technology, Japan

Specific heat measurements in magnetic fields have been performed on α-Dy₂S₃ single

crystal that shows successive magnetic transitions at $T_{\rm N_1}$ = 11.4 K and $T_{\rm N_2}$ = 6.4 K. The

specific heat in no magnetic field exhibits sharp peaks at both temperatures of T_{N_1} and T_{N_2} .

The change of magnetic entropy across each transition is estimated as Rln2/2 per mol-Dy,

which suggests magnetic moments on only one Dy site between two crystallographically

inequivalent Dy sites order at each transition temperature. When the magnetic field is applied

along the b-axis of the orthorhombic system, two peaks of the specific heat shift toward lower

temperatures. On the other hand, the magnetic field perpendicular to the b-axis shifts the

peaks toward higher temperatures. The $T_{\rm N1}$ shifts to 9.6 K (H//b) and 12.5 K ($H \perp b$) under

the magnetic field of 2 T. The peak of T_{N_2} broadens gradually with increasing magnetic field

for each direction, and the peak is consequently obscure under the field of 2 T.

PACS number: 75.40.Cx, 75.30.Gw, 75.50Ee

Keywords: rare-earth sulfides, specific heat, successive magnetic transitions, magnetic entropy

Email: ebisu@mmm.muroran-it.ac.jp

Fax: +81-143-46-5601

1

I. INTRODUCTION

The isostructural compounds of α - R_2S_3 (R = Sm, Gd, Tb and Dy) are fascinating because of their novel physical properties related to magnetic transitions [1-10]. Their crystal structure with an orthorhombic symmetry that belongs to the space group Pnma [1, 11, 12] is shown for α -Dy₂S₃ in Fig. 1. There are two crystallographically inequivalent Dy sites labeled Dy1 and Dy2 in this structure. The Dy1 atoms form planes with buckling in the ab-plane. The Dy2 atoms are connected to these planes. The decorated planes are stacked along the c-axis. The α -Dy₂S₃ shows successive magnetic transitions at $T_{\rm N_1}$ = 11.5 K and $T_{\rm N_2}$ = 6.4 K [7]. The behavior of the magnetization in the vicinities of T_{N_1} and T_{N_2} is extremely anisotropic [7] and the electrical resistivity shows anomalous enlargement between T_{N_1} and T_{N_2} [8]. It has been considered that the antiferromagnetic (AFM) ordering of the magnetic moments on Dy1 site with collinearity along the b-axis occurs at T_{N_1} . On the other hand, non-collinear AFM ordering of the Dy2 site moments takes place at $T_{\rm N_2}$. Moreover, it has been proposed that a weak ferromagnetic (WFM) ordering of the canted moments on Dy2 site occurs at T_{N_1} , because ferromagnetic-like properties are observed at the temperature range of T_{N_2} < $T < T_{\rm N_1}$ [7]. The temperature dependence of the magnetization M(T) under the magnetic field applied along the c-axis becomes steeper just below T_{N_1} , and the magnetization curve M(H) is concave down below T_{N_1} . However, the magnitude of M(H) is not so large as about 1 $\mu_B/(Dy$ atom) at the magnetic field of 0.2 T, and remanent magnetization is not observed in M(H). In this paper, we present the results of the specific-heat measurements in magnetic fields (0, 1 and 2 T), and discuss the effects of the magnetic field on the successive magnetic transitions in α -Dy₂S₃.

II. EXPERIMENTAL

The single crystals of α -Dy₂S₃ were grown by a chemical transport reaction method from the powder sample prepared by sulfurizing method with iodine as a carrier [1]. Identification of the powder sample and determination of the crystal orientation were made by X-ray (Cu K_{α}-radiation)

diffraction measurements. The specific heat was measured by a relaxation method using PPMS (Quantum Design). The measurements were made at the temperature range of 2-300 K without magnetic field and 2-20 K under magnetic fields. The magnetic fields of 1 or 2 T were applied along to the b-axis and perpendicular to the b-axis. The magnetic field of the latter case was applied in the intermediate direction between the a- and the c-axis to ensure thermal contact between the sample and platform of sample holder.

III. RESULTS AND DISCUSSION

The temperature dependence of the molar specific heat; C, and the magnetic entropy change per mol-Dy; $\Delta S_{\rm m}$, in no magnetic field for α -Dy₂S₃ is shown in Fig. 2. The molar specific heat C shows two peaks at $T_{\rm N_1} = 11.4$ K and $T_{\rm N_2} = 6.4$ K which values are consistent with the results of the magnetic experiment [7] within a difference of 0.1 K. Lattice specific heat $C_{\rm lat}$ evaluated by using the Debye model is also shown as dashed curve in Fig. 2. The Debye temperature $\Theta_{\rm D}$ was estimated as 370 K by fitting the data of higher temperatures than 150 K to the Debye's equation. This $\Theta_{\rm D}$ is somewhat higher than the value of 317 K for non-magnetic α -La₂S₃ reported by Gschneidner, Jr. et al. [13]. We assume the specific heat C consists of the lattice contribution $C_{\rm lat}$ and the magnetic one $C_{\rm m}$; $C = C_{\rm lat} + C_{\rm m}$, because the electrical properties are not metallic in lower temperature range [8]. The difference between the $C_{\rm m}$ and the experimental value of C is less than 0.9 % below $T_{\rm N_1}$. Even if we assume the value of $\Theta_{\rm D}$ as 317 K [13], it is less than 1.6 %. The $\Delta S_{\rm m}(T)$ means the magnetic entropy change from the lowest temperature of the measurement T_0 per mol-Dy, and it is estimated from the $C_{\rm m}$ using the equation

$$\Delta S_{\rm m}(T) = \int_{T}^{T} \frac{C_{\rm m}}{2T} dT \,. \tag{1}$$

Here, the $C_{\rm m}$ is divided by 2 because α -Dy₂S₃ has two Dy-atoms in a molecule. As shown in Fig. 2, both values of the magnetic entropy change across the magnetic transitions are estimated as $R\ln 2/2$,

where R is the gas constant. This fact means that the magnetic moments on only one Dy site order at each transition temperature and the ground term of Dy^{3+} on each Dy site is doublet. It looks valid that one of Kramers doublet is ground term of Dy^{3+} in the present crystal structure having low symmetry because Dy^{3+} has 9, odd number, electrons in 4f orbital. It has been found out that the model of spontaneous WFM ordering of the Dy2 moments at T_{N_1} proposed previously [7] is incorrect. Only AFM ordering of the Dy1 moments occurs at T_{N_1} , and spontaneous magnetization due to the Dy2 moments does not exist below T_{N_1} . We consider the AFM ordering of the Dy1 moments eases the Dy2 moments to align along the magnetic field, and it is a reason of the ferromagnetic-like properties observed in magnetic measurements. The C at the lowest temperature 1.9 K has a finite value of about 2 J K⁻¹ mol⁻¹. Although the origin of the remaining specific heat is unidentified at the present stage, a certain freedom might remain on Dy2 site.

Figure 3(a) shows the specific heat in the magnetic field applied along the b-axis. The magnetic field shifts both transition temperatures toward lower temperatures. The magnetic field of 1 T shifts $T_{\rm N_1}$ to 11.1 K from 11.4 K and $T_{\rm N_2}$ to 5.1 K from 6.4 K. Since the magnetic entropy changes across the transitions are almost the same between the field of 0 and 1 T as shown in Fig. 3(b), the same-type magnetic alignment is formed with decreasing temperature across the transitions in both cases. However, it seems that the transition of lower temperature occurs over a wide temperature range because of the broad peak of C(T) around $T_{\rm N_2}$ and the linear increase of $\Delta S_{\rm m}(T)$ in the field of 1 T. On the other hand, the peak of C(T) at $T_{\rm N_1}$ keeps sharpness. Furthermore, the field of 2 T shifts $T_{\rm N_1}$ to 9.6 K with keeping the sharpness of the peak in C(T). The C(T) peak of lower temperature is obscure in the field of 2 T, although an abrupt increase exists at 5.4 K. At the present stage, it is unclear that this increase is owing to a magnetic transition or an experimental error.

The specific heat in the magnetic field perpendicular to the *b*-axis is shown in Fig. 4(a). The magnetic field shifts $T_{\rm N_1}$ to 11.8 K (1 T) or 12.4 K (2 T) from 11.4 K (no field) with keeping the sharpness of the peak in C(T). Such effect, the shift of transition temperature toward higher temperatures by applying magnetic field, has not been observed in α - R_2 S₃ (R = Sm, Gd and Tb). An

AFM transition temperature of α -Gd₂S₃ shifts lower temperatures [2], and two AFM transition temperatures of α -Tb₂S₃ also shift lower temperatures by applying magnetic field regardless of its orientation [14]. In the case for α -Sm₂S₃, two WFM transition temperatures do not affected so much by magnetic field regardless of its orientation [14]. Although the peaks around T_{N_2} are vague in the field of 1 and 2 T, they seem to be in higher temperature range above 6.4 K. The magnetic entropy change $\Delta S_m(T)$ for $H \perp b$ is shown in Fig. 4(b). The $\Delta S_m(T)$ for 1 T increases monotonically with increasing temperature around $T_{N_1} = 6.4$ K for 0 T (no field); however, it rises to almost the same value as the one for 0 T at T_{N_1} . Larger value of $\Delta S_m(T)$ at $T_{N_2} = 12.4$ K for 2 T might originate from excited states of Dy³⁺ on Dy2 site.

IV. CONCLUSION

We have measured specific heat of α -Dy₂S₃ that shows successive magnetic transitions. It has been found out by evaluating the magnetic entropy change across the transitions that the WFM ordering model of the Dy2 moments at T_{N_1} is incorrect. The WFM alignment of the Dy2 moments does not occur spontaneously but was made by applied magnetic field. The magnetic entropy changes across the transitions suggest the ground states of Dy³⁺ on both Dy sites are doublet. The measurements are performed also in magnetic fields to investigated magnetic field effect on the successive magnetic transitions. The higher transition temperature T_{N_1} shifts toward lower temperatures by applying the magnetic field along the *b*-axis, while it shifts toward higher temperatures by the field perpendicular to the *b*-axis, with keeping sharpness of the peak in C(T). Therefore, the present compound has a magnetocaloric effect in only the case of the magnetic field perpendicular to the *b*-axis, although the value of the magnetic entropy change is not so large. The lower transition temperature T_{N_2} also shows the same tendency of the shift by magnetic field as T_{N_1} ; however, the shape of peak in C(T) is broaden and collapsed by the field of 1 T. Consequently, the peak is obscure in stronger field of 2 T. We conclude the ordered state of the Dy2 moments below

 $T_{\rm N_2}$ is not stable, and it is easily influenced by magnetic field. The alignment of the Dy2 moments above $T_{\rm N_2}$ is also affected by magnetic field. We consider that the alignment of the Dy2 moments by magnetic field might assist the shift of $T_{\rm N_1}$ toward higher temperatures.

ACKNOWLEDGEMENT

The present work was partially supported by Grant-in-Aid for Scientific Research (C) (KAKENHI No. 23540384) from the Japan Society for the Promotion of Science and also by a grant for the Kidorui (Rare Earth) Program through Muroran Institute of Technology.

REREFENCES

- [1] S. Ebisu, Y. Iijima, T. Iwasa and S. Nagata, J. Phys. Chem. Solids 65, 1113 (2004).
- [2] A. Kikkawa, K. Katsumata, S. Ebisu and S. Nagata, J. Phys. Soc. Jpn. **73**, 2955 (2004).
- [3] M. Matsuda, A. Kikkawa, K. Katsumata, S. Ebisu and S. Nagata, J. Phys. Soc. Jpn. 74, 1412 (2005).
- [4] K. Katsumata, A. Kikkawa, Y. Tanaka, S. Shimomura, S. Ebisu and S. Nagata, J. Phys. Soc. Jpn. 74, 1598 (2005).
- [5] S. Ebisu, M. Gorai, K. Maekawa and S. Nagata, AIP Conf. Proc. **850**, 1237 (2006).
- [6] M. Matsuda, K. Kakurai, S. Ebisu and S. Nagata, J. Phys. Soc. Jpn. 75, 074710 (2006).
- [7] S. Ebisu, M. Narumi, M. Gorai and S. Nagata, J. Magn. Magn. Mater. **310**, 1741 (2007).
- [8] S. Ebisu, M. Narumi and S. Nagata, J. Phys. Soc. Jpn. 75, 085002 (2006).
- [9] S. Ebisu, K. Koyama, H. Omote and S. Nagata, J. Phys.: Conf. Ser. 150, 042027 (2009).
- [10] S. Ebisu, H. Omote and S. Nagata, J. Phys.: Conf. Ser. 200, 092005 (2010).
- [11] C. T. Prewitt and A. W. Sleight, Inorg. Chem. 7, 1090 (1968).
- [12] A. Aruga, S. Tsujimi and I. Nakai, Anal. Sci. 12, 151 (1996).
- [13] K. A. Gschneidner, Jr., B. J. Beaudry, T. Takeshita and S. S. Eucker, Phys. Rev. B 24, 7187

(1981).

[14] Detail of the specific heat of α - R_2S_3 (R = Sm and Tb) will be given elsewhere.

Figure Captions.

Fig. 1. Crystal structure of α -Dy₂S₃, in which S atoms are omitted for clarity. The orthorhombic

unit cell is shown by a box. Two crystallographically inequivalent Dy sites are shown as Dy1 (filled

circles) and Dy2 (open circles). Thin lines are drawn between the neighbor atoms on the same ac-

plane.

Fig. 2. The temperature dependence of the molar specific heat; C, and the magnetic entropy change

per mol-Dy; $\Delta S_{\rm m}$, in no magnetic field for α -Dy₂S₃. The lattice specific heat $C_{\rm lat}$ is also shown as

dashed curve. The $\Delta S_{\rm m}(T)$ means entropy change between T and T_0 , where T_0 is the lowest

temperature of the measurement. The values of the magnetic entropy change across the transitions

are also shown.

Fig. 3. (a) The temperature dependence of the molar specific heat C in the magnetic field along the

b-axis. (b) The magnetic entropy change per mol-Dy in the magnetic field along the b-axis.

Fig. 4. (a) The temperature dependence of the molar specific heat C in the magnetic field

perpendicular to the b-axis. (b) The magnetic entropy change per mol-Dy in the magnetic field

perpendicular to the *b*-axis.

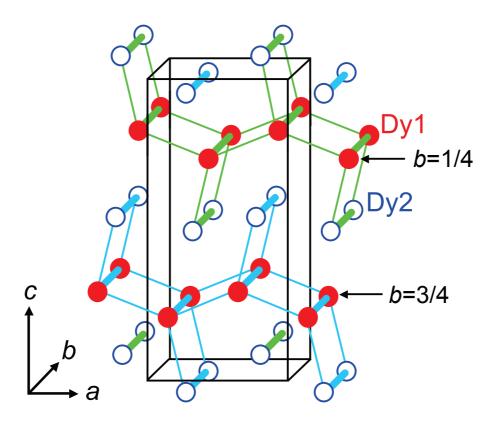


Fig. 1.

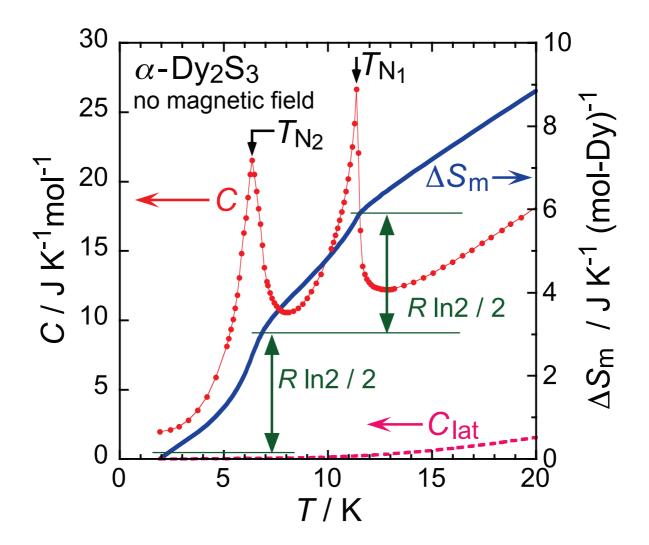


Fig. 2.

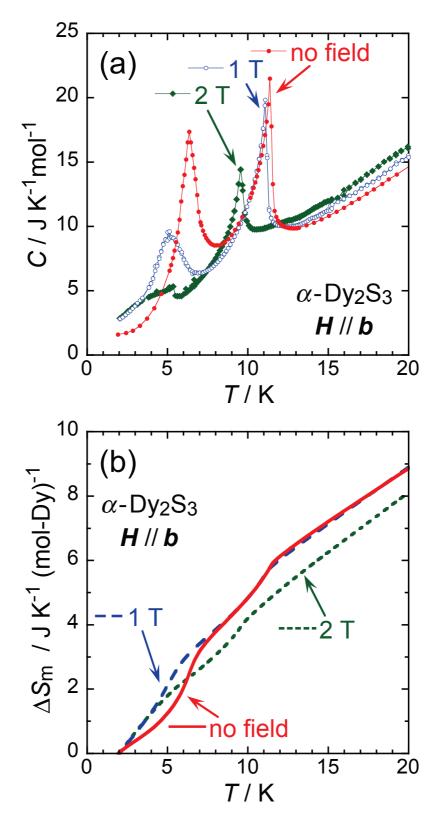


Fig. 3.

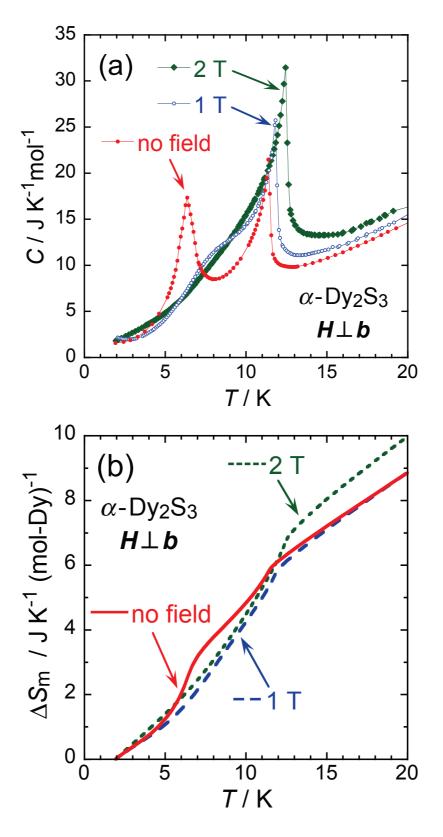


Fig. 4.