Magnetic ordering in the Tb-Co-Sn compounds

K. Iwasa A , Y. Suzuki B , H. Saito C , T. Nakajima C

 A Research and Education Center for Atomic Sciences, Ibaraki Univ., B Faculty of Science, Ibaraki Univ., C ISSP-NSL, Univ. of Tokyo

The Ln₃Co₄Sn₁₃ class of materials show various electronic states accompanied with structural transformations [1, 2]. The Ln = Cecompound shows a Weyl-Kondo semimetal behavior [3], and the Ln = La compound is a superconductor below 2.7 K [4]. The electronic properties are caused by the topological electrons produced from the asymmetric spinorbit effect under the chiral symmetry (the I2₁3 space group). Recently, a heavy-rare-earthbased compound Tb₃Co₄Sn₁₃ was reported for structural and magnetic ordering [5]. The compound shows a structural phase transition at 89.4 K and an antiferromagnetic ordering at 11.3 K. However, various composition ratios were suggested for the ternary *Ln*-Co-Sn materials. In present study, we investigated magnetic behaviors in Tb-Co-Sn, which was synthesized using a molten-Sn flux method with a starting composition Tb: Co: Sn = 1:1:20 and 30-hour heating for reaction [4].

Neutron scattering experiment for the polycrystalline sample were performed at the cold-neutron triple-axis spectrometers PONTA (5G). Measurements were conducted with final neutron energy at 14.7 meV, respectively, chosen by the focusing PG analyzer. Sample temperature was controlled using a closed-helium cryostat.

Figure 1 shows neutron diffraction data measured at chosen temperatures between 6 and 12.5 K. Below approximately 11 K, several peaks emerge, which are attributed to magnetic ordering. However, the peaks are not explained by a single-phase structure. Actually, the nuclear scattering intensity pattern (not shown here) is consistent with a mixture of $Tb_3Co_4Sn_{13}$ and $Tb_7Co_6Sn_{23}$ [7]. The latter material was newly identified to crystallize in a rhombohedral structure revealed using synchrotron x-ray diffraction analysis. Therefore, the synthesis condition used for previous $Ln_3Co_4Sn_{13}$ class of

materials are not optimized for Tb₃Co₄Sn₁₃.

The emergent magnetic peaks are indexed by (0, 0, 0) for $Tb_3Co_4Sn_{13}$ and (1/2, 0, 0) for $Tb_7Co_6Sn_{23}$. These magnetic ordering temperatures were evaluated to be 10.7 and 11.8 K, respectively.

Recently, we confirmed that the starting composition Tb: Co: Sn=1:2:50 is much better for providing a pure $Tb_3Co_4Sn_{13}$, as proposed in previous study [5]. In addition, we succeeded in synthesizing pure $Tb_7Co_6Sn_{23}$ by a longer time reaction at high temperatures. Therefore, it is a further issue to identify the magnetic structures of both compounds precisely.

The present study was performed under the approved proposal No. 24512.

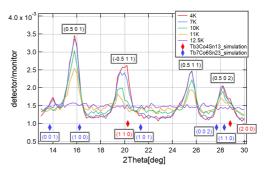


Fig. 1. Neutron diffraction data of the mixture sample of La₃Co₄Sn₁₃ and Tb₇Co₆Sn₂₃ between 6 and 12.5 K.

[1] Y. Otomo et al., Phys. Rev. B **94**, 075109 (2016). [2] J. Welsch et al., Phys. Rev. Mater. **3**, 2 (2019). [3] K. Iwasa et al., Phys. Rev. Mater. **7**, 014201 (2023). [4] E. L. Thomas et al., J. Solid State Chem. **179**, 1642 (2006). [5] C.-W. Wang et al., Chinese J. Phys. **94** (2025) 355. [7] E. K. Okudzeto et al., Physica B **403** (2008) 1628.