ESRF	Experiment title: Study of multiferroic TbMn2O5 through polarization analysis of magnetic resonant x-ray scattering	Experiment number: HE-2598
Beamline: ID20	Date of experiment:from:21th Nov. 2007to:27th Nov. 2007	Date of report : 20/10/08
Shifts: 18	Local contact(s): Dr Javier Herrero-Martin	Received at ESRF:
Names and affiliations of applicants (* indicates experimentalists): Dr. Claudio Mazzoli - ESRF Dr. Thomas Beale – Durham University, Durham, ENGLAND Prof. Peter Hatton – Durham University, Durham, ENGLAND Dr. Stuart Wilkins – Brookhaven National Laboratory, Upton, New York, USA Mr. Roger Johnson – Durham University, Durham, ENGLAND Mr. Stewart Bland – Durham University, Durham, ENGLAND		

Report:

The diverse range of physical properties observed across the multiferroic RMn_2O_3 series (R = rare-earth, Bi or Y) [1] makes clear the significance of the role of the rare-earth ion in this system, an understanding of which is of key importance. For example, $DyMn_2O_5$ exhibits one of the largest magneto-electric couplings [2] and TbMn_2O_5 undergoes a complete electric polarization reversal in applied fields of 2 T [3].

Full x-ray polarization analysis is a novel technique developed at ID20, ESRF, that measures the incident polarization dependence on the polarization of the scattered beam. It has been successful in unravelling multipole resonances in $K_2CrO_{4_1}$ [4] and in modeling competing domain contributions within NpRhGa₅ [5]. We have employed this technique for the first time to measure the spin orientation of electrons in specific bands of the terbium ion in TbMn₂O₅, selected by tuning to the resonances that originate from excitations into both the Tb 5d and 4f states has enabled us to probe the spin polarization (by nearest neighbour manganese ions and the terbium 4f electrons) of the 5d states, and the magnetic structure of the terbium sublattice existing in the partially filled 4f states. This work has culminated in the publication of a Physical Review B article (see below). Not only does this paper describe the magnetic structure of the rare-earth in TbMn₂O₅, key to furthering our understanding of this system; it more importantly has shown a new approach to determining an ion and band specific magnetic structure, excellently complementing well established neutron techniques.

This work is now being considered by ID20 as an ESRF highlight for the forthcoming annual report.

On the following page we present the abstact of the Physical Review B article. The full reference of this article is as follows:

Determination of magnetic order of the rare-earth ions in multiferroic TbMn₂O₅, R. D. Johnson, S. R. Bland, C. Mazzoli, T. A. W. Beale, C-H. Du, C.Detlefs, S. B. Wilkins and P. D. Hatton, Phys. Rev. B **78**, 104407 (2008)

PHYSICAL REVIEW B 78, 104407 (2008) Determination of magnetic order of the rare-earth ions in multiferroic $TbMn_2O_5$ R. D. Johnson,¹ S. R. Bland,¹ C. Mazzoli,² T. A. W. Beale,¹ C-H. Du,³ C. Detlefs,² S. B. Wilkins,⁴ and P. D. Hatton^{1,*} ¹Department of Physics, Durham University, Rochester Building, South Road, Durham, DH1 3LE, United Kingdom ²ESRF, 6 rue Jules Horowitz, BP220, 38043 Grenoble Cedex, France ³Department of Physics, Tamkang University, Tamsui 251, Taiwan ⁴Department of Condensed Matter Physics and Materials Science, Brookhaven National Laboratory, Upton, New York 11973-5000, USA (Received 24 May 2008; revised manuscript received 7 July 2008; published 11 September 2008) We have employed resonant x-ray magnetic scattering to specifically probe the magnetic order of the rare-earth ions in multiferroic TbMn₂O₅. Two energy resonances were observed, one originated from the E1-E1 dipolar transition and the other from the E2-E2 quadrupolar transition. These resonances directly probe the valence 5d band and the partially occupied 4f band, respectively. First, full polarization analysis, which is a measurement of the scattered polarization as a function of incident polarization, confirmed a spin polarization of the terbium valence states (probed by the E1-E1 transition) by the Mn⁴⁺ spin density in the commensurate phase. Second, full polarization analysis data were collected in the low-temperature incommensurate and commensurate phases when tuned to the E2-E2 resonance. By employing a least-squares fitting procedure, the spin orientations of the terbium ion sublattice were refined. DOI: 10.1103/PhysRevB.78.104407 PACS number(s): 75.30.Gw, 78.70.Ck, 75.50.Ee, 75.47.Lx

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- 3. N. Hur et al., Nature 429, 392 (2004)
- 4. C. Mazzoli et al., Phys. Rev. B 76, 195118 (2007)
- 5. B. Detlefs et al., Phys. Rev. B 77, 024425 (2008)