

# Investigation of Terbium with X-ray Resonance Exchange Scattering

J. Strepfner,<sup>1,2</sup> D. Hupfeld,<sup>3</sup> J. Voigt,<sup>3</sup> A. Goldman,<sup>4</sup> Th. Brückel<sup>3</sup>

<sup>1</sup> Northern Illinois University, Department of Physics, DeKalb, IL, U.S.A.

<sup>2</sup> Materials Science Division, Argonne National Laboratory, Argonne, IL, U.S.A.

<sup>3</sup> Institut für Festkörperforschung des Forschungszentrums Jülich, Jülich, Germany

<sup>4</sup> Ames Laboratory, Iowa State University, Ames, IA, U.S.A.

## Introduction

First studies on terbium with x-ray resonance exchange scattering (XRES) have been performed by Perry et al.<sup>1</sup> They observed a splitting in the energy dependence of the intensity diffracted by XRES at the  $L_{III}$ -edge but no such splitting at the  $L_{II}$ -edge. The aim of the series of experiments we conducted first at HASYLAB and then at BESSRC-CAT and MU-CAT at the Advanced Photon Source (APS) at Argonne National Laboratory was to investigate the reason for this splitting in the energy dependence. Instead of measuring the full diffracted signal, we performed a polarization analysis of the diffracted XRES signal. This allows us to distinguish between a contribution of elastic dipolar and quadrupolar transitions.<sup>2</sup>

## Methods and Materials

Terbium has a hexagonal structure and undergoes a transition from a paramagnetic phase to a basal-plane spiral antiferromagnetic phase at  $T_N \sim 230\text{K}$ . At  $T_C \sim 220\text{K}$ , a second phase transition to a ferromagnetic phase occurs. Due to the modulation wave vector  $\tau$  in the antiferromagnetic phase incommensurate satellite reflections appear at positions  $(0\ 0\ 2n \pm \tau)$ . The terbium single crystal had a mosaicity of  $0.18^\circ$  after being etched with concentrated  $\text{HNO}_3$ . For cooling, a closed-cycle He-gas cryostat was used. A resonance enhancement of the magnetic signal is observed at the  $L_{II}$ - and  $L_{III}$ -edges of terbium at  $8252\text{ eV}$  and  $7514\text{ eV}$ , respectively. The polarization analysis at the  $L_{II}$ -edge was done using a pyrolytic graphite crystal. Here the elastic scattering for the  $(0\ 0\ 6)$  reflection occurs at an angle of  $84.5^\circ$ . We performed polarization analysis in the  $\sigma$ - $\pi$ -channel to be able to make definite statements about the nature of the scattered intensity.

## Results

According to the form of the resonant cross section, the contribution of intensity scattered by quadrupolar excitation should be zero for the  $(0\ 0\ 4+\tau)$ -satellite reflection.<sup>1</sup> Therefore, it is possible to conclude from the outcome of the experiment on the nature of the scattered intensity, since the dipole scattering at  $\pm t$  is purely  $\sigma$ - $\pi$ -polarized. The energy dependencies of the magnetic satellites around the  $(0\ 0\ 2)$ ,  $(0\ 0\ 4)$  and  $(0\ 0\ 6)$  reciprocal lattice positions have been measured at the  $L_{III}$ -edge as well as at the  $L_{II}$ -edge. In Fig. 1 the energy dependence of the magnetic intensity at the  $L_{III}$ -edge is shown. The energy dependence of the magnetic intensity at the  $L_{II}$ -edge is shown in Fig. 2. Both energy dependencies are clearly different. At the  $L_{II}$ -edge, there are two peaks of a similar intensity, where one is just below the white line and one just above. At the  $L_{III}$ -edge, there is no peak above the white line. Instead, a peak is growing out with larger momentum transfer 5 eV below the main peak.

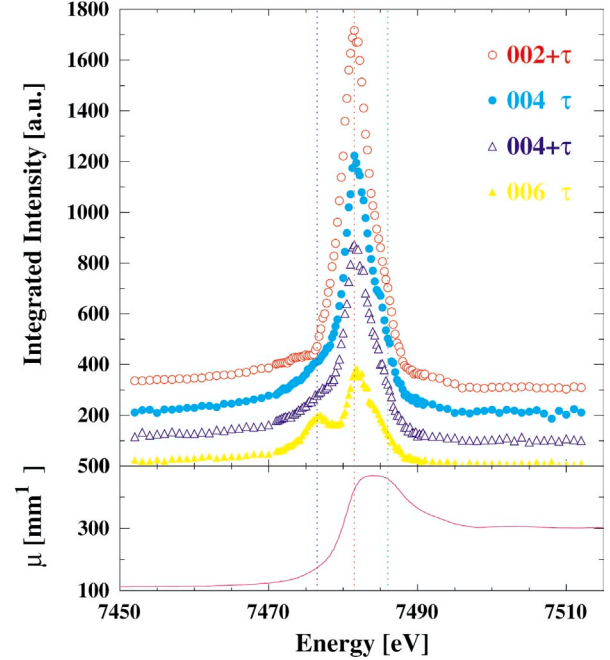


FIG. 1. Absorption-corrected energy dependence at the  $L_{III}$ -edge of terbium of the magnetic  $(0\ 0\ 2+\tau)$ ,  $(0\ 0\ 4-\tau)$ ,  $(0\ 0\ 4+\tau)$ , and  $(0\ 0\ 6-\tau)$  reflections. The graph at the bottom shows the absorption coefficient.

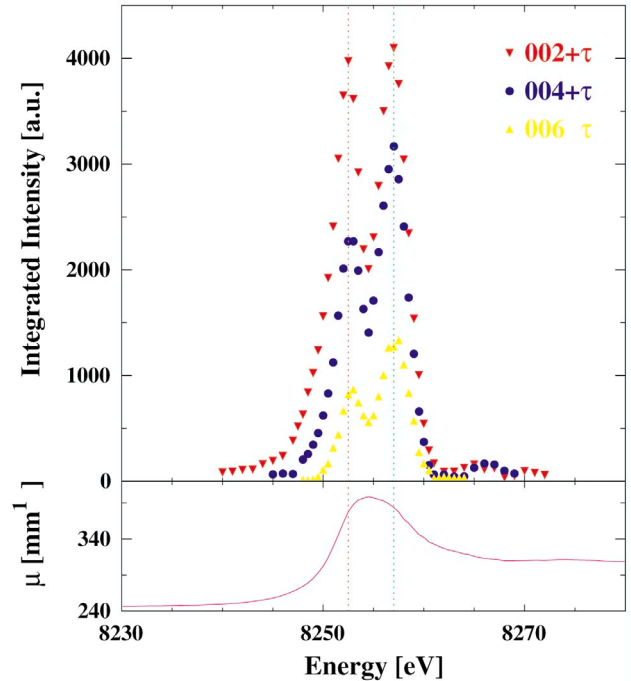


FIG. 2. Absorption corrected energy dependence at the  $L_{II}$ -edge of terbium of the magnetic  $(0\ 0\ 2+\tau)$ ,  $(0\ 0\ 4+\tau)$ , and  $(0\ 0\ 6-\tau)$  reflections.

## Discussion

Based on the calculations provided by Perry et al.,<sup>1</sup> we can conclude that there is no contribution of a quadrupolar transition to the scattering at the L<sub>II</sub>-edge, and that the structure we see in the energy dependence of the intensity may stem from the 5d-band. Although there is no clear peak above the white line at the L<sub>III</sub>-edge, there still is a pronounced slope, which suggests also a small signal here. The second peak below the white line at the L<sub>III</sub>-edge might be of quadrupolar nature.

## Acknowledgment

The Midwest Universities Collaborative Access Team (MU-CAT) sector at the APS is supported by the U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences, through the Ames Laboratory under Contract No. W-7405-ENG-82. Work at BESSRC-CAT and use of the Advanced Photon

Source was supported by the U.S. Department of Energy, Office of Science, Office of Basic Energy Sciences, Division of Materials Sciences, under Contract No. W-31-109-ENG-38 and the State of Illinois under HECA. This work has been supported by the BMBF (German Federal Ministry for Education and Research) under contract 05 NA8CJA 8.

## References:

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