

## Ru-doped $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ thin film as a coercivity tunable material studied by x-ray magnetic circular dichroism

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### Introduction

$\text{La}_{1-x}\text{Sr}_x\text{MnO}_3$  (LSMO) is a typical half-metallic compound. Although LSMO is attracting attention for possible applications to magnetic tunnel junctions, there is a serious problem that the coercivity ( $H_C$ ) is too small. Yamada *et al.*[1] reported that one can enhance the coercivity by substituting Ru for Mn, and attributed this effect to antiferromagnetic coupling between Mn and Ru and charge transfer from  $\text{Mn}^{4+}$  to  $\text{Ru}^{4+}$  ( $\text{Mn}^{4+} + \text{Ru}^{4+} \rightarrow \text{Mn}^{3+} + \text{Ru}^{5+}$ ).

In order to obtain information about the electric and magnetic states of each element in this system and to elucidate the mechanism of the coercivity enhancement, we performed x-ray magnetic circular dichroism (XMCD) measurements.

### Experimental

We fabricated the thin films by pulsed laser deposition (PLD) method. The thickness of these samples was 50nm. The substitution ratio  $y$  of Ru was 0.05. XMCD measurements were performed at BL23SU of SPring-8. The measurement was performed in the total electron yield (TEY) mode with  $T=30\text{K}$  and  $H=1\text{T}$  (out of plane). We used 1Hz helicity-switching mode. [2]

### Results and Discussion

The XMCD spectra of Mn L-edge and Ru M-edge are shown in Fig. 1 and Fig. 2, respectively. One can conclude that the spin direction of Mn is opposite to that of Ru consistent with Yamada *et al.* [1]. By applying the

XMCD sum rules, we found that the orbital moment of Ru is finite and is parallel to the spin moment. This suggests that Ru 4d orbitals are occupied by more than 5 electrons and that the valence of Ru is 3+.

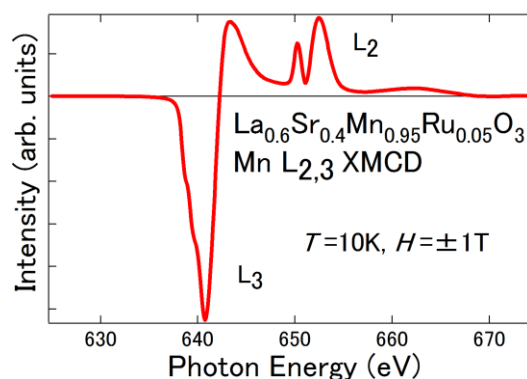


Fig. 1 XMCD spectrum at the Mn L-edge

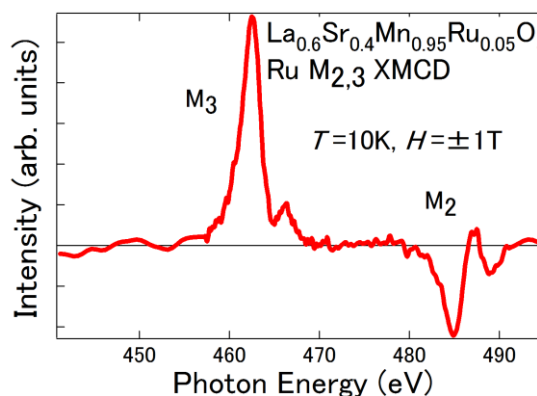


Fig. 2 XMCD spectrum at the Ru M-edge

Reference

- [1] H. Yamada *et al.*, Appl. Phys. Lett. **86**, 192505 (2005).
- [2] Y. Saitoh *et al.*, J. Synchrotron Radiation. **19**, 3, 388 (2012).

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