

広域 X 線吸収微細構造による コランダム型酸化物薄膜の Ir 置換サイトの解析

Ir substitution site analysis of Corundum-type oxide films

by extended X-ray absorption fine structure

東北大¹, JASRI², ImPACT³, 野崎友大¹, S. P. Pati¹, 塩川陽平¹,

鈴木基寛², 伊奈稔哲², M. Al-Mahdawi¹, 葉術軍¹, 佐橋政司^{1,3}

Tohoku Univ.¹, JASRI², ImPACT³, 野崎友大¹, S. P. Pati¹, Y. Shiokawa¹,

M. Suzuki², T. Ina², M. Al-Mahdawi¹, S. Ye¹, and M. Sahashi^{1,3}

E-mail: nozaki@ecei.tohoku.ac.jp

Cr_2O_3 and $\alpha\text{-Fe}_2\text{O}_3$ are Corundum-type oxide antiferromagnets. Cr_2O_3 is a well-known magnetoelectric material and attract much attentions as an electrically controllable antiferromagnet [1]. $\alpha\text{-Fe}_2\text{O}_3$ is a high Néel temperature antiferromagnet ($T_N \sim 950$ K), which has a spin reorientation transition called Morin transition. We've been reported the enhancement of Morin transition temperature (thus enhancement of perpendicular magnetic anisotropy (PMA)) for $\alpha\text{-Fe}_2\text{O}_3$ by tiny amount of Ir-doping [2,3]. In contrast, we didn't observe a distinguish change in PMA for Cr_2O_3 by Ir-doping. To investigate the difference, in this study, we tried to identify the Ir-substitution site in both Cr_2O_3 and $\alpha\text{-Fe}_2\text{O}_3$ from extended X-ray absorption fine structure (EXAFS) analysis.

Ir-doped Cr_2O_3 and $\alpha\text{-Fe}_2\text{O}_3$ films were fabricated on Al_2O_3 (0001) substrate by a reactive sputtering method. We simulated radial distribution function $|\chi(R)|$ of Ir L_3 edge with three different substitution site; 1) cation (Cr or Fe) site, 2) O-site, and 3) interstitial-site, as shown in Fig. 1. Then we fitted experimentally obtained $|\chi(R)|$ by the three simulation results. The fitting results indicate that

Ir most likely substitute cation site of both Cr_2O_3 and $\alpha\text{-Fe}_2\text{O}_3$, rather than interstitial-site and O-site.

This work was partly funded by ImPACT Program of Council for Science, Technology and Innovation (Cabinet Office, Japan Government).

[1] T. Nozaki and M. Sahashi, Jpn. J. Appl. Phys. 57 (2018) 0902A2. [2] N. Shimomura et al., J. Appl. Phys., 117 (2015) 17C736. [3] T. Mitsui et al., J. Phys. Soc. Jpn., 85 (2016) 063601.

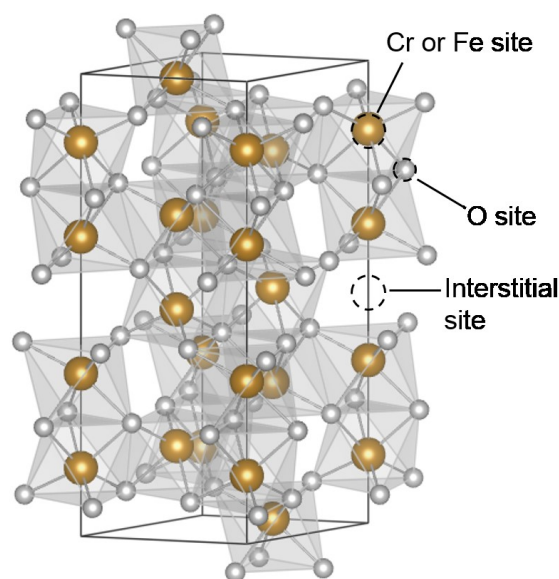


Fig. 1 Schematics of the simulated Ir-substitution site in Corundum-type oxide.