Poster Presentation

Structural and magnetic properties of spinel compound Fe_{1-x}Co_xV₂O₄

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Vanadium spinel oxides AV_2O_4 have attracted much attention for recent years because they show the peculiar physical properties which are caused by competition and cooperation of spin, orbital and lattice degrees of freedom. Among such compounds, FeV_2O_4 is a unique compound showing successive phase transitions: cubic to tetragonal (c < a) at ~140 K, from tetragonal to orthorhombic accompanied by ferrimagnetic transition at ~110 K and from orthorhombic to tetragonal (c > a) at ~70 K with decreasing temperature. It is suggested that these phase transitions originate from the orbital degrees of freedom of both Fe^{2+} ions at A-site (tetrahedral site) and V^{3+} ones at B-site (octahedral site), however, the origin remains controversial. In the present study, we investigate the substitution effect of Fe^{2+} with Co^{2+} having no orbital degrees of freedom to clarify the role of the orbital degree of Fe^{2+} at the A-site. We carried out magnetization and specific heat measurements and synchrotron powder diffraction experiments by the Debye-Scherrer camera at the beamline BL-8B at Photon Factory in KEK. For $x \le 0.1$, the successive structural transitions similar to that observed in FeV_2O_4 occur although the transition temperature of cubic-to-tetraLT transition rapidly decreases with increasing x. For $0.2\le x \le 0.6$, the only structural transition from cubic to tetragonal (c < a) was observed, however, the transition temperatures were somewhat different from the ferrimagnetic transition ones. On the other hand, for $x \ge 0.7$, the crystal structure remains cubic down to 10 K similar to that of CoV_2O_4 . These structural properties are discussed in terms of the orbital states of Fe^{2+} ions obtained by the normal mode analysis, and they are compared with the results of the specific heat and magnetization measurements.

[1] T. Katsufuji et al. J. Phys. Soc. Jpn. 77, 053708 (2008), [2] Y. Nii et al., Phys. Rev. B 86, 125142 (2012)

Keywords: spinel oxide, orbital ordering, x-ray powder diffraction