

QA01

**Mössbauer study of iron ordering in mixed valence system LuFe<sub>2</sub>O<sub>4</sub>**

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LuFe<sub>2</sub>O<sub>4</sub> has shown charge ordering on triangular plane, spontaneous polarization, and sequential phase transition scheme associated with the charge ordering in the mixed valence system[1]. N. Ikeda *et al.*[2] reported that the effect of spontaneous polarization was observed in LuFe<sub>2</sub>O<sub>4</sub> by the ordering of the Fe<sup>3+</sup> and Fe<sup>2+</sup> ions. Single crystalline LuFe<sub>2</sub>O<sub>4</sub> was grown by the floating zone method. The crystallographic and magnetic properties of the sample were measured using X-ray diffractometer (XRD), Mössbauer spectroscopy, and vibrating sample magnetometer (VSM). The crystal structure was found to be a two-dimensional layered-type rhombohedral with space group R3-mi. The magnetic Néel temperature (T<sub>N</sub>) was determined to be 250 K from the M-T curve and Mössbauer spectra. Just below T<sub>N</sub>, the magnetic moment has large value and shows a abrupt change in M-T curve. The Mössbauer spectra have been taken at various temperatures ranging from 4.2 to 360 K as shown in Fig. 1. We confirmed that the charge ordering of Fe<sup>3+</sup> and Fe<sup>2+</sup> ions was begun below 350 K, magnetic superstructure of the different ionic state was formed around 320 K, and Fe ions with different ionic state formed the superstructure around 320 K. The isomer shift value of Fe<sup>3+</sup> doublet increases with decreasing temperature from 320 to 235 K. At low temperature, Mössbauer spectra consisted of four sextets with magnetic ordering. The magnetic hyperfine fields H<sub>hf</sub> as a function of the temperature for sets of Fe<sup>3+</sup> and Fe<sup>2+</sup> on LuFe<sub>2</sub>O<sub>4</sub>, accord with the Brillouin curve B(S) for S=1/2 and S=2, respectively. We interpret that the microscopic electron structure of the Fe<sup>2+</sup> ions is in low spin state.

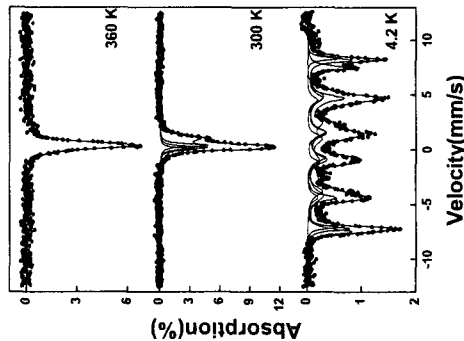


Fig. 1. Mössbauer spectra of LuFe<sub>2</sub>O<sub>4</sub> at 4.2, 300, and 360 K.

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QA02

**Correlation of Magnetic and Electric Properties by Lattice Distortion in YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub> Materials**

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Multiferroic materials have both magnetic and electric properties in single phase. Also, in multiferroic materials, the polarization is induced by the magnetic field or the magnetization is induced by the electric field. In the event, the magnetic and electric properties have interacted with together[1,2]. A single phase of the YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub> (x = 0.01, 0.02, 0.04, 0.1) was obtained by sol-gel method. The lattice constants of YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub> (x = 0.01, 0.1) were determined to be a<sub>0</sub> = 7.276 Å, b<sub>0</sub> = 8.487 Å, c<sub>0</sub> = 5.675 Å and a<sub>0</sub> = 7.282 Å, b<sub>0</sub> = 8.480 Å, c<sub>0</sub> = 5.671 Å, respectively. In other words, the lattice constant a<sub>0</sub> be increased and b<sub>0</sub>, c<sub>0</sub> decreases linearly with increasing Fe concentration. The temperature and the Fe concentration dependence of the dielectric constant (ε) show that electric Curie temperature (TCE) was decreased and the second transition anomaly was smeared from the ε(T) curves. The magnetic properties of YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub> have been measured by vibrating sample magnetometer. It shows that changes of Curie-Weiss temperatures (Θ<sub>W</sub>) and magnetic frustration factors (Γ = |Θ<sub>W</sub>|/T<sub>N</sub>) determined by Θ<sub>W</sub>. This report suggests that electric, magnetic properties and lattice displacement was related with increasing Fe concentrations of YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub>.

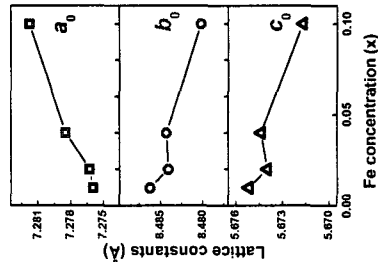


Fig. 1. Fe concentrations dependence of lattice constants for YMn<sub>2-x</sub>Fe<sub>x</sub>O<sub>5</sub> at room temperature.

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