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To cite this version:
hal-00545406

HAL Id: hal-00545406
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Submitted on 10 Dec 2010

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Pressure Induced Structural Transition in LuFe$_2$O$_4$: Towards a New Charge Ordered State

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The electronic ferroelectric lutetium ferrite (LuFe$_2$O$_4$) was studied by X-ray diffraction as a function of pressure. Pressure is shown to induce an irreversible rhombohedral to orthorhombic transition leading to a supercell determined by the combination of electron and synchrotron X-ray diffraction. This new configuration is proposed to be charge-ordered in agreement with the results of resistivity measurements.

PACS numbers: 7  

Coupled phenomena between ferroelectric and ferromagnetic properties are gaining increasing interest from both a fundamental and technological point of view [1]. Although these materials are rare, the mechanisms giving rise to multiferroicity are shown to originate from particular balance between charge-spin-orbital and lattice parameters [2]. Lutetium ferrite (LuFe$_2$O$_4$) appears as an ideal candidate to study as its multiferroic properties were shown to be directly linked to electronically-driven ferroelectricity [3]. At room temperature, the structure exhibits charge ordering [3] and is thus more complex than the usual description, i.e. the stacking along c of triangular lattices of lutetium, iron and oxygen [4] (Figure 1a). Owing to the ordered charge distribution, each iron bi-layer has a dipole moment and the system is ferroelectric below the charge-ordering temperature ($T_{CO}$ $\approx$ 330 K), which was determined based on Mössbauer spectroscopy [5], neutron [6] and electron diffraction [7]. The exact full three-dimensional charge configuration, determining the net electronic polarisation of the material, is however under debate from a theoretical [8] and experimental [6, 7] point of view. Additionally, upon cooling, ferrimagnetism is observed below 240 K giving rise to particular coupling of the sequential charge ordering and spins [9, 10]. Further interesting properties linked with spin-charge order coupling have recently been reported such as magnetoelectric response [11], giant magnetic coercivity [10] and strong insulating behavior [5] for this material. The supercell detected by neutron diffraction is reported to have a charge ordered structure in the ab plane along the [110] direction [6]. Single crystal X-ray scattering studies proposed an antiferroelectric stacking of the charge order associated with an incommensurate propagation close to ($\frac{1}{3} + \frac{2}{3}$), while diffuse scattering at 360 K with ($\frac{1}{3} + \frac{4}{3}$) propagation indicated ferroelectric short-range correlations between neighboring double layers [12].

As the electric polarization arises from the special charge order of Fe$^{2+}$ and Fe$^{3+}$, pressure can be expected to decrease the Fe-Fe distances and hence increase the electronically-driven ferroelectricity. Such a scenario has already been observed in RMn$_2$O$_5$ manganites [13] and the different high pressure effects upon RMnO$_3$ [14] show that pressure is an appropriate parameter to better understand electronically-driven ferroelectricity. A very recent study was reported on the pressure effects on LuFe$_2$O$_4$ up to 2.5 GPa by ac susceptibility and electronic transport measurements showing that pressure tends to suppress the long-range magnetic state [15]. In this Letter, we present a high-pressure structural study of LuFe$_2$O$_4$ by X-Ray diffraction and resistivity measurements. The recovered sample is further characterized by electron and synchrotron X-ray diffraction. We will show that higher pressure induces an irreversible phase transition to a high-pressure ordered structure with an orthorhombic supercell. This new phase is proposed to be due to a new charge ordered state.

LuFe$_2$O$_4$ was prepared by solid state reaction at high temperature, starting from a 10g mixture of 0.485 Lu$_2$O$_3$: 0.815 Fe$_2$O$_3$: 0.37Fe, pressed in the shape of a rod (6mm diameter and several cm length) and heated at 1180°C for 12 h in an evacuated silica ampoule. Neutron powder diffraction (NPD) data (recorded on 3T2 at the Laboratoire Léon Brillouin at room temperature) show that the sample is free of impurities and has the expected oxygen stoichiometry within the accuracy of the refinement. The refined cell parameters are $a_{R\bar{3}m} = 3.44051(2)$ Å and $c_{R\bar{3}m} = 25.2389(2)$ Å in the $R\bar{3}m$ space group (hexagonal setting). Measured Neel and charge ordering transition temperatures occur at $\approx$ 240 K [9, 10] and $\approx$ 330K [5, 7], respectively in agreement with the literature.

Angle-dispersive X-ray diffraction data at high pressure were obtained using a laboratory source and a diamond anvil cell (DAC). The LuFe$_2$O$_4$ powder was placed
in a tungsten gasket along with NaCl as pressure calibrant. A 21:4:1 (volume ratio methanol:ethanol:H₂O) mixture was introduced as a pressure-transmitting medium. X-ray diffraction patterns were obtained with Zr-filtered Mo radiation from a 800-W microfocus tube. X-ray capillary optics were used, giving a 100-µm-diameter beam. Detection was performed with an imaging plate. Exposure times were typically between 24 and 48 hours. The intensities obtained on an imaging plate were integrated as a function of 2θ using FIT2D in order to give conventional, one dimensional diffraction profiles [16]. Pressures were measured based on equation of state of NaCl [17]. Cell parameters were obtained either by Le Bail method or Rietveld refinement using the software Fullprof [18]. Pressure - volume data were fitted to the Birch-Murnaghan equation of state with B\''₀ = 4) [19]. An X-ray diffraction pattern of the recovered sample in the gasket was obtained at the European Synchrotron Radiation Facility on beamline ID27, using 0.3738 Å photons and a Bruker smart CCD 6500. This sample was then studied by electron microscopy, using a JEOL 200CX electron microscope.

Sample resistance was determined in a large volume multianvil press [20]. A pressed pellet of LuFe₂O₄ was placed in a pressure medium consisting of semi-sintered Cr-doped MgO. Resistance across the disc was monitored during compression up to 14 GPa.

In agreement with the NPD study, the X-ray diffraction pattern of LuFe₂O₄ at ambient conditions is consistent with structural data obtained in the literature, Figure 2a (bottom). The hexagonal unit cell parameters could be easily refined up to a pressure of 4.5 GPa beyond which a phase mixture is observed. Above 8 GPa, the initial low-pressure form had almost entirely disappeared. Additional compression steps were made up to 12 GPa to obtain the pure high-pressure phase. The high-pressure phase can be recovered at atmospheric pressure, Figure 2b. The diffraction pattern of the recovered, high-pressure phase of LuFe₂O₄ is more complex than that of the low-pressure starting material; however, there are evident similarities between the two patterns, Figure 2.

A series of sharp reflections are observed systematically shifted to higher 2θ with respect to the 00l (l = 3n) reflections of the low pressure form. These shifts correspond to a decrease in the c lattice parameter by 2.8%. A series of additional, broad low angle reflections are observed in the high-pressure pattern indicating an increase in the size of the unit cell, Figure 2. These new broad peaks are observed at close to twice the d-spacing of a series...
of 10l (l = 5, 8, 11) reflections of the starting material, which appear at close to same 2θ values in the recovered high-pressure form.

FIG. 3. (color online) a) [30 ¯2] electron diffraction pattern of LuFeO4. Only the reflections of the sub-cell are indexed (white numbers): all the satellites are aligned along the intermediate rows indicated by yellow arrows. b) Indexation of the satellites (red numbers in italics), using four indices hklm. c) Reconstitution of [010] pattern, obtained through the rotations and which allows to evidence the modulation vector = ra^2 + sc^2 (with r = 1 and s = 1/2) is given in Figure 3c. The blue spots represent the h0l reflections and the white ones the hIl reflections (located in the layer 1) of the sub-cell; in order to simplify the scheme, only a part of the satellites hIlm are numbered (in red), because the others overlap the reflections of the sub-cell. The new phase can thus be described by a modulated structure: a_{sub} ≈ 5.4 Å, b_{sub} ≈ 17.6 Å and c_{sub} ≈ 32 Å, with α, β, γ ≈ 90° and q = 1a^2 + 1/2c^2. Because of the commensurability of the modulation vector, the high-pressure phase can be described in a 3D lattice with a ≈ 5.4 Å, b ≈ 17.6 Å and c ≈ 64 Å, and α, β, γ ≈ 90°.

The parameters of the sub-cell are consistent with those obtained from Le Bail refinement of the synchrotron X-ray diffraction data (Figure 2b): a_{hp} = 5.400(5) Å, b_{hp} = 17.457(12) Å, c_{hp} = 32.804(25) Å. If the relationship between the a parameters (in high and ambient pressure phases) is evident [a ≈ a_{R3m}√3], the others are more complex. It could be easier by considering the following monoclinic unit cell C2/m with a_{C2/m} = 5.957 Å [ = a_{R3m}√3], b_{C2/m} = 3.436 Å [ = a_{R3m}] and c_{C2/m} = 8.641 Å [ ≈ c_{R3m}/3] with β_{C2/m} = 103.2° (Figure 1b). The cell parameters of the modulated supercell are thus: a_{super} = 3a_{unit}, b_{super} = 5 a_{unit} and c_{super} = 8 c_{unit} (Figure 1c). The (001) plane is obtained from that considered in a hexagonal setting, with a larger surface and a strong distortion 5a_{super}/b_{super} ≈ 1.55 instead of √3. The five-fold multiplicity of the b_{super} parameter is clearly observed in the ED pattern shown in Figure 3a. The value of the c parameter, 8 c_{unit}, confirms the disappearance of the rhombohedral stacking mode of the layers along this direction.

The high-pressure phase is found to be less compressible than the low-pressure phase, Figure 4a, B₀ = 188(3) GPa and B₀ = 148(1) GPa respectively. The recovered sample at atmospheric pressure is only about 0.5% denser than the starting phase, which is evidence that the LuO1.5 layers and Fe2O2.5 layers are retained. The LuO1.5 close packing is similar to that of alumina (Al2O3), which is known to be stable at high pressure [21]. Concerning the Fe2O2.5 bilayers if pressure induced a (partial) increase in the iron coordination number with distinct environments for Fe^{2+} and Fe^{3+}, this would result in a volume change of more than 5 %, which is not the case. The Fe2O2.5 bilayers in agreement with the TEM results retain a similar network of triangular bipyramids and the increase in the size of the unit cell is only due to rearrangements. Accordingly, the initial Fe^{2+}/Fe^{3+} charge order in LuFeO4 is most probably affected and a new charge ordered structure could thus be obtained in the high-pressure form.

Therefore, the observed transition as a function of pressure is due to changes in charge ordered structure of Fe^{2+}
and Fe\textsuperscript{3+}, while retaining five-fold coordination. Pressure enhanced charge ordering has already been reported in literature [22, 23]. As Fe\textsuperscript{2+} and Fe\textsuperscript{3+} in triangular bipyramid arrangements have different ionic radii, \emph{i.e.} about 0.1 Å, they exhibit different bond lengths and local compressibilities. Hence, it is possible to observe a new pressure induced charge-ordered structure (less compressible) that can be recovered at atmospheric pressure. By analogy with the work of Angst \emph{et al.} [12], the orthorhombic high-pressure phase obtained in our study could probably have an antiferroelectric stacking of the dipole moment which explains the large size of the supercell. In this electronically-driven ferroelectricity, such a charge arrangement is probably of greatest interest as the spontaneous polarization arises from the electronic dipole moment which explains the large size of the supercell.

In conclusion, our results show that pressure induces a new charge order in LuFe\textsubscript{2}O\textsubscript{4}, which is recoverable at atmospheric pressure. In accordance with the new large orthorhombic supercell and the conservation of the five fold coordination, the strong X-ray superlattice reflections arise from distinct Fe environments with characteristic Fe-O distances. This work should motivate further studies on the high-pressure form (dielectric, polarization, magnetic properties).

The authors wish to thank Gaston Garbarino from the beamline ID27 of the European Synchrotron Radiation Facilities (ESRF). Financial support for this work was partially provided by the French Agence Nationale de la Recherche, Grant N\textsuperscript{o} ANR-08-BLAN-0005-01 and N\textsuperscript{o} JC08-331297. The multi-anvil apparatus of Laboratoire Magmas et Volcans is financially supported by the Centre National de la Recherche Scientifique (Instrument National de l’INSU).

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