Magnetic and Structural Studies of the Multifunctional Material SrFe$_{0.75}$Mo$_{0.25}$O$_3$–$\delta$

M. Retuerto,† M.-R. Li,‡ Y. B. Go,† A. Ignatov,‡ M. Croft,‡ K. V. Ramanujachary,§ J. Hadermann,∥ J. P. Hodges,⊥ R. H. Herber,# I. Nowik,# and M. Greenblatt*†

†Department of Chemistry and Chemical Biology, Rutgers, The State University of New Jersey, 610 Taylor Road, Piscataway, New Jersey, 08854, United States
‡Department of Physics and Astronomy, Rutgers, The State University of New Jersey, 136 Frelinghuysen Road, Piscataway, New Jersey, 08854, United States
§Department of Chemistry and Physics, Rowan University, 210 Mullica Hill Road, Glassboro, New Jersey, 08028, United States
∥EMAT, University of Antwerp, Groenenborgerlaan 171, 2020 Antwerp, Belgium
⊥Instrument and Source Design Division, Oak Ridge National Laboratory, Oak Ridge, Tennessee 37831, United States
#Racah Institute of Physics, Hebrew University, Jerusalem, 91904 Israel

ABSTRACT: SrFe$_{0.75}$Mo$_{0.25}$O$_3$–$\delta$ has been recently discovered as an extremely efficient electrode for intermediate temperature solid oxide fuel cells (IT-SOFCs). We have performed structural and magnetic studies to fully characterize this multifunctional material. We have observed by powder neutron diffraction (PND) and transmission electron microscopy (TEM) that its crystal symmetry is better explained with a tetragonal symmetry (I$4/mcm$ space group) than with the previously reported orthorhombic symmetry (Pnma space group). The temperature dependent magnetic properties indicate an exceptionally high magnetic ordering temperature ($T_N \sim 750$ K), well above room temperature. The ordered magnetic structure at low temperature was determined by PND to be an antiferromagnetic coupling of the Fe cations. Mössbauer spectroscopy corroborated the PND results. A detailed study, with X-ray absorption spectroscopy (XAS), in agreement with the Mössbauer results, confirmed the formal oxidation states of the cations to be mixed valence Fe$^{3+}$/4+ and Mo$^{6+}$.

INTRODUCTION

The search for multifunctional materials is a promising area of research due to the interest in fields including multiferroicity, fuel cells, and spintronics. Metal oxides with an ABO$_3$ perovskite structure have traditionally attracted much attention since they present a variety of promising physical properties for potential applications, such as superconductivity, ferroelectricity, and magnetoresistance. Recently, the perovskites have been investigated also as electrode materials for the development of intermediate-temperature solid oxide fuel cells (IT-SOFCs).$^{1,2}$

Much attention has been focused on the study of perovskites with two types of cations at the B site of the structure. The interest in these so-called double perovskites of general formula A$_2$B'B'O$_6$ (A = divalent cations or rare earths; B = transition metals) has been renewed due to the room temperature (RT) colossal magnetoresistance (CMR) properties exhibited by Sr$_2$FeMoO$_6$. However, only a few metal oxides were found to show the potentially useful properties of this material, which simultaneously exhibits ferromagnetic (FM) and half-metallic behavior at RT. The modification of the compositional, structural, and magnetic properties of Sr$_2$FeMoO$_6$ has been widely studied to find candidates for technological devices.$^{4,5}$ Similar CMR properties were described for other double perovskites such as A$_2$FeMoO$_6$ (A = Ba, Ca)$^{6,7}$ and A$_2$FeReO$_6$ (A = Sr, Ba, Ca)$^{8,9}$.

In this work, we focused on a double perovskite of stoichiometry Sr$_4$Fe$_2$MoO$_{12}$ (rewritten as Sr$_2$(Fe)$_2$(Fe$_{0.5}$Mo$_{0.5}$)$_2$O$_6$) with intrinsic partial disorder between Fe and Mo over half of the perovskite B positions and full occupation of Fe over the other half of B sites. The objective was to improve the magnetic properties of Sr$_2$FeMoO$_6$ by increasing the amount of the stronger magnetic cation (Fe$^{3+}$, d$^5$) at the B site. This strategy has previously worked for the family of perovskites A$_3$Fe$_2$B'O$_9$ (A = Ca, Sr, Ba; B' = U, Te, Mo)$^{10–12}$.

Recently, the interest in Sr$_2$Fe$_2$MoO$_{12}$ has been renewed, because of its potential application as efficient electrode material for IT-SOFCs$^{13–15}$ both as a cathode and anode simultaneously, with excellent chemical compatibility with the...
electrolyte and chemical stability in both reducing and oxidizing conditions.16

Surprisingly, we found that Sr₄Fe₃MoO₁₂, this interesting material for SOFCs, is also an exciting magnetic material, and therefore in this work we carried out detailed magnetic and crystallographic studies to determine its magnetic and crystallographic structure, as well as further characterizations by X-ray absorption spectroscopy (XAS), Mössbauer spectroscopy, and electron diffraction (ED).

■ EXPERIMENTAL SECTION

Sr₄Fe₃MoO₁₂ was prepared by the sol–gel method. Stoichiometric amounts of analytical grade SrCO₃, FeC₂O₄·2H₂O, and (NH₄)₂MoO₄·4H₂O were dissolved in an aqueous solution of 10% citric acid (10 g of citric acid in 100 mL of water) and 10 mL of nitric acid. The citrate and nitrate solution was slowly evaporated, leading to an organic resin containing a homogeneous distribution of the involved cations. The resin was dried at 120 °C and then decomposed at 600 °C overnight. The organic materials and nitrates were eliminated in a subsequent treatment at 800 °C in the air. This treatment gave rise to a homogeneous precursor material. Then, the precursor was heated in air, in two different steps with intermediate grinding: first at 950 °C for 12 h and finally at 1250 °C for another 12 h. The heating rate was 2 °C/min, and the cooling was inside the furnace after it was shut off. The product was initially characterized by a laboratory powder X-ray diffractometer (PXRD; Bruker D8, Cu Kα, λ = 1.5406 Å) for phase identification and to evaluate the phase purity. For the crystal and magnetic structure refinements, we have collected powder neutron diffraction (PND) data at RT and 14 K at the POWGEN instrument17 in the Spallation Neutron Source at Oak Ridge National Laboratory, Tennessee. The diffraction patterns were analyzed by the Rietveld method, with the Fullprof refinement program.18,19 In the final run, the following parameters were refined: scale factor, background coefficients, unit-cell parameters, positional coordinates, isotropic thermal parameters, occupancies, and magnitudes of the magnetic moments. The space group determination was supported by electron diffraction with a Phillips CM20 transmission electron microscope, for which the powder was ground, dispersed in ethanol, and deposited on a holey carbon grid. The DC magnetic susceptibility was measured with a commercial SQUID magnetometer on a powder sample, in the temperature range of 1.5–800 K and fields up to 5 T. X-ray absorption near edge spectroscopy (XANES) was collected simultaneously in both the transmission and fluorescence modes on powder samples on beamline X-19A at the Brookhaven National Synchrotron Light Source. Mössbauer studies were performed with a ⁵⁷Co/Rh source (50 mCi) and a conventional constant acceleration Mössbauer drive in transmission geometry. The absorber was placed either in a cryostat (85 K) or high temperature oven, in a boron nitride holder.

■ RESULTS

Crystal Structure. We have successfully obtained Sr₄Fe₃MoO₁₂ as a black, well-crystallized polycrystalline sample. A single-phase perovskite was identified by PXD (Figure 1). No superstructure peaks corresponding to long-range Fe and Mo ordering were observed; thus the formula can be written as Sr₀.₇₅Fe₀.₇₅Mo₀.₂₅O₃. The structure of Sr₄Fe₃MoO₁₂ was first reported in a cubic space group Pm₃m,20 and later it was defined by PND in orthorhombic Pnma.15 However, in the later report, possible tetragonal space groups were rejected, because the refinement of the structure by PND did not converge. We reexamined the structure using PND and electron diffraction and found a different crystal structure, although both compounds, ours and the one previously reported, appear to have nearly identical stoichiometries.

In the high-resolution PND data, the splitting observed in some reflections observed can be explained only by the lowering of the symmetry from cubic to tetragonal (Figure 2). The crystal structure was then refined in the tetragonal I4/mcm space group (No. 140). The perovskites with this space group belong to a subgroup of the aristotype perovskite with a space group Pm₃m and may be obtained via an out-of-phase octahedral tilting of BO₆ octahedra along the [001] direction.21 Using the Glazer notation, it can be described as $d^{3}d^{3}c^{−}$.22 No other splitting of the diffraction peaks was observed to indicate additional lowering of the symmetry to an orthorhombic space group, and electron diffraction was used to confirm the tetragonal symmetry. The electron diffraction in Figure 3 shows the existence of the (110)₀, (100)₀, and (111)₀ zones (p stands for indexation in the perovskite parent cell) with and without extra reflections at 1/2(000) (where o indicates odd indices), while the (111)₀, and (100)₀ zones all looked the same and showed no extra reflections compared to the Pm₃m perovskite parent. The 1/
2(000) reflections signify the occurrence of antiphase tilts, while the absence of extra reflections in the other zones signifies the absence of in-phase tilts. Together with the occurrence of (110) patterns in our case. We conclude that we have a tetragonal

$\langle a^p b^p c^p \rangle$, which were not observed in any of the experimental

$\langle e^e o^e o^e \rangle$ due to the combined

$\langle b^a a^a c^a \rangle$ only, with $\langle b^a a^a c^a \rangle$ being the only tetragonal one

$\langle a^e b^e b^e \rangle$ only. Therefore, electron diffraction includes the tilt system proposed from PND as one of the four remaining possibilities. Note that the recently published space group $Pnma\ (a^p b^p c^p)$ for this compound would result in extra refractions (at 1/2(eoo) due to the combined presence of in-phase and antiphase tilts (e means even indices), which were not observed in any of the experimental patterns in our case. We conclude that we have a tetragonal perovskite with the $I4/mmm$ space group. There are other related perovskites with this space group, e.g., SrMn$_{1/2}$Ru$_{1/2}$O$_3$ and Sr(Fe$_{1/2}$Nb$_{1/2}$)O$_3$. This model corresponds to a simple perovskite with only one B site for Fe and Mo cations and two kinds of nonequivalent oxygen atoms (O1 and O2). The O1 and O2 oxygen atoms are located in the ab plane and along the c axis, respectively. The refinement of the possible oxygen deficiency of the sample led to a compound of formula SrFe$_{0.75}$Mo$_{0.25}$O$_{3-d/2}$ with some oxygen vacancies. The oxygen deficiency has been found in the O1 position, and O2 is stoichiometric as previously reported. The structural parameters, oxygen occupancies, main interatomic distances, and angles at RT and 14 K are listed in Tables 1 and 2. The refinements at 300 and 14 K are in good agreement with the calculated patterns (Figure 4, the second set of reflections shown at 14 K corresponds to the magnetic structure that will be explained below). The schematic structure of SrFe$_{0.75}$Mo$_{0.25}$O$_{3-d}$ is shown in the upper inset of Figure 1. As we described earlier, the tetragonal structure occurs due to a tilting of the BO$_6$ octahedra along the c direction.

Table 1. Positional, Thermal Parameters, and Agreement Factors for SrFe$_{0.75}$Mo$_{0.25}$O$_{3-d}$ after the Rietveld Refinement of NPD Data Collected at 300 and 14 K

<table>
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<tr>
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<tr>
<td>Sr</td>
<td></td>
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<tr>
<td>Sr-O1</td>
<td>2.702(2)</td>
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<td>2.877(16)</td>
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<tr>
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<td>Fe=Mo-O</td>
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</tr>
<tr>
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<td>1.9605</td>
</tr>
<tr>
<td>angles</td>
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<td>180.0</td>
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</table>

Table 2. Main Interatomic Distances (Å) and Angles (deg) for SrFe$_{0.75}$Mo$_{0.25}$O$_{3-d}$ at 300 and 14 K

<table>
<thead>
<tr>
<th></th>
<th>300 K</th>
<th>14 K</th>
</tr>
</thead>
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<tr>
<td>Sr-O</td>
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<td>0.62(3)</td>
</tr>
<tr>
<td>Fe/Mo</td>
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<td>0.52(2)</td>
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<td>mag. mom.</td>
<td>0.85(7)</td>
<td></td>
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<tr>
<td>Sr</td>
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<td>B/A</td>
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<td>Fe/Mo</td>
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<td>0.52(2)</td>
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<td>mag. mom.</td>
<td>0.85(7)</td>
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</table>
some oxygen vacancies (as the PND results suggest, i.e., the formal oxidation state of the Fe ions. decrease as the content of Mo decreases due to the increase of di vacancies in the material. The tendency of this compound to Sr$_2$FeMoO$_6$. Also it is important to note that O1 is in the been reported in this compound and in similar ones such as Sr$_2$FeMoO$_6$.15,30 Also it is important to comment the obtained oxygen usivity, has been previously reported.15 It also has been compared with those of Sr$_2$FeMoO$_6$ and Sr$_3$Fe$_2$MoO$_9$: for Sr$_2$FeMoO$_6$ with the I4/m tetragonal space group, a = 5.56863(5) Å, c = 7.9007(1) Å, and V = 244.999(4) Å$^3$ for Sr$_2$Fe$_2$Mo$_2$O$_8$ with the same space group, I4/m with Fe and Fe/Mo ordered over the B sites, a = 5.5608(2), c = 7.8471(4) Å, and V = 242.65(2) Å$^3$; and for Sr$_2$Fe$_2$Mo$_2$O$_8$ (Sr$_2$Fe$_2$Mo$_2$O$_8$), a = 5.54301(15) Å, c = 7.8471(4) Å, and V = 241.103(16) Å$^3$. Also, the parameters obtained using PND are in good agreement with the parameters obtained using PXD of a = 5.54905(7) Å and c = 7.8444(2) Å. The cell parameters decrease as the content of Mo decreases due to the increase of the formal oxidation state of the Fe ions.

It is also important to compare the obtained oxygen vacancies in the material. The tendency of this compound to oxygen vacancies, which gives its facile bulk oxygen diffusivity, has been previously reported.15 It also has been observed before that the vacancies localize over O1 (oxygen in the ab plane) and not over O2 (oxygen over the c axis), which are fully occupied. In Table 2, the (Fe/Mo)−O$_2$ bonds are longer than the (Fe/Mo)−O$_1$ ones, thus the oxygen vacancies are localized over the longest B−O distance—an expected result. This could be related to a preferential localization of Fe ions relative to Mo ions around O1 oxygens, since the Fe−O bonds are longer than Mo−O ones and the preferential localization of the vacancies over this position could be well understood in terms of charge differences, since Fe−O bonds are easier to break than Mo−O bonds. This observation has been reported in this compound and in similar ones such as Sr$_2$FeMoO$_6$.15,30 Also it is important to note that O1 is in the position ($x \; y \; 0$) with the possibility of mobility along $x$ and $y$, while O2 has a fixed position ($0 \; 0 \; 1/4$); thus the structural positions of the oxygens are also consistent with the easier removal of O1 compared to O2.

XANES. XANES was used to further characterize the formal oxidation state of Fe and Mo in SrFe$_{0.75}$Mo$_{0.25}$O$_3$. The L$_{2,3}$ edges of the transition metals (T) manifest intense “white line” (WL) features due to dipole allowed transitions into unoccupied final d states. Specifically, the L$_2$ and L$_3$ WL features respectively involve $2p_{3/2}\rightarrow \delta_{3/2}$ and $2p_{1/2}\rightarrow \delta_{3/2}/\delta_{1/2}$ transitions.31 For 4d and 5d row transition metal compounds, the projected d-symmetry density of states features (weighted by matrix element and the multiplet effects) can be seen in the L$_{2,3}$ spectra.32−34 The octahedral crystal electric field, CEF (or ligand field), splits the d states in perovskite related compounds into a lower energy 6X degenerate, $t_{2g}$, and a higher energy 4X degenerate, $e_g$, state. This CEF splitting is reflected in the splitting of the T−L$_{2,3}$ WL into A ($t_{2g}$-related) and B ($e_g$-related) features.

The CEF splitting is particularly large for the perovskites with large B/B′-site valence disparities.33−34 The Mo−L$_{2,3}$ edges for SrFe$_{0.75}$Mo$_{0.25}$O$_3$ (Figure 5) manifest a particularly large (∼4.3 eV) well resolved CEF split A−B spectral structure. Relative to 4d and 5d row L$_{2,3}$ XAS, the smaller d-bandwidth and core hole line broadening, along with the better beamline energy resolution at lower energy, all make the Mo A−B spectral structure in Figure 5 particularly sharp. It should be noted that matrix element and multiplet effects make the A feature more prominent in the L$_2$ edge relative to the L$_3$ edge.32 Both the A−B feature structure and the chemical shift of the T−L$_{2,3}$ edge spectra provide information to estimate the T-valence/configuration. Since the L$_3$ edge is more intense, it will be used for between-compound comparisons for valence/configuration state interrogation.

Figure 6 illustrates the evolution of the Mo−L$_3$ edge structure and chemical shift for a series of octahedral Mo−oxide compounds and elemental Mo. In the sequence of Mo standards, one can see a consistent upward chemical shift with increasing Mo valence and decreasing d count (e.g., see the ~4d$^0$-Mo$^{6+}$, MoO$_3$; ~4d$^1$-Mo$^{5+}$, SrMo$_{0.5}$Fe$_0.5$O$_3$; ~4d$^2$-Mo$^{4+}$ Sm$_2$Mo$_2$O$_7$, and Mo$^{0}$, elemental-Mo sequence). In the octahedrally coordinated Mo−O spectra, the chemical shift is indicated by the centers of mass (in energy) of the WL features.
Despite the large CEF splitting, the SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$ can be seen to have a large positive chemical shift, comparable to \(\sim 4d^0\)-Mo$^{6+}$ standard MoO$_3$. Thus the \(\sim 4d^0\)-Mo$^{6+}$ state identification is supported by its relative Mo--L$_3$ edge chemical shift.

The spectral shape electronic configuration indicator, the Mo--L$_3$ spectra of the perovskite-related compounds (e.g., \(\sim 4d^0\)-Mo$^{6+}$, MoO$_3$ \(\sim 4d^1\)-Mo$^{5+}$, SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$, and \(\sim 4d^2\)-Mo$^{4+}$ Sm$_2$Mo$_2$O$_7$) will now be considered. The relative A/B-count (increasing 4d electron count) in the decrease of relative A-feature intensity with decreasing 4d hole counts. Indeed, comparison of the spectra shows a clear decrease of relative A-feature intensity with decreasing 4d electron count (in the \(\sim 4d^0\), \(\sim 4d^1\), \(\sim 4d^2\), and \(\sim 4d^3\) sequence of standard spectra). The very intense A-feature intensity in the SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$ spectrum clearly supports a \(\sim 4d^0\)-Mo$^{6+}$ state. Thus, the spectral shape again supports Mo$^{6+}$ valence in our compound.

With respect to the K-edge of Fe, both the spectral shape and chemical shift of the main/ pre-edge K-edge features can be used as valence state indicators. The main edge peak features are related to dipole transitions into final p states. The pre-edge features are related to transitions into final 3d states via quadruple and p–d hybridization dipole matrix elements. The final state attractive 3d-electron/core-hole interaction shifts these transitions down in energy into the pre-edge region. Figure 7 shows the Fe–K main edge spectra for LiFePO$_4$ (Fe$^{3+}$), La$_2$VFeO$_6$ (Fe$^{3+}$), and SrFeO$_3$ (\(\sim \text{Fe}^{4+}\)) standards (all have corner sharing Fe–O octahedra albeit distorted in the LiFePO$_4$ case). The chemical shift of the main edge peak, to higher energy, with increasing valence is clear from the standard spectra. The main edge peak intensity degrades substantially between \(\sim \text{Fe}^{3+}\) and \(\sim \text{Fe}^{4+}\) standards with a more modest chemical shift. Indeed, the loss of intensity and the main edge peak constitute a better indicator of increasing Fe valence in this range than does the chemical shift. This peak degradation is presumably due to strong configuration mixing/covacency causing broadened, split 4p feature in \(\sim \text{Fe}^{4+}\) materials.

The Fe–K pre-edges are shown in Figure 7b with the higher Fe valence compound spectra being displaced downward for clarity. The pre-edges of the sequence Fe$^{3+}$, Fe$^{3+}$, and \(\sim \text{Fe}^{4+}\) manifest, with increasing valence: an overall shift in spectral weight toward higher energy, an increase (in most cases) in spectral intensity, and a change in spectral shape. It should be noted that the integrated spectral weight of the LiFePO$_4$ appears relatively enhanced because of the prominent a1 feature. The noncentrosymmetric distortion of the FeO$_6$ octahedra in this compound could introduce p/d mixing and enhance dipole transitions to the pre-edge features. In general, there are two relatively weak and relatively resolved a1–a2 and b1–b2 features in the Fe$^{3+}$ and Fe$^{4+}$ spectra, respectively. The \(\sim \text{Fe}^{4+}\) standard pre-edge manifests three, poorly resolved c1–c2–c3 features. The pre-edge structure spectra of SrFe$_{0.75}$Mo$_{0.25}$O$_3$ have a centrum of energy between that of Fe$^{3+}$ and \(\sim \text{Fe}^{4+}\) standard pre-edge spectra. It also shows evidence for a c1–c2–c3 structure as seen in the \(\sim \text{Fe}^{4+}\) standard spectrum. Finally, the intensity of the c1 feature occurs at lower energy and with a greater intensity than that of the \(\sim \text{Fe}^{4+}\) standard. Thus, the pre-edge results are consistent with the Fe–K main edge results and supporting a mixed Fe$^{3+}$/Fe$^{4+}$ state in the Mo compound. Taken with the \(\sim 4d^0\)-Mo$^{6+}$ main edge results, the Fe–K edge results are consistent with the formally Fe$^{23+}$ state expected on the basis of the obtained formula SrFe$_{0.75}$Mo$_{0.25}$O$_{2.96(2)}$.

**Magnetic Measurements.** The DC susceptibility vs temperature data of SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$ are displayed in Figure 8a. At low temperatures, below 80 K, the zero field cooling (ZFC) and the field cooling (FC) curves diverge, indicating frustration in the system. This frustration could be well understood if we take into account the different magnetic interactions in the material, as will be discussed later. The magnetic susceptibility decreases slowly with increasing temperature until a large change in the slope is observed around 750 K, which indicates the magnetic ordering temperature. This very high ordering magnetic temperature has been previously observed in several comparable complex perovskites, Sr$_3$M$_2$M’O$_6$: e.g., Sr$_3$Fe$_2$TeO$_6$ (\(T_N = 717\) K) or Ba$_3$Fe$_2$TeO$_6$ (\(T_N = 711\) K) and in other very closely related oxides such as the brownmillerite SrFe$_{2.5}$O$_{6(2)}$ (\(T_N \sim 700\) K).

**Figure 6.** A comparison of the Mo–L$_3$ edges of SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$ with octahedral standards: \(\sim 4d^0\)-Mo$^{6+}$, MoO$_3$; \(\sim 4d^1\)-Mo$^{5+}$, SrMo$_3$Fe$_{0.5}$O$_{3.5}$; \(\sim 4d^2\)-Mo$^{4+}$ Sm$_2$Mo$_2$O$_7$; along with elemental Mo.

**Figure 7.** (a) Fe–K main edges of SrFe$_{0.75}$Mo$_{0.25}$O$_{3.5}$ along with several corner shared octahedral Fe-standard spectra: Fe$^{3+}$, LiFePO$_4$; Fe$^{2+}$, La$_2$VFeO$_6$; and \(\sim \text{Fe}^{4+}\), SrFeO$_3$. (b) Fe–K pre-edges for the same compounds.
The lanthanum ferrite LaFeO$_3$ ($T_N \sim 750$ K).\textsuperscript{41} Since the main magnetic interactions in SrFe$_{0.75}$Mo$_{0.25}$O$_3 - \delta$ are between Fe$^{3+}$ cations, a similar $T_N$ to those found in SrFeO$_{2.5}$ and LaFeO$_3$ is expected.

The onset of magnetic ordering at a high temperature ($\sim 750$ K) might be ascribed to the presence of Fe-containing impurities, even if no Fe-containing impurities were detected by PXD or PND. However, the magnetic ordering temperatures of all the possible iron oxide impurities are different from what was observed; e.g., Fe$_3$O$_4$ (magnetite) is the most probable impurity since the sample was prepared in the air, but it exhibits a very distinct $T_N$ of 850 K.\textsuperscript{42} $\alpha$-Fe$_2$O$_3$ (hematite) is a canted antiferromagnet with an even higher $T_N$ of 950 K.\textsuperscript{43} $\beta$-Fe$_2$O$_3$ (maghemite) has a $T_N$ of 920 K and converts to hematite when heated in the air above 750 K. Other possible impurities include SrFeO$_1$ ($T_N = 134$ K), Sr$_2$FeMoO$_6$ ($T_C \sim 420$ K, which has to be prepared in a reducing condition), or Sr$_2$Fe$_2$MoO$_9$ ($T_C = 330$ K).\textsuperscript{44,45,26,27} Therefore, we conclude that the observed magnetism is intrinsic to the SrFe$_{0.75}$Mo$_{0.25}$O$_{3-\delta}$ perovskite phase, as also will be confirmed by Mössbauer spectroscopy.

The isothermal magnetization curves at 5 K and RT in Figure 8b both exhibit an almost antiferromagnetic response with a small hysteresis characteristic of a weak ferromagnetic component. This ferromagnetic component could be attributed to some canting of the spins or some local ferromagnetic interactions in the compound, since several different magnetic interactions are present between Fe$^{3+}$ and Fe$^{4+}$ cations. For example, some residual ferrimagnetic interactions could occur between Fe$^{3+}$–O–Fe$^{4+}$. The magnetic moment obtained from the magnetization curve at 5 K is 0.4 $\mu_B$/mol, which is consistent with the existence of some ferro- or ferrimagnetic interactions in the system.

**Magnetic Structure.** The establishment of a magnetic ordered phase was studied by PND data measured at 14 K. The low temperature PND pattern shows a weak magnetic contribution of some of the low-angle reflections, e.g., (011) and (211), which could be ascribed to a long-range magnetic ordering between Fe spins. The magnetic intensities have been refined in a model that considers an AFM coupling of Fe$^{3+}$ spins lying along the [100] direction. In the antiferromagnetic structure, a net magnetic moment of 0.85(7) $\mu_B$ (Fe atom) at 14 K is in reasonable agreement with the observed remnant magnetization measured at 5 K, of almost 0.4 $\mu_B$ (Fe atom; Figure 8b). The refinement of the magnetic structure is shown in Figure 4b, for the data at 14 K. At RT, the magnetic reflections are not observed, since the value of the magnetic moment is too low to be detected by PND (below 0.2 $\mu_B$). However, the susceptibility curve and the hysteresis loop indicate that the material is still magnetic at RT.

**Mössbauer Spectroscopy.** The Mössbauer spectra of SrFe$_{0.75}$Mo$_{0.25}$O$_{3-\delta}$ measured at 763, 673, 300, and 85 K are shown in Figure 9. The velocity calibration and zero velocity in these figures correspond to those measured with a Fe foil. At 85 K, the material is below the magnetic ordering temperature with a broadened magnetic Zeeman sextet and a paramagnetic component, indicating large partial magnetic ordering of the material.
sample. At 300 K, the doublet is asymmetric, which is usually observed due to the presence of some magnetism. The high temperature spectra show a broad pure doublet at 763 K and an asymmetric broad doublet at 673 K, indicating the possibility of the presence of some remnant magnetism at 673 K. Therefore, the Mössbauer results agree with the PND data, as well as the M(T) results, indicating that the sample is at least partially magnetically ordered near RT, with a very weak magnetic moment. The broadened lines, observed both at low and high temperatures, indicate that the iron ions are in a distribution of different local environments, as expected for such a mixed valent Fe compound.

The origin of this magnetism has to be rationalized. XANES shows a very clear edge of Mo6+ and Fe valence between +3 and +4. Therefore, the absence of an itinerant Mo electron prevents the magnetic coupling across Fe–O–Mo paths in such a way that the only possible magnetic superexchange mechanism could take place via Fe–O–Fe paths, in areas of the crystal containing neighboring Fe spins, and since only 1/4 of B positions are occupied by Mo, the magnetic interactions between Fe occurs all over the material with high coherence. Figure 10 shows a simplified image of the proposed magnetic ordering in the sample; only one layer of B cations is shown for the sake of simplicity. An antiferromagnetic coupling between Fe3+ spins at both B positions would occur since Fe and Mo cations are completely disordered. Some ferrimagnetic interactions between Fe3+ and Fe4+ also appear, and finally 1/4 are nonmagnetic Mo6+ cations. This disordered magnetic configuration would also explain the observed differences found between ZFC and FC curves (Figure 8a). The many different possible interactions in SrFe0.75Mo0.25O3 were characterized in detail. The crystal symmetry was determined to be tetragonal (I4/mcm) and not orthorhombic (Pnma) as previously reported. SrFe0.75Mo0.25O3 orders antiferromagnetically (TN ∼ 750 K), much higher than room temperature. The magnetic structure determined by analysis of the powder neutron diffraction data shows antiferromagnetic coupling between the Fe cations. Mössbauer spectroscopy corroborates the PND results. X-ray absorption spectroscopy indicates the formal oxidation states of the cations to be mixed valent Fe3+/4+ and Mo6+.

![Figure 10. Schematic view of the magnetic interactions in SrFe0.75Mo0.25O3. Fe3+ cations are drawn as light blue and with the bigger spins, Fe4+ are dark blue and with smaller spins, and the nonmagnetic Mo6+ cations are red and without spin.](image-url)

**CONCLUSIONS**

The multifunctional perovskite SrFe0.75Mo0.25O3 was characterized in detail. The crystal symmetry was determined to be tetragonal (I4/mcm) and not orthorhombic (Pnma) as previously reported. SrFe0.75Mo0.25O3 orders antiferromagnetically (TN ∼ 750 K), much higher than room temperature. The magnetic structure determined by analysis of the powder neutron diffraction data shows antiferromagnetic coupling between the Fe cations. Mössbauer spectroscopy corroborates the PND results. X-ray absorption spectroscopy indicates the formal oxidation states of the cations to be mixed valent Fe3+/4+ and Mo6+.

**AUTHOR INFORMATION**

**Corresponding Author**

*E-mail: martha@rutchem.rutgers.edu.

**Notes**

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