







# Magnetic and ferroelectric properties, crystal and magnetic structures of $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$

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## Abstract

The structural investigations of  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound synthesized by solid-state method have been carried by neutron diffraction method in a wide temperature range. The appearance of spontaneous polarization that coexists with ferrimagnetic ordering has been found out at room temperature in strontium hexaferrite partially substituted with In ions. In order to explain the presence of ferroelectric properties in  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound, its crystal structure has been refined within the framework of both centrosymmetric  $P6_3/mmc$  (No. 194) and non-centrosymmetric  $P6_3mc$  (No. 186) space groups. The analysis of refinement results allows to understand microscopic mechanism of appearance ferroelectric properties in strontium hexaferrites.

Keywords: strontium hexaferrites, spontaneous polarization, crystal and magnetic structures

(Some figures may appear in colour only in the online journal)

## 1. Introduction

Solid solutions of M-type strontium ferrites belong to the extensive class of ferrimagnetic oxides with a rather complex crystalline structure. High values of the Curie temperature [1, 2] and saturation magnetization as well as high electrical

resistivity and corrosion resistance allowed hexaferrites to receive a wide technical application as permanent magnets and magnetic information recording devices and microwave equipment. These materials are successfully used for electromagnetic shielding in the range of decimetric and centimetric radiowaves. Hexaferrite substrates with a high

degree of magnetic texture are used to develop high-Q microband devices for microwave electronics.

It is well known that initial hexaferrites  $\text{MFe}_{12}\text{O}_{19}$  ( $\text{M} = \text{Ba}$  and  $\text{Sr}$ ) are isostructural to  $\text{PbO} \cdot 6\text{Fe}_2\text{O}_3$  magnetoplumbite, which crystal structure was first determined by Adelskold [3] in 1938. There are two formula units ( $Z = 2$ ) per unit cell of M-type hexagonal ferrite, i.e. the unit cell contains 2 strontium ions, 24 iron and 38 oxygen ions. The closest packing of  $\text{O}^{2-}$  ions forms several types of voids: octahedra 2a, 12k and  $4f_{\text{VI}}$ ; tetrahedra  $4f_{\text{IV}}$  and trigonal bipyramid 2b, where magnetic Fe ions are located. Such a complex unit cell is characterized by significant crystalline anisotropy that is reflected in the ratio of the lattice parameters  $c/a \approx 3.96$ .

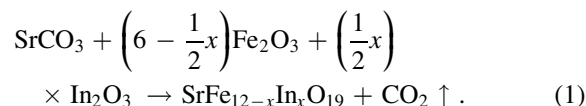
The magnetic structure of hexagonal ferrites is stipulated by superexchange interaction of iron ions through oxygen ions and forms a collinear ferrimagnetic ordering due to strong sublattice exchange.

As a rule, the crystal lattice is described within the framework of the centrosymmetric space group  $\text{P6}_3/\text{mmc}$  (No. 194). However, relatively recent studies of the interactions of barium and strontium hexaferrites with electric fields [4, 5] were discovered the presence of spontaneous electric polarization, its occurrence mechanism remains controversial. In the initial compounds of hexaferrites  $\text{MFe}_{12}\text{O}_{19}$  ( $\text{M} = \text{Ba}$ ;  $\text{Sr}$  or  $\text{Pb}$ ) [4–6], the presence of nonzero spontaneous electric polarization has been explained by the distortion of a single oxygen octahedron  $\text{FeO}_6$  due to the displacement of the iron ion from the center [4, 7]. It was assumed that the distortion of a single oxygen octahedron  $\text{FeO}_6$  occurs below the Curie temperature. The appearance of ferroelectric properties in solid solutions of hexaferrites partially substituted with diamagnetic ions, such as  $\text{BaFe}_{12-x-\delta}\text{Sc}_x\text{Mg}_\delta\text{O}_{19}$  ( $x = 1.6$ ;  $\delta = 0.05$ ) [8], is explained by the presence of a non-collinear magnetic structure. Moreover, a direct link between the electrical and magnetic subsystems was observed in [8]. As a result, taking into account a rather high temperature of the phase transition from ferrimagnetic to paramagnetic order, these compounds can be considered as promising multiferroics.

The present work continues the series our previous investigations [9–13], where we ascertain reasons of appearance spontaneous polarization in hexaferrites partially substituted with different diamagnetic ions ( $\text{In}$ ,  $\text{Ga}$ ,  $\text{Al}$ ,  $\text{Sc}$ ...). The main aim of this paper is to ascertain the microscopic mechanism of appearance of ferroelectric properties in the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound. The difference in the size of ionic radii  $\text{In}$  and  $\text{Fe}$  ions can be observed with diffraction methods that allows to evaluate influence of size effect to distortion internal structural elements. Moreover, the similar small concentration of  $\text{In}$  ions should not lead to appearance of non-collinear magnetic structure that allows to exclude its contribution to origin of spontaneous polarization. The present information is necessary to determine the formation mechanism of ferroelectric properties in hexaferrites.

## 2. Measurement setup

A polycrystalline sample  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  was obtained by the solid-state reaction method. The initial oxides  $\text{Fe}_2\text{O}_3$ ,  $\text{In}_2\text{O}_3$  and carbonate  $\text{SrCO}_3$  were mixed in appropriate proportions and synthesized at  $1200^\circ\text{C}$  (6 h) in air:



The resulting composition was pressed and annealed at  $1300^\circ\text{C}$  (6 h) and then slowly cooled ( $\sim 100^\circ\text{C h}^{-1}$ ). The Curie temperature of the sample was determined by the ponderomotive method by measuring its specific magnetization in the temperature range 300–800 K under external magnetic field of 0.86 T.

Magnetization measurements were carried out on the MPMSXL superconducting quantum SQUID interferometer (DC magnetization mode) in the temperature range from 4 to 300 K. Temperature dependences of magnetic susceptibility were measured by zero field cooled (ZFC) and field cooled warming (FCW) methods. In the first step, the sample was cooled without any magnetic field to 4 K, while its heating was carried out with an applied magnetic field  $H = 100$  Oe. In the second step, the sample was cooled in an applied magnetic field of 100 Oe. In both cases, data were collected during heating the sample.

The geometry of a plane-parallel capacitor was given to the ceramic sample for measuring its polarization. The surface of strontium hexaferrite was coated with silver paste electrodes. Ferroelectric loops were measured using TFAAnalyser 2000 (Aix-ACCT Systems GmbH, Co.). The polarization was determined as the integral of current over time.

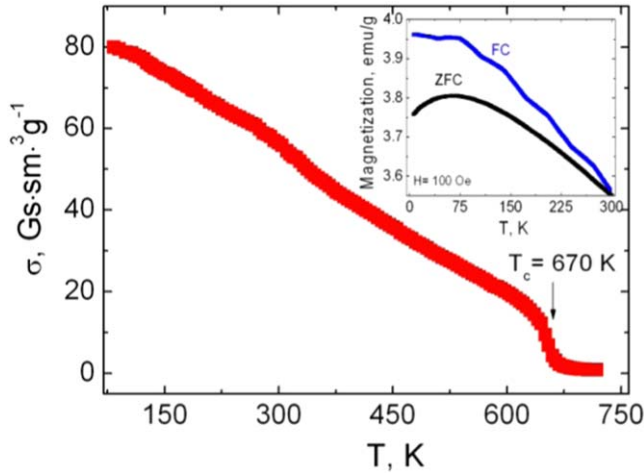
The absence of impurities in the sample was monitored by phase analysis on the x-ray diffractometer Empyrean (firm PANalytical) in  $\text{Co-K}\alpha$ -radiation.

The average crystallite size was determined by the broadening of diffraction peaks using the Scherrer's equation:

$$\langle D \rangle = \frac{K\lambda}{\beta \cos \theta}, \quad (2)$$

where  $K$  is a constant and characterizes the shape and size distribution of crystallites ( $\sim 0.94$  in the approximation of the spherical shape of crystallites),  $\lambda$  is the  $\text{Co-K}\alpha$  radiation wavelength,  $\beta$  is the integral broadening of the diffraction peak at half its height,  $\theta$  is the Bragg angle and  $\langle D \rangle$  is the average particle size in Å.

Neutron diffraction studies of the sample in the temperature range from 1.5 to 350 K were carried out on the diffractometer G 4-1 in the experimental hall of the nuclear center in Laboratoire Léon Brillouin (Saclay, France). The feature of the dual-axis neutron powder diffractometer G 4-1 is the 800-channel detector ( $\text{BF}_3$ ), which allows to cover a range of angles of  $2\theta \sim 80^\circ$  and realize data acquisition at high speed. In addition, it realizes the ability to work in a monochromatic neutron beam with a wavelength of  $2.426$  Å. The measurements were carried out in a helium cryo refrigerator in the temperature range from 1.5 to 350 K.



**Figure 1.** The temperature dependence of the specific magnetization of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound. The Curie temperature is marked. The inset shows temperature dependences of the specific magnetizations of hexaferrite received in ZFC (blue line) and FCW (black line) mode.

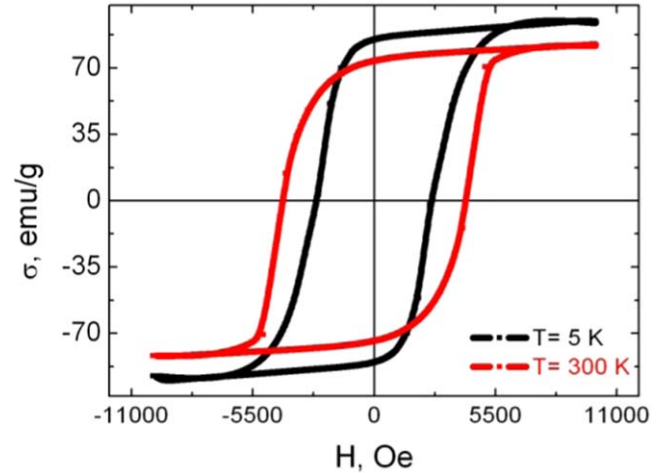
### 3. Experimental results

#### 3.1. Magnetic properties

The temperature dependence of the specific magnetization of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  ferrite is shown in figure 1. The Curie temperature of the compound is  $\sim 670$  K, which is lower than the ferrimagnetic—paramagnet phase transition temperature  $T_c \sim 732$  K for the initial  $\text{SrFe}_{12}\text{O}_{19}$  composition [14, 15]. This behavior of the Curie temperature confirms the replacement of magnetoactive iron ions with diamagnetic indium ions. As a result, the magnetic order is destroyed at lower temperatures, which is consistent with the data [15, 16].

The temperature dependencies of the magnetization of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound collected in ZFC and FCW modes are shown in the insert figure 1. The magnetization curve of ZFC is located below the FCW and first shows growth passing through the maximum, and then decreases when the sample is heated. The distinction between ZFC and FC curves was previously observed in ensembles of superparamagnetic particles [17], in spin glasses [18], as well as in macroscopic magnets with frustration of exchange bonds, with topological disorder, with structural defects, and even in ordered ferromagnets with significant magnetic anisotropy [19]. In our case, the sample can be represented as a system consisting of isolated particles with uniaxial anisotropy, in which the easy magnetization axes are randomly oriented in space. In a zero magnetic field, magnetizations of crystallites are ordered along their easy magnetization axes.

In the process of cooling the sample at  $H = 0$ , the magnetization of crystallites freezes at a certain blocking temperature ( $T_b$ ), while their distribution in the sample remains chaotic. The value of  $T_b$  corresponds to the temperature below which the energy of thermal fluctuations in the crystallite is substantially less than the energy of the magnetic anisotropy. The value of blocking temperature can be



**Figure 2.** The field dependences of the specific magnetization  $\sigma(H)$  of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound measured at 5 (black line) and 300 (red line) K.

determined by the Bean–Livingston equation (3) [20–22]:

$$T_B = \frac{K_V V}{25 k_B}. \quad (3)$$

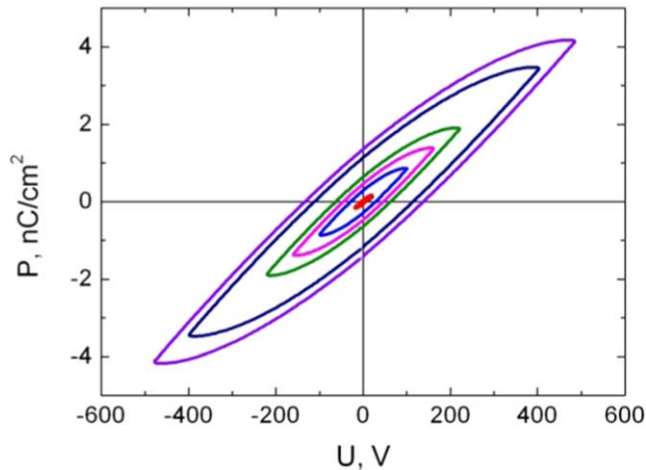
The presence of a broad maximum on the ZFC curve (see inset figure 1) indicates the dispersion of particle sizes and it can be the result of the sum of narrow maxima for particles with various sizes. Therefore, the maximum is determined by the contribution of the volume fractions of particles with a certain size. When the sample is cooled in the magnetic field, the direction of the magnetization in each crystallite is determined by the competition between the magneto crystalline anisotropy energy and Zeeman energy. The magnetization of crystallites tends to be oriented in the direction of the external magnetic field. However, in order to rotate the magnetic moment of a particle in the direction corresponding to the minimum of energy, it is necessary to overcome the energy barrier:

$$\Delta E \approx K_V \times V, \quad (4)$$

where  $K_V$  is the volume anisotropy constant;  $V$  is the volume of the magnetic particle, which should be overcome by magnetic field in order to change the orientation of the magnetic moment of the particles.

At a sufficiently large intensity of the external magnetic field, magnetic moments of all crystallites are aligned along the field, and the shape of the temperature dependence curve in this case is determined by an increase in the spontaneous magnetization of the material.

The magnetic hysteresis loops of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound measured at  $T = 5$  and 300 K are shown in figure 2. The magnitude of the residual magnetization  $M_r$ , measured after switching off the external magnetic field, reaches values of  $85 \text{ emu g}^{-1}$  and  $74 \text{ emu g}^{-1}$  at temperatures 5 K and 300 K, respectively. A similar difference in the residual magnetization values is the result of a decrease in the spin collinearity degree of  $\text{Fe}^{3+}$  at 300 K in particles due to an increase in temperature fluctuations [23, 24].



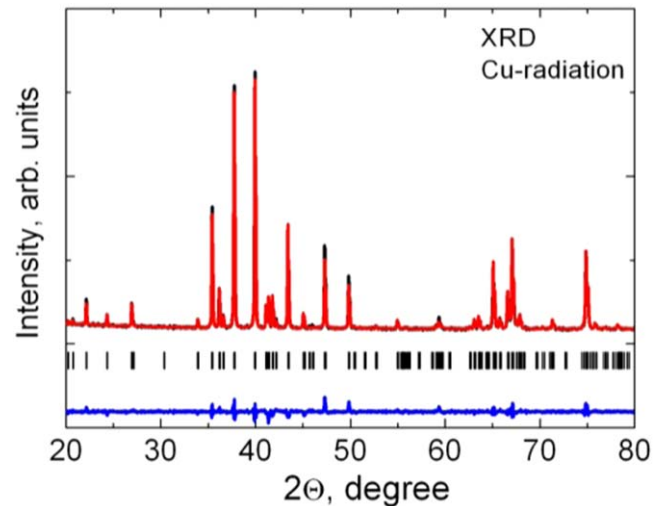
**Figure 3.** Ferroelectric hysteresis loops of  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  ceramic.

Figure 2 shows the unusual behavior of the coercive force ( $H_c$ ) of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound in comparison, for instance, with  $\text{Fe}_3\text{O}_4$  [25] or  $\text{Bi}_{1-x}\text{La}_x\text{FeO}_3$  [26], where the coercive force increases as the temperature is decreased. In this study, the magnitude  $H_c$  decreases in  $\sim 1.6$  times from  $H_c \sim 4185$  Oe to  $H_c \sim 2560$  Oe, when the ambient temperature is decreased from 300 to 5 K. A decrease of coercivity with decreasing temperature was previously observed in  $\text{SrFe}_{12}\text{O}_{19}$  [27] and  $\text{BaFe}_{12}\text{O}_{19}$  [28] ferrites. The similar behavior was explained by the imperfection of the crystal structure due to its proximity to the amorphous state. In our case the sample  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  was annealed at  $1300^\circ\text{C}$  (6 h). This fact and the high quality of x-ray pattern (see below) confirm the good state of the crystal structure. Earlier studies of ferromagnets [20] revealed a significant increase in their coercive force during the transition from a multi-domain to a single-domain state. In addition, according to Weiss's theory, the magnetization in ferro- or ferrimagnetics is accompanied by the growth of some domains at the expense of others. Therefore, in this case, it can be assumed that the multi-domain crystallite has less Bloch walls with increasing temperature due to growing domain size with temperature.

### 3.2. Ferroelectric properties

The electric voltage dependence of the polarization  $P$  for strontium hexaferrite  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  measured at room temperature is shown in figure 3. A rapid increase in the saturation polarization of strontium ceramic was observed under electric field. The presence of hysteresis loops and the appearance of remanent electrical polarization at room temperature, which can be reversed applying an external electric field, allows to characterize strontium ferrite as ferroelectric. The maximal values of polarization ( $P_{\text{max}}$ ) and spontaneous polarization ( $P_r$ ) as well as the coercive force ( $E_c$ ) determined from the hysteresis loops have the following values:  $4.17 \text{ nC cm}^{-2}$ ,  $1.35 \text{ nC cm}^{-2}$  and  $138 \text{ V}$ , respectively.

Ferroelectric properties were earlier discovered in the initial compound  $\text{SrFe}_{12}\text{O}_{19}$  [5], nevertheless, their occurrence mechanism in the solid solutions of hexaferrites remains



**Figure 4.** The general view of x-ray pattern of  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  ferrite refined with the Rietveld method in the framework of the generally accepted centrosymmetric space group  $P6_3/\text{mmc}$  (No. 194).

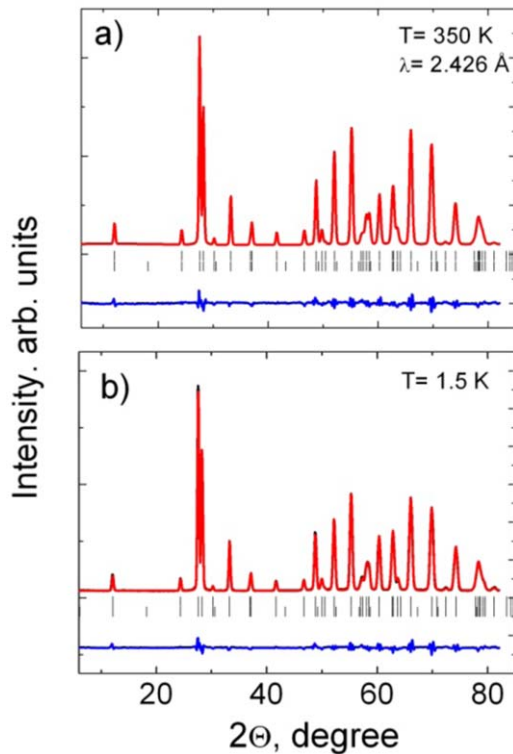
controversial. In particular, the main reason of the generation of ferroelectricity in the initial compounds  $\text{AFe}_{12}\text{O}_{19}$  (where  $\text{A} = \text{Pb}, \text{Ba}, \text{Sr}$ ) [4–6] was suggested the distortion of one of the oxygen octahedral. Meanwhile, in [8] the main reason of the ferroelectric properties in solid solutions was called the occurrence of a non-collinear magnetic structure.

One more explanation of the appearance of spontaneous polarization in solid solutions of barium ferrites was proposed in previously published [29–31], where ferroelectricity is a consequence of non-centrosymmetric distortion of the unit cell. The discussion of possible reasons of the occurrence of ferroelectric properties in the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound will be given below.

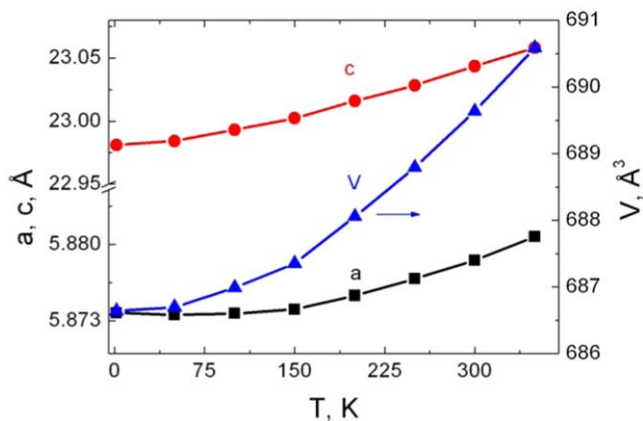
### 3.3. Crystal structure

According to x-ray diffraction data, the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  sample is homogeneous. As was mentioned above, the crystal structure of strontium hexaferrite is well described by the centrosymmetric space group  $P6_3/\text{mmc}$  (No. 194) and correspond ICDD card 00-033-1340. The average crystallite size  $\langle D \rangle \sim 1180 \text{ Å}$  was determined from the broadening of the diffraction peak (0211) by Scherrer's equation (1). Figure 4 shows a general view of the x-ray pattern of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound, which crystal structure was refined by the Rietveld method in the framework of the space group  $P6_3/\text{mmc}$ . However, the description of the crystal structure with the centrosymmetric SG  $P6_3/\text{mmc}$  means that the unit cell has an inversion center, which contradicts the appearance of ferroelectric properties in the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  ferrite shown in figure 3.

It was proposed to consider the crystal structure of hexaferrites in the framework of non-centrosymmetric SG  $P6_3\text{mc}$  (No. 186), according to [29]. However, no information was given about structural parameters such as unit cell parameters, atomic coordinates, etc.



**Figure 5.** Neutron diffraction patterns of  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound measured on diffractometer G 4-1 at temperatures 350–(a) and 1.5 K–(b) and processed by the Rietveld method. Experimental (black curve) and calculated (red curve) curves, as well as the difference function normalized to the statistical error (blue curve) are shown. Vertical bars indicate the calculated positions of the diffraction peaks for the crystalline (top) and magnetic (bottom) structures of strontium hexaferrite, calculated in the framework of the non-centrosymmetric SG  $\text{P6}_3\text{mc}$  (No. 186).



**Figure 6.** Temperature dependence of structural parameters  $a$  and  $c$  (left scale) and volume  $V$  (right scale) of unit cell for  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound.

Therefore, in order to clarify the real crystal structure of strontium ferrite, neutron diffraction patterns were measured in a wide temperature range from 1.5 to 350 K. The general view of neutron diffraction patterns is shown in figure 5. The temperature dependencies of the crystal structure parameters ( $a$  and  $c$ ) and the volume of the unit cell ( $V$ ) are shown in figure 6. The area, where thermal extension of lattice

parameters is absent, was observed in the low-temperature region. For the first time, such behavior was discovered in solid solutions of barium hexaferrite [32]. There is a difference in the temperature dependencies of the lattice parameters  $a$  and  $c$ .

In particular,  $c$  parameter has insignificant variation in the temperature range from 1.5 to 50 K, while  $a$  parameter practically does not change in wider temperature area from 1.5 to 150 K. This difference in the changes of the unit cell parameters  $a$  and  $c$  can be related with strong magneto crystalline anisotropy of strontium hexaferrite. While the presence of a temperature range, where thermal expansion is absent, can be explained by mutual rotations and tilts of the oxygen polyhedra, similarly to [32].

The refinement of the crystal structure of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound was carried out within both the centrosymmetric  $\text{P6}_3/\text{mmc}$  (No. 194) and non-centrosymmetric  $\text{P6}_3\text{mc}$  (No. 186) space groups. The results of the refinement are shown in tables 1 and 2.

The comparison of the relevance factors ( $R_{\text{exp}}$ ,  $R_{\text{wp}}$ ,  $\chi^2$ ) from tables 1 and 2 and difference curves, normalized to the statistical errors from figure 5, indicates that the crystal structure of strontium hexaferrite can be described quite well by both the centrosymmetric and noncentrosymmetric space group in a wide temperature range. However, figures of merit for non-centrosymmetric space group have slightly lower values than for the centrosymmetric space group (see tables 1 and 2). Therefore, observation of ferroelectric properties in the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound shown in figure 3 and less magnitudes figures of merit in the case of the non-centrosymmetric space group (see table 2) allows us to give preference to SG  $\text{P6}_3\text{mc}$  (No. 186).

Fragments of the crystal structure of unit cells for centrosymmetric and non-centrosymmetric space groups are shown in figures 7(a)–(e). In spite of the completely equal values of lattice parameters for unit cells both space groups (see figure 6 and tables 1 and 2), there is a difference in the distortion of individual structural elements.

To evaluate the distortion of the oxygen octahedron ( $S_{\text{BO6}}$ ), the equation was used:

$$S_{\text{FeO}_6}(T) = \frac{100}{\langle d(T) \rangle} \times \sqrt{\frac{\sum_{i=1}^6 (d_i(T) - \langle d(T) \rangle)^2}{5}}, \quad (5)$$

where  $d_i(T)$  is the distance between iron ions and the nearest oxygen ions;  $\langle d(T) \rangle$  is the average  $\langle \text{Fe-O} \rangle$  distance in the oxygen octahedron.

The distortions of the oxygen octahedron at different ambient temperatures are shown in figures 8(a) and (b).

According to figure 8, in the case of the description of the unit cell of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound with the centrosymmetric space group No. 194, the oxygen octahedra of Fe1 ions located in crystallographic positions (2a) have no distortions in the temperature range from 1.5 to 350 K. In the case of a unit cell description in the framework of the non-centrosymmetric space group (No. 186), the distortions of a similar oxygen octahedron achieve  $\sim 5\%$ – $7\%$  (see figure 8(a)) in the studied temperature range.

**Table 1.** Crystal structure parameters, figures of merit and atomic positions refined in the framework of centrosymmetric SG  $P6_3/mmc$  (№ 194) with the Rietveld method for different temperatures. Atoms are located in positions: Sr (2d) (2/3, 1/3, 1/4), Fe1 (2a) (0, 0, 0); Fe2 (2b) (0, 0, 1/4); (Fe/In)3 (4f<sub>IV</sub>) (1/3, 2/3, z); (Fe/In)4 (4f<sub>VI</sub>) (1/3, 2/3, z); (Fe/In)5 (12k) (x, 2x, z); O1 (4e) (0, 0, z); O2 (4f) (1/3, 2/3, z); O3 (6h) (x, 2x, 1/4); O4 (12k) (x, 2x, z); O5 (12k) (x, 2x, z).

Atomic positions	Temperature			
	50 K	150 K	250 K	350 K
<i>a</i> , (Å)	5.8736(1)	5.8741(2)	5.8769(2)	5.8807(2)
<i>c</i> , (Å)	22.9843(6)	23.0023(7)	23.0285(7)	23.0581(8)
<i>V</i> , (Å <sup>3</sup> )	686.69(3)	687.35(3)	688.79(4)	690.58(4)
Fe3/ In3 (4f <sub>IV</sub> )				
<i>z</i>	0.0316(5)	0.0307(6)	0.0311(6)	0.0309(7)
Fe4/ In4(4f <sub>VI</sub> )				
<i>z</i>	0.1919(5)	0.1927(5)	0.1935(5)	0.1936(5)
Fe5/ In5 (12k)				
<i>x</i>	0.1837(11)	0.1840(13)	0.1851(12)	0.1854(12)
<i>z</i>	−0.1087(1)	−0.1088(2)	−0.1088(2)	−0.1089(2)
O1 (4e)				
<i>z</i>	0.1432(5)	0.1442(6)	0.1438(7)	0.1448(7)
O2 (4f)				
<i>z</i>	−0.0374(6)	−0.0384(7)	−0.0371(7)	−0.0369(8)
O3 (6h)				
<i>x</i>	0.1846(34)	0.1802(51)	0.1723(49)	0.1662(49)
O4 (12k)				
<i>x</i>	0.1604(21)	0.1635(29)	0.1646(27)	0.1659(29)
<i>z</i>	0.0546(3)	0.0546(3)	0.0553(3)	0.0555(3)
O5 (12k)				
<i>x</i>	0.4952(23)	0.4947(27)	0.4939(27)	0.4938(30)
<i>z</i>	0.1542(2)	0.1538(3)	0.1538(3)	0.1535(3)
<i>R</i> <sub>wp</sub> , %	5.34	6.09	6.07	6.58
<i>R</i> <sub>exp</sub> , %	3.90	4.01	3.93	4.38
<i>R</i> <sub>B</sub> , %	3.25	3.43	4.32	5.23
<i>R</i> <sub>Mag</sub> , %	3.64	4.26	5.19	6.91
$\chi^2$	1.88	2.31	2.39	2.26

Another behavior is observed in oxygen surroundings of Fe<sub>4</sub> ion (see figures 7(c) and (d)). In the case of centrosymmetric SG No. 194, distortions of the oxygen octahedra that surround crystallographic positions (4f<sub>VI</sub>) monotonously increase from  $\sim 4.6\%$  to  $7.3\%$  as the ambient temperature is increased up to 350 K (see figure 8(b)). In the case of a non-centrosymmetric SG 186, there is a difference in the distortion of neighboring oxygen octahedra (see figure 8(b)). It should be noted that the distortion of oxygen polyhedra can be a reason of the appearance of local electric dipoles for both centrosymmetric and non-centrosymmetric space groups. However, unequal distortions of neighboring oxygen octahedra are the main reason of uncompensated local dipole moments that leads to the appearance of ferroelectric properties. The influence of the temperature factor on the magnitude of local electric dipoles is shown in figures 9(a) and (b).

Therefore, taking into account the presence of an inversion center in the centrosymmetric unit cell, there will be no spontaneous polarization due to the compensation of local electric dipoles. On the contrary, in the non-centrosymmetric unit cell, unequal distortion of neighboring oxygen surroundings (see figures 8(a) and (b)) is the only cause of the occurrence of spontaneous polarization.

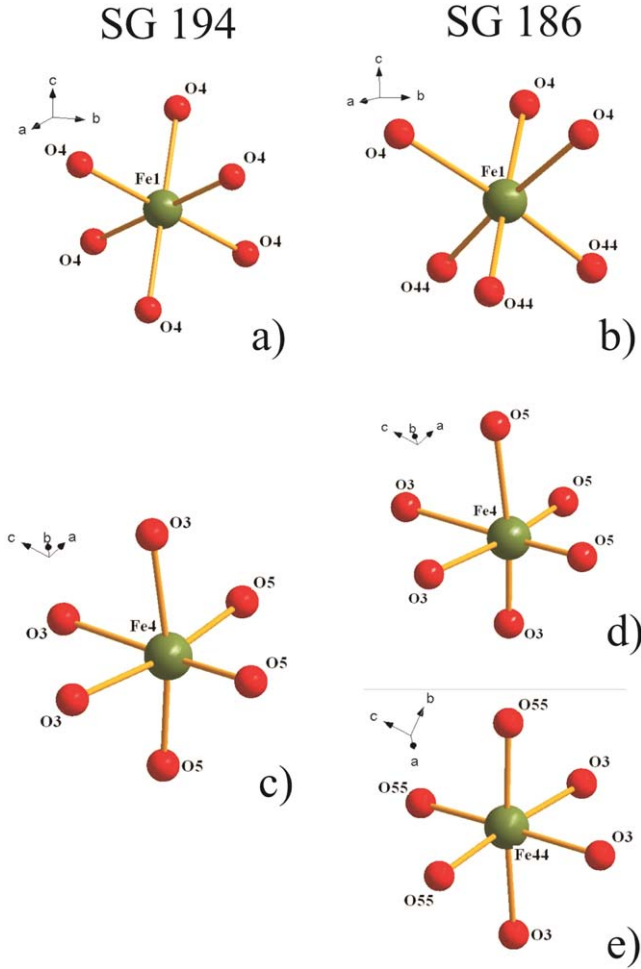
### 3.4. Magnetic structure

As was mentioned above, the main feature of the neutron diffraction method is a direct study of magnetic structure materials. Neutron diffraction patterns were measured in a wide temperature range from 1.5 to 350 K, which is well below the paramagnetic–ferrimagnetic phase transition temperature ( $T_c \sim 670$  K) for a given compound (see figure 1). In figure 5, the bottom row of vertical bars indicates the positions of diffraction peaks calculated by the Rietveld method for the magnetic structure of SrFe<sub>11.9</sub>In<sub>0.1</sub>O<sub>19</sub> hexaferrite. The absence of additional magnetic peaks on neutron diffraction patterns indicates that the wave vector of this ferrimagnetic structure:  $\mathbf{k} = [0, 0, 0]$ .

The magnetic structure of SrFe<sub>11.9</sub>In<sub>0.1</sub>O<sub>19</sub> ferrite satisfies to the model proposed by Gorter [33]. It has ferrimagnetic order, where all magnetic moments of Fe<sup>3+</sup> ions are directed along *c* axis, which is easy magnetization axis [34]. This ferrimagnetic structure is formed by two ferromagnetic sublattices that directed in the opposite direction [35]. The magnetic moments of Fe<sup>3+</sup> ions located in octahedral (2a and 12k) and bipyramidal (2b) oxygen surroundings are oriented in opposite to the magnetic moments of iron ions occupying positions (4f<sub>IV</sub> and 4f<sub>VI</sub>).

**Table 2.** Crystal structure parameters, figures of merit and atomic positions refined in the framework of centrocymmetric SG P6<sub>3</sub>mc (No. 186) with the Rietveld method for different temperatures. Atoms are located in positions: Sr (2b) (2/3, 1/3, z); Fe1 (2a) (0, 0, z); Fe2 (2a) (0, 0, z); (Fe/In)3 (2b) (1/3, 2/3, z); (Fe/In)33 (2b) (1/3, 2/3, z); (Fe/In)4 (2b) (1/3, 2/3, z); (Fe/In)44 (2b) (1/3, 2/3, z); (Fe/In)5 (6c) (x, 2x, z); (Fe/In)55 (6c) (x, 2x, z); O1 (2a) (0, 0, z); O11 (2a) (0, 0, z); O2 (2b) (1/3, 2/3, z); O22 (2b) (1/3, 2/3, z); O3 (6c) (x, 2x, z); O4 (6c) (x, 2x, z); O5 (6c) (x, 2x, z); O55 (6c) (x, 2x, z).

Atomic positions	Temperature			
	50 K	150 K	250K	350 K
<i>a</i> , (Å)	5.8736(1)	5.8741(2)	5.8769(2)	5.8807(2)
<i>c</i> , (Å)	22.9843(6)	23.0023(7)	23.0285(7)	23.0581(8)
<i>V</i> , (Å <sup>3</sup> )	686.69(3)	687.35(3)	688.79(4)	690.58(4)
Sr (2b)				
<i>z</i>	0.2663(14)	0.2618	0.2587	0.2570(39)
Fe1 (2a)				
<i>z</i>	−0.0052(16)	−0.0085(19)	−0.0128(24)	−0.0106(19)
Fe2 (2a)				
<i>z</i>	0.2467(19)	0.2459(21)	0.2443(28)	0.2422(49)
Fe3 (2b)				
<i>z</i>	0.0261(13)	0.0233(19)	0.0232(19)	0.0239(43)
Fe33 (2b)				
<i>z</i>	0.4638(16)	0.4594(26)	0.4573(25)	0.4618(35)
Fe4 (2b)				
<i>z</i>	0.1998(9)	0.1949(17)	0.1929(21)	0.2018(26)
Fe44(2b)				
<i>z</i>	0.3107(10)	0.3049(23)	0.3024(28)	0.3087(23)
Fe5 (6c)				
<i>x</i>	0.1751(32)	0.1759(39)	0.1778(39)	0.1772(45)
<i>z</i>	0.8880(7)	0.8854(12)	0.8845(13)	0.8900(18)
Fe55 (6c)				
<i>x</i>	0.8279(31)	0.8255(34)	0.8209(54)	0.8183(52)
<i>z</i>	0.1089(7)	0.1057(11)	0.1052(9)	0.1104(23)
O1 (2a)				
<i>z</i>	0.1520(19)	0.1482(29)	0.1489(29)	0.1492(44)
O11 (2a)				
<i>z</i>	0.3406(20)	0.3388(28)	0.3404(33)	0.3491(42)
O2 (2b)				
<i>z</i>	0.9375	0.9375	0.9465(69)	0.9487(52)
O22 (2b)				
<i>z</i>	0.5305(23)	0.5273(31)	0.5318(70)	0.5272(69)
O3 (6c)				
<i>x</i>	0.1864(42)	0.1820(68)	0.1758(63)	0.1749(64)
<i>z</i>	0.2517(10)	0.2503(11)	0.2511(31)	0.2532(16)
O4 (6c)				
<i>x</i>	0.1806(47)	0.1810(54)	0.1797(61)	0.1739(83)
<i>z</i>	0.0452	0.0439	0.0414(12)	0.0452
O44 (6c)				
<i>x</i>	0.1420(43)	0.1433(49)	0.1438	0.1521(72)
<i>z</i>	0.4344(9)	0.4332(8)	0.4294(13)	0.4340(13)
O5 (6c)				
<i>x</i>	0.4599(39)	0.4618	0.4637(49)	0.4636(48)
<i>z</i>	0.1581	0.1529(21)	0.1544	0.1609(33)
O55 (6c)				
<i>x</i>	0.5041(57)	0.5035(57)	0.4981(86)	0.4973(111)
<i>z</i>	0.3462(9)	0.3432(21)	0.3433(11)	0.3469(32)
<i>R</i> <sub>wp</sub> , %	4.78	5.39	5.30	5.54
<i>R</i> <sub>exp</sub> , %	3.88	3.99	3.90	4.35
<i>R</i> <sub>B</sub> , %	1.64	1.80	2.49	2.69
<i>R</i> <sub>Mag</sub> , %	1.90	1.96	3.05	3.99
$\chi^2$	1.52	1.83	1.85	1.63



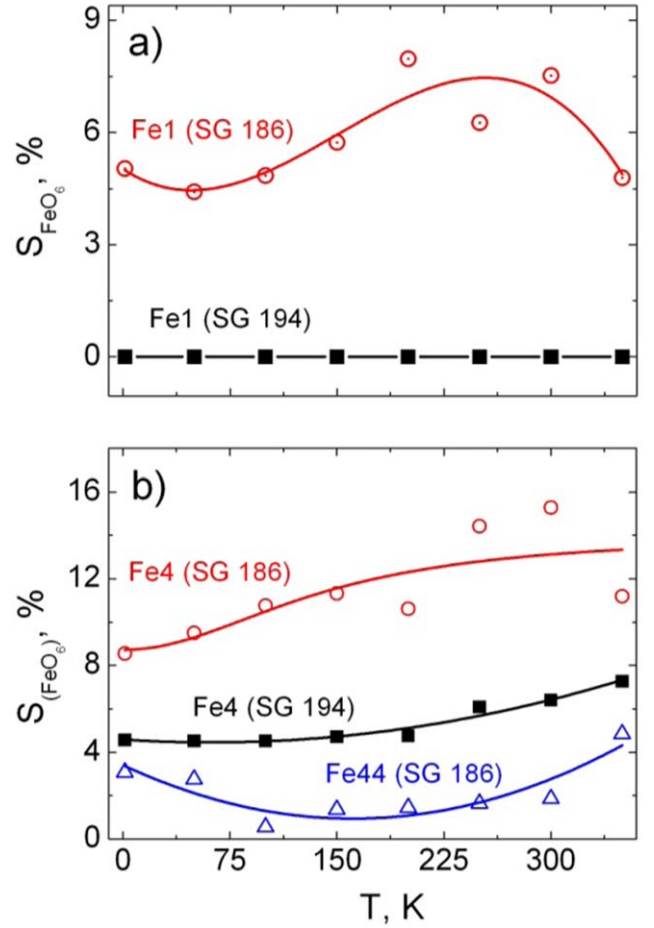
**Figure 7.** Fragments of unit cell: oxygen octahedron around Fe1 ion—undistorted for SG 194 (a) and distorted for SG 186; and oxygen octahedra around Fe4/Fe44 for SG 194 (c) and for SG 186 (d) and (e). (Oxygen ions (O) are designed by red spheres and iron ions (Fe) are designed by dark green spheres.)

As was mentioned above the unit cell was refined in the framework both centrosymmetric ( $P6_3/mmc$ ) and non-centrosymmetric ( $P6_3mc$ ) space groups therefore magnetic moments were also calculated separately for each space group [36]. The temperature dependencies of magnetic moments of  $Fe^{3+}$  ions in various crystallographic positions are shown in figures 10(a) and (b). Values of magnetic moments at different crystallographic positions in the entire temperature range do not exceed  $5 \mu_B$ , which is characteristic for the magnetic moment of  $Fe^{3+}$ . It is an indirect confirmation of the correctness of the calculations [37].

The total magnetic moment per formula unit of strontium ferrite ( $SrFe_{12}O_{19}$ ) with a centrosymmetric structure can be calculated with equation [18]:

$$M_{total}(T) = 1[m_{Fe1}(T)] + 1[m_{Fe2}(T)] - 2[m_{Fe3}(T)] - 2[m_{Fe4}(T)] + 6[m_{Fe5}(T)]. \quad (6)$$

In the case of a non-centrosymmetric structure, the total



**Figure 8.** Temperature dependencies of distortions of oxygen octahedra around: Fe1 ion—(a); and Fe4 and Fe44 ions—(b). (Oxygen octahedra of unit cell for SG 194 are designated with solid squares, and for SG 186—with empty circles and triangles.)

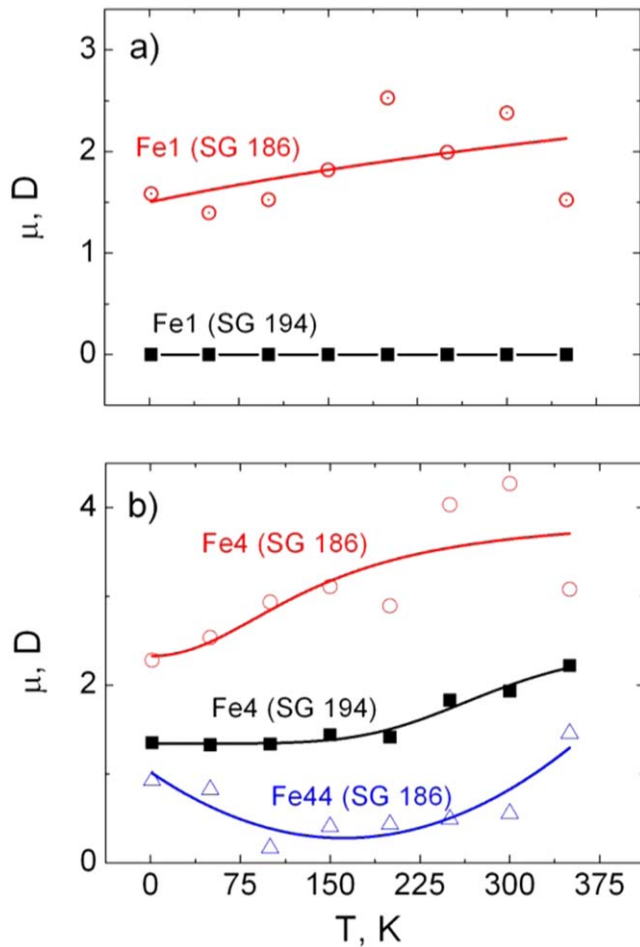
magnetic moment was calculated as:

$$M_{total}(T) = 1[m_{Fe1}(T)] + 1[m_{Fe2}(T)] - 1[m_{Fe3}(T)] - 1[m_{Fe33}(T)] - 1[m_{Fe4}(T)] - 1[m_{Fe44}(T)] + 3[m_{Fe5}(T)] + 3[m_{Fe55}(T)], \quad (7)$$

where  $m_i$  is the magnetic moment of  $Fe^{3+}$  ion in the  $i$ th sublattice.

Temperature dependencies of the total magnetic moments ( $M_{total}$ ) are shown in figure 11. It should be noted that the values of  $M_{total}$ , both for centrosymmetric and non-centrosymmetric structures, have insignificant difference from each other, which does not allow giving the preference to one of them [38]. Moreover, the general view of the magnetic structure is comparable both for SG 194 and SG 186 and is shown in the insert figure 11.

The magnitude of the total magnetic moment has slightly less value than  $20 \mu_B$  per formula unit (f.u.) in the low-temperature region [39]. It is due to the influence of diamagnetic In ions, which disrupts the exchange interaction between neighboring  $Fe^{3+}-O-Fe^{3+}$  ions. The decrease of total magnetic moment is observed with increasing of the ambient temperature. Such behavior is the result of magnetic

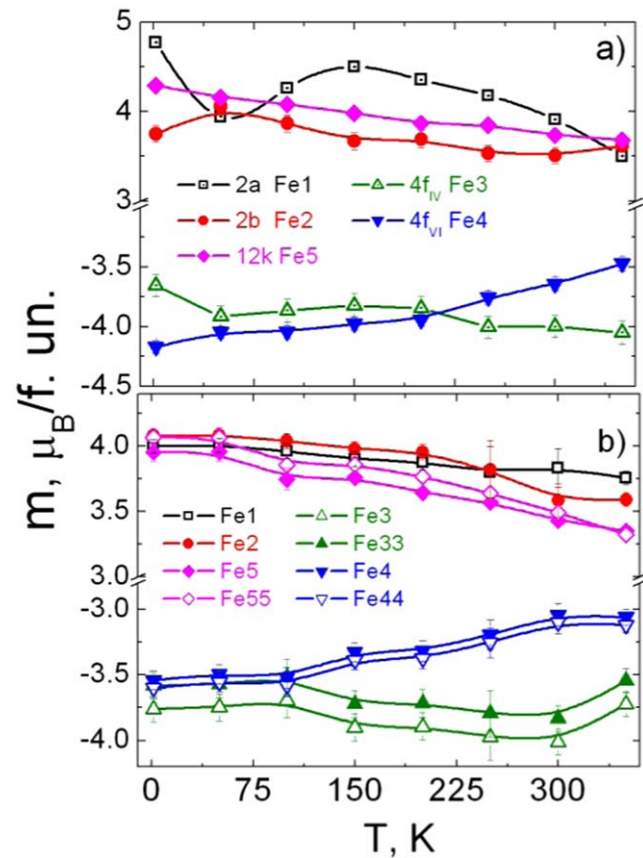


**Figure 9.** Temperature dependencies of electric dipoles as result of distortions of oxygen octahedrons around: Fe1—(a); and Fe4 and Fe44—(b). (Electric dipoles of unit cell for SG 194 are designated with solid squares, and for SG 186—with empty circles and triangles.)

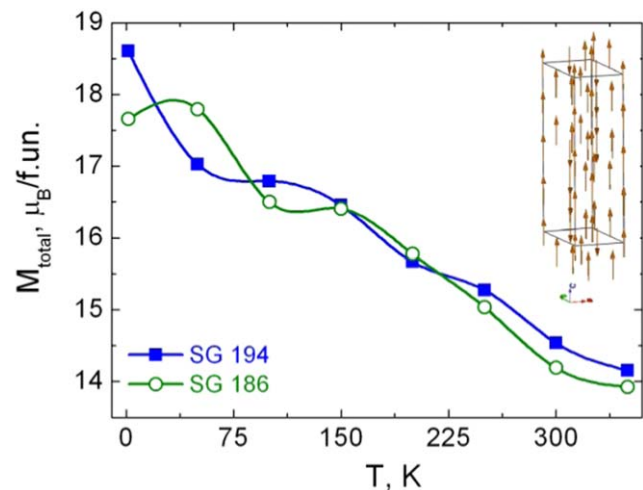
order disrupting due to an increase in thermal oscillations of the ions forming the crystal lattice.

#### 4. Conclusions

The presence of spontaneous polarization at room temperature was observed in strontium ferrite partially substituted with diamagnetic In ions. The appearance of spontaneous polarization in strontium hexaferrite indicates its ferroelectric property that contradicts to the generally accepted description of the crystal structure of similar materials with centrosymmetric SG  $P6_3/mmc$ . The discrepancy between FC and ZFC temperature dependencies of magnetizations indicates the presence of a magnetically inhomogeneous system. It was found that the coercive field decreased in  $\sim 1.6$  times as the ambient temperature increased from 5 to 300 K, which may be the result of growing domain size with temperature. The fact that the ferroelectric properties of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound coexist with ferrimagnetic ordering and its high Curie temperature ( $T_c \sim 670$  K) allows to consider this material as a possible multiferroic.



**Figure 10.** Temperature dependencies of local magnetic moments for suitable crystallographic positions for SG 194—(a) and SG 186—(b). Designations in the figures are follow: Fe1 -  $\square$ ; Fe2 -  $\bullet$ ; Fe3 -  $\triangle$ ; Fe33 -  $\blacktriangle$ ; Fe4 -  $\nabla$ ; Fe44 -  $\nabla$ ; Fe5 -  $\blacklozenge$ ; Fe55 -  $\diamond$ .



**Figure 11.** Temperature dependencies of total magnetic moments for SG 194—(a) and SG 186—(b). In inset: general view of magnetic structure of  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound is shown.

The crystal and magnetic structures of the  $\text{SrFe}_{11.9}\text{In}_{0.1}\text{O}_{19}$  compound was studied by the neutron diffraction method in a wide temperature range from 1.5 to 350 K. Insignificant changes of the lattice parameters were observed in strontium hexaferrites in the temperature range

from 1.5 to 150. It was explained by mutual rotations and tilts of oxygen surroundings.

In order to determine the real type of the crystal structure of strontium ferrite, it was refined within the framework both centrosymmetric  $P6_3/mmc$  (No. 194) and non-centrosymmetric  $P6_3mc$  (No. 186) space groups. It was shown that the crystal structure of the  $SrFe_{11.9}In_{0.1}O_{19}$  compound can be described quite well by both space groups. However, less values figures of merit ( $R_{exp}$ ,  $R_{wp}$  and  $\chi^2$ ) and the presence of ferroelectric properties allows to give preference to the non-centrosymmetric  $P6_3mc$  (No. 186) space group. Moreover, using the non-centrosymmetric  $P6_3mc$  (No. 186) space group for describing crystal structure of the  $SrFe_{11.9}In_{0.1}O_{19}$  allows to explain the appearance of spontaneous polarization as a result of unequal distortions of neighboring oxygen surroundings because of absence inversion center in the unit cell.

The magnetic structure of the  $SrFe_{11.9}In_{0.1}O_{19}$  compound is well described by Gorter's model, according to which magnetic moments of all  $Fe^{3+}$  ions are ordered along hexagonal  $c$  axis. The values of the total and local magnetic moments in various crystallographic positions in a wide temperature range were determined for unit cell described within the framework of both centrosymmetric  $P6_3/mmc$  (No. 194) and non-centrosymmetric  $P6_3mc$  (No. 186) space groups.

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