Once you have Acrobat Reader open on your computer, click on the [Comment](#) tab at the right of the toolbar:

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How to use it

- Highlight a word or sentence.
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standard framework for the analysis of microeconomic behavior. Nevertheless, it also led to the development of a new class of strategic form games. The number of competitors in the market is that the structure of the game is a main component. At the strategic level, are exogenous variables and important works on entry by firms (M. Henceforth) we open the 'black box'.

2. [Strikethrough \(Del\)](#) Tool – for deleting text.

Strikes a red line through text that is to be deleted.

How to use it

- Highlight a word or sentence.
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there is no room for extra profits as mark-ups are zero and the number of firms (set) values are not determined by the market. Blanchard and Kiyotaki (1987), perfect competition in general equilibrium models of aggregate demand and supply in the classical framework assuming monopoly power between an exogenous number of firms.

3. [Add note to text](#) Tool – for highlighting a section to be changed to bold or italic.

Highlights text in yellow and opens up a text box where comments can be entered.

How to use it

- Highlight the relevant section of text.
- Click on the [Add note to text](#) icon in the Annotations section.
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dynamic responses of mark-ups consistent with the VAR evidence.

sation by Markov processes. The number of competitors and the impact of demand shocks on the demand.

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- Type the comment into the yellow box that appears.

standard and supply shocks. Most of the standard framework for the analysis of microeconomic behavior. The number of competitors and the impact of demand shocks on the demand.

5. **Attach File** Tool – for inserting large amounts of text or replacement figures.

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X-ray Absorption Spectroscopy Studies of the Room-Temperature Ferromagnetic Fe-Doped 6H-BaTiO₃

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We investigated the effect of annealing temperature on magnetic properties of 2% and 10% Fe-doped BaTiO₃. To understand the possible structural differences between samples treated at different annealing temperatures, and to correlate them with the magnetic properties, several characterization techniques, such as X-ray diffraction and X-ray absorption spectroscopic methods (XANES and EXAFS) were employed. We found that the 2% Fe-doped BaTiO₃ pseudocubic perovskite is paramagnetic regardless of the heat-treatment conditions. Initially paramagnetic 10% Fe-doped 6H-BaTiO₃, treated at 1250°C, became ferromagnetic after additional annealing at higher temperature. We have crystallographically characterized the cation ordering processes in the 6H-BaTiO₃ that occurred during the high-temperature annealing. The ferromagnetism that is induced in this stage is most probably associated with the observed diffusion processes but it extrinsic character still cannot be fully disregarded.

I. Introduction

SIMULTANEOUS presence of semiconducting and ferromagnetic properties makes dilute magnetic oxide (DMO) materials attractive for application in spintronic and magnetoelectronic devices.^{1–3} The room-temperature magnetism in the wide bandgap semiconductors would be of the immense technological importance only if the magnetic coupling is of intrinsic character and not a consequence of paramagnetic dopant segregation or any other extrinsic reason.

Some of the investigated DMO candidates possess room-temperature ferromagnetism when doped with a small percentage of transition metal ions, for example, Mn-doped ZnO,⁴ Fe-doped SnO₂.⁵ Even for these well investigated systems still some disputes over the intrinsic nature of the magnetism exists.^{6,7} Another very promising DMO material is Fe-doped BaTiO₃. Due to the observed room-temperature magnetism it is very intensively investigated material in a wide range of dopant concentration.^{8–20}

However, the reports on the properties of this material are extremely controversial. The first source of this controversy is in the phase transition that occurs with Fe doping and is often not properly considered. It is known that tetragonal perovskite BaTiO₃ can accommodate about 1% of Fe³⁺ on the perovskite B site.^{13,21} Depending on the synthesis condi-

tions from 6% to 13% of Fe is required to fully stabilize the hexagonal 6H-BaTiO₃ structure.^{17,21–23} The intermediate dopant concentrations yield two phase mixture.^{13,21} However, the literature reports claim the intrinsic room-temperature magnetism for the dopant levels from 2% to 10%, although no BaTiO₃ phase in its bulk form can contain such Fe concentration. Of course, with the increase in the nominal Fe concentration toward 10%, the saturation magnetization of the biphasic mixture increases as the amount of the paramagnetic phase reduces. It would also be expected that an increase in the Fe content in Fe-doped 6H-BaTiO₃ phase would enhance the magnetic ordering. Interestingly, it has been shown the opposite: the saturation magnetization of the low end-member with 10% of Fe is the highest while any increase in Fe content causes a gradual decrease in the magnetism until the material becomes paramagnetic at doping level of 70%.^{11,13}

Despite intensive research and a number of reports that claim the room-temperature ferromagnetism in Fe-doped BaTiO₃ the origin of the magnetic coupling still remains unknown.^{10–13,17,21,24} Some authors claim that the room-temperature ferromagnetism is attributed to some structural characteristics, such as ordering of oxygen vacancies⁸ or Fe pairing on Ti(2) sites^{12,25} whereas others attribute the ferromagnetic behavior to the double exchange interaction, due to the simultaneous presence of Fe³⁺ and Fe²⁺ or Fe⁴⁺ valance states.^{15,16,23} Also, there are assumptions that Ti⁴⁺ partially reduces to Ti³⁺, which can contribute to the ferromagnetic coupling.⁸ In our previous study²⁶ on 20% Fe-doped 6H-BaTiO₃ we showed that Fe³⁺ ions undergo an ordering process with a prolonged high-temperature annealing, which plays a crucial role in induction of the magnetic coupling. We showed that the same batch can be paramagnetic or exhibit some ferromagnetic coupling depending on the degree of the cation ordering. We have described the atomistic mechanism of the ordering and identified the driving force of the ordering to be in reduction in electrostatic repulsion on the Ti(2) sites.

In the light of the newly discovered diffusion processes in Fe-doped 6H-BaTiO₃, which are of crucial importance for its functional properties,^{25,26} we also decided to thoroughly investigate the low-concentration range of this system, that is, the range of ≤10%. First, we made a clear distinction between properties of the Fe-doped paramagnetic tetragonal BaTiO₃ and 6H-BaTiO₃ phases. We performed a microscopic structural analysis on the 6H low end-member, 10% Fe-doped 6H-BaTiO₃, with intention to understand the possible structural differences between samples treated at different annealing temperatures. We correlated this to magnetic properties and to the similar diffusion processes in highly doped Fe-doped 6H-BaTiO₃ to fully describe the crystal chemistry of the entire concentration range that is interesting for the DMO applications.

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Here, we have described the ordering processes which are crucial for the establishing of the long-range ferromagnetic coupling in Fe-doped 6H-BaTiO₃. Our convincing experimental results bring us closer to understanding the origin of magnetic coupling in this system.

II. Experimental Procedure

Fe-doped BaTiO₃ samples with nominal composition of BaTi_{1-x}Fe_xO_{3-δ} ($x = 0.02, 0.1$) were synthesized by conventional solid-state reaction method by stoichiometric mixing of raw oxide materials of Fe₂O₃, TiO₂, and BaCO₃. Before each heat treatment, the samples were mixed by grinding in planetary mill at 200 rpm for 1 h in ethanol ambient, dried at 70°C and pressed into 3–4 g pellets. First, the pellets were calcined at 1000°C for 5 h. Then, 2% Fe-doped BaTiO₃ was treated at 1200°C for 5 h, whereas 10% Fe-doped BaTiO₃ was fired twice at 1250°C for 5 h. After this synthesis step, the obtained powders were divided into two parts. One of these parts was additionally annealed at 1400°C for 5 h for 2% Fe-doped BaTiO₃, and at 1500°C for 10 h for 10% Fe-doped BaTiO₃. All heat treatments were carried out in oxygen atmosphere.

The crystal structure of the samples was analyzed by X-Ray diffractometry (PANalytical X'PRO). The XRD patterns were performed in reflection (Bragg–Brentano) mode using CuK_α radiation. Data were collected in the range $15^\circ \leq 2\theta \leq 90^\circ$ with the scan step of 0.008° and with scan step time of 25 s.

The Fe K-edge X-ray absorption spectroscopy (XAS) spectra were collected at the beamline C of HASYLAB at DESY in Hamburg (Germany) simultaneously in fluorescence and transmission detection modes. Ti K-edge XANES spectra were measured at the XAFS beamline of the ELETTRA synchrotron radiation facility in Trieste (Italy) in transmission detection mode. All experimental details are described in our previous study.²⁶ The Fe and Ti XAS spectra were analyzed with the IFEFFIT program package.²⁷ Magnetization was measured as function of applied magnetic field on vibrating sample magnetometer (VSM, LakeShore, 7404) at room temperature.

III. Results and Discussion

The crystal structure of the samples was analyzed by XRD. 2% Fe-doped BaTiO₃ sample treated at 1200°C has

tetragonal crystal structure (space group $P4mm$) with possible traces of hexagonal phase (space group $P63/mmc$) (Fig. 1). The XRD pattern of 2% Fe-doped BaTiO₃ additionally annealed at 1400°C for 5 h indicates the presence of mixture of hexagonal and tetragonal phases. In addition to prevailing 6H phase the sample with 10% of Fe treated at 1250°C contains minor amount of the tetragonal perovskite phase. The sample additionally annealed at 1500°C shows a single-phase composition with only 6H-BaTiO₃ phase. Our results confirm the main claims of other authors^{8,13,21} that the solid solubility limit of Fe in the tetragonal BaTiO₃ is below 2 at.%, whereas about 10 at.% of Fe is required to fully stabilize 6H-BaTiO₃. In addition, it can be concluded that the low-end stability range of Fe-doped 6H-BaTiO₃ is temperature dependent: more than 10% of Fe are needed for stabilization of the structure at 1250°C and below 10% at higher temperature of 1500°C.

First, we have measured the magnetic properties of the tetragonal perovskite phase with the highest possible Fe concentration (i.e., 2%). Regardless the thermal treatment conditions, the paramagnetic behavior has been determined. Magnetization at room temperature as a function of applied magnetic field is shown on Fig. 2.

As the tetragonal perovskite phase is paramagnetic the magnetic properties of the two-phase region between 2% and

Fig. 2. Room-temperature $M(H)$ loops for 2% and 10% Fe-doped BaTiO₃ samples treated at different annealing temperatures.

Fig. 1. (a) X-ray diffraction patterns of BaTi_{1-x}Fe_xO₃ powder samples treated at different annealing temperatures. The peaks of tetragonal (top) and hexagonal (bottom) BaTiO₃ are labeled with Miller indices (hkl). (b) Magnification of diffraction peaks in the 2θ interval 30° – 34° , which shows the effect of annealing temperature on the phase evolution of Fe-doped BaTiO₃.

10% of Fe are dominated by the ferromagnetic properties of the 6H phase; therefore, this region has not been of our main interest. We analyzed the sample with 10% of Fe that was treated at 1250°C, which has predominant 6H-BaTiO₃ phase with a miniscule amount of the paramagnetic tetragonal phase. The sample exhibits pure paramagnetic behavior, so we can conclude that the dominating 6H matrix phase is also paramagnetic. The same sample was additionally annealed at 1500°C and the magnetic measurements showed that this stage of the thermal treatment has induced the ferromagnetic behavior (Fig. 2). Despite the hysteretic behavior the magnetization increases linearly with the increasing magnetic field above 50 kA/m, indicating that ferromagnetic and paramagnetic phases coexist in the sample.

To correlate the observed variation in the magnetic properties with microscopic processes at atomic level, that occur in the 10% Fe-doped 6H-BaTiO₃ during thermal treatment, we performed a detailed analysis of Fe valence state and local structure around Fe cations in the crystal structure by Fe K-edge XANES and EXAFS.

To verify the possible changes in the valence state of the Fe cations in the 10% Fe-doped 6H-BaTiO₃ samples we analyzed Fe K-edge XANES spectra (Fig. 3). A decrease in

the oxidation state of the absorbing atom shifts the energy position of the absorption edge to the lower energies. A shift of about 4.5 eV is observed between Fe³⁺ and Fe²⁺ in reference compounds with well-established Fe valence states (Fe³⁺ in Fe₂O₃ and NdFeO₃, and Fe²⁺ in FeSO₄), in agreement with previous observations.^{28,29} XANES spectra of

Fig. 3. Normalized Fe K-edge XANES spectra measured on the 10% Fe-doped 6H-BaTiO₃ samples after the different heat treatments and reference compounds [Fe(III)₂O₃, NdFe(III)O₃, and Fe(II)SO₄] with known Fe valence state. Two vertical lines are plotted at the position of the Fe(II) K-edge (7121 eV) and Fe(III) K-edge (7125.7 eV) for comparison of the Fe K-edge positions.

Fig. 4. Normalized Ti K-edge XANES spectra measured on 10% Fe-doped 6H-BaTiO₃ powder samples, treated at different annealing temperatures (1250°C and 1500°C), and undoped BaTiO₃ with tetragonal (space group *P4mm*) crystal structure as a standard for Ti⁴⁺.

Fig. 5. (a) The Fourier-transform magnitudes of 10% Fe-doped BaTiO₃ annealed at different temperatures (dashed line – best fit EXAFS model). (b) and (c) Schematic view of the Fe-distribution within three unit cells of the Fe-doped 6H-BaTiO₃ crystal structure (space group *P63/mmc*): (b) after treatment at 1250°C for 5 h, (c) additionally annealed at 1500°C for 10 h. The polyhedra represent octahedrally coordinated Ti atoms. The Fe atoms incorporated onto Ti(1) and Ti(2) sites and their coordination octahedra are marked black. The octahedra with missing faces illustrate the oxygen vacancy in the corresponding face-sharing plane. The big spheres are Ba atoms. The dark and light smaller spheres are oxygen atoms on the O(2) and O(1) crystallographic sites, respectively. The arrow marks the movements of Fe³⁺ from Ti(1) sites to Ti(2) sites after additional annealing at 1500°C for 10 h.

both samples exhibit similar shape and the same energy position of Fe K-edge and preedge resonances. Small differences in the edge profile and intensity of the preedge resonances can be attributed to changes in average local environment of Fe cations, detected in the Fe K-edge EXAFS analysis.²⁸ Comparison of the Fe K-edge energy positions of the samples and reference compounds reveals that all iron atoms in the samples are in trivalent form.

Comparison of Ti K-edge XANES spectra of the different Fe-doped 6H-BaTiO₃ samples shows that spectra are identical (Fig. 4). The Ti K-edge spectrum of the reference undoped tetragonal perovskite BaTiO₃, which is a well-established reference for Ti⁴⁺ valence state, slightly differs in the shape of the edge and preedge features compared with spectra of the two samples, due to the difference in the crystal structures. However, the energy position of the Ti K-edge in the samples coincides well with that in undoped perovskite BaTiO₃, so we can conclude that all titanium in the 6H-BaTiO₃ samples is in tetravalent state.

We used EXAFS analysis to determine the possible changes in local environment of Fe atoms incorporated onto Ti(1) and Ti(2) crystallographic sites. Ti cations in 6H-BaTiO₃ have two different crystallographic sites: Ti(1) and Ti(2) atoms are situated in the corner-sharing and face-sharing octahedra, respectively. On both crystallographic sites Ti cations in 6H-BaTiO₃ are octahedrally coordinated, however, they have small, but significant differences in the local surroundings.²⁶

The EXAFS signal is a superposition of two signals, one from the Fe(1) atoms located at Ti(1) site and the other, arising from Fe(2) atoms located at Ti(2) site. Contributions of individual shells of atoms around Fe atom are visible up to about 5 Å in Fourier-Transform (FT) magnitudes of representative EXAFS spectra. Fourier Transform (FT) of Fe K-edge EXAFS signals consist of contributions from two Fe coordination shells: first shell is the contribution of the oxygen atoms and one titanium atom, second shell is superimposed contribution of the barium and titanium atoms (Fig. 5). Qualitative EXAFS analysis shows that the local neighborhood of Fe in all samples of Fe-doped BaTiO₃ is similar, with small but significant differences in the first and second coordination shells.

The quantitative information on Fe local structure around the Ti(1) and Ti(2), that is, Fe(1) and Fe(2), crystallographic

sites was obtained from quantitative EXAFS analysis. EXAFS spectra were modeled with an ab initio FEFF³⁰ calculation using simultaneous fit of the Fe EXAFS spectra of 10% Fe-doped BaTiO₃ sample sintered at 1250°C for 5 h and 10% Fe-doped BaTiO₃ sample additionally annealed at 1500°C for 10 h. EXAFS spectra have been fitted using similar model, as in our previous EXAFS study of 20% Fe-doped BaTiO₃.²⁶ Two FEFF models with iron atoms incorporated onto Ti(1) and Ti(2) crystallographic sites were constructed, based on the 6H-BaTiO₃ crystal structure with *P*6₃/*mmc* space group (*a* = 5.7240 Å and *c* = 13.9650 Å).³¹ Measured EXAFS spectra were modeled with a linear combination of both FEFF models, where relative amount of each model signal (*X*_{Fe(1)} and *X*_{Fe(2)})—representing the occupancy of Fe on each Ti site was allowed to vary in the EXAFS fit. Models include two neighbor shells, comprising all single-scattering and significant multiple-scattering paths with length of up to 4.52 Å. The structure parameters (type and neighbor atoms, distances and Debye–Waller factors) of nearest neighbor shells around Fe incorporated onto Ti(1) and Ti(2) sites in 10% Fe-doped BaTiO₃ are listed in the Table I. The fit of the both EXAFS spectra was performed with the background corefinesment.²⁷

In FEFF models we allow to vary separately the common shift of the energy origin (ΔE_0), occupation probability of Fe in Ti(2) crystallographic site (*X*_{Fe(2)}) the distances and Debye–Waller factors for the first four single-scattering [Fe(1)–O(1/1), Fe(2)–O(2/1), Fe(2)–O(2/2), and Fe(2)–Ti(2/1)] paths. For the rest of the paths the general linear expansion coefficient was allowed to vary to compute distances, whereas correlated Debye model was used to determine the Debye–Waller factors by varying the Debye temperature parameter. In addition, Fe(2)–Ba(2/1), Fe(2)–Ba(2/3), and Fe(1)–Ba(1/2) distances were allowed to vary separately. The site occupation probabilities of Fe in Ti(1) and Ti(2) sites were constrained to a unit sum. The amplitude reduction factor was kept fixed at 0.75, the value determined in the Fe K-edge EXAFS fit of Li₂Fe_{0.8}Mn_{0.2}SiO₄ in Ref. [29]. The coordination number of neighboring atoms was kept fixed according to the crystallographic data. According to electroneutrality, substituting Ti⁴⁺ ions by Fe³⁺ ions leads to formation of oxygen vacancies, which are known to be situated mainly in the phase-sharing plane of the hexagonal crystal structure of Fe-doped BaTiO₃ (O(2/2) site).^{17,32–35} Thus, to verify the

Table I. Parameters of the Nearest Neighbors Around Fe Atom on Ti(1) and Ti(2) Crystallographic Sites for the 10% Fe-Doped BaTiO₃ Samples Treated at Different Annealing Temperatures (1250°C and 1500°C)[†]

ΔE_0 Fe(2)		1250°C, 5 h			1500°C, 5 h		
		<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)	<i>N</i>	<i>R</i> (Å)	σ^2 (Å ²)
			<i>X</i> _{Fe(2)} = 0.51(4)			<i>X</i> _{Fe(2)} = 0.61(3)	
			0.7 ± 0.5 (eV)			0.3 ± 0.5 (eV)	
O(2/1)		2.9	1.88 (1)	0.007 (3)	2.9	1.89 (1)	0.002 (1)
O(2/2)		2.9	2.02 (1)	0.004 (2)	1.8 (8)	2.02 (4)	0.004 (2)
Ti(2/1)		1	2.7 (1)	0.003 (1)	0.9	2.59 (2)	0.008 (1)
Fe(2/1)		0	—	—	0.1	2.88 (1)	0.008 (2)
Ba(2/1)		2.9	3.40 (1)	0.005 (1)	2.9	3.31 (1)	0.005 (1)
Ba(2/2)		1	3.50 (1)	0.005 (1)	1	3.45 (1)	0.005 (1)
Ba(2/3)		3	3.57 (1)	0.005 (1)	3	3.51 (1)	0.005 (1)
Ti(2/2)		3	3.94 (1)	0.008 (1)	3	3.88 (1)	0.008 (1)
O(2/3)		3	4.08 (1)	0.016 (1)	3	4.02 (1)	0.016 (1)
Fe(1)		<i>X</i> _{Fe(1)} = 0.49 (4)			<i>X</i> _{Fe(1)} = 0.39 (3)		
O(1/1)		5.8	1.99 (2)	0.010 (2)	5.8	2.04 (1)	0.003 (1)
Ba(1/1)		2	3.49 (1)	0.005 (1)	2	3.44 (1)	0.005 (1)
Ba(1/2)		6	3.65 (1)	0.005 (1)	6	3.70 (1)	0.005 (1)
Ti(1/1)		6	3.94 (1)	0.008 (1)	6	3.88 (2)	0.008 (1)
O(1/2)		12	4.52 (1)	0.016 (1)	12	4.45 (1)	0.016 (1)
O(1/3)		12	4.53 (1)	0.016 (1)	12	4.46 (1)	0.016 (1)

[†]The neighbor atom annotations are based on Fe atoms at two crystallographic sites [Ti(1) and Ti(2)] and Fe neighbor atoms ordered by distance in 6H-BaTiO₃ crystal structure: for example, O(2/1) are oxygen neighbors of Fe(2) atoms located at 1.96 Å; O(2/2) are oxygen neighbors of Fe(2) atoms located at 1.99 Å.

N, number of neighbor atoms; *R*, distance from the Fe atom; σ^2 , Debye–Waller factor.

1 aforementioned statement the O(2/2) coordination number
 2 was allowed to vary separately. Also, the coordination num-
 3 ber of the Fe(2/1) atoms was varied in purpose to verify the
 4 formation of Fe pairs in the Ti₂O₃ polyhedra (i.e., Fe₂O₃).
 5 For Fe(2)–Fe(2/1) and Fe(2)–Ti(2/1) paths the same Debye-
 6 Waller factor was used. Fe(2)–Fe(2/1) distance was allowed
 7 to vary separately. Simultaneous fit of the two EXAFS spec-
 8 tra was carried out with intent to stabilize the fit by increas-
 9 ing the number of independent points and decreasing the
 10 number of variables. Thus, to minimize number of variables
 11 some of the parameters in the simultaneous relaxation were
 12 constrained to the common values for the two samples, in
 13 particular, Debye–Waller factor for Fe(2)–O(2/2) path and
 14 Fe(2)–Ba(2/1) distance and Debye temperature parameter.

15 A very good agreement between the model and the experi-
 16 mental spectrum of 10% Fe-doped BaTiO₃ was found in the
 17 R-range from 1.1 to 3.9 Å and in the *k*-interval 4.3–12 Å^{−1}
 18 using *k*²-weight for sample treated at 1250°C and *k*³-weight
 19 for sample additionally annealed at 1500°C.

20 A unit cell of 6H-BaTiO₃ contains six octahedrally coordi-
 21 nated Ti⁴⁺ ions: two at Ti(1) and four at Ti(2) sites. There-
 22 fore, in a case of random distribution 2/3 of Fe ions occupy
 23 Ti(2) sites, whereas other 1/3 occupy Ti(1) crystallographic
 24 sites. In our case, quantitative EXAFS analysis reveals that
 25 after heat treatment at 1250°C about half of all Fe atoms
 26 preferably occupy the Ti(1) crystallographic sites, whereas
 27 the other half of Fe atoms occupy Ti(2) site (Fig. 5). For
 28 sample fired at 1250°C for 5 h the number of oxygen vacan-
 29 cies in O(2/2) crystallographic site was below detection limit,
 30 so the coordination number of O(2/2) atoms was fixed to
 31 2.94, value, which corresponds to the random distribution of
 32 oxygen vacancies around Fe(2) in the sample. No formation
 33 of Fe pairs has been detected (Table I). For the neighboring
 34 atoms we did not find significant distortion, except a slight
 35 relaxation of interatomic distances.

36 After additional annealing at 1500°C about 10% of iron
 37 atoms redistributed from Ti(1) (X_{Fe(1)} = 40%) to Ti(2) (X_{Fe(2)}
 38 = 60%) site. The redistribution is most probably driven
 39 by the same driving force as established in Ref. [26]: reduc-
 40 tion in electrostatic repulsion within Ti₂O₃ face-sharing octa-
 41 hedral. In contrast to 20% Fe-doped BaTiO₃, during the fit
 42 procedure the number of Fe(2)–Fe(2/1) pairs in 10% Fe-
 43 doped BaTiO₃ sample could not be reliably determined by
 44 varying the coordination number of the Fe(2/1) neighbor
 45 atoms. However, it was found that fixing the coordination
 46 number of the Fe(2/1) neighbor atoms to the 0.1 improves
 47 the fit. For this sample ordering of the oxygen vacancies by
 48 varying the coordination number of the O(2/2) atoms was
 49 detected.

50 Our studies show that very subtle differences exist in the
 51 cation ordering processes of different Fe-doped 6H-BaTiO₃
 52 samples. Despite these differences the ordering process is
 53 always associated with induction of the room-temperature
 54 ferromagnetism. Based on our analysis, the ferromagnetism
 55 appears independently from existence of the Fe–Fe pairs or
 56 the oxygen vacancy segregation at O(2/2) site. These two
 57 crystallographic features are most frequently considered as
 58 an origin of the magnetic coupling in this system. Here, we
 59 showed that in the ferromagnetic 10% Fe-doped 6H-BaTiO₃
 60 no significant formation of Fe–Fe pairs exists while previ-
 61 ously²⁶ has already been shown that the ferromagnetism can
 62 also exist without the segregation of oxygen vacancies in O
 63 (2/2) site. However, it has been undoubtedly seen that the
 64 induction of the ferromagnetism is in some way associated
 65 with the high-temperature annealing and most probably with
 66 the observed diffusion processes but it extrinsic character still
 67 cannot be fully disregarded.

IV. Conclusions

71 2% and 10% Fe-doped BaTiO₃ samples were synthesized
 72 by the solid-state reaction method at different annealing

temperatures. The 2%-doped BaTiO₃ samples were found to
 be paramagnetic at RT after both annealing temperatures. The
 10% Fe-doped BaTiO₃ sample treated at 1250°C is paramag-
 netic, whereas 10% Fe-doped BaTiO₃ sample additionally
 annealed at 1500°C exhibited ferromagnetic behavior. Fe and
 Ti K-edge XANES analysis showed no variations in the
 valance state of Fe³⁺ and Ti⁴⁺ cations in all 10% Fe-doped
 BaTiO₃ samples. A Fe K-edge EXAFS analysis of the 10%
 Fe-doped BaTiO₃ sample treated at 1250°C revealed that Fe³⁺
 ions preferably occupied the Ti(1) sites. At this synthesis step
 no preferential distribution of oxygen vacancies has been
 detected. During the additional annealing at 1500°C for 10 h a
 redistribution of the Fe cations from the Ti(1) sites to Ti(2)
 sites was detected, resulting in a random distribution of the Fe
 cations over these sites. Such redistribution was accompanied
 by segregation of oxygen vacancies on the O(2/2) site. The
 induction of the ferromagnetism coincides with the appearance
 of the observed diffusion processes. However, none of the dif-
 fusion processes can be undoubtedly identified as the one that
 induces the ferromagnetic spin coupling.

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