Effect of Co-Doping on Magnetic Properties of Bismuth Ferrite

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The effect of co-doping on structure and magnetic properties of the BiFeO\textsubscript{3}-based multiferroics with a partial isovalent substitution of bismuth for La\textsuperscript{3+}, Gd\textsuperscript{3+}, Dy\textsuperscript{3+}, and Er\textsuperscript{3+} ions have been experimentally investigated by X-ray diffraction and magnetic methods. The ceramic R\textsubscript{1}R\textsubscript{2}0.2−\textsubscript{x}Bi\textsubscript{0.8}Fe\textsubscript{3}O\textsubscript{3} type (x = 0, 0.05, 0.10, 0.15, 0.20; R\textsubscript{1}, R\textsubscript{2} = La, Gd, Dy, Er) samples have been prepared by a solid-state reaction method under cold pressing at high pressure P = 4 GPa. Temperature dependences of magnetization for the co-doped BiFeO\textsubscript{3} demonstrate magnetic “weak ferromagnetic–antiferromagnetic” phase transitions in a high temperature range T = 550–650 K. The presence of a weak ferromagnetism in all compositions is confirmed by open loops of magnetic field dependences. It has been found out that the magnetic characteristics strongly depend on the degree of substitution, temperature, and magnitude of magnetic field.

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1. Introduction

Multiferroics are the most promising materials, in which two or more types of ferroic ordering phenomena such as ferromagnetism, ferroelectricity, and ferroelasticity coexist [1]. The most well-known multiferroics are compounds obtained by partial substitution in BiFeO\textsubscript{3} (BFO) of bismuth cations by isovalent rare-earth cations. In this case, the growth of the resulting magnetic moment and the enhancement of the magnetoelectric interaction is observed. It is interesting to study the mutual influence of various cations of rare-earth elements on the intensity of magnetic interactions and the type of magnetic ordering that forms in BFO with partial isovalent substitution of bismuth cations. For small (x < 0.10) degrees of substitution of Bi\textsuperscript{3+} cations with rare-earth (R) cations, RBFO samples crystallize in the rhombohedral R3\textit{c} lattice, isostructural to BFO lattice [2]. A relatively small number of works are known in which data on the structure and physical properties of materials are obtained with BFO co-doped by cations La–Gd [3], La–Sm [4], La–Ho [5], La–Dy [6], La–Tb [7], and La–Ce [8]. The aim of this work is to study the crystal structure features and magnetic property correlations in multiferroics of R\textsubscript{1}R\textsubscript{2}0.2−\textsubscript{x}Bi\textsubscript{0.8}Fe\textsubscript{3}O\textsubscript{3} type (x = 0, 0.05, 0.10, 0.15, 0.20; R\textsubscript{1}, R\textsubscript{2} = La, Gd, Dy, Er) depending on the concentrations and types of substitutive rare-earth elements cations.

2. Experimental part

The polycrystalline R\textsubscript{1}R\textsubscript{2}(0.2−\textsubscript{x})Bi\textsubscript{0.8}Fe\textsubscript{3}O\textsubscript{3} (x = 0, 0.05, 0.10, 0.15, 0.20; R\textsubscript{1}, R\textsubscript{2} = La, Gd, Dy, Er) samples were synthesized by a solid-state reaction route [9]. The analysis and refinement of X-ray data were performed by JANA2006 software using the Rietveld method [10]. Magnetic-field dependences of a specific magnetization were studied by SQUID magnetometer LOT-Quantum Design MPMS-XL and a vibration magnetometer OXFORD at temperatures of 5, 10, 100, and 300 K in magnetic fields up to H = 8 T. The temperature dependences of the magnetization for the samples placed in vacuum quartz ampoules were measured by the Faraday method [11] within the temperature range of T = 300–1000 K in a magnetic field of H = 0.86 T.

3. Results and discussions

The results of full-profile analysis of X-ray patterns of LGBFO and DEBFO samples according to the Rietveld method are given in Table I and Table II. Based on the data obtained, it can be concluded that co-doping of BFO with Dy\textsuperscript{3+} and Er\textsuperscript{3+} cations leads to stabilization of crystal lattices and to obtaining samples with a predominant \textit{Pnma} phase. The amount of Bi\textsubscript{2}Fe\textsubscript{2}O\textsubscript{5} impurity, which does not exhibit ferromagnetic, ferrimagnetic, and ferroelectric properties at temperatures higher than 250 K [12], is significantly lower during co-doping than what is observed in pure BFO. It is very difficult to obtain the pure phase in co-doped BiFeO\textsubscript{3} due to kinetics of formation, which is attributed to a loss of the volatile
The lattice parameters (Å) and unit cell volume (Å³) for the LaₓGd₁₋ₓBi₀.₈FeO₃ (x = 0, 0.05, 0.10, 0.15, 0.20) samples obtained by the Rietveld refinement of X-ray patterns.

| Composition         | Space group | a [Å]   | b [Å]   | c [Å]   | V [Å³] | GOF        | R_p [%] | R_wp [%] |
|---------------------|-------------|---------|---------|---------|--------|------------|---------|---------|
| Gd₀.₂Bi₀.₈FeO₃     | Pnma        | 5.6200  | 7.7992  | 5.4239  | 237.7  | 1.08       | 11.83   | 17.23   |
| La₀.₀₅Gd₀.₁₅Bi₀.₈FeO₃ | Pnma        | 5.6300  | 7.8004  | 5.4462  | 239.3  | 1.16       | 13.25   | 19.50   |
| La₀.₁₀Gd₀.₁₀Bi₀.₈FeO₃ | Pnma        | 5.5812  | 7.9021  | 5.4541  | 240.5  | 1.13       | 16.16   | 23.40   |
| La₀.₁₅Gd₀.₀₅Bi₀.₈FeO₃ | Pnma        | 5.6058  | 7.8234  | 5.5755  | 244.5  | 0.99       | 10.70   | 16.76   |
| La₀.₂Bi₀.₈FeO₃      | C222        | 5.7666  | 7.9306  | 5.5871  | 255.5  | 1.17       | 13.14   | 17.78   |

The lattice parameters (Å) and unit cell volume (Å³) for the DyₓEr₁₋ₓBi₀.₈FeO₃ (x = 0, 0.05, 0.10, 0.15, 0.20) samples obtained by the Rietveld refinement of X-ray patterns.

| Composition         | Space group | a [Å]   | b [Å]   | c [Å]   | V [Å³] | GOF        | R_p [%] | R_wp [%] |
|---------------------|-------------|---------|---------|---------|--------|------------|---------|---------|
| Er₀.₂Bi₀.₈FeO₃      | Pnma        | 5.7666  | 7.9306  | 5.5871  | 255.5  | 1.15       | 13.14   | 17.78   |
| Dy₀.₀₅Er₀.₁₅Bi₀.₈FeO₃ | Pnma        | 5.5715  | 7.9401  | 5.6126  | 257.2  | 0.99       | 12.05   | 16.12   |
| Dy₀.₁₀Er₀.₁₀Bi₀.₈FeO₃ | Pnma        | 5.5175  | 7.9154  | 5.5908  | 253.0  | 0.96       | 11.89   | 16.84   |
| Dy₀.₁₅Er₀.₀₅Bi₀.₈FeO₃ | Pnma        | 5.4186  | 7.9379  | 5.6026  | 241.0  | 0.96       | 11.67   | 15.39   |
| Dy₀.₂Bi₀.₈FeO₃      | Pnma        | 5.4196  | 7.8977  | 5.6216  | 240.6  | 1.09       | 12.64   | 16.96   |

bismuth during the calcination process. The content of Bi₂FeO₄ impurity in the La₀.₂Bi₀.₈FeO₃ sample is maximum and amounts to 9.2%. Traces of the complex oxide Bi₂FeO₅ were detected, the content of which does not exceed 3% in any of the studied samples. Although the obtained values of the lattice parameter, differ in absolute magnitude from known literature data, the concentration dependences of their values agree well with the results obtained in independent studies [3].

The introduction of La³⁺ cations with increase in substitution degree causes a gradual structural distortion, and for x > 0.15, to a transformation from the orthorhombic to rhomboic C222 lattice. As can be seen in Table I, with an increase in the content of Gd³⁺ cations in the LGBFO system, the values of the volumes of unit cells decrease. This is due to the replacement of larger La³⁺ cations (1.032 Å) by smaller Gd³⁺ (0.938 Å) ions [13]. The criteria of fit La₀.₂Bi₀.₈FeO₃ sample for Pnma, R3c space group were GOF = 1.83, R_p = 18.17%, R_wp = 27.92%, GOF = 1.86, R_p = 19.04, R_wp = 28.45, respectively. In compositions with x < 0.15, the Pnma phase predominates. Accordingly, the decrease in the volume of unit cells is more pronounced in samples with high concentrations of Gd³⁺ ions. The results of the data analysis presented in Table II indicate that the magnitudes of the constant lattices of the DEBFO samples are systematically changed due to the 4f-compression effect, but the structural factors are slightly sensitive to small changes in the composition. As the degree of substitution increases, the diffraction maxima shift to higher angles. However, since the radii of the Dy³⁺ and Er³⁺ cations are close, the values of the parameters of the crystal lattices of the samples vary slightly depending on the substitution degree. This circumstance suggests that changes in the magnetic properties of DEBFO samples will be more determined by the magnetic characteristics of the substituting cations than by changing the parameters of their crystal lattices. The absolute magnitude of the intrinsic magnetic moment of the Dy³⁺ cation (10.7 µ_B) is larger than that of Er³⁺ (9.5 µ_B) ones [14]. Therefore, it is logical to expect the observation of higher values of specific magnetizations with increasing degree of substitution at least at low temperatures. The magnetization curves of the samples exhibit hysteresis characteristic of ferromagnetic materials. Unlike pure BiFeO₃, which shows a linear dependence of the specific magnetization on the magnitude of the magnetic field typical of antiferromagnets, the magnetic field dependences of cation-substituted samples indicate the presence of a weak ferromagnetic moment. In the case of substitutions of Bi³⁺ ions, quantitative changes in the values of specific magnetizations and values of coercive fields recovered from particular magnetic hysteresis loops are observed, as can be seen in Fig. 1a–d.

Such behavior is revealed by the concentration dependences of the magnetic characteristics of the samples of the DyₓEr₁₋ₓBi₀.₈FeO₃ system, except for the values of the specific residual magnetizations of the composition Er₀.₂Bi₀.₈FeO₃, which takes values less than those in compositions with 0.05 < x < 0.15 (Fig. 2). An increase in the absolute values of the coercive forces and residual specific magnetizations at all substitutions with respect to the initial matrix (BFO) is revealed. A change in the R–O, Fe–O bond lengths and, as a result, the values of the Fe–O-Fe bond lengths, leads to a change in the intensities of indirect exchange interionic interactions. With an increase in the content of Er³⁺ cations in the DyₓEr₁₋ₓBi₀.₈FeO₃ system, the average effective radius...
of the rare-earth cation in the A sublattice decreases. This leads to a change in the distances between the ions of iron and oxygen and, as a consequence, to a change in the magnitudes of the “Fe–O–Fe” angles. To study the wFM-FM transition, respectively, the \( M(T) \) dependence was calculated. Within the framework of the phenomenological model proposed by Hamad [15], the temperature dependence of magnetization \( (M) \) in the vicinity of the Néel point \( (T_N) \) can be written as

\[
M = \frac{1}{2} (M_i - M_f) \tanh [A (T_N - T)] + BT + C, \tag{1}
\]

where \( M_i \) and \( M_f \) are initial and final points of temperature range of wFM-AFM transition. The other coefficients are defined as

\[
A = \frac{2B - S_N}{M_i - M_f}, \quad B = \left( \frac{dM}{dT} \right)_{T_i},
\]

\[
S_N = \left( \frac{dM}{dT} \right)_{T_N}, \quad C = \frac{1}{2} (M_i + M_f) - BT_N.
\]

To a first approximation, the value of the Néel point was determined by at least the first derivative \( dM/dT \). Then its value was changed to obtain the best agreement between the model and experimental data. The “Fe–O–Fe” angles decrease with an increase in the replacement cations concentration. This leads to a decrease in the \( T_N \) values, the non-monotonic behavior of which in the \( x > 0.10 \) region is due to magnetization frustration caused by the coexistence of various phases. However, in general, there is an almost linear decrease in the Néel point with an increase in the degree of substitution.

4. Conclusions

X-ray diffraction and magnetic methods have been used for investigating the structure and magnetic properties of the ceramic \( R_1_xR_2_{0.2-x}Bi_{0.8}FeO_3 \) (\( R_1, R_2=La, \ Gd, \ Dy, \ Er \)) samples with a concentration of \( x = 0.2 \) prepared by a solid-state reaction method under cold pressing at high pressure \( P = 4 \) GPa. According to magnetic data, the temperature dependences of magnetization for the \( R_1_xR_2_{0.2-x}Bi_{0.8}FeO_3 \) multiferroics show magnetic “weak ferromagnetic–antiferromagnetic”
at $T_N = 550$–$650$ K phase transitions. The appearance of a weak ferromagnetism is caused by suppression of the spatial spin-modulated structure in course of an isovalent substitution of Bi$^{3+}$ with R$^{3+}$ cations in the $R_1R_2_{0.2-x}Bi_{0.8}FeO_3$. It has been established that the magnetic characteristics in the vicinity of the $T_N$ point strongly depend on the concentration, temperature, and magnetic field. The results of the investigations of the magnetic properties for the $R_1R_2_{0.2-x}Bi_{0.8}FeO_3$ samples indicate the possibility of its practical application at temperatures above room temperature. The obtained data can be used to study the features of phase magnetic transitions in similar materials.

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