Magnetoelectric coupling and phase transition in BiFeO$_3$ and (BiFeO$_3$)$_{0.95}$(BaTiO$_3$)$_{0.05}$ ceramics


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In situ high-resolution synchrotron x-ray diffraction reveals a local minimum in rhombohedral distortion angle $\gamma_R$ (associated with an inflection in the lattice constant $a_R$ near 400 and 350 °C in BiFeO₃ (BFO) and (BiFeO₃)₀.95(BaTiO₃)₀.05 (BFO–5%BT), respectively. It suggests a coupling between ferroelectric and magnetic parameters near the antiferromagnetic–paramagnetic transition, which is responsible for the broad frequency-dependent dielectric maxima. A rhombohedral (R)–orthorhombic (O)–cubic (C) transition sequence takes place near 820 and 850 °C in BFO upon heating. BFO–5%BT exhibits a R–C transition near 830 °C. The BaTiO₃ substitution can enhance dielectric and ferromagnetic responses and reduce electric leakage. The dielectric loss of BFO–5%BT remains less than 0.04 below 150 °C. © 2011 American Institute of Physics. [doi:10.1063/1.3551578]

I. INTRODUCTION

The multiferroic bismuth ferrite BiFeO₃ (BFO) materials have attracted much attention not only because they possess ferroelectric (FE) and ferromagnetic (FM) properties, but also because the coupling between the electric and magnetic ordering leads to additional functionalities. BFO can provide an alternate choice as a “green” FE/FM material because the coupling between the electric and magnetic properties, which is responsible for the broad frequency-dependent dielectric maxima. A rhombohedral (R)–orthorhombic (O)–cubic (C) transition sequence takes place near 820 and 850 °C in BFO upon heating. BFO–5%BT exhibits a R–C transition near 830 °C. The BaTiO₃ substitution can enhance dielectric and ferromagnetic responses and reduce electric leakage. The dielectric loss of BFO–5%BT remains less than 0.04 below 150 °C. © 2011 American Institute of Physics.

II. EXPERIMENTAL PROCEDURES

The BFO ceramic was prepared by the solid state reaction method. The dried starting powders of Bi₂O₃ and Fe₂O₃ (purity ≥ 99.0%) were weighed in a 1.1:1 ratio to compensate the low melting point of Bi, and then mixed in an agate mortar for more than 24 h using alcohol as a medium. The mixture was dried and mixed with polyvinyl acetate as a binder for granulation. The ground mixture was pressed into a 1.0-cm-diameter disk. The pressed BFO disk was sintered in the region of 850–870 °C for 1–3 h. The optimal sintering condition for BFO ceramic is 860 °C for 2 h and the density is ~90% of the theoretical density. For synthesis of BFO–5%BT ceramic, the dried starting powders of BFO and BT (purity ≥ 99.0%) were weighed in a 0.95:0.05 molar ratio...
and then a similar process was followed as for the BFO ceramic. The optimal sintering condition for BFO–5%BT ceramic is 960 °C for 3 h and the density is ~93% of the theoretical density.

In situ high-resolution synchrotron XRD was performed at the National Synchrotron Radiation Research Center (in Taiwan) with a photon energy of 8.0 keV (λ = 1.550 Å). A Wayne–Kerr Analyzer PMA3260A was used to obtain the real (ε′) and imaginary (ε″) parts of dielectric permittivity. Hysteresis loops of polarization versus E field were taken by using a Sawyer-Tower circuit at f = 46 Hz.

III. RESULTS AND DISCUSSION

Figure 1 shows XRD spectra of BFO and BFO–5%BT ceramics at room temperature. A two-peak splitting occurs in the (110), (111), (210), and (220) reflections, suggesting a rhombohedral structure. Some minor second phases (impurity) of possible Bi12FeO39 (Ref. 21) and Bi2Fe2O7 (Ref. 33) were observed as indicated by the symbols “*” and “#,” respectively. BFO–5%BT has less impurity phases. The XRD peaks of BFO–5%BT occur at lower 2θ than those in BFO, mainly due to the larger atomic radius of the Ba2+ ion (1.35 Å) compared with the Bi3+ ion (1.03 Å) on the perovskite A site. The Fe3+ ion (0.60 Å) and Ti4+ ion (0.65 Å) have similar atomic radii on the perovskite B site.

Figure 2 shows frequency-dependent dielectric permittivity (ε′) and dielectric loss (tan δ = ε″/ε′) upon heating. The temperature (Tm) corresponding to the maximum in ε′ exhibits a broad frequency dispersion. For BFO, Tm shifts from ~420 °C at 50 kHz to ~490 °C at 1 MHz. BFO–5%BT does not exhibit obvious frequency dispersion in ε′ below 200 °C and its Tm shifts to lower temperatures. This dielectric response is likely activated by the antiferromagnetic–paramagnetic transition, which takes place at the Néel temperature (TN). A similar but rather pronounced frequency dispersion in dielectric maxima was observed in BFO–10%BT ceramic, associated with a local minimum in rhombohedral distortion angle xR near TN. It was attributed to the changes in relative positions of Bi3+ and Fe3+ ions in the perovskite structure as temperature approaches TN. In addition, the neutron scattering result of BFO also revealed changes of distortion and strain in oxygen octahedra (FeO6) near TN caused by the magnetoelectric and/or magnetoelastic couplings, which can change electric polarization.

The dielectric loss (tan δ) of BFO and BFO–5%BT are respectively about 0.2–0.4 and 0.02–0.04 at room temperature, indicating that 5 mol% BaTiO3 substitution can efficiently reduce electric conductivity. The tan δ of BFO–5%BT remains less than 0.04 below 150 °C, which is close to the tan δ ≡ 0.02 of the soft PZT-5 ceramic. The dielectric losses of BFO and BFO–5%BT exhibit an exponential upturn above ~300 °C with magnitude proportional to 1/f, indicating conductivity activated by thermal energy.

Figure 3 shows temperature-dependent (110) synchrotron XRD spectra of BFO and BFO–5%BT upon heating. The insets are the XRD spectra near structural transitions. The (110) XRD spectrum of BFO–5%BT is much broader than in BFO, likely caused by random distributions of Bi and Ba ions on the A site, and Fe and Ti ions on the B site. The random displacements can cause different 2θ reflections and result in a broadening effect in XRD. Upon heating, two (110) R-phase reflections of BFO exhibit a triple splitting near 820 °C and then merge into a single peak near 850 °C, indicating a transition sequence of rhombohedral (R)–tetragonal (T)–orthorhombic (O)
orthorhombic (O)–cubic (C) upon heating. A R–O–C phase sequence was also revealed in bulk BFO near 820 and 925°C from Raman and thermal studies. Our result is consistent with the dielectric result of BFO–PZT solid solutions, which predicted a ferroelectric Curie temperature near 850°C for BFO. Additional XRD peaks appear above 860°C in BFO, implying possible decompositions of Fe₂O₃ and Bi₂Fe₄O₉ (Ref. 33) due to the loss of Bi in high-temperature region.

For BFO–5%BT, two (110) R-phase reflections merge into a broad single peak near 830°C, indicating a rhombohedral (R)–cubic (C) transition upon heating. This result may imply that the orthorhombic phase is not favored by replacing BaTiO₃ in the perovskite structure. Decomposition does not occur in BFO–5%BT below 900°C, implying that 5%BT substitution can stabilize the perovskite structure.

Figure 4 shows temperature-dependent lattice parameters calculated from the (110) XRD peaks. A local minimum in rhombohedral distortion angle $\alpha_R$ (associated with a slight inflection in lattice constant $a_R$) occurs near 400 and 350°C in BFO and BFO–5%BT, respectively. A similar minimum of $a_R$ was reported near $T_N$ in BFO and BFO–10%BT. This local minimum in $a_R$ suggests that the position shift (or distortion) of Bi³⁺ cation gradually reaches a maximum as temperature approaches $T_N$. This confirms a coupling between ferroelectric and magnetic order parameters near $T_N$, which causes the broad frequency-dependent dielectric maxima as shown in Fig. 2.

Electric hysteresis loops at room temperature are given in Fig. 5 and show attainable remanent polarizations of $\sim$0.01 and $\sim$0.1 μC/cm² in BFO and BFO–5%BT, respectively. However, the saturated polarization cannot be achieved in either case. A similar hysteresis loop with an unsaturated remanent polarization of $\sim$0.001–0.015 μC/cm² was observed from the BFO ceramic. As evidenced in Fig. 5, BFO–5%BT can sustain a higher measuring field than BFO, confirming that 5%BT substitution can reduce electric leakage and thus enhance polarization.

Figure 6 shows the magnetic hysteresis loops (magnetization versus magnetic field) of BFO and BFO–5%BT at...
room temperature. The magnetization curve of BFO is linear with the field, which is typical for an antiferromagnetic arrangement of the Fe$^{3+}$ magnetic moments. BFO–5%BT exhibits a rather weak ferromagnetic nature, which is similar to the magnetization curve observed in the BFO–10%BT ceramic. These results suggest that BT substitution can enhance the FM feature.

IV. CONCLUSIONS

A local minimum in rhombohedral distortion angle $z_R$ (associated with an inflection in the lattice constant $a_D$) was revealed near 400 and 350 $^\circ$C in BFO and BFO–5%BT, respectively. The local minimum in $z_R$ indicates that the position of Bi$^{3+}$ reaches the largest distortion near $T_N$. This suggests a magnetoelectric coupling near $T_N$, which is responsible for the broad frequency dispersion in dielectric maxima. This work suggests that the Néel temperatures of BFO and BFO–5%BT at room temperature. The magnetization curve of BFO is linear with the field, which is typical for an antiferromagnetic arrangement of the Fe$^{3+}$ magnetic moments. BFO–5%BT exhibits a rather weak ferromagnetic nature, which is similar to the magnetization curve observed in the BFO–10%BT ceramic. These results suggest that BT substitution can enhance the FM feature.

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