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# Composition-induced phase evolution and high strain response in Ba(Zn<sub>1/3</sub>Nb<sub>2/3</sub>)O<sub>3</sub>-modified (Bi<sub>0.5</sub>Na<sub>0.5</sub>)TiO<sub>3</sub>-based lead-free ferroelectrics

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In this work, a paraelectric complex perovskite Ba(Zn<sub>1/3</sub>Nb<sub>2/3</sub>)O<sub>3</sub> (BZN) was introduced into the morphotropic boundary composition (Bi<sub>0.5</sub>Na<sub>0.5</sub>)<sub>0.93</sub>Ba<sub>0.07</sub>TiO<sub>3</sub> (BNBT) to modulate the phase structure and electrical properties as well as the field-induced strain behavior. Using a columbite route, the ceramics with pure perovskite structure were successfully fabricated. The structure and electrical measurements showed that the introduction of BZN into BNBT results in the decrease of the rhombohedrality 90-γ, and promotes the evolution from ferroelectric to antiferroelectric (AFE) relaxor. Besides, the introduction of BZN induces the volume increase and dimension reduction of the nanosized AFE relaxor domains. A large field-induced strain of 0.39% with good stability against frequency, field, and temperature was obtained at the BNZ addition of composition  $x = 0.01$ , which locates at the critical composition boundary between ferroelectric state and AFE relaxor state.

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## I. Introduction

Lead-based piezoelectric materials cause serious environmental concerns due to the toxicity of lead, even though they have been widely used in several industries.<sup>1</sup> Tremendous effort has been devoted to developing competitive lead-free piezoelectric ceramics.<sup>2</sup> Among all the lead-free materials developed, Bi<sub>0.5</sub>-Na<sub>0.5</sub>TiO<sub>3</sub> (BNT) is considered to be one of the most promising candidates due to its high electric field-induced strain level.<sup>3,4</sup> Further, to enhance the field-induced strain behavior of BNT-based materials is not only scientifically interesting but also technologically important for practical application. Especially, the BNT-based compositions located in morphotropic phase boundary (MPB) attract extensive interest of researchers due to their relatively large strains originating from the nanotwins as an intermediate bridging phase between tetragonal *P4mm* and rhombohedral *R3c* phases.<sup>5,6</sup> By incorporating strong ferroelectric compounds as modifier, the ferroelectric order of the BNT-based MPB compositions could be disrupted,<sup>7-9</sup> such as K<sub>0.5</sub>Na<sub>0.5</sub>NbO<sub>3</sub>,<sup>10</sup> LiNbO<sub>3</sub>,<sup>11</sup> thus obtaining a large and hysteresis-free strain.

Ba(Zn<sub>1/3</sub>Nb<sub>2/3</sub>)O<sub>3</sub> (BZN) is paraelectric perovskite with a cubic structure.<sup>12</sup> BZN could form a stable solid solution with BNT due to its adequate tolerance factor close to 1 (1.0285). Moreover, the polarization of a ferroelectric may be enhanced by

weak ferroelectric species,<sup>13</sup> which is beneficial to obtain a large field-induced strain. Besides, the complex B-site cations could effectively induce the local inhomogeneity that destroy the ferroelectric order and induce the polar nanoregions (PNRs). So far, the incorporation of BZN into BNT-based solid solutions could disrupt the stability of ferroelectric domains and make the AFE relaxor percolating into low temperatures, which will construct the another base of a large field-induced strain.

In this work, (1- $x$ )(Bi<sub>0.5</sub>Na<sub>0.5</sub>)<sub>0.93</sub>Ba<sub>0.07</sub>TiO<sub>3</sub>- $x$ Ba(Zn<sub>1/3</sub>Nb<sub>2/3</sub>)O<sub>3</sub> ceramics was designed and fabricated by a columbite route suggested by Swartz and Shrout.<sup>14</sup> Crystal structure, dielectric and ferroelectric response as well as field-induced strain behavior were investigated in details. The experiments demonstrated that a large strain of 0.39% was obtained at composition  $x = 0.01$ , which provides the potential in application of environmental friendly solid-state actuators.

## II. Experimentals

Samples based on (1- $x$ )(Bi<sub>0.5</sub>Na<sub>0.5</sub>)<sub>0.93</sub>Ba<sub>0.07</sub>TiO<sub>3</sub> (BNBT)- $x$ Ba(Zn<sub>1/3</sub>Nb<sub>2/3</sub>)O<sub>3</sub> (BZN) for  $0 \leq x \leq 0.02$  were fabricated through the columbite route using Bi<sub>2</sub>O<sub>3</sub> (purity  $\geq 99.0\%$ ), Ba<sub>2</sub>CO<sub>3</sub> (purity  $\geq 99.0\%$ ), Na<sub>2</sub>CO<sub>3</sub> (purity  $\geq 99.8\%$ ), TiO<sub>2</sub> (purity  $\geq 99.0\%$ ), ZnO (purity  $\geq 99.0\%$ ), and Nb<sub>2</sub>O<sub>5</sub> (purity  $\geq 99.0\%$ ) as raw materials. First, the columbite ZnNb<sub>2</sub>O<sub>6</sub> was prepared by the solid synthesis method: the predetermined amounts of ZnO and Nb<sub>2</sub>O<sub>5</sub> were mixed in ethanol (purity  $\geq 99.7\%$ ) for 12 h using ZrO<sub>2</sub> balls, and then calcined at 1000 °C for 4 h to obtain ZnNb<sub>2</sub>O<sub>6</sub>. Then, the precursor ZnNb<sub>2</sub>O<sub>6</sub> was mixed with predetermined amounts of Bi<sub>2</sub>O<sub>3</sub>, Ba<sub>2</sub>CO<sub>3</sub>, TiO<sub>2</sub>, and Na<sub>2</sub>CO<sub>3</sub>, and

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calcined at 850 °C for 4 h. The calcined BNBZN100x powders were subsequently ball-milled in ethanol again for 12 h, and pressed into pellets of 11.5 mm diameter under 300 MPa using polyvinyl butyral as binder. After removing the binder at 560 °C for 12 h, the green pellets were sintered at 1150 °C for 4 h in a covered alumina crucible. The ceramic samples were ground and polished to diameter of 9.5 mm and thickness of 0.8 mm.

The phase structure was identified by powder X-ray diffraction (XRD, Bruker D8 Advance, Karlsruhe, Germany) in the  $\theta$ - $2\theta$  configuration at room temperature, using Cu K $\alpha$  (1.54178 Å) radiation (scan range = 20.00–80.00°, scan speed = 2.00° min<sup>-1</sup>, sampling pitch = 0.02°). Before the XRD measurement, the samples were poled under 5 kV mm<sup>-1</sup> in silicone oil bath at 65 °C for 30 min and field-cooled to room temperature, and then crushed to powder. To determine the cell parameters, the cell refinement was performed on the XRD data using JADE 6.5 software (Materials Data, Livermore, CA). To observe the atomic arrangement and local symmetry, a high-resolution transmission electron microscope (HR TEM, JOEL JEM ARM300F) were used. Besides, the micromorphology was observed on the thermally etched fracture surface using a scanning electron microscopy (SEM, Hitachi S4800, Tokyo, Japan). To measure the electric properties, the silver paste was coated on both surfaces of the ceramic samples and fired at 700 °C for 20 min to form electrodes. The dielectric permittivity of poled or unpoled samples was measured with a LCR meter (E4980, Agilent, Santa Clara, CA) equipped with an automated temperature controller at heating ramp rates of 3 °C min<sup>-1</sup>. Ferroelectric polarization ( $P$ )-field ( $E$ ) loops and field-induced strain were measured in silicone oil using a ferroelectric tester (Premier II, Radiant Technologies, Albuquerque, NM) equipped with a photonic sensor (2000, MTI, Albany, NY) at 1 Hz. Electrical impedance spectrum was measured by an impedance analyzer (Concept 400, Novocontrol Technologies, Montabaur, Germany). The local structure was characterized by a Raman scattering spectrometer (HR800, Horiba Jobin-Yvon, France) using the 532.08 nm Ar-ion laser as the excitation source.

### III. Results and discussion

Fig. 1 shows the SEM images of compositions  $x = 0$ –0.02. The grains show a polyhedral morphology, and no apparent pores were observed in grain internals or boundaries, which indicate the good sintering ability of the samples. Moreover, the average grain size varies in 0.92–0.99  $\mu\text{m}$  estimated by a freeware SmileView, which is almost independent from BZN content, implying that BZN addition exact a weak impact on the grain growth.

Fig. 2 shows the powder XRD patterns of the unpoled and poled BNBZN100x ceramics. As seen from Fig. 2a, all the unpoled samples present a pseudocubic perovskite structure, and no apparent second phase is observed, which implies that BZN completely diffuse into the lattice of BNBT and form a stable solid solution. To give an insight into the phase evolution induce by BZN addition, the fine scanning XRD patterns of sample before and after poling were recorded in the  $2\theta$  range of 39–41° and 67–68.5°, as shown in Fig. 2b and c. For

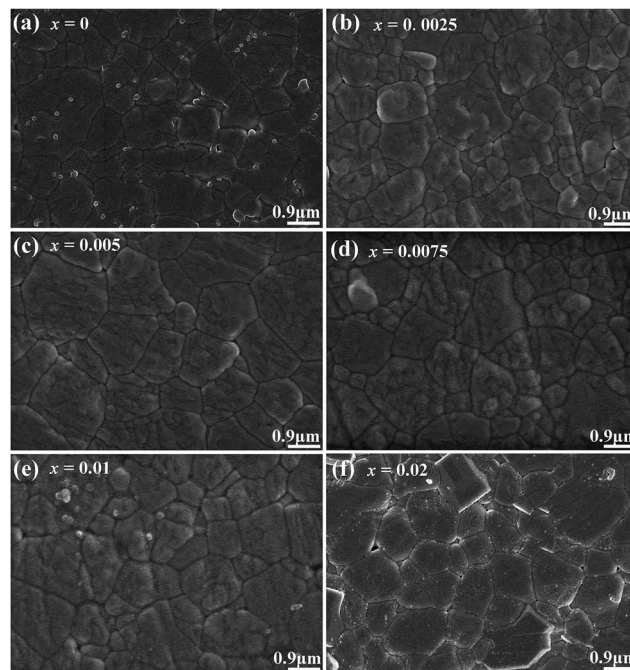


Fig. 1 SEM images of compositions  $x = 0$ –0.02.

the poled compositions  $x \leq 0.01$ , the (111)<sub>PC</sub> diffraction around 40° splits, and a shoulder assigned to the (002) diffraction of rhombohedral  $R3c$  occurs, verifying their structural nature of rhombohedral  $R3c$  symmetry. Meanwhile, the (220)<sub>PC</sub> diffraction around 68° also splits after poling, further indicating the asymmetric structure. However, when  $x > 0.01$ , the (111)<sub>PC</sub> and (220)<sub>PC</sub> diffractions keep unchanged after the poling, which means that the compositions  $x > 0.01$  possesses the pseudocubic structure, and should be related to the percolation of AFE relaxor phase with  $P4bm$  structure.<sup>15</sup> Moreover, to elucidate the effect of BZN incorporation on the crystal lattice, Fig. 2d depicts the lattice constant  $a_{pc}$  and rhombohedrality  $90-\gamma$  fixed by a freeware JADE 6.5 based on the XRD data. As seen from Fig. 2d, the incorporation of BZN into BNBT brings in the lattice expansion, which could be ascribed to the larger effective ionic radii of Zn<sup>2+</sup> (0.74 Å) and Nb<sup>5+</sup> (0.69 Å) than that of Ti<sup>4+</sup> (0.605 Å). Especially, as BZN added, the  $90-\gamma$  value decreases gradually to zero when  $x$  arrives to 0.075, affirming the composition induced phase evolution from rhombohedral ferroelectric ( $R3c$ ) and pseudocubic AFE relaxor ( $P4bm$ ), which may be the result of the disruption of the long-range ferroelectric order.

To verify the percolation of AFE relaxor phase with  $P4bm$  structure, the HR TEM observation was carried out on the composition  $x = 0.01$ . It can be seen from Fig. 3 that the atomic arrangement is quite orderly along the [001]<sub>c</sub> zone axis, which indicates its perfect perovskite structure. As given in the inset, the selected-area diffraction pattern (SADP) shows a  $P4bm$  symmetry as revealed by the presence of  $1/2\{00e\}$  superlattice diffraction spots by fast fourier transformation software analysis.

To verify the effect of BZN addition on the long-range ferroelectric order, the electric impedance spectroscopy was



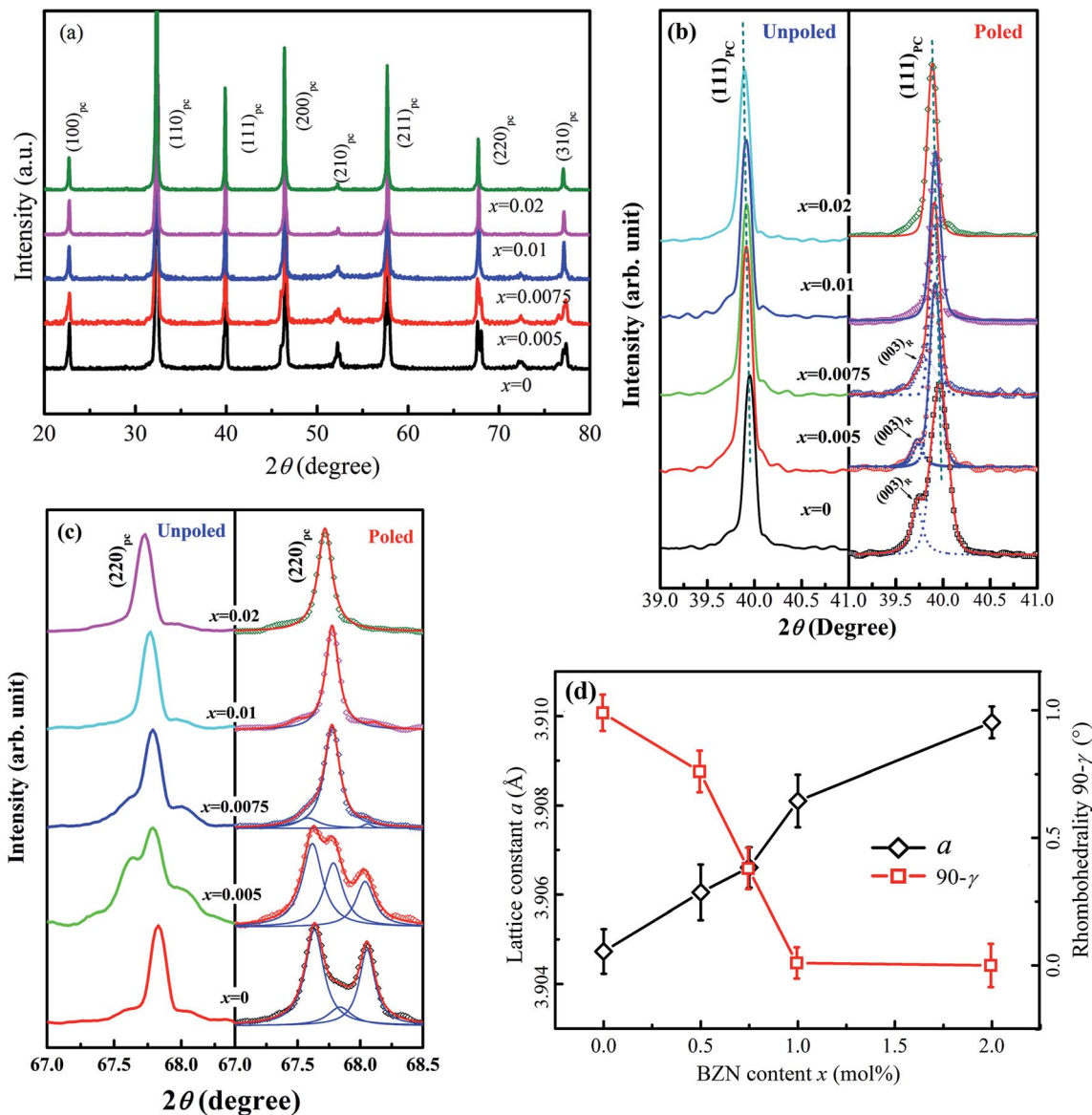


Fig. 2 Powder XRD patterns (a), fine-scanning XRD patterns around  $2\theta = 40^\circ$  (b) and  $68^\circ$  (c) as well as the cell refinement result (d) of compositions  $x = 0-0.02$ .

carried out. It is reported that the relaxor phase with  $P4bm$  symmetry demonstrates a nanosize domain configuration.<sup>16</sup> And the nanosized AFE relaxor domains will bring in a shoulder response in the modulus  $M''(f)$  spectra at high temperature.<sup>17</sup> Fig. 4 gives the  $M''(f)$  spectra of the compositions  $x = 0, 0.005$ , and  $0.01$  at  $300^\circ\text{C}$ . Obviously, the shoulder response enhances gradually with the BZN addition, affirming that the addition of BZN brings in the volume growth of the AFE relaxor domains. Especially, as the BZN added, the frequency  $f_{\text{max}}$  corresponding to maximum  $M''$  of the shoulder response shifts to high frequency, which infers the dimension reduction of the nanosized AFE relaxor domains.<sup>18</sup> In a word, the BZN addition not only increases the volume but also reduces the dimension of the nanosized AFE relaxor domains.

Fig. 5 shows the temperature dependence of dielectric permittivity ( $\epsilon_r$ ) of unpoled and poled compositions ( $x = 0-$

$0.02$ ). As seen from Fig. 5, two anomalies are observed for the composition  $x = 0$ . A maximum of dielectric permittivity with significant frequency dispersion occurs at about  $280^\circ\text{C}$  ( $T_{\text{max}}$ ), which is the relaxation of AFE relaxor phase with  $P4bm$  structure without any measurable structural transition.<sup>19</sup> Besides, a frequency-independent divergence of the  $\epsilon_r$  before and after poling is observed at  $90^\circ\text{C}$  for the composition  $x = 0$ , which corresponds to the transition from ferroelectric to AFE relaxor<sup>20,21</sup> and defined as the temperature  $T_{\text{F-R}}$ . As BZN added, the  $T_{\text{F-R}}$  lowers to about  $45^\circ\text{C}$  for composition  $x = 0.005$ , indicating the composition-induced ferroelectric to relaxor phase transition.<sup>21</sup> Besides, the lowering of the  $T_{\text{F-R}}$  is also related to the decrease of the rhombohedrality  $90-\gamma$ <sup>22,23</sup> and reveals that the BZN addition weakened the stability of ferroelectric domains. However, for the compositions  $x > 0.005$ , the divergence could not be distinguished in the dielectric responses,



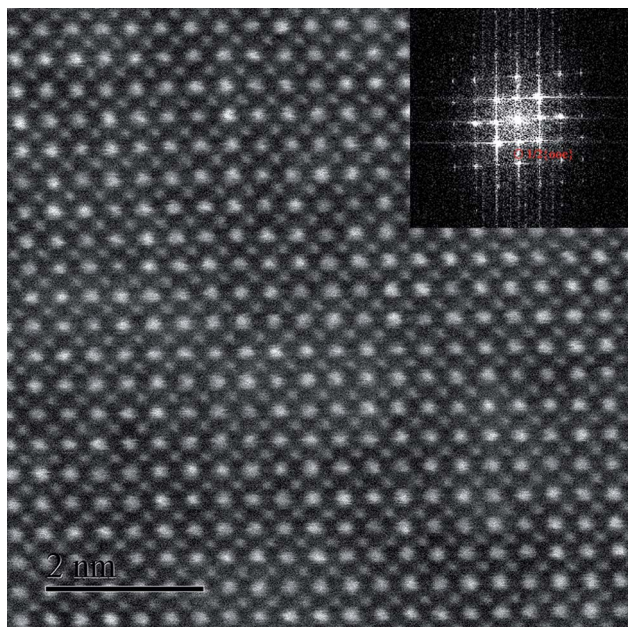


Fig. 3 HR TEM image and the selected-area diffraction pattern of composition  $x = 0.01$ .

which is associated with the fact that the relaxation process of the remanent ferroelectric phase is almost swamped in that of AFE relaxor.<sup>24</sup>

Raman scattering spectroscopy could yield important information about the local phase structure. Fig. 6a exhibits the temperature-variable Raman scattering spectra of the composition  $x = 0.01$  from 0 °C to 100 °C and the deconvolutions by LabSpec5 software using Gaussian shape lines. It is reported

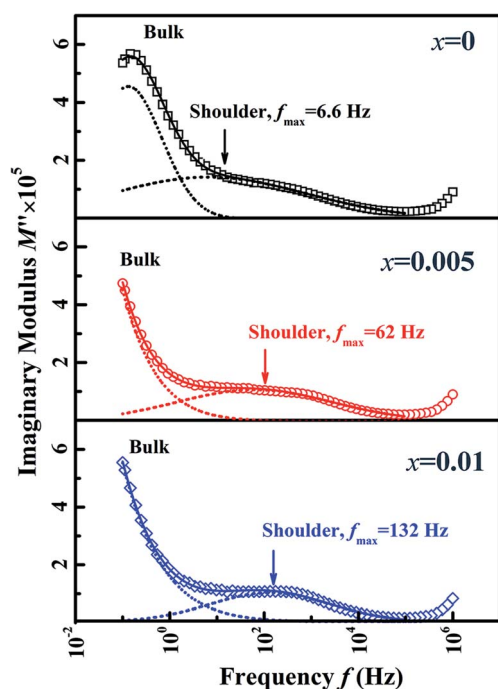


Fig. 4  $M''(f)$  spectra of compositions  $x = 0, 0.005$ , and  $0.01$  at 300 °C.

that the Raman spectra of BNBZT could be divided into four regions A, B, C and D.<sup>25</sup> Among them, modes  $C'$  and  $C''$  in the range of 560–620  $\text{cm}^{-1}$  present double splitting peaks of the oxygen octahedra. Fig. 6b shows the temperature dependence of the Raman shift of the modes  $C'$  and  $C''$  as well as their intensity ratio  $I_{C'}/I_{C''}$ . It can be seen that the modes  $C'$  and  $C''$  shift to low wavenumbers, which could be ascribed to the enlargement of average B–O bond length induced by the lattice expansion. The softening of modes C will result in the weakening of the coupling between  $\text{BO}_6$  and induce the phase transition.<sup>26–28</sup> Meanwhile, a sudden turning point is observed at 50 °C in the  $I_{C'}/I_{C''}$  curve, a sign of variation of cation ordering since the B–O bonds are influenced by the dynamics of the nanodomain phases.<sup>25</sup> As mentioned above (Fig. 5), for composition  $x = 0.01$ , no obvious dielectric divergence could be observed that the  $T_{\text{F-R}}$  could not be distinguished. The inflection at 50 °C appeared in Raman spectra (Fig. 6b) approves the existence of the ferroelectric to AFE relaxor transition for the composition  $x = 0.01$ , implying that the remanent ferroelectric domains exist, similar to the composition with low BZN addition.

Fig. 7 gives the ferroelectric  $P$ - $E$  loops of BNBZN100x samples. As seen from Fig. 7, the saturated  $P$ - $E$  loops were observed for compositions  $x < 0.005$ , a typical sign of a ferroelectric. As  $x$  increased to 0.005, the  $P$ - $E$  loops began to be pinched, which is the result that the AFE relaxor phase with tetragonal  $P4bm$  structure percolates down to room temperature.<sup>15,29</sup> Further increase of BZN addition induces the  $P$ - $E$  loop pinched much severely, inferring the contribution of nanosized AFE relaxor phase on the ferroelectric response. Generally, the nanosized AFE relaxor phase can reversibly evolve into a long-range ferroelectric order under an electric field, and return to original 'nonpolar' state at zero field.<sup>30</sup> Considering that the remanent polarization  $P_r$  is the contribution from ferroelectric phase while the spontaneous polarization  $P_s$  is the glossary contribution originating both ferroelectric and nanosized AFE relaxor domains, the ratio  $(P_s - P_r)/P_s$  could be regarded as the indication of the polarization contribution originating from the

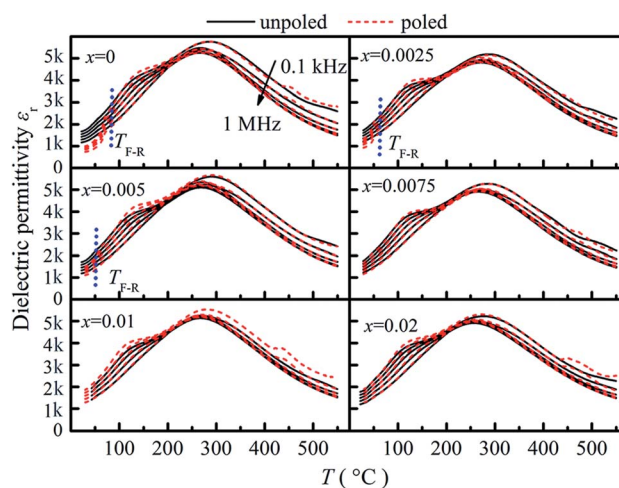


Fig. 5 Temperature dependent dielectric permittivity of compositions  $x = 0–0.02$  before and after poling.



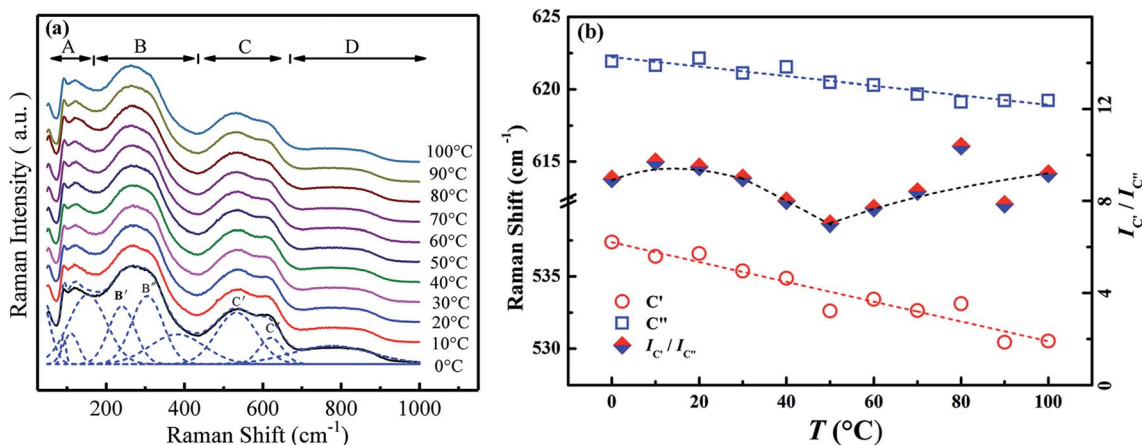


Fig. 6 Temperature-dependent Raman scattering spectra (a) and the deconvolutions results (b) of composition  $x = 0.01$  from 0 °C to 100 °C.

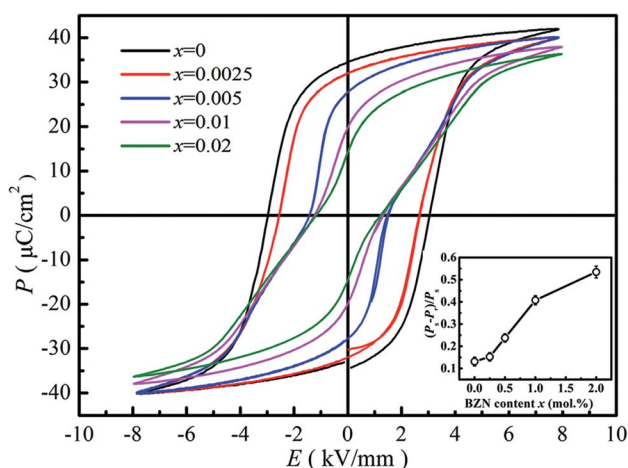


Fig. 7  $P$ - $E$  hysteresis loops of compositions  $x = 0$ – $0.02$  at 1 Hz. The inset shows the dependence of  $(P_s - P_r)/P_s$  on BZN content.

nanosized AFE relaxor phase. The inset of Fig. 7 gives the  $(P_s - P_r)/P_s$  value as a function of composition  $x$ . Obviously, the ratio  $(P_s - P_r)/P_s$  increases monotonously with BZN content,

indicating that the contribution of nanosized AFE relaxor domains to polarization enhances. In a word, the addition of BZN destroys the ferroelectric order and increases the volume of nanosized AFE relaxor domains, which will affect its field-induced strain behaviors.

Fig. 8 shows the field-induced strain behaviors of BNBZN100 $x$  ceramics. As mentioned above, BZN addition results in the percolation of AFE relaxor phase with  $P4bm$  structure down to room temperature. As seen from Fig. 8a, the compositions  $x \leq 0.0075$  exhibit the butterfly shaped  $S$ - $E$  loops, which is consistent with their nature of ferroelectricity as mentioned in previous section; hence, the back switching of ferroelectric domains results in a visible negative strain  $S_{\text{neg}}$  during bipolar cycle.<sup>31</sup> However, the compositions  $x \geq 0.01$ , the  $S$ - $E$  loops become into a sprout shape since the samples are dominated by the AFE relaxor behaviors. Fig. 8b depicts the positive strain  $S_{\text{pos}}$ , negative strain  $S_{\text{neg}}$  and the dynamic piezoelectric coefficient  $d_{33}^*$  ( $=S_{\text{max}}/E_{\text{max}}$ ) as a function of BZN content. For compositions  $x \leq 0.0075$ , the samples present a relatively large negative strain  $S_{\text{neg}}$ . Further increase  $x$  to 0.01, the negative strain  $S_{\text{neg}}$  approaches to zero. A critical composition boundary between ferroelectric response and AFE relaxor

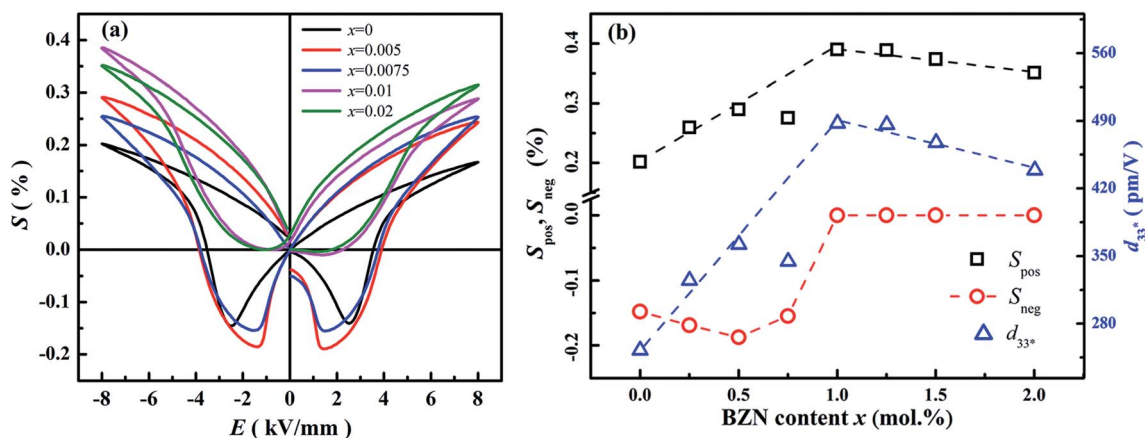


Fig. 8  $S$ - $E$  strain curves of compositions  $x = 0$ – $0.02$  at 1 Hz (a) and the composition dependence of positive strain  $S_{\text{pos}}$ , negative strain  $S_{\text{neg}}$  and the  $d_{33}^*$  as a function of BZN content (b).



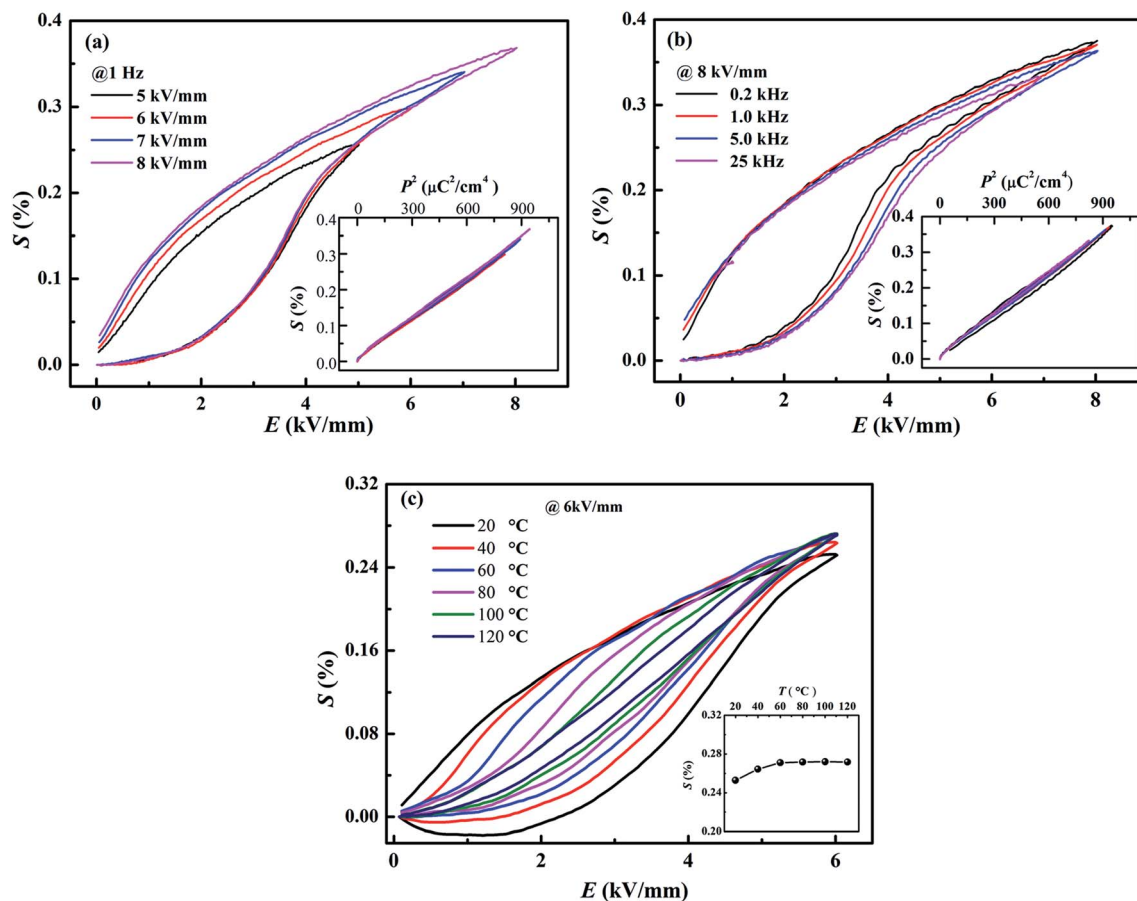


Fig. 9 Unipolar strain loops of the composition  $x = 0.01$  at different fields (a), frequencies (b) and temperatures (c). The insets in (a) and (b) show the  $S$ - $P^2$  profiles, and the inset in (c) shows the temperature-dependence of field-induced strain.

response appears at composition  $x = 0.01$ , and dynamic piezoelectric  $d_{33}^*$  up to  $487 \text{ pm V}^{-1}$  are achieved at this critical composition boundary.

Besides of the large strain, the composition  $x = 0.01$  demonstrates well stability against frequency, field and temperature. Fig. 9a and b shows the field-induced strain of the composition  $x = 0.01$  as a function of electric field and frequency, respectively. Well-defined field-induced strains at different fields and frequencies could be observed in Fig. 9a and b, and the  $S$ - $P^2$  curves almost overlap for different fields and frequencies, indicating the high stability against voltage and frequency. Besides, it is interesting that, within the temperature range from  $20 \text{ }^\circ\text{C}$  to  $120 \text{ }^\circ\text{C}$  in Fig. 9c, the field-induced strain fluctuates in the range from  $0.25\%$  to  $0.27\%$  by a slight deviation less than  $8\%$ . All in all, the composition  $x = 0.01$  shows a giant field-induced strain with excellent stability against frequency, applied field and temperature, which is of importance in application as lead-free actuators and micro-positioners.

## IV. Conclusion

Lead-free  $(1-x)(\text{Bi}_{0.5}\text{Na}_{0.5})_{0.93}\text{Ba}_{0.07}\text{TiO}_3$  (BNBT)- $x\text{Ba}(\text{Zn}_{1/3}\text{Nb}_{2/3})\text{O}_3$  (BZN) ferroelectric ceramics with perovskite structure were

successfully fabricated by a columbite route. It is found that BZN addition causes the decrease of the rhombohedrality  $90\text{-}\gamma$  and the expansion of the crystal lattice, thus inducing the phase evolution from ferroelectric ( $R3c$ ) to AFE relaxor (pseudocubic,  $P4bm$ ). Moreover, the incorporation of BZN into BNBT lattice leads to the nanosize domains of AFE relaxor reducing in the dimension and percolating to low temperature. A large field-induced strain of  $0.39\%$  and dynamic piezoelectric  $d_{33}^*$  of  $487 \text{ pm V}^{-1}$  were achieved at the composition  $x = 0.01$ , which locates at the critical composition boundary between ferroelectric state and AFE relaxor state. To be highlighted, the composition  $x = 0.01$  shows a stable field-induced strain against voltage, frequency, and temperature, making it of potential application in lead-free solid-state actuators.

## Conflicts of interest

The authors declare there is no conflicts of interest regarding the publication of this paper.

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## References

- 1 K. Uchino, *Sci. Technol. Adv. Mater.*, 2015, **16**, 046001.
- 2 J. Rödel, K. G. Webber, R. Dittmer, W. Jo, M. Kimura and D. Damjanovic, *J. Eur. Ceram. Soc.*, 2015, **35**, 1659–1681.
- 3 N. J. Donnelly, T. R. Shrout and C. A. Randall, *J. Am. Ceram. Soc.*, 2009, **92**, 1203–1207.
- 4 J. E. Daniels, W. Jo, J. Rödel, V. Honkimäki and J. L. Jones, *Acta Mater.*, 2010, **58**, 2103–2111.
- 5 E. A. Rogacheva, *Phys. B*, 2000, **291**, 359–367.
- 6 V. Dorcet, G. Trolliard and P. Boullay, *Chem. Mater.*, 2009, **20**, 5061–5073.
- 7 Y. Yu and R. N. Singh, *J. Appl. Phys.*, 2003, **94**, 7250–7255.
- 8 S.-T. Zhang, F. Yan and B. Yang, *J. Appl. Phys.*, 2010, **107**, 114110.
- 9 X. Liu, B. Liu, F. Li, P. Li, J. Zhai and B. Shen, *J. Mater. Sci.*, 2018, **53**, 309–322.
- 10 S. Zhang, A. B. Kounga, E. Aulbach, H. Ehrenberg and J. Rödel, *Appl. Phys. Lett.*, 2007, **91**, 1804.
- 11 J. Chen, Y. Wang, Y. Zhang, Y. Yang and R. Jin, *J. Eur. Ceram. Soc.*, 2017, **37**, 2365–2371.
- 12 S. Zhang, A. Devonport, N. Newman and P. K Davies, *J. Am. Ceram. Soc.*, 2015, **98**, 1188–1194.
- 13 Q. Wei, M. Zhu, L. Li, Z. Guo, M. Zheng and Y. Hou, *J. Alloys Compd.*, 2018, **731**, 631–635.
- 14 Y.-J. Siao and X. Qi, *J. Alloys Compd.*, 2017, **691**, 672–682.
- 15 S.-T. Zhang, A. B. Kounga, W. Jo, C. Jamin, K. Seifert, T. Granzow, J. Rodel and D. Damjanovic, *Adv. Mater.*, 2009, **21**, 4716–4720.
- 16 C. Ma, X. Tan, E. Dul'kin and M. Roth, *J. Appl. Phys.*, 2010, **108**, 104105.
- 17 S. Praharaaj, D. Rout, S. Anwar and V. Subramanian, *J. Alloys Compd.*, 2017, **706**, 502–510.
- 18 J. Zang, M. Li, D. C. Sinclair, W. Jo, J. Rödel and S. Zhang, *J. Am. Ceram. Soc.*, 2014, **97**, 1523–1529.
- 19 W. Jo, S. Schaab, E. Sapper, L. A. Schmitt, H. J. Kleebe, A. J. Bell and J. Rodel, *J. Appl. Phys.*, 2011, **110**, 074106.
- 20 D. I. Woodward, R. Dittmer, W. Jo, D. Walker, D. S. Keeble, M. W. Dale, J. Rödel and P. A. Thomas, *J. Appl. Phys.*, 2014, **115**, 114109.
- 21 X. Liu, F. Li, P. Li, J. Zhai, B. Shen and B. Liu, *J. Eur. Ceram. Soc.*, 2017, **37**, 4585–4595.
- 22 Y. Hiruma, K. Yoshii, H. Nagata and T. Takenaka, *J. Appl. Phys.*, 2008, **103**, 084121.
- 23 X. Liu, J. Zhai, B. Shen, F. Li, Y. Zhang, P. Li and B. Liu, *J. Eur. Ceram. Soc.*, 2017, **37**, 1437–1447.
- 24 T. Rödig, A. Schönecker and G. Gerlach, *J. Am. Ceram. Soc.*, 2010, **93**, 901–912.
- 25 B. Wylie-van Eerd, D. Damjanovic, N. Klein, N. Setter and J. Trodahl, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **82**, 104112.
- 26 L. Luo, W. Ge, J. Li, D. Viehland, C. Farley, R. Bodnar, Q. Zhang and H. Luo, *J. Appl. Phys.*, 2011, **109**, 6.
- 27 J. Kreisel, A. M. Glazer, P. Bouvier and G. Lucazeau, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2001, **63**, 145–155.
- 28 S. V. Kalinin, B. J. Rodriguez, J. D. Budai, S. Jesse, A. N. Morozovska, A. A. Bokov and Z. G. Ye, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **81**, 064107.
- 29 C. Moure and O. Peña, *Prog. Solid State Chem.*, 2015, **43**, 123–148.
- 30 A. Ullah, M. Alam, A. Ullah, C. W. Ahn, J.-S. Lee, S. Cho and I. W. Kim, *RSC Adv.*, 2016, **6**, 63915–63921.
- 31 D. L. West and D. A. Payne, *J. Am. Ceram. Soc.*, 2003, **86**, 192–194.

