Composition design, electrical properties, and temperature stability in (1 – x) K_{0.44}Na_{0.56}Nb_{0.96}Sb_{0.04}O_{3}–xBi_{0.45}La_{0.05}Na_{0.5}ZrO_{3} lead-free ceramics

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In this work, we designed a new system of (1 – x)K_{0.44}Na_{0.56}Nb_{0.96}Sb_{0.04}O_{3}–xBi_{0.45}La_{0.05}Na_{0.5}ZrO_{3} (KNNS–xBLNZ, 0 ≤ x ≤ 0.06) ceramics, and systematically investigated both their electrical performance and temperature stability. Through optimizing the composition, a relatively good comprehensive performance (e.g., \(d_{33} \approx 455 \pm 10 \) pC N\(^{-1}\), \(k_p \approx 0.47 \pm 0.02\), \(T_c \approx 266\) °C, strain \(\approx 0.148\%), and \(d'_{31} \approx 493\) pm V\(^{-1}\)) is obtained in the ceramics with \(x = 0.040\), which is attributed to the construction of a rhombohedral–orthorhombic–tetragonal (R–O–T) phase boundary. Moreover, a good temperature stability of remnant polarization \(P_r\) as well as strain value \(P_{180} = C/P_{RT} \approx 89.6\%, P_{180} = C/P_{RT} \approx 73.2\%\). \(S_{100} = C/S_{RT} \approx 92.6\%\), \(S_{180} = C/S_{RT} \approx 74.1\)%) is gained in KNNS–0.040BLNZ ceramics with a broad temperature range from room temperature to 180 °C. Hence, we believe that KNNS–xBLNZ ceramics opens a window for the practical application of lead-free ceramics.

1. Introduction

Piezoelectric materials play an important role in electronic devices because they can interconvert electrical energy into mechanical energy. Lead zirconate titanate (PbZr\(_{1-x}\)Ti\(_x\)O\(_3\), PZT) ceramics have been the dominating material in the piezoelectric market during the last 60 years.\(^1\) However, in order to protect the environment and human sustainable development due to the toxicity of lead in the compositions, a lot of effort has been recently devoted to develop high-performance lead-free piezoelectric ceramics which can eventually replace the lead-based ones.\(^2\)–\(^26\)

Among all the lead-free piezoelectric materials, (K, Na) NbO\(_3\) has been regarded as a promising candidate due to its good comprehensive electrical properties especially its flexibility in terms of compositional modification.\(^2\)–\(^26\) Saito et al. reported that alkali niobate ceramics based on Li, Ta, and Sb-modified KNN show high density and excellent performance (\(d_{33} \approx 416\) pC N\(^{-1}\); \(T_c \approx 253\) °C) by the reactive-templated grain growth method in 2004.\(^2\) Ever then, researchers paid much attention to promote the electricity by constructing the multiphase boundary which has been extensively applied in KNN-based lead-free materials.\(^2\)–\(^26\) Recently, some breakthroughs

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2. Experimental procedure

(1 − x)K0.44Na0.56Nb0.96Sb0.04O3·xBi0.45La0.05Na0.5ZrO3 (0 ≤ x ≤ 0.06) piezoelectric ceramics were prepared by the conventional solid-state reaction process. Raw materials [Na2CO3 (99.8%), K2CO3 (99%), Nb2O5 (99.5%), Sb2O3 (99%), ZrO2 (99%), Bi2O3 (99.99%), and La2O3 (99.99%)] were weighed according to chemical formula above. The powder dried and calcined at 850 °C for 6 h after being mixed with ZrO2 balls for 24 h with alcohol as the dispersion medium, and then pressed into the pellets (≈10 mm diameter, ≈1 mm thickness) under a pressure of ≈10 MPa when the calcined powders were mixed with 8 wt% polyvinyl alcohol (PVA). After the PVA binder was burnt out, the samples were sintered at 1110–1130 °C for 3 h in air, and then printed with silver paste on both sides as the electrode. All specimens were polarized in a silicone oil bath at room temperature under a dc field of 3 kV mm⁻¹ for 30 min.

The phase structure of the sintered cylindrical pellets is identified by X-ray diffraction meter (Cu, DX-2700, Dandong, China), measured at 40 kV and 30 mA. The field-emission scanning electron microscope (FE-SEM) (JSM-7500, Japan) has been used to measure their surface micro-structure and chemical compositions. The dielectric properties of the sintered samples were measured using an LCR analyzer (HP 4980, Agilent, U.S.A.) with varied temperatures in the range of −150 to 200 °C and 25–450 °C. The P–E loops and S–E curves of the cylindrical pellets were investigated by the ferroelectric tester (aixACCT, TF Analyzer 2000E, Germany). The kₚ was measured by a resonance-antiresonance method with the impedance analyzer (Impedance Analyzer, PV70A, Beijing, China). The d₃₃ was measured by the commercial Berlincourt-type of d₃₃ meter, (ZJ-3A, China) for the poled samples.

3. Results and discussion

The XRD patterns of the KNNS-xBLNZ (0 ≤ x ≤ 0.06) ceramics in the 2θ range of 20–60° is shown in Fig. 1(a). All samples exhibit a pure perovskite structure, indicating that BLNZ has completely diffused into the KNNS lattice to form a solid solution. The O phase is characterized by (220)/(020) peaks splitting at about 45°, that is, the intensity ratio (2 : 1) between the left peak and the right one means the O symmetry. From the Fig. 1(b), we can find that an O phase was formed in the ceramics (0 ≤ x < 0.020), and other ceramics (0.02 ≤ x < 0.060) demonstrate multiphase structure. To further explore their phase structure of the ceramics, the εₛ–T curves (−150–200 °C) tested at 10 kHz have been plotted in Fig. 2(a)–(h). There are two obvious abnormal dielectric peaks in the ceramics with x ≤ 0.040, which respectively correspond to their T₉–O at low temperature and T₉–O at high temperature, as shown in Fig. 2(a)–(e). The T₉–O (≈136 °C) and T₉–O (≈2 °C) is far to room temperature in the ceramics with 0 ≤ x < 0.020, showing that the O phase was observed [see Fig. 2(a)]. For the ceramics with 0.02 ≤ x ≤ 0.030, the T₉–O (≈87 → 75 °C) is close to room temperature and T₉–O (≈3 → 7 °C) is below room temperature, confirming that the O–T phase was obtained [see Fig. 2(b) and (e)]. As for BLNZ content increasing from 0.035 to 0.040, T₉–O (≈9 → 13 °C) and T₉–O (≈51 → 40 °C) are close to room temperature, suggesting that the R–O–T phase was constructed in the ceramics [see Fig. 2(d) and (e)]. Subsequently, only one abnormal dielectric peak was observed in the ceramics with 0.045 ≤ x ≤ 0.060, indicating that both T₉–O and T₉–O were shifted to the same temperature, and formed the R–T phase boundary [see Fig. 2(f) and (g)]. In addition, the εₛ–T curve of x = 0.060 becomes flat to some degree, leading to the R phase boundary [see Fig. 2(h)], which can be explained by the fine-grain [see Fig. 4(d)]. Eventually, through systematical analysis of the phase structure by XRD and εₛ–T curves, we concluded that: the ceramics with 0 ≤ x < 0.020 are an O symmetry, the ceramics with 0.020 ≤ x ≤ 0.030 belong to be O–T phase coexistence, the ceramics with 0.035 ≤ x ≤ 0.040 have R–O–T phase boundary, the ceramics with 0.040 < x < 0.060 are R–T phase boundary, the ceramic with x = 0.060 is R phase boundary.
Fig. 3(a) shows the $\varepsilon_r-T$ curves (25–450 °C) of the KNNS-xBLNZ ceramics. We have concentrated mainly on the variation of ferroelectric–paraelectric phase transition temperature ($T_c$) versus BLNZ content. One can see that the $T_c$ decreases with increasing the BLNZ content. Fig. 3(b) plots the corresponding phase diagrams of the ceramics as a function of BLNZ content. One can see that the $T_c$ and $T_{O-T}$ values have a similar variation trend, that is, they reduce monotonously with increasing BLNZ content, but the $T_{R-O}$ rises. It is reported that both (Bi$_{0.45}$La$_{0.05}$Na$_{0.5}$)$_{2+}$ and Zr$^{4+}$ can simultaneously drop $T_{O-T}$ and raise $T_{R-O}$ in KNN-based ceramics.\cite{3,20} Finally, the O symmetry gradually disappears with increasing of the BLNZ content, resulting in the appearance of multiphase coexistence (O-T for 0.020 ≤ x ≤ 0.030, R-O-T for 0.035 ≤ x ≤ 0.040, R-T for 0.040 < x < 0.060, and R for x = 0.060).

Fig. 4 shows the FE-SEM micrographs of the KNNS-xBLNZ ceramics. One can see that the grain size strongly depends on the BLNZ content in this ceramics. From Fig. 4(a), a relatively...
homogeneous grain size distribution is found in the ceramics with \( x = 0 \). The grain size of the ceramics \( (0.020 \leq x \leq 0.040) \) is increasing with introducing BLNZ content, and the inhomogeneous grains appear in the ceramics [see Fig. 4(b) and (c)]. It may be caused by the addition of Bi\(^{3+}\), which can improve the migration rate of the grain boundary, so as to accelerate the grain growth of the KNN-based ceramics due to the formation of liquid phase.\(^{23}\) Moreover, the small grains stuff the aperture of the large ones can lead to the formation of the dense microstructure in the ceramics. One can find that the grain size sharply decreases as BLNZ content increases to 0.060 [see Fig. 4(d)], which can be interpreted as the inhibited grain growth due to the aggregation at grain boundaries of both excessive Zr\(^{4+}\) and Bi\(^{3+}\). In addition, the fine-grain plays a key role in the flat dielectric peak in the ceramics of \( x = 0.060 \). The element mapping (K, Na, Nb, Sb, Bi, La, Zr, O) of KNNS-\( x \)BLNZ ceramics with \( x = 0.040 \) is shown in Fig. 5 shows. All chemical elements in the ceramics distribute uniformity.

Fig. 6 shows \( \varepsilon_r \) and tan \( \delta \) values of KNNS-xBLNZ ceramics. One can find that the \( \varepsilon_r \) value is sensitive to the BLNZ content, that is, \( \varepsilon_r \) value rises sharply with increasing BLNZ content, getting a high value (2003–2376) in the ceramics with \( 0.035 \leq x \leq 0.050 \), which can be ascribed to the existence of multiphase boundary. The tan \( \delta \) value almost fluctuates in the range of 0.035–0.044 in the ceramics due to the dense microstructure, as is shown in Fig. 6.

Fig. 7(a) plots the \( P-E \) loops of KNNS-xBLNZ ceramics. We can see that all ceramics exhibit typical ferroelectric loops in Fig. 7(a). The remnant polarization \( (P_r) \) and the coercive field \( (E_c) \) as a function of BLNZ content were shown in Fig. 7(b). As is known, the \( P_r \) value is the remained aligned dipoles after removing electric field, and the \( E_c \) value is the electric field required for domain switching. The \( P_r \) value slowly decreases with increasing BLNZ content to 0.045, and then reduces sharply with further increasing the BLNZ content [see Fig. 7(b)]. The \( E_c \) value almost fluctuates in the range of 7.8–9.5 kV cm\(^{-1}\) in the ceramics [see Fig. 7(b)]. The piezoelectric properties of KNNS-xBLNZ ceramics is plotted in Fig. 8(a). One can see that the \( d_{33} \) value firstly increases and then decreases as increasing the BLNZ content. The \( k_p \) value firstly keeps stable with increasing the BLNZ content to 0.045, and then decreases sharply with further increasing the BLNZ content. Owing to the coexistence multiphase of R–O–T, an optimum piezoelectric \( (d_{33} \sim 455 \pm 10 \text{ pC N}^{-1}, k_p \sim 0.47 \pm 0.02) \) properties were obtained in the ceramics with \( x = 0.040 \). It can be explained by the following two aspects: firstly, the multiphase boundaries have more equivalent spontaneous polarization vectors, indicating that it can provide more polarization vectors than the one with a pure phase structure. Secondly, the polarization anisotropic energy of the ceramics can be decreased in multiphase coexistence, which provides a shortcut for polarization rotation in anisotropically flattened free energy profile.\(^{26–28}\)

As is known, the formula of \( d_{33} \sim \alpha \varepsilon_r P_r \) appears to be well established in KNN-based ceramics,\(^{29,30}\) namely, the higher \( d_{33} \) value should be closely correlated with the higher \( \varepsilon_r P_r \) value in the ceramics. To explore the relationship between \( d_{33} \) and \( \varepsilon_r P_r \), a diagram of \( d_{33} \) and \( \varepsilon_r P_r \) vs. BLNZ content is shown in Fig. 8(b). One can see that there is a similar trend for the \( d_{33} \) and \( \varepsilon_r P_r \) vs. BLNZ content. Therefore, the improvement of piezoelectric properties can be ascribed to the multiphase coexistence, dielectric and ferroelectric properties.

Bipolar electric field induced strain curves of KNNS-xBLNZ ceramics is shown in Fig. 9(a). A typical butterfly shape strain

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**Fig. 4** FE-SEM surface images of \((1-x)\)K\(_{0.44}\)Na\(_{0.56}\)Nb\(_{0.96}\)Sb\(_{0.04}\)O\(_3\)-xBi\(_{0.45}\)La\(_{0.05}\)Na\(_{0.5}\)ZrO\(_3\) ceramics with (a) \( x = 0 \), (b) \( x = 0.020 \), (c) \( x = 0.040 \), (d) \( x = 0.060 \).
curves as a characteristic of the ferroelectric is exhibited in all ceramics. It was reported that the electric field-induced strain comes from intrinsic and extrinsic contributions. The intrinsic contributions mainly include the piezoelectric effect and electrostriction, and the extrinsic contributions are attributed to domain wall movement. In addition, the perfectly linear $S$–$E$ response is the main contribution by the piezoelectric effect is, and the domain switching mainly produce “butterfly shape” of $S$–$E$ loops.\textsuperscript{31} In this work, one can see that the “butterfly shape” of $S$–$E$ loops is obtained in ceramics with $0 \leq x \leq 0.050$. It can be explained by the intrinsic effect which is aligned with the lattice response and the extrinsic effect which is aligned with domain switching; Moreover, the nearly linear response of strain is observed in the ceramics with $x = 0.060$, indicating that both the lattice response and domain wall movement contribute to the strain. Fig. 9(b) shows the strain and $d_{33}^*$ values vs. the compositions of BLNZ. The strain and $d_{33}^*$ value is similar variation with increasing BLNZ content, that is, the strain value firstly increases, and then decreases with increasing the BLNZ, obtaining the maximum value (strain $\sim 0.148\%$, $d_{33}^* \sim 493$ pm V$^{-1}$) at $x = 0.040$, which is benefit from the multiphase zone.

Fig. 10(a) represents the temperature-dependent $P$–$E$ curves of the ceramics with $x = 0.040$, measured at $f = 1$ Hz and room temperature to 180 °C. One can find that all are statured $P$–$E$ loops with increasing the measured temperature. The remnant polarization $P_r$ changed slightly with high $P_{rT}/P_{rRT}$ ($P_{rT}$ is the $P_r$ value which measured at the temperature of $T_r$; $P_{rRT}$ is the $P_r$ value which measured at room temperature) values ($P_{r100}^\circ C/P_{rRT} \sim 89.6\%$, $P_{r180}^\circ C/P_{rRT} \sim 73.2\%$) in the wide temperature range

Fig. 5 Elemental (K, Na, Nb, Sb, Bi, La, Zr, and O) mapping on the surface of $(1 - x)K_{0.44}Na_{0.56}Nb_{0.96}$Sb$_{0.04}$O$_3$–xBi$_{0.45}$La$_{0.05}$Na$_{0.5}$ZrO$_3$ ceramics with $x = 0.040$.

Fig. 6 Dielectric ($\varepsilon$, tan $\delta$) properties of $(1 - x)K_{0.44}Na_{0.56}Nb_{0.96}$–Sb$_{0.04}$O$_3$–xBi$_{0.45}$La$_{0.05}$Na$_{0.5}$ZrO$_3$ ceramics.
It indicates that the switched polarization state along the direction of the external electric field was of little variation. As another important characteristic for ferroelectric property, the coercive field ($E_c$) which indicates the energy barrier between the equivalent states of the order parameter decreased on increasing temperature. Fig. 10(d) shows bipolar electric field induced strain curves vs. temperature of KNNS-xBLNZ ceramics, measured at room temperature to 180 °C. All the strain curves show the typical butterfly shape. Different from the hysteresis loops, which involved all domain switching, the bipolar strain curves only reflected the population of non-180° domain switching. For ferroelectrics, the non-180° polarization switching is always coupled to the lattice strain, an instability in polarization orientation (i.e., isotropy of polarization) will necessarily induce a softening in elastic modulus, and hence a high electrostrain. Therefore, low polarization anisotropy and elastic softening go hands in hands, both contributing to a high electrostrain. As summarized in Fig. 10(b), the $S_{\text{neg}}$ (the strain between minimum strain and strain under zero electric field) of the ceramics decreased. The changes in $S_{\text{neg}}$ can be explained by the quantity of 90°/180° domain increased and tetragonality c/a decreased with increasing temperature approaching its $T_C$. While the $S_{\text{pos}}$ (the strain between the maximum strain and the
strain under zero electric field) showed little change with high $S_{RT}/S_{RT}$ ($S_{RT}$ is the $S$ value which measured at the temperature of $RT$; $S_{RT}$ is the $S$ value which measured at room temperature) values ($S_{100 \degree C}/S_{RT} \sim 92.6\%$, $S_{180 \degree C}/S_{RT} \sim 74.1\%$) in the wide temperature range [see Fig. 10(e) and (f)]. The excellent temperature stability is due to the high $T_C$ ($\sim 266 \degree C$) and relatively stable $\varepsilon_r$ in this composition, which has been demonstrated elsewhere. Therefore, the $P_r$ and strain values of KNNS-xBLNZ ceramics with $x = 0.040$ exhibit excellent temperature stability ($P_{1100 \ degree C}/P_{RT} \sim 89.6\%$, $P_{180 \ degree C}/P_{RT} \sim 73.2\%$, $S_{100 \ degree C}/S_{RT} \sim 92.6\%$, $S_{180 \ degree C}/S_{RT} \sim 74.1\%$) in the wide temperature range.

4. Conclusion

$(1-x)K_{0.44}Na_{0.56}Nb_{0.96}Sb_{0.04}O_3-xBi_{0.45}La_{0.05}Na_{0.5}ZrO_3$ lead-free ceramics were developed by the normal sintering, and the effects of BLNZ content on their phase structure and electrical properties was investigated. The R–O–T coexistence phase has been confirmed by both XRD patterns and $\varepsilon_r$–$T$ curves in the ceramics with $0.035 \leq x \leq 0.040$. Through optimizing the composition, the relatively good comprehensive performance (e.g., $d_{33} \sim 455 \pm 10 \mu C \cdot cm^{-2}$, $K_p \sim 0.47 \pm 0.02$, $T_C \sim 266 \degree C$, strain $\sim 0.148\%$, and $d_{33}^e \sim 493 \text{ pm} \cdot \text{V}^{-1}$) is obtained in the ceramics with $x = 0.040$ which is attributed to the construction of rhombohedral–orthorhombic–tetragonal (R–O–T) phase boundary. Moreover, a good temperature stability of remnant polarization ($P_r$) as well as strain value ($P_{1100 \ degree C}/P_{RT} \sim 89.6\%$, $P_{180 \ degree C}/P_{RT} \sim 73.2\%$, $S_{100 \ degree C}/S_{RT} \sim 92.6\%$, $S_{180 \ degree C}/S_{RT} \sim 74.1\%$) is gained in KNNS-0.040BLNZ ceramics with a broad temperature range from room temperature to 180 $\degree C$. Hence, such a material is promising for practical applications in the piezoelectric material markets.

Conflicts of interest

There are no conflicts to declare.

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