

A-site size effect in a family of unfilled ferroelectric tetragonal tungsten bronzes: $\text{Ba}_4\text{R}_{0.67}\text{Nb}_{10}\text{O}_{30}$ ($\text{R} = \text{La}, \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}$ and Y)[†]

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The effect of A-cation size on the structural and electrical properties in a family of ferroelectric tetragonal tungsten bronzes (TTBs) $\text{Ba}_4\text{R}_{0.67}\square_{1.33}\text{Nb}_{10}\text{O}_{30}$ ($\text{R} = \text{La}, \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}$ and Y ; $\square = \text{vacancy}$) was investigated. In each case, the crystal structure, as determined from lab-based ambient powder X-ray diffraction (PXRD), is metrically tetragonal and can be refined in the $P4bm$ space group. XRD data show an increased splitting of $hk0$ $00l$ reflections with decreasing R cation size indicating an increasing tetragonal distortion (measured by tetragonality c/a). Dielectric data and ferroelectric measurements indicate that the ferroelectric Curie temperature, T_C , increases with decreasing R size and so a direct correlation between T_C and tetragonality/ionic radius of R is demonstrated. Rietveld refinements show that the large A2-site is fully occupied by Ba^{2+} and, in addition to the R cation size, the presence of vacancies at the A1-site (perovskite-like site) is also shown to strongly affect the stability of ferroelectricity in this structure type.

Introduction

Modern applications for ferroelectrics and dielectrics are still dominated by materials with the perovskite structure, largely due to the possibility of structure/property manipulation rationalised *via* geometrical tolerance factors and tilt systems.^{1–3} The development/growth of the electronics industry has driven the investigation of other structures in the search for new materials to *e.g.* replace existing lead-based compounds. Materials with the tetragonal tungsten bronze (TTB) structure were reported to display ferroelectric properties by Goodman in 1953^{4,5} and offer great scope for materials development due to their similar compositional flexibility compared to the closely related perovskite structure.

The TTB structure has general formula $(\text{A1})_2(\text{A2})_4(\text{B1})_2(\text{B2})_8(\text{C})_4(\text{X})_{30}$ (where X = fluorine or more commonly oxygen and A, B, C are metal cations) and consists of square A1-site, perovskite-like, units linked by B1-site BO_6 units, which together demarcate a large pentagonal tunnel, the A2-site (Fig. 1).⁵ A1- and A2-sites are crystallographically distinct but both are occupied by larger cations: alkaline,⁶ alkaline earth,⁷ rare earth,⁸ or lead,⁵ however, substitutions of heavier cations such as bismuth⁹ or thorium¹⁰ have been reported. Where two (or more) A-cation types are present, the

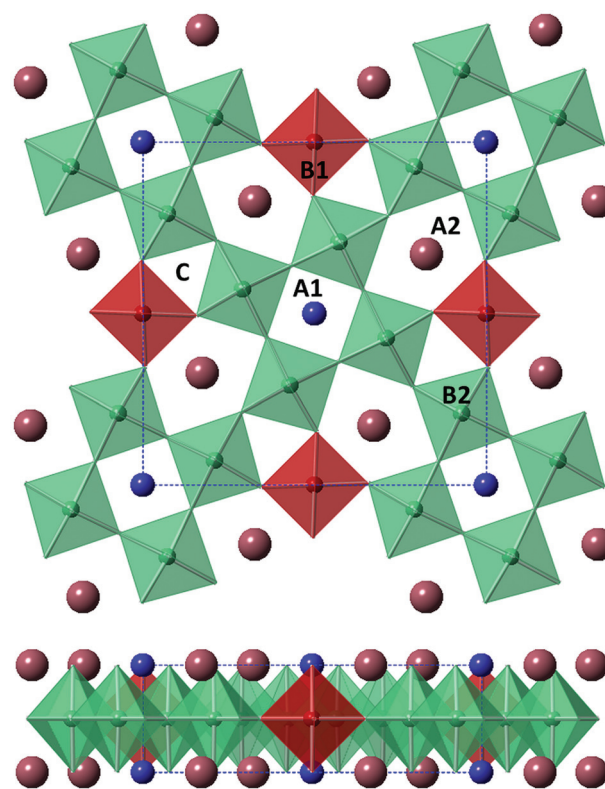


Fig. 1 Tetragonal tungsten bronze (TTB) structure (space group $P4/mbm$) viewed along $[001]$ (top) and $[100]$ (bottom). Oxygen octahedra contain B1-cations (red) and B2-cations (green), respectively, with the unit cell indicated by blue dashed lines.

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larger A2-site is generally preferentially occupied by the larger cations.^{4,11} The B1- and B2-sites are non-equivalent BO₆ octahedral sites and usually occupied by small and highly charged cations: Nb, Ti, Ta or W,¹² although minimal amount of doping of first row transition metals and others is possible.^{13–15} The C-site is of sufficient size to only accommodate the smallest cations, *e.g.* Li⁺, and consequently is vacant in most reported TTB compounds. When all cation sites are occupied the structure is described as being ‘stuffed’, while in ‘filled’ structures A- and B-sites are fully occupied but the C-site is vacant.^{16,17} When vacancies are present within the A-sites, the structure is described as ‘unfilled’.¹⁷ Alternative expressions of the general TTB formula are common in the literature. Filled oxide compounds are commonly expressed in the reduced form (A1)(A2)₂(B1)(B2)₄O₁₅ (A₃B₅O₁₅) *e.g.* Ba₂NaNb₅O₁₅⁶ while unfilled compounds are described by the further reduced forms AB₂O₆ *e.g.* (Ba, Sr)Nb₂O₆.¹²

The cell dimensions of the TTB structural aristotype are typically $a = b \approx 12.5 \text{ \AA}$ and $c \approx 4 \text{ \AA}$ (and is assigned space group $P4/mbm$).⁴ An orthorhombic distortion, leading to an expansion in the ab plane (such that $a_o = \sqrt{2}a$, $b_o = \sqrt{2}b$) as well as further cell doubling in a , b and/or c is common.¹⁸ Incommensurate superstructures, primarily originating from modulations of the oxygen sub-lattice, have been reported for several TTB systems.¹⁸ These features are often subtle and therefore difficult to characterise with conventional powder X-ray diffraction methods; consequently most studies concerning these features employ electron, single crystal or neutron diffraction techniques. Polarisation in TTBs is, in most instances, constrained within the c axis, although in lead (and similar cations *e.g.* Bi³⁺)-based compounds the polar axis is rotated into the ab plane.⁴ This may also be observed in charge ordered systems.¹⁹

Previous investigation into the properties of this structure has been predominately focused on filled TTBs, specifically mixed titanate–niobates. Stennett and co-workers observed an increase in T_C with decreasing M ionic radius at the A1-site in the filled TTB, Ba₂MTi₂Nb₃O₁₅ (M = Bi, La, Pr, Nd, Sm, Eu, Gd and Dy), with a transition between normal ferroelectric (M = Pr to Dy) to relaxor ferroelectric (M = Bi and La) behaviour.^{20,21} A superstructure in which the c -axis of the regular TTB structure is doubled was identified, with commensurate and incommensurate structures observed, correlating with normal ferroelectric and relaxor ferroelectric states, respectively. This has been attributed to the average rare earth size at the A1-site driving tilting of the BO₆ octahedra, leading to an incommensurate modulation of the structure.²² The influence of A2-site size was shown to be important by Zhu *et al.* in (Ba_{*x*}Sr_{1–*x*})₄Nd₂Ti₄Nb₆O₃₀, who suggested that the difference in the mean size of A1-site cations and A2-site cations, rather than absolute A1-site cation size, was responsible for inducing octahedral tilting, with decreasing size difference destabilising ferroelectric ordering and leading to the onset of incommensurate periodicity.²³

B-site substitution has also been shown to affect ferroelectric properties of TTBs in compounds in which A-site size is

constant.^{24–26} Increasing M⁴⁺ size in Ba₆M⁴⁺₂Nb₈O₃₀ shows a decrease in T_C (or T_m), while increasing M³⁺ in Ba₆M³⁺Nb₉O₃₀ increases T_C (or T_m). A relationship between the structural distortion, quantified by tetragonality (c/a), originating from these substitutions and T_C has been shown to reconcile these apparently conflicting trends.²⁷

The properties of unfilled TTBs have been comparatively less well examined, with the exception of SBN.¹² Masuno reported a series of unfilled ferroelectric barium niobate TTBs, (Ba_{1–*x*}RE_{2*x*/3})Nb₂O₆, (R = Y, Sm or La) with single phase compounds in the approximate range $x = 0.2$ to 0.4 , depending on R.⁸ Higher dopant levels depressed the Curie temperature, T_C , with a maxima in T_C occurring for the composition corresponding to $x = 0.2$, Ba₄R_{0.67}Nb₁₀O₃₀. Higher values of T_C were reported with decreasing R cation size for each value of x . A similar relationship between T_C and ionic radius was shown to exist by Wakiya *et al.*²⁸ in Ba_{3.75}R_{0.833}Nb₁₀O₃₀, R = Nd, Sm, Eu, Gd, Y, Ho and Er (corresponding to $x = 0.25$ in Masuno’s compounds).

Wakiya and co-workers assigned A1- and A2-site distribution in Ba_{3.75}R_{0.833}Nb₁₀O₃₀ (R = Nd, Sm, Eu, Gd, Y, Ho and Er) based on analysis of powder X-ray diffraction, however, did not report lattice parameters.²⁸ We have produced a series Ba₄R_{0.67}Nb₁₀O₃₀ TTB compounds of various R³⁺ size to examine if size difference and tetragonality concepts are applicable to unfilled systems. Previous reports on both unfilled and filled rare earth-doped TTBs, have indicated that when cations of different sizes occupy the A1- and A2-site, the larger cation preferentially occupies the larger A2-site.^{4,11} In filled rare earth-doped TTBs of A-site composition Ba₄RE₂, the size difference appears to be sufficiently large such that Ba²⁺ and RE³⁺ are restricted to A2- and A1-sites respectively.²⁹ It is expected that Ba₄R_{0.67}Nb₁₀O₃₀ compounds display similar cation ordering with Ba²⁺ fully occupying the larger A2-site and with R³⁺ (and vacancies) on the A1-site. This allows the influence of A-site-size differences and tetragonality/structural distortion on the ferroelectric properties of this unfilled TTB system to be tested while excluding B-site effects and avoiding mixed A-site cation distribution. The R-cations reported here were specifically selected by two criteria: (i) they only adopt +III valence state and hence avoids the possibility of alternative charge compensation mechanism(s) on doping and (ii) they are of sufficiently large ionic radius to occupy the A-sites only.

It is important to re-emphasise that many TTB compounds adopt larger unit cells due to an orthorhombic distortion and/or incommensurate modulations in the ab -plane. These distortions are frequently subtle and require advanced characterisation techniques such as synchrotron X-ray, neutron or selected area electron diffraction methods. Such techniques, however, are not routinely available and either involve long lead times to access central facilities or are destructive (*e.g.*, SAED of sintered ceramics). The majority of structural variations of the TTB structure, however, involve $\sqrt{2}$ or $2\sqrt{2}$ extension of the a and b lattice parameters, and/or doubling in the c -axis and so can also be described in terms of reduced cell parameters as for perovskites. In many cases such reduced cell



Table 1 Lattice parameters, tetragonality (c/a), unit cell volume and goodness-of-fit statistics, obtained from refinements of ambient temperature PXRD data in space group $P4bm$, and T_C for $Ba_4R_{0.67}\square_{1.33}Nb_{10}O_{30}$

R	a (Å)	c (Å)	c/a	Volume (Å ³)	χ^2	R_{wp}	$R[F^2]$	T_C (K)
Y	12.45172(14)	3.95503(6)	0.3176	613.209(14)	5.76	0.193	0.23	537
Dy	12.45845(16)	3.95434(6)	0.3174	613.765(14)	3.81	0.157	0.169	524
Gd	12.46285(18)	3.95474(7)	0.3173	614.260(17)	4.06	0.159	0.121	491
Sm	12.46758(17)	3.95548(7)	0.3172	614.843(16)	3.80	0.151	0.151	459
Nd	12.47453(7)	3.95439(4)	0.3170	615.358(7)	3.96	0.148	0.147	406
La ^a	12.48735(13)	3.95577(5)	0.3168	616.839(12)	2.91	0.145	0.127	297 ^b

^a Refined in $P4bm$ and $P4/mbm$ with near identical values. ^b T_m at 1 MHz.

parameters result in a metrically tetragonal representation. Our approach in this study is to utilise this observation allowing widely accessible lab-based PXRD analysis to simply determine tetragonality (without the need for complex techniques or larger units cells) and which can then be empirically correlated with T_C (Table 1).

Experimental methods

Polycrystalline ceramic samples of $Ba_4R_{0.67}\square_{1.33}Nb_{10}O_{30}$ (R = La, Nd, Sm, Gd, Dy, Ho, Er, Yb and Y; \square = vacancy), were synthesised using a conventional solid state method. Precursor powders (all >99% purity) of: $BaCO_3$, Nb_2O_5 , La_2O_3 , Nd_2O_3 , Sm_2O_3 , Gd_2O_3 , Gd_2O_3 , Dy_2O_3 , Ho_2O_3 , Er_2O_3 , Yb_2O_3 and Y_2O_3 (Aldrich); Er_2O_3 (Apollo Scientific); and Yb_2O_3 (Alfa Aesar), were dried prior to use. $BaCO_3$ and Nb_2O_5 were dried at 600 °C, the remainder at 1000 °C. The appropriate stoichiometric ratios of the precursor reagents were homogenised as an acetone slurry in an agate mortar and pestle. The resulting dry powder was formed into a cylindrical body by manual compression using a 12 mm diameter steel die press then heated in an alumina boat lined with platinum foil for 2 hours at 1000 °C, immediately followed by 15 hours at 1250 °C. Samples were air quenched, ground to a powder then milled in ethanol by planetary ball mill for 1 hour (600 r.p.m.). A 10 mm die press was used to form pellets of approximately 0.5–2 mm thickness by uniaxial pressing. Pellets were heated in a tube furnace for 6 hours at temperatures of 1350, 1375 or 1400 °C (heating/cooling rates of 20 °C min⁻¹) resulting in ceramics of typically greater than 95% theoretical density as calculated from refined lattice parameters. R = Y required additional sintering to yield single phase ceramics, with optimum duration found to be 12 hours for the above temperatures. Despite repeated attempts and successive sintering stages single phase TTB analogues of R = Ho, Er and Yb were not obtained.

Structural characterisation was carried out using powder X-ray diffraction. Data were collected over a range of 10–90° 2-theta with a PANalytical Empyrean diffractometer operating in reflection mode, Cu K α_1 radiation (1.540598 Å) and step size of 0.0167 degrees 2-theta. The General Structure Analysis System (GSAS)³⁰ and associated graphical user interface, EPGUI³¹ were used for structural (Rietveld) refinements. For

PXRD data the following parameters were refined: lattice parameters, background, scale, peak profile and sample displacement. Atomic positions and isotropic displacement parameters were refined for the cations, but fixed for oxygen. Sample microstructure was investigated using a Joel JSM 5600 scanning electron microscope (SEM). Micrographs were taken of the thermally etched (1215 °C, 10 h) fracture surfaces of sintered pellets. Platinum electrodes were sputtered onto the polished faces of sintered pellets for electrical characterisation. Dielectric and immittance measurements were taken over a frequency range of 0.025 and 2000 kHz with Agilent 4294A and Wayne Kerr 6500B impedance analysers between 50 and 873 K using a closed cycle cryocooler and Carbolite MTF 10/25/130 tube furnace. Polarisation–electric field (P – E) measurements were recorded using an AixACCT TF Analyzer 2000, TREK 609E-6 high voltage amplifier, variable temperature (ambient to 473 K) AixACCT piezo sample holder (model TFA 317-7 with sample immersed in Dow Corning 200/100 cs silicone fluid (BDH Ltd)) or closed cycle cryocooler.

Results and discussion

A. Crystal structure

Ceramic samples of $Ba_4R_{0.67}\square_{1.33}Nb_{10}O_{30}$ (R = La, Nd, Sm, Gd, Dy, Ho, Er, Yb and Y), were prepared as described above and powder X-ray diffraction (PXRD) used to establish the formation of the crystalline TTB structure and ensure the absence of impurity phases. Compositions with R = La to Dy and also Y, resulted in nominally single phase TTBs. Attempts at producing R = Ho, Er and Yb analogues yielded phase mixtures of a TTB main phase and significant quantities of $BaNb_2O_6$ (see ESI†) suggesting that these R cations are too small to be incorporated in these amounts. It is interesting to note that Wakiya *et al.* also reported that they could not obtain single phase $Ba_{3.75}Ho_{0.833}Nb_{10}O_{30}$, although they reported success for the Er analogue.²⁸ The incorporation of Yb into the TTB structure has been reported but to our knowledge only in minimal concentrations.^{32,33} Based on these observations the smaller rare earth Lu was not investigated.

As mentioned previously, TTB structures can adopt several different symmetries, predominantly in the tetragonal and orthorhombic crystal systems, although distortions are usually sufficiently subtle that it is difficult to distinguish between



them. Rietveld refinements were carried out on PXRD data using structural models with the non-centrosymmetric $P4bm$ and centrosymmetric $P4/mbm$ tetragonal space groups. These space groups are commonly reported for the ferroelectric ($P4bm$) and paraelectric ($P4/mbm$) polymorphs of TTBs.^{34–37} including those of similar compositions.^{11,28,38} Refinements in both space groups yield near identical lattice parameters, but differing atomic positions. Refinements with other space groups reported for TTBs, notably orthorhombic $Pba2$,²⁴ resulted in poor, divergent or unstable refinements and so, based on the laboratory XRD data obtained, there was no evidence to warrant further lower of symmetry from $P4bm$.

All observed reflections were indexed by the $P4bm$ model, however, a number of calculated peak intensities deviate slightly for data collected on both powder and pellets. These differences may result from preferred orientation associated with the columnar grain morphology (described later), but this was not investigated in detail. Attempts at refining A-cation occupancies both with Ba^{2+} confined to the A2-site and also allowing cross substitution of Ba^{2+} and R^{3+} between the A1- and A2-sites, resulted in unstable, unrealistic, or inferior refinements. Refinements where A-site cation occupancies were consequently fixed such that the A2-site is fully occupied by Ba^{2+} and the two A1-sites partially occupied by $0.67R^{3+}$ and $1.33\Box$ to achieve the nominal stoichiometry, $Ba_4R_{0.67}\Box_{1.33}Nb_{10}O_{30}$ resulted in significantly improved (and stable) fits. (It is important to note that, especially for the early period rare earths, the X-ray scattering factors are not significantly different to Ba.)

The lattice parameters indicate increasing tetragonality, c/a , with decreasing size of R-cation which nominally occupies the A1-site, Fig. 2a. The increase in c/a is predominantly driven by decreasing a as the ionic radius of R decreases, whilst the c parameter is comparatively invariant. The resultant decrease in cell volume with decreasing R radius is thus also largely driven by contraction in the ab plane (see ESI†).

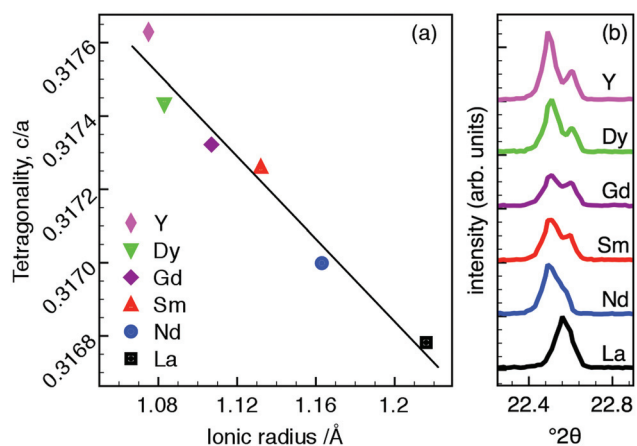


Fig. 2 (a) Tetragonality (c/a), as a function of ionic radius for $Ba_4R_{0.67}\Box_{1.33}Nb_{10}O_{30}$ (all ionic radii are for IX coordination³⁹) and (b) splitting of ambient temperature PXRD 310 and 001 reflections ($P4bm$) showing increasing distortion.

Peak splitting of differing magnitude was observed in 4 pairs of $hk0\ 00l$ reflections in the PXRD data of each compound. This is most clearly evident for the 001 and 310 reflections as shown in Fig. 2b. The magnitude of the splitting increases with decreasing R radius and consequently increasing tetragonality, as expected. This response to changing A1-site cation size has previously been reported in both filled²⁰ and unfilled⁴⁰ TTBs. While these trends appear to be consistent, they are comparatively subtle; PXRD patterns for each composition are otherwise visually indistinguishable with the exception of the aforementioned peak splitting.

In addition to the previously described splitting, $Ba_4La_{0.67}\Box_{1.33}Nb_{10}O_{30}$ displays several peaks which exhibit slightly different peak profiles. These peaks do not coincide with known impurity phases and possibly indicate splitting due to lowering of symmetry. However, it was not possible to determine this conclusively given the resolution of the laboratory PXRD data. Nevertheless, the pattern can still be indexed in a tetragonal setting.

SEM micrographs of the thermally etched fracture surfaces of the sintered pellets indicate elongated and relatively uniform sized crystallites in the range of 10–15 μm . Relatively few pores were observed which is consistent with calculated pellet densities (see ESI†).

B. Electrical properties

Polycrystalline ceramics consist of crystalline grains (grain bulk) connected by intergranular material (grain boundaries) and may include a level of porosity, depending on the compound and synthetic conditions. These various electroactive regions typically display differing electrical properties; standard capacitance measurements of polycrystalline samples measure the total, macroscopic sample response and are therefore a convolution of the contributions of these individual electroactive regions within the sample. The electrical properties of a ceramic are typically dominated by the grain bulk and grain boundary responses, however, other contributions such as those from electrodes and surface effects may be observed.^{41,42} The ‘brickwork’ model of discrete grains separated by thin grain boundaries (which therefore are characterised by high capacitances) is not universally observed in all cases as ceramic microstructure is heavily influenced by processing conditions.⁴³ Dissimilar electrical responses can therefore be observed in nominally identical compositions due to microstructural differences such as grain size, thickness/definition of grain boundary regions, and density.

High temperature immittance spectroscopy was carried out to determine the number of electroactive regions present and to ensure fixed frequency capacitance measurements carried out at lower temperatures represented the bulk response.

All samples display single semi-circular arcs in both impedance, Z^* , and modulus, M^* , complex plane plots, and single peaks in the Z'' and M'' spectroscopic plots, Fig. 3(a) and (b). The Z'' and M'' peaks are spectroscopically coincident, Fig. 3(b), and so belong to the same electroactive component. The magnitude of the capacitance values obtained from the



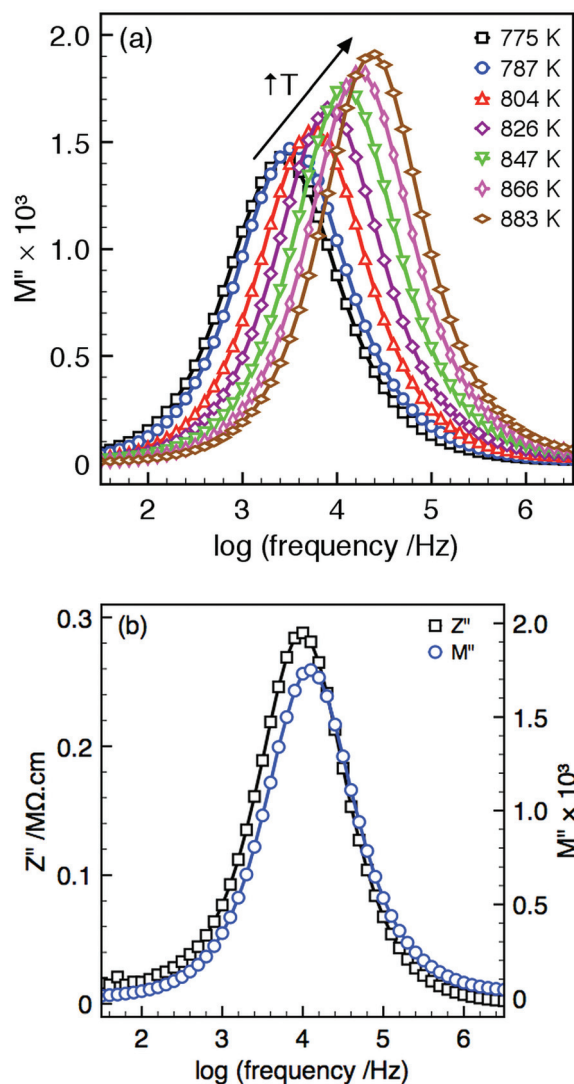


Fig. 3 High temperature impedance spectroscopy data for $\text{Ba}_4\text{Dy}_{0.67}\square_{1.33}\text{Nb}_{10}\text{O}_{30}$ showing: (a) M'' spectroscopic plot (775–883 K), and (b) combined M'' and Z'' spectroscopic plots (847 K) all showing the dominant bulk response.

complex plane plots allow this response to be attributed to the bulk region. Fixed frequency (1 MHz) capacitance measurements can therefore be used as an approximation of the bulk response at all temperatures presented in Fig. 4.

Fixed frequency capacitance sweeps were carried out for all compositions as detailed in the Experimental section. Compositions $R = \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}$ and Y display single anomalies in the relative permittivity within the temperature range in which data were collected (50–890 K), Fig. 4. Composition $R = \text{La}$ displays a low temperature frequency dependent relaxor-type peak with maximum, T_m (1 MHz), at 297 K and a frequency independent high temperature peak. This unusual behaviour will be discussed later. Peak maxima for all other compositions display little frequency dependence; T_C differs by less than 5 K over measured frequency range (10^2 – 10^7 Hz). Relaxation in the dielectric loss was observed to occur coincident to the permit-

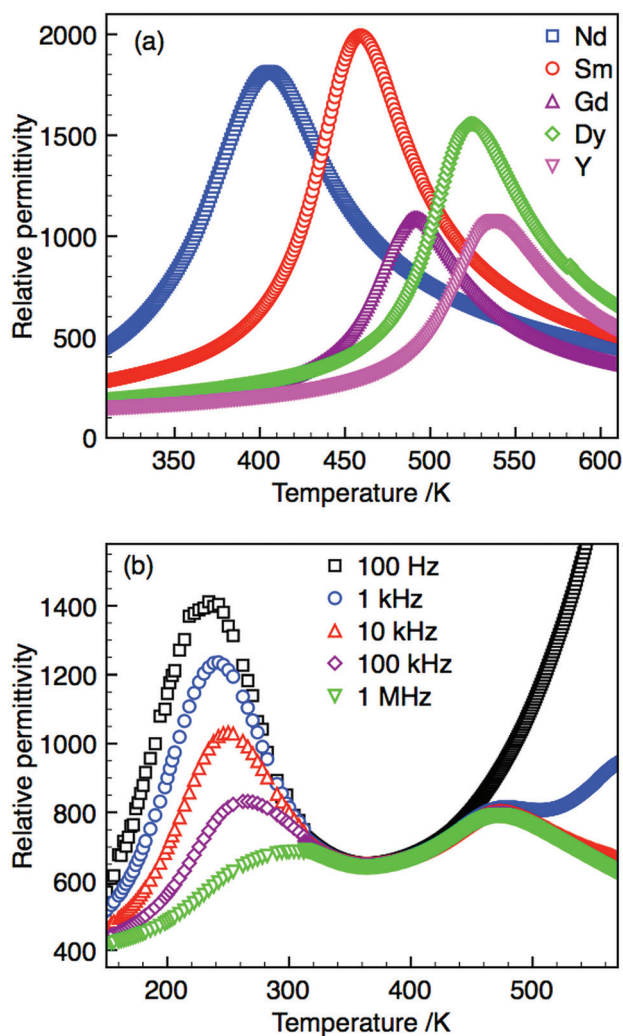


Fig. 4 (a) Relative permittivity (at 1 MHz) of $\text{Ba}_4\text{R}_{0.67}\square_{1.33}\text{Nb}_{10}\text{O}_{30}$ ($R = \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}$ and Y) and (b) $\text{Ba}_4\text{La}_{0.67}\square_{1.33}\text{Nb}_{10}\text{O}_{30}$ at selected frequencies.

tivity maxima; all losses ($\tan \delta$) were less than $\sim 10\%$. The diffuse nature of the permittivity peaks in comparison to many perovskite structured ferroelectrics is a common feature of ceramic TTB ferroelectrics and is likely due to the nature of the ferroelectric transition and the intrinsic anisotropy within the TTB structure.⁴ It should be noted that despite being multi-phase the Ho and Er samples were also ferroelectric with T_C of ca. 269 and 264 °C, respectively. The lattice parameters of the TTB phase in these samples also have tetragonality ratios which correlate with the observed T_C , however as the exact compositions of these phases are unknown and are not likely to contain the same stoichiometric ratios of $\text{Ba} : \text{R} : \text{vacancies}$ we prefer not to include them in the discussion presented here (see ESI† for data regarding $R = \text{Ho}$ and Er).

All samples display saturated P - E hysteresis loops with clear associated switching current, Fig. 5, demonstrating that all compositions are ferroelectric up to T_C (or up to the maximum temperature measured, 473 K). Linear lossy dielectric P - E



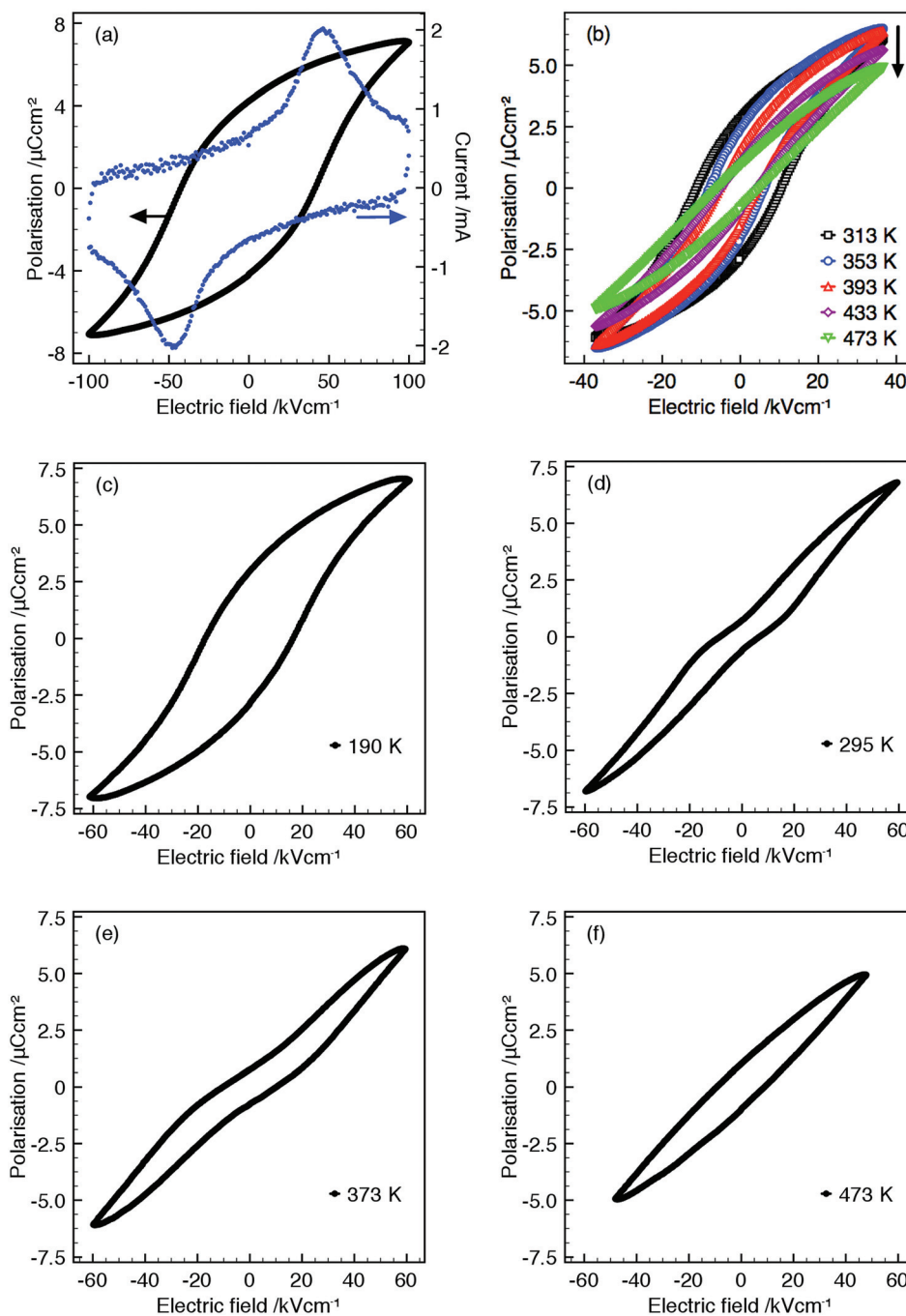


Fig. 5 (a) P - E hysteresis loop for Ba₄Dy_{0.67}□_{1.33}Nb₁₀O₃₀ (296 K) and (b) variable temperature P - E loops for Ba₄Nd_{0.67}□_{1.33}Nb₁₀O₃₀ (arrow indicates increasing temperature); (c)-(f) shows evolution of P - E behaviour for Ba₄La_{0.67}□_{1.33}Nb₁₀O₃₀ over the temperature range 190 to 473 K. All data were collected at 100 Hz.

loops above T_C for compositions R = La, Nd and Sm demonstrate the high temperature paraelectric phase, Fig. 5(b) and confirm that the peak maxima in the dielectric data corresponds to a polymorphic phase transition between ferroelectric and paraelectric states and justifies the assignment of the non-centrosymmetric $P4bm$ space group over the centrosymmetric $P4/mbm$ for ambient PXRD data.

Ba₄La_{0.67}□_{1.33}Nb₁₀O₃₀ displays two anomalies in the relative permittivity: a frequency dependent relaxor-type peak, $T_m =$

297 K (at 1 MHz) and a high temperature, frequency independent peak, $T_m = 471$ K, Fig. 4(b). Normal ferroelectric P - E hysteresis loops are observed at temperatures below the maxima in the relaxor peak and gradually transform into constricted or 'pinched' non-ferroelectric loops between the maxima of the dielectric anomalies with (slightly lossy) linear dielectric loops above 473 K, Fig. 5(c-f). Similar dielectric behaviour has been observed in other TTBs although varying rationalisations as to the origin of these observations have been proposed. These



include: onset of incommensurate structures⁴⁴ affected by octahedral rotation²² which has been suggested as being driven by decreasing smaller relative sizes of cations occupying A1- and A2-sites,²³ variation of dipole correlation lengths due to charge imbalance from mixed valence cations,⁴⁵ a frustrated ferroelectric/ferroelastic transition,⁴⁶ polar ordering disruption due to 5 coordinate B-cation.³⁴ P - E loops were not reported for these TTBs, however, the coexistence of multiple dielectric anomalies and associated pinched P - E loops has been reported in perovskite systems.^{47,48}

We are currently investigating the possible origins of these observations in our materials and these features will be discussed more fully in a future communication. Nevertheless, the observation that it is only possible to observe polarisation switching, *i.e.* ferroelectricity, below the lower temperature permittivity peak allows us to associate it with the ferroelectric T_C .

The Curie temperature, T_C , increases with decreasing ionic radius; the simultaneous increase in tetragonality (c/a) observed suggests a successively larger structural distortion in the TTB structure shifting T_C to higher temperatures (Fig. 6). This relationship between Curie temperature and tetragonal distortion appears to be common within other series of rare earth-doped barium niobate and barium titanium niobate TTBs. It also occurs when cation size in the A1-site is controlled by means of solid solution between two rare earth cations (*e.g.* in the series $(1-x)\text{Ba}_{3.75}\text{Sm}_{0.833}\text{Nb}_{10}\text{O}_{30}$ - $x\text{Ba}_{3.75}\text{Y}_{0.833}\text{Nb}_{10}\text{O}_{30}$ ³⁸). As in the compositional series presented here, the increased structural distortion generally originates in contraction of the ab plane, while the c -axis remains comparatively invariant; this is seen for both filled *e.g.*, $\text{Ba}_4\text{M}_2\text{-Ti}_4\text{Nb}_6\text{O}_{30}$ ($\text{M} = \text{La}, \text{Pr}, \text{Nd}, \text{Sm}, \text{Eu}, \text{Gd}, \text{Dy}, \text{and Bi}$)²⁰ and $\text{Sr}_4(\text{La}_{1-x}\text{Sm}_x)\text{Ti}_4\text{Nb}_6\text{O}_{30}$,⁴⁹ and unfilled TTBs *e.g.*, $(1-x)\text{Ba}_{3.75}\text{Sm}_{0.833}\text{Nb}_{10}\text{O}_{30}$ - $x\text{Ba}_{3.75}\text{Y}_{0.833}\text{Nb}_{10}\text{O}_{30}$ ³⁸ and $\text{Ba}_4\text{RE-TiNb}_9\text{O}_{30}$ ($\text{R} = \text{La}, \text{Pr}, \text{Nd}, \text{Sm}, \text{Eu}, \text{Gd}$).⁴⁰ Sun *et al.*⁵⁰ also

reported a number compositional series with barium and rare earth (La, Sm and Nd) on the A-sites as described by the general formulae: $\text{Ba}_5\text{RTi}_3\text{Nb}_7\text{O}_{30}$, $\text{Ba}_4\text{R}_2\text{Ti}_4\text{Nb}_6\text{O}_{30}$ and $\text{Ba}_3\text{R}_3\text{-Ti}_5\text{Nb}_5\text{O}_{30}$. In the latter series, there was significantly more variation in the c -axis, presumably due to the rare earth now also occupying the A2-site.

It has been suggested that structural distortions lead to an increase in the c lattice parameter generating greater atomic shifts, Δz , of the cation in the ferroelectrically active barycentre, raising T_C .⁵¹ The necessity to additionally consider distortion within the ab plane has been shown^{27,52} and the invariant nature of the c -axis within this series suggests that the tetragonal distortion, and thus increase in T_C , originates from contraction in the ab plane with decreasing RE ionic radius. The magnitude of the difference in the mean size between the cations occupying the A1- and A2-sites influences the onset of relaxor behaviour in filled TTBs studied by Chen and co-workers²³ with classical ferroelectric transitions occurring with larger size difference. Based on Rietveld refinements of PXRD data the barium solely occupies the larger A2-site,¹³ with R^{3+} confined to the A1-site in this series of compounds. The effect of the vacancy 'size' is difficult to quantify but is likely to decrease A1-site size relative to A2-site. Unfilled TTBs with A1-site vacancies are therefore likely to have a larger effective A1 : A2 size difference compared to filled TTBs. This is consistent with the observation that filled compounds with Ba^{2+} occupying the A2-site display relaxor behaviour with smaller cations such as Nd and Sm in the A1-site²³ compared to only the La analogue in the unfilled series reported here. The presence of vacancies at the A1 site therefore stabilises the ferroelectric distortion in this structure type.

Conclusions

TTB structured materials with varying A1-site cation size, $\text{Ba}_4\text{R}_{0.67}\square_{1.33}\text{Nb}_{10}\text{O}_{30}$ ($\text{R} = \text{La}, \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}, \text{Y}$) were produced obtaining nominally single phase ceramics. From lab-based XRD data, these compositions are metrically tetragonal and exhibit an increasing tetragonal distortion as R ionic radius decreases as indicated by refined lattice parameter values and increasing peak splitting of $00l$ $hk0$ reflections. Compounds $\text{R} = \text{Nd}, \text{Sm}, \text{Gd}, \text{Dy}, \text{and Y}$ display single peaks in the relative permittivity with Curie temperatures increasing with decreasing ionic radius and therefore, concurrently with increasing tetragonal distortion. A contraction in the ab plane drives the structural distortion, with the c -axis largely invariant. This empirical observation of a correlation between tetragonality and T_C appears to be universal within A-site (R) doped barium niobate-based TTB compositional series of comparable Ba : R stoichiometric ratio.

The ferroelectric nature of all compositions has been established with P - E hysteresis measurements. The presence of vacancies at the A1-site are important in inducing the ferroelectric instability in these materials, with ferroelectricity observed for Nd compositions compared to relaxor behaviour

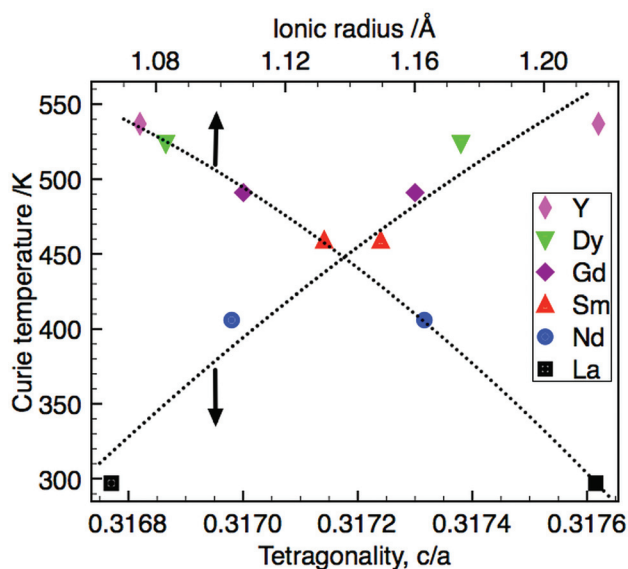


Fig. 6 Correlation of Curie temperature with tetragonality and ionic radius (the data point for La is T_m at 1 MHz).



in corresponding filled analogues. The amount of vacancies incorporated could therefore provide an additional degree of freedom for tuning of properties.

A low temperature relaxor-type peak with strong frequency dispersion and a non-frequency dependent high temperature peak were detected in the relative permittivity of $\text{Ba}_4\text{La}_{0.67}\text{Nb}_{10}\text{O}_{30}$. Low temperature P - E hysteresis loops, indicating ferroelectric behaviour, transform into a 'pinched' or antiferroelectric-type loop in the temperature range between the dielectric peaks, and finally to a linear dielectric-type P - E response at higher temperature. The precise origins of this behaviour are not yet known.

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References

- 1 A. S. Bhalla, R. Guo and R. Roy, *Mater. Res. Innovations*, 2000, **4**, 3.
- 2 A. M. Glazer, *Acta Crystallogr., Sect. B: Struct. Crystallogr. Cryst. Chem.*, 1972, **28**, 3384.
- 3 V. M. Goldschmidt, *Die Naturwissenschaften*, 1926, **4**, 477.
- 4 M. E. Lines and A. M. Glass, *Principles and Applications of Ferroelectrics and Related Materials*, Oxford University Press, Oxford, 2001.
- 5 M. H. Francombe and B. Lewis, *Acta Crystallogr.*, 1958, **11**, 696.
- 6 P. B. Jamieson, S. C. Abrahams and J. L. Bernstein, *J. Chem. Phys.*, 1969, **50**, 4352.
- 7 M. H. Francombe, *Acta Crystallogr.*, 1960, **13**, 131.
- 8 K. Masuno, *J. Phys. Soc. Jpn.*, 1964, **19**, 323.
- 9 T. Ikeda, T. Haraguchi, Y. Onodera and T. Saito, *Jpn. J. Appl. Phys.*, 1971, **10**, 987.
- 10 J. Thoret, W. Freundlich, J. Ravez, A. Simon and P. Hagenmuller, *Ferroelectrics*, 1983, **56**, 1061.
- 11 N. Wakiya, J. Wang, K. Shinozaki and N. Mizutani, *Korean J. Ceram.*, 2000, **6**, 380.
- 12 P. B. Jamieson, S. C. Abrahams and J. L. Bernstein, *J. Chem. Phys.*, 1968, **48**, 5048.
- 13 M. Josse, O. Bidault, F. Roulland, E. Castel, A. Simon, D. Michau, R. Von der Muhill, O. Nguyen and M. Magilone, *Solid State Sci.*, 2009, **9**, 1118.
- 14 Y.-Q. Tan, Y. Yu, Y.-M. Hao, S.-Y. Dong and Y.-W. Yang, *Mater. Res. Bull.*, 2013, **48**, 1934.
- 15 P. P. Liu, X. L. Zhu and X. M. Chen, *J. Appl. Phys.*, 2009, **106**, 074111.
- 16 Y. Yuan, X. M. Chen and Y. J. Wu, *J. Appl. Phys.*, 2005, **98**, 084110.
- 17 Y. B. Yao, C. L. Mak and B. Ploss, *J. Eur. Ceram. Soc.*, 2012, **32**, 4353.
- 18 M. Smirnov and P. Saint-Grégoire, *Acta Crystallogr., Sect. A: Fundam. Crystallogr.*, 2014, **70**, 283.
- 19 K. Yamauchi and S. Picozzi, *Phys. Rev. Lett.*, 2010, **105**, 107202.
- 20 M. C. Stennett, I. M. Reaney, G. C. Miles, D. I. Woodward, A. R. West, C. A. Kirk and I. Levin, *J. Appl. Phys.*, 2007, **101**, 104114.
- 21 M. C. Stennett, G. C. Miles, J. Sharman, I. M. Reaney and A. R. West, *J. Eur. Ceram. Soc.*, 2005, **25**, 2471.
- 22 I. Levin, M. C. Stennett, G. C. Miles, D. I. Woodward, A. R. West and I. M. Reaney, *Appl. Phys. Lett.*, 2006, **89**, 122908.
- 23 X. L. Zhu, S. Y. Wu and X. M. Chen, *Appl. Phys. Lett.*, 2007, **91**, 162906.
- 24 E. O. Chi, A. Gandini, K. M. Ok, L. Zhang and P. S. Halasyamani, *Chem. Mater.*, 2004, **16**, 3616.
- 25 A. Rotaru, D. C. Arnold, A. Daoud-Aladine and F. D. Morrison, *Phys. Rev. B: Condens. Matter*, 2011, **83**, 184302.
- 26 R. R. Neurgaonkar, J. G. Nelson and J. R. Oliver, *Mater. Res. Bull.*, 1992, **27**, 677.
- 27 D. C. Arnold and F. D. Morrison, *J. Mater. Chem.*, 2009, **19**, 6485.
- 28 N. Wakiya, J. Wang, A. Saiki, K. Shinozaki and N. Mizutani, *J. Eur. Ceram. Soc.*, 1999, **19**, 1071.
- 29 G. C. Miles, M. C. Stennett, I. M. Reaney and A. R. West, *J. Mater. Chem.*, 2005, **15**, 798.
- 30 A. C. Larson and R. B. V. Dreele, Los Alamos National Laboratory Report LAUR 86-748, 1994.
- 31 B. H. Toby, *J. Appl. Crystallogr.*, 2001, **34**, 210.
- 32 K. Lebbou, H. Itagaki, A. Yoshikawa, T. Fukuda, F. Carillo-Romo, G. Boulon, A. Brenier and M. T. Cohen-Adad, *J. Cryst. Growth*, 2000, **210**, 655.
- 33 B. Boudour, A. Karek, A. El Hassouni, C. Goutaudier and G. Boulon, *Afr. Rev. Phys.*, 2011, **6**, 181.
- 34 M. Prades, N. Maso, H. Beltran, E. Cordoncillo and A. R. West, *Inorg. Chem.*, 2013, **52**, 1729.
- 35 E. Castel, P. Verber, M. Albine, M. Velazquez, S. Pechev, D. Denux, J. P. Chaminade, M. Magilone and M. Josse, *J. Cryst. Growth*, 2012, **340**, 156.
- 36 M. X. Cao, X. L. Zhu, X. Q. Liu and X. M. Chen, *J. Am. Ceram. Soc.*, 2010, **93**, 782.
- 37 Y. J. Wu, Z. J. Hong, Y. Q. Lin, S. P. Gu, X. Q. Liu and X. M. Chen, *J. Appl. Phys.*, 2010, **108**, 014111.
- 38 J.-K. Wang, N. Wakiya, K. Shinozaki and N. Mizutani, *J. Ceram. Soc. Jpn.*, 2000, **108**, 36.
- 39 R. D. Shannon, *Acta Crystallogr., Sect. A: Cryst. Phys., Diffraction, Theor. Gen. Cryst.*, 1976, **32**, 751.
- 40 C. Hu, L. Hou, L. Fang and L. Liu, *J. Alloys Compd.*, 2013, **581**, 547.
- 41 A. R. West, D. C. Sinclair and N. Hirose, *J. Electroceram.*, 1997, **1**, 65.
- 42 D. C. Sinclair, *Bol. Soc. Esp. Ceram. Vidrio*, 1995, **34**, 55.
- 43 J. T. S. Irvine, D. C. Sinclair and A. R. West, *Adv. Mater.*, 1990, **2**, 132.
- 44 M. Venet, J. M'Peko, F. L. Zabetto, F. Guerrero, D. Garcia and J. A. Eiras, *Appl. Phys. Lett.*, 2009, **94**, 172901.



- 45 E. Castel, M. Josse, D. Michau and M. Magilone, *J. Phys.: Condens. Matter*, 2009, **21**, 452201.
- 46 A. Torres-Pardo, R. Jimenéz, J. M. González-Calbet and E. García-González, *Inorg. Chem.*, 2011, **50**, 12091.
- 47 Z. Xu, X. Dai, J. Li and D. Viehland, *Appl. Phys. Lett.*, 1996, **68**.
- 48 W. Jo, S. Schaab, E. Sapper, L. A. Schmitt, H.-J. Kleebe, A. J. Bell and J. Rödel, *J. Appl. Phys.*, 2011, **110**, 074106.
- 49 X. L. Zhu, Y. Bai, X. Q. Liu and X. M. Chen, *Appl. Phys.*, 2011, **110**, 114101.
- 50 Y. H. Sun, X. M. Chen and X. H. Zheng, *J. Appl. Phys.*, 2004, **96**, 7435.
- 51 S. C. Abrahams, S. K. Kurtz and P. B. Jamieson, *Phys. Rev.*, 1968, **172**, 551.
- 52 K. Li, X. L. Zhu, X. Q. Liu and X. M. Chen, *Appl. Phys. Lett.*, 2013, **102**, 112912.

