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Normal ferroelectric to relaxor behavior in laser ablated Ca-doped barium titanate thin films

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$\text{Ba}_{1-x}\text{Ca}_x\text{TiO}_3$ thin films ($x=0.05$ to 0.17) were deposited on Pt-coated Si substrates using a pulsed excimer laser ablation technique. X-ray diffraction and scanning electron microscope studies of the $\text{Ba}_{1-x}\text{Ca}_x\text{TiO}_3$ targets exhibit a polycrystalline nature and thin films also show the same but with a significant orientation along the (111) direction. Secondary ion mass spectrometer analysis reveals the presence of a sharper interface existing at the thin film substrate. The dielectric phase transition temperature of $(\text{Ba}_{1-x}\text{Ca}_x)\text{TiO}_3$ targets were sharp and the transition temperature was found to decrease from 140°C to 110°C with an increase in the values of x ($x>0.05$ at. %). The laser ablated Ca-doped BaTiO_3 thin films deposited at 100 mTorr exhibited a higher dielectric constant, lower dielectric loss, and an anomalous decrease in phase transition was observed. The anomalous phase transition decrease was ascribed to the occupancy of the Ca^{2+} in the Ti^{4+} site. There was a cross over from the sharp to diffused phase transition for a higher composition of Ca (>9 at. %) in BaTiO_3 thin films. The diffuse transition behavior might be due to the larger number of the Ca^{2+} ions occupying the Ti^{4+} site, eventually introducing larger compositional and structural disorder and this occupancy leads to the generation of oxygen vacancies. The activation energy obtained from impedance spectroscopy was 1.05 eV, and was attributed to the oxygen vacancy motion. © 2003 American Institute of Physics. [DOI: 10.1063/1.1618914]

I. INTRODUCTION

Ferroelectric materials are gaining importance in technology especially in the areas of the memory storage devices,^{1,2} microelectromechanical systems,³ multilayer ceramic capacitors,⁴ and recently in the areas of optoelectronic devices.⁵ Specifically, barium titanate and isovalent substituted BaTiO_3 are the promising candidates for potential use in the applications of microwave communications and optoelectronic devices. Barium calcium titanate (BCT) crystals show promising applications in advanced laser systems, optical interconnects, and optical storage devices.⁶ Calcium acts as a reduction inhibitor in BaTiO_3 and reduces the possibility of formation of the unwanted hexagonal phase. $\text{Ba}_{(1-x)}\text{Ca}_x\text{TiO}_3$ ceramics ($x=0$ to 22 at. %) prepared via a conventional solid-state reaction technique was observed to be ferroelectric at room temperature with point symmetry of 4 mm.^{7,8} The dielectric phase transition temperature (~ 140 – 120°C) from the tetragonal to cubic phase was found to marginally affect the substitution of the Ca^{2+} in Ba^{2+} site of the $\text{Ba}_{1-x}\text{Ca}_x\text{TiO}_3$ ceramics. In the case of Ca^{2+} occupying the Ti^{4+} site, the transition temperature was found to decrease drastically to 320 K for $\text{BaCa}_x\text{Ti}_{(1-x)}\text{O}_{(3-x)}$ ($x=0.15$) ceramics.⁹ There are various routes adopted to prepare the BCT ceramics and, interestingly, the BCT is strongly processing dependent especially on the occupancy of Ca^{2+} ions in the Ti^{4+} site of the BaTiO_3 . There are questions yet to be clearly answered on the possibility of Ca^{2+} occupying the Ti^{4+} site because of their differences in the

ionic radii and valences, in spite of there being sufficient experimental evidence in literature based on Raman spectroscopy,¹⁰ neutron diffraction,¹¹ and conductivity⁹ studies. Based on the existing reports in BCT ceramics, the occupancy of Ca^{2+} in the Ti^{4+} site gives rise to doubly charged acceptor centers Ca_{Ti}'' .

Han *et al.*,¹² Zhang *et al.*,¹³ and Hennings *et al.*,¹⁴ suggested that there is a probability of Ca^{2+} occupying the Ti^{4+} site in the BaTiO_3 lattice, provided the atomic concentration ratio of $(\text{Ba}+\text{Ca})/\text{Ti}>1$. This ratio leads to excess barium, thereby forcing Ca^{2+} to occupy the Ti^{4+} site and eventually leading to a drastic decrease in the dielectric phase transition temperature. Zhuang *et al.*¹⁵ proposed that the occupancy of Ca^{2+} in the Ti^{4+} site will create a compressional stress and eventually leads to the distortion of the lattice in BaTiO_3 .

There exist very few reports on BCT thin films in open literature and BCT is considered to be one of the foremost potential candidates for the electro-optic modulators and memory devices. Hence, in this article, we report on the growth of Ca-doped barium titanate thin films using a pulsed laser ablation technique and analyzed its structural, dielectric, and dielectric phase transitions in detail.

II. EXPERIMENT

$\text{Ba}_{(1-x)}\text{Ca}_x\text{TiO}_3$ targets [$x=0.05$ (denoted as BCT5C), 0.08, 0.10 (BCT12C), 0.13 and 0.15] were prepared via conventional solid-state reaction technique. High-purity starting powders of BaCO_3 , TiO_2 , and CaCO_3 were weighed in stoichiometric proportion, ball milled for 10 h, dried, and calcined at 1100°C for 3 h. The calcined powders were pressed at 100 kN/cm^2 into a 15 mm diameter disk-shaped pellet and

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the pellets were sintered at 1400 °C for 4 h. The densities of the targets were 96% and the targets were used for the laser ablation. The BCT thin films were deposited on the Pt(111)-coated Si substrate using a KrF excimer laser ablation technique (Lambda Physik, Germany, 248 nm). The substrate temperature during the deposition was maintained at 650 °C, with the laser fluence maintained at 3 J/cm² and substrate–target distance at 3.5 cm. The base pressure were initially brought down to 1 × 10^{−6} Torr and prior to the deposition, high-purity oxygen gas was introduced into the chamber and the deposition were carried at 100 mTorr for all the compositions of Ba_{1−x}Ca_xTiO₃ targets. The *in situ* grown thin films and targets were structurally characterized by x-ray diffraction [(XRD) Scintag XDS 2000]. The composition and distribution of the elements in thin films across the thickness were identified using energy dispersive analysis (Oxford Instruments) and secondary ion mass spectroscopy (SIMS), respectively. The surface morphology and cross section were analyzed using scanning electron microscope (SEM). Circular gold dots of 500 μm diameter were deposited by thermal evaporation for the top electrodes (Au/BCT/Pt configuration). The electrode annealing was carried out at 325 °C for 15 min and the metal–ferroelectric–metal (MFM) capacitor was used for the electrical characterization. Polarization hysteresis was analyzed using a RT-66A ferroelectric test system (RT 66A, test pulse width of 8 ms). Dielectric and capacitance–voltage (*C–V*) measurement was measured using a programmable Keithley LCZ 3330 meter (frequencies ranging from 0.1 to 100 kHz at an oscillating amplitude of 100 mV) coupled with a Keithley 230 voltage source. All of the measurements on MFM capacitors were carried out at a temperature ranging from 200 K to 550 K.

III. RESULTS

A. Structural determination

Figure 1 shows the XRD of the Ba_(1−x)Ca_xTiO₃ targets (denoted as BCT) and these patterns show a polycrystalline nature with distinct tetragonal splitting at (002/200). The distinct tetragonal splitting is observed for the lower concentration of *x* in the Ba_{1−x}Ca_xTiO₃ targets and the tetragonal splitting merges into a single peak for *x* > 0.21 eventually leading to the cubic phase at room temperature. There were additional peaks observed for the higher composition of Ca (*x* > 0.13) in Ba_{1−x}Ca_xTiO₃ targets, showing the evolution of secondary phases (Ca-rich BaTiO₃ phase and is denoted as # in the Fig. 1) and these peaks grow stronger with higher Ca content. Figure 2 shows the calculated *c/a* ratio plotted as a function of the *x* in the Ba_{1−x}Ca_xTiO₃ targets. The *c/a* ratio was observed to decrease with an increase in the incorporation of the Ca dopant (Ca > 5 at. %) in the lattice site of the BaTiO₃ targets. The lattice constant in the unit cell is observed to shrink with the increase in Ca²⁺ ions at the Ba²⁺ site, due to the smaller ionic radii of Ca²⁺ over Ba²⁺. It appears that the replacement of a²⁺ with smaller Ca²⁺ “tightens” the structure making it difficult to remove the oxygen from the lattice and renders it more favorable to fill the extrinsic oxygen vacancies. There is an increase in the

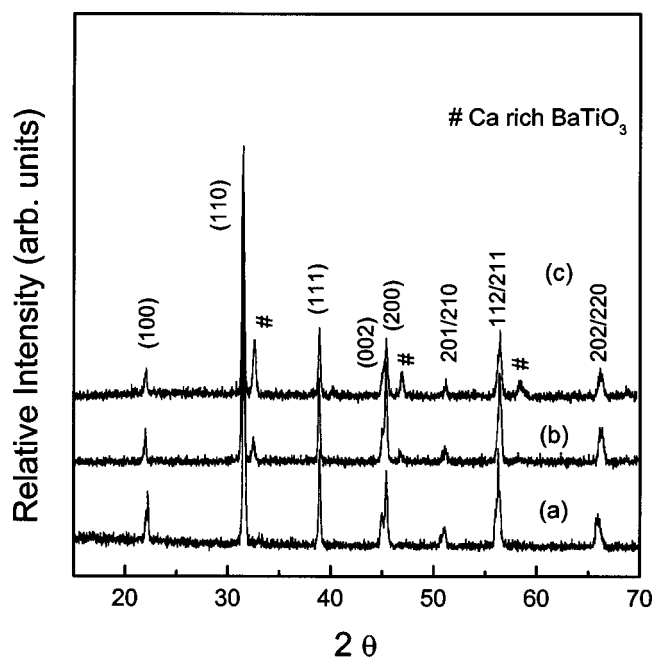


FIG. 1. The XRD pattern of the Ba_(1−x)Ca_xTiO₃ targets: (a) Ba_{0.95}Ca_{0.05}TiO₃, (b) Ba_{0.88}Ca_{0.12}TiO₃, and (c) Ba_{0.85}Ca_{0.15}TiO₃.

broadening of the diffracted peaks and a significant decrease in the grain size observed with increasing the Ba_{1−x}Ca_xTiO₃ targets.

The XRD of the BCT thin films deposited on a Pt(111)-coated Si substrate is shown in Fig. 3. All of the thin films exhibited the (111) orientation with little orientation along (110), and the orientations of the thin films can be tailored accordingly to the processing conditions. Kim *et al.*¹⁶ reported on the tailoring of the orientation of the laser ablated BaTiO₃ thin films on Si substrates, based on the growth pressure. The evolution of the secondary phases was observed for

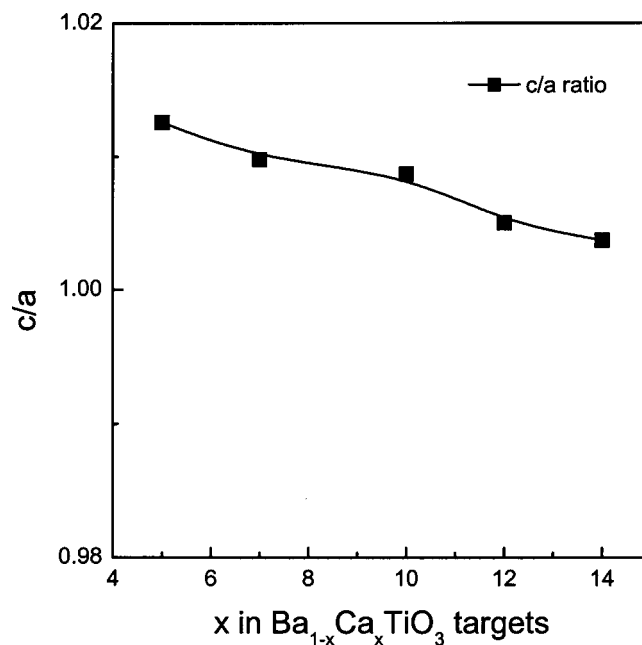


FIG. 2. The calculated *c/a* ratio as a function of *x* in Ba_{1−x}Ca_xTiO₃ targets.

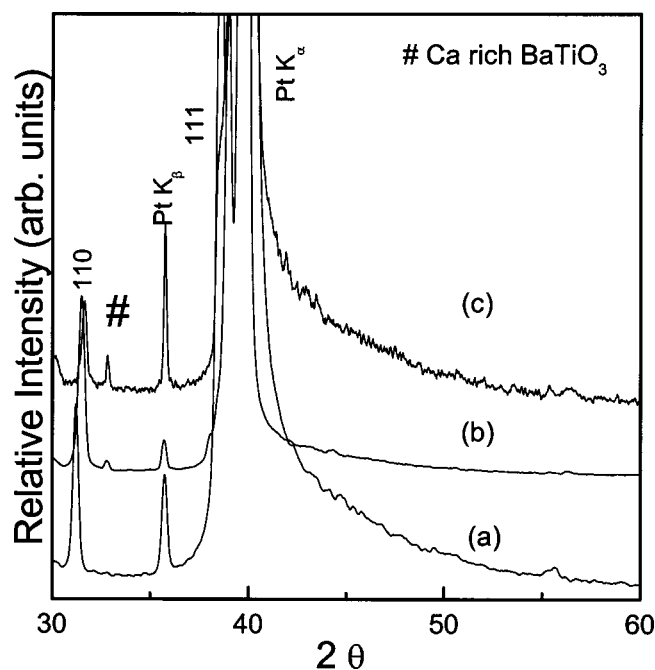


FIG. 3. The XRD of the BCT thin films deposited on Pt(111)-coated Si substrate: (a) 5 at. % Ca-doped BaTiO₃, (b) 12 at. % Ca-doped BaTiO₃, and (c) 15 at. % Ca-doped BaTiO₃ thin films.

higher Ca concentrations and is in accordance with the XRD pattern of BCT targets. Figures 4(a) and 4(b) shows the SEM micrographs of BCT5C and BCT12C thin films grown at 100 mTorr with $T_{\text{sub}} = 700^\circ\text{C}$. The surface morphology of micrographs shows significant faceted growth orientation along (111), with little orientation along (110), for the present given deposition conditions and is evident from our XRD pattern (Fig. 3). The cross-sectional view of the BCT12C thin films is shown in Fig. 4(c), and display mostly a columnar structure with little presence of the other planes, and the grains size varies from 45 to 80 nm exhibiting well defined densely packed grains. The three-dimensional tetrahedron-like structures are observed in the deposited BCT thin films as shown in Figs. 4(a) and 4(b) and a detailed study has not been carried out. The thickness of the thin films varied from 500 to 800 nm as measured by thin-film thickness monitor system (Filmetrics) and stylus method (Dektek). Figure 5 shows the SIMS profile of BCT12C thin films showing that the distribution of the elements is found to be uniform across the film and the interface at the film—Pt is sharper without any interdiffusion. These results establish that the grown thin films were of excellent quality.

B. Dielectric studies

The dielectric constant (ϵ') was observed to decrease with an increase in the Ca concentration of Ba_{1-x}Ca_xTiO₃ ceramics and, likewise, the phase transition temperature also correspondingly decreased. The dielectric constant (ϵ') and $\tan \delta$ dependence on the temperature for the ferroelectric phase from the tetragonal to cubic phase at varied measuring frequencies on different BCT targets are shown in Figs. 6(a) and 6(b). The $\tan \delta$ is also found to peak near the phase transition temperature as observed in the $\epsilon' - T$ and exhibit a

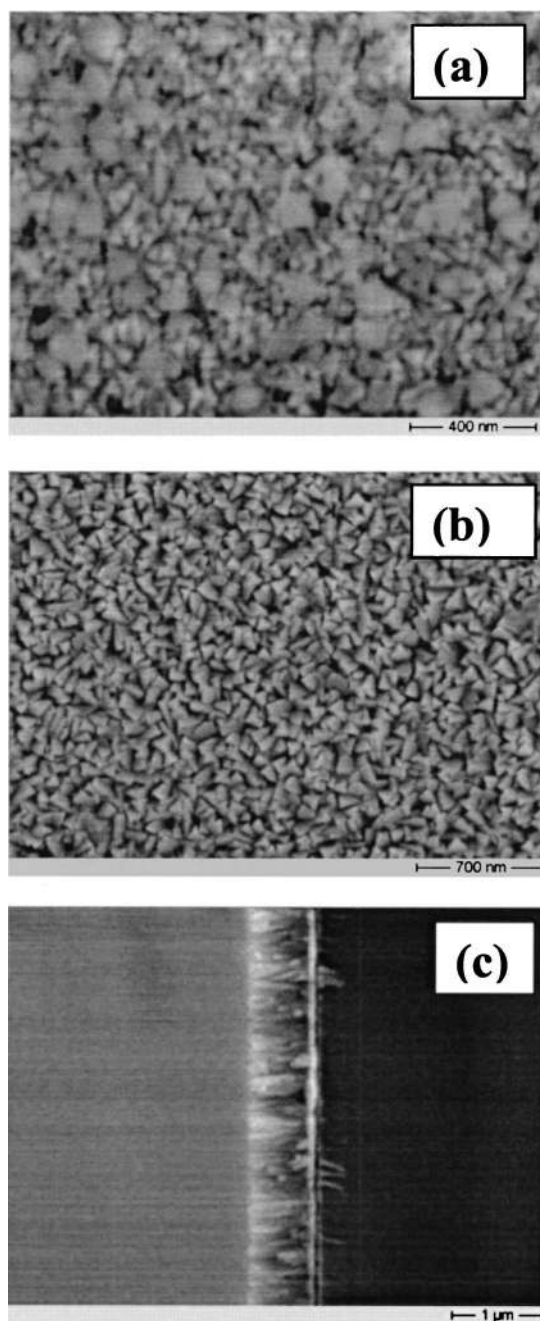


FIG. 4. (a) and (b) show the SEM micrographs of BCT5C and BCT12C thin films, respectively, grown at 100 mTorr with $T_{\text{sub}} = 700^\circ\text{C}$. (c) The cross-sectional view of the BCT12C thin films.

sharper phase transition, even for higher Ca²⁺ ions substituting in the Ba²⁺ site of BCT targets. This observed behavior is in accordance with the Kramers–Kronig relationship. Mitsui and Westphal¹⁷ also reported the existence of the sharper ferroelectric to paraelectric phase transition temperature for Ba_{1-x}Ca_xTiO₃ ceramics ($x < 20$ at. %). Pandey *et al.*¹⁸ observed a diffuse phase transition for higher calcium concentration in the barium titanate ceramics and attributed it to the purity of processing condition of ceramics and observed to be highly route dependent. The dielectric constant and $\tan \delta$ dependence on the temperature of BCT5C and BCT12C thin films are shown in Figs. 7(a) and 7(b). The phase transition was sharper for BCT5C thin films ($T_c \sim 178$ K and ~ 272 K

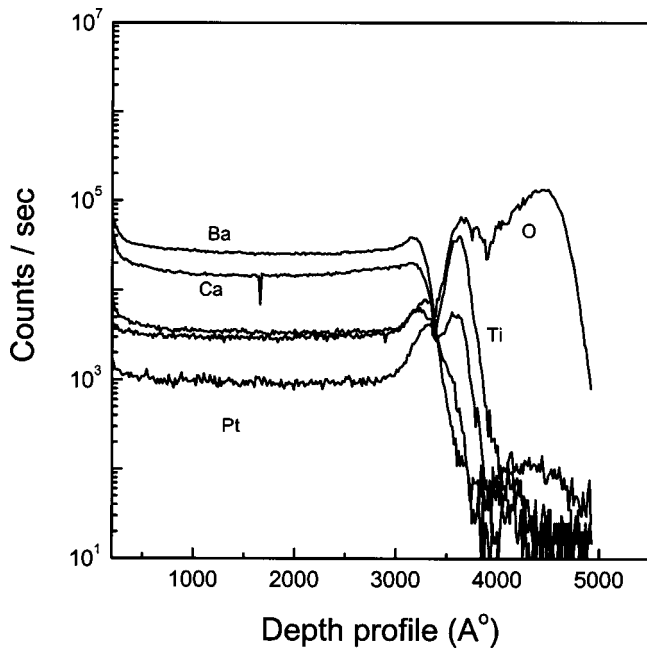


FIG. 5. The SIMS profile of BCT12C thin films showing the distribution of the elements to be uniform across the film and the interface of film–Pt/Si substrate.

at 1 kHz), while a diffuse phase transition behavior was observed for the BCT12C thin films ($T_c \sim 268$ at 1 kHz). A similar diffuse phase transition behavior trend was also observed for higher Ca content ($Ca > 9$ at. %) in BaTiO₃ thin films. In all laser ablated BCT thin films, there was an anomalous decrease in the phase transition temperature as opposed to the observed bulk phase transition temperature. In the case of the BCT12C thin films, the dielectric constant showed a broad maximum and the T_c peak in $\tan \delta$ was significantly lower than observed in ϵ' as shown in Fig. 7(b). The phase transition behavior in BCT12C thin films exhibited the signature of the typical diffuse phase transition behavior, i.e., relaxor-type behavior. In general, the decrease in the phase transition temperature is attributed to several reasons, such as intergranular stresses, ferroelectric domain sizes,¹⁹ interface between the substrate thin film, possibility of Ca²⁺ occupancy in the Ti⁴⁺ site, and compositional inhomogeneity. In our case, the grain size effects have been ignored as the grain size in our grown thin films was quite large (~45 nm to 80 nm) and the only possibility for the decrease in the T_c is the occupancy of Ca²⁺ in the Ti⁴⁺ site and is dealt with in detail in Sec. IV. The $\tan \delta$ versus temperature plot is shown in Fig. 7(a) and it shows two distinct phase transition peaks observed in BCT5C thin films. The lower phase transition temperature ($T_c \sim 178$ K) can be due either to the transition from the rhombohedral to tetragonal phase directly unlike the BaTiO₃ in the same temperature regime, has the phase transition from the rhombohedral to orthorhombic phase or the lower phase transition can also be from the rhombohedral to orthorhombic phase and the second phase transition to follow can be the coexistence of the tetragonal and orthorhombic phases. Beyond the second phase transitions, there is no phase transition behavior been observed. These investigations are to be carried out in detail

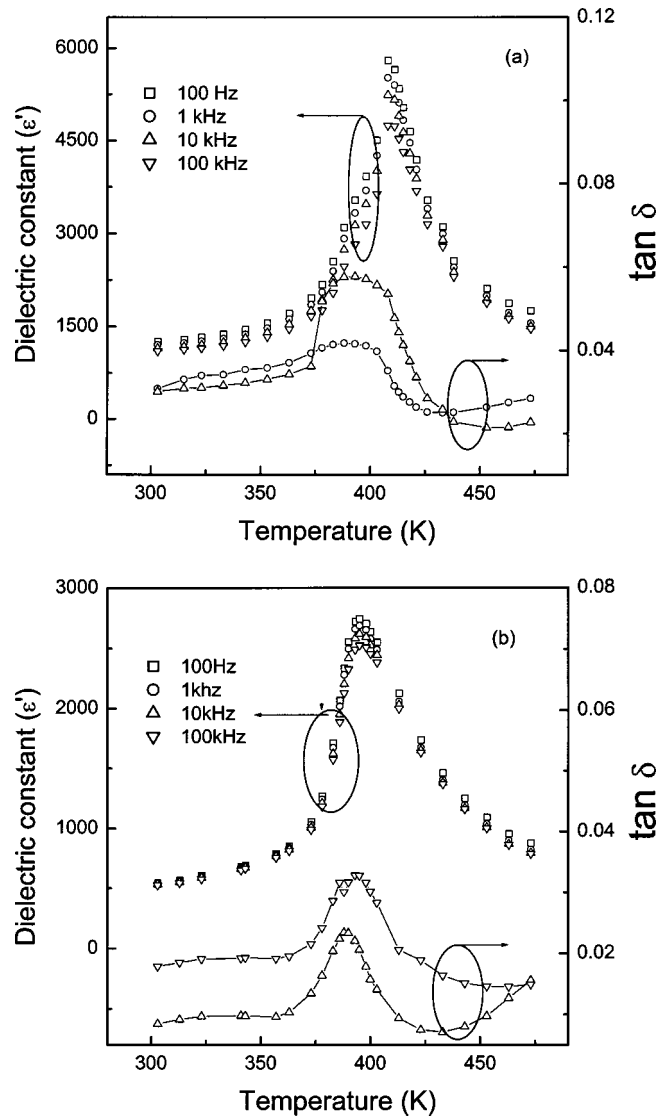


FIG. 6. (a) and (b) The dielectric constant and the $\tan \delta$ vs temperature of Ba_{0.95}Ca_{0.05}TiO₃ and Ba_{0.90}Ca_{0.10}TiO₃ targets, respectively.

and are not discussed in this article. The lower phase transition temperature for BCT12C thin films is not clearly seen and its diffuse phase transition behavior is relaxor type. The dielectric response of the relaxor ferroelectric is modeled as a response of an ensemble of the relaxors with a spectrum of the relaxation times. These are observed irrespective of the physical nature of these relaxators, exhibiting a broad maximum in the temperature dependence of ϵ' , with the relaxation dispersion around and below T_m . This type of behavior is attributed to the temperature evolution of this spectrum and the strong expansion of the spectrum occurs at a certain temperature T_f . All of the $\epsilon'(T)$ curves measured at various frequencies namely show the reduction of the ϵ_{max} and shift of the corresponding temperature T_m with an increase in frequency. This observed behavior are the distinct features of the relaxors. The accountability of DPT behavior in the BCT12C thin films is dealt in detail as follows.

Assuming the distribution of the local transition temperatures of microregions in a diffuse phase transition material, with microscopic compositional fluctuations and on as-

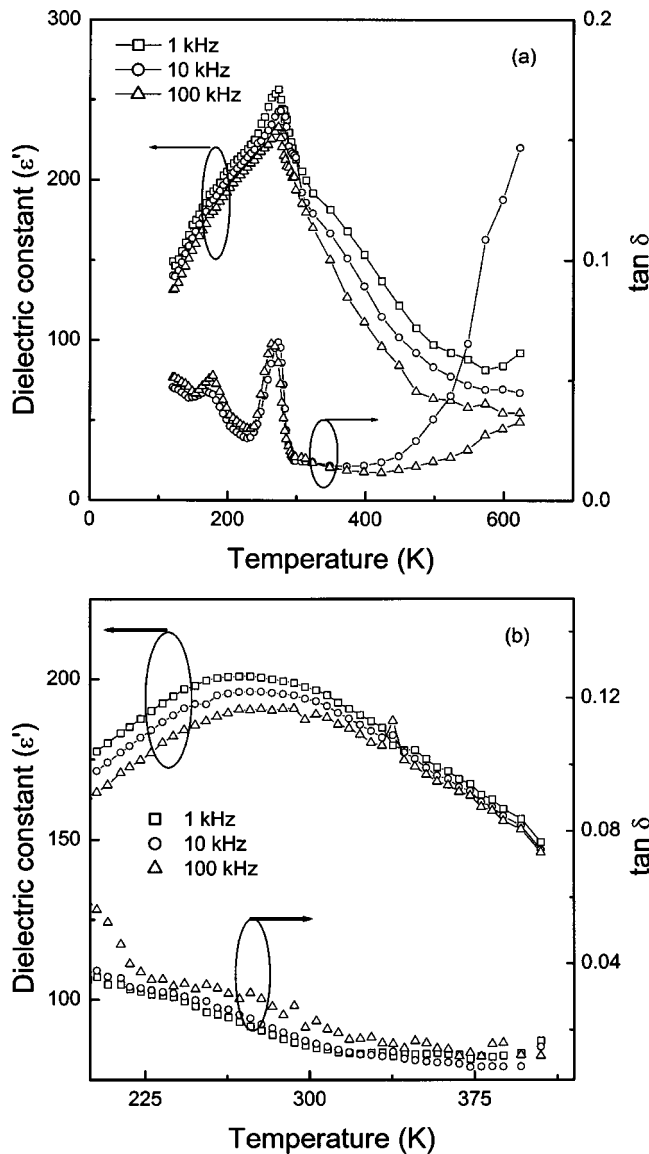


FIG. 7. (a) and (b): The dielectric constant and the $\tan \delta$ dependence on the temperature of the BCT5C and BCT12C thin films.

suming it to be Gaussian around the mean Curie point T'_m , Kirillov and Isupov²⁰ have shown that the

$$\left(\frac{1}{\epsilon' - \epsilon_\infty}\right) = \left(\frac{1}{\epsilon'_m}\right) \exp((T - T'_m)2\delta^2) \quad (1)$$

with 2δ is the width of the Gaussian and determines the extent of diffuseness of the phase transition. If $\epsilon' \gg \epsilon_\infty$, then, in series expansion, higher than second order can be neglected yielding $\gamma=2$. The limiting values $\gamma=1$ and $\gamma=2$ reduce expression (1) to the Curie–Weiss law valid for the case of a normal ferroelectric and the quadratic dependence (relaxor), respectively. The quadratic dependence of $1/\epsilon'$ on temperature has been claimed to be obeyed by several materials with diffuse phase transition behavior. The γ value was obtained from the least-squares fit of $\ln((1/\epsilon) - (1/\epsilon'_m))$ versus $\ln(T - T'_m)$ plots and is shown in Fig. 8(a) and the calcu-

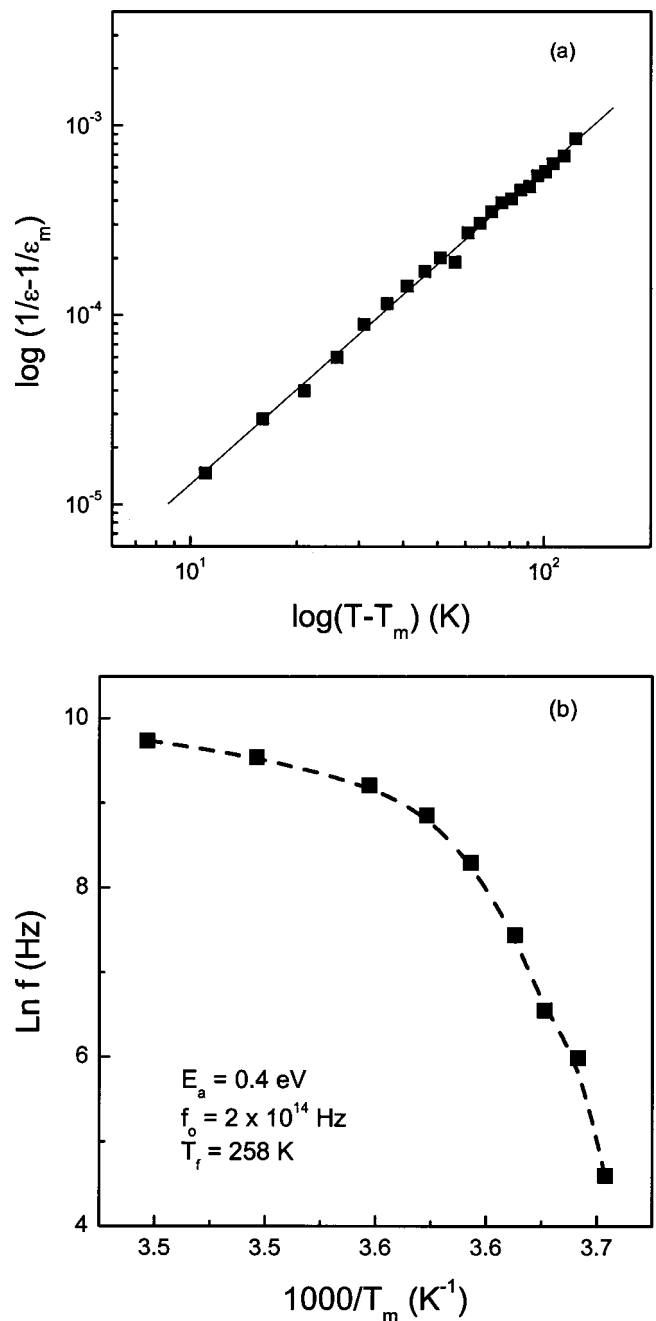


FIG. 8. (a) The plot of $\ln((1/\epsilon) - (1/\epsilon'_m))$ vs $\ln(T - T'_m)$ to determine γ value for the BCT12C thin films. (b) Frequency dependence of T_m for BCT12C thin films. The symbols and solid line indicate data points and fit to Vogel–Fulcher relationship, respectively.

lated γ value is 1.78 for the BCT12C thin films. The diffuseness of the phase transition can be described by an empirical parameter^{21–23}

$$\Delta T_{\text{diff}} = T_{0.9\epsilon}(100 \text{ Hz}) - T_{\epsilon m}(100 \text{ Hz}), \quad (2)$$

i.e., the difference between $T_{0.9\epsilon}(100 \text{ Hz})$ (the temperature corresponding to 90% of the maximum dielectric constant (ϵ_m) in the high-temperature side) and $T_{\epsilon m}$ (maximum ϵ' at a temperature for 100 Hz). The calculated diffuseness ΔT_{diff} for the BCT12C thin films is 70 K. On the other hand, the degree of relaxation behavior can be described by a parameter

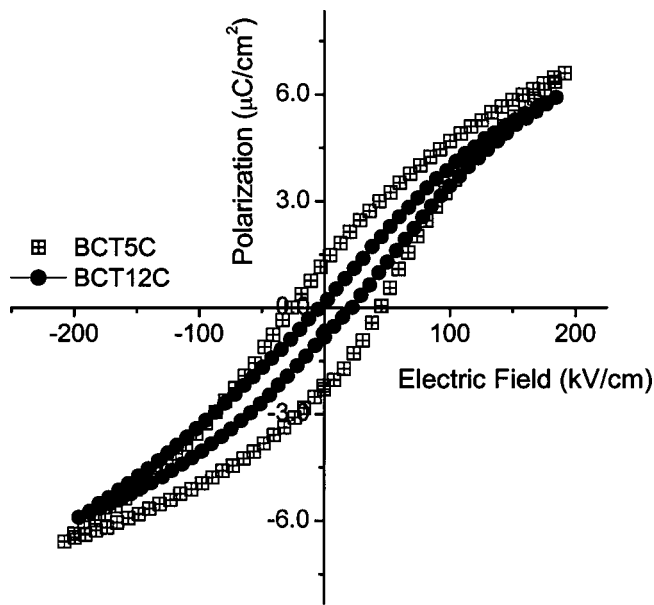


FIG. 9. The P - E field curves of BCT5C and BCT12C thin films measured at 260 K.

$$\Delta T_{\text{relax}} = T_{\epsilon_m}(100 \text{ kHz}) - T_{\epsilon_m}(100 \text{ Hz}), \quad (3)$$

with $T_{\epsilon_m}(100 \text{ kHz})$ and $T_{\epsilon_m}(100 \text{ Hz})$ denotes the temperature at which maximum ϵ' was measured at 100 kHz and 100 Hz, respectively. The degree of the relaxation behavior measured in BCT12C thin films was 10 K.

The above empirical characterizations with the Curie-Weiss law and the estimated parameters (ΔT_{relax} , ΔT_{diff} , and γ) show that the ϵ of the BCT targets and BCT5C thin films follows the normal ferroelectric Curie-Weiss law. In the case of BCT12C thin films, the phase transition behavior at the ϵ'_{max} , calculated parameters ΔT_{diff} and ΔT_{relax} show significant diffuseness of the phase transition and frequency dispersion. The γ value is close to 2 for BCT12C thin films, implying that these thin films exhibit a typical relaxor behavior.

The frequency dependence of T_m is shown in Fig. 8(b) in the form of $1/T_m$ versus $\ln f$. The observed frequency dependence of T_m was empirically evaluated using the Vogel-Fulcher relationship, adopted from the theory of magnetic relaxation in spin-glass systems²⁴ and is given as

$$f = f_o \exp\left(-\frac{E_a}{k(T_m - T_f)}\right), \quad (4)$$

where E_a is the activation energy, T_f is the freezing temperature of polarization fluctuations, and f_o is the pre-exponential factor. The values of E_a , T_f , and f_o obtained for our BCT12C thin films were 0.4 eV, 257.8 K, and 2×10^{14} Hz, respectively.

C. Polarization and capacitance-voltage measurements

Figure 9 establishes the signature of ferroelectricity in both the BCT5C and BCT12C thin films, obtained from the polarization-electric field (P - E) curves measured at 260 K. The spontaneous polarization of BCT5C and BCT12C thin

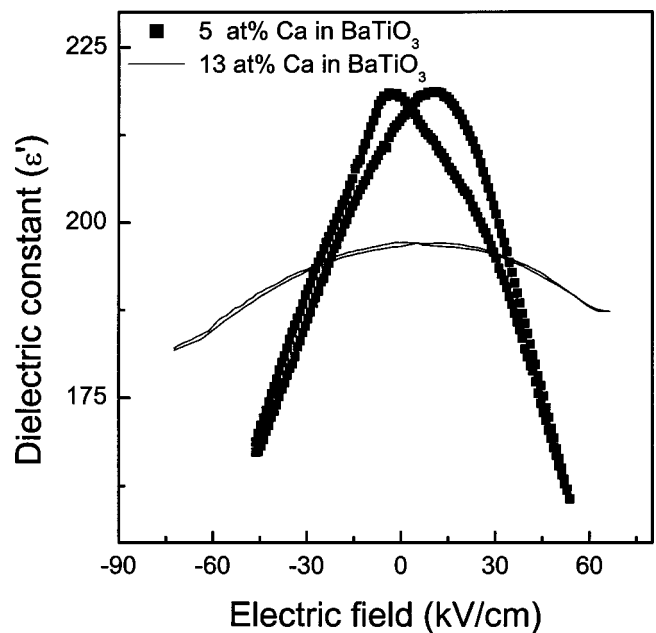


FIG. 10. The C - V measurements of BCT thin films in MFM configuration measured at 260 K.

films was found to be 6.8 and 5.8 $\mu\text{C}/\text{cm}^2$, respectively, at 0.2 MV/cm. The spontaneous polarization was found to decrease with the increase in the Ca concentration in BaTiO_3 thin films. The C - V measurements of BCT thin films are shown in the Fig. 10. A typical butterfly-shaped loop had been traced as seen in the C - V curve for MFM configuration measured at 260 K and is consistent with that observed for ferroelectric thin films. The C - V curve displays more diffusiveness for the BCT12C thin films signifying the presence of either compositional inhomogeneity or strains between the films and substrate. The asymmetric loop in the C - V curve may also be ascribed to the different types of electrodes (Au/BCT/Pt) present at the both ends. The C - V curve measured at 310 K exhibited a linear paraelectric behavior.

D. Impedance spectroscopy

The impedance spectroscopy plot provides insight into the types of carriers contributing to the conduction mechanisms. Figure 11 shows the impedance spectroscopy plot of BCT12C thin films and the inset shows the Arrhenius plot of resonant frequency versus $1000/T$ obtained from the impedance plot of BCT12C thin films. The calculated activation energy from the Arrhenius plot was 1.05 eV and is attributed to the motion of the oxygen vacancies,²⁵ which could have been created due to the substitution of Ca^{2+} ions in the Ti^{4+} site.²⁶

IV. DISCUSSION

The possibility of an anomalous drastic decrease in the phase transition temperature of the BCT thin films has been attributed to the possibility of forcing Ca^{2+} ions to occupy the Ti^{4+} site, thereby introducing compressional stress in the lattice.¹² The laser ablated BCT thin films exhibited a drastic

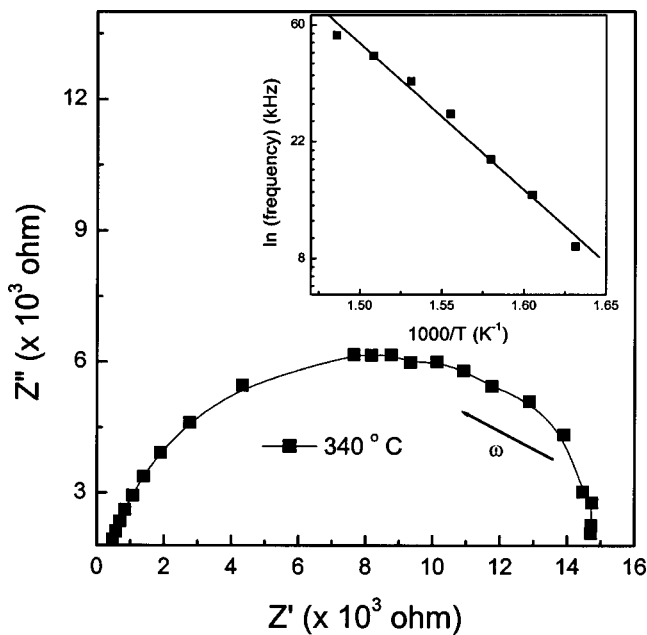
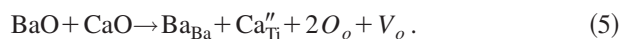


FIG. 11. Impedance spectroscopy plot of BCT12C thin films and the inset shows the Arrhenius plot of f_r vs $1000/T$.

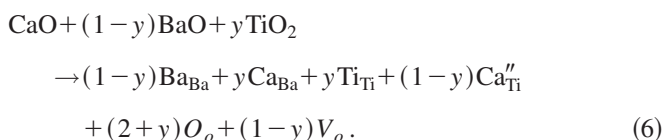
decrease in the phase transition temperature to 285 K for BCT5C and, for higher Ca^{2+} ions ($x > 0.09$) occupying the Ti^{4+} site, results in a still lower T_c and a broad diffused-type behavior. The diffuse phase transition was broad and appeared to be a relaxor type, which could have arose from either to the coexistence of the two phases in thermal equilibrium or the local compositional fluctuations arising from the excess Ca^{2+} ions occupying the Ti^{4+} site. There are earlier reports on the decrease of the dielectric phase transition temperature to 293 K for $\text{Ba}_{0.95}\text{Ti}_{1-x}\text{Ca}_{0.05+x}\text{O}_{3+x}$ ($x = 0.05$) ceramics and 320 K for the $\text{BaTi}_{1-y}\text{Ca}_y\text{O}_{3-y}$ ($y = 0.05$)²⁷ ceramics. The occupancy of Ca^{2+} ions either in the Ba^{2+} or Ti^{4+} site of BaTiO_3 , results in the following.^{9,10}

- (1) The simple substitution of Ca^{2+} with Ba^{2+} does not generate a charged defect; i.e., $\text{Ba}_{1-x}\text{Ca}_x\text{TiO}_3$.
- (2) Hans *et al.*¹⁰ and Zhang *et al.*¹¹ suggested that in case Ca^{2+} substitutes for Ti^{4+} in the BaTiO_3 lattice, it will act as an acceptor to trap the unlocalized electrons. It gives rise to a doubly charged acceptor center Ca_{Ti}'' , which compensates for the oxygen vacancies formed [Eq. (5)], during sintering in a reducing atmosphere and results in the formation of high resistivity dielectrics; i.e., $\text{BaTi}_{1-x}\text{Ca}_x\text{O}_{3-x}$,



The possibility of the Ca in interstitials has been excluded as these defects are unlikely.

An alternative general equation for the incorporation reactions is summarized as follows



The limiting cases for $y=1$ and $y=0$ correspond to the incorporation solely on the Ba or Ti sites, respectively. In the case of Ca^{2+} substituting for Ti^{4+} ions in the BaTiO_3 thin films, it results in a compositional and structural disorder due to the symmetry breaking. This is brought about by the difference in valences ($4+$ versus $2+$) and ionic radii (0.69 versus 0.99 \AA) which is believed to be responsible for the relaxor behavior and result in the drastic decrease in T_c behavior. The cross over from the ferroelectric to diffuse phase transition arises when the higher composition of the Ca content enters the B site (Ti^{4+}) disrupting the dipoles, creating the higher degree of disorder, and eventually inducing relaxor behavior. Such behavior introduces structural randomness, lattice defects, lattice strains, and altered forces. The symmetry breaking occurs at the nanometer scale leading to the formation of polar nanodomains and increases in size upon cooling, but never become large enough to precipitate a long-range-ordered ferroelectric state.²⁸ The activation energy obtained from the impedance spectroscopy plot is 1.05 eV and is attributed to the oxygen vacancy motion, which could have been created due to Ca occupying the Ti site as shown in Eq. (5). This also further confirms that the oxygen vacancy is present in these thin films, which could have resulted from the substitution of Ca^{2+} in the Ti^{4+} site. $P-E$ and $C-V$ curves further confirm that these BCT thin films are ferroelectric in nature below the observed T_c .

V. CONCLUSIONS

$\text{Ba}_{1-x}\text{Ca}_x\text{TiO}_3$ targets were prepared by a conventional solid-state reaction technique and the thin films were deposited on Pt-coated Si substrates by a pulsed excimer laser ablation technique. The anomalous drastic decrease in T_c was attributed to the substitution of Ca^{2+} ions in the Ti^{4+} site. The dielectric constant and dielectric phase transition temperature of the target and thin films was observed to decrease with an increase in the calcium content. There was a significant cross over observed from the normal ferroelectric to diffuse phase transition behavior on the substitution of the higher Ca^{2+} ions in the Ti^{4+} site.

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