



HAL
open science

Domain structure sequence in ferroelectric $\text{Pb}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ thin film on MgO

P.-E. Janolin, B. Fraisse, B. Dkhil, Françoise Le Marrec, E. Ringgaard

► **To cite this version:**

P.-E. Janolin, B. Fraisse, B. Dkhil, Françoise Le Marrec, E. Ringgaard. Domain structure sequence in ferroelectric $\text{Pb}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ thin film on MgO. Applied Physics Letters, 2007, 90 (16), pp.162906. 10.1063/1.2727563 . hal-00382765

HAL Id: hal-00382765

<https://hal.science/hal-00382765>

Submitted on 3 Jun 2021

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

See discussions, stats, and author profiles for this publication at: <https://www.researchgate.net/publication/224404786>

Domain structure sequence in ferroelectric $\text{Pb}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ thin film on MgO

Article in Applied Physics Letters · May 2007

DOI: 10.1063/1.2727563 · Source: IEEE Xplore

CITATIONS

11

READS

282

5 authors, including:



Pierre-Eymeric Janolin

CentraleSupélec

65 PUBLICATIONS 1,266 CITATIONS

[SEE PROFILE](#)



Bernard Fraisse

Université de Montpellier

106 PUBLICATIONS 1,854 CITATIONS

[SEE PROFILE](#)



Brahim Dkhil

CentraleSupélec

342 PUBLICATIONS 11,289 CITATIONS

[SEE PROFILE](#)



F. Le Marrec

Université de Picardie Jules Verne

47 PUBLICATIONS 538 CITATIONS

[SEE PROFILE](#)

Some of the authors of this publication are also working on these related projects:



Photovoltaic and photocatalytic potential of ferroelectrics [View project](#)



Multiferroic Materials [View project](#)

Domain structure sequence in ferroelectric $\text{Pb}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ thin film on MgO

Pierre-Eymeric Janolin,^{a)} Bernard Fraisse, and Brahim Dkhil
*Laboratoire Structures, Propriétés et Modélisation des Solides, CNRS, École
 Centrale Paris, 92295 Châtenay-Malabry Cedex, France*

Françoise Le Marrec
*Laboratoire de Physique de la Matière Condensée, Université de Picardie Jules Verne,
 80039 Amiens, France*

Erling Ringgaard
Ferroperm Piezoceramics A/S, Hejreskovvej 18A, DK-3490 Kvistgård, Denmark

(Received 17 February 2007; accepted 19 March 2007; published online 19 April 2007)

The structural evolution of a polydomain ferroelectric $\text{Pb}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ film was studied by temperature-dependent x-ray diffraction. Two critical temperatures were evidenced: $T^* = 740$ K, corresponding to a change in the domain structure ($a/c/a/c$ to $a_1/a_2/a_1/a_2$), and $T_C^{\text{film}} = 825$ K, where the film undergoes a ferroelectric-paraelectric phase transition. The film remains tetragonal on the whole range of temperature investigated. The evolutions of the domain structure and lattice parameters were found to be in very good agreement with the calculated domain stability map and theoretical temperature-misfit strain phase diagram, respectively. © 2007 American Institute of Physics. [DOI: 10.1063/1.2727563]

Among the ferroelectric materials, $\text{Pb}(\text{Zr}_{1-x}\text{Ti}_x)\text{O}_3$ (PZT $1-x/x$) is used in most of the optical and microelectronic devices because of its exceptional properties. With temperature this solid solution undergoes a paraelectric (cubic) to ferroelectric (slightly distorted unit cell) phase transition. For Ti-rich compositions, the symmetry of the ferroelectric phase is tetragonal, whereas the Zr-rich composition is rhombohedral (except for a narrow region close to PbZrO_3). The two phases are separated by a morphotropic phase boundary (MPB) at $x \approx 0.48$, which symmetry at low temperature has been extensively discussed.¹⁻⁹ Because of the anisotropy of their properties, controlling the domain structure of ferroelectric materials is desirable; this can be achieved through epitaxial growth of thin films.

An extension of the tetragonal phase toward the MPB was observed,¹⁰ and different domain structures have been observed for tetragonal PZT films deposited on MgO: epitaxial PZT 52/48 has been deposited by different techniques including sol-gel,¹¹ pulsed laser deposition (PLD),¹² and sputtering.¹⁰ For the three techniques a 100% c -domain structure has been reported with thickness ranging from 90 to 700 nm. PZT 40/60 has also been deposited by PLD (Ref. 13) and the domain structure was found to be 100% c domains. A polydomain structure has been reported by Lee *et al.*,¹⁴ with a decrease of the volume fraction of c domains (α^c) with increasing Ti concentration from 90% for PZT 32/68 down to 75% for pure PbTiO_3 . PbTiO_3 , the end member of PZT, has been studied as a thin film deposited on MgO by many authors. α^c was reported to vary from 57% to 90%, depending on the thickness of the film and to increase¹⁵ as well as to decrease¹⁶ with film thickness. It should be underlined that as these volume fractions have been calculated in different ways, the comparison is not always straightforward (see Ref. 17). In addition Roemer *et al.* showed that α^c is

strongly dependent on the deposition temperature and partial pressure of O_2 for thin film pulsed laser deposited.¹⁸ The reduction of α^c with temperature was predicted^{19,20} and measured for PbTiO_3 films deposited on MgO.^{15,21-23}

Lattice parameter evolution with temperature for PbTiO_3 thin films deposited on MgO was reported by Kim *et al.*²³ and Batzer *et al.*²¹ The transition temperature for these films has been reported to exhibit different behaviors and remains unclear: Batzer *et al.*²¹ reported the transition temperature of the film T_C^{film} to be lowered by 10 K (35 K) when heating (cooling) as a consequence of the compressive stress arising from the difference in the thermal expansion coefficients between the film and the substrate. Kim *et al.*²³ reported T_C^{film} to be 10 K higher (40 K lower) than the bulk when heating (cooling). Temperature evolution of $\text{Pb}(\text{Zr}_{1-x}\text{Ti}_x)\text{O}_3$ ($x = 1, 0.92, 0.84, 0.76, 0.68$) thin films deposited on MgO was reported by Lee *et al.*¹⁴ The out-of-plane lattice parameters as well as T_C^{film} were equal to the corresponding bulk values, i.e., the films considered were strain-free.

We have performed x-ray diffraction measurements to follow the evolution with temperature from 14 up to 925 K of both the lattice parameters and fraction of c domains (α^c) of PZT 20/80 film deposited on MgO(001). These evolutions allow direct comparison with the calculations done on multidomain PZT film as well as the discussion of the symmetry of the paraelectric phase. At room temperature, bulk PZT is tetragonal with $a_b = 3.950$ Å, $c_b = 4.150$ Å, and $c_{\text{MgO}} = 4.213$ Å. 100 nm PZT 20/80 film was deposited by PLD with a KrF excimer laser ($\lambda = 248$ nm), under 0.2 mbar O_2 with a laser repetition rate of 2 Hz at a fixed fluence of 1.6 J/cm². Ceramic $\text{Pb}_{1.1}(\text{Zr}_{0.2}\text{Ti}_{0.8})\text{O}_3$ target was used to deposit the 100 nm film at a substrate temperature of 650 °C. The precise thickness of a PZT 20/80 film deposited on SrTiO_3 was determined through finite size oscillations, and the thickness of the PZT 20/80 film deposited on MgO was estimated from the calculated deposition rate. The composi-

^{a)}Electronic mail: pierre-eymeric.janolin@ecp.fr

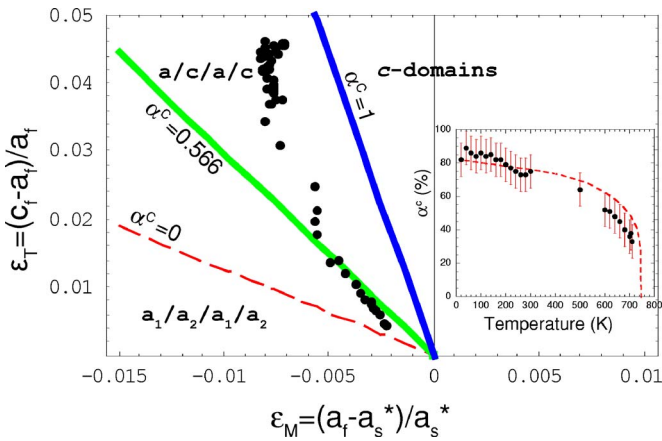


FIG. 1. (Color online) Calculated domain stability map for PZT (20/80)/MgO. The solid circles are the experimental points from $T = 14$ to 740 K from top to bottom. For each region, delimited by thick solid lines, the most stable phase is indicated. The $a/c/a/c$ phase is metastable in the region between the lines $\alpha^c = 0.566$ (solid) and $\alpha^c = 0$ (dashed). Inset: α^c vs temperature with the fit using $\alpha^c = [(1 - 0.5\eta)/(1 - \eta)][(1 + \nu)/(1 - \eta)]\epsilon_M/\epsilon_T$ with $\eta = 0.1$ and $\nu = 0.36$.

tion of the PZT film has been checked through volume calculation.⁵ Lattice parameter determination was carried out on a high-precision diffractometer using Cu K_β wavelength issued from a 18 kW rotating anode generator.

At room temperature, the out-of-plane lattice parameters of the films were calculated from $(00l)_{l=2,3,4}$ Bragg reflections to improve accuracy and to correct any misalignment of the sample and are $a_f^\perp = 3.974$ Å and $c_f^\perp = 4.129$ Å. It has not been possible to determine the in-plane lattice parameters even though the out-of-plane ones give a good approximation of their value.²⁴ The film is polydomain ($a/c/a/c$) and, from the intensity ratio of the (002) and (200) PZT 20/80 rocking curves, composed of $\alpha^c \approx 75\%$ of c domains and $\approx 25\%$ of a domains. These values are typical for PbTiO₃ films deposited on MgO (Refs. 15 and 16) but somewhat smaller than the interpolated value from Ref. 14 which is consistent with our film being not strain-free.

Intriguingly, the film is mainly composed of c domains even though it experiences a tensile stress at room temperature. This tensile stress results from firstly the tensile stress developed at the deposition temperature (T_d) because of both the lattice mismatch between PZT and MgO and the deposition process itself, and secondly from the intrinsic stress associated with the phase transition. This tensile stress is only partially compensated by the compressive thermal stress ($\alpha_{\text{MgO}} = 12.4 \times 10^{-6} \text{ K}^{-1}$ and $\alpha_{\text{PZT}} = 9.2 \times 10^{-6} \text{ K}^{-1}$) experienced when cooling down from T_d to RT.

We have calculated the domain stability map for PZT (20/80) on MgO(001) following Alpay and Roytburd.²⁰ The experimental points have been plotted on the calculated domain stability map (see Fig. 1) and it confirms the x-ray observations: a film being slightly under tension (i.e., $\epsilon_M < 0$) is composed of majority of c domains as long as the tetragonality (ϵ_T) is high enough. Here $\epsilon_T = (c_f - a_f)/a_f$ is equal to 0.039 at room temperature.

For $1 > \alpha^c > 0.566$ (i.e., $14 < T(\text{K}) < 600$) the observed ($a/c/a/c$) domain structure is in agreement with the calculated domain stability map. When α^c becomes smaller than 0.566 (which corresponds to $T > 600$ K), the $a/c/a/c$ phase becomes metastable and disappears for $\alpha^c = 0$, at T^* . There-

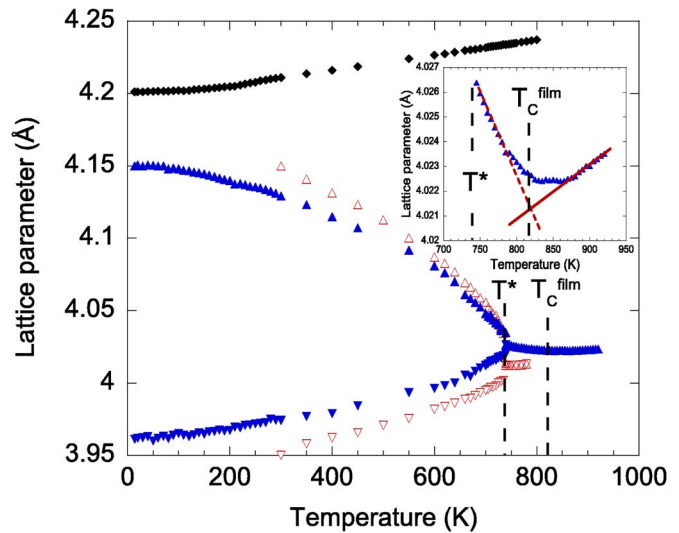


FIG. 2. (Color online) Lattice parameters of PZT 20/80 bulk (c_f : Δ , a_f : ∇) and thin film (c_f^\perp : \blacktriangle , a_f^\perp : \blacktriangledown) deposited on MgO (\blacklozenge). Inset: High-temperature evolution of the out-of-plane lattice parameter. Full line arises from a calculation based on an elastic behavior of the film (see text).

fore, from 600 to 740 K the $a/c/a/c$ and $a_1/a_2/a_1/a_2$ phases should coexist. The temperature evolution of α^c has also been determined (see inset of Fig. 1). The decrease of α^c with temperature has been predicted by Pertsev and Zembilgotov¹⁹ and is consistent with the evolution of the experimental points toward the $\alpha^c = 0$ line (see Fig. 1) with increasing temperature (i.e., decreasing ϵ_T). This decrease toward zero indicates that c domains nucleate for $T = T_C^{\text{film}}$ and then grow continuously as T decreases, a mechanism proposed by Kim *et al.*²³ The measured α^c is, however, overestimated by the fit for $T > 600$ K. We will address this point later.

Figure 2 shows the temperature evolution of the lattice parameters of both PZT 20/80 thin film and MgO between 14 and 925 K. For comparison the lattice parameters of the PZT 20/80 target used for deposition are also plotted from 300 to 800 K. Assuming that the overall evolution is depicted by the out-of-plane lattice parameters, the lattice parameters of the film exhibit a bulklike evolution with temperature and they become equal when heating at T^* close to $T_C^{\text{bulk}} = 740$ K. However, the most intriguing feature in the evolution is the out-of-plane lattice parameter above T^* . Indeed when heating, it begins to decrease linearly with temperature for $T^* < T < 825$ K and then increases linearly (see inset of Fig. 2) for $T > 825$ K.

This peculiar behavior may be explained considering the following evolution, from the deposition temperature T_d : at T_d the film is in equilibrium with an effective substrate (MgO^{*}), i.e., with the misfit-dislocation-modified MgO ($a_f^\parallel = a_{\text{MgO}^*}$ at T_d). In the paraelectric phase, we consider the film as purely elastic with its in-plane parameter driven by the effective substrate. It is then possible to calculate the temperature evolution of the out-of-plane parameter.²⁵ This calculation gives a straight line (represented by the full line on the inset of Fig. 2) that fits very well the high-temperature evolution of the out-of-plane parameter. At ~ 825 K the film cannot be considered as purely elastic anymore. The turning point in the evolution is due to a phase transition. This brings $T_C^{\text{film}} = 825$ K which corresponds to the paraelectric to ferroelectric phase transition

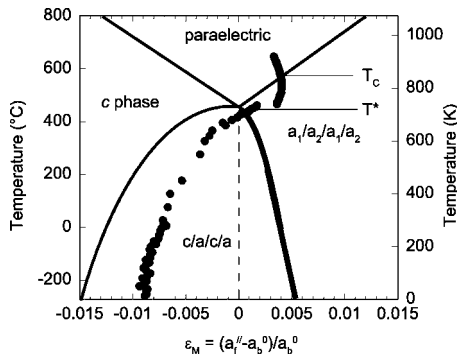


FIG. 3. Experimental misfit strain experienced by the film on the temperature-misfit strain diagram calculated by Kukhar *et al.* (Ref. 26).

temperature. When cooling from T_C^{film} to T^* , the film is tetragonal and ferroelectric and the evolution of the out-of-plane parameter is imposed by the phase transition. It becomes equal to the in-plane parameter at T^* , only temperature where the film adopts a pseudocubic symmetry remaining, however, ferroelectric.

This proposed evolution is supported by the temperature-misfit strain phase diagram. It is indeed possible to calculate the misfit strain experienced by the film for each temperature, $\epsilon_M = (a_f^0 - a_b^0)/a_b^0$, with a a_b^0 the pseudo-cubic parameter of the bulk and to compare it with the recent calculation by Kukhar *et al.*²⁶ of the temperature-misfit strain phase diagram for polydomain epitaxial $\text{Pb}(\text{Zr}_{1-x}\text{Ti}_x)\text{O}_3$ thin films with dense domain structure (see Fig. 3). First of all, the domain structure observed is in agreement with the temperature-misfit strain phase diagram: from 14 K up to T^* , the film is polydomain and the evolution of the experimental points with temperature is in agreement with the decreasing α^c on heating, as the experimental points tend toward the $a_1/a_2/a_1/a_2$ phase which consists of a domains only. Then the experimental points cross the line separating the $a/c/a/c$ and $a_1/a_2/a_1/a_2$ phases at T^* , confirming the ferroelectric nature of the phase above T^* . Moreover, the $a/c/a/c \rightarrow a_1/a_2/a_1/a_2$ transition is of first order, according to the phase diagram. This explains the apparent underestimation of α^c (see inset of Fig. 1) when compared to the fit: the measured volume fraction of c domains in the temperature range where the two structures coexist takes into account not only the a domains of the $a/c/a/c$ structure but also the a domains of the $a_1/a_2/a_1/a_2$ structure, resulting in an overestimation of the a domains and hence an underestimation of the c domains. In other words, when the two structures coexist (600–740 K), we measured α^c for a $(a/c/a/c + a_1/a_2/a_1/a_2)$ structure, which is necessarily smaller than for the $a/c/a/c$ structure alone. Finally, the phase (or structure) sequence proposed and the corresponding transition temperatures are in very good agreement with the temperature-misfit phase diagram. It is worth noting that by shifting the calculated phase diagram toward more positive

misfit strains, we would get a better fit to our experimental data. This shift would cause the phase diagram to be not centered anymore with respect to the zero misfit value.²⁷

In conclusion, we have shown that, despite its domain structure, a polydomain ferroelectric thin film is not necessarily strain-free and exhibits a complex evolution with temperature; a change in the domain structure takes place at T^* , close to T_C^{bulk} from a polydomain $a/c/a/c$ phase to a $a_1/a_2/a_1/a_2$ phase. The Curie temperature of the film is shifted toward higher temperature by ~ 60 K and the symmetry of the lattice remains tetragonal from 14 K up to temperatures higher than the deposition temperature. These results are in full agreement with both the domain stability map we have calculated and with the temperature-misfit strain phase diagram.

This work was partially funded by the MINUET European Program.

- ¹B. Noheda, D. Cox, G. Shirane, J. Gonzalo, L. Cross, and S.-E. Park, *Appl. Phys. Lett.* **74**, 2059 (1999).
- ²B. Noheda, D. Cox, G. Shirane, R. Guo, B. Jones, and L. Cross, *Phys. Rev. B* **63**, 014103 (2000).
- ³K. Lima, A. S. Filho, A. Ayala, J. M. Filho, P. Freire, F. Melo, E. Araujo, and J. Eiras, *Phys. Rev. B* **63**, 184105 (2001).
- ⁴D. Vanderbilt and M. Cohen, *Phys. Rev. B* **63**, 094108 (2001).
- ⁵J. Frantti, S. Ivanov, S. Eriksson, H. Rundlöf, V. Lantto, J. Lappalainen, and M. Kähkönen, *Phys. Rev. B* **66**, 064108 (2002).
- ⁶A. Souza Filho, K. Lima, A. Ayala, I. Guedes, P. Freire, F. Melo, J. Mendes Filho, E. Araujo, and J. Eiras, *Phys. Rev. B* **66**, 132107 (2002).
- ⁷B. Noheda, L. Wu, and Y. Zhu, *Phys. Rev. B* **66**, 060103(R) (2002).
- ⁸D. Hatch, H. Stokes, R. Ranjan, Ragini, S. Mishra, D. Pandey, and B. Kennedy, *Phys. Rev. B* **65**, 212101 (2002).
- ⁹I. A. Kornev, L. Bellaiche, P.-E. Janolin, B. Dkhil, and E. Suard, *Phys. Rev. Lett.* **97**, 157601 (2006).
- ¹⁰S. H. Oh and H. M. Jang, *Appl. Phys. Lett.* **72**, 1457 (1998).
- ¹¹K. Nashimoto, D. K. Fork, and G. B. Anderson, *Appl. Phys. Lett.* **66**, 822 (1995).
- ¹²J. Lee, A. Safari, and R. L. Pfeffer, *Appl. Phys. Lett.* **61**, 1643 (1992).
- ¹³H. J. Kim, S. H. Oh, and H. M. Jang, *Appl. Phys. Lett.* **75**, 3195 (1999).
- ¹⁴K. Lee, Y. Kang, and S. Baik, *J. Mater. Res.* **14**, 132 (1999).
- ¹⁵K. S. Lee and S. Baik, *J. Appl. Phys.* **87**, 8035 (2000).
- ¹⁶B. Kwak, A. Erbil, J. Budai, M. Chisholm, L. Boatner, and B. Wilkens, *Phys. Rev. B* **49**, 14865 (1994).
- ¹⁷W.-Y. Hsu and R. Raj, *Appl. Phys. Lett.* **67**, 792 (1995).
- ¹⁸A. Roemer, E. Millon, B. Vincent, A. Boudrioua, O. Pons-Y-Moll, R. M. Defourneau, and W. Seiler, *J. Appl. Phys.* **95**, 3041 (2004).
- ¹⁹N. A. Pertsev and A. G. Zembilgotov, *J. Appl. Phys.* **80**, 6401 (1996).
- ²⁰S. Alpay and A. Roytburd, *J. Appl. Phys.* **83**, 4714 (1998).
- ²¹R. S. Batzer, B. M. Yen, D. Liu, H. Chen, H. Kubo, and G. R. Bai, *J. Appl. Phys.* **80**, 6235 (1996).
- ²²K. S. Lee and S. Baik, *J. Appl. Phys.* **85**, 1995 (1999).
- ²³S. Kim, Y. Kang, and S. Baik, *J. Am. Ceram. Soc.* **79**, 1105 (1996).
- ²⁴K. S. Lee, J. H. Choi, J. Y. Lee, and S. Baik, *J. Appl. Phys.* **90**, 4095 (2001).
- ²⁵P.-E. Janolin, Ph.D. thesis, École Centrale Paris, 2006.
- ²⁶V. G. Kukhar, N. A. Pertsev, H. Kohlstedt, and R. Waser, *Phys. Rev. B* **73**, 214103 (2006).
- ²⁷B.-K. Lai, I. Kornev, L. Bellaiche, and G. Salamo, *Appl. Phys. Lett.* **86**, 132904 (2005).