Strain-gradient-induced switching of nanoscale domains in free-standing ultrathin films

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We report first-principle atomistic simulations on the effect of local strain gradients on the nanoscale domain morphology of free-standing PbTiO$_3$ ultrathin films. First, the ferroelectric properties of free films at the atomic level are reviewed. For the explored thicknesses (10 to 23 unit cells), we find flux-closure domain structures whose morphology is thickness dependent. A critical value of 20 unit cells is observed: thinner films show structures with 90° domain loops, whereas thicker ones develop, in addition, 180° domain walls, giving rise to structures of the Landau-Lifshitz type. When a local and compressive strain gradient at the top surface is imposed, the gradient is able to switch the polarization of the downward domains, but not to the opposite ones. The evolution of the domain pattern as a function of the strain gradient strength consequently depends on the film thickness. Our simulations indicate that in thinner films, first the 90° domain loops migrate towards the strain-gradient region, and then the polarization in that zone is gradually switched. In thicker films, instead, the switching in the strain-gradient region is progressive, not involving domain-wall motion, which is attributed to less mobile 180° domain walls. The ferroelectric switching is understood based on the knowledge of the local atomic properties, and the results confirm that mechanical flexoelectricity provides a means to control the nanodomain pattern in ferroelectric systems.

I. INTRODUCTION

Flexoelectricity, the linear coupling between electric polarization $P$ and inhomogeneous deformation, is always allowed by symmetry in all insulators since strain gradients break inversion symmetry. This effect, found 50 years ago, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motivated a revival interest in flexoelectricity, which has become an attractive topic in material science in the recent experimental breakthroughs, has traditionally received little attention owing to its relatively weak effect in macroscopic samples. At the nanoscale, however, strain gradients are typically much larger than at macroscopic scales, as reduced dimensions imply larger gradients. The impressive progress in the design and control of nanoscale structures has motiva...
ent affect the domain pattern in free-standing ultrathin films? and How does this depend on the film thickness and the strain gradient strength? To answer these, using a first-principles-based atomistic model we study the flexoelectric switching in ultrathin films of the stereotypical ferroelectric PTO, when a longitudinal strain gradient is applied locally over the center of one domain. Our simulations indicate that for the explored films, thinner than 10 nm, the domain patterns in the unstrained films are thickness dependent and responsible for the film being locally deformed. In agreement with this, the film response to a local and longitudinal strain gradient varies according to the nature of the domain pattern and as a function of the gradient intensity. The paper is organized as follows. Section II describes the computational details for modelling the film and the conditions for mimicking a strain gradient. Section III presents the results in two parts. First, the local properties of strain-free films are reported for two thicknesses in order to understand later the gradient-induced changes. Second, the domain pattern evolution for each of these films is investigated as the strain gradient strength is enhanced. Finally, the paper concludes with a summary in Sec.IV.

II. METHODOLOGY: COMPUTATIONAL DETAILS

Prototype ferroelectric PTO is modeled using an atomistic core-shell model fitted completely to first-principles calculations (i.e., no explicit experimental data were used as input) as described in detail in ref. 18. The model contains fourth-order core-shell couplings, long-range Coulombic interactions, and short-range interactions. It has shown to correctly reproduce the cubic-tetragonal phase transition in bulk20 and also was able to describe the surface properties and interface effects of ultrathin films.21

Here we study free-standing ultrathin films of PTO under ideal open-circuit electric boundary conditions performing classical molecular dynamics (MD) simulations using the DL_POLY package20 at a constant temperature of 50 K, where PTO presents a tetragonal ferroelectric phase. Our films have two PbO-terminated surfaces and are modeled using supercells of \( N_x \times N_y \times N_z \) unit cells. Periodic boundary conditions are applied along the in-plane pseudocubic [100] and [010] directions (axes \( x \) and \( y \), respectively) but not along the [001] one (axis \( z \)) defined as the growth direction. The long-ranged electrostatic energy and forces are calculated by a direct sum method20 which is a computationally efficient alternative to the Ewald summation. In the \( xy \) plane, the simulated supercell is kept constant and constrained to form a simple square lattice (\( N_x = N_y \)) of edge \( N_x \times a \) with \( a = 0.3866 \text{ nm} \), the in-plane lattice parameter in bulk computed at 50 K with the present model. The film thicknesses range from 4.0 nm (10c) to 9.2 nm (23c), where \( c \) is the lattice parameter in the growth direction. Atoms are free to relax along all directions, except when mentioned, as long as they remain inside the simulation box.

Local polarization is defined here as the dipole moment per unit volume of a perovskite unit cell centered on the Ti atom and delimited by the eight Pb nearest neighbors at the corners of the box. We consider contributions from all atoms in the conventional cell and atomic positions with respect to that center:

\[
p = \frac{1}{v} \sum_i \frac{1}{\omega_i} q_i (r_i - r_{Ti})
\]

where \( v \) is the volume of the cell, \( q_i \) and \( r_i \) denote the charge and the position of the \( i \) particle, respectively, and \( \omega_i \) is a weight factor equal to the number of cells to which the particle belongs. The reference position \( r_{Ti} \) corresponds to the core of the Ti atom, and the sum runs over 29 particles, including cores and shells of the surrounding ions (8 Pb and 6 O atoms) plus the shell of the Ti atom. Note that \( p \) is independent of the origin for the \( r_i \) vectors and vanishes when atoms are at their high-symmetry positions. The charge values \( q_i \) are given in Ref. 18.

The initial configuration of the strain-free film for all the considered thicknesses is set up by creating 180° stripe domains with polarizations pointing upward and downward normal to the surface, alternated along the \( x \) axis, as schematically shown in the inset of Fig. 1. The simulation supercell, indicated by dashed lines, holds information for two thicknesses in order to understand later the gradient-induced changes. Second, the domain pattern evolution for each of these films is investigated as the strain gradient strength is enhanced. Finally, the paper concludes with a summary in Sec.IV.
depend on the film thickness as described below.

Strain gradient description. As we are interested in the effect of switching the out-of-plane polarization, we apply strain gradients in the direction normal to the surface, that is, a gradient of the out-of-plane strain component \( \eta_{zz} = c/c_0 - 1 \) along the [001] direction denoted as \( \partial \eta_{zz}/\partial z \). We impose a linear gradient localized on an in-plane square region of \( 4 \times 4 \) unit cells that penetrates until the central plane, which compresses the upper plane by a given strain \( -\eta_{zz}^{\text{max}} \) and then decreases linearly in the intermediate planes until vanishing at the central cells. One advantage of using the present atomistic model to treat flexoelectric effect\(^{22}\) is that the interatomic potential does not need to be modified; it is sufficient to place the system under appropriate mechanical-boundary conditions that mimic the required gradient. To do that, one first has to establish the strain gradient in the film, noting that its presence implies a stress gradient which produces net forces acting on the atoms inside each unit cell. During MD simulations, the atoms will try to relax in order to vanish these net forces. Thus, in order to preserve the gradient a force pattern must be applied to the atoms in the unit cell; i.e., certain atoms must be fixed along the full simulation. Specifically, to set up the gradient, we start from the relaxed free film (which already presents a domain structure) and displace rigidly the atomic planes of the chosen unit cells until the average distance between consecutive Pb-O lattice planes equals the required local strains \( \eta_{zz} \) in order to have \( \partial \eta_{zz}/\partial z \). We choose to freeze out the \( z \) components of the Pb-core positions, allowing all other internal degrees of freedom (the Pb shell and other atoms, cores plus shells) to relax along all directions. Other choices are possible, for instance, the force pattern could be distributed between all atoms in the unit cell in different proportions\(^{23}\). Finally, note that the local strain gradient induces an internal electric field which can compensate the depolarization field allowing that part of the free-standing film is poled.

III. RESULTS

A. Free-standing ultrathin films of PbTiO\(_3\)

We focus first on strain-free films under ideal open circuit boundary conditions and revisit their local features. To begin with, the optimum lateral size of the simulation supercell (i.e., the number of cells \( N_x \)) is determined. For each considered thickness, we build up supercells of different lateral sizes, relax their structures, compare their total energies, and choose the lowest-energy one. The optimized structures are described below; here we just present the optimum width \( \omega \) of one domain period as a function of the film thickness \( t \). Figure 1 displays the linear relation between \( \omega \) and the root square of \( t \), showing the fulfillment of the Kittel law. We found that the optimum lateral sizes of the supercells are \( 26a, 30a, 34a, \) and \( 36a \) for thickness of \( 10c, 16c, 20c, \) and \( 23c \), respectively.

Hereafter we present results only for films of two thicknesses \( 4.0 \) nm and \( 9.2 \) nm because of their representative domain structures. Figure 2 shows a side view of cell-by-cell polarization map for the two thicknesses, where colors indicate the \( z \)-component values in the left panels and the \( c/a \) cell tetragonality in the right ones. Starting with the left panels, the 10c ultrathin film (top panel) shows a flux-closure arrangement around central cores or vortices. This vortex state, where the local \( P \) orientation rotates continuously around one geometrical core, can also be described as a 90\(^\circ \)-domain loop. Indeed, we can demarcate upward and downward \( P \) domains (hereafter UP and DOWN, respectively) with a rhombus shape surrounded by triangular caps (where \( P \) is almost parallel to the surface) that complete an overall closed loop. Our simulations show that this domain morphology holds for thicker films until thicknesses of \( \sim 20 \) unit cells. Above that critical value, the central vortices enlarge in the normal direction, giving rise to 180\(^\circ \) DWs that grow together with the film thickness. Consequently, the shape of the UP and DOWN domains changes to hexagonal and a domain structure as envisioned by Landau-Lifshitz is formed, as shown in the bottom panels for a 23c-thickness film. In this structure, in particular, we see that the 180\(^\circ \) DWs end up in vortices or confluences of three DWs as predicted by Srolovitz and Scot\(^{25}\) and recently confirmed with high-resolution transmission electron microscopy images of Pb(Zr\(_{1-x}\)Ti\(_x\))O\(_3\)\(^{20}\).

As regards the thicknesses of the DWs, they vary depending on the type and location inside the film, the difference being more evident in the thicker films. In the inner region, the orientation of the \( P \) changes abruptly from up to down over a distance of two lattice constants, creating very narrow 180\(^\circ \) DWs, in agreement with previous theoretical results for defect-free walls\(^{11}\). The 90\(^\circ \) DWs instead differ according to their position with respect to the out-of-plane \( P \) orientation, as indicated schematically in the bottom-right panel in Fig. 2 for the UP domain as an example, but the same behavior is repeated in the DOWN domain. In the top part, the \( P \) rotates 90\(^\circ \) in only three unit cells, leading to narrow 90\(^\circ \) DWs. In the bottom part, instead, the \( P \) rotates more continuously, requiring a few unit cells to change from the in-plane \( P \) at the surface until pointing perpendicular, making these 90\(^\circ \) DWs much broader. This behavior can be attributed to the electrostatic repulsion between the head-to-head in-plane Ps that confront each other at the junction of two triangular domains.

The cell-by-cell tetragonal \( c/a \) ratio is indicated by colors in the right panels of Fig. 2 where \( c \) and \( a \) were obtained by taking the distances between pairs of nearest-neighbor Pb atoms along \( z \) and \( x \), respectively. The cells inside the triangular domains with \( P \) parallel to the surfaces are almost cubic, whereas those at the cores of the domains with \( P \) perpendicular are tetragonal and their values are thickness dependent. Further, and in agreement with the high strain-\( P \) interaction in PTO, these tetragonality maps demonstrate the difference be-
between the 90° DWs. The narrow upper ones are well defined, presenting an abrupt change in the c/a ratio as the tetragonal axis rotates from perpendicular to parallel to the surface; however, the broader ones at the bottom are characterized by a large tetragonality (reaching, for instance, ~8% in the 23c film) which attenuates gradually in a few unit cells towards both sides. Concerning the 180° DWs in the thicker films, the tetragonality does not change notably, rather the wall is defined by an abrupt relative displacement or offset of the Pb atoms along z.

Finally, we emphasize the significant lattice deformation observed at the surfaces (shown by different colors in the right panels of Fig. 2) as well as across the film thickness specially in the 90° DWs.

To obtain further insights, we monitor those properties in detail through the film thickness. Figure 3 shows the in-plane a and out-of-plane c lattice parameters across the thickness for both films. Averaging these values over complete planes parallel to the surface [open (green) circles], we find that the average local a’s remain constant across the film due to the mechanical boundary conditions imposed on the simulation supercell. The average c values show the typical symmetric profile determined by the two free surfaces: they are compressed at both surfaces and gradually increase towards the inner film, consistent with previous result. However, examining closely at the central region of each domain, the local c’s present an asymmetric distribution: they are highly compressed in the two unit cells of the top surface (with respect to the P orientation) and slightly enlarged at the bottom ones. The strain change along [001] is compensated in the plane only by the local lattice parameters a, which behave oppositely to c, but not by the b parameters, which remain almost constants (not shown here).
corroborating that the unit cells at the surface neighborhood are highly distorted. This particular asymmetry through the thickness was also predicted in superlattices of PTO/SrTiO$_3$ from first-principles calculations$^{29}$ and even in the limit case of PTO films of 1c thickness whose unit cell is distorted$^{19}$ Thus, this information together with the above c/a description shows that the free surfaces induce spontaneous strain gradients that break the inversion symmetry in the film, in agreement with previous work$^{15,16}$ Furthermore, knowledge of the local strain maps is essential to predict and understand the effect of an imposed strain gradient.

Moving to the right panels of Fig. 3, the profiles of the local $P_z$ modulus show that the averaged values per lattice plane are symmetric. Remarkably, inside each domain, opposite to the highly asymmetric profiles of local c, the $P_z$ profiles are almost symmetric (in thinner films, actually they are slightly enhanced towards the top part). Then we note that the modulus of $P$ and not $P_z$ follows the c behavior across the film.

TABLE I. Oxygen octahedra rotations (in degree) of the AFD$_z$ distortions for two thicknesses (t) in different atomic layers starting at the top surface, given at the center of the UP, DOWN and triangular domains, and averaging per full TiO$_2$ atomic layer. Abinitio results after Ref. 10 are reported in parenthesis. The last row of data corresponds to our calculations performed in the bulk material with two domains separated by 180° domain walls.

| t  | Atomic layer | UP     | DOWN   | Triangular | Average |
|----|--------------|--------|--------|------------|---------|
| 10c| 1            | 14.2   | 13.3   | 13.0       | 13.0 (11.4) |
|    | 2            | 1.0    | 2.9    | 3.5        | 2.8 (2.9) |
|    | 3            | 6.0    | 6.2    | 6.4        | 6.5 (3.9) |
|    | 4            | 4.9    | 5.1    | 5.2        | 4.9     |
|    | 5            | 4.8    | 5.0    | 5.8        | 5.2     |
| 23c| 1            | 14.3   | 13.0   | 13.0       | 13.1 (11.4) |
|    | 2            | 0.9    | 2.8    | 4.1        | 3.0 (2.9) |
|    | 3            | 5.7    | 6.3    | 6.5        | 6.6 (3.9) |
|    | 4            | 3.1    | 4.3    | 6.4        | 5.16    |
|    | 5            | 3.0    | 4.1    | 6.1        | 5.1     |
|    | 11           | 0.8    | 1.0    | 4.0        | 1.6     |
| bulk| 1.0          | 1.0    | 3.9    | 1.0        | 3.3     |

To end up this section, we analyze the local distribution of the antiferrodistortive distortion which has been shown$^{10,15,29}$ to coexist with the polar distortion at the surface, even though it essentially vanishes in bulk. It involves the oxygen octahedron rotation around the central Ti ion, where the rotation axis is the z axis (AFD$_z$) with out-of-phase movements between neighbor cells on the (001) plane but successive in-phase movements along the [001]. Its strength is described by the rotation angle of TiO$_2$ in the normal layer at each unit cell. Table I lists the AFD$_z$ rotation angle dependence of the atomic layers starting at the top surface, for the two films of interest and in different regions of the film by entering perpendicular to the surface: in the central cells of the UP and DOWN domain, and crossing the center of one triangular domain (reaching the central vortex of the 90° DW loop or the 180° DWs, depending on the film thickness). The last column reports the values averaged for each complete TiO$_2$ atomic layer parallel to the surface. Moving along [001] in the center of one domain, it is found that the rotation angle of $\sim 13°$ in the superficial cells decays towards the internal atomic layers: although it almost vanishes in the second layer, recovers in the third one, and then keeps decreasing slowly. It reaches the bulk value of $\sim 1°$ just at the eleventh layer of the 23c film, while in the thinner film, the superficial effects predominate. Furthermore, we note that cells in opposite surfaces differ in $\sim 1°$, which is also related to the different natures of the junction of the 90° DWs as explained before. On the other hand, when entering the film through the triangular domains, the AFD$_z$ distortions from the second layer towards the inner ones are a bit larger than those in the center of the UP and DOWN domains. Specifically, in the 180° DWs the AFD$_z$ distortion is about 4° (the octahedra rotate to allow the abrupt P change, in addition to the offset), versus the $\sim 1°$ in the domain cores, where the cells are tetragonal with ferroelectric distortions mostly along z. The averaged values show a behavior similar to that in the cores of the out-of-plane P domain but with values $\sim 1°$ larger than in the inner cells due to the values in the DWs. Finally, note that our results for the first three atomic layers can be compared, and indeed fully agree with previous calculations reported using the present model$^{11}$ as well as first-principles method$^{10}$ Although those calculations have provided evidence of AFD$_z$ variation in the near-surface region, they were performed using smaller supercells than here and without any reference to the effect of the domain pattern. Hence, our inspection of the AFD$_z$ local distribution gives insights into the dependence of the domain morphology, especially in the DW neighborhood.

B. Film response to a local strain gradient

Now we investigate how the PTO film responds to a compressive strain gradient applied locally as its intensity increases. We are particularly interested in following the evolution of the domain configuration until eventually the polarization switches in the strain-gradient zone. As the first step, we have to choose where to apply the strain gradient. Based on the observed local strain distribution in the central region of each domain (Fig. 3), we suppose that a compressive strain gradient on the top surface of the film would only accentuate the already existent inhomogeneous compression in the UP domain; on the contrary, it would alter the almost-constant c distribution in the DOWN domain. Therefore, we apply to the latter a linear gradient on the top half of the film along the z axis as described in Sec. II and shown in Fig. 3(a) where the local c profile is plotted before and after the application for a particular $\eta_{zz}^{max}$ of $-4.25\%$. The same profile is established on a region of
FIG. 4. (Color online) Results for a 10c-thickness film. (a) Local lattice parameter $c$ in the central region of the DOWN domain for a strain-free film (triangles) and after the strain gradient is applied (circles). (b-d) Cell-by-cell $P$ distributions for an unstrained film (b) and when a strain gradient of $-1.53 \times 10^7$ m$^{-1}$ (c) and $-2.18 \times 10^7$ m$^{-1}$ (d) imposed in the zone delimited by a parallelepiped. For all cases, the 3D map and its projection on the central $xy$ plane are shown. In (b) and (c) only one half of the supercell projection on that plane is plotted, for clarity, and the lines mark the position of the central vortices. In (d), the line in the 3D map also indicates the positions of the central vortices, and the projections on three planes are presented: (d,i) on the central $xy$ plane, where dashed lines indicate the positions of two lateral planes, $L_x$ (d,ii) and $L_y$ (d,iii). Colors denote the $z$ components of the polarization.

$4 \times 4$ unit cells that extends from the top surface to the central plane, as schematically indicated in the figure by a parallelepiped. The strain gradient strength is increased progressively by varying the compressive strain in the superficial cells $\eta_{zz}^{\max}$ in intervals of $-0.25\%$. For each strain gradient value, we carry out an MD simulations until the system rearranges to that strain condition.

Thinner films. The results for the 10c film are illustrated in Figs. 4(b) 4(d) using three-dimensional (3D) maps of the local $P$ and projections of them on different planes. The unstrained film is shown in Fig. 4(b), and how it changes when the imposed local strain gradient $g$ takes two particular values, namely, $g_1 = -1.53 \times 10^7$ m$^{-1}$ and $g_2 = -2.18 \times 10^7$ m$^{-1}$ in Figs. 4(c) and 4(d), respectively. Starting with the unstrained film, which is repeated here so as to have a reference (same as the top panel of Fig. 2), the 3D map displays a domain structure of 90° domain loops. Its projection on the central $xy$ plane shows only two domains with the $P$ downward and upward perpendicular to the paper plane separated by the central vortices, whose positions aligned along the $y$ axis are indicated by a line in the figure. The square in the DOWN domain identifies the region where the strain gradient will be applied. Thus, when the local strain gradient is imposed and as its strength $g$ increases, we find that the response of the film presents three different behaviors.

(I) For $g < g_1$, no appreciable changes are observed and the local $P$ distribution remains similar to that of the unstrained film.

(II) For $g_1 \leq g < g_2$, when $g = g_1$, corresponding to an $\eta_{zz}^{\max}$ of $-3.00\%$, the whole domain structure is displaced, almost without changing either the size or the shape of the domains as shown in the 3D map of Fig. 4(c). The projection on the $xy$ plane clearly shows that the central vortices have been shifted in $-x$ towards the gradient region and sit just before its border. $P$ distributions similar to this one are observed as the gradient is further increased.

(III) For $g = g_2$ or $\eta_{zz}^{\max} = -4.25\%$, the dipoles rearrange and the local $Ps$ in the gradient region, initially downward, are switched [3D map of Fig. 4(d)]. This region has merged with the original UP domain, extending its size and therefore changing its shape. The new domain distribution can be appreciated more easily in the projection on the central $xy$ plane [Fig. 4(d,i)]. Also, we examine two lateral planes that cross the gradient region. The projection on a $yz$ plane shown in Fig. 4(d,ii) (where all cells pointed downward before the gradient
was applied) reveals that the new domain pattern has developed a flux-closure arrangement around the central vortices analogous to that found in the unstrained film. The projection on the $L_y$ plane [Fig. 4(a,iii)] shows that most cells, including those in the gradient region, are poled upward. The asymmetric distribution indicates that the switching process has started in the vortices that were brought close to the gradient region and now have disappeared, while the other vortices remain on the right side.

Note that due to the supercell in-plane periodicity, it is equally likely that in (II) the vortices had shifted in $+x$ towards the gradient region and therefore the switching process would have started at the other side. Further, we note that the $c$ profile in the gradient region, whose inferior part [open circles in Fig. 4(a)] has been allowed to relax, has adopted a profile quite similar to the one observed in the UP domains of the unstrained films, as was assumed initially. Finally, we want to remark that the observed movements of domains in our free-standing films are allowed because of the absence of a substrate. Its presence, however, can hamper the movement of domains as, for instance, in piezoelectric force microscopy images of Pb(Zr$_{0.2}$Ti$_{0.8}$)O$_3$ patterns in epitaxial thin films.

**Hypertoroidal moment.** In order to better describe the evolution of these complex dipolar configurations, we use the recently defined order parameter $h$ named hypertoroidal moment, which provides a measure of subtle microscopic features. It involves the double cross product of the local dipoles $p_i$ with the vectors $r_i$ locating their positions,

$$h = \frac{1}{4V} \sum_i r_i \times (r_i \times p_i)_t$$  \hspace{1cm} (2)

where $V$ denotes the supercell volume, the subindex $t$ labels the transversal component, and the summation runs over all unit cells within the supercell. The transversal component can be approximated as $p_i \approx p - \langle p \rangle$, $(p)$ being the average of the individual dipoles over all the sites. This approximation is also valid for the expression in parenthesis. We refer the reader to Ref. 34 for a detailed discussion of this parameter. For 2D systems (i.e., systems periodic in two Cartesian directions and finite in the third one) such as the films considered here, once the simulation supercell is chosen, the $h$ value is independent of the origin used for $r_i$. However, the magnitude and even the sign of $h$ do depend on the choice made for the supercell, leading to the multivaluedness of the $h$ parameter. We compute the hypertoroidal moment for different supercells whose centers scan the complete central $xy$ plane of the original supercell. Therefore we explore for the first time a surface of $h$ and find that the changes in value and/or sign reflect modifications in the full $P$ distribution.

Although $P$ switching is a dynamical and complex process, we may gain some insight by examining the surface of only the z-component of the hypertoroidal moment $h_z$ (the other two Cartesian components are almost null) as the imposed strain gradient increases. Beginning with the unstrained film shown in Fig. 4(a), we corroborate that the $h_z$ surface displays a sinusoidal dependence along the $x$ direction with the same period as the nanodomains and remains constant along the $y$ direction. Specifically, the maxima (minima) of $h_z$ extend along $y$, coinciding with the positions of the central cores of the UP (DOWN) domain, while $h_z$ vanishes when rotation around the perpendicular $y$ axis exists, certainly where the vortex centers sit. Then as the strain-gradient intensity increases and following the $P$ distributions, the $h_z$ response also presents three behaviors separated by the gradient values $g_1$ and $g_2$.

(I) For $g < g_1$, the sinusoidal shape remains almost unchanged.

(II) For $g_1 \leq g < g_2$, when $g = g_1$, in agreement with the $P$ distribution, the $h_z$ surface shifts in $-x$ as shown by its projection on a transverse cut that crosses the central gradient region [see the central curve in Fig. 5(b)]. $h_z$ behaves similarly as the gradient increases.

(III) For $g = g_2$, needed to fully invert the $P$ in the affected region, the $h_z$ surface shown in Fig. 5(c) has...
FIG. 6. (Color online) Two-dimensional projection of the polarization patterns for an $xz$ plane that crosses the gradient region in a 23$c$-thickness film subject to a strain gradient of (a) $g_1^s = -1.01 \times 10^7$ m$^{-1}$ and (b) $g_2^s = -1.57 \times 10^7$ m$^{-1}$, just before and after the switching, respectively. The gradient region and the immobile 180° DW are indicated.

changed dramatically, indicating that the 90°-domain loop configuration has been modified. Two global extrema are observed, one maximum and one minimum, revealing that the periodicity of the domain structure and thus the extended cores of the each domain have vanished. From the definition, we interpret that the $h_z$ minimum represents the part of the DOWN domain farthest away from the gradient region and therefore the less affected one, whereas the $h_z$ maximum is located in the region of the UP domain farthest away from the distribution of $-P_z$. Fig. 5(b) shows that the $h_z$ profile in this particular direction has slightly gained height in the positive branch, versus an important reduction in the negative one, reflecting the size enhancement of the UP domain at expenses of the DOWN domain.

Summarizing, the evolution of the hypertoroidal moment as a function of the strain gradient strength provides information of the domain-structure evolution localizing the domain cores and the vortices in a regular structure, as well as their redistribution when the $P$s in the strain region are finally switched.

**Thicker films.** Figure 6 displays the $P$-map projection on a transverse cut crossing the gradient region of a 23$c$-thickness film for two strain-gradient values of $g_1^s = -1.01 \times 10^7$ m$^{-1}$ and $g_2^s = -1.57 \times 10^7$ m$^{-1}$, corresponding to $\eta_{zz}^{max}$ of -5.00 and -7.50 %, respectively. As the gradient increases, starting at the surface the local $P$s near and inside the gradient region rotate progressively until pointing parallel to the surface, i.e., their $P_z$ components decrease gradually [as shown in Fig. 6(a) for $g_1^s$] until they fully vanish. A lateral view in a $yz$ plane, not shown here, indicates that the zone with $P$ parallel to the surface starts to grow and expands towards the bottom of the film as the gradient increases, then they are switched. For the $g_2^s$ value [Fig. 6(b)] the affected region is already poled upward and the asymmetric distribution shows that one of the 180° DWs is still immobile. Remarkably, as the gradient is increased further, the switched zone moves forward, generating vortices at the bottom as it continues growing laterally. Thus, unlike the DW movements observed in thinner films, we find that the 180° DWs never move forward to the gradient region. This fact can be attributed to the relatively high energy barrier for the coherent motion of one entire 180° DW compared to the 90° DW case.

In summary, we investigate from an atomistic point of view flexoelectricity-induced domain evolution. In free-standing PbTiO$_3$ ultrathin films with thickness-dependent domain patterns, we study the ferroelectric switching induced by a strain gradient acting locally in the direction normal to the surface. We first review the properties of strain-free films and confirm that the inhomogeneity of polarization, strain (strain gradient), and antiferrodistortive rotational angles appears not only among the surface cells but also in the neighborhood of the DWs. For the explored thicknesses (10 to 23 unit cells), we find flux-closure domain structures whose morphology is thickness dependent. A critical value of 20 unit cells is found: thinner films show structures with 90° domain loops, whereas thicker ones develop, in addition, 180° DWs giving rise to structures of the Landau-Lifshitz type. When a compressive strain gradient normal to the surface acts locally at the top surface of the film, we find that it is able to reverse the normal polarization in DOWN domains but not in UP domains, which is understood based on the knowledge of the local strain distribution before imposition of the gradient. The evolution of the domain pattern as a function of the strain gradient strength depends, accordingly, on the film thickness. In thinner films, first one of the center of the 90° DW loops moves towards the gradient region and then the affected region is gradually switched. In thicker films, instead, no 180° DW movement is observed and the local polarizations in the gradient region are progressively reversed as the gradient strength is enhanced. This work contributes from an atomistic point of view to the understanding of the mechanical flexoelectricity in tuning domains in ultrathin films.

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We have checked that the present atomistic model for PTO is able to reproduce the longitudinal flexoelectric coefficient in the range of nC/m\(^2\).