Phase coexistence and electric-field control of toroidal order in oxide superlattices

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Systems that exhibit phase competition, order parameter coexistence, and emergent order parameter topologies constitute a major part of modern condensed-matter physics. Here, by applying a range of characterization techniques, and simulations, we observe that in PbTiO3/SrTiO3 superlattices all of these effects can be found. By exploring superlattice period-, temperature- and field-dependent evolution of these structures, we observe several new features. First, it is possible to engineer phase coexistence mediated by a first-order phase transition between an emergent, low-temperature vortex phase with electric toroidal order and a high-temperature ferroelectric α/α′ phase. At room temperature, the coexisting vortex and ferroelectric phases form a mesoscale, fibre-textured hierarchical superstructure. The vortex phase possesses an axial polarization, set by the net polarization of the surrounding ferroelectric domains, such that it possesses a multi-order-parameter state and belongs to a class of gyrotropic electrotoroidal compounds. Finally, application of electric fields to this mixed-phase system permits interconversion between the vortex and the ferroelectric phases concomitant with order-of-magnitude changes in piezoelectric and nonlinear optical responses. Our findings suggest new cross-coupled functionalities.

High-temperature superconductivity1, colossal magnetoresistance2,3 and multi-ferroism12–17 (that is, materials simultaneously exhibiting magnetic and ferroelectric order) represent some of the key discoveries of condensed-matter physics in the past few decades. In all cases, emergent phenomena arise from the interplay of spin, charge, orbital and/or lattice degrees of freedom and related coupling that leads to a rich spectrum of competing phases and novel effects4–10. Understanding the nature of spontaneous symmetry breaking and ordering in these phases, and the phase transition pathway under applied stimuli is of fundamental significance. At such phase boundaries, there is often the opportunity to induce colossal field-driven effects (for example, colossal magnetoresistance11,12 in mixed-valent manganites, giant electromechanical responses at morphotropic phase boundaries13–16 and so on). At the same time, considerable interest has been given to exploring new phases with non-trivial, real-space order parameter topologies and emergent continuous symmetry that can occur at the mesoscale (for example, complex spin textures such as vortices and skyrmions17–19).

In the case of ferroelectrics, several theoretical studies20–24 have proposed complex polarization topologies (reminiscent of rotational spin topologies) in low-dimensional structures. The models have predicted the formation of polarization waves, vortices and so on that can be characterized by an emergent order parameter, a so-called electric toroidal moment \( G = \frac{1}{2} \int r \times \mathbf{P}(r) \mathrm{d}r \) (refs 20,22,25). Such toroidal moments can also give rise to a range of novel phenomena (for example, pyrotoroidic, piezotoroidic and so on effects) in response to thermodynamic stimuli. Experimental realization of such structures occurred only recently26, wherein superlattices of (PbTiO3)/(SrTiO3) \( n \approx 12–20 \) unit cells) exhibiting novel polarization-based topologies (namely polarization vortices) were achieved by inducing a competition between electrostatic and strain boundary conditions that inhibit the formation of a homogeneous ferroelectric ground state. The result was nanometre-scale vortex arrays that match the predictions for structures exhibiting an electric toroidal moment.

By exploring superlattice period-, temperature- and field-dependent evolution using phase-field models, second-principle calculations, synchrotron X-ray diffraction, piezoresponse force microscopy (PFM), X-ray photoemission electron microscopy (PEEM)-based linear dichroism, and optical second-harmonic generation (SHG) and SHG nano-imaging, we have discovered that these superlattices exhibit a rich diversity of physical phenomena. Most important is the observation of phase coexistence mediated

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Temperature (°C)

250
350

3

1

DyScO

3

2

θ − 2θ (°)

22.2 22.4 22.6 22.8 23.0 23.2 23.4 23.6 23.8

44 46 47 48

Figure 1 | Structural evolution of ferroelectric and vortex phases with superlattice periodicity. a, High-resolution θ − 2θ scans of 100-nm-thick (PbTiO$_3$)$_n$/(SrTiO$_3$)$_{1−n}$ superlattices grown on DyScO$_3$ (001)$_{pc}$ substrates for $n = 4, 6, 10, 14$ and $16$ (top to bottom, respectively) revealing evolution from a single $a_1/a_2$ alloy peak and corresponding superlattice peaks at short periodicities to mixed-phase films revealing an extra set of alloy and superlattice peaks corresponding to the vortex (V) phase. b, c, Three-dimensional, synchrotron-based symmetric reciprocal space maps about the 002$_{pc}$-diffraction condition of the DyScO$_3$ substrate containing the in-plane [010]$_{pc}$ and out-of-plane [001]$_{pc}$ confirming (b) single $a_1/a_2$ alloy peak and corresponding superlattice peaks at $n = 4$, and (c) mixed-phase coexistence for $n = 16$ showing additional set of alloy and superlattice peaks corresponding to the vortex phase along with the appearance of characteristic satellite peaks corresponding to an in-plane periodicity of ~11 nm along the in-plane [010]$_{pc}$.

d, 2D intensity colour-map summarizing temperature-dependent synchrotron X-ray diffraction θ − 2θ scans about the 002$_{pc}$-diffraction condition of the DyScO$_3$ substrate, and e, a plot of the ratio of diffracted beam intensity from a vortex superlattice peak $(I_{V(−1)})$ to the $a_1/a_2$ superlattice peak $(I_{F(−1)})$ revealing hysteresis indicative of a first-order phase transition from the vortex phase to the $a_1/a_2$ phase at ~200 °C.

by a first-order phase transition between an emergent, low-temperature vortex phase with electric toroidal order and a high-temperature ferroelectric $a_1/a_2$ phase. At room temperature, the coexisting vortex and ferroelectric phases spontaneously assemble in a mesoscale, fibre-textured, hierarchical superstructure. The vortex phase is further found to possess an axial polarization ($P_a$) that is set by the net polarization of the surrounding ferroelectric domains such that it possesses a multi-order-parameter state and is set by the net polarization of the surrounding ferroelectric phase concomitant with order-of-magnitude changes in piezoelectric and nonlinear optical responses and promise for new cross-coupled functionalities.

(PbTiO$_3$)$_n$/(SrTiO$_3$)$_{1−n}$ superlattices ($n = 4, 6, 10, 14, 16$) were synthesized on 5 nm SrRuO$_3$-buffered DyScO$_3$ (110) ((001)$_{pc}$, where pc indicates pseudocubic indices) substrates using established procedures (Methods)$^{28}$. X-ray diffraction θ − 2θ studies about the 002$_{pc}$-diffraction condition of the DyScO$_3$ substrate reveal an interesting evolution in structure with superlattice periodicity (Fig. 1a). Superlattices with $n = 4–10$ reveal a single alloy film peak ($c \approx 3.915$ Å) along with first- and higher-order superlattice reflections (labelled as $F(±m)$). Synchrotron-based three-dimensional (3D)-reciprocal space mapping (RSM) (Supplementary Fig. 1) and high-resolution PFM (Supplementary Fig. 2a) studies show that the single alloy film peak corresponds to a phase with in-plane polarized ferroelectric $a_1/a_2$ stripe domains running along the [110]$_{pc}$ (henceforth referred to as the $a_1/a_2$ phase). Moreover, the $a_1/a_2$ domain width progressively reduces with increasing superlattice periodicity (Supplementary Fig. 2c); indicative of a thickness regime where interface effects dominate and Kittel’s law scaling is invalid.$^{27,28}$

On transitioning from short- to intermediate-period superlattices (that is, $n = 12–18$), where interface effects are less dominant, the $a$ domain size decreases to its lowest value (~6 nm). In addition to the diffraction peaks corresponding to the $a_1/a_2$ phase, X-ray θ − 2θ scans also reveal an extra set of alloy ($c \approx 3.944$ Å, labelled V) and superlattice (labelled V (±m)) peaks indicative of a new phase (Fig. 1a); thus suggesting the possibility of an emergent mixed-phase structure. Additional insights into this new phase are obtained from 3D-RSM studies (Supplementary Fig. 3) and, for brevity, we compare a planar section of the 3D-RSM about the 002$_{pc}$-diffraction condition for the $n = 4$ (Fig. 1b) and $n = 16$ (Fig. 1c) superlattices. Consistent with the θ − 2θ scans, the $n = 16$ superlattices show extra peaks corresponding to the new phase (V) and associated V (±m).
The nanoscale phase separation between ferroelectric and vortex phases in intermediate-period superlattices also manifests itself in PFM (Fig. 2a) as a complex pattern that is dramatically different from the \(a_1/a_2\) twin structure (for example, see the \(n=4\) superlattices; Supplementary Fig. 2a). The lateral piezoresponse image for an \(n=16\) superlattice reveals a disordered order in which alternating stripes exhibit high (chequered white and black) and low or zero (brown) piezoresponse with a periodicity of \(\sim 300 \text{ nm}\) along the \([010]_{pc}\). The high-response regions are the ferroelectric \(a_1/a_2\) phase that exhibits a net polarization along the in-plane \([110]_{pc}\) (from vector PFM studies, Supplementary Fig. 7) and individual domains with widths of only \(6–7 \text{ nm}\) (from X-ray studies, Fig. 1c) and the low-response regions are the vortex structures. This is further confirmed from height modulation of the sample surface since the difference in the unit cell lattice parameters between the vortex (\(c=3.94 \text{ Å}\)) and \(a_1/a_2\) (\(c=3.915 \text{ Å}\)) phases gives rise to systematic variations in the surface topography (Fig. 2b) and correlated changes in the lateral piezoresponse amplitude (Fig. 2c). This correlation is highlighted by line traces along a trench step of the film (dashed lines, Fig. 2h,c) that shows opposite trends in height (green, Fig. 2d) and lateral piezoresponse amplitude (red, Fig. 2d). Ultimately, these data indicate that at room temperature, the coexisting vortex and ferroelectric phases spontaneously assemble in a mesoscale, fibre-textured, hierarchical superstructure.

To understand the microscopic details of how the ferroelectric phase evolves into the vortex phase, we first focus on a single PtTiO\(_2\) layer within the superlattice near the phase boundary as extracted from phase-field models (Fig. 3a). As expected, the \(a_1/a_2\) domains (labelled FE (\(a_1/a_2\)) wherein the colour scale shows the polarization component along the \([100]_{pc}\) exhibit uniform polarization along the \([100]_{pc}\) and \([010]_{pc}\) resulting in a net polarization along the \([110]_{pc}\) (white arrow, Fig. 3a). As one transitions into the vortex phase (labelled Vortex wherein the colour scale corresponds to the magnitude of \(\nabla \times \mathbf{P}\)) the polarization evolves into tracking the evolution of the polarization components (\(P_{[001]}\), \(P_{[001]}\) and \(P_{[001]}\)) as well as the magnitude of the axial electric toroidal moment (\(|\langle G \rangle|\), averaged across the section) as one transitions from the ferroelectric to vortex structure (Fig. 3b). Starting with the ferroelectric phase, we observe constant in-plane polarization components and near-zero (\(|\langle G \rangle|\)). The transition to the vortex phase occurs over \(\sim 20\) unit cells and is characterized by complex polarization rotation pathways (reminiscent of Bloch- and Neel-like walls in ferromagnets) along with a continuously increasing (\(|\langle G \rangle|\) that eventually saturates to a large value of \(0.2 \text{ e Å}^{-1}\) that arises from the continuous rotation of polarization within the vortex structures (blue box, Fig. 3a and in colour-maps of the polarization components \(P_{[001]}\) (Fig. 3c), \(P_{[001]}\) (Fig. 3d) and \(P_{[001]}\) (Fig. 3e) for the same section). We also note that the clockwise and anticlockwise
vortices are arranged in an alternating fashion, such that the toroidal moment $G$ periodically switches direction (akin to what has been called an antiferrotoroidal phase)\textsuperscript{21}.

Also notable is that while the spatial averages of the nonaxial polarization components ($\langle P_a \rangle$, $\langle P_b \rangle$) go to zero within the vortex phase, a sizable axial component ($\langle P_c \rangle \approx 0.1 \text{ C m}^{-2}$) is observed (Fig. 3b); the direction of which is related to the component of net polarization in the adjacent ferroelectric phase. In turn, this produces a screw-like character to the vortex phase and suggests that the vortex structure is characterized by a multi-order-parameter state consisting of axial ferroelectric polarization that coexists with $G$. This is also observed in second-principles calculations (Supplementary Fig. 8) for an $n = 14$ superlattice (Methods and Supplementary Fig. 9) and is consistent with previous predictions for low-dimensional ferroelectrics\textsuperscript{23,30–33}. Such axial polarization can be experimentally observed by focusing on a region of phase coexistence (that is, alternating ferroelectric $a_1/a_2$ and vortex stripes, Fig. 3f). Using PFM, it is again observed that the amplitude of the piezoresponse (Fig. 3g) is higher in the ferroelectric $a_1/a_2$ regions than in the vortex regions, along with the topography, allows for clear demarcation of the regions. The phase of the piezoresponse (Fig. 3h) shows two contrasts across the same area. By combining the amplitude and phase data, the fact that the cantilever axis is aligned along the $[010]_{pc}$ and our understanding of the $a_1/a_2$ structure, the net in-plane polarization for each area can be extracted (arrows, Fig. 3g). Consistent with the phase-field predictions, the axial polarization component of the vortex phase is set by the direction of the polarization in the adjacent $a_1/a_2$ stripe (for example, region I, Fig. 3g,h). There are situations, however, when adjacent $a_1/a_2$ stripes on either side of a vortex region possess opposite net in-plane polarization, and in this case we observe an inversion of the lateral phase signal in the vortex region indicative of a ‘domain-wall-like’ feature (for example, region II, Fig. 3g,h). Either way, both the phase-field models and PFM data suggest that the vortex structures possess an axial component of polarization that is correlated with the polarization in the surrounding ferroelectric phase.

The presence of this axial polarization component is interesting because it suggests the formation of a multi-order-parameter state. In turn, electric-field control of toroidal order is of great interest because it offers the potential for new cross-coupled functions\textsuperscript{20–24}. To explore this concept, the time-dependent evolution of both the
vortex and \(a_1/a_2\) regions of a mixed-phase structure was explored under an applied out-of-plane voltage (15 V) using phase-field models. Specifically, we have collected the evolution of energy and polarization under bias (1,500 time steps) and during relaxation after the bias is turned off (16,000 time steps; where each time step is \(\approx 40\) fs). After turning on the voltage, the various energies (green and red data, respectively, for the vortex and \(a_1/a_2\) phases; Fig. 4a) and polarization states (Fig. 4b–d) are observed to evolve. Starting from the as-grown, mixed-phase state (bias time step 1, Fig. 4a,b), the out-of-plane voltage drives a rotation of the polarization towards the field direction resulting in the electric energy of the system (bias time steps \(\sim 1–100\), Fig. 4a). Within the \(a_1/a_2\) regions, this corresponds to a rotation of the polarization in the applied field direction (not shown) and within the vortex regions (Fig. 4c), the polarization in the direction of the applied voltage is found to be enhanced and that opposite to the voltage is reduced thereby quenching the vortex structure. Both the \(a_1/a_2\) and the vortex structures ultimately trend to the same final state, a uniformly poled, monoclinic state with large out-of-plane polarization (\(\sim 70 \mu C cm^{-2}\)), time steps \(>200\), Fig. 4a,d). We note that this evolution is almost entirely driven by a reduction in the electrical energy that overcomes barriers in the Landau and elastic energies to align the polarization in the field direction.

After turning off the applied bias and allowing the system to relax, a complex evolution of the energies (Fig. 4e) and polarization states (Fig. 4f–h) is observed wherein the entire sample relaxes in a similar fashion. Immediately after turning off the applied bias (time step 1, Fig. 4e), there is a large jump in the electric energy driven by depolarization effects that destabilizes the out-of-plane polarization. In turn, there is a rapid reduction in the electric energy and a corresponding increase in the Landau energy of the system (time steps 1–200, Fig. 4e) due to a reduction of the magnitude of the polarization (from \(\sim 70\) to \(35 \mu C cm^{-2}\)) in the out-of-plane direction. This marks the beginning of a series of events driven by a competition between Landau and elastic energies wherein fast changes in the polarization (that is, the Landau energy) are counteracted by slow changes in the elastic energy as it attempts to reach a new equilibrium (time steps 200–8,000, Fig. 4e). In this range, the Landau energy first begins to dominate and works to restore the total polarization by driving the polarization to increase in the in-plane direction resulting in a fully in-plane polarized monodomain structure (by time step 200–300, Fig. 4e). This in-plane polarized phase, however, forms at the expense of enhanced elastic energy and effectively launches an elastic shockwave that drives twinning in the in-plane polarized phase to create an \(a_1/a_2\)-like structure (by time step 3,000–4,000, Fig. 4e,f). At this point, the system must activate other means to reach equilibrium since it cannot be accomplished with the Landau and elastic energies alone. Thus, small increases in the electric and gradient energies (corresponding to the introduction of a wave-like instability; time step \(\sim 8,000\), Fig. 4e,g) break the oscillatory nature of the Landau and elastic energy variations and drive the formation of the vortex phase across the entire sample (time step 16,000, Fig. 4e,h). Once this phase is formed, no further evolution is observed. Thus, from phase-field simulations, it appears that there is a pathway—albeit a complex one—by which we can not only manipulate the vortex structure, but deterministically control the relative fraction of vortex and \(a_1/a_2\) phases with d.c. electric fields thus making possible effective cross-control of ferroelectric and electric toroidal order that could give rise to a rich spectrum of effects.

To realize this predicted electric-field control of the multi-order-parameter state, we focus on an \(n = 16\) superlattice and explore electric-field evolution of the phases under applied d.c. electric fields using PFM. Beginning with an as-grown superlattice in the mixed-phase form (Fig. 5a), application of a positive electrical bias (via a scanning probe tip within the orange box, Fig. 5a) results in the mixed-phase structure being converted to one with a uniformly low piezoresponse corresponding to the vortex state (Fig. 5b). Similar effects are observed following application of negative bias, which can also give rise to conversion to pure vortex phase (Supplementary Fig. 9). Within this pure vortex region, subsequent application of a negative electrical bias to selected locations (blue lines, Fig. 5a) restores the region back into a mixed-phase structure with similar stripe-like features to the as-grown superlattice (Fig. 5c). Closer inspection reveals that, after applying the negative bias to the tip,
the regions directly under the tip have contrast consistent with the low piezoelectric response of the vortex phase and only adjacent to these written lines (where the in-plane fields from the tip are the strongest) can the high response consistent with the $a_1/a_2$ phase be observed (Supplementary Fig. 10). This change in phase structure is confirmed via synchrotron X-ray nanodiffraction studies (spatial resolution of $\sim 200 \text{nm}$, inset of Fig. 5b,c and Supplementary Fig. 11) and PEEM-based X-ray linear dichroism studies (spatial resolution of $30 \text{nm}$, Supplementary Fig. 12) that provide structural and spectroscopic confirmation, respectively, of the proposed electric-field-induced manipulation of phases. Such reversible electric-field control of the vortex and ferroelectric phases is indicative of a strong coupling between the ferroelectric polarization and electric toroidal moment, the potential for interconversion using a variety of stimuli, and highlights the possibility to control function by tuning the phase fractions. For instance, the SHG response from the as-grown regions (orange data, Fig. 5d) is found to be consistent with that observed for the purely ferroelectric $a_1/\alpha_2$ phase in a single-layer PbTiO$_3$ film (Supplementary Fig. 13). The ability to electrically switch to a purely vortex state results in a strong reduction of the second-harmonic intensity (blue data, Fig. 5d). In turn, realizing that the presence of the vortex phase results in a reduction of SHG intensity and by normalizing the as-grown signal based on the volume of the two phases, we estimate nearly an order of magnitude reduction in the SHG signal in the vortex phase as compared with the ferroelectric phase. Such effects are also resolved at the nanoscale in near-field SHG nano-imaging (inset, Fig. 5d) obtained using shear-force atomic force microscopy-based scanning near-field optical microscopy (Methods) where the ferroelectric and vortex regions provide alternating strong and weak signals, respectively.

All told, these observations have broad-ranging implications for a range of novel physical phenomena and next-generation applications. First, the establishment of a true electric toroidal moment is important since such structures have not been widely observed experimentally. Second, the demonstrated coexistence and reversible electric-field control of ferroelectric and electric toroidal order provides for a number of novel opportunities for applications. In the simplest form, the ability to ‘turn-off’ ferroelectric response via conversion to the vortex phase allows for large tunability and susceptibilities. Potentially more interesting is the possibility to both explore and, ultimately, utilize the nature of emergent phenomena made possible by the presence of the electric toroidal moment including electrotoroidic, piezotoroidic and pyrotoroidic susceptibilities; something that has not been possible to date. Finally, the vortex structure itself also hold tremendous potential for other properties including emergent chirality derived from the screw-like nature of the vortex that could enable electrically switchable optical rotation and dichroism$^{25}$. In fact, it has been shown that the electric-field derivative of the electric toroidal moment$^{22,25}$ is related to the imaginary part of the magnetoelectric susceptibility, and thus can allow for electric-field control of natural optical activity$^{25}$. Ultimately, this work represents an important milestone towards the utilization of complex electrical polarization topologies for next-generation applications.

In summary, this work has demonstrated the possibility to control and engineer phase coexistence between an emergent vortex phase with electric toroidal order and a ferroelectric $a_1/a_2$ phase. This phase coexistence also exhibits classical temperature-dependence (that is, a first-order phase transformation and hysteretic behaviour) and spontaneous self-assembly into a mesoscale, fibre-textured, hierarchical superstructure at room temperature. Closer inspection of the emergent vortex phase reveals an axial polarization, set by the net polarization of the vortex state results in a strong reduction of the second-harmonic intensity (blue data, Fig. 5d). In turn, realizing that the presence of the vortex phase results in a reduction of SHG intensity and by normalizing the as-grown signal based on the volume of the two phases, we estimate nearly an order of magnitude reduction in the SHG signal in the vortex phase as compared with the ferroelectric phase. Such effects are also resolved at the nanoscale in near-field SHG nano-imaging (inset, Fig. 5d) obtained using shear-force atomic force microscopy-based scanning near-field optical microscopy (Methods) where the ferroelectric and vortex regions provide alternating strong and weak signals, respectively.

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Methods

Methods, including statements of data availability and any associated accession codes and references, are available in the online version of this paper.

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Author contributions
A.R.D., J.D.C., R.R. and L.W.M. conceived of the central concepts and designed the experiments. A.R.D., H.L. and M.R.M. conducted the synchrotron and laboratory X-ray diffraction studies. J.D.C. and A.R.D. conducted the scanning probe-based PEM measurements. Z.H. performed and analysed the phase-field simulations. A.K.Y. and M.R.M. synthesized the materials. C.T.N. and S.H.L. performed the TEM-based characterization of the superlattice samples, along with the detailed polarization vector analysis. K.-D.P. and V.K. performed the near- and far-field SHG measurements. A.F. contributed to analysis, discussions and understanding of the data and development of the manuscript. A.R.D., R.R. and L.W.M. wrote the manuscript. All authors discussed the results and implications of the work and read, edited and commented on the manuscript at all stages.

Additional information
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Competing financial interests
The authors declare no competing financial interests.
Methods

Synthesis of superlattices using RHEED-assisted pulsed-laser deposition. Methods for superlattice synthesis have been performed according to previous work and are reproduced here for completeness. Superlattices of (PbTiO
3
)_x/(SrTiO
3
)_y, were grown on 5 nm SrRuO
3
/DyScO
3
(110)_z, [001]_z substrates via pulsed-laser deposition. All depositions were performed at a laser fluence of 1.5 J cm
−2
in an on-axis geometry that was found to provide for stoichiometric transfer of all materials from target to substrate. The SrRuO
3
 was grown at a heater temperature of 700 °C and a dynamic oxygen pressure of 50 mtorr. The superlattices were grown at a heater temperature of 630 °C and a dynamic oxygen pressure of 100 mtorr. The growth was monitored in situ using reflection high-energy electron diffraction (RHEED) and revealed layer-by-layer growth throughout the growth process (Supplementary Fig. 1), thus enabling growth of the superlattices with different periodicities while achieving the same total thickness of 100 nm. RHEED oscillations were monitored for specular spots along the [110]_z of the DyScO
3
(110)_z, substrate. Following growth, the samples were cooled in 50 torr of oxygen to room temperature to promote oxidation.

Structural characterization using laboratory-based X-ray diffraction. High-resolution \( \theta \rightarrow 2 \theta \) symmetric scans using a Pananalytical X’Pert Pro X-ray diffractometer (Cu source) were used to confirm the superlattice period achieved via RHEED-controlled growth. The presence of superlattice peaks, as well as Pendellosung fringes about both the central Bragg and the superlattice peaks (for example in superlattices with periodicity \( p = 6 \times 10^{2} \) [Fig. 1(a)], attest to the high crystallinity of the superlattice structure, and the precise nature and smoothness of the interfaces in these structures.

Structural characterization using synchrotron X-ray diffraction. Synchrotron X-ray diffractography experiments were employed to study the evolution of structure and complex phase coexistence in the superlattice structures. The high flux from a synchrotron source delivered at the Sector 33-BM-C beamline of the Advanced Photon Source, Argonne National Laboratory was optimal to extract lattice modulations associated with the nanoscale ferroelectric domains and vortex structures. A double-crystal monochromator in conjunction with two mirrors was used to obtain a highly monochromatic beam with negligible higher-order harmonics. The excellent accuracy of a Huber 4-circle diffractometer in symmetric scans using a Panalytical X’Pert Pro X-ray diffractometer was optimal to extract lattice modulations associated with the nanoscale ferroelectric domains and vortex structures. The main effects of the stacking are modulations associated with the nanoscale ferroelectric domains and vortex structures. The excellent accuracy of a Huber 4-circle diffractometer in symmetric scans using a Panalytical X’Pert Pro X-ray diffractometer was optimal to extract lattice modulations associated with the nanoscale ferroelectric domains and vortex structures. The main effects of the stacking are modulations associated with the nanoscale ferroelectric domains and vortex structures.

Topography and domain structure characterization. Study of the superlattice structures was carried out using an atomic force microscope (MFP-3D, Asylum Research). Dual a.c. resonance tracking PFM was conducted using a conductive Pt/Ir-coated probe tip (NanoSensor PPP-EFM) to image the domain structures and complete vectorial PFM studies.

Phase-field simulations. Phase-field simulations were performed to simulate the polar structure and switching behaviour of the (PbTiO
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)_x/(SrTiO
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)_y superlattices grown on DyScO
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(110)_z substrates. The spontaneous polarization vector \( P \) is chosen as the order parameter, governed by the time-dependent Ginsburg–Landau equations:

\[
\frac{\partial P(x,t)}{\partial t} = -L \frac{\partial E}{\partial P(x,t)} \quad (i = 1, 2, 3)
\]

(1)

where \( i, L \) and \( F \) denote the evolution time step, kinetic coefficient, and total free energy of the system, respectively. The free energy has contributions from the Landau, elastic, electric (electrostatic) and gradient energies, that is,

\[
F = \int \left( E_{\text{elastic}} + E_{\text{electric}} + E_{\text{friction}} + E_{\text{friction}} \right) \text{d}V
\]

(2)

The detailed expressions of these energies can be found in previous reports. A 3D mesh of 200 × 200 × 250 is used, with each grid representing 0.4 nm, the thickness dimension (from bottom to top) is made up of 30 layers of substrate, 176 layers of film and 22 layers of air. In the film layer, 9 unit cells of PbTiO
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, and 9 unit cells of SrTiO
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, as well as 1 layer of transition layer between in the PbTiO
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 and SrTiO
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 layers are deposited periodicaly. Periodic boundary conditions are applied along the x and y direction and the mixed elastic boundary condition is used in the thickness dimension. Fixed potential electric boundary conditions are applied where the electric potentials on the bottom and top are fixed to 0 and the applied bias, respectively. During the poling process, 15 V is applied on the top electrode. A mixed elastic boundary condition is used where the displacement on the bottom of the substrate is zero while the out-of-plane stress on the film top is free. All of the material constants are adopted from previous literature. Random noise with small magnitude (<0.1 μC cm
−2
) is added as the initial nuclei.

Second-principles simulations. Methods for second-principles simulations have been performed according to previous work and are reproduced here for completeness. Interactions within the PbTiO
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 or SrTiO
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 layers are addressed with previously introduced potentials for the bulk compounds. These potentials give qualitatively correct descriptions for the lattice dynamical properties and structural phase transitions of both materials. A simple numerical average was used for interactions of the ion pairs touching or crossing the interface. Within the superlattice periodicities studied in this work, the main effects of the stacking are electrostatic wherein long-range dipole–dipole interactions are governed by the Born effective charges of the parent compounds and a bare electronic dielectric constant \( e_{\infty} \) is taken as a weighted average (with weights reflecting the composition of the superlattice periodicity \( n \)). To preserve the electrostatic energy within each material as close as possible to the bulk parent compounds, the Born effective charge tensors of the inner atoms were rescaled by \( \sqrt{\frac{1}{n}} e_{\infty} \) (where \( ABO \) stands for PbTiO
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 or SrTiO
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 depending on the layer) and, as noted in equation (23) of ref. 44, the dipole–dipole interactions were maintained to be the same as bulk. For atoms touching the interfaces, the Born effective charge tensors were left untouched. In plane lattice constants of \( a = b = 3.901 \) Å were assumed. An external expansive hydrostatic pressure of \(-11.2 \) GPa was imposed to counterunderestimation of the local density approximation of the cubic-lattice constant that was taken as the reference structure for the fitting of the second-principles parameters. Such approximations and adjustments allow for the construction of models for superlattices of arbitrary \( n \) stacking. Simulations applied periodically repeated supercells containing \( 2 \times 2 \times n \) elemental ferroelectric units in-plane, an one full supercell period in the out-of-plane direction.

Local polarizations were computed within a linear approximation of the product of the Born effective charge tensors times the atomic displacements from the reference structure positions divided by the volume of the unit cell. The effect of external electric fields was incorporated in the effective Hamiltonian simulation by including the leading coupling term between the field and the local dipoles. The models were solved via Monte Carlo simulations comprising 10,000 thermalization sweeps followed by 40,000 sweeps to compute thermal averages. If the Metropolis Monte Carlo samplings are performed at low temperature, then only random configurations along the Markov chains that produce a lowering in the energy are accepted, so this procedure can be used as an equivalent to structural relaxations to find the ground state.

Conventional and scanning transmission electron microscopy. Methods for TEM-based studies have been performed according to previous work and are reproduced here for completeness. Samples for TEM studies were produced via mechanical polishing on an an Aligned High Tech MultiPrep. Cross-sectional samples were polished at a 0.5° wedge, and planar samples with a flat polish. Following polishing, the samples were Ar-ion milled using a Gatan Precision Ion Milling System from a starting energy of 4 keV to a final cleaning energy of 200 eV.

Scanning TEM was performed on a Cs-corrected TEAM0.5 FEI Titan microscope at 300 kV using a high-angle annular detector resulting in ‘Z-contrast’ images. Raster distortion was minimized using a pair of HR-STEM images of the same region taken with orthogonal scan orientations. These orthogonal scan orientation images were used to calculate a corrected ‘pre-distortion’ image to compensate for the slow scan axis. Sampling and extraction of the high-angle annular dark-field images using local A- and B-site sublattice offset measurements. This process is based on the fact that for many displacive perovskite ferroelectrics the electric dipole moment is manifested by relative shifts of the cations, as is the case for PbTiO
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, and this offset can be used to infer the polarization (P). This is helpful since the anions (oxygen) have relatively weak electron scattering resulting in low signal-to-noise ratios. Beam sampling and extraction of the high-angle annular dark-field images, limiting the impact they can have on characterizing the electric dipole. In bulk PbTiO
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, at room temperature, the Ti and O ions are shifted by 16.2 pm and 47–49 pm in the direction opposite to the polarization, respectively, relative to the Pb lattice. In turn, we can use the Ti ions to infer the offset of the Ti-centred oxygen octahedra, which can be used to determine the electric dipole. Displacement vectors \( (P_{n}) \) corresponding to local offsets between the A- and
B-site sublattices were calculated in the following three steps. First, the atomic positions were determined by fitting each atom site in the image to four-parameter, spherical Gaussians using a trust-region algorithm in Matlab. Such fits were performed independently for five-atom clusters of A-site-centred perovskite unit cell. Second, the displacement for each atom is then calculated as the difference between its atom position and the mean position (that is, the centre of mass) of the surrounding four opposite-type cations. Using a pseudocubic grid indexing (wherein A-site defines the grid) and defining the cross-section image to lie in the x–z Cartesian plane (x ≈ [100], z ≈ [001]₀), for an atom at grid position i this corresponds to mean neighbour positions (MNP) of:

\[
\text{MNP}_{x,i} = \frac{1}{4} [x_{i} - x_{i+\frac{1}{2}} + x_{i+\frac{1}{2}} + x_{i+1} - x_{i-\frac{1}{2}} - x_{i} + x_{i-\frac{1}{2}}],
\]

\[
\text{MNP}_{z,i} = \frac{1}{4} [z_{i} - z_{i+\frac{1}{2}} + z_{i+\frac{1}{2}} + z_{i+1} - z_{i-\frac{1}{2}} - z_{i} + z_{i-\frac{1}{2}}].
\]

Finally, the polar displacement vector is defined with reversed order for A-site and B-site-centred atoms to maintain a consistent displacement vector sign (otherwise it would alternate directions):

\[
P_{2 \beta,\text{tot}} = (x_{i} - \text{MNP}_{x,i}) \hat{x} + (z_{i} - \text{MNP}_{z,i}) \hat{z}.
\]

This convention defines a displacement vector direction nominally parallel to the full electrical polarization of the bulk crystal structure.

The curl vector (or vorticity) of the displacement vector field \( P_{\text{tot}} \) in the TEM plane is the \( y \) component of \( \nabla \times P_{\text{tot}} \) or \( (\partial P_{\text{tot},z}/\partial x) - (\partial P_{\text{tot},x}/\partial z) \) (again the TEM image is defined as the \( x-z \) plane, \( y \) is the direction of the electron beam). The polarization gradient terms \( \partial P_{\text{tot},z}/\partial x \) and \( \partial P_{\text{tot},x}/\partial z \) were calculated by a weighted least-squares fit of a two-variable \((x \text{ and } z)\) first-order polynomial to \( P_{\text{tot},x} \) and \( P_{\text{tot},z} \) at each atom site with a fit area extending to 12 surrounding atoms. Said a different way, the local \( P_{\text{tot},x} \) and \( P_{\text{tot},z} \) values were fitted as a plane, and the derivatives are taken as the \( x \) and \( z \) slopes of the plane. A Gaussian function with a \( \sigma \) of the mean grid spacing was used as the weighting function; that is, the central atom was given a weight of 1 and decreased to a weight of \( \approx 0.1 \) for the most distant of the 12 neighbour atoms considered. The polynomial fit was used as a smoothing function for the vector fields displayed in the figures, effectively smoothing the first derivative, and the gradients input into the curl equation.

**Photoemission electron microscopy (PEEM)-based X-ray linear dichroism.** X-ray linear dichroism in the near-edge X-ray absorption fine structure senses the local anisotropy of the electron charge cloud, and is sensitive to the local ferroelectric polarization axis. PEEM images and spectra were taken at the Ti L₂,₃ edge using s- (along [100]) and p- (30° from [001]) polarized X-rays. Vortex and ferroelectric regions are distinguished in X-ray linear dichroism PEEM images mainly by their different [100] polarization components.

**Second-harmonic generation (SHG) and SHG nano-imaging.** A typical reflection mode geometry was used the far-field SHG anisotropy measurement, with combinations of input and output polarizations selected by an achromatic half-wave plate and a polarizer to probe specific tensor components. The backscattered SHG intensity was then measured while keeping the polarization optics stationary and rotating the sample. For near-field SHG nano-imaging, a shear-force atomic force microscopy-based scattering near-field optical microscopy set-up was used in a tilted tip geometry as described previously. The Au tip was illuminated from the top by broadband ~10 fs pulses derived from a Ti:sapphire oscillator (Femtolasers, 800 nm centre wavelength, 78 MHz repetition rate), with tip-parallel polarization and average incident power of 3 mW. The tip-enhanced SHG signals were collected by a microscope objective (100 ×, NA = 0.8) in the back-reflection geometry and detected with a spectrometer and a CCD (charge-coupled device; Princeton Instruments).

**Data availability.** All relevant data are available from the authors, and/or are included with the manuscript as source data or Supplementary Information including that in Figs 1–5 and Supplementary Figs 1–13.

**References**


