Direct Observation of the Proliferation of Ferroelectric Loop Domains and Vortex-Antivortex Pairs

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We discovered stripe patterns of trimerization-ferroelectric domains in hexagonal REMnO₃ (RE = Ho, \cdots , Lu) crystals (grown below ferroelectric transition temperatures (T_c), reaching up to 1435 °C), in contrast with the vortex patterns in YMnO₃. These stripe patterns roughen with the appearance of numerous loop domains through thermal annealing just below T_c , but the stripe domain patterns turn to vortex-antivortex domain patterns through a freezing process when crystals cross T_c even though the phase transition appears to not be Kosterlitz-Thouless-type. The experimental systematics are compared with the results of our six-state clock model simulation and also the Kibble-Zurek mechanism for trapped topological defects.

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The subtle phase transitions straddling the boundary between long-range order and disorder have attracted significant attention due to the fundamental science and the technological perspective [1–5]. For example, the ground state of two-dimensional (2D) spins with planar continuous degrees of freedom undergoes only quasi-long-range order with spin-spin correlations falling off algebraically in space, and becomes a high-temperature disorder state with exponentially decaying correlations through the so-called Kosterlitz-Thouless (KT) transition by the proliferation of unbound topological vortices [6–8]. Not only the KT transition in 2D planar spins but also various phenomena such as 2D melting and roughening transition at surface are associated with the emergence of a topological order, resulting from the binding of topological defects [9,10]. Even though the ordering issue of 2D condensed matters is a time-honored topic, the topological ordering process in "large-scale real space" has been little investigated experimentally. Furthermore, it is a profound question how the topological KT order is influenced by the 3rd-dimensional coupling and quenched disorder that exists often in real systems.

Layered hexagonal YMnO₃ is an improper ferroelectric where the size mismatch between Y and Mn-O layers induces a trimerization-type structural phase transition, and this structural transition leads to three antiphase domains (α, β, γ) , each of which can support two directions (+, -) of ferroelectric polarization [11–13]. The antiphase and ferroelectric domains of YMnO₃ meet in cloverleaf arrangements that cycle through all six domain configurations. Occurring in pairs, the cloverleaves can be viewed as vortices and antivortices, in which the cycle of domain configurations is reversed [14]. Large-scale arrangements of topological vortices and antivortices in

YMnO₃ reveal intriguing real space domain patterns with mathematical simplicity, which can be analyzed with graph theory [15]. The six possible characteristics of domains, combined with the layered structure of hexagonal YMnO₃, suggest the analogy between the 2D six-state clock model and the physics of YMnO₃. In the six-state clock model, the presence of three phases has been claimed: long-range ordered (LRO) phase, intermediate KT phase, and high-temperature disordered phase [16]. However, vortex-antivortex domain patterns have been observed in YMnO₃ at room temperature far below the structural transition temperature, which suggests the KT phase, rather than a LRO phase, as the ground state.

Herein, in order to unveil the origin of this inconsistency, we have studied the systematics of domain configurations in a series of hexagonal REMnO₃ (RE = rare earths) crystals grown with a flux method, and also the thermal evolution of the domain configurations. In order to reveal domain patterns, thin-plate-like crystals with optically flat surfaces were etched chemically in phosphoric acid at ~130 °C. The domain patterns of chemically etched crystals were investigated using optical microscopy (OM), transmission electron microscopy (TEM) and atomic force microscopy (AFM). Note that preferential chemical etching of surface areas with upward polarization enables the observation of ferroelectric domain patterns on a crystal surface using OM or AFM. (See the detailed experimental method in the Supplemental Material [17].)

Unlike vortex-antivortex domain patterns in YMnO₃, stripe domain patterns with large downward (-) polarization domains are discovered in most of REMnO₃ crystals. The OM image of a stripe domain pattern on one entire surface of an ErMnO₃ crystal is shown in Fig. 1(a). These stripe lines in the OM image are identified as narrow

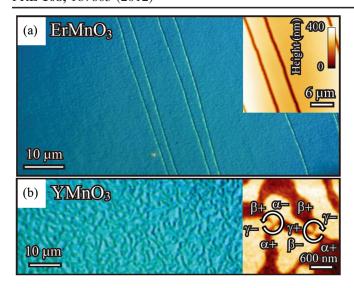


FIG. 1 (color online). Two distinct domain patterns of REMnO₃ crystals: stripe vs vortex patterns. (a) Optical microscope image of a chemically etched ErMnO₃ crystal surface. The inset shows the atomic force microscope (AFM) image of a stripe domain pattern after chemical etching. These domain patterns for a long-range ordered phase are also observed in HoMnO₃, TmMnO₃, YbMnO₃, LuMnO₃ crystals, as shown in Fig. S1 in the Supplemental Material [17]. (b) Optical microscope image of a chemically etched YMnO₃ crystal surface. The inset shows the AFM image of a YMnO3 surface after chemical etching, showing a vortex-antivortex domain pattern. Vortex and antivortex are distinguished by the arrangement of trimerization antiphase and ferroelectric domains with the opposite sense of rotation around a core. Note that the entire surfaces of all as-grown crystals REMnO₃ (RE = Ho, Er, Tm, Yb, Lu) exhibit no hint of the presence of any vortices or antivortices.

trenches with a depth of ~500 nm as revealed in AFM scans (see the inset of Fig. 1(a)], and tend to be along the [110] direction (the hexagonal $P6_3cm$ notation). These trenched lines correspond to narrow upward (+) polarization domains [15]. These stripe domain patterns are distinct from the topological vortex-antivortex domain pattern with small domains in YMnO₃, shown in Fig. 1(b). In the vortex-antivortex domain pattern in YMnO₃, a vortex consists of six trimerization antiphase (α, β, γ) and ferroelectric (+, -) domains merging at the center of the vortex, and is paired with an antivortex (or antivortices) with the opposite vorticity in terms of structural antiphase and ferroelectric relationship as shown in the inset of Fig. 1(b) [15]. On the other hand, the stripe domain pattern in Fig. 1(a) spans the entire crystal surface, and we have, in fact, observed only these stripe domain patterns without any hint of the presence of vortices in all REMnO₃ (RE = Ho, Er, Tm, Yb, Lu, but not Y) crystals. We also note that the stripe domain patterns with large downwardpolarization domains appear to be consistent with an LRO phase, rather than the topological KT phase. As discussed below, this striking difference between stripe and vortex-antivortex domain patterns depends entirely on whether the crystal growth temperature is above the trimerization-structural transition temperature (T_c) or not.

In order to explore the thermal evolution of these stripe domain patterns, we cooled down crystals from various annealing temperatures to room temperature and then etched them chemically. Figure 2(a) displays the AFM image of a chemically etched ErMnO₃ surface after cooling it down fast from 1120 °C. Even though the equilibrating temperature is very high, the LRO stripe domain pattern changes little. However, when the temperature is raised by only 20 °C, the pattern changes significantly, and exhibits highly curved lines with the appearance of many curved closed loops, as shown in Fig. 2(b). Dark stripe lines, conserved robustly up to 1120 °C, start to wiggle heavily at 1140 °C, but never cross to each other; i.e., there is no hint of the presence of vortices in the entire crystal

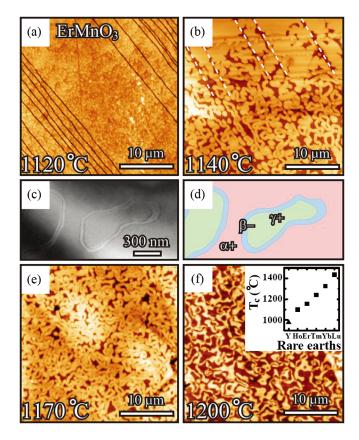


FIG. 2 (color online). The thermal evolution of stripe domain patterns in ErMnO₃. (a),(b),(e), and (f) The AFM images of chemically etched ErMnO₃ crystals annealed at 1120 °C, 1140 °C, 1170 °C, and 1200 °C, respectively. A stripe domain pattern with thermal roughening are evident in (b), but (e) and (f) display vortex-antivortex patterns. The inset displays T_c 's of REMnO₃ estimated from the formation temperature of vortex-antivortex domain patterns. T_c of YMnO₃ is from Ref. [18]. (c) and (d) TEM image of loop domains of ErMnO₃ quenched from 1140 °C and a possible corresponding schematic, respectively. The loop domains observed in the TEM image are assigned with three antiphase domains ($\alpha + , \beta - , \gamma +$).

surface (see the Supplemental Material, section 2 and Fig. S2 [17]). The more-or-less straight parts of dark stripe lines are indicated with white dashed lines in the upper region of Fig. 2(b). A TEM image of closed loop domains is shown in Fig. 2(c), and the corresponding possible schematic is shown in Fig. 2(d). Therefore, when the system approaches T_c from below, thermal fluctuations induce roughening of the stripe domain walls and the appearance of a large number of loop domains.

When the annealing temperature was further raised by 30 °C up to 1170 °C, a complicated pattern was observed after chemical etching as shown in Fig. 2(e). This pattern, in fact, shows the crossing of lines and the presence of a large number of vortices. This vortex-antivortex domain pattern formation was more evident when the annealing temperature was raised to 1200 °C, as shown in Fig. 2(f). We have determined the characteristic temperatures for all REMnO₃ (RE = Ho, Er, Tm, Yb, Lu) at which stripe domain patterns turn into vortex-antivortex domain patterns after the annealing experiment. The obtained characteristic temperatures are plotted in the inset of Fig. 2(f), and the reported T_c of YMnO₃ is also plotted in the inset [18]. The rough linear dependence in the inset strongly suggests that the characteristic temperatures are, indeed, the trimerization-structural T_c of REMnO₃. Note that T_c 's of REMnO₃ (RE = Ho, Er, Tm, Yb, Lu) have never been reliably determined because of the very high temperature nature. We also point out that the drastic increase of T_c with decrease of RE size is consistent with the notion that the structural transition is induced by the mismatch between small RE layers and large Mn-O layers in the REMnO₃ structure. Note that REMnO₃ crystals were grown by slow cooling of the materials with Bi₂O₃ flux in the temperature range of 1200 °C and 950 °C, but the real growth through nucleation occurs probably slightly above 950 °C. Thus, YMnO₃ crystals are likely grown above T_c , but other REMnO₃ crystals below T_c . Therefore, it appears that stripe domain patterns form when the crystal growth temperature is below T_c , while vortex-antivortex domain patterns are realized when crystals are exposed to temperatures above T_c .

Interestingly, we found that once vortex-antivortex domain patterns, spanning the entire crystal surface (see the Supplemental Material, section 3 and Fig. S3 [17]), form by crossing T_c , they are conserved with various thermal treatments, but the domain size of vortex-antivortex domain patterns or the distance between vortices and antivortices can vary in a systematic manner. In order to find out the thermal evolution of vortex-antivortex domain patterns and the domain growth kinetics, the cooling rate near T_c was changed from $0.5\,^{\circ}$ C/h to $300\,^{\circ}$ C/h. In addition, we cooled one specimen from $1220\,^{\circ}$ C to $677\,^{\circ}$ C with a cooling rate of $5\,^{\circ}$ C/h, followed by quenching to room temperature. Figures 3(a)–3(d) show the AFM images of etched ErMnO₃ crystals with the above thermal

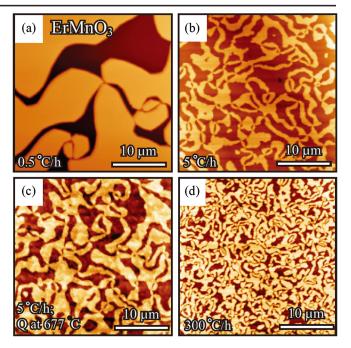


FIG. 3 (color online). The evolution of vortex-antivortex domain patterns with varying cooling rate. The AFM images of chemically etched ErMnO₃ crystals: (a) cooled from 1220 °C to 890 °C with a rate of 0.5 °C/h, followed by furnace cooling, (b) cooled from 1200 °C to room temperature with a rate of 5 °C/h, (c) from 1200 °C to 677 °C with a rate of 5 °C/h, followed by quenching, and (d) from 1200 °C to room temperature with a rate of 300 °C/h.

treatments. With the large variation of cooling rate from 0.5 °C/h to 300 °C/h, vortex-antivortex domain patterns remain intact, but the domain size of vortex-antivortex domain patterns changes systematically.

We emphasize that the vortex-antivortex domain patterns in Figs. 3(b) and 3(c) are basically identical, indicating that the cooling rate below 677 °C does not influence the domain patterns. This is an important result for the origin of the mysterious second transition of REMnO₃ near 600 °C reported in many early publications [12,18–25]. This second transition at ~600 °C was identified as the ferroelectric transition from centrosymmetric $P6_3/mmc$ to low-temperature polar P6₃cm structures via an intermediate $P6_c/mcm$ structure [19,20] whereas other results argued for the nonexistence of intermediate $P6_3/mcm$ state, but the presence of an isosymmetric phase transition with Y-O hybridization [12,18,21,22,24]. We, in fact, directly compared vortex-antivortex domain patterns at room temperature and 730 °C from TEM dark-field experiments using the 131 diffraction spot as shown in Figs. 4(a) and 4(b). Basically there is little difference between two vortex-antivortex domain patterns, which, combined with no difference in vortex-antivortex domain patterns on the cooling rate across the second transition temperature, are consistent with the possibility of an isosymmetric change at the second transition if it exists.

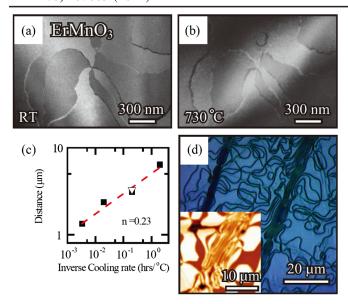


FIG. 4 (color online). The pinning of vortex-antivortex domains. (a) and (b) TEM images of vortex-antivortex domain patterns of ErMnO₃ at room temperature and 730 °C, respectively. (c) The cooling rate dependence of the average distance of vortex-antivortex pairs, exhibiting a power law dependence with the power of 0.23. [The open triangle is from Fig. 3(c).] (d) The optical microscope and AFM (inset) images of a chemically etched ErMnO₃ crystal surface, suggesting the strong pinning of vortex-antivortex domains by surface defects.

The analysis of the evolution of domain size of vortexantivortex domain patterns with varying cooling rate demonstrates slow growth kinetics associated with the topological vortex-antivortex domain patterns. The average distance of vortex-antivortex pairs vs inverse cooling rate, t, is plotted in Fig. 4(c). The cooling rate dependence of the average vortex-antivortex pair distance, D, can be described by $D \propto t^n$ with n = 0.23 (see the Supplemental Material, section 4 and Fig. S4 [17]). This value of $n \approx$ 1/4 is rather different from the typical parabolic domain growth value of n = 1/2 [26,27]. This slow growth kinetics with vortex-antivortex domain patterns seems consistent with the Kibble-Zurek mechanism for spontaneous trapping of topological defects in a system undergoing a continuous phase transition [28,29]. For the average vortex-antivortex pair distance $\propto t^n$ with $n \approx 1/4$, the density of vortex should vary like $t^{1/2}$, which is precisely the prediction of the Kibble-Zurek mechanism with mean field critical exponents [29]. We also note that chemical- or structural-defect-induced pinning may play an important role for the variation of the vortex-antivortex distance with cooling rate. In fact, a strong pinning tendency of domain patterns has been observed in many of our results. For example, the domain patterns in Figs. 2(a) and 4(b) change little with expected large thermal fluctuations at temperatures such as 1120 °C and 730 °C. In addition, we have directly observed vortex-antivortex domain patterns pinned by large-scale surface defects, as shown in Fig. 4(d). Strong pinning, slow kinetics, and the different topology between vortex-antivortex and stripe domain patterns probably induce an "astronomical" time scale for the conversion of vortex-antivortex domain patterns to stripe domain patterns, even though the true ground state corresponds to stripe domain patterns.

The concept of topological KT order with binding vortices and antivortices was originally developed for the quasi-long-range order in 2D XY systems. In the KT phase, algebraically decaying correlation can accompany topological defects, i.e., vortices. As the order parameter possesses a finite value, q, representing the number of evenly spaced possible order orientations, there exists a longrange ordered phase at low temperatures when q is small. Convincing evidence has been amassed, based both on qualitative and quantitative considerations, to indicate that for large enough q (thought to be $5 \le q < 8$), there exists an intermediate phase similar to the KT phase [16]. The 2D q-state clock model with $q \ge 8$ precisely reproduces the KT transition [30,31]. In the case of the 2D sixstate clock model, q = 6, the topological order exists only in an intermediate temperature range, and the ground state is a LRO phase (see the Supplemental Material, section 5 and Fig. S5 [17]). Any 3rd-dimensional coupling will destabilize the intermediate KT phase in the six-state clock model. Therefore, one expects theoretically that a LRO phase will be the ground state of hexagonal REMnO₃ with 6 degrees of freedom, and the intermediate KT order may or may not exist, depending on the strength of 3rd-directional coupling. Experimentally, our results indicate that the LRO ground state with stripe domain patterns could be observed, but only when crystals were grown below T_c . When crystals were grown from temperatures above T_c or crystals with stripe domain patterns were exposed to a temperature above T_c , domain patterns were always vortex-antivortex domain patterns. In addition, we have observed convincing evidence for strong pinning and also slow kinetics associated with domain patterns. Furthermore, our results do not show any indication for two step transitions from a long-range order to a KT order to disorder. Therefore, it is convincing that there exists only one LRO transition associated with 6 degrees of freedom. However, in any real situations without astronomical-time-scale annealing, slow kinetic creation of topological defects, which is well manifested in the Kibble-Zurek mechanism, and their strong pinning induce KT-type vortex-antivortex domain patterns when crystals undergo the structural transition at T_c . In other words, a KT order is only implicit in hexagonal REMnO₃ and should not occur as the ground state, but it is arrested by slow kinetic and strong pinning. This "arresting" scenario is fully consistent with our simulation of the 2D six-state clock model with different cooling rates (see the Supplemental Material, section 5 and Fig. S5 [17]).

In summary, the true ground state with stripe domain patterns can be realized when hexagonal $REMnO_3(RE = Ho, Er, Tm, Yb, Lu)$ crystals are grown below T_c , and is consistent with the long-range ordered ground state of the six-state clock model with a significant 3rd-dimensional coupling. When crystals cross T_c , domain patterns with topological vortices and antivortices are realized. These vortex domains can be enlarged with the time exponent of 1/4, which is much smaller than the typical domain growth exponent of 1/2. It is conceivable that the topological KT-type vortex-antivortex domain patterns eventually turn into long-range-ordered stripe domain patterns through astronomical-time-scale annealing. Therefore, the presence of stripe domain patterns in REMnO₃ crystals grown below T_c is an experimental marvel, enabling the observation of what happens in the true ground state of REMnO₃.

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