Dynamical Multiferroicity and Magnetic Topological Structures Induced by the Orbital Angular Momentum of Light in a Nonmagnetic Material

Lingyuan Gao[®], Sergei Prokhorenko, Yousra Nahas, and Laurent Bellaiche^{*}

Physics Department and Institute for Nanoscience and Engineering, University of Arkansas, Fayetteville, Arkansas 72701, USA

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Recent studies have revealed that chiral phonons resonantly excited by ultrafast laser pulses carry magnetic moments and can enhance the magnetization of materials. In this work, using first-principlesbased simulations, we present a real-space scenario where circular motions of electric dipoles in ultrathin two-dimensional ferroelectric and nonmagnetic films are driven by orbital angular momentum of light via strong coupling between electric dipoles and optical field. Rotations of these dipoles follow the evolving pattern of the optical field and create strong on-site orbital magnetic moments of ions. By characterizing topology of orbital magnetic moments in each 2D layer, we identify the vortex type of topological texture— magnetic merons with a one-half topological charge and robust stability. Our study thus provides alternative approaches for generating magnetic fields and topological textures from light-matter interaction and dynamical multiferroicity in nonmagnetic materials.

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Spin, as an intrinsic angular momentum of electron, possesses a magnetic dipole moment and has no counterpart in classical mechanics [1]. Spin can be in one of two states, corresponding to either an "up" or "down" magnetic field, leaving it as the main source of ferromagnetism. As a contrast, orbital angular momentum of electrons, resembling orbital motion of classical charges, contributes to another part of the magnetic dipole moment. The definition of orbital angular momentum has gone beyond the motion of electrons and been extended to other charged systems, such as rotating molecules [2,3].

Recently, many efforts have been made in studying effective magnetization caused by ionic motions in solids: as a duality to electric polarization induced by spatially varying spin textures [4,5], emergent magnetization results from the temporally varying electric polarization $M \propto P \times$ $\partial_t \mathbf{P}$ [6–11]. Such dynamical polarization can be produced by driving collective motions of ions, usually termed as "phonons" in solid-state language. The cross product between **P** and its time-derivative $\partial_t \mathbf{P}$ further requires ions to move on a closed orbit and to carry an orbital angular momentum, and that corresponds to chiral or circularly polarized phonons [12-17]. In reciprocal space, chiral phonons are located at high symmetry points either at the center or the boundary of the Brillouin zone [14,17-20]. At the zone center, chiral phonons exist as a superposition of two orthogonal, degenerate phonon modes; while at the zone edges, recent experiments show that nondegenerate chiral phonons are observed in crystals with hexagonal symmetries [14,19].

Moreover, numerous studies show that light can control the magnetization in crystals by driving optical phonon modes and tuning structural distortions [21–28]. At the zone center, infared (IR)- and Raman-active modes, related to electric polarization and polarizability, respectively, can be excited with an ultrafast terahertz (THz) laser pulse. Specifically, a degenerate chiral phonon can be excited by a circularly polarized light [17,29]. At the zone edge, chiral phonon is not permitted to be excited by light, due to the large mismatch between momenta of phonons and photons. Recent studies predicted and demonstrated that cerium trihalides are capable of generating strong effective magnetic fields by exciting chiral phonons with circularly polarized THz pulses [28,30].

In this work, we propose an alternative approach to induce effective magnetic field established on the mechanism of dynamical multiferroicity. Instead of focusing on coherently excited chiral phonons, we drive circular motions of ions with a particular type of light-optical vortex (OV) beam. Such light carries an orbital angular momentum (OAM) [31-33], and can exert OAM to manipulate the motion of microparticles as an optical tweezer [34-36]. Here we demonstrate that OAM can be transferred from the light to the motion of ions via the coupling between electric dipoles and optical field. As a prototype example, we use first-principles-based simulations and demonstrate that in ultrathin ferroelectric Pb(Zr, Ti)O₃ films, microscopic electric dipoles indeed rotate with the time-varying OAM field. As a result, orbital magnetic moments of ions are effectively generated resulting in a non-negligible magnetic field even in nonmagnetic materials. This differs from previously proposed mechanisms in that in a confined system, the real-space circular motion of ions at each site directly follows the pattern of the optical field, rather than move along the phonon eigenvectors. Here the resulting displacements have much larger amplitudes than the displacements caused by phonons, which result in much larger dynamical magnetic moments. This largely broadens the range of candidate materials restricted by crystal symmetry as well as light frequencies, so that the effect of dynamical multiferroicity can be tuned more freely. More intriguingly, at specific times, microscopic magnetic moments on a single layer of this quasi-2D system are arranged in a vortex-type configuration, proving topological magnetic structures—magnetic merons—can also be produced out of orbital magnetic moments of ions and dynamical multiferroicity.

We perform first-principles-based effective Hamiltonian $(H_{\rm eff})$ molecular dynamics (MD) calculations to conduct this time-dependent study [37,38]. $H_{\rm eff}$ represents loworder energy expansions in structural distortions relative to a highly symmetric configuration, and it depends on the following specific variables: local soft mode u_i where *i* refers to the atomic site index, homogeneous strain tensor η , and inhomogeneous strain variable v_i . All parameters of $H_{\rm eff}$ are fitted from first principles from calculations, especially for the acoustic phonon and soft optical phonon branches that are most responsible for structural distortions during phase transitions. The effect of external electric field $\boldsymbol{E}(\vec{r})$ is incorporated by including a $-\sum \boldsymbol{p}_i \cdot \boldsymbol{E}_i$ term in the $H_{\rm eff}$ energy [39,40], where p_i is the local electric dipole moment that is directly proportional to the local soft mode u_i by a factor of Born effective charge Z^* . The effective Hamiltonian has been well developed for Pb(Zr, Ti)O₃ systems and has made a series of successful predictions about topological structures and phase transitions, which were then confirmed by experiments [41-49]. Here, we model $Pb(Zr_{0.4}Ti_{0.6})O_3$ (PZT) ferroelectric ultrathin films grown along the [001] direction with an $80 \times 80 \times 8$ supercell, which is periodic along the two in-plane directions [100] and [010] but finite along the out-of-plane [001] direction with one substrate layer (layer 1) on the bottom and two vacuum layers on the top (layers 7, 8). Note that the local heating effect from field with large amplitudes could also reshape the structural textures, but at a much slower speed [50,51]. Furthermore, with a current set of parameters, we estimate the increasing temperature brought by the OV beam and find that the heating effect is negligible (see Supplemental Material [52], which includes Refs. [53–81]).

The OV beam is introduced as the lowest order (l = 0) of the "Laguerre-Gaussian" mode [31,82], and the electric field is written as

$$\vec{E}(\vec{r},t) = E_0 \left(\frac{\sqrt{2}\rho}{w}\right)^{|m|} e^{-\frac{\rho^2}{w^2}} (\cos(m\phi - \omega t)\vec{e}_x - \sigma\sin(m\phi - \omega t)\vec{e}_y).$$
(1)



FIG. 1. Illustration of ferroelectric PZT films illuminated by OV beam. (a) OV beam with a helical wavefront is normally incident on PZT films. Current loops form at each site due to the rotation of electric dipoles, which induce emergent magnetic moments (black arrows pointing up). (b) The distribution of inplane components of local soft mode vectors $u_{x,y}$ (denoted by arrows) on each site (gray sphere) at t = nT after the system establishes cyclic motion. We draw polar coordinates with its origin coinciding with the center of the (001) plane to denote the thin film plane and three circles in radius of 0.5w, w, and 2w, colors of which indicate the variation of the *E* field intensity along the radial direction. The bottom part depicts the change of the dipole's orientation with time at a selected site.

In the equation above, \vec{e}_x and \vec{e}_y are two light polarization vectors pointing along the [100] and [010] in-plane directions, respectively. E_0 and ω denote the magnitude and frequency of the electric field, respectively; considering the response time of electric dipoles is typically at the picosecond level, we set $\omega = 1$ THz, and we chose $E_0 = 18 \times 10^9$ V/m. ρ and ϕ are radial distance and azimuth angle in cylindrical coordinates (ρ , ϕ , z), and w is the beam width at the wavefront z = 0; considering the dimension of the system, we set w = 7 unit cells (u.c.). The integer *m* and σ characterize the phase twist of the field and the handedness of the polarization, corresponding to orbital and spin angular momentum, respectively. Their product $-m \cdot \sigma$ gives the winding number of the field [83]. In this work, we choose $\sigma = 1$ and m = -1. We describe other computational details and explain the feasibility of designing such optical field in experiment in the Supplemental Material [52].

Figure 1(a) illustrates the interaction between the OV beam and the PZT films. The light is normally incident onto the (001) plane, and we let the beam pass through the center of the plane. In contrast with linearly and circularly polarized lights for which the polarized electric field has a line-shape profile, the electric field of the OV beam at the wavefront has a circular shape; also, the field vector $\vec{E}(\vec{r}, t)$ changes its orientation with both azimuth angle ϕ and time *t*. According to Eq. (1), the factor $\rho e^{-(\rho^2/w^2)}$ dictates that, along the radial

direction, the field intensity $E(\rho)$ first increases and then decreases.

The initial configuration in PZT film without OV beam is a homogeneously polarized state (i.e., a monodomain) due to good screening and compressive misfit strain, where all electric dipoles are aligned and point down with out-ofplane p_{τ} components. After turning on the light and when the cyclic dynamics is well established, the local soft modes develop an in-plane component $u_{x,y}(\vec{r},t)$ parallel to the local $E_{xy}(\vec{r}, t)$. Such local in-plane polarization is due to the $-P \cdot E$ coupling and appears in addition to the original p_z component reminiscent of the original poled domain. As an example, Fig. 1(b) illustrates that at the moment t = nT (where *n* is an integer and *T* is the period of the OV beam), $u_{x,y}(\vec{r})$ exhibits a divergent distribution pattern with all $u_{x,y}(\vec{r})$ directed radially outward, similar to the synchronous distribution of $E_{x,y}(\vec{r})$ on the film's plane. As demonstrated in the bottom part of Fig. 1(b), at each site, $u_{x,y}(\vec{r}, t)$ will rotate in an anticlockwise way following the evolution of $E_{x,y}(\vec{r},t)$. In the classical point-charge model [3,84], individual rotating electric dipole induces the current loop formed at each site. This rotation produces microscopic magnetic moments and these latter overall add up to a macroscopic magnetic field [see illustration in Fig. 1(a)].

Figure 2 shows plots of $u_{x,y}(t)$ versus time at characteristic sites. Figures 2(a) and 2(b) show $u_{x,y}(t)$ of sites at $\phi = 0^{\circ}$ and 180° [namely, the x axis of the coordinates displayed in Fig. 1(b)]: both $u_{x,v}(t)$ at sites 7 u.c.(1w) and 4 u.c.(0.5w) from the center display sine or cosine pattern with close amplitudes, resulting from similar electric field magnitudes at these two distances. The periodicity is 1 ps consistent with the frequency of 1 THz OV beam, and the initial displacement of $u_{x,y}$ at t = 2 ps is settled by the azimuth angle ϕ . Note the maxima or minima of $u_{x,y}(t)$ have a 0.05 ps delay relative to the maxima or minima of local $E_{x,y}(t)$ due to the response time of electric dipole to local field. At sites 14 u.c.(2w) from the center, periodic profiles of $u_{x,y}(t)$ deviate from the sine or cosine pattern with largely reduced amplitudes, indicating that at small local field, $u_{x,y}(t)$ do not closely follow $E_{x,y}(t)$. According to symmetry, by interchanging $u_x \leftrightarrow u_y$, $u_{x,y}(t)$ of sites at $\phi = 90^{\circ}$ and 270° [the y axis of the coordinates displayed in Fig. 1(b)] in Figs. 2(c) and 2(d) are just replica of Figs. 2(a) and 2(b), only up to a shift of 0.5 ps to compensate the phase difference. These mode displacements indicate that the winding pattern and orbital angular momentum are indeed transferred from light to local soft modes owing to their mutual couplings.

Now we start to analyze the effective magnetic field from the rotation of local soft modes or, equivalently, electric dipoles. Enlightened by previous derivations about the magnetic moment induced by steady current, moving



FIG. 2. (a)–(d) The variation of $u_{x,y}(\vec{r}, t)$ at different sites with time. In accordance with the coordinates displayed in Fig. 1(b), $u_{x,y}$ associated with 6 sites on the *x* axis of the coordinates are shown in (a),(b), while $u_{x,y}$ associated with 6 sites on the *y* axis are shown in (c),(d). In the legend, the "+" and "–" signs denote whether sites are located on the positive or negative side of the axes, respectively. "*xw*" ($x = 0.5 \times, 1 \times, 2 \times,$ and *w* is the beam radius of 7 u.c.) denotes the distance from the site to the center. $u_{x,y}$ is in the unit of lattice constant (3.99 Å as 1 u.c.). (e) The effective magnetic field generated in the PZT system illuminated by the OV beam.

charges and circular phonons [3,84], the on-site orbital magnetic moment is given by

$$\boldsymbol{m}_i = \frac{Z^* e}{2} \boldsymbol{u}_i \times \frac{d \boldsymbol{u}_i}{d t},\tag{2}$$

where e is the elementary charge of the electron (see Supplemental Material [52] for detailed derivation). This equation can be directly used in the effective Hamiltonian approach to compute microscopic m_i .

Histograms in the Supplemental Material [52] show that at all times, on-site m_i are distributed in the range of $5 \times 10^{-8} \sim 3 \times 10^{-3} \mu_B$. By approximating B_i by $\mu_0 m_i/V$, where V is the volume of the unit cell and μ_0 is the vacuum permeability, and summing over all sites in the system, we compute the z component of the effective magnetic field $B_z(t)$ and plot it versus time in Fig. 2(e). As $B_z(t) \sim u(t)^2$, the frequency is doubled and it has a period about 0.5 ps. The magnitude of the total magnetic field varies between 0.55*T* and 0.65*T*, much more dramatic compared to the dynamical magnetic field from a moving charged domain wall where the magnetic moment and magnetic field per cell is $2.5 \times 10^{-5} \mu_B$ and 4.5 µT, respectively [80]. We note that a very recent work studied dynamical multiferroicity in SrTiO₃ illuminated by circularly-polarized THz field and reported a simulation value of m_i around $0.01\mu_B$ [28], larger than the maximal m_i in our calculation. Though with different origins, both works demonstrate that generating strong magnetic field with dynamical multiferroicity via THz light is feasible.

In contrast to $B_z(t)$, the net in-plane magnetic field $B_{x,y}(t)$ vanishes, owing to a symmetric distribution of $m_i(t)$ within each plane. As an example, Figs. 3(a) and 3(b) display full in-plane views of the $m_{x,y}$ distribution at time 3.3 ps on layer 2 and layer 6, respectively. Both layers present vortex ring structures with left-hand circulation, and $m_{x,y}$ spans a full 360° when it goes along ϕ for a full circle. Nevertheless, the vortex on layer 6 is divergent with the central $m_{x,y}$ pointing outwards, while the vortex on layer 2 is convergent with central $m_{x,y}$ pointing inwards. As revealed in our previous study [85], the out-of-plane component of electric dipoles p_z vary with time as an electrostatic response to the time-varying $p_{x,y}$: at time 3.3 ps, all layers have negative p_z and share a similar anticlockwise pattern for $p_{x,y}$, while layer 2 has negative dp_z/dt but layer 6 has positive dp_z/dt in the central region. Mutual products between dp_z/dt and $p_{x,y}$, and between $dp_{x,y}/dt$ and dp_z induce magnetic merons with their configuration differing between different layers. Figure 3(c) gives a section view when the system is cut through y = 40, and it presents m_r and m_z at sites on the x axis for all five layers. With an increasing ρ , m_x decreases while the out-of-plane m_z develops to be the major component; for ρ larger than 7 u.c.(w), m_z starts to decrease, and m_i becomes much smaller at sites 14 u.c.(2w) from the center. The positive m_{τ} at each site and the chirality of vortices are settled by the left-hand polarization of the light and the counterclockwise rotating $p_{x,y}$.

We characterize the topology of magnetic vortex ring on each layer, by computing the Pontryagin charge density ρ_{sk} and then integrating it over the full plane:

$$N_{sk} = \int \rho(\vec{r}) d^2 \vec{r} = \int \frac{1}{4\pi} \vec{n}(\vec{r}) \cdot \left(\frac{\partial \vec{n}(\vec{r})}{\partial x} \times \frac{\partial \vec{n}(\vec{r})}{\partial y}\right) d^2 \vec{r}, \quad (3)$$

where $\vec{n}(\vec{r})$ is the normalized magnetic moment vector at site \vec{r} . Note at some sites, especially far from the center, m_i are several orders of magnitudes smaller compared to m_i within beam radius; therefore, we round up each component of m_i at an accuracy of $10^{-5}\mu_{\beta}$, so that m_i smaller than



FIG. 3. (a) In-plane view of the magnetic moment configuration on layer 2 at t = 3.3 ps, with arbitrary units. The horizontal and vertical axes denote the site number along the x and y directions, respectively. In-plane arrows denote $m_{x,y}$ and the color denotes m_{z} . (b) Similar to (a), but on layer 6 at t = 3.3 ps. (c) Section view of the magnetic moment configuration in the y = 40 plane at t = 3.3 ps. The horizontal and vertical axes denote the site number along the x direction and the layer number along the z direction, respectively. The arrows denote m_x and m_z components. (d) Topological charge density map ρ_{sk} on layer 2 at t = 3.3 ps. (e) Spherical triangles set by three orbital magnetic moments m_i on sites (40, 40), (41, 40), and (41, 41), respectively. Vertices S_1 , S_2 , and S_3 are projections of three m_i on the sphere. An enlarged in-plane view of these moments are presented in the inset. (f) The variation of topological number N with time for each layer during 3-4 ps.

the threshold value are excluded in contributing to field topology.

The computed topological charge N_{sk} is 0.5 for all five layers at 3.3 ps, consistent with direct observation in Figs. 3(a)–3(c) that m_i on each layer cover the whole hemisphere in three-dimensional order parameter space. This demonstrates that "meronlike" objects, as one half of skyrmions, are created [86–89]. Distinguished from spin magnetic moment and asymmetric magnetic exchange interactions, where usually merons and antimerons appear in pairs [90–93], here antimerons in the shape of antivortex and with an opposite charge $N_{sk} = -0.5$ are not observed; also, the merons do not emerge from a uniform background field. The distribution of ρ_{sk} on layer 2 is shown in Fig. 3(d), and we identify a sharp charge density peak localized at the center. In a discrete lattice model, ρ_{sk} is computed as the area of spherical triangle whose vertices are projected field vectors at three neighboring sites [81]. The large ρ_{sk} at the center corresponds to a swirling configuration where $m_{x,y}$ at site (40, 40) are almost antiparallel to $m_{x,y}$ at site (41, 41) along the diagonal [110] direction, and are almost orthogonal to $m_{x,y}$ at site (41, 40) [and $m_{x,y}$ at (40, 41)], as shown in the inset of Fig. 3(e). When unit vectors of m_1, m_2 , and m_3 (or m_1, m_3 , and m_4) are projected onto the surface of the 3D sphere [Fig. 3(e)], great arcs passing these vertices enclose spherical triangles with large area and result in large ρ_{sk} , as shown in Fig. 3(e).

We also compute topological charges N_{sk} at other times and plot them in Fig. 3(f). Stabilities of magnetic merons and $N_{sk} = 0.5$ for all five layers are well maintained between 3.15-3.35 and 3.6-3.85 ps, while between 3.0–3.1 and 3.5–3.55 ps, N_{sk} for some layers deviate from 0.5 slightly. Interestingly, N_{sk} of layer 6 at 3.4–3.45 ps and N_{sk} of layer 2 at 3.9–3.95 ps significantly differ from 0.5. As revealed in the Supplemental Material [52], this is associated with more than one sharp ρ_{sk} peak and swirling configuration of m_i residing at different places within the layer. Since m_i originates from variation of electric dipoles p_i , this demonstrates that at these moments, local p_i vary differently at interfacial layers compared to other layers. Note that ferroelectric skyrmions only form at layer 2 and layer 6 and the inversion of p_z occurs when N_{sk} of the respective layer is far from 0.5 [85]. We present the detailed variation of p_i resolved on each layer in the Supplemental Material [52].

In conclusion, using first-principles-based atomistic methods, we demonstrate that magnetic fields can be induced in ferroelectric but nonmagnetic PZT ultrathin films under illumination by an optical vortex beam. Orbital angular momentum is transferred from the optical field to local soft modes, and rotations of electric dipoles produce loop current and the resultant orbital magnetic moment of ions. We further identify formation of magnetic merons by characterizing orientations of magnetic moments on different layers. Our study compliments the frequently discussed "excitations of chiral phonon" as an alternative way to put dynamical multiferroicity into practice. In experiment, the dynamical multiferroicity induced in PZT thin films can be quantified by measuring the time-resolved Kerr ellipticity [28,30], while the induced magnetic merons can be visualized by Lorentz transmission electron microscopy [94–97].

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*Corresponding author: laurent@uark.edu

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