Stationary Charge Radiation in Anisotropic Photonic Time Crystals

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Time metamaterials offer a great potential for wave manipulation, drawing increasing attention in recent years. Here, we explore the exotic wave dynamics of an anisotropic photonic time crystal (APTC) formed by an anisotropic medium whose optical properties are uniformly and periodically changed in time. Based on a temporal transfer matrix formalism, we show that a stationary charge embedded in an APTC emits radiation, in contrast to the case of isotropic photonic time crystals, and its distribution in momentum space is controlled by the APTC band structure. Our approach extends the functionalities of time metamaterials, offering new opportunities for radiation generation and control, with implications for both classical and quantum applications.

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The electrodynamics of space-time metamaterials, i.e., artificial materials whose parameters are modulated in space and time, has become in recent years a very active field of research [1-7]. In the special case of pure-time media [2], a wave propagating in a spatially homogeneous medium changes its propagation features as the material properties change uniformly in time [8–11]. The simplest example is a plane wave subject to a sudden change of permittivity ε and/or permeability μ . This problem was studied in the last century by Morgenthaler [12], who showed that the wave, with given frequency ω and wave vector \vec{k} , splits into forward and backward propagating waves. Both waves retain the original value of \vec{k} due to preserved spatial uniformity of the medium, but oscillate at a new frequency corresponding to the new dispersion of the modified medium. An insightful discussion of this timereflection effect, with its experimental demonstration for both water and electromagnetic waves, appears in Refs. [13-15]. In such a context, a variety of different platforms for temporal modulation have been considered in the last few years, enabling intriguing phenomena and/or mechanisms, such as the inverse prism [16], temporal aiming [17], complete polarization conversion [18], extreme energy transformations [19], temporal disorder [20–22], and nonreciprocity [23].

Of great interest are also problems concerning radiation by sources (charges, electric dipoles, quantum emitters) embedded in time-varying media [24–30]. Already in the 1970s it was pointed out by Ginzburg and co-workers that a charge moving with constant velocity in a homogeneous medium is bound to radiate if the refractive index changes in time [24–27]. This early work focused mostly on a single switching event of the material properties. Recently, it has been shown that photonic time crystals (PTCs) created by periodically modulating the scalar permittivity $\varepsilon(t)$ of a homogeneous isotropic material [31,32] can induce amplified coherent radiation associated with momentum-gap modes, of both free moving electrons [28] and oscillating dipoles [30], based on which the interesting concept of nonresonant PTC lasers has been proposed.

So far, PTCs have been limited to isotropic media. In this Letter, we introduce anisotropic photonic time crystals (APTCs), in which the periodic temporal modulation pertains to the (relative) permittivity tensor $\overline{\overline{\varepsilon}}(t)$ characterizing uniform anisotropic media. We construct a generalized temporal transfer matrix formalism to deal with sources embedded in APTCs, showing that the anisotropy plays a fundamental role in enriching the radiation features of these time metamaterials. In particular, we show that a stationary charge embedded in an APTC emits radiation, in stark contrast with light emission of free electrons in PTCs, for which charge motion is required [28]. Based on our results, we unveil an implicit relation between light emission of the fixed charge and the band structure of the APTC, demonstrating exciting opportunities to control the radiative energy distribution in momentum space for steering the direction of light emission in space.

Anisotropic photonic time crystals and their band structure.—We start with the discussion of APTCs without sources, focusing on the simplest form of an APTC. Specifically, our APTC is formed by alternating periodically in time a pair of lossless nonmagnetic media, one being isotropic and the other one being a uniaxial crystal [labeled by (m), m = 1, 2, respectively] [see Fig. 1(a)]. For

simplicity, we neglect material dispersion and use the principal axes of the uniaxial crystal as our coordinate system. In these axes, the dielectric tensor of the uniaxial crystal is $\bar{\varepsilon}^{(2)} = \text{diag}(\varepsilon_{\perp}, \varepsilon_{\perp}, \varepsilon_{\parallel})$, while the isotropic crystal is described by $\overline{\bar{\epsilon}}^{(1)} = \epsilon \overline{\bar{I}}_3$ with $\overline{\bar{I}}_N$ being the identity matrix of order *N*. In the APTC, the displacement field $\vec{D}^{(r)}(\vec{r}, t)$ of the total radiation can be written in terms of the space Fourier transform as $\vec{D}^{(r)}(\vec{r},t) = [1/(2\pi)^3] \times$ $\int_{k,>0} \vec{D}^{(r)}(\vec{k},t) e^{-j\vec{k}\cdot\vec{r}} d^3\vec{k}$ + c.c., where the wave vector $\vec{k} = [k_x, k_y, k_z]^T$, with c.c. standing for complex conjugation, and superscripts (r) and T denoting radiation waves and the transpose operation, respectively. Within each temporal interval in which the medium is uniaxial, for given wave vector \vec{k} , the dispersion relation allows for two different positive frequencies $\omega_1^{(2)} = k c_0 / \sqrt{arepsilon_\perp}$ and $\omega_2^{(2)} = kc_0 \sqrt{(\hat{k}_x^2 + \hat{k}_y^2)/\varepsilon_{\parallel} + \hat{k}_z^2/\varepsilon_{\perp}}$, where c_0 is the speed of light in free space, and $\hat{k}_{\alpha} \equiv k_{\alpha}/k$, $\alpha = x, y, z$ (with wave number $k \equiv |\vec{k}|$) are the direction cosines of \vec{k} . The two frequencies correspond to two waves, i.e., the ordinary and extraordinary waves, propagating in the \vec{k} direction (forward) and having mutually orthogonal electric displacements, $\vec{D}_1 = [-\hat{k}_y, \hat{k}_x, 0]^T$ and $\vec{D}_2 = [\hat{k}_x \hat{k}_z, \hat{k}_y \hat{k}_y \hat{k}_z, \hat{k}_y \hat{k}_y \hat{k}_y \hat{k}_z, \hat{k}_y \hat{k}_y \hat{k}_y \hat{k}_y \hat{k}_y \hat{k}_z, \hat{k}_y \hat{k}_y$ $-(\hat{k}_x^2 + \hat{k}_y^2)]^T$. These two vectors, $\vec{D}_1(\vec{k})$ and $\vec{D}_2(\vec{k})$, together with the vector \vec{k} , form an orthogonal triplet. In each temporal interval in which the medium is isotropic, the pair D_1 and D_2 can be chosen as a basis supporting two independent forward waves with the same positive frequency $\omega_1^{(1)} = \omega_2^{(1)} = kc_0/\sqrt{\epsilon}$, essentially treating the medium as a degenerate uniaxial crystal.

Considering the additional two backward waves $(-\vec{k})$ direction) oscillating at the corresponding negative frequencies $-\omega_{1,2}^{(m)}$ within each temporal slab of the isotropic (m = 1) or the uniaxial (m = 2) crystal, the total displacement of the radiation field in momentum space $\vec{D}^{(r)}(\vec{k}, t)$ for the APTC [see Fig. 1(a)] can be written compactly as

$$\vec{D}^{(r)}(\vec{k},t) = \sum_{n=1}^{2} \vec{D}_{n}(\vec{k}) [1, 1] \psi_{n}(t), \qquad (1)$$

where the vector $\psi_n(t) \equiv [f_n(t), b_n(t)]^T$ consists of the forward [and backward] time-dependent components $f_n(t) \propto e^{j\omega_n^{(m)}t}$ [and $b_n(t) \propto e^{-j\omega_n^{(m)}t}$] of the first (n = 1) and the second (n = 2) wave in medium *m*.

We proceed to describe the evolution of the radiation in the APTC, which boils down to studying the time dependence of the total state vector $\psi(t) \equiv \begin{bmatrix} \psi_1(t) \\ \psi_2(t) \end{bmatrix}$ of the two waves in \vec{k} space, since momentum is conserved upon each temporal switching event. Free wave propagation in each temporal slab of the medium m = 1, 2 in the APTC is described by the block-diagonal matrix

$$F^{(m)} = \text{diag}\{F_1^{(m)}, F_2^{(m)}\}$$
 (2)

with blocks $F_n^{(m)} = \text{diag}\{e^{j\omega_n^{(m)}T_m}, e^{-j\omega_n^{(m)}T_m}\}, n = 1, 2,$ which describes the temporal evolution of the state $\psi(t)$ within the slab of duration T_m . At a switching event of the APTC, say, at switching time $t_s = 0$ [see Fig. 1(a)], as the medium changes abruptly from m = 1 to 2, the state vector $\psi(t)$ experiences an instantaneous variation $\psi(t_s^+) = J^{(2,1)}\psi(t_s^-)$ with matching matrix [33]



FIG. 1. (a) Schematic of the APTC described in the text. (b),(c) Band structure of the APTC [with $\varepsilon = 1$, $\varepsilon_{\perp} = 25$, $\varepsilon_{\parallel} = 4$, and $T_1 = T_2 = T/2$ in (a)] in momentum \vec{k} (normalized to $k_0 = 2\pi/Tc_0$) for (b) ordinary and (c) extraordinary waves. In (c), the red [and the blue] vertical dashed line intersects the $k_x - k_z$ plane at \vec{k} with the spherical coordinates (2.5 k_0 , 0.094, 0) [and (2.5 k_0 , 1.477, 0)].

$$J_{n}^{(2,1)} = \begin{bmatrix} J_{1}^{(2,1)} & 0\\ 0 & J_{2}^{(2,1)} \end{bmatrix},$$
$$J_{n}^{(2,1)} \equiv \frac{1}{2\omega_{n}^{(2)}} \begin{bmatrix} \omega_{n}^{(2)} + \omega_{n}^{(1)}, & \omega_{n}^{(2)} - \omega_{n}^{(1)} \\ \omega_{n}^{(2)} - \omega_{n}^{(1)}, & \omega_{n}^{(2)} + \omega_{n}^{(1)} \end{bmatrix}$$
(3)

following from the continuity of \vec{D} and its time derivative $\partial \vec{D}/\partial t$ (or equivalently, the magnetic-flux density \vec{B}) [5] at the time interface t_s . Similarly, we obtain the matching matrix $J^{(1,2)} = [J^{(2,1)}]^{-1}$ accounting for the rapid evolution of the state $\psi(t)$ at the other switching event, say, at time $t_s = T_2$ [see Fig. 1(a)] when the medium changes from m = 2 to 1. To study the evolution of $\psi(t)$ in the APTC, the matching matrix $J^{(2,1)}$ in Eq. (3) and the propagation matrix $F^{(m)}$ in Eq. (2) constitute building blocks.

Of primary interest for our purpose is to study the transfer matrix $M \equiv F^{(1)}[J^{(2,1)}]^{-1}F^{(2)}J^{(2,1)}$, which describes the evolution of the state $\psi(t)$ across one temporal unit cell of the APTC, of duration $T = T_1 + T_2$ [an example of such a cell is indicated in Fig. 1(a) by the red dashed box, and $\psi(3T^-) = M\psi(2T^-)$ correspondingly]. This matrix M fully describes the long-time dynamics of the APTC, since the transfer matrix for N periods is simply given by M^N . Using Eqs. (2) and (3), the matrix M turns out to be block diagonal, i.e., $M = \text{diag}\{M_1, M_2\}$, with each block being $M_n = F_n^{(1)}(J_n^{(2,1)})^{-1}F_n^{(2)}J_n^{(2,1)}$, n = 1, 2. The first and the second blocks M_1 and M_2 are associated with the ordinary and the extraordinary waves of the uniaxial crystal, respectively. Using the blocks M_n , we obtain their band structure separately by solving the secular equations

$$\det\{M_n(\vec{k}) - e^{j\Omega_{n\sigma}T}\bar{I}_2\} = 0, \qquad n = 1, 2, \qquad (4)$$

where $\Omega_{n\sigma}$ is the Floquet frequency, with the subindex $\sigma =$ \pm labeling different eigenvalues. In Figs. 1(b) and 1(c), we consider the APTC shown schematically in Fig. 1(a) with parameters $\varepsilon = 1$, $\varepsilon_{\perp} = 25$, $\varepsilon_{\parallel} = 4$, and $T_1 = T_2 = T/2$ [38], and plot the complex dispersion relation of the ordinary [Fig. 1(b)] and extraordinary [Fig. 1(c)] waves propagating in the $k_x - k_z$ plane. To understand these band structures, we note that the blocks $M_n(\vec{k})$, n = 1, 2 in Eq. (4) can be parametrized as $M_n = \begin{bmatrix} a_n & d_n \\ d_n^* & a_n^* \end{bmatrix}$ and satisfy det $M_n = 1$ based on Eqs. (2) and (3) (see also Ref. [21]). Correspondingly, their eigenvalues are given by $\lambda_{n\pm} \equiv$ $e^{j\Omega_{n\pm}T} = \operatorname{Re}a_n \pm \sqrt{(\operatorname{Re}a_n)^2 - 1}$. If $(\operatorname{Re}a_n)^2 < 1$, we have $|\lambda_{n+}| = 1/|\lambda_{n-}| = 1$ and thus $\operatorname{Im}(\Omega_{n\pm}T) = 0$, leading to oscillations in time without systematic growth or decay. On the other hand, when $(\text{Re}a_n)^2 > 1$, the eigenvalues $\lambda_{n\pm}$ are real and $|\lambda_{n+}| = 1/|\lambda_{n-}| \neq 1$. In this case, $\operatorname{Im}(\Omega_{n\pm}T) \neq 0$ and one eigenstate grows exponentially in time while the other decays. Generally, which of these two cases is realized depends on \vec{k} producing bands and gaps in momentum space.

For ordinary waves (block M_1), the band structure is consistent with the one of PTCs, and it does not depend on the propagation direction of the waves [see Fig. 1(b)], following from the direction cosines \hat{k}_a -independent $\omega_1^{(m)}$ for M_1 . In contrast, in the case of extraordinary waves, the band structure determined by the block M_2 is anisotropic, since $\Omega_{2\sigma}$ depends not only on the magnitude of the wave vector \vec{k} but also on its direction [see the direction dependence of $\omega_2^{(2)}$ and Fig. 1(c)]. This feature implies remarkable features unique to APTCs: For instance, whether the waves in APTCs are amplified (and how strongly) depends on their momentum directions. Interestingly, the gaps in the APTC band structure can close when fixing the wave number k [for instance, at k = $2.5k_0$ with $k_0 = 2\pi/Tc_0$ in Fig. 1(c)] and varying the direction cosines \hat{k}_{α} , or more precisely, only the polar angle θ , since, due to the rotational symmetry of the considered APTC, the band structure is invariant with respect to the azimuthal angle φ .

Radiation by a stationary charge in APTCs.—Next, we study the effect of a stationary point charge Q in the considered APTCs. Clearly, the existence of a stationary charge does not affect the free propagation of radiation fields in static media, so that the evolution of the state $\psi(t)$ dictating the radiation $\vec{D}^{(r)}(\vec{k}, t)$ in Eq. (1) is determined by the same propagation matrix $F^{(m)}$ in Eq. (2) within each temporal slab of the APTC [see Fig. 1(a)]. Nevertheless, due to the presence of the charge Q, the effects of the temporal interfaces on the radiation fields can change dramatically. Assume for instance that the charge Q is fixed at the origin $\vec{r} = 0$. As a result, the total displacement field in momentum space becomes $\vec{D}(\vec{k}, t) = \vec{D}_0(\vec{k}, t) + \vec{D}^{(r)}(\vec{k}, t)$, which includes also a particular solution due to the source Q,

$$\vec{D}_0(\vec{k},t) = \frac{jQ}{\vec{k} \cdot \bar{\vec{e}}(t)\vec{k}} \bar{\vec{e}}(t)\vec{k}$$
(5)

with the matrix $\overline{\overline{e}}(t) = \overline{\overline{e}}^{(m)}$ yielding electrostatic parts $\vec{D}_0(\vec{k}, t) = \vec{D}_0^{(m)}(\vec{k})$, respectively, for time *t* within the temporal slab of the crystal m = 1 or 2 [see Fig. 1(a)]. Correspondingly, based on the same temporal boundary conditions for Eq. (3), the matching relation for the state $\psi(t)$ of the radiated fields becomes [33]

$$\psi(t_s^+) = S^{(2,1)} + J^{(2,1)}\psi(t_s^-) \tag{6}$$

with the extra source term $S^{(2,1)} = s_{21}[0, 0, 1, 1]^T$ [where the prefactor $s_{21} = jQ\Delta_{\varepsilon}\cos\theta/[k(1 + \Delta_{\varepsilon}\cos(2\theta))]$ with $\Delta_{\varepsilon} \equiv (\varepsilon_{\parallel} - \varepsilon_{\perp})/(\varepsilon_{\parallel} + \varepsilon_{\perp})$] for the temporal interface at time t_s when the medium switches from m = 1 to 2. Indeed, the source term $S^{(2,1)}$ appears in Eq. (6) due to the mismatch between the electrostatic solutions $\vec{D}_0^{(1)}(\vec{k})$ and $\vec{D}_0^{(2)}(\vec{k})$ of the charge Q in m = 1 and 2 [see also Ref. [39]], producing spontaneous emission of extraordinary waves even when the radiation fields vanish before switching, i.e., $\psi(t_s^-) = 0$ [25]. We note that the source term $S^{(2,1)} = 0$ when $\varepsilon_{\parallel} = \varepsilon_{\perp}$, which corresponds to a conventional (isotropic) photonic time crystal. Indeed, stationary charges are not expected to radiate in such a scenario. Similarly, an extra source term $S^{(1,2)} = -S^{(2,1)}$ emerges at the other switching event of the APTC in Fig. 1(a) when the medium switches from m = 2 to 1.

Now, we are ready to study the radiation by a stationary charge Q embedded in the APTC. We assume that initially, i.e., prior to time t = 0, there is no radiation field, so that the initial state of radiation $\psi(0^-) = 0$. Then, we study the effects of the first modulation cycle of the APTC. At the first switching event at time t = 0, the medium switches from the crystal m = 1 to 2, producing a radiation field $\psi(0^+) = S^{(2,1)}$ [see Eq. (6)]. This field freely propagates in the medium m = 2 for a time interval T_2 , resulting in $\psi(T_2^-) = F^{(2)}S^{(2,1)}$. Next comes the switching event back to the medium m = 1 at time $t = T_2$, which multiples $\psi(T_2^-)$ by the matching matrix $J^{(1,2)}$, and, in addition, it creates a new radiation field $S^{(1,2)}$. Thus, we have $\psi(T_2^+) = S^{(1,2)} + J^{(1,2)}\psi(T_2^-)$. The cycle is completed by free propagation for duration T_1 in medium m = 1, so that $\psi(T^{-}) = F^{(1)}\psi(T_{2}^{+})$. Putting these pieces together, we can write the radiated field at the end of the first modulation cycle as $\psi(T^-) = (F^{(1)} - M)S^{(1,2)}$, or equivalently, $\psi_1(T^-) = 0$ and $\psi_2(T^-) = -s_{21}(F_2^{(1)} - M_2)[{}_1^1],$ where $M = F^{(1)}J^{(1,2)}F^{(2)}J^{(2,1)}$ is the same unit-cell transfer matrix as introduced before. Indeed, it is generally true that the radiation field produced by the APTC in Fig. 1(a), when $\psi(0^-) = 0$, does not involve ordinary waves, i.e., $\psi_1(t > 0) = 0$, due to the fact that the source term creates only extraordinary waves [see Eq. (6)], and that they propagate independently from the ordinary waves [see Eqs. (2) and (3)]. Repeating the above recursion n_t times, we obtain the total radiation field after n_t modulation cycles [33]

$$\psi_2(n_t T^-) = -s_{21} \frac{\bar{\bar{I}}_2 - M_2^{n_t}}{\bar{\bar{I}}_2 - M_2} (F_2^{(1)} - M_2) \begin{bmatrix} 1\\1 \end{bmatrix}, \quad (7)$$

which connects the radiation field of the APTC to the band structure of extraordinary waves via M_2 [see Eq. (4)].

The APTC with a stationary charge not only supports radiation fields, but it also controls their energy distribution in momentum space. To see this, we study the timedependent evolution of the energy distribution in momentum space of the radiation field based on Eq. (7). At each

time $t = n_t T^-$, the fields are present in the isotropic medium m = 1, and thus the total electromagnetic energy $W_{\rm EM}(t) =$ $[1/(2\pi)^3] \int_{k_{-}>0} w_{\rm EM}(\vec{k},t) d^3\vec{k}$ with energy density in momentum space $w_{\text{EM}}(\vec{k},t) = w_{\text{EM}}^{(s)}(\vec{k},t) + w_{\text{EM}}^{(r)}(\vec{k},t)$, which involves the constant electrostatic part $w_{\rm EM}^{(s)}(\vec{k},t) = Q^2/$ $(\varepsilon_0 \varepsilon k^2)$ and the radiation part $w_{\rm EM}^{(r)}(\vec{k},t) = 2\sin^2\theta$ $\psi_2(t)|^2/(\varepsilon_0\varepsilon)$. In Fig. 2, we choose the same parameters as in Figs. 1(b) and 1(c), and show the normalized radiative energy density $\hat{w}_{\text{EM}}^{(r)}(\vec{k},t) \equiv w_{\text{EM}}^{(r)}(\vec{k},t) / \max_{\vec{k}} w_{\text{EM}}^{(r)}(\vec{k},t)$ after different modulation cycles. As shown, the radiative energy density $\hat{w}_{\rm FM}^{(r)}(\vec{k},t)$ spreads over a relatively large momentum space near k = 0 initially [Fig. 2(a)] due to the prefactor $s_{21} \propto 1/k$ of the source in Eq. (7). With increasing modulation cycles, the modes in the band gaps with nonzero imaginary frequency grow exponentially and start to dominate in momentum space [Fig. 2(b)]. When the number of modulation cycles becomes larger [Figs. 2(c)-2(f)], the radiative energy $\hat{w}_{\rm EM}^{(r)}(\vec{k}, t)$ increasingly localizes around the momentum with $k \approx 2.5k_0$ and $\theta \approx 0.094$ [33], where the gap modes have the relatively larger imaginary frequency $|\text{Im}(\Omega_{2\sigma}T)|$ (and thus growing rate) [Fig. 1(c)] within the k range of initial excitations [Fig. 2(a)], forming a highly directional nonresonant laser source tunable by manipulating the momentum band gaps [33]. In contrast with the nonresonant tunable laser based on PTCs proposed in Refs. [28,30], the extreme lasing directionality in APTCs is achieved due to their inherent anisotropic band structure, which does not require source manipulation, such as precise control of the movement of free charges for delicate phase matching required to sustain the radiation process [28].

To illustrate better the evolution of the radiation field in momentum space, we study the distinct dynamics of the magnetic-flux density $\vec{B}(\vec{k}, t)$ in the band gap and band,



FIG. 2. Normalized radiative energy $\hat{w}_{\text{EM}}^{(r)}(\vec{k}, t)$ in momentum space at various time instants (a) $t = T^-$, (b) $t = 4T^-$, (c) $t = 7T^-$, (d) $t = 10T^-$, (e) $t = 13T^-$, and (f) $t = 16T^-$ of the APTC in Fig. 1(a) due to an embedded stationary charge. The initial radiative energy at time $t = 0^-$ is zero, and the other parameters are the same as in Figs. 1(b) and 1(c).



FIG. 3. Magnitude of the magnetic-flux intensity component $B_y(\vec{k}, t)$ (in units of $Q\eta_0/k_0$) as a function of the time *t* for the APTC with an embedded stationary charge *Q*, at the momentum \vec{k}_l , l = 1, 2 of the spherical coordinates (a) $(2.5k_0, 0.094, 0)$ and (b) $(2.5k_0, 1.477, 0)$. The empty circles and the solid lines are the theoretical and numerical results, respectively. The lower inset in (a) shows the initial stage of the main panel on a linear scale, and the upper insets of both (a) and (b) exhibit the corresponding spatial distribution of the B_y field at time t = 6T. The other parameters are the same as Figs. 1(b) and 1(c).

respectively. For the APTC with a stationary charge, $\vec{B}(\vec{k}, t)$ is associated only with the radiation, and it reads $\vec{B}(\vec{k}, n_t T^-) = \vec{B}_2^{(1)}(\vec{k})[1, -1]\psi_2(n_t T^-)$ [with the basis state $\vec{B}_2^{(1)}(\vec{k}) = \vec{D}_1(\vec{k})\eta_0/\sqrt{\epsilon}$ after n_t modulation cycles, where η_0 is the characteristic impedance of free space. In Fig. 3, we choose two representative momenta $\vec{k} = \vec{k}_l$, l =1, 2 with spherical coordinates $(k, \theta, \varphi) = (2.5k_0, 0.094, 0)$ and $(2.5k_0, 1.477, 0)$ in the band gap and band [see Fig. 1(c)] and show the only nonzero component $B_{\nu}(\vec{k}_l, t)$ as a function of the time calculated both solving Maxwell's equations numerically (solid lines) and using Eq. (7) (empty circles). As shown, starting from zero initial values without radiation, the gap mode grows exponentially [Fig. 3(a) and lower inset] and takes over the bounded band mode [Fig. 3(b)] after a few modulation cycles, steering the direction of light emission in space; see upper insets for individual snapshots at time t = 6T and Ref. [33] for full animations.

Conclusions.—In this Letter, we introduced the concept of APTCs formed by photonic time crystals involving anisotropic media and developed a generalized temporal transfer matrix formalism to study APTCs with sources. We showed that APTCs enable radiation of a stationary charge and manage emission features due to their reconfigurable (anisotropic) band structure in momentum space. Our approach greatly extends the concept of nonresonant tunable PTC lasers proposed in Refs. [28,30] to the scenario of stationary charges, and it showcases unique opportunities to leverage time interfaces, anisotropy, and electrostatic fields to efficiently produce directional dynamic nonresonant lasing. This concept may be further generalized by incorporating non-Hermitian elements [19] to push its boundaries toward the full control of light emission via time metamaterials without spatial structures, with implications for laser technologies and classical and quantum photonic applications. Furthermore, these concepts may be fruitfully implemented in other physical domains, such as for acoustic waves.

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