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Orbital magnetization of Floquet topological systems

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A general expression for the orbital magnetization of a Floquet system is derived. The expression holds for a clean system and is valid for any driving protocol and arbitrary occupation of the bands. The orbital magnetization is shown to be large not only for Chern insulators, but also for anomalous phases where the Chern number does not fully account for the topology. In addition, the orbital magnetization is shown to take significant values both for a thermal equilibrium occupation of the Floquet bands and for occupations determined by a quantum quench from an initial state with zero orbital magnetization. For the latter case, the orbital magnetization is shown to be highly sensitive to van Hove singularities of the Floquet bands.

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I. INTRODUCTION

A frontier topic in condensed matter physics and atomic, molecular, and optical systems is the engineering of new states of matter by Floquet driving [1–6]. Such driving can give rise to myriad topological [7–24] and ordered phases [25–33] that have no analog in static systems. For example, in two spatial dimensions (2D), the Chern number does not fully characterize the topology under Floquet driving. An extreme case is one where the Chern number is zero and yet chiral edge modes exist in the system [9]. These edge modes can give rise to a quantized charge pumping [34] and a quantized orbital magnetization [35,36] when the bulk states are fully localized by spatial disorder, with the corresponding system known as an anomalous Floquet Anderson insulator [34].

Despite this progress, understanding the linear response properties for Floquet topological systems is a largely open question. While many new topological invariants have been constructed for Floquet systems, how these manifest when external probe fields are applied is mostly unexplored. As a first step, a topological quantum field theory for the driving protocol of Ref. [9] was derived in Ref. [37], and the appearance of an orbital magnetization as a linear response to an external magnetic field, under spatially periodic boundary conditions, was shown. However, the orbital magnetization for general Floquet driving and when the bulk states are not fully localized is an open and important question both from a theoretical point of view and for experiments where bulk states may be conducting. In contrast, orbital magnetization of static systems has been actively studied [38–40], with recent applications to twisted bilayer materials [41–43] that have been shown to exhibit large orbital magnetic moments [44,45].

We derive a general formula for the orbital magnetization of 2D Floquet systems in the absence of disorder. Our formula holds for any driving protocol as well as any filling of the bands. We apply our formula to graphene under Floquet driving, and we present results for two filling profiles, one where the Floquet states are occupied according to a thermal distribution, and the second where the occupation of the Floquet states is set by a quantum quench from an initial state with zero orbital magnetization. For the latter case, the orbital magnetization is shown to be highly sensitive to van Hove singularities of the Floquet bands.
T is the period of the drive, the Hamiltonian may be written as 
\[ H(k, t) = H(k, t + T) \]
According to Floquet theory [46, 47], the Floquet eigenstate \( |\psi_{n,k}(t)\rangle \) can be decomposed as 
\[ |\psi_{n,k}(t)\rangle = e^{-i\epsilon_n \tau \hat{A}} |\phi_{n,k}(t)\rangle \]
where \( |\phi_{n,k}(t)\rangle \) is the time-periodic Floquet quasimode and \( \epsilon_n \) is the quasienergy, with \( n \) labeling the bands. The first Floquet Brillouin zone corresponds to \( \epsilon_n \in [-\pi/T, \pi/T] \). It is convenient to define the Floquet Hamiltonian \( H_F = H(k, t) - i\partial_t \), which obeys
\[ H_F |\phi_{n,k}(t)\rangle = \epsilon_n |\phi_{n,k}(t)\rangle. \tag{1} \]

Our goal is to study the linear response of the above system to a small magnetic field \( B \) applied in the perpendicular \( \hat{z} \) direction and varying slowly with an in-plane wave vector \( \mathbf{q} = q\hat{y}, B = \dot{z}B\cos(qy) \) [40]. Using a gauge where the vector potential is \( \mathbf{A} = -\dot{B}\sin(qy)/q \), the perturbing term corresponding to this magnetic field is (we set \( e = 1, h = 1 \); see Appendix A)
\[ V(t) = \sum_\mathbf{p} \mathbf{A}_\mathbf{p} \cdot \mathbf{J}_\mathbf{p}(t) = \mathbf{A}_q \cdot \mathbf{J}_\mathbf{q}(t) + \text{c.c.}, \tag{2} \]
where the current operator is \( \mathbf{J}_\mathbf{q}(t) = \sum_{\mathbf{k}} e^{i\mathbf{q}\cdot\mathbf{k}/2} \mathbf{v}(\mathbf{k}, t)c_{\mathbf{k}-\mathbf{q}/2} \) and \( \mathbf{v}(\mathbf{k}, t) = \partial_\mathbf{k} H(k, t) \), with \( \mathbf{A}_q = -\dot{B}/(2iq) \). Denoting \( \delta|\psi_{n,k}(t)\rangle \) as the change in the eigenstate to \( O(V) \), and assuming that the perturbation is switched on at time \( t = 0 \), we have (see Appendix A)
\[ \delta|\psi_{n,k}(t)\rangle = -i \int_0^t dt' \int_{\mathbf{k}_a} e^{-i\epsilon_n \tau} |\phi_{\mathbf{k}_a}(t)\rangle e^{-i\epsilon_n \tau} |\phi_{\mathbf{k}_a}(t')\rangle |\phi_{\mathbf{k}_a}(t')\rangle \tag{3} \]
Since the quasienergy does not change to \( O(V) \) [or, equivalently, to \( O(B) \); cf. Appendix A], the leading change to the quasimode \( \delta|\phi_{n,k}(t)\rangle = e^{i\epsilon_n \tau} |\phi_{n,k}(t)\rangle \).

We are interested in studying the linear response of the above system to a weak external magnetic field. It was shown that the orbital magnetization in a Floquet eigenstate time averaged over one drive cycle equals the rate at which the quasienergy of that eigenstate changes due to the applied magnetic field (see, for example, Ref. [35]). With this as the starting point, we will compute the change in the average quasienergy of each Floquet eigenstate due to the applied magnetic field, weighting the result by the occupation probability of each Floquet eigenstate.

The average quasienergy for a system described by the density matrix \( \rho(t) = \sum_{n,k} f_{n,k} |\phi_{n,k}(t)\rangle \langle \phi_{n,k}(t) | \) is
\[ E = \sum_{n,k} f_{n,k} \langle \phi_{n,k}(t) | H_F |\phi_{n,k}(t)\rangle = \sum_{n,k} f_{n,k} \epsilon_n. \tag{1} \]

![Figure 1](image-url)  
FIG. 1. (a) Schematic showing system irradiated by a circularly polarized laser. The yellow (red) loops denote chiral edge modes at the Floquet zone center (boundaries). A perturbing perpendicular magnetic field induces an orbital magnetization (tiny loops with arrows indicate induced magnetic moments). (b) and (c) Floquet bands of driven graphene on a cylinder with periodic boundary conditions in the x direction. The bands have a Chern number of 3 (b) and 0 (c). Edge states at the Floquet zone center (boundary) are indicated by yellow (red) color. In (b), the insets show two edge states at the Floquet zone boundary (red) and one edge state at the Floquet zone center (yellow) [see schematic in (a)]. The chirality of the edge modes changes from 2 to −1 from the zone center to the zone boundary consistent with \( C = 2 − (−1) = 3 \). In (c), the zone center and the zone boundary each host two edge modes, with no change in the chirality of the edge modes from the zone center to the zone boundary, consistent with \( C = 0 \). In (c), for orientation, the bands of static graphene are shown as black dashed lines. We take 50 sites [(b) and (c)] in the y direction and keep Floquet harmonics up to \( |m_{\text{max}}| = 5 \) (b) and \( |m_{\text{max}}| = 20 \) (c).
to another. At $O(B)$, only virtual processes are allowed, and therefore to this order, there is no change in the occupation.

Thus the change in the average quasienergy due to the external magnetic field is

$$
\delta E(t) = \sum_{n,k} f_{n,k} \{ \phi_{n,k}(t)|H_F|\delta \phi_{n,k}(t) \} + \text{c.c.} = M(t) B,
$$

(4)

where $M(t)$ is the induced magnetization. We are interested in the limit of long wavelengths, $q \to 0$, and long times $t \to \infty$. Even in the long-time limit, the time periodicity of the Floquet states gives rise to a residual time dependence to the magnetization. We will explore the magnetization averaged over one drive cycle, i.e., $\overline{M} = \lim_{t \to \infty} \langle \phi_{n,k}(t)|H_F|\phi_{n,k}(t) \rangle / T$, where $\overline{O}$ indicates the average of a quantity over one drive cycle. We introduce the following notation for the $n$th Fourier component of various matrix elements:

$$
[\overline{O}]_{nk}^{m+q} = \frac{1}{T} \int_0^T dt' e^{-i\omega_0 t'} \langle \phi_{n,k}(t)|\overline{O}(t')|\phi_{n',k+q}(t') \rangle.
$$

(5)

We find (see Appendix A for intermediate steps)

$$
\overline{M} = \lim_{q \to 0} \frac{1}{2q} \sum_{n',k,m=\text{int}} \left( f_{n,k} - f_{n',k+q} \right) \left[ [H_F]_{nk}^{m+q} \right]
\times \left[ v^r(k) \right]_{nk}^{m+q} \frac{1}{\epsilon_{n,k} + 2\hbar^2 - \epsilon_{n',k+q}} + \text{c.c.},
$$

(6)

Above, $v^r$ is the velocity operator along the $\hat{x}$ direction.

Taking the long-wavelength limit is subtle, and the final expression depends on the distribution function $f_{n,k}$. In particular, we find (restoring $e, \hbar$; see Appendix A)

$$
\overline{M} = -\frac{e}{2\hbar} \text{Im} \sum_{n,k}
\times \left( f_{n,k} \{ \delta_k \phi_{n,k}(t) \} (\epsilon_{n,k} + H_F) \times [\delta_k \phi_{n,k}(t)] - \lim_{\epsilon \to 0} \left( \frac{f_{n,k} - f_{n,k+q}}{\epsilon_{n,k} - \epsilon_{n',k+q}} \right) \epsilon_{n,k}
\times \{ \delta_k \phi_{n,k}(t) \} (\epsilon_{n,k} - H_F) \times [\delta_k \phi_{n,k}(t)] \right).
$$

(7)

The cross product indicates that the orbital magnetization depends on the Berry curvature of the bands. Since the orbital magnetization is related to the operator $\mathbf{r} \times \mathbf{v}$, with $\mathbf{v} = \partial_t \mathbf{r} = -i[\mathbf{r}, H(t)]$, and since $\mathbf{r}$ in the momentum basis corresponds to $\partial_k$, that explains why we have two partial derivatives in momentum weighted by $H_F, \epsilon_{n,k}$ [39].

In order to explore the physics, we need to make some assumptions about the occupations $f_{n,k}$. This requires us to think carefully about the initial state before the periodic driving was switched on and to also account for relevant dissipative processes. When the system is coupled to an ideal reservoir described by a Fermi-Dirac distribution function with a chemical potential $\mu$ and temperature $\beta^{-1}$, and when the Floquet system is driven at frequencies that are large compared with the bandwidth, it has been shown that the Floquet system acquires the ideal distribution function of the reservoir [48,49]. However, for driving frequencies that are resonant, i.e., comparable to the bandwidth, the occupation of the Floquet states can be complicated and highly dependent on the details of the system-reservoir coupling [48,49]. Nevertheless, one can imagine performing careful reservoir engineering to obtain desired results for the Floquet occupation. For this reason, we first consider the case where all states are occupied by a Fermi-Dirac distribution function at a temperature $\beta^{-1}$ and a chemical potential $\mu$. Moreover, we study corrections to the combination $E - \mu/N$, where $N$ is the number operator. This involves shifting $\epsilon_{n,k}, H_F$ by $\epsilon_{n,k} - \mu, H_F - \mu$, thus arriving at the following expression for the orbital magnetization (see Appendix A):

$$
\overline{M}_1 = -\frac{e}{2\hbar} \text{Im} \sum_{n,k}
\times \{ \delta_k \phi_{n,k}(t) \} (\epsilon_{n,k} + H_F - 2\mu) \times [\delta_k \phi_{n,k}(t)],
$$

(8b)

$$
\overline{M}_2 = \frac{e}{2\hbar} \text{Im} \sum_{n,k}
\times [\delta_k \phi_{n,k}(t)] (\epsilon_{n,k} - H_F) \times [\delta_k \phi_{n,k}(t)].
$$

(8c)

Above, $f'_{n,k} = \partial_{\epsilon_{n,k}} f$. Since $\overline{M}_1$ is proportional to the occupation, it survives at zero temperature; $\overline{M}_2$, being proportional to the derivative of the occupation times the energy, contributes only at nonzero temperature. Equations (8a)–(8c) reduce to the equation for a static system [40], with $H_F$ being replaced by the static Hamiltonian $H$. However, for static systems in thermal equilibrium, the orbital magnetization is determined as a linear response correction to the free energy $F = E - \mu/N - \beta^{-1} S$, $S$ being the entropy and $\beta^{-1}$ being the temperature. In contrast, here we are considering the linear response of a closed quantum system described by a density matrix $\rho$. For the orbital magnetization, this corresponds to determining corrections to the quasienergy of each Floquet eigenstate weighted by the occupation of that state [35]. Thus the natural object that appears here is corrections to $E$ or $E - \mu N$, where $\mu N$ is a convenient shift.

For a two-band Floquet system with particle-hole symmetry, the orbital magnetization per unit area $A$ when one of the bands is fully occupied and the other is empty reduces to $\overline{M}/A = -\langle e/\hbar \rangle C \mu / 2\pi$ (see Appendix A). Thus the magnetization vanishes for this ideal filling for anomalous phases with $C = 0$. Therefore, in order to probe the $C = 0$ phase, we need to either break particle-hole symmetry or raise the temperature in order to occupy the other band, or we could simply have a nonequilibrium occupation of the Floquet bands. As discussed above, the occupation of Floquet states is not guaranteed to be in thermal equilibrium even when they are coupled to an ideal reservoir [48–50] except for some limiting cases of high-frequency driving or for very careful reservoir engineering. The most natural distribution is one arising from a quantum quench from an initial state where a drive was...
absent, and then the drive was switched on following a certain protocol. For a sudden switch-on of the drive at a time \( t = 0 \), the quench distribution function is

\[
f_{n,k} = \sum_{\alpha} |\langle \phi_{n,k}(0) | \psi^{\text{in}}_{\alpha,k} \rangle|^2 f^{\text{in}}_{\alpha,k},
\]

where \( |\psi^{\text{in}}_{\alpha,k}\rangle \) are the eigenstates of the Hamiltonian in the absence of drive and \( f^{\text{in}}_{\alpha,k} \) is the occupation of these states, which we take to be a Fermi-Dirac distribution and a chemical potential \( \mu \).

The orbital magnetization following a quantum quench and obtained from corrections to the average quasienergy \( E \) is (see Appendix A)

\[
\mathcal{M}_Q = -\frac{e}{2\hbar} \text{Im} \sum_{n,k} \left[ f_{n,k} \langle \partial_t \phi_{n,k} | (\epsilon_{n,k} + HF) \times | \partial_t \phi_{n,k} \rangle \right. \\
\left. - \left( \sum_{\alpha} |\langle \phi_{n,k} | \psi^{\text{in}}_{\alpha,k} \rangle|^2 \partial_t f^{\text{in}}_{\alpha} \frac{\nu_n(\alpha, k)}{\nu_k(\alpha, k)} \right) \right. \\
\left. + f^{\text{in}}_{\alpha} \left| \frac{\partial_t \langle \psi^{\text{in}}_{\alpha,k} | \phi_{n,k} \rangle}{\nu_k(\alpha, k)} \right|^2 \right]
\times \epsilon_{n,k} \langle \partial_t \phi_{n,k} | (\epsilon_{n,k} - HF) \times | \partial_t \phi_{n,k} \rangle \right].
\]

There are two main differences between Eq. (10) and Eqs. (8a)–(8c). One is that the occupation probabilities entering in Eq. (10) are given by Eq. (9) rather than a Fermi-Dirac distribution function. Second, the term coming from \( \lim_{t' \to 0} f_{n,k} - f_{n,k+q} \) \( |(\epsilon_{n,k} - \epsilon_{n,k+q})| \) [cf. Eq. (7)] is quite subtle for the quench as the energies entering in \( f_{n,k} \) are those of the quench Hamiltonian before driving was switched on. Thus a ratio of velocities of the pre- and postquench systems, \( \frac{\nu^{\text{in}}(\alpha, k)}{\nu_k(\alpha, k)} \), appears in the formula for the quench, with the orbital magnetization becoming sensitive to van Hove singularities (vHs) \( |\psi^{\text{in}}_{\alpha,k} | = 0 \) of the Floquet bands. We emphasize that vHs are always important whenever a sum on all momenta is involved. Thus vHs play a role even for a thermal occupation of the bands. However, as Eq. (10) shows, the role of vHs, especially those coming from \( \nu_k = 0 \), becomes more important for a quench occupation of the bands than for a thermal equilibrium occupation of the bands.

### III. APPLICATION TO DRIVEN GRAPHENE

We now apply the above formulas for the orbital magnetization to periodically driven (solid-state or artificial) graphene. We choose a driving protocol where \( \mathbf{k}_0 \to \mathbf{k}_0 + A_0 a_0 (\cos(\Omega t) \mathbf{e} - \sin(\Omega t) \mathbf{y}) \) (corresponding to a circularly polarized laser), where \( a_0 \) is the spacing between nearest-neighbor sites, \( A_0 \) is the dimensionless drive amplitude, and \( \Omega \) is the drive frequency. The chosen driving protocol effectively breaks time-reversal symmetry, with the high-frequency limit [52–55] corresponding to the Haldane model [56]. Denoting the hopping amplitude of graphene by \( t_0 \) and the velocity at the Dirac points by \( v_F \), the gap at the Dirac points in the high-frequency limit is \( \approx 2(A_0 v_F)^2/\Omega \) [2], where \( v_F = (3/2)c_0a_0 \) [57].

On varying the drive frequency and amplitude, a rich phase diagram is obtained which includes conventional Chern insulators with edge modes completely characterized by the Chern number, as well as anomalous phases where the Chern number is insufficient to characterize the topology [49,58–61]. The bandwidth of graphene is \( D \approx 6t_0 \). In order to highlight the main physics, we will study three qualitatively different cases.

One is that of high-frequency driving \( \Omega \gg D \), the second is that of a weak \( A_0a_0 \ll 1 \) but resonant drive \( \Omega \lesssim D \), while the third is that of a strong \( A_0a_0 \gg 1 \) and resonant drive \( \Omega < D \) where the Floquet bands bear little resemblance to that of graphene. The high-frequency case shows physics identical to the Haldane model. The other two cases, for our parameters, correspond to anomalous phases.

In particular, the case of weak amplitude but resonant drive [Fig. 1(b), \( \Omega = 5t_0, A_0a_0 = 0.5 \)] has Chern number \( C = 3 \). On a cylinder, the system hosts three chiral edge modes, with two of them of the same chirality and traversing the Floquet zone boundary (red), while the third is of the opposite chirality and traversing the zone center (yellow). The Chern number of 3 measures the change in the chirality from the zone center to the zone boundary but cannot determine the number of edge modes at the two gaps. The second anomalous phase [Fig. 1(c), \( \Omega = 0.5t_0, A_0a_0 = 10 \)] is an example of a low-frequency and high-amplitude drive for which \( \mu = 0 \), but the system still hosts chiral edge modes on the cylinder. In particular, the Floquet zone center (yellow) and boundary (red) each host two chiral edge modes, with all edge modes of the same chirality.

While Fig. 1 is for a system with boundaries, the rest of the paper presents results for a spatially periodic system. The results for the orbital magnetization for the three cases are plotted in Fig. 2. The first, second, and third columns correspond to \( C = 1, 3, \) and 0, respectively, with results presented in units of \( t_0a_0^2/\hbar \) on the left axes, and in Bohr magnetons \( \mu_B = e\hbar/(2m_e) \) on the right axes, where \( m_e \) is the mass of the electron and \( t_0, a_0 \) are for solid-state graphene. We choose a temperature of \( \beta^{-1} = 0.05t_0 \), with the Floquet zone boundaries \( \pm \Omega/2 \) indicated by vertical dashed lines and the Floquet band edges indicated by vertical solid lines. The magnetization \( \mathcal{M} \) for a thermal equilibrium occupation is plotted in the first row, while the second row shows the quench orbital magnetization \( \mathcal{M}_Q \).

As expected, the high-frequency case of \( C = 1 \), for a thermal occupation of the bands, agrees with the Haldane model [Fig. 2(a)], with the orbital magnetization showing structure as \( \mu \) traverses the gap at zero quasienergy. The steep linear rise in \( \mathcal{M} \) across \( \mu = 0 \) is proportional to \( C \). The quench magnetization \( \mathcal{M}_Q \) [Fig. 2(b)], in addition to changing rapidly around \( \mu = 0 \), also shows sharp features due to the dependence of \( \mathcal{M}_Q \) on vHs [see coincidence with sharp structures in the density of states (DOS), purple dashed lines].

The \( C = 3 \) case shows the same behavior as \( C = 1 \) close to \( \mu = 0 \) [Fig. 2(c)]; however, the largest orbital magnetization appear near the Floquet zone boundaries \( \mu = \Omega/2 \) (in contrast, the \( C = 1 \) case has zero orbital magnetization near \( \mu = \pm \Omega/2 \)). We associate the large peaks visible in \( \mathcal{M}, \mathcal{M}_Q \) (see Fig. 4 for the orbital magnetization over the full range) at \( \mu \approx \pm \Omega/2 \) for \( C = 3 \) as an example of bulk-boundary correspondence as two additional edge modes of the opposite
FIG. 2. Orbital magnetization per unit dimensionless area $A/a_0^2$ of the spatially periodic system as a function of chemical potential $\mu$ for Chern numbers $C = 1, 3, 0$, with the latter two being anomalous phases. Floquet bands filled according to a thermal distribution function with $\beta^{-1} = 0.05 t_0$ [(a), (c), and (e)] and according to a sudden quench from graphene at thermal equilibrium with $\beta^{-1} = 0.05 t_0$ [(b), (d), and (f)]. Up to $|m_{\text{max}}| = 3$ [(a) and (b)], $|m_{\text{max}}| = 5$ [(c) and (d)], and $|m_{\text{max}}| = 20$ [(e) and (f)] Floquet harmonics were kept. A $1001 \times 1001$ $k$ grid was used for all panels. The vertical axes on the right indicate values in units of the Bohr magneton $\mu_B = e\hbar/2m_e$. Vertical black dotted lines indicate the Floquet zone boundary $\mu = \pm \Omega/2$. Vertical black solid lines indicate Floquet band edges. (a) The black dashed lines indicate reference results obtained from the Haldane model with next-nearest-neighbor hopping $t_2 = 0.01 t_0$, flux $\phi = 0.5\pi$, and the same temperature. (c) and (e) Black triangles indicate the chemical potential values used in Figs. 3 and 6. (b), (d), and (f) The renormalized Floquet DOS is indicated by purple dashed lines (Fig. 5 gives more details of the DOS). The magnetization changes sign around $\mu = 0$ (all panels) and shows strong peaks at the zone boundaries $\mu = \pm \Omega/2$ for the anomalous case $C = 3$ [(c) and (d)] (see Fig. 4 for orbital magnetization over the full range). The orbital magnetization after the quench [(b), (d), and (f)] is very sensitive to the vHs, making (d) and (f) depend on the $k$ grid close to momenta where vHs are found.

Chirality appear at $\mu = \pm \Omega/2$ relative to $\mu = 0$, enhancing the orbital magnetization. In addition, the sharp features in $M_Q$ coincide with the vHs [Fig. 2(d)].

The final case is $C = 0$ [Figs. 2(e) and 2(f)], where the bands have a nonzero Berry curvature, although it integrates to zero. The nonzero orbital magnetization arises from the nonzero Berry curvature. Since the chirality does not change from the Floquet zone centers to the zone boundaries, the strength of the orbital magnetization is of the same magnitude

FIG. 3. Total magnetization density [integrands of Eqs. (8a)–(8c)] as a function of quasimomentum for $C = 3$ (a) and $C = 0$ (b) and fixed chemical potential $\mu = 2.3 t_0$ (a) and $\mu = 0.1 t_0$ (b) (as indicated by black triangles in Fig. 2). Floquet bands are filled according to a thermal distribution with $\beta^{-1} = 0.05 t_0$. We employed (a) $|m_{\text{max}}| = 5$ and (b) $|m_{\text{max}}| = 20$ Fourier modes and a $1001 \times 1001$ $k$ grid. Cyan (a) and black (b) contours indicate regions where the Berry curvature has significant contributions.

FIG. 4. Same as Fig. 2(d), i.e., the orbital magnetization for a quench occupation of the bands for $C = 3$, but with the full magnetization range shown.
FIG. 5. Floquet bands and Brillouin-zone-averaged density of states (DOS) for the three different driving cases with Chern number $C = 1 \ (a) \text{ and } (b)$, $C = 3 \ (c) \text{ and } (d)$, and $C = 0 \ (e) \text{ and } (f)$] within the first Floquet Brillouin zone, for periodic boundary conditions in $x$ and $y$. Black dashed lines indicate the respective values for the static case (undriven graphene). The inset in (a) shows the driving-induced gap. We assume a Lorentzian broadening of the energy levels with a width of $0.05t_0$ (b), $0.025t_0$ (d), and $0.005t_0$ (f). Numerical parameters such as number of Floquet harmonics and $k$-grid size are the same as for Fig. 2.

at these points. Note that a temperature of $\beta^{-1} = 0.05t_0$ is "high" for this case and as a result the orbital magnetization is smoother and gives nonzero contributions even when $\mu$ is outside the band edges. Despite the Chern number being zero, the orbital magnetization takes values comparable to those of the $C = 1$ case, making it a good probe of anomalous Floquet phases.

The integrand of Eqs. (8a)–(8c), namely, the orbital magnetization density, is shown in Fig. 3 [and that for Eq. (10) is shown in Fig. 6] for a chemical potential where the orbital magnetization takes large values [black triangles in Figs. 2(c) and 2(e)]. The plots highlight the connection between peaks in the orbital magnetization density and peaks in the Berry curvature. A nonzero Berry curvature is a prerequisite for a nonzero orbital magnetization density. However, the distribution functions, defined by the chemical potential $\mu$ and the temperature $\beta^{-1}$, determine which parts of the band Berry curvature actually contribute. In Fig. 3(a) the Dirac points (small cyan circles) do not contribute to the magnetization density because a chemical potential of $\mu = 2.3t_0$ was chosen so that the Dirac points are fully occupied and the Berry curvature contributions from both Dirac bands cancel out (see also Fig. 5). The area around the $\Gamma$ point (big cyan circle), however, is only partly occupied, which together with the peaks in Berry curvature in this region leads to a strong magnetic response. The same reasoning applies to Fig. 3(b), where we chose a different chemical potential, $\mu = 0.1t_0$.

IV. CONCLUSIONS

We derived a general expression for the orbital magnetization of bulk Floquet systems and applied it to periodically driven graphene. The orbital magnetization has a lot more information than the Chern number, showing a significant response even for anomalous phases with $C = 0$. Thus orbital magnetization is a more sensitive probe of Floquet induced topology than traditional metrics such as the Chern number. Reference [35] showed that when the bulk states are localized, the orbital magnetization is quantized and related to a 3D winding number [9]. However, in our setting where the
FIG. 6. The orbital magnetization density, but with occupations according to a sudden quench [Eq. (9)] and with the magnetization density corresponding to the integrand of Eq. (10). Black contours in both panels indicate regions where the Berry curvature has significant contributions. White dots in both panels indicate regions that have vanishing Floquet band velocity in the \( y \) direction and therefore are related to the vHs. The initial state, i.e., the state of undriven graphene, is taken to be in thermal equilibrium at a temperature \( \beta^{-1} = 0.05 t_0 \) and a chemical potential \( \mu = 2.3 t_0 \) (a) and \( \mu = 0.1 t_0 \) (b). Grid size and number of Floquet harmonics are the same as in Fig. 3.

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\(\mathbf{p} = q \hat{y} \) and \(\mathbf{p} = -q \hat{y}\). Thus we arrive at Eq. (2), which we rewrite below for convenience:

\[
V(t) = \mathbf{A}_q \cdot \mathbf{J}_q(t) + \text{c.c.,} \quad \text{(A7)}
\]

with \(\mathbf{A}_q = -\hat{\varepsilon} B/(2iq)\).

\[
\delta |\psi_{nk}(t)\rangle = -i \int_0^t dt' U(t, t') V(t') U(t', 0) |\psi_{nk}(0)\rangle
\]

\[
= -i \sum_{n', k'} |\psi_{n', k}(t)\rangle \int_0^t dt' |\psi_{n', k}(t')\rangle U(t, t') V(t') U(t', 0) |\psi_{nk}(0)\rangle
\]

\[
= -i \sum_{n', k'} |\psi_{n', k}(t)\rangle \int_0^t dt' |\psi_{n', k}(0)\rangle U(0, t) U(t, t') V(t') U(t', 0) |\psi_{nk}(0)\rangle
\]

\[
= -i \sum_{n', k'} e^{-i\varepsilon_{n', k'} t} |\psi_{n', k}(t)\rangle \int_0^t dt' |\phi_{n', k}(t')\rangle V(t') |\phi_{nk}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n', k'} t'}
\]

\[
= -i \sum_{n', k'} e^{-i\varepsilon_{n', k'} t} |\psi_{n', k}(t)\rangle \int_0^t dt' |\phi_{n', k}(t')\rangle V(t') |\phi_{nk}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n', k'} t'} \tag{A9}
\]

Above we have used that the Floquet states constitute a complete set of states, and we have employed the identity \(U(0, t) U(t, t') = U(0, t')\).

Note that to \(O(B)\) the change in the quasienergy is given by \(\delta \varepsilon_{n, k} = \langle \phi_{nk} | V(t) | \phi_{nk} \rangle\). \(V(t)\) is proportional to the velocity operator, and the average velocity vanishes in an eigenstate, \(\langle \phi_{nk} | V(t) | \phi_{nk} \rangle = 0\). Therefore the change in the quasienergy comes entirely from the change in the Floquet eigenstate, i.e., \(\delta |\phi_{nk}(t)\rangle = e^{i\varepsilon_{nk} t} \delta |\psi_{nk}(t)\rangle\) to \(O(B)\). Substituting for \(V(t)\), the change in the quasienergy to \(O(B)\) is

\[
\delta |\phi_{nk}(t)\rangle = \frac{B}{2q} \sum_n |\phi_{n, k+q}(t)\rangle \int_0^t dt' |\phi_{n, k+q}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n, k+q} t'}
\]

\[
- \frac{B}{2q} \sum_n |\phi_{n, k-q}(t)\rangle \int_0^t dt' |\phi_{n, k-q}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n, k-q} t'} \tag{A10}
\]

Above,

\[
v_{k+q/2}^x(t) = v^x(k_x, k_y + q/2, t) = \frac{\partial}{\partial k_x} H(k_x, k_y + q/2, t). \tag{A11}
\]

Thus the change in the average quasienergy to \(O(B)\) is

\[
\delta E(t) = \sum_{n, k} f_{nk} \left[ \langle \phi_{nk}(t) | H_F | \delta \phi_{nk}(t) \rangle + \text{c.c.} \right]
\]

\[
= \frac{B}{2q} \sum_{n, n', k, k'} f_{nk} \left[ \langle \phi_{nk}(t) | H_F | \phi_{n', k+q}(t) \rangle \int_0^t dt' |\phi_{n', k+q}(t')\rangle |v_{k+q/2}^x(t')\rangle |\phi_{nk}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n', k+q} t'}
\]

\[
- \langle \phi_{nk}(t) | H_F | \phi_{n', k-q}(t) \rangle \int_0^t dt' |\phi_{n', k-q}(t')\rangle |v_{k-q/2}^x(t')\rangle |\phi_{nk}(t')\rangle e^{-i\varepsilon_{nk} t + i\varepsilon_{n', k-q} t'}
\]

\[
+ \langle \phi_{n', k+q}(t) | H_F | \phi_{nk}(t) \rangle \int_0^t dt' |\phi_{n', k+q}(t')\rangle |v_{k+q/2}^x(t')\rangle |\phi_{nk}(t')\rangle e^{i\varepsilon_{nk} t - i\varepsilon_{n', k+q} t'}
\]

\[
- \langle \phi_{n', k-q}(t) | H_F | \phi_{nk}(t) \rangle \int_0^t dt' |\phi_{n', k-q}(t')\rangle |v_{k-q/2}^x(t')\rangle |\phi_{nk}(t')\rangle e^{i\varepsilon_{nk} t - i\varepsilon_{n', k-q} t'} \right] \tag{A12}
\]
In the second and last terms we find it convenient to shift $k \to k + q$, and interchange $n, n'$. This gives

$$\delta E(t) = \frac{B}{2q} \sum_{n,n',k} \left( f_{nk} - f_{n',k+q} \right) \left[ \langle \phi_{nk}(t) | H_F | \phi_{n'k+q}(t) \rangle \right] \int_0^t dt' \langle \phi_{nk+q}(t') | v'_{k+q/2}(t') \rangle \langle \phi_{n'k}(t') e^{-i\epsilon_{nk}(t' - t) + i\epsilon_{n'k+q}(t' - t)} + c.c. \rangle = BM(t). \tag{A13}$$

Note that the term multiplying $B/(2q)$ is $O(q)$ because it is a product of a term which is symmetric under exchange of $n, n'$ when $q = 0$ (the term in square brackets for $q = 0$) and another which is antisymmetric under exchange of $n, n'$ when $q = 0$ [the term $(f_{nk} - f_{n',k})$]. Thus the prefactor of $B/2q$ vanishes when $q = 0$. Now the goal is to expand this prefactor to $O(q)$ and determine $M(t)$. When $n = n'$, the $O(q)$ correction comes from Taylor-expanding $(f_{nk} - f_{n',k})$ to $O(q)$, while for the $n \neq n'$ term, the $O(q)$ correction comes from Taylor-expanding $(\phi_{nk}(t)|H_F|\phi_{n'k+q}(t))$. Thus, in what follows, we may replace $v'_{k+q/2}$ by $v'_k$.

We introduce the following notation for the Fourier transform of matrix elements between Floquet quasimodes:

$$[\hat{O}^{m}_{nk,n'k+q}] = \frac{1}{T} \int_0^T dt e^{-im\Omega t} \langle \phi_{nk}(t') \hat{O} \phi_{n'k+q}(t') \rangle. \tag{A14}$$

In Eq. (A13) we perform the $t'$ integral by including a small imaginary part and extending the integral to $\infty$. This gives the following expression for the average orbital magnetization:

$$\overline{M} = \lim_{q \to 0} \frac{1}{2q} \sum_{n,n',k,m} \left[ (f_{nk} - f_{n',k+q}) | H_{F} | v^{*}(k) |_{n'k+q,nk}^m \frac{i}{\epsilon_{nk} + m\Omega - \epsilon_{n'k+q}} + c.c. \right], \tag{A15}$$

where

$$| H_{F} | v^{*}(k) |_{n'k+q,nk}^m = \frac{1}{T} \int_0^T dt e^{im\Omega t} \langle \phi_{nk}(t') \hat{O} \phi_{n'k+q}(t') \rangle = \frac{1}{2} \left[ \epsilon_{n'k+q} + \epsilon_{nk} + m\Omega \right] \sum_{m} \langle \phi_{nk}^{m} | \phi_{n'k+q}^{m} \rangle. \tag{A16}$$

and

$$[v^{*}(k)]_{n'k+q,nk}^m = \frac{1}{T} \int_0^T dt e^{im\Omega t} \langle \phi_{nk}(t') \hat{O} \phi_{n'k+q}(t') \rangle = \frac{1}{2} \left[ \epsilon_{n'k+q} + \epsilon_{nk} + m\Omega \right] \sum_{m} \langle \phi_{nk}^{m} | \phi_{n'k+q}^{m} \rangle. \tag{A17}$$

Equation (A15) corresponds to Eq. (6).

In order to Taylor expand in $q$, it is convenient to separate the terms into $n \neq n'$ and $n = n'$ as follows:

$$\overline{M} = \lim_{q \to 0} \frac{1}{2q} \sum_{n,n',k,m} \left[ (f_{nk} - f_{n',k+q}) | H_{F} | v^{*}(k) |_{n'k+q,nk}^m \frac{i}{\epsilon_{nk} + m\Omega - \epsilon_{n'k+q}} + c.c. \right]$$

$$+ \lim_{q \to 0} \frac{1}{2q} \sum_{n,k,m} \left[ (f_{nk} - f_{nk+q}) | H_{F} | v^{*}(k) |_{nk,nk}^m \frac{i}{\epsilon_{nk} + m\Omega - \epsilon_{nk+q}} + c.c. \right]. \tag{A18}$$

Substituting for matrix elements of $v_k$ and $H_F$, we obtain

$$\overline{M} = \lim_{q \to 0} \frac{1}{2q} \sum_{n,n',k,m} \left[ (f_{nk} - f_{n',k+q}) \left[ \frac{1}{2} \left[ \epsilon_{n'k+q} + \epsilon_{nk} + m\Omega \right] \sum_{m} \langle \phi_{nk}^{m} | \phi_{n'k+q}^{m} \rangle \right] \right.$$

$$\times \left. \left( \epsilon_{nk} - \epsilon_{nk+q} + m\Omega \right) \sum_{m} \langle \phi_{nk}^{m} | \phi_{n'k+q}^{m} \rangle \right] \left( \overline{M} \frac{i}{\epsilon_{nk} + m\Omega - \epsilon_{nk+q}} + c.c. \right]. \tag{190}$$
\[
\lim_{q \to 0} \frac{1}{2q} \sum_{n,k,m} (f_{nk} - f_{nk+q}) \left[ \frac{1}{2} (\epsilon_{n,k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m)}|\partial_k \epsilon_{nk}^{(m)}) + \frac{i}{\epsilon_{nk} + m\Omega - \epsilon_{n,k} + q} + c.c. \right].
\]

(A19)

When \(n \neq n'\), the term in the first line above \(\sum_{m}(\phi_{nk}^{(m-m)|\phi_{nk}^{(m)}})\) is such that when \(q = 0\), this term vanishes, i.e., \(\sum_{m} (\phi_{nk}^{(m-m)|\phi_{nk}^{(m)}}) = 0\). This is because \(\langle \phi_{a}(t)|\phi_{b}(t)\rangle = 0\) and therefore all the Fourier components of this overlap should be zero. Thus, for the \(n \neq n'\) term, we need to Taylor expand only \(\sum_{m} (\phi_{nk}^{(m-m)|\phi_{nk}^{(m)}})\) to \(O(q)\). In addition, for the \(n = n'\) term, the expression

\[
\sum_{m_1} (\phi_{nk}^{(m-m)}|\phi_{nk}^{(m)}) \times \sum_{m_2} (\phi_{nk+q}^{(m+m)}|\phi_{nk}^{(m)}) \times \sum_{m_2} (\phi_{nk+q}^{(m)}|\partial_k \epsilon_{nk}^{(m)}) = \left[ \frac{1}{T} \int_0^T dt e^{-i\Omega t} \langle \phi_{nk}(t)|\phi_{nk+q}(t)\rangle \right] \left[ \frac{1}{T} \int_0^T dt' e^{i\Omega t'} \langle \phi_{nk+q}(t')|\phi_{nk}(t')\rangle \right]
\]

(A20)

and is purely real. Thus the term proportional to \(\langle \partial_k \epsilon_{nk}\rangle\) does not contribute due to the \(i\) factor multiplying it. This leads to

\[
\begin{align*}
\mathcal{M} &= \frac{1}{2} \sum_{n \neq n', k, m} (f_{nk} - f_{nk'}) \left[ \frac{1}{2} (\epsilon_{n',k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) \left\{ \sum_{m_2} (\phi_{nk}^{(m+m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} + c.c. \right] \\
&+ \lim_{q \to 0} \frac{1}{2q} \sum_{n,k,m} (f_{nk} - f_{nk+q}) \left[ \frac{1}{2} (\epsilon_{n,k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) \left\{ \sum_{m_2} (\phi_{nk+q}^{(m+m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} + c.c. \right].
\end{align*}
\]

(A21)

As before, since \(\langle \phi(t)|\phi(t)\rangle = 1\), the term \(\sum_{m_1} (\phi_{nk}^{(m-m)}|\phi_{nk}^{(m)}) = \delta_{m=0} + O(q)\delta_{m\neq0}\). This is because, when \(q = 0\), all time dependence should vanish (i.e., \(m = 0\)). So to leading order in \(q\), only the \(m = 0\) term in the second line survives, giving

\[
\begin{align*}
\mathcal{M} &= \frac{1}{2} \sum_{n \neq n', k, m} (f_{nk} - f_{nk'}) \left[ \frac{1}{2} (\epsilon_{n',k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) \left\{ \sum_{m_2} (\phi_{nk}^{(m+m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} + c.c. \right] \\
&+ \lim_{q \to 0} \frac{1}{2q} \sum_{n,k,m} (f_{nk} - f_{nk+q}) \left[ \frac{1}{2} (\epsilon_{n,k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) + c.c. \right].
\end{align*}
\]

(A22)

Now we divide and multiply by \((\epsilon_{nk} - \epsilon_{nk+q})\) in the second term and use the identity

\[
(\epsilon_{nk} - \epsilon_{nk+q}) \sum_{m_1} (\phi_{nk+q}^{(m)}|\partial_k \epsilon_{nk}^{(m)}) = (\phi_{nk+q}|\epsilon_{nk} - H_F|\partial_k \epsilon_{nk}) \approx q [\partial_k \epsilon_{nk}|\epsilon_{nk} - H_F|\partial_k \epsilon_{nk}]
\]

(A23)

to obtain

\[
\begin{align*}
\mathcal{M} &= \frac{1}{2} \sum_{n \neq n', k, m} (f_{nk} - f_{nk'}) \left[ \frac{1}{2} (\epsilon_{n',k} + \epsilon_{n,k} + m\Omega) \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) \left\{ \sum_{m_2} (\phi_{nk}^{(m+m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} + c.c. \right] \\
&+ \lim_{q \to 0} \frac{1}{2q} \sum_{n,k,m} (f_{nk} - f_{nk+q}) \epsilon_{nk} \left[ i [\partial_k \epsilon_{nk}|\epsilon_{nk} - H_F|\partial_k \epsilon_{nk}] + c.c. \right].
\end{align*}
\]

(A24)

Since \(\langle \phi_{n,k}(t)|\phi_{n,k}(t)\rangle = \text{constant in time}, \partial_k [(\phi_{n,k}(t)|\phi_{n,k}(t))] = 0\). This must hold for each Fourier component giving \(\partial_k \sum_{m_1} (\phi_{nk}^{(m+m)}|\phi_{nk}^{(m)}) = 0\) for each \(m\). Using this, we can move the derivatives between the bras and kets to show that

\[
\sum_{n \neq n', k, m} (f_{nk} - f_{nk'}) m\Omega \left[ \left\{ \sum_{m_1} (\phi_{nk}^{(m-m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} \left\{ \sum_{m_2} (\phi_{nk}^{(m+m)}|\partial_k \epsilon_{nk}^{(m)}) \right\} + c.c. \right] = 0.
\]

(A25)

Similar manipulations lead to

\[
\mathcal{M} = -\frac{1}{2} \text{Im} \left[ \sum_{n',n,k,m} f_{n',k} (\epsilon_{n',k} + \epsilon_{n,k}) \left\{ \sum_{m_1} (\phi_{n,k}^{(m-m)}|\phi_{n,k}^{(m)}) \right\} \right] \\
+ \frac{i}{2} \text{Im} \left[ \sum_{n',n,k,m} f_{n',k} (\epsilon_{n',k} + \epsilon_{n,k}) \left\{ \sum_{m_1} (\phi_{n,k}^{(m-m)}|\phi_{n,k}^{(m)}) \right\} \right] \\
+ \text{Im} \left[ \sum_{n,k} \lim_{q \to 0} (f_{nk} - f_{nk+q}) \epsilon_{nk} \left[ \partial_k \epsilon_{nk}(t)|\epsilon_{nk} - H_F|\partial_k \epsilon_{nk}(t) \right] \right].
\]

(A26)
The above can be written as
\[ M = -\frac{1}{2} \text{Im} \left[ \sum_{n,n',k} f_{nk} \langle \partial_k \phi_{nk}(t) | \epsilon_{nk} + H_F | \partial_k \phi_{nk}(t) \rangle \right] + \frac{1}{2} \text{Im} \left[ \sum_{n,n',k} f_{nk} \langle \partial_k \phi_{n'k}(t) | \epsilon_{n'k} + H_F | \partial_k \phi_{n'k}(t) \rangle \right] \]
\[ + \text{Im} \left[ \sum_{n,k} \lim_{q \to 0} \left( f_{nk} - f_{nk+q} \right) \frac{\epsilon_{nk} - \epsilon_{n,k+q}}{\epsilon_{nk} - \epsilon_{n,k+q}} \langle \partial_k \phi_{nk}(t) | \epsilon_{nk} - H_F | \partial_k \phi_{nk}(t) \rangle \right] \] (A28)
\[ + \text{Im} \left[ \sum_{n,k} \lim_{q \to 0} \left( f_{nk} - f_{nk+q} \right) \frac{\epsilon_{nk} - \epsilon_{n,k+q}}{\epsilon_{nk} - \epsilon_{n,k+q}} \langle \partial_k \phi_{nk}(t) | \epsilon_{nk} - H_F | \partial_k \phi_{nk}(t) \rangle \right] \] (A29)

Now we remove the complete set of states, \( \sum_n | \phi_{nk}(t) \rangle \langle \phi_{nk}(t) | = 1 \), and by noting that the second line is simply the complex conjugate of the first, we obtain
\[ M = - \text{Im} \left[ \sum_{n,k} f_{nk} \langle \partial_k \phi_{nk}(t) | \epsilon_{nk} + H_F | \partial_k \phi_{nk}(t) \rangle \right] + \text{Im} \left[ \sum_{n,k} \lim_{q \to 0} \left( f_{nk} - f_{nk+q} \right) \frac{\epsilon_{nk} - \epsilon_{n,k+q}}{\epsilon_{nk} - \epsilon_{n,k+q}} \langle \partial_k \phi_{nk}(t) | \epsilon_{nk} - H_F | \partial_k \phi_{nk}(t) \rangle \right] \] (A30)

Multiplying by \(-e/\hbar\), and writing \( \text{Im}(a) = (a - a^*)/2i \), we obtain
\[ M = -\frac{e}{2\hbar} \text{Im} \left[ \sum_{n,k} f_{nk} \langle \partial_k \phi_{nk}(t) | (\epsilon_{nk} + H_F) \times | \partial_k \phi_{nk}(t) \rangle - \lim_{q \to 0} \left( f_{nk} - f_{nk+q} \right) \frac{\epsilon_{nk} - \epsilon_{n,k+q}}{\epsilon_{nk} - \epsilon_{n,k+q}} \langle \partial_k \phi_{nk}(t) | (\epsilon_{nk} - H_F) \times | \partial_k \phi_{nk}(t) \rangle \right] \] (A31)

The above is Eq. (7).

### Simplified formulas for the two-band model

When there are only two bands \( n = u, d \) as in the case of periodically driven graphene, the formulas for the orbital magnetization can be simplified. For the orbital magnetization under assumptions of thermal equilibrium, and taking into account corrections from \( \mu(N) \), we have
\[ M = -\frac{e}{2\hbar} \text{Im} \left[ \sum_k (f_{dk} - f_{uk})(\epsilon_{d,k} + \epsilon_{u,k} - 2\mu)F_{xy}(k, t) - (\epsilon_{d,k} - \epsilon_{u,k})F_{xy}(k, t) \sum_{n=d,u} f_{nk} (\epsilon_{nk} - \mu) \right] \] (A32)

For the orbital magnetization for a quench occupation of the bands, and looking at corrections only to \( E \), we have
\[ M_Q = -\frac{e}{2\hbar} \text{Im} \left[ \sum_k (f_{dk} - f_{uk})(\epsilon_{d,k} + \epsilon_{u,k})F_{xy}(k, t) \right. \]
\[ - (\epsilon_{d,k} - \epsilon_{u,k})F_{xy}(k, t) \sum_{\alpha} \left[ \langle \phi_{d,k} | \psi_{uk}^{\text{in}}(\alpha, k) \rangle^2 \partial_{\alpha} f_{uk}^{\text{in}} \frac{v_{\alpha}(d, k)}{v_{\alpha}(u, k)} \epsilon_{d,k} + \langle \phi_{uk} | \psi_{uk}^{\text{in}}(\alpha, k) \rangle^2 \partial_{\alpha} f_{uk}^{\text{in}} \frac{v_{\alpha}(u, k)}{v_{\alpha}(u, k)} \epsilon_{u,k} \right. \]
\[ + \partial_{\alpha} f_{uk}^{\text{in}} \langle \phi_{d,k} | \psi_{uk}^{\text{in}}(\alpha, k) \rangle^2 \epsilon_{d,k} + \partial_{\alpha} f_{uk}^{\text{in}} \langle \phi_{uk} | \psi_{uk}^{\text{in}}(\alpha, k) \rangle^2 \epsilon_{u,k} \left. \right] \] (A34)

Above, \( F_{xy} \) is the time average of the Berry curvature, and \( \alpha \) are the bands of the system before the quench. Note that when there is particle-hole symmetry \( \epsilon_{d,k} = -\epsilon_{u,k} \), then the first term in the above equations proportional to \( \epsilon_{d,k} + \epsilon_{u,k} \) does not contribute. In addition, when the system is in thermal equilibrium at zero temperature, with one band fully occupied, and the other empty, then Eq. (A32) gives \( M_A = -(e/\hbar)C\mu/2\pi \).

### APPENDIX B: VAN HOVE SINGULARITIES

For the sake of completeness, in Fig. 4 we show the same data as in Fig. 2(d) but with the full magnetization range. Figure 5 shows the Floquet band structure within the first Floquet Brillouin zone along the Dirac point for the three driving cases with \( C = 1 \) [Fig. 5(a)], \( C = 3 \) [Fig. 5(c)], and \( C = 0 \) [Fig. 5(e)]. For better orientation the static bands are shown by black dashed lines. The inset in Fig. 5(a) shows the Dirac point with the light-induced gap opening. The corresponding density of states is presented in Figs. 5(b), 5(d), and 5(f).

### APPENDIX C: ORBITAL MAGNETIZATION DENSITY FOR THE QUENCH

In Fig. 6 we present the \( k \)-resolved orbital magnetization density after a sudden quench, as described by Eq. (10), for the driving schemes corresponding to \( C = 3 \) [Fig. 6(a)] and \( C = 0 \) [Fig. 6(b)]. We use the same parameters for \( \mu \) and \( \beta^{-1} \) as for Fig. 3, where \( \mu \) and \( \beta^{-1} \) now characterize
the state before the quench, i.e., the temperature and occupation of undriven graphene. Due to the nonequilibrium occupations, the magnetization shows more complexity, for example, clouds of nonzero magnetization density at the Dirac point position [Fig. 6(a)]. In contrast, for thermal equilibrium occupation of the bands, and for the same chemical potential, the magnetization density at the Dirac points is zero [cf. Fig. 3(a)].


