

Particle dynamics and ergodicity breaking in twisted-bilayer optical latticesGanesh C. Paul ¹, Patrik Recher,^{1,2} and Luis Santos³¹*Institut für Mathematische Physik, Technische Universität Braunschweig, D-38106 Braunschweig, Germany*²*Laboratory for Emerging Nanometrology Braunschweig, D-38106 Braunschweig, Germany*³*Institut für Theoretische Physik, Leibniz Universität, 30167 Hannover, Germany*

(Received 20 June 2023; accepted 26 September 2023; published 2 November 2023)

Recent experiments have realized a twisted-bilayer-like optical potential for ultracold atoms, which in contrast to solid-state setups may allow for an arbitrary ratio between the inter- and intralayer couplings. For commensurate moiré twistings, a large-enough interlayer coupling results in particle transport dominated by channel formation. For incommensurate twistings, the interlayer coupling acts as an effective disorder strength. Whereas for weak couplings the whole spectrum remains ergodic, at a critical value part of the eigenspectrum transitions into multifractal states. A similar transition may be observed as well as a function of an energy bias between the two layers. Our theoretical study reveals atoms in a twisted-bilayer system of square optical lattices as an interesting platform for the study of ergodicity breaking and multifractality.

DOI: [10.1103/PhysRevA.108.053305](https://doi.org/10.1103/PhysRevA.108.053305)**I. INTRODUCTION**

Twisted-bilayer graphene [1–5] has attracted broad attention owing to the observation of unconventional superconducting [6–8] and correlated insulating behavior [9–12]. A small rotation of one of the layers leads to the vanishing of Fermi velocity around the Dirac point, giving rise to an almost flat band [1,2,13,14]. These quasi-flat bands, ideal to observe strongly correlated phenomena, are obtained only for very small twist angles $\lesssim 1^\circ$ in solid state systems, as the interlayer coupling is much smaller than the intralayer one [14–16].

Ultracold gases in optical potentials may provide an interesting highly controllable platform for the study of the physics of twisted-bilayer lattices. These systems allow for a basically arbitrary ratio between interlayer and intralayer couplings. In addition, ultracold gases are, in principle, defect-free, although suitable impurities can be added in a controllable way, making this platform ideal for understanding the effects of disorder. Different proposals have been recently put forward [17–20] to simulate twisted-bilayer-like potentials using ultracold atoms. In particular, Ref. [17] has proposed the use of two internal states that (in a synthetic dimension) play the role of the two layers. The twisted lattices result from a state-dependent optical potential, such that one state experiences an optical lattice tilted at an angle from the lattice experienced by the other state. A microwave or two-photon Raman coupling induces an effective interlayer hopping. An alternative proposal was introduced in Ref. [18], also using a synthetic dimension for the bilayer geometry, with lattices without twisting but with a spatially dependent interlayer hopping. Very recently, ideas similar to those of Ref. [17] have been employed to realize experimentally for the first time a twisted-bilayer optical potential [21], in which a Bose-Einstein condensate was loaded, opening an interesting novel platform for the study of superfluids in twisted-bilayer lattices.

In recent years, atoms in optical lattices have been shown to provide a suitable platform to study experimentally both single- and many-body localization. In particular, the use of bichromatic lattices has allowed for the realization of the one-dimensional quasi-disordered (Aubry-André) model [22–24], characterized by a phase transition at a critical quasi-disorder strength between a fully localized and a fully ergodic eigenstate spectrum. Recent experiments have also realized two-dimensional optical quasi-crystals [25]. An alternative, also highly controllable setup for the study of localization, is provided by photonic lattices, in which a spatial direction plays the role of an effective time dimension. Interestingly, in addition to one-dimensional geometries [26], very recent experiments have analyzed the wave dynamics in two-dimensional (monolayer) photonic moiré-like lattices [27], revealing a localization-to-delocalization transition [28].

The abovementioned realization of twisted-bilayer-like optical lattices opens intriguing questions concerning particle dynamics and ergodicity breaking in these potentials, which we theoretically address in this paper for the case of coupled square lattices. Whereas in solid-state setups the interlayer coupling is very small compared to the intralayer one (typically 5 to 10 times smaller), it may be potentially dominant in optical lattice platforms, resulting for commensurate twistings in particle transport dominated by the formation of channels. Moreover, for incommensurate twistings and due to the finite spatial range of the interlayer coupling, this coupling acts as an effective quasi-disorder strength. Whereas below a given coupling threshold the whole eigenspectrum remains ergodic, at a critical coupling part of the spectrum experiences a transition into nonergodic extended (multifractal) states. A similar transition may be observed, alternatively, by employing an energy bias between the two layers. Our results show, hence, that the combination of moderately strong interlayer coupling and incommensurate twist angles makes twisted-bilayer optical lattices an alternative suitable platform for the study of multifractality.

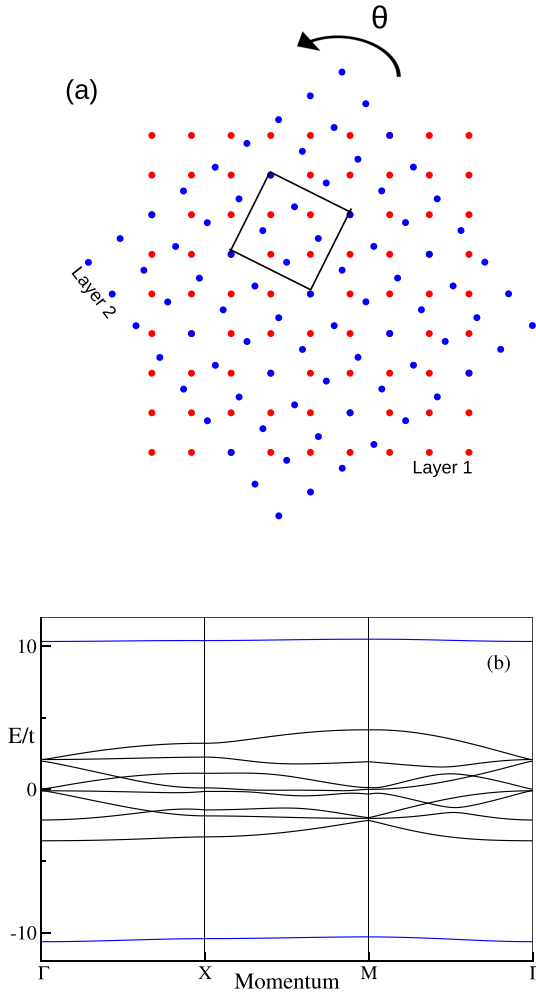


FIG. 1. (a) Twisted-bilayer structure for a twisting angle $\theta = \theta(2, 1)$. Red (blue) sites correspond to layer 1 (2). The elementary cell is indicated by a black square. (b) Band structure along the high-symmetry line Γ -X-M- Γ for the same lattice with $t_{\perp}/t = 10$, $l_0/a = 0.15$, and $\Delta = 0$. The quasi-flat bands are indicated in blue.

The remainder of the paper is organized as follows. In Sec. II, we describe the twisted-bilayer model. Section III is devoted to the particle dynamics in the case of commensurate twistings, while the dynamics in incommensurately twisted bilayers is presented in Sec. IV. The impact of interlayer bias is discussed in Sec. V. Finally, we summarize our results in Sec. VI.

II. OPTICAL TWISTED BILAYERS

In the following, we consider two layers of square optical lattices [see Fig. 1(a)], where one layer is twisted by an angle θ with respect to the other. A possible way of implementing such an optical potential, recently proposed in Ref. [17] and realized experimentally in Ref. [21], employs an atom in two different internal states. In this scenario, which we assume below, the bilayer structure is provided by the synthetic dimension given by the two internal states, whereas a state-dependent potential results in the twisted-bilayer

geometry. For a square lattice, a moiré pattern is achieved for $\theta = \theta(m, n)$, where $\theta(m, n) = \arccos(\frac{2mn}{m^2+n^2})$ ($n, m \in \mathbb{Z}$).

We are interested in the dynamics of a single particle, and hence we do not account for interaction terms. The results should remain, however, valid as long as the lattice filling is sparse enough. The system is characterized by the Hamiltonian:

$$H = -\Delta \sum_{\alpha, j} (-1)^{\alpha} |\alpha, j\rangle \langle \alpha, j| - t \sum_{\alpha=1,2} \sum_{\langle j, j' \rangle} |\alpha, j\rangle \langle \alpha, j'| - \sum_{j, j'} t_{\perp}(j, j') [|1, j\rangle \langle 2, j'| + \text{H.c.}], \quad (1)$$

where $|\alpha, j\rangle$ is the state in which the particle is in layer α in site $j = (j_x, j_y)$. The site in layer 1 (2) is located at the position $\vec{R}_{1,j} = j_x \vec{e}_x + j_y \vec{e}_y$ ($\vec{R}_{2,j} = j_x \vec{e}_{x'} + j_y \vec{e}_{y'}$), with $\vec{e}_{x'} = \cos \theta \vec{e}_x + \sin \theta \vec{e}_y$ and $\vec{e}_{y'} = -\sin \theta \vec{e}_x + \cos \theta \vec{e}_y$. The first term of Eq. (1) denotes an uniform bias between the layers, characterized by the bias strength Δ , which in the synthetic dimension scenario amounts for a level-dependent shift (e.g., using a magnetic field or an optically imposed Stark-shift). The intralayer hopping, given by the rate t , occurs only to nearest neighbors, denoted by $\langle j, j' \rangle$.

The rate $t_{\perp}(j, j')$ characterizes the hopping between a site j in layer 1 and site j' in layer 2. In the considered scenario, such a coupling occurs between different internal states, and it is given either by a microwave or two-photon optical Raman coupling. In stark contrast to regular cubic lattices, the sites in both layers are generally not on top of each other. For a sufficiently strong lattice, we may approximate the on-site Wannier functions in each layer as a Gaussian of $l_0 = \frac{a}{\pi s^{1/4}}$, with a being the lattice spacing characterizing the square lattice in both layers and s being the lattice depth in recoil units. As a result, the interlayer hopping acquires the following form [17]:

$$t_{\perp}(j, j') = t_{\perp} e^{-|\vec{R}_{1,j} - \vec{R}_{2,j'}|^2 / (4l_0^2)}. \quad (2)$$

Reference [17] considered for simplicity the case of $l_0 = 0$, in which only sites exactly on top of each other may undergo interlayer coupling with a rate t_{\perp} . We show below that the finite Gaussian width l_0 plays a crucial role in the actual particle dynamics in the bilayer-like optical potential. Note as well, that in contrast to solid-state scenarios where typically $t_{\perp}/t \ll 1$ and is hardly tunable, in the optical lattice platform t_{\perp} is easily tunable and may be much larger than the intralayer hopping t . As shown below, this opens interesting possibilities for the dynamics for both commensurate and incommensurate twisting angles.

In the following, we consider for simplicity, unless otherwise indicated, a twist angle θ in the vicinity of the magic angle $\theta(2, 1) = 36.87^\circ$, although our conclusions are general for the dynamics in the vicinity of any commensurate twist angle. The choice of $\theta(2, 1)$ is justified by the small number of sites (five in each layer) per moiré unit cell, which greatly simplifies the analysis of the system, compared to solid-state platforms, where due to the very small twisting angle, a moiré cell may contain tens of thousands of sites [13]. For $\theta = \theta(2, 1)$, only two of the ten sites are on top of each other (directly connected sites) [see Fig. 1(a)].

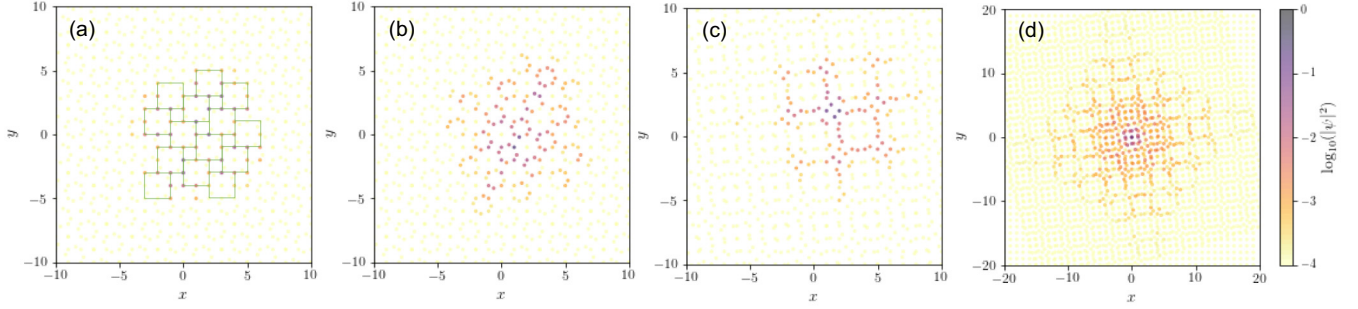


FIG. 2. Commensurate twisting. Panels (a) and (b) show the probability density at time $\tau = 2/t$ for $\theta = \theta(2, 1)$, $t_{\perp}/t = 100$, and $l_0 = 0$ and $l_0/a = 0.15$, respectively, for a particle initially at site $j = (1, 0)$ of layer 1. In panel (a), green lines indicate the effective lattice with out-projected sites mentioned in the text. Panel (c) depicts the probability density at $\tau = 6/t$ for $\theta = \theta(4, 3)$ for $l_0/a = 0.15$, and at $t_{\perp} = 50t$ for a particle initially at site $j = (2, 2)$ of layer 1. Panel (d) shows, at $\tau = 10/t$, the particle distribution for an initial Gaussian wave packet in layer 1, $\frac{1}{\sqrt{2\pi\sigma^2}} e^{-|\vec{R}_{1,j} - \vec{R}_{1,j_0}|^2/2\sigma^2}$, of width $\sigma/a = 1$, centered at site $j_0 = (0, 0)$, in a lattice with $t_{\perp} = 10t$ and $\theta(4, 3)$. In all the figures, the x and y axes are in units of the lattice spacing a .

III. PARTICLE DYNAMICS IN MOIRÉ LATTICES

Figure 1(b) shows the corresponding band structure for $\theta = \theta(2, 1)$, $\Delta = 0$, and $t_{\perp} = 10t$, along the symmetry line Γ - X - M - Γ , with $\Gamma = (0, 0)$, $X = \frac{2\pi}{5a}(\frac{1}{2}, 1)$, and $M = \frac{2\pi}{5a}(-\frac{1}{2}, \frac{3}{2})$. As shown in Fig. 1(b), when $t_{\perp}/t \gg 1$, the spectrum presents an uppermost and a lowest band (in blue), with an energy $\simeq \pm t_{\perp}$, which originate from directly connected sites. The large interlayer coupling projects out those sites, which build a separate grid. Since directly connected sites are far apart, hopping between them only occurs in high order in perturbation theory, resulting in almost flat bands. Particles initially placed in those sites remain, hence, basically localized.

The rest of the not directly connected sites form separated bands around zero energy. For $l_0 = 0$, they cannot participate directly in the interlayer hopping. For $t_{\perp} \rightarrow \infty$, a particle starting in one of those sites in the upper (lower) layer would be disconnected from the lower (upper) layer. The particle would experience an effective square lattice with out-projected sites (the directly connected ones), characterized by four sites in the elementary cell [see Fig. 2(a)]. As a result, the band spectrum would present two degenerate sets of four bands. This degeneracy is lifted, even for $l_0 = 0$, at finite $t_{\perp}/t \gg 1$ due to processes of order $O(t^2/t_{\perp})$ induced by virtual couplings between directly and nondirectly connected sites.

A single particle in the twisted optical lattice is described by the state $\sum_{\alpha,j} c_{\alpha,j}(t) |\alpha, j\rangle$, with the probability amplitudes given by the Schrödinger equations:

$$i\hbar \frac{\partial c_{\alpha,j}}{\partial \tau} = -t \sum_{(j')} c_{\alpha,j'} - \sum_{j'} t_{\perp}(j, j') c_{\bar{\alpha},j'}, \quad (3)$$

with $\bar{\alpha} = 2$ (1) for $\alpha = 1$ (2). We solved these equations by standard Runge-Kutta methods, using absorptive boundary conditions, which allow for the analysis of the particle expansion and eventual localization without the need of very large lattices.

We consider the evolution of a particle initially placed in a not directly connected site. As mentioned above, if $l_0 = 0$, a sufficiently large t_{\perp}/t results in the motion of the particle in an effective square lattice with out-projected sites [see Fig. 2(a) for $t_{\perp}/t = 50$]. The situation is radically different for a finite

$l_0 = 0.15a$, corresponding to a lattice depth of $s = 20$. For $\theta = \theta(2, 1)$, due to the particularly simple elementary cell, particles move amongst all the nondirectly connected sites of both layers [see Fig. 2(b)], and the expansion dynamics is independent of the chosen initial not directly connected site. For smaller moiré angles, the dynamics for $t_{\perp}/t \gg 1$ strongly depends on the initial site, because the central bands break into separate subbands characterized by very different transport properties. Whereas some sites form quasi-isolated islands, other sites connect efficiently to a net of sites building transport channels [see Fig. 2(c)]. Hence, for finite times, of practical relevance in typical experiments, the formation of channels dominates particle transport in the moiré lattice. As a result, a particle initially distributed amongst various sites generally undergoes a bimodal expansion dynamics, characterized by partial channel-like expansion and partial quasi-localization in poorly connected sites [see Fig. 2(d)].

IV. DYNAMICS IN INCOMMENSURATE BILAYERS

As shown above, for general $\theta = \theta(m, n)$, the particle dynamics is characterized by the splitting of the spectrum into separate bands and the corresponding formation for large t_{\perp}/t of quasi-isolated regions and effective lattice channels that dominate the (partial) particle expansion. This picture is distorted when the tilting angle departs from commensurability. We introduce at this point the departure angle ϕ , such that $\theta = \theta(m, n) + \phi$. This incommensurability, together with the finite Gaussian range l_0 , results in a spatial quasi-disorder of the interlayer coupling, which may severely affect the particle dynamics, resulting in ergodicity breaking and eventually localization. In this section, we analyze in detail the role played by the interlayer coupling, whereas Sec. V is devoted to the effect of the bias Δ .

A. Eigenstates

We first focus on the localization properties of the lattice eigenstates, $|\psi\rangle = \sum_{\alpha,j} c_{\alpha,j}(\psi) |\alpha, j\rangle$, which are well characterized by the moments

$$I_q(\psi) = \sum_{\alpha,j} |c_{\alpha,j}(\psi)|^{2q} \propto N^{-D_q(\psi)(q-1)}, \quad (4)$$

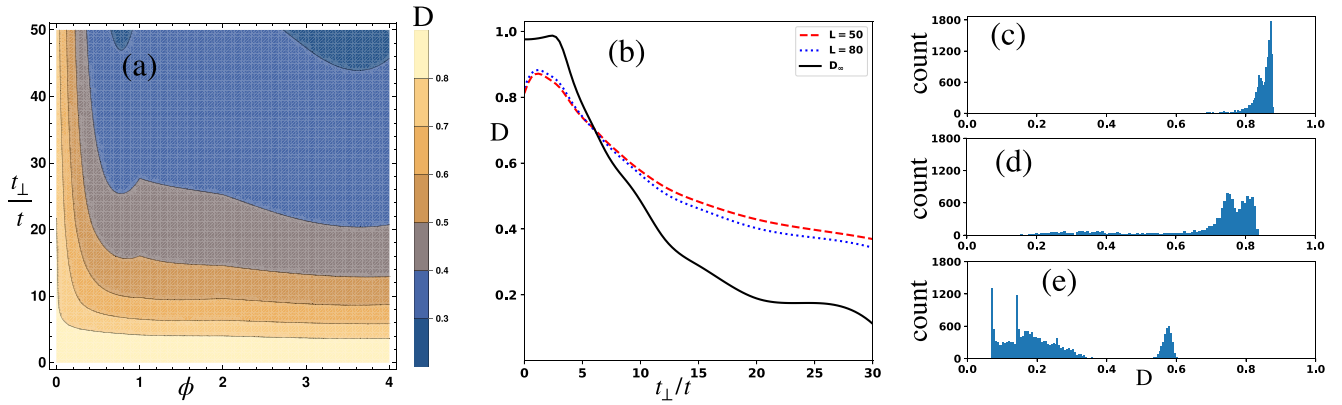


FIG. 3. Eigenstate properties for incommensurate twisting angles. (a) Averaged fractal dimension $D(N)$ for $N = 80 \times 80$ sites, as a function of t_{\perp}/t and ϕ . (b) $D(N)$ for $\phi = 3^{\circ}$, and a lattice with $N = 50 \times 50$ (red dashed) and 80×80 (blue dotted) sites. The black solid curve depicts D_{∞} (see text). (c–e) Histogram of the distribution of $D_2(\psi)$ values for $t_{\perp}/t = 3$ (c), 12 (d) and 100 (e). In all the figures we consider $\theta = \theta(2, 1) + \phi$, $l_0/a = 0.15$, and $\Delta = 0$.

where N is the total number of sites in each layer, and $D_q(\psi)$ are the fractal dimensions. In particular, the inverse participation ratio (IPR) [29–31] $I_2(\psi)$ is given by the inverse of the number of sites in which the eigenstate $|\psi\rangle$ has significant support. Localized (ergodic) states are characterized by $D_q(\psi) = 0$ ($D_q(\psi) = 1$), whereas intermediate q -dependent $0 < D_q(\psi) < 1$ implies an extended but nonergodic character, and a multifractal geometry [32–40].

In the following, we focus our analysis on the IPR, determining for a lattice with N sites, up to 80×80 , and open boundary conditions, the fractal dimension of each eigenstate $D_2(N; \psi) = \log I_2(\psi) / \log(N)$. In order to provide a global characterization of the system, we evaluate the averaged value of the fractal dimension over all eigenstates $D(N) = \frac{1}{N} \sum_{\psi} D_2(N; \psi)$.

Figure 3(a) shows $D(N)$ for $N = 80 \times 80$ sites, as a function of the interlayer hopping t_{\perp}/t and the departure angle ϕ (note that we limit to $0 < \phi < 4^{\circ}$ in order to avoid nearby commensurate tilting angles; the results for $\phi < 0$ are very similar). For $t_{\perp} = 0$, each layer constitutes an independent disorder-free square lattice, characterized by ergodic eigenstates, and correspondingly band expansion. The eigenstates are also strictly speaking ergodic for a moiré lattice ($\phi = 0$) irrespective of the value of t_{\perp}/t . However, as mentioned above, the corresponding bands may be significantly flat, leading to potentially very long timescales for the band-expansion dynamics. Note as well that the formation of separated bands characteristic of large-enough t_{\perp}/t values reduces the lattice support of the eigenstates, resulting for large t_{\perp}/t in a finite-size-induced deviation from the expected value $D = 1$. We discuss this finite-size effect below.

For a nonzero tilting deviation ϕ , a finite t_{\perp} results in an effective two-dimensional quasi-disordered spatial dependence of the interlayer hopping amplitude. Note that this is so, crucially, because $l_0 > 0$. A vanishing l_0/a would result for a finite ϕ in the almost complete decoupling of the layers, and hence in ergodic, basically monolayer, dynamics. As a result, t_{\perp}/t would not play the role of quasi-disordered strength discussed below. Note as well that the quasi-disorder is, in principle, established even for very small angle deviations. However, a lattice with a very small $\phi < 1^{\circ}$ is barely

distinguishable from a moiré lattice for the system sizes considered in our numerics (and for typical experimental sizes), resulting in the enhancement of the value of D observed in Fig. 3(a). For large $\phi > 1^{\circ}$ values, the results are approximately ϕ independent. As seen in Fig. 3(a), the ergodic character of the whole eigenspectrum is maintained at low-enough interlayer couplings, with $D(N) > 0.85$ for $t_{\perp}/t < 4$. Beyond that value, D decays markedly, reaching values $D(N) \lesssim 0.3$ already for $t_{\perp}/t \simeq 25$.

Finite-size effects pose a major difficulty when studying the localization properties and, in particular, the fractal dimension. These effects may be to a large extent mitigated using the following argument. Note that, for a given eigenstate $|\psi\rangle$ in a system with N sites, $I_2(\psi) = \gamma(\psi)/N^{D_2(\psi)}$. Hence the evaluated fractal dimension $D_2(N; \psi) = D_2(\psi) + \frac{\log \gamma(\psi)}{\log N}$. Assuming that the deviation averaged over all eigenstates $\frac{1}{N} \sum_{\psi} \log \gamma(\psi)$ is approximately N independent, we may then employ the following ansatz for the relation between the averaged fractal dimension $D(N)$ for the case of N sites and the value D_{∞} expected for an infinitely large system:

$$D(N) = D_{\infty} + \alpha / \log(N), \quad (5)$$

where D_{∞} and α are determined by fitting our results for different system sizes, up to 80×80 sites. Figure 3(b) shows our results for D_{∞} for $t_{\perp}/t = 10$, $l_0/a = 0.15$, and a deviation $\phi = 3^{\circ}$ from the moiré angle $\theta(2, 1)$.

The extrapolated results confirm the qualitative picture observed in Fig. 3(a). The spectrum shows a clear change of character at $t_{\perp}/t \simeq 3$. For weaker interlayer coupling $D_{\infty} \simeq 1$, and hence the whole spectrum is ergodic. Figure 3(c) shows the distribution of $D_2(N; \psi)$ for $N = 80 \times 80$ sites, with the expected peak at large D values (only limited by finite-size effects). At $t_{\perp}/t \simeq 3$, whereas part of the spectrum remains ergodic, the rest undergoes an ergodic-to-nonergodic transition, resulting in intermediate D_{∞} values. These intermediate values are not a finite-size effect. Note in this sense, that for $t_{\perp}/t \simeq 6$, the $D(N)$ curves with different N cross, indicating that around that value the spectrum is N independent ($\alpha \simeq 0$ in the expression above). Note as well that the nonergodic eigenstates do not have a localized nature, but rather an

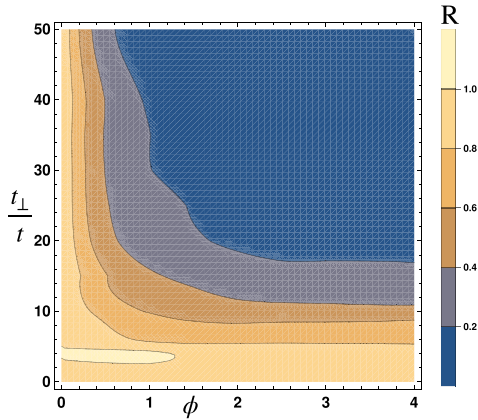


FIG. 4. Expansion dynamics for incommensurate twist angles. Expansion radius $R(\tau = 16/t)$ as a function of ϕ and t_{\perp}/t for the same case as in Fig. 3(a).

extended nature. This is evident from Fig. 3(d), where we depict the distribution of $D_2(N; \psi)$ for $t_{\perp}/t = 12$. This distribution shows, in addition to basically ergodic states, a large number of states well within the intermediate regime of D_2 values. Further increasing t_{\perp}/t [see Fig. 3(e) for $t_{\perp}/t = 100$] results in the displacement of the bulk of the spectrum to low D_2 , indicating localization (although part of the states remain with a clear nonergodic extended character even for such a strong interlayer coupling). While the behavior of D around the twist angle $\theta(2, 1)$ is presented in Fig. 3(a), we have verified that the physics is very similar for other twist angles.

B. Expansion dynamics

The change in the nature of the eigenstates when increasing t_{\perp}/t and ϕ translates into a marked modification of the expansion dynamics of an initially localized wave packet (at $\tau = 0$). We characterize the particle expansion at a given time $\tau > 0$ by means of the average distance \bar{r} , from the initial central position \bar{R}_{in} of the particle wave packet:

$$\bar{r}(\tau)^2 = \sum_{\alpha=1,2} \sum_j |\bar{R}_{\alpha,j} - \bar{R}_{\text{in}}|^2 |c_{\alpha,j}(\tau)|^2. \quad (6)$$

In order to assess the effect of twisting incommensurability we compare this radius with the one, $\bar{r}_0(\tau)$, expected for $\phi = 0$, defining $R(\tau) = \bar{r}(\tau)/\bar{r}_0(\tau)$. This normalization is necessary, since, as mentioned above, the dynamics in a moiré lattice may slow down very significantly with t_{\perp}/t due to the appearance of quasi-flat bands. Figure 4 shows, for different values of ϕ and t_{\perp}/t , the normalized radius $R(\tau = 16/t)$, for an initial wave packet centered at $(0,0)$ with width $\sigma/a = 1$. The qualitative behavior of R mirrors that of the fractal dimension D in Fig. 3(a). For $t_{\perp}/t < 4$ the expansion dynamics is basically identical to that of the commensurate case. In the vicinity of $\phi = 0$, we observe again that, due to finite-size (and also finite-time) limitations, a small ϕ is almost indistinguishable from a moiré bilayer. In contrast, the results are only weakly dependent on ϕ for $\phi > 1^\circ$. In that regime, $R(\tau)$ decreases very markedly with growing t_{\perp}/t , indicating the onset of nonergodic dynamics and eventually localization.

V. INTERLAYER BIAS

The previous section has shown that the interlayer coupling acts as an effective quasi-disorder strength that controls the degree of ergodicity breaking in the system. As discussed in this section, a similar role may be played as well by the bias Δ , i.e., by the energy offset between the two layers in Eq. (1). Figure 5(a) depicts the averaged fractal dimension D as a function of ϕ and Δ/t_{\perp} , for an 80×80 lattice, with $\theta = \theta(2, 1) + \phi$, $l_0/a = 0.15$, and $t_{\perp}/t = 10$.

As in the previous section, the region of small ϕ presents an anomalously large D due to the finite-size quasi-indistinguishability from the moiré bilayer case. Increasing the bias Δ renders the interlayer hopping off-resonant, eventually resulting for a large-enough Δ/t_{\perp} in an effective decoupling of the layers. Since a decoupled layer is a regular square lattice, a large-enough bias leads to the retrieval of ergodicity irrespective of ϕ . The extrapolated value D_{∞} [see Fig. 5(b)] shows that the whole spectrum remains ergodic for $\Delta/t_{\perp} > 1.5$. Below that value, part of the spectrum becomes multifractal. Note that also in this curve we may identify a point, at $\Delta/t_{\perp} \simeq 0.5$, at which the curves $D(N)$ for different N cross.

As in the previous section, the time dependence of the normalized averaged distance R mirrors the spectral properties. Similar to the previous section, we normalize R to the value expected for $\phi = 0$. Figure 5(c) depicts $R(\tau = 20/t)$ as a function of ϕ and Δ/t , showing a good qualitative agreement with Fig. 5(a). The average distance is almost independent of ϕ for large bias, implying an extended phase in the effective single-layer regime. A smaller bias leads to a markedly nonergodic dynamics. Interestingly, the largest deviation from ergodicity occurs for $\phi > 1^\circ$ not at $\Delta = 0$, but rather at a finite $\Delta/t \simeq 2$.

VI. CONCLUSIONS

Particle dynamics in a twisted-bilayer optical lattice presents a nontrivial dependence on the lattice depth (which controls the hopping t and the width l_0 of the interlayer Gaussian coupling), the interlayer coupling strength t_{\perp}/t , the interlayer bias Δ/t , and the tilting angle θ . Crucially, in contrast to solid-state twisted-bilayer setups, the interlayer coupling can be controlled basically at will and may be much larger than the intralayer one. As a result, the interlayer coupling may play a crucial role in the particle dynamics. For commensurate moiré-like lattices, the eigenstates are in any case ergodic, but due to the flatness of the moiré bands, for relevant experimental timescales transport for $t_{\perp}/t \gg 1$ is dominated by channel formation. For incommensurate lattices, and due to the finite range of the interlayer coupling, the coupling rate t_{\perp} acts as an effective quasi-disorder strength. Whereas the spectrum remains fully ergodic for small $t_{\perp}/t \lesssim 3$ (for $\Delta = 0$), a larger interlayer coupling induces the transition of part of the spectrum into nonergodic (but still extended) states. Similarly, ergodicity is recovered for a sufficiently large interlayer bias, whereas reducing the bias induces again a partial ergodic-to-nonergodic transition. The spectral properties may be revealed from the analysis of the expansion dynamics of an initially localized wave packet. Our results hence show that twisted-bilayer optical lattice

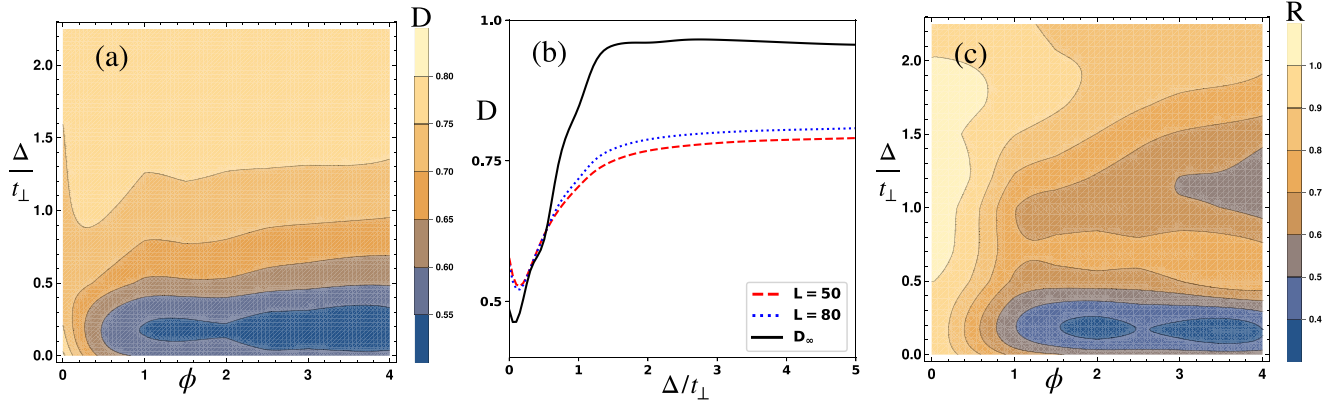


FIG. 5. Effect of the interlayer bias. (a) Averaged fractal dimension $D(N)$ for $N = 80 \times 80$ sites as a function of Δ/t_{\perp} and ϕ . (b) $D(N)$ for $\phi = 3^{\circ}$, and a lattice with $N = 50 \times 50$ (red dashed) and 80×80 (blue dotted) sites. The black solid curve depicts D_{∞} (see text). (c) Expansion radius $R(\tau = 20/t)$ as a function of ϕ and t_{\perp}/t for the same case as in panel (a). In all the figures, $t_{\perp}/t = 10$, $l_0 = 0.15a$, and $\theta(2, 1) + \phi$.

setups provide an interesting controllable platform for the study of multifractality. Furthermore, multilayer setups may allow for the study of particle transport in the two-dimensional-to-three-dimensional cross-dimensional regime, an intriguing possibility for future studies.

ACKNOWLEDGMENT

We acknowledge support of the Deutsche Forschungsgemeinschaft (DFG, German Research Foundation) under Germany's Excellence Strategy – EXC-2123 Quantum-Frontiers – 390837967.

- [1] R. Bistritzer and A. H. MacDonald, *Proc. Natl. Acad. Sci. USA* **108**, 12233 (2011).
- [2] J. M. B. Lopes Dos Santos, N. M. R. Peres, and A. H. Castro Neto, *Phys. Rev. Lett.* **99**, 256802 (2007).
- [3] E. Y. Andrei and A. H. MacDonald, *Nat. Mater.* **19**, 1265 (2020).
- [4] L. Balents, C. R. Dean, D. K. Efetov, and A. F. Young, *Nat. Phys.* **16**, 725 (2020).
- [5] P. Törmä, S. Peotta, and B. A. Bernevig, *Nat. Rev. Phys.* **4**, 528 (2022).
- [6] Y. Cao, V. Fatemi, S. Fang, K. Watanabe, T. Taniguchi, E. Kaxiras, and P. Jarillo-Herrero, *Nature (London)* **556**, 43 (2018).
- [7] M. Yankowitz, S. Chen, H. Polshyn, Y. Zhang, K. Watanabe, T. Taniguchi, D. Graf, A. F. Young, and C. R. Dean, *Science* **363**, 1059 (2019).
- [8] M. Oh, K. P. Nuckolls, D. Wong, R. L. Lee, X. Liu, K. Watanabe, T. Taniguchi, and A. Yazdani, *Nature (London)* **600**, 240 (2021).
- [9] Y. Cao, V. Fatemi, A. Demir, S. Fang, S. L. Tomarken, J. Y. Luo, J. D. Sanchez-Yamagishi, K. Watanabe, T. Taniguchi, E. Kaxiras *et al.*, *Nature (London)* **556**, 80 (2018).
- [10] E. Codecido, Q. Wang, R. Koester, S. Che, H. Tian, R. Lv, S. Tran, K. Watanabe, T. Taniguchi, F. Zhang *et al.*, *Sci. Adv.* **5**, eaaw9770 (2019).
- [11] K. P. Nuckolls, M. Oh, D. Wong, B. Lian, K. Watanabe, T. Taniguchi, B. A. Bernevig, and A. Yazdani, *Nature (London)* **588**, 610 (2020).
- [12] Y. Cao, D. Rodan-Legrain, O. Rubies-Bigorda, J. M. Park, K. Watanabe, T. Taniguchi, and P. Jarillo-Herrero, *Nature (London)* **583**, 215 (2020).
- [13] G. Trambly de Laissardière, D. Mayou, and L. Magaud, *Nano Lett.* **10**, 804 (2010).
- [14] S. Lisi, X. Lu, T. Benschop, T. A. de Jong, P. Stepanov, J. R. Duran, F. Margot, I. Cucchi, E. Cappelli, A. Hunter *et al.*, *Nat. Phys.* **17**, 189 (2021).
- [15] E. S. Morell, J. D. Correa, P. Vargas, M. Pacheco, and Z. Barticevic, *Phys. Rev. B* **82**, 121407(R) (2010).
- [16] G. Tarnopolsky, A. J. Kruchkov, and A. Vishwanath, *Phys. Rev. Lett.* **122**, 106405 (2019).
- [17] A. González-Tudela and J. I. Cirac, *Phys. Rev. A* **100**, 053604 (2019).
- [18] T. Salamon, A. Celi, R. W. Chhajlany, I. Frérot, M. Lewenstein, L. Tarruell, and D. Rakshit, *Phys. Rev. Lett.* **125**, 030504 (2020).
- [19] X.-W. Luo and C. Zhang, *Phys. Rev. Lett.* **126**, 103201 (2021).
- [20] J. Lee and J. Pixley, *SciPost Phys.* **13**, 033 (2022).
- [21] Z. Meng, L. Wang, W. Han, F. Liu, K. Wen, C. Gao, P. Wang, C. Chin, and J. Zhang, *Nature (London)* **615**, 231 (2023).
- [22] S. Aubry and G. André, *Ann. Isr. Phys. Soc.* **3**, 133 (1980).
- [23] G. Roati, C. D'Errico, L. Fallani, M. Fattori, C. Fort, M. Zaccanti, G. Modugno, M. Modugno, and M. Inguscio, *Nature (London)* **453**, 895 (2008).
- [24] M. Schreiber, S. S. Hodgman, P. Bordia, H. P. Lüschen, M. H. Fischer, R. Vosk, E. Altman, U. Schneider, and I. Bloch, *Science* **349**, 842 (2015).
- [25] K. Viebahn, M. Sbroscia, E. Carter, J.-C. Yu, and U. Schneider, *Phys. Rev. Lett.* **122**, 110404 (2019).
- [26] Y. Lahini, R. Pugatch, F. Pozzi, M. Sorel, R. Morandotti, N. Davidson, and Y. Silberberg, *Phys. Rev. Lett.* **103**, 013901 (2009).
- [27] P. Wang, Y. Zheng, X. Chen, C. Huang, Y. V. Kartashov, L. Torner, V. V. Konotop, and F. Ye, *Nature (London)* **577**, 42 (2020).
- [28] C. Huang, F. Ye, X. Chen, Y. V. Kartashov, V. V. Konotop, and L. Torner, *Sci. Rep.* **6**, 1 (2016).

- [29] F. Wegner, *Z. Physik B* **36**, 209 (1980).
- [30] F. Evers and A. D. Mirlin, *Phys. Rev. Lett.* **84**, 3690 (2000).
- [31] N. C. Murphy, R. Wortis, and W. A. Atkinson, *Phys. Rev. B* **83**, 184206 (2011).
- [32] A. Jagannathan, *Rev. Mod. Phys.* **93**, 045001 (2021).
- [33] W. Pook and M. Janßen, *Z. Phys. B* **82**, 295 (1991).
- [34] C. d. C. Chamon, C. Mudry, and X.-G. Wen, *Phys. Rev. Lett.* **77**, 4194 (1996).
- [35] X. Deng, B. L. Altshuler, G. V. Shlyapnikov, and L. Santos, *Phys. Rev. Lett.* **117**, 020401 (2016).
- [36] N. Macé, F. Alet, and N. Laflorencie, *Phys. Rev. Lett.* **123**, 180601 (2019).
- [37] X. Deng, S. Ray, S. Sinha, G. V. Shlyapnikov, and L. Santos, *Phys. Rev. Lett.* **123**, 025301 (2019).
- [38] M. Sarkar, R. Ghosh, A. Sen, and K. Sengupta, *Phys. Rev. B* **105**, 024301 (2022).
- [39] M. Reisner, Y. Tahmi, F. Piéchon, U. Kuhl, and F. Mortessagne, *Phys. Rev. B* **108**, 064210 (2023).
- [40] A. De Luca, B. L. Altshuler, V. E. Kravtsov, and A. Scardicchio, *Phys. Rev. Lett.* **113**, 046806 (2014).