

Flat-Band Localization in Creutz Superradiance Lattices

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(Received 18 August 2020; accepted 2 February 2021; published 8 March 2021)

Flat bands play an important role in diffraction-free photonics and attract fundamental interest in many-body physics. Here we report the engineering of flat-band localization of collective excited states of atoms in Creutz superradiance lattices with tunable synthetic gauge fields. Magnitudes and phases of the lattice hopping coefficients can be independently tuned to control the state components of the flat band and the Aharonov-Bohm phases. We can selectively excite the flat band and control the flat-band localization with the synthetic gauge field. Our study provides a room-temperature platform for flat bands of atoms and holds promising applications in exploring correlated topological materials.

DOI: [10.1103/PhysRevLett.126.103601](https://doi.org/10.1103/PhysRevLett.126.103601)

Flat bands are characterized by the zero bandwidth over the whole Brillouin zone. Owing to the destructive interference between the hopping pathways [1,2], the group velocity of excitations vanishes, and hence the diffusion in flat bands is inhibited. The resulting compact localized eigenstates (CLSs) [3,4] have been experimentally realized in photonics [5–13] and polariton-exciton condensates [14,15]. Immune to environmental noises, localized states in flat bands are promising candidates for realizing quantum networks [16] and diffraction-free photonics [17–19]. Flat bands are also of fundamental interest in many-body physics because of their high degeneracy. The density of states is divergent such that even weak interactions lead to strong correlations and exotic topological phases [20–23].

Many-body interactions can be engineered to realize correlated topological phases in atoms [24,25]. However, in previous realizations of the flat bands in optical lattices, the underlying lattices [26,27] are gapless and topologically trivial. A feasible model that integrates both band flatness and topology is the two-leg ladder in a uniform magnetic field with cross-linked couplings, i.e., the Creutz lattice [28,29] [see Fig. 1(a)]. Despite theoretical proposals in photonic waveguides [9] and ultracold atoms [21], flat bands in the Creutz lattice have never been experimentally realized [30–32].

Here we report the synthesis of a Creutz ladder with tunable tight-binding parameters in a momentum-space lattice, i.e., the superradiance lattice [33,34], in room-temperature cesium atoms. The bipartite ladder consists of timed Dicke states with different momenta [35]. We find

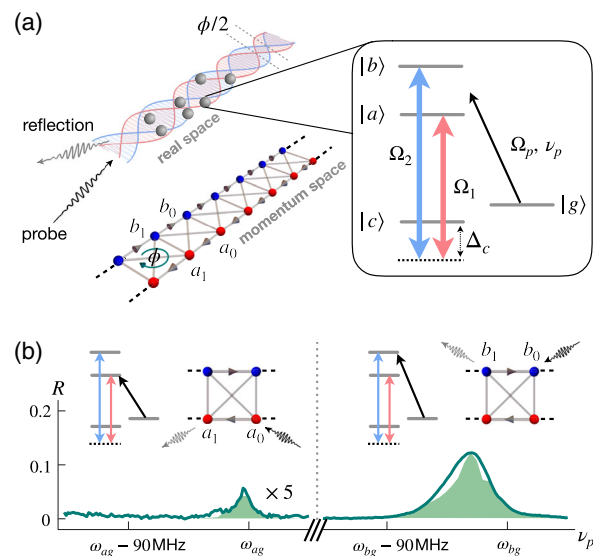


FIG. 1. (a) Schematic configuration of the experiment. The amplitude envelopes of the two standing waves have a $\phi/2$ phase difference. In the Creutz ladder, the arrows indicate the phase $\phi/2$ attached to the transitions. Inset is the configuration of atomic levels and laser fields. (b) Typical reflection spectrum. The insets show the configurations of the fields and the lattice responses when ν_p is near resonant with either atomic transition. The AB flux $\phi = \pi$. The power of the probe field is 24 μW . The powers of each plane wave component of the two standing waves are $P_1 = 29$ mW and $P_2 = 215$ mW with effective Rabi frequencies $\Omega_1 = 15$ MHz and $\Omega_2 = 68$ MHz. The thick dark lines (light filling areas) are the experimental data (numerical simulation).

that the corresponding energy band structure exhibits a flat band and a dispersive band, which are distinguished by localized and delocalized excitations, respectively. In the experiment, we excite one site in the ladder with a weak probe field and measure the optical response of the adjacent site. The hopping strengths and the Aharonov-Bohm (AB) phases in the lattice are carefully tuned, which enables us to excite a particular band and control the flat-band localization. We observe that the optical response is significantly suppressed when the flat band is selectively excited. By controlling the AB phases, we reveal the relation between the flat-band localization and gauge fields [36]. Our work demonstrates a versatile platform for flat bands of atoms with multiple tunable parameters, which holds promising applications in exploring correlated topological phases.

We first introduce the experimental scheme implemented in the hyperfine levels of the ^{133}Cs $D1$ line in a bichromatic standing-wave-coupled configuration in Fig. 1(a). Two continuous standing-wave fields couple two excited states $|a\rangle \equiv |6^2P_{1/2}, F=3\rangle$ and $|b\rangle \equiv |6^2P_{1/2}, F=4\rangle$ to the same metastable state $|c\rangle \equiv |6^2S_{1/2}, F=3\rangle$. The frequencies of the two standing-wave coupling fields ν_1 and ν_2 fulfill the two-photon resonance condition $\Delta_c = \nu_1 - \omega_{ac} = \nu_2 - \omega_{bc}$, where ω_{ij} being the atomic transition frequency between $|i\rangle$ and $|j\rangle$. The envelopes of the Rabi frequency amplitude of the two standing waves are $2\Omega_1 \cos(k_c x - \phi/4)$ and $2\Omega_2 \cos(k_c x + \phi/4)$, where k_c is the x component of the wave vectors and $\phi/2$ is the phase difference. The standing-wave-coupled atoms are probed with a weak continuous-wave field with the wave vector k_p . The reflection of the probe field reveals the steady-state response of the atoms. The frequency of the probe field ν_p is scanned to couple the ground state $|g\rangle \equiv |6^2S_{1/2}, F=4\rangle$ to either $|a\rangle$ or $|b\rangle$. Featured signals can be observed when the probe field is near resonant with each atomic transition. A typical spectrum is shown in Fig. 1(b).

In order to show that our experiment constructs a Creutz ladder and reveal the connection between the reflection signal and the excitation transport in the ladder, we write the Hamiltonian $H = H_s + H_p$ in momentum space [37], with H_s and H_p being the parts of the Hamiltonian involving the standing-wave coupling fields and the probe field. Here, we set $\hbar = 1$ and H_s reads

$$H_s = \sum_n [2t_1 a_n^\dagger a_n + 2t_2 b_n^\dagger b_n + (2t_3 \cos \frac{\phi}{2} a_n^\dagger b_n + t_3 a_n^\dagger b_{n+1} + t_3 b_n^\dagger a_{n+1} + t_1 e^{-i\phi/2} a_n^\dagger a_{n+1} + t_2 e^{i\phi/2} b_n^\dagger b_{n+1} + \text{H.c.})], \quad (1)$$

which gives a tight-binding superradiance lattice [33] and $d_j^\dagger = \sqrt{1/N} \sum_m |d_m\rangle \langle g_m| \exp[i(k_p - 2jk_c)x_m]$ ($d = a, b$), where m labels the m th atom at the position x_m , j is an

integer, and N is the total number of atoms. Applying d_j^\dagger to the ground state of the atoms $|G\rangle = |g_1 g_2 \dots g_N\rangle$, we obtain the single-photon timed Dicke states [35,40–42]. $t_{1(2)} = -\Omega_{1(2)}^2/\Delta_c$ are the hopping amplitudes along each leg. Here, we can adiabatically eliminate the state $|c\rangle$, since Δ_c is much larger than all relevant Rabi frequencies ($\Delta_c \gg \Omega_j$). The two hoppings acquire a phase $\phi/2$ in opposite directions. The loop transition along a plaquette accumulates an AB phase ϕ , such that the lattice is effectively in a uniform magnetic field. $t_3 = -\Omega_1 \Omega_2 / \Delta_c$ and $2t_3 \cos \phi/2$ are the hopping strengths along the diagonals and the rungs of each plaquette. The on-site energies of $a_j(b_j)$ are $2t_1(2t_2)$, respectively. The relationship between the current ladder model and previously studied one-dimensional lattices based on Λ -type electromagnetically induced transparency configuration [33,34,43] is discussed in the Supplemental Material [37].

The interaction Hamiltonian due to the probe field is $H_p = \sqrt{N}\Omega_p e^{-i\Delta_p t} a_0^\dagger + \sqrt{N}\Omega_p' e^{-i\Delta_p' t} b_0^\dagger + \text{H.c.}$, where Ω_p (Ω_p') and $\Delta_p = \nu_p - \omega_{bg}$ ($\Delta_p' = \nu_p - \omega_{ag}$) are the Rabi frequency and the frequency detuning between the probe field and the atomic transition between $|a\rangle$ ($|b\rangle$) and $|g\rangle$. Hence, H_p shows that the excitation is prepared by the probe field to the site a_0 or b_0 in the ladder. When we probe the site a_0 (b_0), the phase-matching condition $k_p - 2k_c \approx -k_p$ is only satisfied for the excitation on the site a_1 (b_1), which results in a superradiant backward emission, which is collected by a photodetector. In the weak probe field limit, the coherent and collective effects of the atoms are manifested by the phase-matched unidirectional radiation of the single-photon superradiant state (a_1 or b_1) [35,40–42]. Substantially different from the superfluorescence in population inverted systems [44,45], where atomic coherence is induced by the mutual stimulation, the coherence between atoms of the single-photon superradiant state is established by the probe field and coherently controlled by the standing-wave coupling fields. Meanwhile, we can also distinguish our signal from population-inverted superfluorescence by noticing that the observed line widths (see Fig. 2) are determined by the energy bands of the superradiance lattice [43]. The spectrum in the left (right) of Fig. 1(b) characterizes the excitation transport from a_0 to a_1 (b_0 to b_1) in the ladder of Eq. (1). In the experiment, the probe field is weak ($\Omega_p \ll t_i$) such that only a small fraction of the atoms are excited. In this condition, a_j, b_j are approximately bosonic annihilation operators [33].

We diagonalize H_s in real space [37] and the band structures are shown in Fig. 2. We define $\eta \equiv t_2/t_1$ as the relative hopping strength along the two legs. All three band structures in Fig. 2(a)–2(c) with different η are composed of a flat and a dispersive band. In general, the dynamics of the excitation is governed by both bands and cannot be distinguished. Probing only one band by controlling the

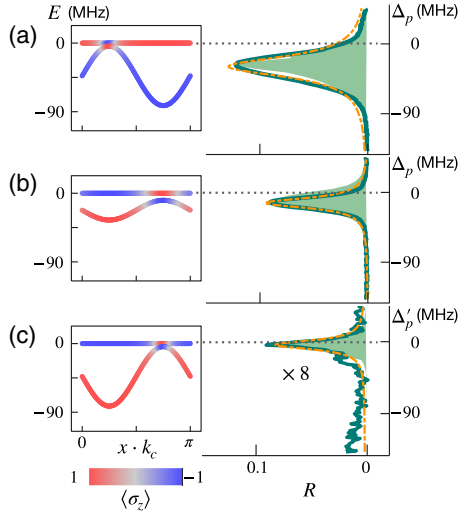


FIG. 2. Band structures (left) and the corresponding reflection spectra (right) with (a) $\eta = 20.7$, (b) $\eta = 3.8$, and (c) $\eta = 1/20.7$. The reflection spectrum is mainly contributed by the dispersive (flat) band for $\eta \gg 1$ ($\eta \ll 1$). The data in (c) is obtained by measuring the a -leg response owing to the symmetry of the Hamiltonian (see the text). $\sqrt{P_1 P_2} \approx 78$ mW is kept as a constant and other parameters are the same as in Fig. 1(b). Dotted gray lines mark the energy of the flat band. The thick dark lines (light filling areas) are the experimental data (numerical simulation). The spectra are fitted with Lorentzian functions (orange dashed lines) with FWHM = 32.4, 15.7, and 13.7 MHz and peaks at -29.2 , -13.6 , and -4.0 MHz from top to bottom.

excitation energy [46–48] is inapplicable since the band gap closes when ϕ approaches zero [37].

Nevertheless, the tunability of the hopping strengths enables us to determine which band to excite by controlling the state component of the bands. In the experiment, we tune η , which is proportional to P_2/P_1 , where P_1 and P_2 are the powers of each plane wave component of the two standing waves. An interesting correlation can be noticed between the parameter η and the band components, where the color represents the polarization $\langle \sigma_z \rangle \equiv \langle |a\rangle\langle a| - |b\rangle\langle b| \rangle$ of the eigenstates. In particular, $\langle \sigma_z \rangle = +1$ or -1 means the band fully locates on the a or b leg. In Fig. 2(a), we see that $\langle \sigma_z \rangle \approx -1$ for almost the whole dispersive band, meaning that the dispersive band supports a large excitation component on the b leg for $\eta \gg 1$ (we take $\eta = 20.7$ according to the experiments). Therefore, the $b_0 \rightarrow b_1$ transport is governed by the dispersive band. On the other hand, for $\eta \ll 1$ [$\eta = 1/20.7$ in Fig. 2(c)], $\langle \sigma_z \rangle \approx -1$ for almost the entire flat band, so the $b_0 \rightarrow b_1$ transport dynamics is governed by the flat band.

This band selection is manifested in the bandwidths, the central frequencies, and the magnitudes of the reflection spectra in Fig. 2. In the experiment, we change η and keep $t_3 \propto \sqrt{P_1 P_2}$ a constant. In Fig. 2(a) for $\eta \gg 1$, the reflection spectrum of the dispersive band has a larger bandwidth and a lower central frequency. As a comparison,

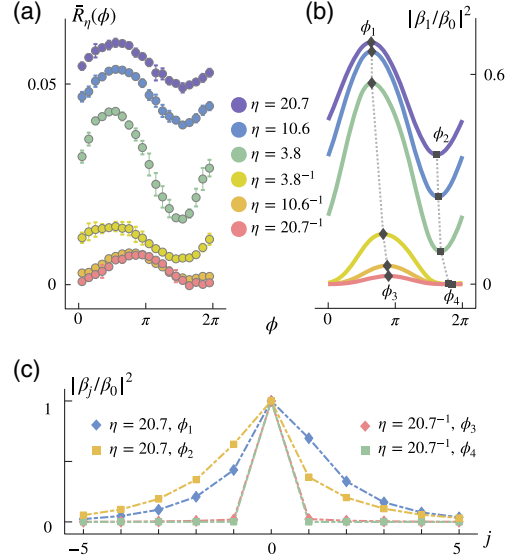


FIG. 3. Response of the dispersive and flat bands with different gauge fields. (a) The averaged reflectivity \bar{R} versus ϕ with different η . (b) The normalized probability on the site b_1 versus ϕ . The diamonds (squares) indicate where the curves reach their maxima (minima). $\Delta_p(\Delta'_p) = -2$ MHz and $\Delta_c = 233.5$ MHz. The gray dashed lines are plotted to guide the extrema of the curves. (c) The probability distribution of the steady state $|\psi_s\rangle$ on the b leg with $\phi_1 = 0.64\pi$, $\eta = 20.7$ (blue diamonds), $\phi_2 = 1.62\pi$, $\eta = 20.7$ (yellow squares), $\phi_3 = 0.9\pi$, $\eta = 1/20.7$ (red diamonds), and $\phi_4 = 1.86\pi$, $\eta = 1/20.7$ (green squares). The powers of each plane wave component of the two standing waves are $P_1 = 67$ mW, $P_2 = 90$ mW for $\eta = 3.8$ and $1/3.8$; $P_1 = 40$ mW, $P_2 = 153$ mW for $\eta = 10.6$ and $1/10.6$; and $P_1 = 29$ mW, $P_2 = 215$ mW for $\eta = 20.7$ and $1/20.7$. Other experimental parameters are the same as in Fig. 1. The points are simply connected for clarity. Error bars are obtained from four independent data sets.

in Fig. 2(c) for $\eta \ll 1$, the reflection spectrum due to the flat band has a much narrower bandwidth and the peak locates near the predicted frequency of the flat band. The localization in the flat band is demonstrated by the decrease of the reflection peak when we decrease η , during which the reflection is more contributed by the flat band. As a side note, in obtaining Fig. 2(c), we use the symmetry that lattice Hamiltonian H_s is invariant when we exchange the sublattices a and b , inverse η , and flip the flux ϕ [37].

The reflection spectrum is mostly contributed by the slowly moving atoms that have Doppler shifts smaller than the lattice bandwidth [43]. We take the average of reflectivity over the bands $\bar{R} = \int R d\nu_p / \int d\nu_p$ to investigate the localization and its ϕ dependence. In Fig. 3(a), the flat-band localization is demonstrated by the suppression of \bar{R} when η decreases. Furthermore, we notice that the ϕ dependence of \bar{R} changes with η . The sinusoidal curve of averaged reflectivity $\bar{R}_\eta(\phi)$ is shifted from top to bottom in Fig. 3(a).

The ϕ -dependent shift shows the distinct responses to the gauge fields of the two bands. If the excitation is completely prepared in the dispersive band, the transport dynamics is determined by the flux-dependent unidirectional chiral edge current [43,49–53] of the dispersive band. The unidirectional chiral current breaks the symmetry between the transitions from b_0 to $b_{1(-1)}$. When the magnetic flux $\phi \in (0, \pi)$, the chiral current enhances the probability in b_1 , and hence the reflectivity increases (vice versa). On the other hand, flat-band response to the gauge field can be understood by the CLSs $|F_j\rangle$ [3]. When $\phi = 2n\pi$ (n is an integer), $|F_j\rangle \propto (\eta a_j^\dagger - b_j^\dagger)|G\rangle$ is localized within the j th unit cell. Therefore, only $|F_0\rangle$ is excited when we probe the site b_0 , leading to the maximum localization. Otherwise, $|F_j\rangle \propto (\eta a_{j+1}^\dagger - e^{i\phi/2} b_{j+1}^\dagger + \eta e^{i\phi/2} a_j^\dagger - b_j^\dagger)|G\rangle$ is localized within two unit cells. Probing site b_0 leads to a coherent superposition between $|F_0\rangle$ and $|F_{-1}\rangle$, and hence results in a finite overlap with $b_{\pm 1}$ [9,10].

The two responses can be further investigated by the steady state of the collective excited states of atoms, where the wave function $|\psi_s\rangle \approx [1 + \sum_j (\alpha_j a_j^\dagger + \beta_j b_j^\dagger)]|G\rangle$ ($\alpha_j, \beta_j \ll 1$) in the weak excitation approximation. In the steady state, we obtain the probability amplitude $\beta_j = \sqrt{N}\Omega_p \langle G|b_j(\Delta_p + i\hat{\gamma} - H_s)^{-1}b_0^\dagger|G\rangle$ [54], where $\hat{\gamma} = \sum_j (\gamma_a a_j^\dagger a_j + \gamma_b b_j^\dagger b_j)$ and γ_a (γ_b) is the decoherence rate between the excited state $|a\rangle$ ($|b\rangle$) and $|g\rangle$. Since the reflectivity is approximately proportional to the $|\beta_1/\beta_0|^2$ [55], we plot $|\beta_1/\beta_0|^2$ in Fig. 3(b) to demonstrate both features of the \bar{R} in Fig. 3(a), including the magnitude suppression and the ϕ dependence. We also plot four typical probability amplitude distributions along the b leg in Fig. 3(c). For $\eta = 20.7$, the distributions of $|\beta_j|^2$ with $\phi_1 = 0.64\pi$ and $\phi_2 = 1.62\pi$ are almost symmetric to each other with respect to the 0th site, implying flux-dependent unidirectional chiral edge currents of the dispersive band. However, the distributions with $\phi_3 = 0.9\pi$ and $\phi_4 = 1.86\pi$ are symmetrically localized at b_0 with different localization lengths. $|\beta_1/\beta_0|^2$ for $\eta = 1/20.7$ is minimized at ϕ_4 , which means that the eigenstates are maximally localized when the synthetic gauge field almost vanishes. We notice that ϕ_4 is slightly different from $2n\pi$, which comes from the suppressed but finite contribution from the dispersive band.

In Fig. 4(b), the different ϕ dependences of the dispersive and flat bands are illustrated with Lissajous curves. The spatial phase difference between the envelopes of the two standing waves is slowly tuned to cover all values of ϕ . We obtain the data sets $\{\bar{R}_\eta(\phi), \bar{R}_{1/\eta}(-\phi)\}$ and fit them with ellipses to reconstruct the Lissajous curves. The shape of the ellipse elucidates the phase differences between the two underlying functions. For example, a Lissajous curve composed of two parametric equations with argument u , e.g., $x = \sin(u)$ and $y = \sin(-u + \phi)$, is a line (circle) when $\phi = 0$ ($\pi/2$). In Fig. 4(a), ϕ obtained by fitting the

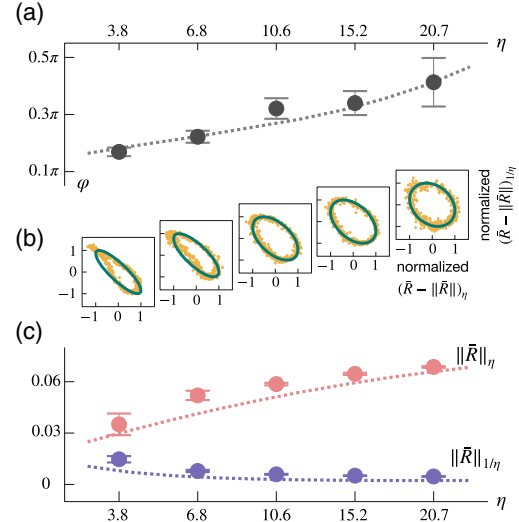


FIG. 4. Gauge field dependence of the reflectivities from the dispersive band and flat band. (a) The phase difference ϕ between the ϕ dependences of \bar{R}_η and $\bar{R}_{1/\eta}$. (b) The Lissajous curves composed of $\bar{R}_\eta(\phi)$ and $\bar{R}_{1/\eta}(-\phi)$. The phase difference between \bar{R}_η and $\bar{R}_{1/\eta}$ increases and approaches $\pi/2$ when η increases. The yellow dots are experimental data and the green lines are fitted elliptic curves. The powers of each plane wave component of the two standing waves are $P_1 = 50$ mW, $P_2 = 124$ mW for $\eta = 6.8$ and $1/6.8$; $P_1 = 33$ mW, $P_2 = 183$ mW for $\eta = 15.2$ and $1/15.2$. Other experimental parameters are the same as in Fig. 3. Each plot contains 400 data points. (c) The mean averaged reflectivity $\|\bar{R}\|_\eta$ versus η . $\|\bar{R}\|_\eta$ increases with η monotonically. The dotted lines are the simulated results.

Lissajous curve approaches $\pi/2$ when η increases, indicating the different types of the excitations on the flat and dispersive bands. In Fig. 4(c), we notice that $\|\bar{R}\|_\eta$ increases with η monotonically, where $\|\cdot\|$ stands for the mean value of the averaged reflectivity \bar{R} over all ϕ . The numerical simulation agrees with the data.

In conclusion, we experimentally realize Creutz ladders with tunable gauge fields, where the flat band can be selectively excited and the interplay between the flat-band localization and the AB phases is investigated. We study the flat-band localization in an open system, where the steady state balanced by pumping, driving, and dissipation exhibits the dynamics in the corresponding closed system [54]. We also need to emphasize that our scheme is substantially different from the incoherently pumped polariton-exciton condensates [14,15,56–58], where coherence is not accessible between multiple CLSs. It is interesting to notice that both bands of the Creutz ladder are topologically nontrivial [10,31,59,60] provided that $\phi \neq 2n\pi$ [37]. It is a step towards the simulation of the strong correlated quantum phases [20,21,61–63]. An interaction term between the sites in momentum space can be introduced by weakly coupling the excited atomic level to a Rydberg state [25] or by s -wave interactions

[39]. With a negative Δ_c , the flat band has the lowest energy and can be used to study the many-body ground states of ultracold atoms [25,37].

We acknowledge the support from the National Natural Science Foundation of China (Grants No. 11934011, No. 11874322, No. 91736209, and No. U1330203), the National Key Research and Development Program of China (Grants No. 2019YFA0308100 and No. 2018YFA0307200), Zhejiang Province Key Research and Development Program (Grant No. 2020C01019) and the Fundamental Research Funds for the Central Universities. Y.L. was supported by National Natural Science Foundation of China (Grants No. 12074431 and No. 11774428) and L. Y. was supported by Natural Science Foundation of Shanghai (19ZR1475700).

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