Percolating states in the topological Anderson insulator

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We investigate the presence of percolating states in disordered two-dimensional topological insulators. In particular, we uncover a close connection between these states and the so-called topological Anderson insulator, which is a topologically nontrivial phase induced by the presence of disorder. The decay of this phase could previously be connected to a delocalization of bulk states with increasing disorder strength. We identify this delocalization to be the result of a percolation transition of states that circumnavigate the hills of the bulk disorder potential.

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I. INTRODUCTION

Topological insulators are recently discovered materials with promising electronic properties [1–3]. The first physical system that was identified as a topological insulator was the HgTe/CdTe quantum well [4] featuring one-dimensional edge states that are protected from backscattering by time-reversal symmetry, strongly stabilizing them against nonmagnetic disorder. The existence of such conducting edge states is hallmarkd by a quantized conductance plateau that has meanwhile been verified experimentally [5]. As discovered in a successive numerical study [6], such edge states are not only immune from backscattering but can even be elicited by disorder in systems that have no topologically distinct properties in the clean limit. This disorder–induced topological phase was first believed to be caused by Anderson localization, and was thus named a “topological Anderson insulator” (TAI). A later study found, however, that the phase boundaries at the transition from an ordinary insulator to the TAI can be explained by an effective medium theory in which the presence of disorder leads to a renormalization of the medium parameters [7]. In this sense, the TAI appears due to a change of topology in the effective medium.

While the transition from an ordinary insulator to the TAI could be explained by the aforementioned effective description [7], the transition from the TAI phase back to an ordinary insulating phase at very strong disorder values proves more involved: The bulk states localize at intermediate disorder strength, allowing for unimpeded edge transport in the TAI phase, yet delocalize when disorder becomes even stronger [7–9]. So far, the resulting breakdown of the TAI phase could be attributed to the coupling of counterpropagating edge states on opposing edges through these delocalized bulk states, resulting in a suppression of the edge states’ immunity from backscattering [10,11]. This mechanism is responsible for an increased sensitivity to finite size effects [12,13], making the transition hard to explore numerically and leaving the true nature of this counterintuitive delocalization unclear. While first studies [9,13,14] interpreted the bulk delocalization as an intermediate metallic phase, a later study [11] considering larger systems pointed out that only a single bulk state is probably responsible for the delocalization and an intermediate metallic phase is not present. In addition, a spatially correlated potential and the associated pronounced bulk delocalization turn out to destroy the TAI phase entirely [15]. In this paper, we resolve the puzzle associated with these different observations by identifying the emergence of percolating states as the origin of the delocalization and by clarifying the general connection between such states and the TAI phase.

II. MODEL AND METHODS

As a starting point, we choose the well-studied disordered HgTe/CdTe quantum well as described by the Bernevig-Hughes-Zhang (BHZ) model [4] in terms of an effective four-band Hamiltonian

$$H_{\text{eff}}(k_x, k_y) = \begin{pmatrix} h(\vec{k}) & 0 \\ 0 & h^*(-\vec{k}) \end{pmatrix},$$

(1)

with

$$h(\vec{k}) = \epsilon(\vec{k}) + d_i(\vec{k})\sigma^i, \quad \epsilon(\vec{k}) = C - D(k_x^2 + k_y^2),$$

$$d_i = (A k_x, -A k_y, M(\vec{k}))^T, \quad M(\vec{k}) = m - B(k_x^2 + k_y^2),$$

(2)

and $\sigma^i$ representing the Pauli matrices. Following Ref. [16], we choose the following set of realistic quantum well parameters in all our computations: $A = 364.5$ nm meV, $B = -686.0$ nm$^2$ meV, and $D = -512.0$ nm$^2$ meV. The topology of the system is determined by the sign of the topological mass $m$: For $m < 0$, the bulk band gap of size $2|m|$ is topologically nontrivial and thus filled with gapless edge states characterizing a two-dimensional topological insulator. On the other hand, for $m > 0$, the bulk band gap is topologically trivial and does not contain any states, leaving us with a system that is an ordinary insulator.

Using the advanced modular recursive Green’s function method [17–19] we calculate the conductance $G$ through two-dimensional rectangular ribbons of HgTe/CdTe quantum wells discretized on a square grid with discretization constants $\Delta x$ and $\Delta y$, width $W = (n_x + 1)\Delta x$, and length $L = n_x \Delta x$. In accordance with previous studies [6,7,9–13,15,20], the discretization constants are set to $\Delta x = \Delta y = a = 5$ nm. Two clean semi-infinite leads are attached to the left and right ends of the ribbon. Following the Landauer-Büttiker formalism, the conductance $G$ in the limit of vanishing temperature is given by the total transmission $T$ at the Fermi energy $E_F$. Our method also allows for a calculation of the scattering wave functions $\psi(x, y)$ as well as the density of states $\rho(E) = -\text{Tr} [\text{Im} G^*(\vec{x}, E)] / \pi$, where $G^*$ is the retarded Green’s function. Disorder is modeled by static on-site energy values.
V(x_i, y_i) at each grid point (x_i, y_i) randomly chosen from the interval [−U/2, U/2], with U the disorder strength. In most studies [6,7,9–13,20], the values of V(x_i, y_i) are chosen without any spatial correlations between neighboring grid points [see Fig. 1 (left panel)]. Here, we also consider spatial correlations in V(x, y) [see Fig. 1 (right panel)], characterized by a finite correlation length ξ, which can significantly affect the conduction properties [15].

III. RESULTS

In our simulations we first consider a quadratic region of width W = L = 1000 nm and an uncorrelated disorder potential (i.e., ξ → 0). Two geometries will be studied that only differ in their boundary conditions: a ribbon for which hard wall boundary conditions along the edges are applied and a cylinder with periodic boundary conditions in the y direction. A comparison between the disorder-averaged conductance ⟨G⟩ through these two geometries has been used previously to distinguish between bulk and edge phenomena as the periodicity of the cylinder eliminates the edges of the geometry [10]. The results for ⟨G⟩ in a topological insulator with m = −10 meV at Fermi energy E_F = 16 meV are shown in Fig. 2(a) as a function of the disorder strength U for both geometries. The value of E_F is chosen such that for uncorrelated disorder the TAI conductance plateau with ⟨G⟩ = 1 e^2/h clearly appears in the ribbon [8,10–12] for disorder strength 80 meV ≤ U ≤ 280 meV [see the red curve in Fig. 2(a)]. In the cylinder this plateau is clearly absent, since no edge states can exist in this edgeless geometry. While for disorder values beyond this plateau the conductance drops monotonically in the ribbon, the conductance through the cylinder geometry [blue dashed curve in Fig. 2(a)] shows a renewed increase at the same disorder strength. This is the signature of the aforementioned bulk delocalization that has already been observed in uncorrelated potentials [7–9]. A physical intuition for this transition is, however, still lacking, but will become clear by considering disorder potentials that are spatially correlated (illustrations for uncorrelated and correlated disorder potentials are shown in Fig. 1).

In a previous work [15] we demonstrated that spatial correlation in the disorder potential can destroy the TAI phase entirely. Here we consider a situation, in which the correlations all but dissolve the plateau in the ribbon geometry [see the red curve in Fig. 2(b)]. For these parameter values it is best
To get a better insight into this scenario, we now consider the scattering wave functions \( \psi(x_i, y_i) \) during this delocalization transition [see Figs. 3(a) and 3(c) for two such states at \( U = 220 \) meV]. A first optical inspection of these wave function images suggests that the associated flux is circumnavigating the hills of the underlying correlated potential [15], reminiscent of percolation states found in the quantum Hall effect [21] and in antidot topological insulator lattices [22]. To make this first impression more quantitative, we analyze how the intensities of the wave functions \( \psi(x_i, y_i) \) shown in Fig. 3 are correlated with the values \( V(x_i, y_i) \) of the underlying potential landscape. For this purpose we compute the weighted probability \( P(V) \) for a wave to encounter a potential value \( V \) with the weights of this probability distribution being given by the intensity \( |\psi(x_i, y_i)|^2 \) of the wave function at a grid point \( (x_i, y_i) \) with a potential value \( V \). The distribution \( P(V) \) resulting from an average over 1000 disorder realizations shows a surprisingly pronounced enhancement at positive disorder values \( V \) approximately situated between \( V_{\text{min}} \approx 25 \) meV and \( V_{\text{max}} \approx 75 \) meV (see Fig. 4), suggesting that disorder values from this interval give rise to clearly enhanced wave function intensities. Apparently the states responsible for the bulk delocalization tend to reside primarily at relatively high values of the disorder potential, i.e., in a certain altitude interval of the hills in the correlated potential landscape. Correspondingly, we find that the wave function intensities shown in Figs. 3(a) and 3(c) strongly resemble contour plots of the associated disorder potential, when we truncate the latter to the interval \( V \in [25, 75] \) meV [see Figs. 3(b) and 3(d)].

To identify the origin of this curious behavior, we first point to the fact that the above interval bounds, i.e., \( V_{\text{min}} \approx 25 \) meV and \( V_{\text{max}} \approx 75 \) meV, are astonishingly close to the minimal and maximal distances \( E_{\text{min}} = 26 \) meV and \( E_{\text{max}} = 78 \) meV of the Fermi energy \( E_F = 16 \) meV to the energy range \( E_{\text{bulk}} = [-10, -62] \) meV in which the valence band states are situated in the clean system [see the band structure of Fig. 5(a)]. This observation suggests that the flux in our correlated potential is carried mostly by the disorder analogs of these valence band states. Further evidence for this correspondence can be deduced when considering the rescaled probability distribution \( P(E_F - V) \), which measures, as above, the probability for a scattering state to reside at a potential value \( V \), but now shifted by the Fermi energy \( E_F \). We find that this distribution, quite remarkably, stays almost invariant with respect to a change...
of the Fermi energy [see a comparison between two different values of $E_F$ in Fig. 5(c)]. This observation reflects the fact that a change of $E_F$ just shifts the corresponding wave functions to different disorder values, but that the origin of states in the valence bands stays unchanged. Furthermore, the density of states $\rho(E)$, shown in Fig. 5(b), and the distribution $P(E_F - V)$, shown in Fig. 5(c), are very similar—even the small kinks in $\rho(E)$ are clearly reproduced in $P(E_F - V)$. Such kinks in the density of states are merely van Hove singularities resulting from the flat bands in the band structure. We may thus conclude that the flat valence band states, especially at the Brillouin zone (BZ) edges [where the maximum of $P(E_F - V)$ is located], represent the most significant contribution to the intensity of the scattering wave functions. In addition, we find that the distribution $P(E_F - V)$ is not at all sensitive to the boundary conditions since it is almost exactly equal for the ribbon and for the cylinder [see Fig. 5(c)]. The distribution $P(E_F - V)$ thus turns out to be quite fundamental in that it has its origin in basic system properties, which are given here by the band structure in Fig. 5(a) and by the flat band states contained in it.

These observations allow us to construct a comprehensive picture of the physics in the cylindrical geometry with a correlated potential [see the blue dashed curve in Fig. 2(b)]: While in the clean limit pure bulk conduction is observed, the conductance drops down to a minimum at disorder strength of $U \approx 60$ meV due to the increasing localization of the bulk states [see Fig. 2(b)]. When at $U \approx 100$ meV the hills of the correlated potential are high enough to locally shift the Fermi energy into the valence band, the bulk is filled with localized states deriving from clean valence band bulk states spatially located around the hills of the underlying disorder potential.

With growing disorder strength, these localized states connect with each other and go through a percolation threshold, which is responsible for the delocalization transition and the increased bulk conductance. Only at very strong disorder $U \approx 300$ meV does the connection between these percolating states weaken and the conductance again decreases. This picture is also strongly supported by previous studies of the TAI in the uncorrelated case (see Refs. [20,23]): Considering the arithmetic and geometric average of the local density of states, it was shown there that the states carrying the flux in the TAI are not single extended states throughout the whole TAI phase (as would be expected for edge states), but rather for very strong disorder formed by clusters of well-localized states. Our percolating wave functions deriving from clean valence band bulk states spatially located around the hills of the underlying disorder potential.

The flatness of the states in the effective band structure is, in fact, also important for the theory put forward in the aforementioned studies [20,23]: Considering disordered supercell structures, it was argued that flat and localized bands develop small subgaps that can be of topological nontrivial type. Hence these gaps have to be filled with edge states in the same way as the inverted band gap of a clean topological insulator [4,5,16]. In this picture the TAI phase
is thus characterized by edge states that appear in the energy gaps between localized bulk states and are consequently again immune from backscattering.

At this point the question arises how the above results can be reconciled with our own model, which so far does not contain any reference to edge states in the percolation transition. To investigate this issue in detail, we performed additional calculations for a system where no edge effects can be present due to a topological mass, which we choose to take on the positive value of \( m = 10.0 \) meV. As shown in Fig. 2(e), this sign change of \( m \) significantly modifies the conductance properties. While previously for \( m < 0 \) and moderate disorder strength \( U \) the conductance in the ribbon was clearly enhanced in comparison to the cylinder [see Fig. 2(b)], the conductance of the ribbon for \( m > 0 \) is even smaller than in the cylinder [see Fig. 2(c)]. This behavior can be attributed to the absence of edge states at the sample edges for positive topological mass \( m > 0 \). In the cylindrical geometries we find that the delocalization transition is less pronounced for \( m > 0 \) than it was for \( m < 0 \) [compare the blue dashed lines in Figs. 2(b) and 2(c)]. On the other hand, the fact that the delocalization transition still exists for \( m > 0 \) supports our model of flat bulk states undergoing a percolation transition. On the other hand, however, the more pronounced nature of the transition for \( m < 0 \) suggests that edge states propagating along the edges of the potential hills provide an additional link between localized states leading to a larger conductance. This picture, indeed, agrees very well with the analyses of Refs. [20,23], since the connecting local edge states in our model can directly be identified with the edge states that were predicted to form in the nontrivial subgaps of the localized flat bands.

We would thus be in a perfect position to complement the theory of Refs. [20,23] with the intuitive explanation that these subgap edge states exist locally and connect bulk states localized around hills of the potential landscape to form a percolating network of internal bulk states that lead to the decay of the TAI phase. The missing piece to complete our argument is to show that the picture we derived for the case of correlated disorder holds also for the uncorrelated case considered in Refs. [20,23].

We check this point explicitly, by verifying that our model can explain the appearance of the TAI as well as the observed delocalization-localization transition of the bulk states for the case of uncorrelated disorder. Consider, in this context, that the TAI conductance plateau in the ribbon geometry between \( U \approx 80 \) and \( 280 \) meV [see the red curve in Fig. 2(a)] is destroyed by the onset of the bulk delocalization at \( U \approx 280 \) meV in the cylinder [see the blue dashed curve Fig. 2(a)] which happens for much larger \( U \) than in the correlated case. Still, when the delocalization transition is in full effect (at \( U = 370 \) meV), the corresponding scattering states show a similar weighted distribution \( P(E_F - V) \) in the now spatially uncorrelated potential, as already observed in the correlated case [see Fig. 5(d)]. Again, the peak of this distribution fits nicely to the band structure of the clean limit, indicating that our picture of local edge states percolating around internal edges of strong disorder holds also for the uncorrelated case. Last but not least, we mention that such a percolating state corresponds exactly to the “single bulk state” that is held responsible for the delocalization in Ref. [11].

IV. DISCUSSION

Our results suggest that the emergence as well as the decay of the TAI phase depend strongly on the energy offset and on the flatness of the bulk valence bands in the clean limit. These flat bands feature an enhanced contribution to the density of states and occur in the center as well as at the boundaries of the BZ. Yet, the underlying BHZ model is only valid for small \( k_x \) close to the \( \Gamma \) point and thus does not yield a good approximation for the valence bands at the BZ boundaries of a real HgTe/CdTe quantum well [4]. Correspondingly, we find that when changing the grid spacing \( \alpha \) in our discretized lattice from the value conventionally used in the literature (\( \alpha = 5 \) nm) to different values, the position of the BZ boundaries \( k_x^{BZ} = \pm \pi/\alpha \) and the energy offset of these states at the BZ boundaries also change significantly. We also verified that the flatness of the bands at the BZ boundary is a direct consequence of the discreteness of the underlying lattice used for the numerical solution of the transport problem (see also Refs. [6,7,10] where discretized models were first employed to describe the TAI). As a result, the localization-delocalization transition and possibly even the TAI phase itself associated with these states at the BZ boundary will not occur in real HgTe/CdTe quantum wells as the strong-disorder limit in these devices will be different from the predictions of the discretized model. Quite remarkably, however, realizations of topological insulators have recently also been considered based on photonic systems [24,25]. These so-called Floquet topological insulators are based on a discretized lattice of sites, just as in the numerical model used above to approximate the physics in HgTe/CdTe quantum wells. The strong-disorder physics, which we have discussed here, may thus well be realized in experiments based on effective model systems in optics [24,25] as well as in acoustics [26] or in other fields where wave scattering parameters can be tuned appropriately.

V. CONCLUSION

In this paper we uncover the existence of percolating states in two-dimensional topological insulators. In particular, we show how these states affect the phase boundaries of the topological Anderson insulator, which is a topologically nontrivial phase caused by disorder. While the reason for the emergence of this phase has already been understood, its breakdown could so far only be vaguely connected to a delocalization of bulk states. Here we show that in a spatially correlated potential, this delocalization is caused primarily by bulk states that are localized when circumnavigating the hills of the disorder potential, but that become connected with each other when passing a percolation threshold. These connections and thus also the delocalization transition are consolidated by local edge states that can internally form in the disordered sample. By showing how the localized bulk states derive from flat bands in the valence band structure of the clean sample without disorder, we clarify that the same physics is at work also in the well-studied case of an uncorrelated disorder potential.
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