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Landau level spectroscopy of the PbSnSe topological crystalline insulator

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We report on an infrared magnetospectroscopy study of $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$, a topological crystalline insulator. We have examined a set of samples, all in the inverted regime of electronic bands, with the tin composition varying from $x = 0.2$ to 0.33 . Our analysis shows that the observed response, composed of a series of interband inter-Landau-level excitations, can be interpreted and modeled using the relativistic-like Hamiltonian for three-dimensional massive Dirac electrons, expanded to include diagonal quadratic terms that impose band inversion. In our data, we have not found any clear signature of massless electron states that are present on the surface of $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$ crystals in the inverted regime. Reasons for this unexpected result are discussed.

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I. INTRODUCTION

Topological crystalline insulators (TCIs) are systems that belong to the broader class of materials with a band structure endowed with nontrivial topological properties. TCIs are characterized by a well-defined bulk band gap and two-dimensional (2D) conical bands on the surface. To a great extent, they resemble the better known topological insulators (TIs) [1–3]. Nevertheless, the surface states of TCIs are not protected by time-reversal symmetry, but instead by the mirror or rotational symmetry of the crystal lattice. The existence of TCIs was confirmed experimentally in angle-resolved photoemission spectroscopy (ARPES) studies of lead-tin selenide [4], thus verifying pending theoretical predictions [5–8].

Lead-tin selenide is a mixed crystal widely explored in the past. Intensive investigations of this compound started in the 1960s (see, e.g., Refs. [9–14] or the reviews [15,16]), and they were often motivated by potential applications in thermoelectrometry and optoelectronics—in particular, by a possibility to use this material as an active medium in infrared lasers. THz/infrared magnetospectroscopy played an important role in these pioneering studies by providing us with useful insights into the electronic band structure [17–20].

The current consensus implies that $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$ is a narrow-gap semiconductor/semimetal. The fundamental energy band gap is direct and located at four nonequivalent L points, the centers of hexagonal facets of the first Brillouin

zone (Fig. 1; see also the beginning of Sec. IV for further details). The energy band gap drops with the increasing tin concentration x , and at liquid-helium temperatures, it closes at $x \approx 0.15$ [14,22,23]. For higher tin compositions, the band gap reopens but it becomes inverted, and 2D conical bands appear on the surface [4,24]. Such surface/interface states were predicted theoretically by Volkov and Pankratov [5,6], and later on they were classified using the Z_2 invariant [7]—this topological index takes on nontrivial values in a class of materials related to SnTe [8]. Nowadays, they continue to be a subject of intensive theoretical [25–28] and experimental studies [23,29–32], including the search for their magneto-optical signature [33–35].

In this work, we explore a series of $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$ samples using infrared magnetospectroscopy, and we observe a relatively rich magneto-optical response due to electronic excitations between pairs of bulk Landau levels. The analysis of the collected data allows us to extract the band-structure parameters, and eventually to get a quantified theoretical estimate for the magneto-optical response due to surface electrons. Nevertheless, our experiments performed in high magnetic fields—strong enough to approach the quantum limit of surface electrons and to probe their fundamental cyclotron mode—do not show a clear signature for the magneto-optical response originating from the surface states.

II. PREPARATION AND CHARACTERIZATION OF PbSnSe SAMPLES

A series of lead-tin selenide ($\text{Pb}_{1-x}\text{Sn}_x\text{Se}$) samples (denoted here by letters A, B, C, D, E, and F) was grown

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FIG. 1. Bulk Brillouin zone for a rocksalt lattice. Red points mark the positions of the four nonequivalent L points at the border of the zone. The blue hexagon shows the surface Brillouin zone in the (111) direction. The relativistic-like conical bands appear at the $\bar{M}$ and $\bar{\Gamma}$ points, which are surface projections of the bulk L points [21].

using the hot-wall-epitaxy method on a (111)-oriented BaF_2 substrate, with a weak doping by bismuth that ensures n -type conductivity. The thickness of epilayers ranged from 1 to 2.55 μm , and the performed x-ray analysis indicated their (111)-orientation. The nominal tin content was chosen at $x = 0.20, 0.25$, and 0.33 . The composition of samples D and F was cross-checked using energy-dispersive x-ray spectroscopy (EDS). The obtained results— $x = 0.22 \pm 0.02$ and 0.34 ± 0.02 , respectively—show a weak deviation from the nominal values. With the chosen tin contents, the samples are expected to have an inverted band structure at liquid-helium temperatures, but they are still away from the crossover from the cubic (rocksalt) to orthorhombic phase that occurs around $x \approx 0.4$ [4]. All prepared samples were characterized using the magnetotransport technique at room temperature, which indicated n -type conductivity (Table I).

III. MAGNETO-OPTICAL EXPERIMENTS

The optical response of our PbSnSe samples was probed in the middle infrared spectral range using the magnetotransmission technique in the Faraday configuration, with the wave vector of light perpendicular to the sample surface. To measure magnetotransmission, unpolarized radiation of a global

TABLE I. List of explored samples with the nominal Sn content x , carrier density n , and electron mobility μ deduced from room-temperature transport experiments:

Sample	x	d (μm)	μ [$\text{cm}^2/(\text{V s})$]	n (10^{18} cm^{-3})
A	0.20	0.8	720	10
B	0.20	0.7	580	9.0
C	0.25	2.5	1200	1.4
D	0.25	2.55	970	5.0
E	0.25	2.5	1200	2.5
F	0.33	1	350	11

was modulated in a Fourier-transform spectrometer and delivered to the sample using light-pipe optics. Typically, several mm^2 of the surface was exposed to the radiation, which was then detected by a composite bolometer placed directly below the sample. The measured transmission spectra T_B were corrected for the field-induced changes in the response of the bolometer. During experiments, the samples were kept in helium exchange gas at $T = 2 \text{ K}$ and placed either in a superconducting (below 13 T) or resistive (above 13 T) coil.

To illustrate the observed magneto-optical response, we have chosen the data collected on sample C and plotted them in Fig. 2. In panel (a), we show a stack plot of relative magnetotransmission spectra, T_B/T_0 , for selected values of the magnetic field. The false-color plot of relative magnetoabsorbance, $A_B = -\ln(T_B/T_0)$, is presented in panel (b). At $B > 2 \text{ T}$, the transmission of sample C becomes modulated by a series of excitations that follow a sublinear dependence in B . These excitations may be straightforwardly identified as bulk inter-Landau-level (inter-LL) transitions that promote electrons from the valence to the conduction band.

To read out the positions of excitations, we have examined the differential magnetotransmission spectra $T_B/T_{B-\delta B}$, where δB is the field step, and we searched for the inflection points in regions with a negative slope. These can be associated, with reasonable precision, with the positions of excitations that shift relatively fast with the applied magnetic field. This procedure was implemented to suppress the impact of a pronounced Fabry-Pérot interference pattern, which appears in the transmission spectra of epilayers at wavelengths comparable to their thicknesses and which are still partly visible in the T_B/T_0 spectra. The deduced fan chart of inter-LL transitions observed in the magnetotransmission of sample C is presented in Fig. 3(a). Up to 10 individual transitions can be distinguished in the magneto-optical response of this particular sample.

IV. MAGNETO-OPTICAL RESPONSE OF BULK STATES—THEORETICAL MODEL

The parent compound of $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$, rocksalt structure lead selenide, is a semiconductor with a band gap around 0.15 eV at liquid helium temperatures [16,36]. Close to the gap, both cation and anion atomic orbitals contribute significantly to the electronic states whereas Se (Pb) dominates in the doubly spin-degenerate valence(conduction) band which transforms according to the Γ_{+6} (Γ_{+6}) irreducible representation. An effective Hamiltonian can be derived using standard $\mathbf{k} \cdot \mathbf{p}$ perturbation theory, applied to the first order and to a finite number of bands, to describe the electronic states around the L point of the Brillouin zone [9]: $\mathbf{q} = (q_x, q_y, q_z) = \mathbf{k} - \mathbf{k}_L$, being the small parameter.

With a full electron-hole symmetry, which holds up to almost 1 eV away from the gap in PbSe [36], we therefore assume that the low-energy excitations in the bulk of $\text{Pb}_{1-x}\text{Sn}_x\text{Se}$ can be described by

$$\hat{H}_0 = \begin{pmatrix} \Delta + Mq^2 & \hbar v q_+ & 0 & -\hbar v q_z \\ \hbar v q_- & -\Delta - Mq^2 & \hbar v q_z & 0 \\ 0 & \hbar v q_z & \Delta + Mk^2 & \hbar v k_- \\ -\hbar v q_z & 0 & \hbar v q_+ & -\Delta - Mq^2 \end{pmatrix}, \quad (1)$$

FIG. 2. (a) The stacked-plot of relative magnetotransmission data, T_B/T_0 , collected on sample C at $T = 2$ K at magnetic fields up to $B = 33$ T with steps of 1 T. For clarity, individual spectra were shifted vertically by B (T) $\times 0.24$. (b) False color plot of relative magnetoabsorbance, $A_B = -\ln(T_B/T_0)$, measured on sample C. The white lines separate the areas of the data collected using a superconducting and resistive coil.

where $q_{\pm} = q_x \pm iq_y$, and v , Δ , and M are band-structure parameters that depend on x . This Hamiltonian implies an isotropic dispersion for both conduction and valence bands

($\beta = 1$ and -1 , respectively):

$$E_{\beta}(\mathbf{q}) = \beta \sqrt{(\Delta + Mq^2)^2 + \hbar^2 v^2 q^2} \quad (2)$$

FIG. 3. (a) Fan chart of inter-LL excitations determined experimentally for sample C using a superconducting (squares) and resistive coil (circles). The theoretically calculated energies of inter-LL excitations are plotted as solid lines for parameters deduced for sample C: $2\Delta = 55$ meV, $M = -20$ eV $\text{\AA}^2$, and $v = 4.4 \times 10^5$ m/s (Table II). (b) Landau level spectrum (at $q_z = 0$) calculated for the parameters deduced for sample C. The levels plotted in dark cyan and violet correspond to the $\alpha = 1$ and -1 LL series, respectively. The blue line shows the calculated position of the Fermi energy assuming that the number of electrons is independent of B . The vertical arrows show the interband inter-LL excitations active in the electric-dipole approximation. The color coding in both panels has been chosen to facilitate the identification of corresponding transitions.

with both dispersion branches doubly degenerate for spin.

The Hamiltonian (1) is equivalent to the three-dimensional Dirac Hamiltonian for massive electrons, where the velocity parameter v can be viewed as an effective speed of light. It is expanded to include dispersive diagonal elements, Mq^2 , which account for the impact of remote bands as well as the band inversion. For a relatively small inversion parameter, $|M| < \hbar^2 v^2 / \Delta$, the energy band gap of 2Δ appears at $\mathbf{q} = \mathbf{0}$. Notably, the Hamiltonian closely resembles the one proposed by Liu *et al.* [37,38] for the electronic states around the Γ point in the Bi_2Se_3 family of topological insulators.

The applied magnetic field ($B \parallel z$) transforms each electronic band into Landau levels. When the motion of electrons along the applied magnetic field is neglected ($q_z = 0$), the LL spectrum reads

$$E_{\alpha,\beta}^n|_{q_z=0} = \alpha \frac{M}{l_B^2} + \beta \sqrt{\left(\Delta + 2\frac{M}{l_B^2}n\right)^2 + 2\frac{v^2\hbar^2}{l_B^2}n}, \quad (3)$$

where $n = 1, 2, 3, \dots$, and $l_B = \sqrt{\hbar/eB}$ is the magnetic length. This spectrum describes LLs in the conduction and valence bands ($\beta = 1$ and -1 , respectively) as well as their splitting with respect to spin ($\alpha = \pm 1$). In addition, there exists a pair of $n = 0$ LLs that are spin-polarized and that disperse linearly with B : $E_{1,1}^0 = \Delta + M/l_B^2$ and $E_{-1,-1}^0 = -\Delta - M/l_B^2$. In the inverted regime, $M < 0$, these so-called zero-mode LLs (anti)cross each other, which is behavior typical of all topological (crystalline) insulators [3,30,31,39].

Knowing the LL spectrum and the corresponding eigenstates, one may proceed with the calculation of the dynamical magnetoconductivity that encodes the response in magneto-optical experiments. In the linear-response theory, the dynamic conductivity—for the right (+) and left (−) circularly polarized light—is given by

$$\sigma^\pm(\omega, B) = 2 \frac{iG_0\zeta}{l_B^2\omega} \sum_{n,m,q_z} \frac{(f_m - f_n)|\langle m|\hat{v}_\pm|n\rangle|^2}{E^n - E^m - \hbar\omega + i\gamma}, \quad (4)$$

where $G_0 = e^2/(2\pi\hbar)$ is the quantum of conductance, γ is the broadening parameter, and ζ stands for the valley degeneracy ($\zeta = 4$; see Fig. 1). The occupation of LLs is given by the Fermi-Dirac distribution f , and the indices m and n run over all available initial and final LLs, including states with nonzero momenta q_z . The velocity operators $\hat{v}_\pm = \hat{v}_x \pm i\hat{v}_y$ are defined as $\hbar\hat{v}_x = \partial\hat{H}/\partial q_x$ and $\hbar\hat{v}_y = \partial\hat{H}/\partial q_y$, respectively, and are rewritten in the representation of ladder operators $a^\pm = l_B(q_x \mp iq_y)$.

The matrix elements $\langle m|\hat{v}_\pm|n\rangle$ contain the selection rules for electric-dipole excitations between various pairs of LLs. The dominant contribution to the magneto-optical response arises from the states with vanishing momenta q_z for which those rules have a particularly simple form. The spin parameter α is always conserved, and, for photon energies not exceeding the band gap considerably, only two series appear: the $\alpha = 1$ series with the selection rule $n \rightarrow n - 1$, and the $\alpha = -1$ series with $n \rightarrow n + 1$. These are active for left and right circularly polarized radiation, respectively (see, e.g., Refs. [40,41]).

TABLE II. The energy band gap 2Δ , velocity v , and inversion parameter M extracted from the magneto-optical response of the samples A, B, C, D, E, and F.

Sample	Sn content	2Δ (meV)	v (10^5 m s $^{-1}$)	M (eV Å 2)
A	0.20	30 ± 10	4.5 ± 0.1	-12 ± 5
B	0.20	23 ± 10	4.3 ± 0.2	-15 ± 2
C	0.25	55 ± 5	4.4 ± 0.1	-20 ± 2
D	0.25	60 ± 10	4.4 ± 0.1	-19 ± 5
E	0.25	36 ± 10	4.4 ± 0.2	-21 ± 6
F	0.33	104 ± 5	4.2 ± 0.1	-24 ± 2

V. MAGNETO-OPTICAL RESPONSE OF BULK STATES—DISCUSSION

Let us now confront our experimental data with predictions of the sketched theoretical model. In the first step, we compare the positions of inter-LL excitations deduced from our relative magnetotransmission/absorbance data with theoretical expectations. Such a comparison is illustrated using the experimental data collected on sample C [Fig. 3(a)]. Individual observed transitions were interpreted based on the response in high magnetic fields when bulk electrons supposedly reach their quantum limit. The interband transition with the lowest energy may then be assigned to the fundamental electric-dipole-active interband mode, which promotes electrons from the holelike $n = 0$ level to the spin-up $n = 1$ level [Fig. 3(b)].

Very good agreement between our simple theory and experimental data presented in Fig. 3(a) corroborates the validity of the model. The positions as well as the field dependence of all observed transitions are reproduced by using only three adjustable parameters: Δ , M , and v . Notably, our model accounts for the response of both longitudinal and oblique valleys (L_1 versus $L_{2,3,4}$ in Fig. 1). Therefore, no anisotropy of the velocity parameter v had to be considered in order to explain the magneto-optical data collected on sample C (Figs. 2 and 3) nor on the other explored samples. This conclusion agrees with a relatively low anisotropy of the electronic bands at the L point in PbSe (expressed, e.g., in terms of the anisotropic effective masses [16,19]), which further decreases upon incorporation of tin [23]. This is in contrast to some other lead-salt compounds, such as PbSnTe, in which the anisotropy is profound [29,42], thus giving rise to well-resolved splitting of interband inter-LL resonances due to oblique and longitudinal valleys.

The deduced values of parameters for all six explored samples are listed in Table II and visualized in Fig. 4. We observe that the energy band gap ($E_g = 2\Delta$) increases roughly linearly with tin content. The same behavior is observed for the absolute value of the inversion parameter $|M|$. The difference in the extracted band gap for samples with presumably the same tin concentration points toward a certain variation of x as compared to the declared nominal values, in line with the EDS results mentioned before. The deduced velocity parameters show a weak tendency to decrease with x . Such behavior is perfectly in line with previous studies [19,23,43].

Notably, the characteristic shape of the conduction and valence bands described by the Hamiltonian (1) can be assessed using the parameter $\eta = 4|M|/\Delta/(\hbar^2 v^2)$. For $\eta = 1$, the

FIG. 4. Band-structure parameters extracted from the magneto-optical data as a function of the nominal tin concentration: (a) energy band gap $E_g = 2\Delta$, (b) velocity parameter v , and (c) inversion parameter M . The symbols connected by dashed lines indicate the band gap and velocity parameter values extracted in Refs. [22] and [23].

electronic bands described by Eq. (2) are strictly parabolic: $E_{\pm}(\mathbf{q}) = \pm(\Delta + |\mathbf{M}|q^2)$, and the LL spectrum as well as the whole magneto-optical response then scale linearly with B . This occurs, for instance, at the Γ point of the well-known Bi₂Se₃ topological insulator. For all PbSnSe samples investigated in this work, we obtained $\eta < 1$. This implies subparabolic profiles of bands, and consistently with our experiments, magneto-optical excitations following a sublinear dependence in B (Figs. 2 and 3). An extrapolation of band-structure parameters as a function of the tin content [see Figs. 4(a)–4(c)] suggests that the condition $\eta = 1$ might be achieved in Pb_{1-x}Sn_xSe at concentrations slightly above $x = 0.35$.

In the second step, we compare the experimentally determined relative magnetotransmission spectra, T_B/T_0 , with those calculated theoretically; see Fig. 5. To obtain the theoretical spectra, we have numerically diagonalized the full Hamiltonian (1) in which the magnetic field was introduced using the standard Peierls substitution. The calculated eigenstates (Landau levels/bands) and the corresponding spectrum were then used to evaluate the optical conductivity (4).

FIG. 5. Relative magneto-transmission spectra T_B/T_0 at selected values of the applied magnetic field compared to those calculated using the linear-response theory (4). To calculate T_B and T_0 , the Landau levels up to the index 20 and 200 were included, respectively. The contribution of excitation at nonzero momenta were taken into account up to $q_z = 1.25 \text{ nm}^{-1}$. Two cases have been considered to account for the disorder in the system: $\gamma = 2 \text{ meV}$ (red curve) and $\gamma = 0.05 \text{ meV} + 0.0125\hbar\omega$ (meV) (blue curve).

Afterward, magnetotransmission spectra $T_B^{\pm}$ of the slabs with given thicknesses d (Table I) were calculated for both circular polarizations of light [44]. The background dielectric constant, $\epsilon = 20$, was estimated from the Fabry-Pérot interference pattern (cf. Ref. [45]). The average transmission spectrum $T_B = (T_B^+ + T_B^-)/2$ was then normalized by T_0 , approximated by T_B calculated at the magnetic field below the onset of the LL quantization in the theoretical and also experimental response ($B = 1 \text{ T}$).

To reproduce the measured magnetotransmission spectra, the broadening parameter γ had to be adjusted. The simplest choice—a parameter independent of the energy and magnetic field—led to acceptable results (red curves in Fig. 5). The used value of $\gamma = 2 \text{ meV}$ is consistent with the onset of the LL quantization observed in the magneto-optical data ($B \approx 2 \text{ T}$). The agreement is further improved by taking an effective value of γ that increases linearly with the photon energy: $\gamma = \gamma_0 + \xi\hbar\omega$, with $\gamma_0 = 0.05 \text{ meV}$ and $\xi = 0.0125$. This latter choice resembles the empirical rule deduced for interband inter-LL excitations in graphene [46,47]. Notably, for relatively weak disorder, the width of excitations is not entirely determined by the γ parameter. The pronounced high-energy tail of interband inter-LL excitations appears due to excitations at nonzero q_z momenta. Moreover, even though the electron-hole asymmetry was not directly evidenced in our data, it may still influence the width of transitions. This latter contribution would be roughly equal to the difference between the cyclotron energies of electrons and holes.

The carrier concentration, or alternatively the Fermi energy, is another tunable parameter in our model. In our calculations, we considered the electron density to be constant

as a function of B . The electron densities, tuned to achieve the best possible agreement with the magneto-optical data, were smaller by roughly a factor of 2 as compared to the results of room-temperature Hall experiments (Table I). The sample F was the only exception—a carrier density almost one order of magnitude smaller than the one deduced from room-temperature characterization had to be considered to reproduce our data. To adjust the Fermi energy, the appearance of the lowest bulk interband inter-LL excitation ($0 \rightarrow 1$) was taken as a main reference point.

The comparison of the experimental and theoretical magnetotransmission spectra in Fig. 5 indicates that fairly good agreement is achieved at higher photon energies, both for the shape and intensity of excitations. At lower energies, a certain deviation may be observed in the range corresponding to the excitations of electrons from the valence band to the vicinity of the Fermi energy in the conduction band. The relatively pronounced maximum in the theoretical T_B/T_0 spectra (around 120 meV in Fig. 5) is not observed experimentally. It appears in the calculated spectra as a result of B -induced splitting of the interband absorption edge for opposite circular polarization of the incident light. In our experimental data, this feature seems to be smeared out due to the inhomogeneity of the electron density (fluctuation of the Fermi energy) across the explored sample.

VI. MAGNETO-OPTICAL RESPONSE OF SURFACE STATES

Let us now focus on possible signatures of surface states in our magneto-optical data. In contrast to the (001)-oriented surface of PbSnSe samples [48], the topologically nontrivial states on the (111) surface have, according to the existing theoretical models, a relatively simple structure [8,21]. Four conical bands are expected at the $\bar{M}$ and $\bar{\Gamma}$ points, which are projections of the bulk L points where the electronic bands are inverted (Fig. 1).

The particular shape of conical bands—their slope and anisotropy—is given by the properties of the bulk velocity parameter. As discussed above, we have found no differences in the magneto-optical response of the longitudinal and oblique valleys. Therefore, we consider the velocity parameter to be isotropic in our PbSnSe samples in the explored range of tin content. In this approximation, both the surface and the interface of our PbSnSe epilayers host four identical conical bands with the slope defined by the bulk velocity parameter v (Table II).

When a strong enough magnetic field is applied, 2D cones split into a series of non-equidistantly-spaced LLs:

$$E_{2D}^n = \pm v \sqrt{2e\hbar n B}. \quad (5)$$

The magneto-optical response of systems with such a LL spectrum has been widely studied, both theoretically and experimentally, on graphene and other systems [46,47,49–55]. The response is dominated by a series of interband and intraband inter-LL excitations that follow the standard electric-dipole selection rules in isotropic systems: $n \rightarrow n \pm 1$.

When the quantum limit of electrons in a conical band is approached, the fundamental cyclotron resonance (CR)

mode emerges in the optical response. This mode comprises excitations of electrons among the $n = 0$ and 1 levels and becomes—in terms of the integral strength and absolute absorption—the most pronounced line in the transmission spectrum. This simple empirical rule can be justified using the standard linear-response theory. Therefore, we focused on this particular excitation in order to find a magneto-optical signature of the surface states in PbSnSe. We applied high magnetic fields to our samples to drive surface electrons close to their quantum limit. At the same time, their fundamental CR mode is expected to move above the reststrahlen band of BaF₂. Hence, it should be traceable using the infrared magnetotransmission technique.

The latter fact is illustrated in Fig. 6. The LL spectra of bulk and surface electrons, expected for sample C, are shown in panel (a), along with the estimated Fermi energy. In panel (b), we present a false-color plot of magnetoabsorbance of sample C overlaid with the expected energies of bulk and surface inter-LL resonances. Panel (b) shows that the fundamental CR mode of surface electrons should be accompanied by two bulk resonances at nearby energies—the fundamental bulk CR mode and the lowest bulk interband inter-LL transition. These three excitations are marked by green, cyan, and black vertical arrows in Fig. 6(a). The same color-coding is used for the curves in Fig. 6(b). The dotted parts of the theoretical curves indicate the magnetic fields at which the final-state Landau level is occupied, and therefore the transition should not appear due to Pauli blocking. Here we considered the bulk Fermi energy to determine the occupation of the surface LLs.

To explore the magneto-optical response in greater detail, the differential magnetotransmission spectra, $T_B/T_{B-\delta B}$, were analyzed [Figs. 7(a)–7(f)]. As discussed in Sec. III, the impact of a weakly field-dependent Fabry-Pérot interference pattern is partly suppressed in the differential spectra, and we may better follow individual inter-LL resonances. Each transition manifested as a minimum in the relative magnetotransmission, T_B/T_0 , is now expected to be seen as a derivative-like feature (a maximum followed by a minimum) in the differential spectra $T_B/T_{B-\delta B}$. To facilitate our analysis, the theoretically expected positions of the three lowest excitations are marked by vertical arrows in Figs. 7(a)–7(f), using the color-coding introduced in Fig. 6. The transparent arrows correspond to excitations blocked by the occupation effect. Again, we assume here that the occupation of the surface LLs is given by the bulk Fermi energy. Following these assumptions, the fundamental CR mode of surface electrons should be observable above the reststrahlen band of BaF₂ in all explored epilayers, except sample D.

Prior to continuing the analysis of the experimental data, let us discuss the strength of the searched contribution from surface electrons. We performed a simple calculation using the standard transfer-matrix formalism [56]. The PbSnSe epilayer was modeled as a dielectric slab with a thickness d (Table I) and a refractive index of $n_{\text{PbSnSe}} = 5$. The optical response of the surface/interface states was described by the 2D dynamical magnetoconductivity. It comprised a single mode at $\hbar\omega = E_{2D}^{n=1}$, with the broadening parameter $\gamma = 2$ meV extracted from the bulk response, and the integral strength corresponding to the fourfold degeneracy. We assumed that this mode emerges when the bulk Fermi energy drops below the

FIG. 6. (a) Landau level spectrum of surface and bulk states and the expected Fermi energy in sample C as a function of B . The dashed and solid lines used for the LLs reflect the spin parameter $\alpha = -1$ and 1 , respectively. The vertical arrows show three inter-LL excitations with lowest energies expected in the spectrum: the bulk interband $0 \rightarrow 1$ transition, and fundamental CR resonances of bulk and surface electrons. (b) False-color plot of relative magnetoabsorbance measured on sample C compared to the theoretically expected positions of bulk (black and green) and surface (dark cyan) inter-LL resonances. The dotted lines correspond to the values of the magnetic field at which the transitions are not active due to occupation of the final-state LL (Pauli blocking). The vertical white line separates the data measured at low magnetic fields using a superconducting coil from the high-field data obtained with a resistive coil.

$n = 1$ LL of surface electrons. The substrate was represented by a dielectric slab with a thickness of $500 \mu\text{m}$ and a refractive index of $n_{\text{BaF}_2} = 1.4$. The calculated differential spectra (cyan dotted lines) were then plotted along with experimental data in Fig. 7. The inflection point in the calculated curves matches the energy of the fundamental CR mode of surface electrons, marked by vertical cyan arrows. When the fundamental CR mode is Pauli-blocked, the arrows are transparent. Due to the interference effect, the calculated amplitude of the surface CR mode varies with the thickness of the epilayers. Nevertheless, the theoretically expected signal clearly exceeds the noise level for all explored samples. For the sake of completeness, let us note that the strength of the CR mode is much lower as compared to free-standing graphene [47]—the difference in the velocity parameters reduces the strength roughly by a factor of 4; further reduction occurs via the dielectric effect, due to PbSnSe bulk slab and BaF₂ substrate.

From the experimental viewpoint, the differential spectra measured on samples C and F reveal no signatures attributable to the signal from the surface states. For other samples, some features develop just above the reststrahlen band of BaF₂. Nevertheless, their position, amplitude, and shape do not correspond to the expectations for surface electrons. The relatively pronounced minimum observed for samples A and E (around $\hbar\omega \approx 100 \text{ meV}$ at 30 T) is most likely due to bulk interband inter-LL excitations ($n = 0 \rightarrow 1$). Their position

seems to be slightly overestimated in our bulk data modeling (black vertical arrows). The shape—a minimum instead of a derivative-like profile—may suggest the presence of some additional weaker excitations. However, there is no solid argument connecting them with the surface states. Below, we discuss three different scenarios that allow us to explain why no clear signatures of surface electrons were found in the magneto-optical response: (i) an occupation effect, (ii) disorder, and (iii) too simplistic theoretical modeling.

(i) The occupation of surface states, or in other words the distribution of electrons among the surface LLs, may differ considerably from the one expected from the position of the bulk Fermi energy. The band bending close to the surface or interface of bulk crystals is a well-established phenomenon, often discussed in the context of topological (crystalline) insulators and other narrow-gap semiconductors [57–59]. This effect is governed by standard rules of electrostatics, and it may lead to the appearance of both depletion or accumulation of bulk electrons near the surface/interface. The latter possibility could explain the lack of signatures of surface states in our data. Nevertheless, for our samples with relatively large electron doping, and consequently large density of positively charged defects, a relatively narrow depletion layer and only a weak band-bending effect are expected.

(ii) The topological protection makes surface states robust against the opening of a band gap, but they still remain

FIG. 7. Differential magnetotransmission spectra, $T_B/T_{B-\delta B}$, plotted for all explored samples A, B, C, D, E, and F in parts (a)–(f), respectively, using the difference $\delta B = 1$ T. The spectra are shifted vertically, with the offset scaling linearly with B . The energies of three excitations having the lowest energy in the quantum limit of bulk and surface states are marked by the vertical arrows [cf. Fig. 6(b)]: green, the bulk fundamental CR mode; dark cyan, the fundamental CR mode of surface electrons; and black, the lowest interband inter-LL transition ($n = 0 \rightarrow 1$). Transitions are expected to emerge in the spectra (following the occupation effect) when transparent arrows change into solid ones. The dotted cyan line is the theoretically expected response, the differential magnetotransmission, due to the fundamental CR of surface electrons, calculated using the transfer-matrix method; see the text. Neither bulk inter-LL resonances nor the presence of the reststrahlen band is included in this model for simplicity.

sensitive to disorder, which is ubiquitous in a random alloy (note that Sn atoms do not replace Pb atoms in any periodic or regular manner). In magneto-optical experiments on 2D systems, the strength of disorder is directly reflected by the widths of inter-LL excitations. In our rule-of-thumb model for the fundamental CR mode of surface electrons, we considered the characteristic broadening equal to that of bulk states. In this way, we might have neglected additional sources of disorder, such as the surface/interface roughness or surface point defects. In fact, our transfer-matrix simulations suggest that an increase of γ by a factor of 3–5 brings the calculated magneto-optical signal of the surface states down to the experimental noise level. Disorder in the surface states thus may represent a plausible explanation of the apparently missing signal from the surface electrons.

(iii) The accuracy of the existing theoretical models represents another possible explanation. Clearly, there is no apparent reason to question their general validity [8,21], notably, when they are convincingly corroborated by various experimental studies [4]. At the same time, however, it is not obvious to what extent these models account for the surface states quantitatively; strictly speaking, the concept of band structure does not even apply to a random alloy. Cushioning such fundamental doubts, the dispersion of the surface states does not need to have its Dirac point in the midgap position (reflecting the weak but present electron-hole asymmetry at the L point). This would effectively correspond to a higher Fermi level for the surface states, thus pushing the quantum limit to larger magnetic fields and making the fundamental CR mode of surface electrons hardly observable in our experiments. In addition, one may expect a certain deviation from the linearity of the conical band on the surface, which is not included in the simplest model [21]. The fundamental CR mode of surface electrons could thus have a lower energy and stay hidden within the reststrahlen band of the BaF₂ substrate even in the highest magnetic fields applied in our experiments. Hybridization of surface and bulk states is another possible effect. Sticking strictly to the standard theoretical model [21], the bulk and surface states are truly eigenstates of a semi-infinite crystal, and they do not mix with each other. At the same time, the discrete surface LLs coexist—apart from the interval in between zero-mode levels—with the quasicontinuum of bulk electronic states, and one may speculate about mechanisms that mediate their coupling. The hybridization

of surface and bulk states may represent an explanation—for an alternative to plain disorder—for spectrally broad features observed above the reststrahlen band of BaF₂ in the differential spectra of the A, B, D, and E samples.

VII. CONCLUSIONS

A series of PbSnSe topological crystalline insulator samples has been explored using the infrared spectroscopy technique in high magnetic fields. We show that the fairly rich response observed, comprising a sequence of transitions between bulk Landau levels, can be successfully explained using a simple model of massive Dirac electrons, expanded to include the effects of the band inversion. Despite expectations, and perhaps surprisingly for PbSnSe epilayers with a relatively high electronic quality, we do not find any clear signature of symmetry-protected surface states, which are the most salient features of topological crystalline insulators, and which could—similar to topological insulators—give rise to appealing magneto-optical effects such as universal Faraday and Kerr rotation [60–64]. We discuss several scenarios that may account for this: the band-bending effect, surface disorder, and the hybridization between bulk and surface electronic states.

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