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THz response and colossal Kerr rotation from the surface states of the topological insulator Bi₂Se₃

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We report the THz response of thin films of the topological insulator Bi_2Se_3 . At low frequencies, transport is essentially thickness independent showing the dominant contribution of the surface electrons. Despite their extended exposure to ambient conditions, these surfaces exhibit robust properties including narrow, almost thickness-independent Drude peaks, and an unprecedentedly large polarization rotation of linearly polarized light reflected in an applied magnetic field. This Kerr rotation can be as large as 65° and can be explained by a cyclotron resonance effect of the surface states.

Ordered states of matter are typically categorized by their broken symmetries. With the ordering of spins in a ferromagnet or the freezing of a liquid into a solid, the loss of symmetry distinguishes the ordered state from the disordered one. In contrast, topological states are distinguished by specific *topological* properties that are encoded in their quantum mechanical wavefunctions [1]. Frequently, a consequence of these properties is that there are robust "topologically protected" states on the sample's boundaries. The edge states of the quantum Hall effect (QHE) are the classic example [2]. In the last few years, it was realized that another class of such topological matter may exist in 3D band insulators with large spin-orbit interaction [3–5]. These so-called topological insulators are predicted to host robust surface states, which exhibit a number of interesting properties including spin helicity, immunity to back-scattering, and weak anti-localization. There are predictions of a number of unusual phenomena associated with these surface states, including a proximity-effect-induced exotic superconducting state with Majorana fermions bound to a vortex [6, 7] and an *axion* electromagnetic response [8, 9], and proposals for applications, such as their use in terahertz (THz) devices [10].

Most of the signatures of topological behavior in these materials thus far have come from surface probes such as angle resolved photoemission (ARPES) and scanning tunneling spectroscopy [11–16]. These experiments have revealed that the surface states indeed show signatures of the predicted topological properties, such as a Diraclike dispersion, chiral spin textures, and the absence of backscattering. Direct observation of the topological behavior in transport has been hampered by the lack of a true bulk insulating state. Only recently have transport experiments started to distinguish the surface contribution from the bulk [17–22].

As opposed to the case of the quantum Hall effect, in topological insulators, the quantization of the offdiagonal conductivity is not a requirement for the existence of the topological state. This, along with the problem of bulk conduction, has made finding a unique signature of this state difficult. It has been proposed that topological insulators may be characterized by their electrodynamic properties [8] due to the existence of an axionic term in the action $\Delta \mathcal{L} = \alpha \theta \int dx dt \mathbf{E} \cdot \mathbf{B}$, where α is the fine structure constant. θ is a modulo 2π number that distinguishes between topologically non-trivial $(\theta = \pi)$ and trivial insulators $(\theta = 0)$. There have been several proposals of how to measure this θ parameter using polarized electromagnetic radiation at low frequencies [8, 23–27]. All these proposals predict that linearly polarized THz range electromagnetic waves reflected from (transmitted through) a sufficiently low disorder topological insulator in a magnetic field, will undergo a Kerr (Faraday) rotation of the polarization plane, the magnitude of which depends on the experimental configuration, but it is set by the fine structure constant α . Thus, such experiments can directly probe the topological nature of the surface states for samples with true insulating bulk and low sheet carrier density. With the current available samples one is able to access the semi-classical regime of large Landau level index.

In this Letter, we report the THz response of the topological surface states (TSS) in thin films of the topological insulator Bi₂Se₃. We measured thin films of several thicknesses grown on sapphire (Al₂O₃) substrates by molecular beam epitaxy. The time elapsed between growth and THz measurements was at least 2 weeks for each sample [28]. Due to their small thickness, low bulk carrier density and high mobilities, these films have been recently shown to exhibit 2D behavior dominating the DC transport for film thicknesses of less than 200 QL [21]. We find clear signatures of the protected surface states in the 2D behavior of the THz conductivity obtained using time domain terahertz spectroscopy (TDTS). In addition we find a colossal Kerr rotation almost entirely due to the TSS. The Kerr rotation measurement allows us to extract the effective mass of the 2D Dirac electrons.



FIG. 1. **a**). Real part of the conductance of four films of different thicknesses (16, 32, 64 and 100 QL, 1 QL=9.4Å). **b**). Spectral weight of the Drude and the phonon contributions obtained from the fit to the conductance (symbols). Lines are guides to the eye. **c**). Schematic of experiment, where multiple echoes may be separated in time (top). The bottom figure shows the polarizer arrangement for experiments in a magnetic field, the first and last polarizers can be set either parallel or perpendicular to each other, and the middle one can be rotated. **d**). Time domain trace of the transmission through a bare Al₂O₃ (top, displaced vertically for clarity), and a thin film sample (bottom) at 2 K.

In Fig. 1a we show typical data of the real part of the longitudinal conductance ($G_{xx} = \sigma_{xx}t$) of a number of samples of different thicknesses (t) at 6 K and at zero magnetic field. The data show a clear signature of free electron behavior with a reasonably narrow (≈ 1.2 THz wide) Drude peak centered at zero frequency. In addition, we observe a contribution from an optical phonon close to 2 THz. The data are qualitatively similar to ones reported by others in single crystals [29, 30]. We fit these data with a model consisting of two identical Drude terms (one for each surface), and a Drude-Lorentz term for the bulk phonon. These fits allow an essentially perfect parameterization of the data (see ref.[28]), showing that the low frequency transport is dominated by a single charge species. The surface Drude term has an almost thicknessindependent integrated spectral weight as shown by Fig. 1b. The phonon spectral weight shows linear dependence with thickness, typical of bulk response. In this fashion we conclude that the surface transport dominates the free electron response at these thicknesses. A similar conclusion has been reached through DC studies of these films [21]. Using the Fermi velocity from ARPES [13], our observed scattering rate gives a mean free path of approximately 0.5 microns. The small scattering rate and long mean free path are remarkable considering that these films have no capping layers and the 'active area' has been directly exposed to atmosphere for several days.



FIG. 2. **a**). Color map of the electric field magnitude of the 2^{nd} THz pulse transmitted through the 16QL film in the cross polarizers configuration, for fields between -10 and 10 T applied along the film's *c* axis (positive fields are antiparallel to the THz pulse propagation direction). **b**). Real part of the conductance of Bi₂Se₃ 100 QL film for magnetic fields between 0 and 10 T at 5 K.

We now take advantage of a unique aspect of the time structure of TDTS and use the sapphire substrate itself as an optical resonator to measure both the Kerr and Faraday rotation angles in the same setup (further details in [28]). In the experimental geometry of a thin film on a dielectric substrate, after transmitting through the sample, the THz pulse partially reflects back from the substratevacuum interface and returns to the film, where it reflects and travels back to the detector. In principle this process of internal reflection inside the sapphire substrate is repeated ad infinitum. These echoes are illustrated in Figs. 1c and d, where we show a schematic of the experimental configuration and typical time domain scans of a transmitted pulse. In the presence of an external magnetic field **B**, the first time the pulse is transmitted the wave polarization may be rotated an angle φ_F , the Faraday angle (labeled "Faraday" in fig. 1c). As the 2nd pulse reflects back to the film and then reflects from it, the polarization may be rotated by an additional angle φ_K . the Kerr angle (labeled "Kerr"). The fact that TDTS measurements are resolved in time allows the separation of the different contributions to the rotation angle; this type of separation is generally not possible with standard continuous wave techniques.

In Fig. 2a we note the first evidence of an anoma-

lously large Kerr rotation induced by the surface states in applied field. The figure shows the amplitude of the transmitted electric field from the Kerr pulse with crossed polarizers P1 and P3 at $\pm 45^{\circ}$ respectively (see [28]). As we increase \mathbf{B} , we observe a rise in the transmitted amplitude as the electric field apparently undergoes a large rotation, since there should be no transmission for the crossed polarizers unless the polarization is rotated by the sample. In contrast, the Faraday rotation was always small – at the level of the experimental sensitivity ($\sim 5^{\circ}$) - because it is proportional to the optically active layer thickness, which in this case is only a few nanometers. We neglect its contribution to the total Kerr + Faradayrotation of the 2nd peak in the analysis below. We gain insight into the origin of this apparent large Kerr rotation by studying the longitudinal conductance G_{xx} in an applied magnetic field **B**. Within a conventional picture of cyclotron resonance, we expect that the spectral weight of the Drude term moves to higher frequency as the field is increased. The shift in spectral weight is also accompanied by the increase in the off-diagonal conductance G_{xy} with magnetic field; this increase is responsible for inducing rotation to polarized light impinging on the sample. One qualitatively observes such behavior in Fig. 2b with a very weak maximum at finite frequency in the real part of the conductivity for the 100QL sample at 5 K and for several magnetic fields up to 10 T, and with the rotation indicated in Fig. 2a. Films at other thicknesses show the same behavior.

We quantify the rotation angle in two different experimental configurations as described in further detail in ref. [28]. In the *collinear* polarizer configuration we expect the amplitude of the electric field to be proportional to $|\cos(\phi - \varphi)\cos(\phi)|$, and in the cross polarizer mode $|\cos(\phi - \varphi)\sin(\phi)|$, where ϕ is the angle of polarizer P2 and φ is the polarization induced by the sample. Fig. 3 shows precisely this behavior. Panel (a) shows a polar plot of the amplitude of the first and second transmitted pulses in the collinear arrangement through a bare substrate at 10 T and 5 K. It is clear that no rotation is observed and $\varphi=0$. In Figs. 3b and c, we show polar plots for both collinear and cross polarizer geometries for a 16 QL sample also at 10 T and 5 K. Both sets of data are consistent with a rotation angle of $\varphi_K = 65 \pm 3^\circ$. This is an extremely large value, both in its absolute scale and when normalized by the field and active thickness of the surface state. To the best of our knowledge, it is a world record for the Kerr rotation of a thin film.

In Fig. 4, we show the result of an experiment in a third configuration, where P2 is placed before the sample and is rotated at a high angular speed and the inand out-of-phase outputs of a lock-in amplifier give the electric field components \tilde{X} and \tilde{Y} (more details in [28]). This method allows us to perform fast scanning of the magnetic field and frequency dependence of the Kerr angle. In Fig. 4, we show as a function of magnetic field



FIG. 3. Polarizer angle ϕ dependence of the transmitted electric field amplitude normalized to its maximum value at each displayed frequency for the 16 QL sample. All displayed data were taken at 10 T and 5 K. **a**). Amplitude of the transmitted electric field through a bare sapphire (Al₂O₃) substrate in the collinear polarizers configuration. Electric field of the Faraday pulse multiplied by 2 (open circles), and normalized amplitude of the Kerr pulse (closed) are shown. **b**). Normalized amplitude for the Kerr pulse in the collinear polarizer configuration. A rotation angle of $65\pm3^{\circ}$ is observed. **c**). Same as in **b**) in the cross polarizers configuration, also shows a rotation angle of $65\pm3^{\circ}$.

and frequency the absolute value of the Kerr angle for the 32 QL sample measured at 5 K. It is clear that a similar frequency dependence exists as in Fig. 2b up to a maximum Kerr rotation of 49.5° for this sample. The variation in the values of the Kerr rotation with the films thickness, are consistent with the variations of the carriers concentration and mobility as found in ref. [21]

Such a large Kerr rotation can be qualitatively explained using the physics of cyclotron resonance, if one correctly takes into account the enhancement that occurs when the reflection off the film happens from *within* the substrate. The complex Kerr angle can be written as



FIG. 4. Absolute value of Kerr rotation $(|\varphi_K|)$ as a function of frequency and magnetic field for the 32 QL sample at 5 K. A maximum rotation of order 49.5° exists at low frequencies and finite fields.

 $\tan(\varphi_K) = \frac{2nZ_0 \mathcal{G}_{xy}}{n^2 - 1 - 2Z_0 \mathcal{G}_{xx} - Z_0^2(\mathcal{G}_{xx}^2 + \mathcal{G}_{xy}^2)}, \text{ where } n$ is the refractive index of the substrate, $Z_0 = 377$ Ohms is the vacuum impedance and \mathbf{G}_{xy} is the Hall conductance. We use parameters of the total conductance and scattering rate appropriate for our films (see [28] for details) and carriers with negative charge, and obtain from the fit an effective cyclotron mass, $m^* \sim 0.35 \text{ m}_e$. An estimate of the Dirac fermion cyclotron mass, $m^* = E_f / v_f^2$, using the Fermi energy from the carrier density $n \sim 3.3 \times 10^{13} cm^{-2}$, $E_f \sim 0.5 \text{eV}$, and velocity from ARPES [13] $v_f \sim 5 \times 10^5$ m/s, gives only a 10% difference from our value. We find that this formula reproduces the main features of the data in Fig. 4, as shown in Fig. S2 of ref. [28]. A similar analysis using bulk parameters [18, 31, 32] cannot reproduce the frequency and field dependence, which further confirms that the observed effect comes from the 2D surface states. Small values of Kerr and Faraday rotations have been found recently on single crystals of Bi₂Se₃, which have an appreciable bulk contribution [33]. Therefore, we can again conclude that the low frequency THz response is largely independent of the bulk contribution to the conductance and the observed colossal Kerr rotation is intrinsic to the surface metallic states. Its large value is due to a near cancellation of terms in the denominator (because of the large G_{xy}), and the additional enhancement when the reflection occurs inside the substrate, i.e. at the substrate-TI interface (see ref. [28]).

It has recently been found [34–36] through ARPES that non-topological 2D states induced by band-bending effects can exist at the surfaces of topological insulator crystals cleaved in ultra-high vacuum. The thickness of these states was found to be ~ 20 nm and the sheet carrier concentration of the order of 10^{13} cm⁻². This implies

that in thin films of less than 40 nm, transport could appear as effectively three dimensional. This is contrary to what we found in the THz conductivity, and to what is reported in DC transport [21]. In addition, using the effective masses reported for these carriers ($m^* \sim 0.11 \text{ m}_e$ [35]), it is not possible to reproduce the frequency and field dependence of the Kerr angle as shown in ref. [28]. We speculate that these states, if they exist in our thin films have mobilities so low that they don't make appreciable contributions to transport.

In our work, we have shown clear evidence for the robust THz response of 2D topological surface states in thin films of topological insulator Bi₂Se₃. In magnetic fields we find a colossal Kerr rotation with angles up to $\varphi_K \approx 65^\circ$ that is due to the cyclotron resonance of the surface electrons. This measured Kerr rotation is larger by an order of magnitude than rotations typically found on high mobility GaAs heterostructures [33], but it is not quantized. In order to reach a regime where the quantized topological magnetoelectric effect can be distinguished from semiclassical cyclotron resonance physics at accessible fields, it appears that the films will have to have an even larger mobility and the chemical potential must be tuned closer to the Dirac point. As the Fermi energy is tuned to the Dirac point, measurements like ours would have the clear signature of a reduction of the effective Dirac mass, and would eventually give way to the predicted signatures of *axion* electrodynamics [8]. In general, the Kerr rotation we observe represents a benchmark for the intrinsic magnetoelectric effect predicted to exist in topological insulators. We also think that the ideas and techniques demonstrated in this work will be useful in studying effects of the interaction of polarized THz radiation with other novel states of matter.

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- [1] D. Thouless, Phys. Rev. Lett. 49, 405 (1982).
- [2] K. v. Klitzing, G. Dorda, and M. Pepper, Phys. Rev. Lett. 45, 494 (1980).
- [3] L. Fu and C. L. Kane, Phys. Rev. B 76, 045302 (2007).
- [4] J. E. Moore and L. Balents, Phys. Rev. B 75, 121306

(2007).

- [5] R. Roy, Phys. Rev. B **79**, 195322 (2009).
- [6] L. Fu and C. L. Kane, Phys. Rev. Lett. 100, 096407 (2008).
- [7] A. R. Akhmerov, J. Nilsson, and C. W. J. Beenakker, Phys. Rev. Lett. **102**, 216404 (2009).
- [8] X.-L. Qi, T. L. Hughes, and S.-C. Zhang, Phys. Rev. B 78, 195424 (2008).
- [9] A. M. Essin, J. E. Moore, and D. Vanderbilt, Phys. Rev. Lett. **102**, 146805 (2009).
- [10] X. Zhang, J. Wang, and S.-C. Zhang, Phys. Rev. B 82, 245107 (2010).
- [11] D. Hsieh, D. Qian, L. Wray, Y. Xia, Y. Hor, R. Cava, and M. Hasan, Nature 452, 970 (2008).
- [12] Y. Xia, D. Qian, D. Hsieh, L. Wray, A. Pal, H. Lin, A. Bansil, D. Grauer, Y. Hor, R. Cava, et al., Nature Physics 5, 398 (2009).
- [13] D. Hsieh, Y. Xia, D. Qian, L. Wray, J. Dil, F. Meier, J. Osterwalder, L. Patthey, J. Checkelsky, N. Ong, et al., Nature 460, 1101 (2009).
- [14] Y. L. Chen, J. G. Analytis, J.-H. Chu, Z. K. Liu, S.-K. Mo, X. L. Qi, H. J. Zhang, D. H. Lu, X. Dai, Z. Fang, et al., Science **325**, 178 (2009).
- [15] P. Roushan, J. Seo, C. Parker, Y. Hor, D. Hsieh, D. Qian, A. Richardella, M. Hasan, R. Cava, and A. Yazdani, Nature 460, 1106 (2009).
- [16] Z. Alpichshev, J. Analytis, J.-H. Chu, I. Fisher, Y. Chen, Z. Chen, A. Fang, and A. Kapitulnik, Phys. Rev. Lett. 104, 016401 (2010).
- [17] D.-X. Qu, Y. S. Hor, J. Xiong, R. J. Cava, and N. P. Ong, Science **329**, 821 (2010).
- [18] N. P. Butch, K. Kirshenbaum, P. Syers, A. B. Sushkov, G. S. Jenkins, H. D. Drew, and J. P. Paglione, Phys. Rev. B 81, 241301 (2010).
- [19] J. Chen, X. Y. He, K. H. Wu, Z. Q. Ji, L. Lu, J. R. Shi, J. H. Smet, and Y. Q. Li, Phys. Rev. B 83, 241304 (2011).
- [20] J. Xiong, A. C. Petersen, D. Qu, R. J. Cava, and N. P.

Ong, Physica E (2011), in press.

- [21] N. Bansal, Y. Kim, M. Brahlek, E. Edrey, and S. Oh (2011), arxiv:1104.5709.
- [22] H. Steinberg, J.-B. Laloë, V. Fatemi, J. S. Moodera, and P. Jarillo-Herrero, Phys. Rev. B 84, 233101 (2011).
- [23] W.-K. Tse and A. H. MacDonald, Phys. Rev. Lett. 105, 057401 (2010).
- [24] W.-K. Tse and A. H. MacDonald, Phys. Rev. B 82, 161104 (2010).
- [25] J. Maciejko, X.-L. Qi, H. D. Drew, and S.-C. Zhang, Phys. Rev. Lett. **105**, 166803 (2010).
- [26] Y. Lan, S. Wan, and S.-C. Zhang, Phys. Rev. B 83, 205109 (2011).
- [27] G. Tkachov and E. M. Hankiewicz, Phys. Rev. B 84, 035405 (2011).
- [28] See Supplementary Material for further information.
- [29] A. LaForge, A. Frenzel, B. C. Pursley, T. Lin, X. Liu, J. Shi, and D. N. Basov, Phys. Rev. B 81, 125120 (2010).
- [30] A. B. Sushkov, G. S. Jenkins, D. C. Schmadel, N. P. Butch, J. Paglione, and H. D. Drew, Phys. Rev. B 82, 125110 (2010).
- [31] K. Eto, Z. Ren, A. A. Taskin, K. Segawa, and Y. Ando, Phys. Rev. B 81, 195309 (2010).
- [32] J. G. Analytis, J.-H. Chu, Y. Chen, F. Corredor, R. D. McDonald, Z. X. Shen, and I. R. Fisher, Phys. Rev. B 81, 205407 (2010).
- [33] G. S. Jenkins, A. B. Sushkov, D. C. Schmadel, N. P. Butch, P. Syers, J. Paglione, and H. D. Drew, Phys. Rev. B 82, 125120 (2010).
- [34] M. Bianchi, D. Guan, S. Bao, J. Mi, B. B. Iversen, P. D. C. King, and P. Hofmann, Nature Communications 1, 128 (2010).
- [35] P. D. C. King, R. C. Hatch, M. Bianchi, R. Ovsyannikov, C. Lupulescu, G. Landolt, B. Slomski, J. H. Dil, D. Guan, J. L. Mi, et al., Phys. Rev. Lett. **107**, 096802 (2011).
- [36] H. M. Benia, C. Lin, K. Kern, and C. R. Ast, Phys. Rev. Lett. 107, 177602 (2011).