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 wave functions [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 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 backscattering, and weak antilocalization.

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 Dirac-like 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 off-diagonal 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 $\Delta L = \alpha \theta \int dx dt \mathbf{E} \cdot \mathbf{B}$, where $\alpha$ is the fine structure constant. $\theta$ is a modulo $2\pi$ number that distinguishes between topologically nontrivial ($\theta = \pi$) and trivial insulators ($\theta = 0$). There have been several proposals of how to measure this $\theta$ 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 $\alpha$. 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 semiclassical regime of large Landau level index.

In this Letter, we report the THz response of the topological surface states in thin films of the topological insulator Bi$_2$Se$_3$. We measured thin films of several thicknesses grown on sapphire (Al$_2$O$_3$) substrates by molecular beam epitaxy. The time elapsed between the growth and THz measurements was at least 2 weeks for each sample [28]. Because of 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 300 QL, where QL means...
quintuple layer and equals 9.4 nm [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 topological surface states. The Kerr rotation measurement allows us to extract the effective mass of the 2D Dirac electrons.

In Fig. 1(a) 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 ($\approx 1.2$ 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 parametrization 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 thickness-independent integrated spectral weight as shown by Fig. 1(b). 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 $\mu$m. 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.

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 substrate-vacuum 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. 1(c) and 1(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 by an angle $\varphi_F$, the Faraday angle (labeled “Faraday” in Fig. 1(c)). As the 2nd pulse reflects back to the film and then reflects from it, the polarization may be rotated by an additional angle $\varphi_K$, 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. 2(a) we note the first evidence of an anomalously large Kerr rotation induced by the surface states in an applied field. The figure shows the amplitude of the transmitted electric field from the Kerr pulse with crossed polarizers $P_1$ and $P_3$ at \(\pm 45^\circ\) respectively (see [28]). As we increase $B$, 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 + Faraday rotation 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 a 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 the magnetic field; this increase is responsible for inducing

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**FIG. 1** (color online). (a) Real part of the conductance of four films of different thicknesses (16, 32, 64, and 100 QL, 1QL = 9.4 Å). (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 the 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 $\text{Al}_2\text{O}_3$ (top, displaced vertically for clarity), and a thin film sample (bottom) at 2 K.
rotation to polarized light impinging on the sample. One qualitatively observes such behavior in Fig. 2(b) with a very weak maximum at finite frequency in the real part of the conductivity for the 100 QL sample at 5 K and for several magnetic fields up to 10 T, and with the rotation indicated in Fig. 2(a). 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 \(\phi\) is the angle of polarizer \(P2\) and \(\varphi\) is the polarization induced by the sample. Figure 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. 3(b) and 3(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 in- and out-of-phase outputs of a lock-in amplifier give the electric field components \( \hat{X} \) and \( \hat{Y} \) (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 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. 2(b) up to a maximum Kerr rotation of 49.5° for this sample. The variations 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

\[
\tan(\varphi_K) = \frac{2nZ_0G_{xy}}{n^2 - 1 - 2Z_0G_{xx} - Z_0^2(G_{xx} + G_{yy})},
\]

where \( n \) is the refractive index of the substrate, \( Z_0 = 377 \text{ Ohms} \) is the vacuum impedance, and \( G_{xy} \) 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 \( n \sim 3.3 \times 10^{13} \text{ cm}^{-2} \), \( E_F \sim 0.5 \text{ eV} \), and velocity from ARPES [13] \( v_f \sim 5 \times 10^5 \text{ m/s} \), 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 Bi\(_2\)Se\(_3\), 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 nontopological 2D states induced by band-bending effects can exist at the surfaces of topological insulator crystals cleaved in ultrahigh vacuum. The thickness of these states was found to be \( \sim 20 \text{ nm} \) and the sheet carrier concentration of the order of \( 10^{13} \text{ cm}^{-2} \). 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 do not 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 the topological insulator Bi\(_2\)Se\(_3\). In magnetic fields we find a colossal Kerr rotation with angles up to \( \varphi_K \sim 65^\circ \) 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 magneto-electric 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.

The authors would like to thank H. D. Drew, J. Hancock, Z. Hao, G. S. Jenkins, A. Kuzmenko, A. MacDonald, N. A. Mecholsky, A. J. Pearson, O. Tchernyshyov, W-K. Tse, and Y. Wan for helpful discussions and/or correspondences.

FIG. 4 (color online). Absolute value of the Kerr rotation \(|\varphi_K|\) as a function of the 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.
Support for the measurements at JHU was provided under the auspices of the “Institute for Quantum Matter” DOE DE-FG02-08ER46544 and the Gordon and Betty Moore Foundation. The work at UB was supported by NSF MRI-R2 0959989 and NSF DMR-1006078. The work at Rutgers was supported by IAMDN of Rutgers University, NSF DMR-0845464 and ONR N000140910749.

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