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Band resolved imaging of photocurrent in a topological insulator

H. Soifer,¹ A. Gauthier,^{1,2} A. F. Kemper,³ C. R. Rotundu,¹ S.-L. Yang,^{1,2,*} H. Xiong,^{1,2} D. Lu,⁴ M. Hashimoto,⁴ P. S. Kirchmann,¹ J. A. Sobota,^{1,2,5,†} and Z.-X. Shen^{1,2,‡}

¹*Stanford Institute for Materials and Energy Sciences, SLAC National Accelerator Laboratory, 2575 Sand Hill Road, Menlo Park, California 94025, USA*

²*Geballe Laboratory for Advanced Materials, Departments of Physics and Applied Physics, Stanford University, Stanford, California 94305, USA*

³*Department of Physics, North Carolina State University, Raleigh, North Carolina 27695, USA*

⁴*Stanford Synchrotron Radiation Lightsource, SLAC National Accelerator Laboratory, 2575 Sand Hill Road, Menlo Park, California 94025, USA*

⁵*Advanced Light Source, Lawrence Berkeley National Laboratory, Berkeley, California 94720, USA*

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We study the microscopic origins of photocurrent generation in the topological insulator Bi_2Se_3 via time- and angle-resolved photoemission spectroscopy. We image the unoccupied band structure as it evolves following a circularly polarized optical excitation and observe an asymmetric electron population in momentum space, which is the spectroscopic signature of a photocurrent. By analyzing the rise times of the population we identify which occupied and unoccupied electronic states are coupled by the optical excitation. We conclude that photocurrents can only be excited via resonant optical transitions coupling to spin-orbital textured states. Our work provides a microscopic understanding of how to control photocurrents in systems with spin-orbit coupling and broken inversion symmetry.

Topological insulators are bulk insulators with robust surface conductivity mediated by topological surface states (TSS) [1–4]. Much of the excitement surrounding topological insulators stems from the rich phenomena enabled by the spin-orbital texture of the TSS, in which the electron angular momentum is locked perpendicular to its crystal momentum k [5–7]. However, accessing the intrinsic properties of the TSS via conventional transport methods has been hindered by the high residual conductivity of the bulk [8–10]. In contrast, light-based probes have proven fruitful because optical selection rules permit direct coupling to the spin-orbital degrees of freedom, and are therefore less obstructed by the bulk response [11–14]. The circular photogalvanic effect bridges optics and transport phenomena: circularly-polarized light generates photocurrents which can be measured in conventional transport configurations [15–19]. The correlation of the photocurrent direction with the light helicity has been taken as evidence that the current is carried by the TSS [17, 20–23].

Microscopically, the total photocurrent density j parallel to the surface can have contributions from all bands n , in the Brillouin zone ($-k_{\text{BZ}}, k_{\text{BZ}}$):

$$j = \sum_n j_n = -\frac{e}{\hbar} \sum_n \int_0^{k_{\text{BZ}}} [\rho_n(+k) - \rho_n(-k)] \frac{\partial \varepsilon_n}{\partial k} dk, \quad (1)$$

which shows that a momentum asymmetry in the electron density $\rho_n(k) \neq \rho_n(-k)$ is required for generating current [24]. Individual band contributions j_n can be directly measured using time- and angle-resolved photoemission spectroscopy (trARPES) which resolves the photoexcited populations along the electronic band disper-

sions $\varepsilon_n(k)$. Early trARPES works have demonstrated that circularly-polarized light generates an asymmetric electron distribution in the TSS [14, 25–28]. The question remains whether this phenomenon is exclusive to topological states or can be generalized to other spin-orbit coupled states. It is therefore important to characterize the photoexcited carriers over a broader energy range, including photoholes injected below the Fermi level [19]. A detailed understanding of optical coupling to spin-orbital degrees of freedom is required for a microscopic picture of photocurrent generation in topological insulators.

In this Letter, we use trARPES to study the momentum-asymmetric optical transitions responsible for photocurrent generation in Bi_2Se_3 . We excite a wide range of states, and analyze the population dynamics to simultaneously yield the momentum distributions of electrons and holes. Our results vividly highlight the key role of resonant conditions in generating asymmetric distributions using circularly-polarized light. Moreover, we find that population asymmetries are not exclusive to topological states, and broadly reexamine the ingredients required for generating such asymmetries.

In our trARPES experiments we used circularly-polarized 3.02 eV pump photons to populate unoccupied bands above the Fermi level, E_F , and s -polarized 6 eV probe pulses to photoemit electrons at a variable delay Δt . The s -polarization of the probe was chosen to eliminate additional photoexcitation channels associated with an image potential state [29], which spectrally overlaps the bands measured here. The overall time resolution of 100 fs is extracted from cross-correlations of pump and probe pulses. $\Delta t = 0$ refers to both pulses overlapping in time. The incidence plane of light is along the mir-

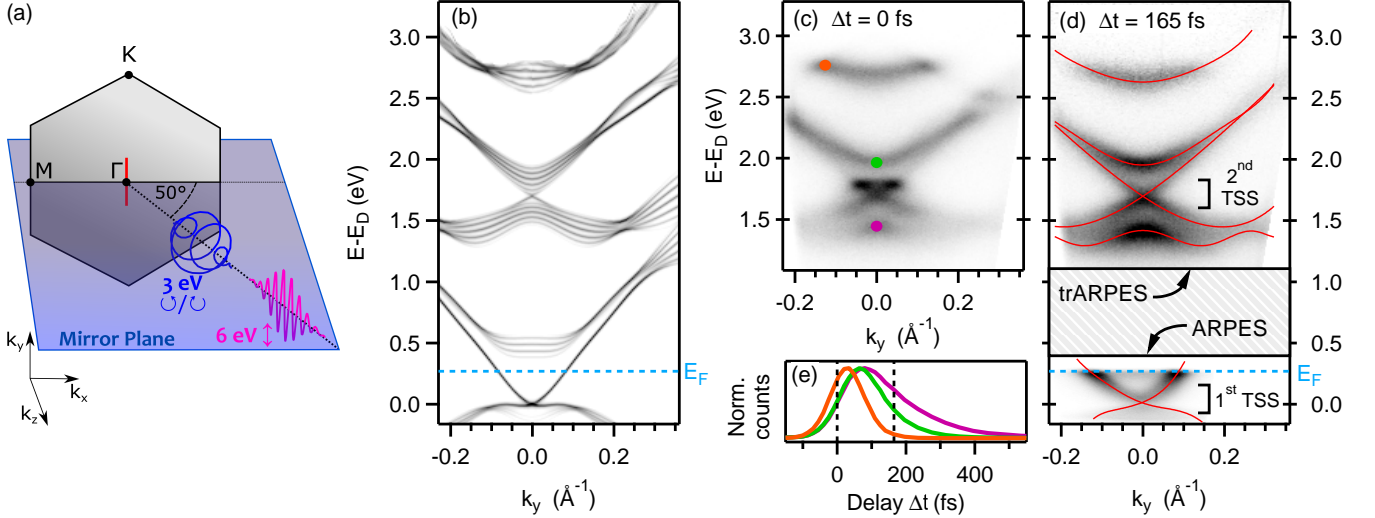


FIG. 1. **Unoccupied band structure of Bi_2Se_3 .** (a) Sketch of the experimental geometry. The cut direction is shown in red. (b) DFT calculation of the band structure. Energies are referenced to the 1st Dirac point. The blue dashed line marks the Fermi level ($E_F - E_D = 270$ meV). (c) and (d) Population of unoccupied bands from trARPES at two delays (average of population excited by σ_+ and σ_-). The bottom panel of (d) shows the the first Dirac cone, measured with 6 eV ARPES. The intensities in the top panel of (d) were rescaled exponentially as a function of energy for better visibility. Red lines are guides to the eye that trace the bands. (e) Transient intensity for 3 spectral regions, as marked by dots of the same color in (c). Dashed vertical lines mark the delays at which the trARPES spectra in (c) and (d) were taken.

ror plane of the sample, see Fig. 1(a). Photoelectrons are collected in a hemispherical analyzer along the Γ -K direction [30]. Additional ARPES data of the equilibrium band structure was recorded at beamline 5-2 of the Stanford Synchrotron Radiation Lightsource using 35 eV photons. We perform density functional theory (DFT) calculations using the VASP [31–36] package using PAW PBE [36–39] GGA-type pseudopotentials. The calculation used a 7 quintuple layer slab geometry based on the experimental coordinates and with a 30 Å vacuum layer. We used a $15 \times 15 \times 1$ momentum grid centered on Γ , and a wave function cutoff of 250 eV. Spin-orbit coupling was included through a non-self-consistent calculation after a converged density was obtained. The energy scale was renormalized by a factor of 1.11 to match the experimental energy difference between the 1st and 2nd Dirac points of 1.7 eV.

Figure 1(b) shows the calculated band structure of Bi_2Se_3 with the cone-shaped dispersion of the 1st TSS visible at the bottom [2]. Throughout the paper we reference energies to the 1st Dirac point, E_D . The blue dashed line marks E_F , which is 270 meV above E_D in our experiments. A second cone-shaped dispersion at 1.7 eV corresponds to the 2nd TSS above E_F [40, 41].

The population in the unoccupied bands measured at $\Delta t = 0$ and 165 fs is shown in Fig. 1(c) and (d) respectively. The spectrum at $\Delta t = 165$ fs clearly corresponds to the calculated bands. Red lines are guides to the eye that trace the band dispersions and are used through-

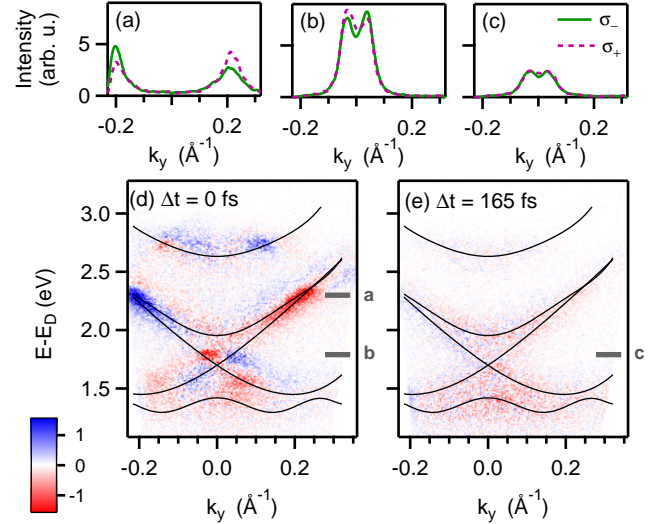


FIG. 2. **Helicity Dependent Population Asymmetry.** (a)-(c) Momentum Distribution Curves (MDCs) of the unoccupied band structure, excited with σ_+ (dashed-purple) and σ_- (solid-green) polarized pulses. (a) and (b) were taken at $\Delta t = 0$, and (c) at $\Delta t = 165$ fs, at the energies marked by the short gray lines in (d)-(e). (d),(e) Asymmetry image: difference between the populations of the unoccupied bands when excited by σ_- and σ_+ polarized pulses, taken at $\Delta t = 0$ and 165 fs respectively. Black lines are guides to the eye that follow the dispersions of the unoccupied bands.

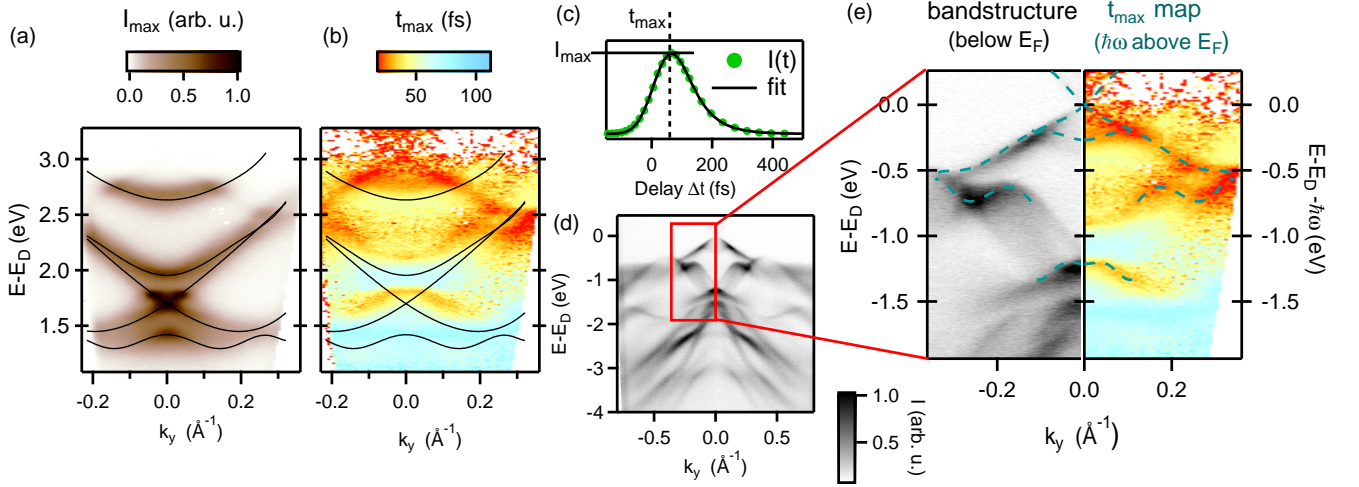


FIG. 3. **Time Mapping: E and k resolved analysis of the population dynamics.** (a) $I_{\max}(k, E)$: Maximum intensity from fits of each energy-momentum bin. (b) $t_{\max}(k, E)$: delay at which the intensity is maximal for each bin. Solid lines in (a) and (b) represent the dispersions of the unoccupied bands. Bins where the fit did not converge appear white in both images. (c) Prototypical transient population (green dots, same as green curve in Fig. 1(e)), and fit (solid black). I_{\max} and t_{\max} are shown. (d) Occupied band structure from ARPES, using s-polarized 35 eV photons. (e) Comparison between t_{\max} map (right) and the occupied states one pump photon energy below ($\hbar\omega = 3.02$ eV) (left). The dashed lines are guides to the eye based on the early-rise times in the t_{\max} map.

out the paper as reference to the unoccupied bands. The first TSS is shown at the bottom of Fig. 1(d), measured via 6 eV ARPES. The spectral intensity distribution at $\Delta t = 0$ is different, in particular the increased intensity around the 2nd TSS at $E - E_D = 1.8$ eV. The intensity distribution at $\Delta t = 0$ is determined by the joint density of states between the initial and final states of the electronic excitation [41, 42], whereas the distribution at later delays, after the electrons have scattered, reflects more directly the unoccupied band structure. The normalized, transient intensity in three different spectral regions is shown in Fig. 1(e).

Next, we explore how the momentum distribution of the photoexcited electrons depends on the light helicity. Figure 2(a-c) displays momentum distribution curves (MDCs) of the unoccupied states at two energies and two delays, when excited by σ_+ (dashed purple) and σ_- (solid green) circular polarizations. At $\Delta t = 0$ and $E - E_D = 2.3$ eV [Fig. 2(a)] circular polarization excites an asymmetric population distribution, indicating that at this energy there is a net flow of electrons in one direction. This asymmetry reverses direction when switching the light helicity. Similarly, in Fig. 2(b) ($\Delta t = 0$, $E - E_D = 1.8$ eV) we observe the signature of a helicity dependent asymmetry, whereas the MDCs at the same energy and $\Delta t = 165$ fs display symmetric populations [Fig. 2(c)]. We note that this experiment is distinguished from conventional Circular Dichroism (CD) in equilibrium ARPES [43–46] which is typically used as a spectroscopic tool for determining band character. In our

experiment the polarization of the pump and not the photoemission probe is modified, allowing us to study non-equilibrium optically-induced population asymmetries above E_F [47].

The difference plots between the σ_+ and σ_- excited spectra are shown in Fig. 2(d)-(e), where the red/blue colors signify a helicity-dependent population asymmetry. MDCs of the difference signal are shown in the Supplementary Material [47]. The difference map at $\Delta t = 0$ in Fig. 2(d) reveals a complex pattern of band- and momentum-dependent asymmetries which decay within 165 fs [Fig. 2(e)]. These results show that multiple bands contribute to the photocurrent, and can contribute with opposite directions.

It is important to note that the photo-excited electrons do not represent all of the charge carriers contributing to the current, the remainder being the corresponding holes injected below E_F . Though we do not have direct access to the hole population, the conservation of energy and momentum in optical transitions links the electron and hole distributions. Therefore, we can infer the hole distribution by determining which part of the excited population corresponds to direct optical transitions, and which to filling via secondary scattering processes. These two processes can be distinguished by their timescales, with direct optical transitions causing a faster population rise than secondary scattering [48, 49].

We analyze the rise times of the photoexcited population to identify which parts of the unoccupied bands are populated via direct optical transitions. This time-

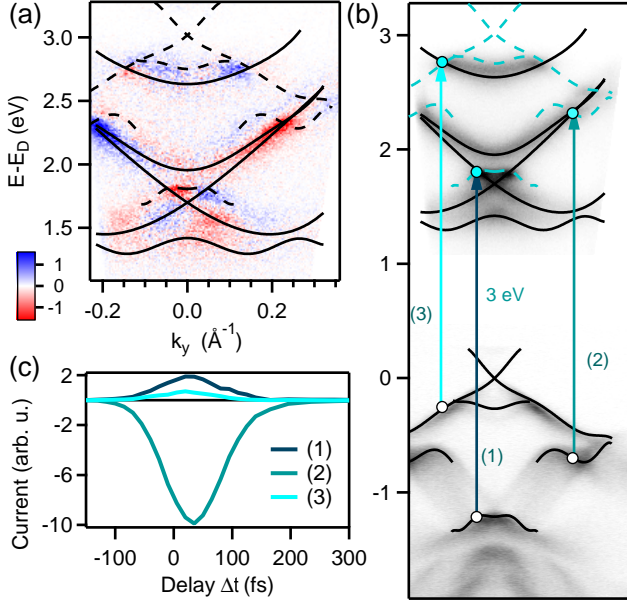


FIG. 4. **Resonant transitions.** (a) Asymmetry image: difference between the populations of the unoccupied bands when excited by σ_- and σ_+ polarized pump, at $\Delta t = 0$ fs. [same as Fig. 2(d)]. The solid lines follow the dispersions of the unoccupied bands. Dashed lines follow the early-rise times from the t_{\max} map, marking the dispersions of the initial states $\hbar\omega = 3.02$ eV below. (b) Schematic visualization of the resonant optical transitions between occupied and unoccupied states of Bi_2Se_3 with a 3 eV excitation. The solid black lines mark the bands, and the blue-dashed lines mark the initial states upshifted by the excitation. (c) Time dependent contributions to the photocurrent from each of the resonant optical transitions in (b) according to Eq. 1.

mapping analysis is based on energy- and momentum-resolved fits of the intensity dynamics: for each spectral bin ($0.007 \text{ \AA}^{-1} \times 12 \text{ meV}$), we fit the transient signal with a Gaussian convoluted with an exponential decay to model the filling and subsequent decay of the optically excited population. A typical fit is shown in Fig. 3(c). We focus on two fitted quantities: the maximum intensity, I_{\max} , and the delay at which the maximum intensity is reached, t_{\max} , which are plotted in Fig. 3(a) and (b) respectively. Other fitted quantities are shown in [47]. The intensity I_{\max} bears a close similarity to the spectral intensity at $\Delta t = 0$ [Fig. 1(b)], which however is not the case for the t_{\max} map. It can be separated into two types of regions: where the intensity rises early ($t_{\max} \lesssim 50$ fs, yellow-red regions), and regions with later rise times (blue-white). This observation underlies our approach to distinguish between direct optical transitions and secondary scattering using population rise times.

Though the early-rise regions do not follow the dispersion of the unoccupied bands (solid black lines), we find that they reflect the initial states of the excitation (occupied states below E_F). To demonstrate this link, we mea-

sure the occupied band structure of Bi_2Se_3 via ARPES as shown in Fig. 3(d). The t_{\max} map agrees remarkably well with the occupied bands, when shifted down by one pump photon energy ($\hbar\omega = 3.02$ eV), see Fig. 3(e). Guides to the eye (dashed blue) trace the early-rise features in the t_{\max} map and are reflected around $k = 0$ for an easier comparison. Notably, for every early-rise feature in the time-map there is a corresponding band in the initial state spectrum, while the opposite is not true - there are occupied bands without a corresponding early-rise feature in the t_{\max} map (see e.g., electronic band at $E - E_D = -1.5$ eV). We conclude that the rise-time mapping reveals only the occupied states which participate in the optical excitation process.

This mapping of initial states allows us to characterize the conditions which result in asymmetric populations. Figure 4(a) shows the asymmetry image at $\Delta t = 0$, overlaid with the dispersions of the excited states (solid lines), as well as the initial states obtained from our time-mapping analysis (shifted up by $\hbar\omega$, dashed lines). These three elements together reveal that the regions characterized by the strongest electron population asymmetry are those where the initial and excited states overlap in momentum and energy. Only resonant optical transitions lead to asymmetric populations and consequently contribute to photocurrent generation. Scattering processes subsequently fill non-resonant states. Intra-band small- k scattering processes conserve the population asymmetry and spread it to a small extent beyond the resonant energies (see e.g. asymmetry at $E - E_D \sim 1.6$ eV), whereas inter-band and $k \rightarrow -k$ scattering lead to a loss of asymmetry, in agreement with the symmetric populations observed at later delays [Fig. 2(e)].

We can use this information to describe the contribution of the photoholes to the current. Since the measured asymmetric population is associated with resonant energy- and momentum-conserving direct optical transitions, it also reflects the asymmetry of the photohole distribution, shifted by the pump photon energy. Despite having similar populations, the magnitudes of the electron and hole contributions to the photocurrent are not equal, since they originate in bands with different velocities $\frac{\partial \epsilon_n}{\partial k}$ (see Eq. 1). The power of our approach is that we directly extract the band velocities of the electrons from the trARPES spectrum at later delays and of the holes from the dispersions via time-mapping. For example, the contribution to the photocurrent from the resonant optical transition into the second TSS [arrow (1) in Fig. 4(b), for σ_+ excitation] is due to electrons populating the left, negatively dispersing, branch of the unoccupied Dirac cone, whereas holes reside in a weakly dispersing region of the bulk band structure and contribute negligibly to the photocurrent. We therefore expect that the photocurrent contribution from this particular transition is dominated by electrons in the unoccupied TSS flowing in the $-y$ direction. Figure 4(c) shows the time-

dependent contribution to the current from each of the three resonant optical transitions calculated by Eq. 1 [47].

Not each asymmetric distribution is associated with a TSS, as evidenced by transition (2). This observation motivates us to examine the fundamental mechanism for driving k -asymmetric optical excitations. Circularly polarized light couples to orbital degrees of freedom by incrementing or decrementing the orbital angular momentum m_L along the quantization axis defined by the light propagation vector. Though spin-orbit coupling entangles m_L with the electron momentum k [6, 7], this alone is insufficient for enabling helicity-dependent optical transitions because $\pm m_L$ are degenerate in the presence of inversion symmetry. This degeneracy is lifted at the surface [50, 51], while time-reversal symmetry ensures that states at $\pm k$ have opposite signs of m_L . Thus, the key ingredients for enabling k -asymmetric optical excitations are spin-orbit coupling and inversion-symmetry breaking. Though transitions (1) and (3) involve a TSS, it is not obvious that transition (2) involves states of broken inversion symmetry. We hypothesize that the final state includes a topologically trivial surface resonance with a spin-orbital texture oriented opposite to that of the TSS [47], as was recently observed for states accompanying the first TSS [52, 53].

This detailed view of optical transitions can help us understand photocurrent in other scenarios as well. For example, photo-population of the second TSS in Bi_2Se_3 was observed with 1.5 eV excitation [54], and might therefore contribute to 1.5 eV-driven photocurrent [16–19, 21]. Similarly, surface resonance states are generally related to the strong spin-orbit coupling in TIs [53, 55, 56] and could contribute to photocurrent in other TI materials [19]. Since both topological and trivial states participate in asymmetric optical excitations, this phenomenon should be abundant in many families of spin-orbit coupled materials with broken inversion symmetry. Indeed, the same concepts underlie valley-dependent optical transitions in transition metal dichalcogenides [57], as well as proposals to manipulate the chirality of Weyl fermions in semimetals [58, 59]. In fact, the photocurrent associated with these transitions in Weyl semimetals is of central importance, since it is currently the only quantized observable predicted for this class of materials [60].

This work establishes trARPES as a powerful toolset for identifying the optical transitions which form the microscopic basis for photocurrent generation via the circular photogalvanic effect. We account for contributions from photoelectrons and photoholes, including both topological and trivial states. An important finding is that different bands contribute to the net current with opposing signs. Therefore, there are promising opportunities to optimize the magnitude of the photocurrent for applications by exploring resonance conditions achieved by other photon energies. We also point out that we excite photocurrents in the 2nd TSS using eas-

ily generated light in the visible range. These results lay the foundation for future measurements of the spin-polarization of the excited carriers, which would be a milestone toward establishing complete control over the degrees of freedom which make topological materials attractive for spintronic applications. The concepts exemplified here apply universally to materials with spin-orbit coupling and inversion-symmetry breaking, and are therefore likely to find relevance for optical investigations of transition-metal dichalcogenides and Weyl semimetals.

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* Present address: Kavli Institute at Cornell for Nanoscale Science, Laboratory of Atomic and Solid State Physics, Department of Physics, and Department of Materials Science and Engineering, Cornell University, Ithaca, NY 14853, USA

† sobota@stanford.edu

‡ zxshen@stanford.edu

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