

RAPID COMMUNICATION

Perpendicular magnetization switching by large spin–orbit torques from sputtered $Bi_2Te_{3-}^*$

To cite this article: Zhenyi Zheng et al 2020 Chinese Phys. B 29 078505

View the article online for updates and enhancements.

You may also like

- <u>Spin-orbit torque induced magnetization</u> <u>switching in Pt/Co/Ta structures with</u> <u>perpendicular magnetic anisotropy</u> Jijun Yun, Dong Li, Baoshan Cui et al.
- Effect of seed and interlayer Pt thickness on spin-orbit torque efficiency in Co/Pt multilayer with perpendicular magnetic anisotropy
 Gerard (assent Lim, Weillang Gan and
- Gerard Joseph Lim, Weiliang Gan and Wen Siang Lew
- <u>Annealing effect on the magneto-electric</u> properties of SOT-MTJs from micro to nano-sized dimensions Peiyue Yu, Lei Zhao, Jianfeng Gao et al.

This content was downloaded from IP address 3.17.184.90 on 06/05/2024 at 15:07

RAPID COMMUNICATION

Perpendicular magnetization switching by large spin–orbit torques from sputtered Bi₂Te₃*

Zhenyi Zheng(郑臻益)^{1,2,3}, Yue Zhang(张悦)^{1,†}, Daoqian Zhu(朱道乾)¹, Kun Zhang(张昆)¹, Xueqiang Feng(冯学强)¹, Yu He(何宇)¹, Lei Chen(陈磊)¹, Zhizhong Zhang(张志仲)^{1,2}, Dijun Liu(刘迪军)², Youguang Zhang(张有光)^{1,2}, Pedram Khalili Amiri³, and Weisheng Zhao(赵巍胜)^{1,‡}

¹ Fert Beijing Research Institute, BDBC, School of Microelectronics, Beihang University, Beijing 100191, China
 ² School of Electronics and Information Engineering, Beihang University, Beijing 100191, China
 ³ Department of Electrical and Computer Engineering, Northwestern University, Evanston, Illinois 60208, USA

(Received 2 April 2020; revised manuscript received 13 May 2020; accepted manuscript online 19 May 2020)

Spin–orbit torque (SOT) effect is considered as an efficient way to switch the magnetization and can inspire various high-performance spintronic devices. Recently, topological insulators (TIs) have gained extensive attention, as they are demonstrated to maintain a large effective spin Hall angle (θ_{SH}^{eff}), even at room temperature. However, molecular beam epitaxy (MBE), as a precise deposition method, is required to guarantee favorable surface states of TIs, which hinders the prospect of TIs towards industrial application. In this paper, we demonstrate that Bi₂Te₃ films grown by magnetron sputtering can provide a notable SOT effect in the heterostructure with perpendicular magnetic anisotropy CoTb ferrimagnetic alloy. By harmonic Hall measurement, a high SOT efficiency ($8.7 \pm 0.9 \text{ Oe}/(10^9 \text{ A/m}^2)$) and a large θ_{SH}^{eff} (3.3 ± 0.3) are obtained at room temperature. Besides, we also observe an ultra-low critical switching current density ($9.7 \times 10^9 \text{ A/m}^2$). Moreover, the low-power characteristic of the sputtered Bi₂Te₃ film is investigated by drawing a comparison with different sputtered SOT sources. Our work may provide an alternative to leverage chalcogenides as a realistic and efficient SOT source in future spintronic devices.

Keywords: spin-orbit torque, sputtered topological insulator, ferrimagnet, magnetization switching

PACS: 85.70.-w, 75.60.Jk, 75.70.Tj

1. Introduction

Physical mechanisms leading to controllable and stable magnetization switching are considered as the core of spintronic devices with high performance.^[1-4] In recent years, researchers have found that, by applying a current through the cross-section of a non-magnetic metal (NM)/ferromagnetic metal (FM) bilayer structure, the spin–orbit interactions (SOIs) can provide strong spin torques to switch the adjacent magnetic layer via the spin–orbit torque (SOT) effect.^[5-7] This method serves as a more efficient way to switch the magnetization than the spin-transfer torque (STT) effect and is expected to inspire a variety of high-speed and multifunctional spintronic devices.^[8–13]

The most common spin generators in NM/FM bilayers are heavy metals (HMs), such as Ta, W, Pt, Hf, since bulk SOIs in heavier elements are much more significant.^[14,15] However, the effective spin Hall angle θ_{SH}^{eff} of HM is limited, leading to a relatively high critical magnetization switching current density j_{sw} on the order of 10^{11} – 10^{12} A/m.^[5–7,15] Furthermore, with regard to certain high-performance bulk **DOI:** 10.1088/1674-1056/ab9439

magnets, such as ferrimagnetic alloys,^[8] Heusler alloys,^[16] magnetic insulators,^[17] featuring a larger thickness, j_{sw} is expected to be further increased to get over the augmenting anisotropic energy barrier. These facts seriously hinder the realization of low power consumption. Therefore, numerous studies have been conducted to investigate novel NM materials with larger θ_{SH}^{eff} such as topological insulators (TIs),^[18-20] two-dimensional electron gas (2DEG),^[21,22] oxidized HMs,^[23,24] and so on. Among those materials, TIs (including Bi₂Te₃, Bi₂Se₃, Bi_{0.9}Sb_{0.1}, (BiSb)₂Te₃) exhibit great potential for application, thanks to the large SOTs induced by the spin-momentum locking on their surfaces at room temperature. For instance, a several-nm-thick magnetic layer can be switched by SOTs from Bi_2Se_3 or $Bi_{0.9}Sb_{0.1}$ with a j_{sw} of 10^{10} A/m,^[19,20,25] while with (BiSb)₂Te₃, j_{sw} could even be reduced to the order of 10⁹ A/m.^[26] Note that all those reported TI samples are grown on single-crystal substrates with an ultra-precise growing method of molecular beam epitaxy (MBE) to form excellent topological surface states (TSS). This method is acceptable at laboratory level, but incompatible with

[†]Corresponding author. E-mail: yz@buaa.edu.cn

[‡]Corresponding author. E-mail: weisheng.zhao@buaa.edu.cn

Corresponding aution. E-main. weisneng.znao@buaa.edu.en

http://iopscience.iop.org/cpb http://cpb.iphy.ac.cn

^{*}Project supported by the National Natural Science Foundation of China (Grant Nos. 61971024 and 51901008), Young Elite Scientist Sponsorship Program by CAST (Grant No. 2017QNRC001), the International Mobility Project (Grant No. B16001), and National Key Technology Program of China (Grant No. 2017ZX01032101). P.K.A. acknowledges support by a grant from the National Science Foundation, Division of Electrical, Communications and Cyber Systems (NSF ECCS-1853879).

 $[\]ensuremath{\mathbb{O}}$ 2020 Chinese Physical Society and IOP Publishing Ltd

industrial memory or logic fabrication process. To resolve this obstacle, sputtered Bi₂Se₃ films on oxidized silicon substrate are recently reported to possess a great θ_{SH}^{eff} and can efficiently switch the adjacent ferromagnet at room temperature.^[27,28] Magnetron sputtering is commonly used in semiconductor industry and therefore offers a possibility to introduce typical TI-related chalcogenides with enhanced SOT into large-scale application. Nevertheless, few similar materials are reported so far to maintain large θ_{SH}^{eff} through this method.

In this paper, we report that Bi₂Te₃ films sputtered on oxidized silicon substrate can exhibit a remarkable SOT effect. By applying harmonic Hall measurement in Bi₂Te₃/CoTb bilayer, θ_{SH}^{eff} in this system is determined as large as 3.3 ± 0.3 at room temperature. Then, current-induced switching of a 6-nm-thick perpendicular magnetic anisotropy (PMA) CoTb ferrimagnetic alloy is also realized in the mentioned structure. Compared with traditional HMs, j_{sw} as well as the power consumption to switch the magnetic layer can be reduced by more than one order.

2. Method

The samples in our work are deposited on thermally oxidized Si substrates using DC or RF magnetron sputtering under a base pressure better than 2×10^{-8} Torr. We first deposited a series of substrate/MgO (2 nm)/Bi2Te3 (t nm)/MgO (1 nm) stacks with t varying from 5 nm to 40 nm for the fourpoint resistivity measurements. Figure 1(a) shows that the resistivity of the sputtered Bi2Te3 has a downward trend versus its thickness t. This trend is similar to those in previously reported TI materials^[20] and can be explained by the combined effect of surface conduct and bulk conduct properties in Bi₂Te₃.^[29] To guarantee a relatively good θ_{SH}^{eff} as well as enough passing current through the cross-section, 8-nm-thick Bi₂Te₃ layer with a resistivity of 3011 $\mu\Omega$ cm is chosen for the SOT experiments in our work. This Bi₂Te₃ layer should have a polycrystalline structure, and the roughness characterized by atomic force microscope (AFM) is shown in the inset of Fig. 1(a). The root mean square value of the surface roughness is 0.35 nm, which is smooth enough for the following heterostructure deposition.



Fig. 1. (a) Thickness-dependent resistivity change measured by four-point method in MgO/Bi₂Te₃/MgO trilayers. Inset is an AFM image of Bi₂Te₃ (8 nm) sample. (b) Stack schematic of the investigated Bi₂Te₃/CoTb heterostructure. (c) Dependence of the magnetization direction of CoTb layer on *x*. When *x* is in the range of 6-8 nm, CoTb layers show excellent PMA properties.

In addition, a ferrimagnetic rare-earth-transition-metal alloy CoTb, in which Co and Tb sub-lattices are coupled antiferromagnetically, is used as the adjacent magnetic layer for two reasons: 1) CoTb alloy possesses robust bulk PMA within a large thickness range, and bulk PMA magnets present better switching performance and thermal stability in real device applications;^[19,30] 2) CoTb alloy has a relatively high resistivity (206 μ Ω·cm), allowing more current to pass through the Bi₂Te₃ layer. Here, CoTb is deposited by the co-sputtering of Co and Tb targets. By tuning the chemical composition or temperature, CoTb could approach angular momentum compensation point or magnetization compensation point, and exhibits plenty of charming magnetic properties.^[30–32] However, few evidences yet support that CoTb approaching compensation point could contribute to increase θ_{SH}^{eff} in the whole system. Therefore, we use a Co-rich chemical composition, i.e., Co_{0.83}Tb_{0.17}, throughout our whole work, which is estimated by the independent deposition speed of Co and Tb. As

shown in Fig. 1(b), a series of substrate/MgO (2 nm)/Bi₂Te₃ (8 nm)/CoTb (x nm)/Ta (1.5 nm) stacks are deposited. It can be found that the capping Ta layers are fully oxidized and not involved in the SOT switching process.^[33] Vibration sample magnetometry is utilized to determine the magnetic anisotropies of all the samples. As illustrated in Fig. 1(c), when x ranges from 3 nm to 12 nm, the magnetization direction of the CoTb varies from in-plane to out-of-plane, and tends to go slightly back to the in-plane state when x reaches 12 nm.

For electrical measurements, we choose the 6-nm-thick CoTb sample, since it possesses good PMA when the magnetic layer remains relatively thin. The net magnetization M_s of this 6-nm-thick CoTb layer is 215 emu/cc. By typical lithography and ion beam etching, the sample is fabricated into 5-µm-width Hall bar device. Keithley 6221 current source and 2182 voltage-meter are used to implement AHE and SOT switching experiments. To measure the SOT efficiency, harmonics Hall measurement is then implemented by injecting an AC current of 133.33 Hz along *x* axis. Two SR830 lock-in amplifiers are used to detect the harmonic signals.

3. Result and discussion

Figure 2(a) shows the fabricated device with gold electrodes and the basic electrical measurement schematic. The

anomalous Hall effect (AHE) resistance versus the applied out-of-plane magnetic field is plotted in Fig. 2(b). Squared loop and similar coercive field confirm that the stack maintains its as-deposited magnetic properties. Figure 2(c) plots the curves of the first and the second harmonic voltages, i.e., $R_{1\omega}$ and $R_{x(y),2\omega}$ versus the in-plane magnetic field $H_{x(y)}$. A ratio coefficient $B_{x(y)}$ could be firstly calculated with the respective curvature of $R_{1\omega}$ and the slope of $V_{x(y),2\omega}$ ^[34,35]

$$B_{x(y)} = \left(\frac{\partial R_{x(y),2\omega}}{\partial H_{x(y)}}\right) \times \left(\frac{\partial^2 R_{1\omega}}{\partial H_{x(y)}^2}\right)^{-1}.$$
 (1)

Then, the damping-like H_{DL} and field-like effective field H_{FL} can be determined by the following expression:^[34,35]

$$H_{\rm DL(FL)} = -2 \times \frac{B_{x(y)} \pm 2\xi B_{y(x)}}{1 - 4\xi^2},$$
(2)

where ξ is the ratio of planar Hall resistance R_{PHE} and AHE resistance R_{AHE} . The measured R_{PHE} in our samples is about 0.06 Ω , which is much smaller than R_{AHE} and thus can be negligible. In this occasion, $H_{\text{DL}(\text{FL})} \approx -2B_{x(y)}$. Moreover, it also should be noticed that $R_{x,2\omega}$ induced by the field along *x* axis is much larger than $R_{y,2\omega}$ induced by the field along *y* axis, even if $R_{y,2\omega}$ is amplified by 3 times in Fig. 2(c). This phenomenon indicates that the induced H_{FL} is negligible compared to the induced H_{DL} ; for this reason, we concentrate on the dampinglike SOT effect in the following discussion.



Fig. 2. (a) Image of 5- μ m-width Hall bar device with electrical measurement setup. (b) Anomalous Hall resistance curve of Bi₂Te₃ (8 nm)/CoTb (6 nm) sample. (c) Harmonics resistance curve with applied in-plane magnetic field. H_x -induced damping-like signal (red points) is much larger than H_y -induced field-like signals (blue circles, amplified by 3 times). (d) Effective damping-like field as a linear function of current density in Bi₂Te₃ layer.

Figure 2(d) shows the calculated H_{DL} as a function of applied charge current density j_c in the Bi₂Te₃ layer. Here, j_c is calculated by a parallel circuit model of Bi₂Te₃/CoTb bilayer, which can be written as

$$j_{\rm c} = \frac{\rho_{\rm CoTb}}{\rho_{\rm Bi_2Te_3} t_{\rm CoTb} + \rho_{\rm CoTb} t_{\rm Bi_2Te_3}} \frac{I}{w_{\rm HallBar} t_{\rm Bi_2Te_3}},\qquad(3)$$

where ρ_{CoTb} , $\rho_{\text{Bi}_2\text{Te}_3}$, t_{CoTb} , $t_{\text{Bi}_2\text{Te}_3}$, w_{HallBar} and I represent the resistivity of CoTb, resistivity of Bi2Te3, thickness of CoTb, thickness of Bi₂Te₃, width of Hall bar, and injected current, respectively. H_{DL} augments linearly with the increasing current density, revealing that Joule heating effect is negligible in the measured current density range.^[32] In the investigated Bi₂Te₃/CoTb sample, the determined current-induced SOT efficiency χ (defined by $H_{\rm DL}/j_{\rm c}$) is 8.7 ± 0.9 Oe/(10⁹ A/m²). Although the applied CoTb is far from its magnetic compensation point, the obtained value is surprisingly high. For comparison, χ in W/CoTb/AlO_x structure in our previous work is about 0.4 ± 0.04 Oe/(10⁹ A/m²),^[8] while χ in a conventional Ta/CoFeB/MgO structure is 0.2–0.9 Oe/(10⁹ A/m²).^[36,37] This outstanding χ indicates that strong SOTs are expected in the sputtered Bi_2Te_3 layer. We also calculate the θ_{SH}^{eff} in this system by the following expression:^[30–33]

$$\theta_{\rm SH}^{\rm eff} = \frac{2e}{\hbar} \frac{M_{\rm s} t_{\rm CoTb} H_{\rm eff}}{J_{\rm c}},\tag{4}$$

where t_{CoTb} is thickness of CoTb. The obtained $\theta_{\text{SH}}^{\text{eff}} \approx 3.3 \pm$

0.3, which exceeds 100% and is competitive with the results in MBE-grown TIs.^[18–20] In addition, it is worth mentioning that there might be spin loss at the interface between Bi_2Te_3 and CoTb. Given this fact, even a larger intrinsic spin Hall angle could be expected in our sputtered Bi_2Te_3 films.

Current-induced magnetization switching at room temperature is demonstrated as well and plotted in Figs. 3(a) and 3(d). Here, a series of 0.1-ms-width current pulses are applied to switch the magnet, while a 400 Oe in-plane magnetic field is used to break the magnetization precession symmetry. The j_{sw} to switch 6-nm-thick CoTb is about 9.7×10^9 A/m². This value is one or two orders lower than that in typical HM-based heterostructure,^[5–8] confirming again the existence of strong SOTs in the sputtered Bi₂Te₃. When the direction of the applied in-plane field is reversed, the switching polarity of the curve changes as expected in the theoretical SOT switching framework. We also notice that the variation of SOT-driven $R_{\rm AHE}$ (~ 1.6 Ω) is smaller than that of magnetic-field-driven $R_{\rm AHE}$ (~ 2.1 Ω), revealing a partial switching phenomenon (76% magnetization). This difference is also observed in previous switching results with MBE-grown TIs and can be explained by the Joule-heating-caused demagnetization in CoTb layer.^[26] We are convinced that reducing shunting current in CoTb layer by reasonable device design could efficiently eliminate the problem.



Fig. 3. Current-induced switching curves of Bi₂Te₃/CoTb under an in-plane magnetic field $H_x = 400$ Oe (a) and -400 Oe (e). (b)–(c) The AHE curves of Pt/CoTb, Ta/CoTb, and W/CoTb samples, respectively. (f)–(h) The current-induced switching curves of Pt/CoTb, Ta/CoTb, and W/CoTb samples under $H_x = 400$ Oe. The different switching polarities reveal different spin Hall angle signs.

To better evaluate the SOT performance of the sputtered Bi_2Te_3 sample, some control samples based on conventional sputtered SOT sources (Pt, Ta, W) are also studied with a stack structure of HM (5 nm)/CoTb (4 nm). While maintaining the same chemical composition and M_s , the thickness of CoTb is reduced in these samples, since SOTs originating from common HMs can hardly switch magnets with large thickness. As shown in Figs. 3(b)–3(d), all the control samples have squared R_{AHE} loops versus perpendicular mag-

netic field, indicating that they possess good PMA properties. Besides, because of the different shunting effects of Bi₂Te₃, Pt, Ta, and W, R_{AHE} in the four samples are not consistent. Figures 3(f)–3(h) illustrate the current-induced magnetization switching curves of Pt/CoTb, Ta/CoTb, and W/CoTb samples. In these experiments, the applied in-plane magnetic field is fixed to +400 Oe to guarantee a more reasonable comparison. We observe similar R_{AHE} in current-induced switching and field-induced switching for each sample, revealing that the aforementioned partial switching phenomenon barely exists in HM based samples. Regarding the calculation of current density, the resistivity of Pt, Ta, and W is measured to be 27 $\mu\Omega$ ·cm, 112 $\mu\Omega$ ·cm, and 151 $\mu\Omega$ ·cm, respectively. We find that the values of j_{sw} to switch 4-nm-CoTb for Pt, Ta, and W are approximately 5.5×10^{11} A/m², 2.3×10^{11} A/m², and 1.7×10^{11} A/m². All those critical current densities are consistent with previously reported values^[5,7,8] and are obviously much larger than those in our Bi₂Te₃/CoTb system, although CoTb in Bi₂Te₃ based sample is even thicker, i.e., 6 nm. Note that, under an in-plane field with fixed direction, the switching polarity in Bi₂Te₃ sample is in line with that in Ta or W samples, which is opposite to that in Pt sample. This SOT polarity characteristic indicates that our sputtered Bi2Te3 possesses the same spin Hall angle sign as Ta and W, but is opposite to Pt. Nevertheless, this phenomenon is in contradiction with SOTs in previous-reported Bi-based materials: both MBE-grown TIs and sputtered Bi₂Se₃ films are demonstrated to have the same sign as Pt.^[19,20,25–27] The origin of this difference will be discussed subsequently. In our work, values of θ_{SH}^{eff} of Pt, Ta, and W in Fig. 4(a) are determined as 0.07 ± 0.007 , 0.14 ± 0.01 , and 0.18 ± 0.02 , through previously mentioned harmonic measurement and Eq. (4). In accordance with the varying j_{sw} values in those SOT materials, these calculated values are quite reliable and accepted in our following discussion.

Essentially, improving the magnetization switching energy efficiency is the eventual goal of obtaining higher θ_{SH}^{eff} . Considering that the resistivity of our sputtered Bi₂Te₃ is similar to that of MBE-grown TIs and can be dozens, even hundreds of times higher than that of conventional HMs, we think that it is significant to figure out whether the sputtered Bi₂Te₃ indeed offers energy superiority by calculating the switching power consumption *P*. Basically, *P* to switch the magnetization of CoTb per unit volume with different SOT sources can be analyzed by a heat dissipation formula^[27]

$$P \propto \rho_{\text{SOTsource}} \cdot j_{\text{sw}}^2,$$
 (5)

where $\rho_{\text{SOTsource}}$ is the resistivity of the certain SOT source material. As shown by orange bars in Fig. 4(b), the power consumption for Bi₂Te₃ in this model is 3.7%, 3.2%, and 3.9% of that for Pt, Ta, and W, indicating that sputtered Bi₂Te₃ is a low-power SOT source. Furthermore, since $j_{\text{sw}} \propto \theta_{\text{SH}}^{\text{eff}-1}$ theoretically, another formula is considered as well to confirm the results^[19]

$$P \propto \rho_{\text{SOTsource}} \cdot \theta_{\text{SH}}^{\text{eff}-2}.$$
 (6)

Black bars in Fig. 4(b) reveal that, although the calculated power of Pt, Ta, W with Eq. (5) varies in a possible range compared to those values with Eq. (4), the sputtered Bi_2Te_3 material still shows the best energy efficiency performance (2.8%, 3.5%, and 4.3% of that for Pt, Ta, and W).



Fig. 4. (a) Effective spin Hall angles of sputtered Pt, Ta, W, and Bi₂Te₃. (b) Normalized power consumption to switch the magnetization of CoTb per unit volume by different SOT source materials. The black bars and orange bars correspond respectively to the calculation formulas related to θ_{SH}^{eff} and j_{sw} .

Last but not least, we will discuss about the origin of the strong SOTs in the sputtered Bi2Te3 films. In MBE-grown TIs, SOTs basically come from the spin-momentum locking in TSS. Thus, great surface quality is the basic precondition to gain large θ_{SH}^{eff} . However, good TSS definitely will not appear in the deposited Bi2Te3 films, since magnetron sputtering is a rapid and rough growing method. Generally, there are two possible reasons to answer this question: 1) sputtered Bi chalcogenides could possibly possess nanoscale grain structure. Some ab initio calculation results suggest that this reduced dimensionality may contribute to the non-equilibrium spin accumulation driven by intraband Edelstein effect.^[27] 2) It is also possible that remarkable SOIs already exist in the sputtered Bi₂Te₃ films with relatively high resistivity and heavy elements. To further analyze the possible origin, an important point is that, the observed spin Hall angle sign of our sputtered Bi₂Te₃ films is opposite to that in MBE-grown TIs (same as Pt). Another recent work also reports a similar result: sputtered $W_x Te_{1-x}$ films possess opposite spin Hall angle sign to WTe₂ single crystals.^[38–40] However, for related Se-based material, the sputtered Bi₂Se₃ films have the same spin Hall angle to the MBE-grown Bi₂Se₃.^[27,28] Those evidences indicate that sputtered Te-based films probably exhibit different SOT mechanisms to sputtered Se-based films, although they both belong to the Bi chalcogenide class. Besides, the opposite spin Hall angle signs of sputtered Bi₂Te₃ and MBE-grown Bi₂Te₃ also exclude the possibility of major origin from TSS. Therefore, considering that both sputtered Te-based materials^[38] and Bi-based chalcogenides^[27] are reported to have obvious thickness-dependent spin Hall angle, the second reason attached to intrinsic bulk spin Hall effect is more convincing from our perspective.

4. Conclusion

In summary, we investigate SOTs in Bi₂Te₃ films grown by an industry-compatible deposition method magnetron sputtering. A high SOT efficiency χ (8.7±0.9 Oe/(10⁹ A/m²)) and a remarkable θ_{SH}^{eff} (3.3±0.3) are obtained in Bi₂Te₃/CoTb heterostructure as shown by harmonic Hall measurement. In addition, an ultra-low current switching current density (9.7× 10⁹ A/m² to switch 6-nm-thick PMA CoTb) is achieved as well. Compared with other sputtered SOT sources, the sputtered Bi₂Te₃ in our work shows much higher energy efficiency, indicating that it is a promising candidate in future SOT spintronic devices. Our work may also provide an alternative route to introduce more laboratory-level high-performance chalcogenides in industry-level spintronic applications.

References

- [1] Slonczewski J C 1996 J. Magn. Magn. Mater. 159 L1
- [2] Wadley P, Howells B, Železný J, Andrews C, Hills V, Campion R P, Novák V, Olejník K, Maccherozzi F, Dhesi S S, Martin S Y, Wagner T, Wunderlich J, Freimuth F, Mokrousov Y, Kuneš J, Chauhan J S, Grzybowski M J, Rushforth A W, Edmonds K W, Gallagher B L and Jungwirth T 2016 *Science* 351 587
- [3] Chen X, Zhou X, Cheng R, Song C, Zhang J, Wu Y, Ba Y, Li H, Sun Y, You Y, Zhao Y and Pan F 2019 Nat. Mater. 18 931
- [4] Yang T, Kimura T and Otani Y 2008 Nat. Phys. 4 851
- [5] Liu L, Pai C F, Li Y, Tseng H W, Ralph D C and Buhrman R A 2012 Science 336 555
- [6] Miron I M, Garello K, Gaudin G, Zermatten P G, Costache M V, Auffret S, Bandiera S, Rodmacq B, Schuhl A and Gambardella P 2011 *Nature* 476 189
- [7] Yu G, Upadhyaya P, Fan Y, Alzate J G, Jiang W, Wong K L, Takei S, Bender S A, Chang L T, Jiang Y, Lang M, Tang J, Wang Y, Tserkovnyak Y, Amiri P K and Wang K L 2014 *Nat. Nanotechnol.* 9 548
- [8] Zheng Z, Zhang Y, Feng X, Zhang K, Nan J, Zhang Z, Wang G, Wang J, Lei N, Liu D, Zhang Y G and Zhao W S 2019 *Phys. Rev. Appl.* 12 044032
- [9] Sato N, Xue F, White R M, Bi C and Wang S X 2018 Nat. Electron. 1 508
- [10] Zhang Z, Zhu Y, Zhang Y, Zhang K, Nan J, Zheng Z, Zhang Y G and Zhao W S 2019 *IEEE Electron. Device. Lett.* 40 1984
- [11] Zhang K, Zhang Y, Zhang Z, Zheng Z, Wang G, Zhang Y G, Liu Q, Yan S and Zhao W S 2019 Adv. Electron. Mater. 5 1800812

- [12] Wang M, Cai W, Zhu D, Wang Z, Kan J, Zhao Z, Cao K, Wang Z, Zhang Y, Zhang T, Park C, Wang J P, Fert A and Zhao W S 2018 Nat. Electron. 1 582
- [13] Zheng C, Chen H, Zhang X, Zhang Z and Liu Y 2019 Chin. Phys. B 28 037503
- [14] Feng X, Zhang Q, Zhang H, Zhang Y, Zhong R, Lu B, Cao J and Fan X 2019 Chin. Phys. B 28 107105
- [15] Ramaswamy R, Qiu X, Dutta T, Pollard S D and Yang H 2016 Appl. Phys. Lett. 108 202406
- [16] Finley J, Lee C H, Huang P Y and Liu L 2019 Adv. Mater. 31 1805361
- [17] Li P, Liu T, Chang H, Kalitsov A, Zhang W, Csaba G, Li W, Richardson D, DeMann A, Rimal G, Dey H, Jiang J S, Porod W, Field S, Tang J, Marconi M C, Hoffmann A, Mryasov O and Wu M 2016 Nat. Commun. 7 12688
- [18] Mellnik A R, Lee J S, Richardella A, Grab J L, Mintun P J, Fischer M H, Vaezi A, Manchon A, Kim E A, Samarth N and Ralph D C 2014 *Nature* 511 449
- [19] Han J, Richardella A, Siddiqui S A, Finley J, Samarth N and Liu L 2017 Phys. Rev. Lett. 119 077702
- [20] Wang Y, Zhu D, Wu Y, Yang Y, Yu J, Ramaswamy R, Mishra R, Shi S, Elyasi M, Teo K L, Wu Y and Yang H 2017 Nat. Commun. 8 1
- [21] Yang H, Zhang B, Zhang X, Yan X, Cai W, Zhao Y, Sun J, Wang K L, Zhu D and Zhao W S 2019 Phys. Rev. Appl. 12 034004
- [22] Ohtomo A and Hwang H Y 2004 Nature 427 423
- [23] Kageyama Y, Tazaki Y, An H, Harumoto T, Gao T, Shi J and Ando K 2019 Sci. Adv. 5 eaax4278
- [24] An H, Ohno T, Kanno Y, Kageyama Y, Monnai Y, Maki H and Ando K 2018 Sci. Adv. 4 eaar2250
- [25] Khang N H D, Ueda Y and Hai P N 2018 Nat. Mater. 17 808
- [26] Wu H, Xu Y, Deng P, Pan Q, Razavi S A, Wong K, Huang L, Dai B, Shao Q, Yu G, Han X, Sánchez J C R, Mangin S and Wang K L 2019 Adv. Mater. 31 1901681
- [27] Mahendra D C, Grassi R, Chen J, Jamali M, Hickey D R, Zhang D, Zhao Z, Li H, Quarterman P, Lv Y, Li M, Manchon A, Mkhoyan K A, Low T and Wang J P 2018 *Nat. Mater* 17 800
- [28] Zhang X, Cui B, Mao J, Yun J, Yan Z, Chang M, Zuo Y and Xi L 2020 Phys. Status Solidi-Rapid Res. Lett. 14 2000033
- [29] Sourabh, B, Rajeev and K P 2014 AIP Adv. 4 017135
- [30] Finley J and Liu L 2016 Phys. Rev. Appl. 6 054001
- [31] Ueda K, Mann M, de Brouwer P W P, Bono D and Beach G S D 2017 *Phys. Rev. B* 96 064410
- [32] Je S G, Sánchez J C R, Pham T H, Vallobra P, Malinowski G, Lacour D, Fache T, Cyrille M C, Kim D Y, Choe S B, Belmeguenai M, Hehn M, Mangin S, Gaudin G and Boulle O 2018 Appl. Phys. Lett. 112 062401
- [33] Woo S, Mann M, Tan A J, Caretta L and Beach G S D 2014 Appl. Phys. Lett. 105 212404
- [34] Hayashi M, Kim J, Yamanouchi M and Ohno H 2014 Phys. Rev. B 89 144425
- [35] Garello K, Miron I M, Avci C O, Freimuth F, Mokrousov Y, Blügel S, Auffret S, Boulle O, Gaudin G and Gambardella P 2013 Nat. Nanotechnol. 8 587
- [36] Zhang S, Su Y, Li X, Li R, Tian W, Hong J and You L 2019 Appl. Phys. Lett. 114 042401
- [37] Cao J, Chen Y, Jin T, Gan W, Wang Y, Zheng Y, Lv H, Cardoso S, Wei D and Lew W S 2018 Sci. Rep. 8 1355
- [38] Li X, Li P, Hou V D H, Mahendra D C, Nien C H, Xue F, Yi D, Bi C, Lee C M, Lin S J, Tsai W, Suzuki Y and Wang S X 2020 arXiv:2001.04054[cond-mat.mes-hall]
- [39] Shi S, Liang S, Zhu Z, Cai K, Pollard S D, Wang Y, Wang J, Wang Q, He P, Yu J, Eda G, Liang G and Yang H 2019 Nat. Nanotechnol. 14 945
- [40] Li P, Wu W, Wen Y, Zhang C, Zhang J, Zhang S, Yu Z, Yang S A, Manchon A and Zhang X 2018 Nat. Commun. 9 3990