## Tunable Sign Change of Spin Hall Magnetoresistance in Pt/NiO/YIG Structures

Dazhi Hou,<sup>1,2</sup> Zhiyong Qiu,<sup>1,2,\*</sup> Joseph Barker,<sup>3</sup> Koji Sato,<sup>1</sup> Kei Yamamoto,<sup>3,4,5</sup> Saül Vélez,<sup>6</sup> Juan M. Gomez-Perez,<sup>6</sup> Luis E. Hueso,<sup>6,7</sup> Fèlix Casanova,<sup>6,7</sup> and Eiji Saitoh<sup>1,2,3,8</sup>

<sup>1</sup>WPI Advanced Institute for Materials Research, Tohoku University, Sendai 980-8577, Japan

<sup>2</sup>Spin Quantum Rectification Project, ERATO, Japan Science and Technology Agency, Sendai 980-8577, Japan

<sup>3</sup>Institute for Materials Research, Tohoku University, Sendai 980-8577, Japan

<sup>4</sup>Institut für Physik, Johannes Gutenberg Universität Mainz, D-55128 Mainz, Germany

<sup>5</sup>Department of Physics, Kobe University, 1-1 Rokkodai, Kobe 657-8501, Japan

<sup>2</sup>CIC nanoGUNE, 20018 Donostia-San Sebastian, Basque Country, Spain

<sup>7</sup>IKERBASQUE, Basque Foundation for Science, 48011 Bilbao, Basque Country, Spain

<sup>8</sup>Advanced Science Research Center, Japan Atomic Energy Agency, Tokai 319-1195, Japan

(Received 24 October 2016; revised manuscript received 9 February 2017; published 3 April 2017)

Spin Hall magnetoresistance (SMR) has been investigated in Pt/NiO/YIG structures in a wide range of temperature and NiO thickness. The SMR shows a negative sign below a temperature that increases with the NiO thickness. This is contrary to a conventional SMR theory picture applied to the Pt/YIG bilayer, which always predicts a positive SMR. The negative SMR is found to persist even when NiO blocks the spin transmission between Pt and YIG, indicating it is governed by the spin current response of the NiO layer. We explain the negative SMR by the NiO "spin flop" coupled with YIG, which can be overridden at higher temperature by positive SMR contribution from YIG. This highlights the role of magnetic structure in antiferromagnets for transport of pure spin current in multilayers.

DOI: 10.1103/PhysRevLett.118.147202

Magnetoresistance plays essential roles in providing both a fundamental understanding of electron transport in magnetic materials and in various technological applications. Anisotropic magnetoresistance (AMR) [1,2], giant magnetoresistance [3,4], and tunneling magnetoresistance [5-8] underpin technologies in sensors, memories, and data storage. Recent studies of thin film bilayer systems comprised of a normal metal (NM) and a ferromagnetic insulator (FI) revealed a new type of magnetoresistance called spin Hall magnetoresistance (SMR) [9–11], originating from the interplay between the spin accumulation at the NM/FI interface and the magnetization of the FI layer. When the NM layer has a significant spin-orbit interaction, e.g., Pt, an in-plane charge current  $j_c$  induces a spin current via the spin Hall effect, which in turn generates a spin accumulation near the NM/FI interface. At the same time, this spin accumulation is affected by the orientation of the magnetization in the ferromagnet. The conductivity of the NM layer is thus subject to a magnetization dependent modification to the leading order in  $\theta_{\text{SHE}}^2$ , where  $\theta_{\text{SHE}}$  is the spin Hall angle in the NM layer.

Since the discovery of SMR, experimental studies were instigated in various systems [12–19]. The amplitude of SMR is defined as the difference of the resistivities with an in-plane magnetic field, H, parallel ( $\rho_{\parallel}$ ) and perpendicular  $(\rho_{\perp})$  to  $j_c$ :  $\rho_{\text{SMR}} = \rho_{\parallel} - \rho_{\perp}$ . This is predicted to be always positive because when  $H || j_c$ , the FI can absorb more spin current, by which the backflow required to ensure the stationary state is reduced at the FI/NM interface, in turn causing less secondary forward charge current, and therefore gives  $\rho_{\parallel} > \rho_{\perp}$  [9,10]. Positive  $\rho_{\text{SMR}}$  is found in most experimental observations.

Very recently, a negative SMR ( $\rho_{\parallel} < \rho_{\perp}$ ) was reported when an antiferromagnetic (AFM) insulator, in this case NiO, is inserted between Pt and YIG [20]. The negative SMR was also found to revert to the conventional positive sign at higher temperatures. Signal contamination from other magnetoresistances such as AMR was excluded by a systematic field angle dependence measurement. This result challenges the present understanding of SMR. Since the SMR does not change its sign in the Pt/YIG bilayer structure, the NiO layer must be the cause. However, it is not clear why NiO should give a negative SMR since antiferromagnets are thought only to affect the efficiency of the spin communication between Pt and YIG [21-26].

In this Letter, we report the temperature dependence of SMR in Pt/NiO/YIG structures with different thicknesses of NiO. The temperature at which the SMR becomes negative is found to depend on the NiO thickness. The anomalous negative SMR at low temperatures is explained from a spin-flop configuration whereby the Néel order of the NiO is perpendicularly coupled to the magnetization of YIG [27]. As the spin conductivity of NiO increases with increasing temperature [24-26], the moments of the YIG beneath have an increasing influence on the total SMR signal. The positive SMR contribution from YIG competes with the negative SMR from NiO. At the sign change, the competition leads to a vanishing SMR. Above, in the high temperature regime, the positive SMR of the YIG dominates. We introduce a phenomenological model to describe the competition between the positive and negative SMR contributions, which reproduces the NiO thickness dependent SMR sign change behaviors in Pt/NiO/YIG.

An epitaxial YIG film of thickness 3  $\mu$ m was grown on a gadolinium gallium garnet (111) substrate prepared by the liquid phase epitaxy. NiO films of different thicknesses were grown by sputtering onto the YIG at 400 °C. The film was then covered with 4 nm of sputtered Pt. The *x*-ray diffraction patterns of a 50 nm NiO film on YIG are plotted in Fig. 1, which only shows (111) and (222) NiO peaks of narrow line width. It suggests that the NiO film is of high crystallinity and a (111) preferred orientation. The inset in Fig. 1 shows a representative cross-section TEM picture for a Pt/NiO/YIG sample, which confirms a good thickness uniformity and clean interface.

Figure 2(a) shows the illustration of the magnetoresistance (MR) measurement setup and the definition of magnetic field angles. The standard four-probe method is employed for the MR observation at current density  $\sim 10^8$  A/m<sup>2</sup>, and MR can be detected either by sweeping H along a fixed direction or by rotating H of the same magnitude [9]. Figure 2(b) shows the MR measured by H sweeping in a Pt/NiO(2.5 nm)/YIG sample at field angle  $\alpha = 0^{\circ}$  for various temperatures. The range of magnetic field over which the magnetoresistance occurs coincides with that of the switching process of YIG [28]. The MR data for T > 140 K are consistent with the prediction  $\rho_{\parallel} > \rho_{\perp}$  of the SMR theory. When T = 140 K, the MR nearly vanishes. For T < 140 K, a sign change of MR is observed and the MR amplitude increases with decreasing temperature. The MR data from the same sample at field angle  $\alpha = 90^{\circ}$  are plotted in Fig. 2(c), which shows the same feature of the sign change. The SMR ratio  $\Delta \rho_{\text{SMR}} / \rho_{xx}$ extracted from Figs. 2(b) and 2(c) is plotted in Fig. 2(d). Figures 2(e) and 2(f) show the field angle dependence of resistance in Pt/NiO(2 nm)/YIG at 260 and 20 K, which not only reproduces the MR sign change behavior, but confirms the SMR-type field angle dependence symmetry as well [20]. Thus, it looks reasonable to claim that SMR is



FIG. 1. X-ray diffraction patterns of a 50 nm NiO film on YIG (111). The inset shows the cross-section TEM photo for a Pt/NiO/YIG trilayer measured in the transport experiment.

the dominant contribution for the MR in Pt on NiO/YIG, since other mechanisms such as anisotropic magnetoresistance cause a different field angle dependence [29]. However, the sign change of the magnetoresistance in the low temperature regime seems to be at odds with SMR, which, conventionally, can only be positive [10].

Figure 3(a) shows the temperature dependence of the SMR ratio measured in Pt/NiO/YIG devices with different NiO thicknesses,  $d_{\text{NiO}}$ . The change in sign of the SMR occurs at higher temperatures in larger  $d_{\text{NiO}}$  samples. The  $d_{\text{NiO}}$  dependence proves to be a key piece of information for understanding the negative SMR. Furthermore, the SMR ratios have (positive) maxima at higher temperatures for thicker NiO samples. These  $d_{\text{NiO}}$  dependent characteristics show a quantitative effect of the NiO on the SMR modulation, rather than a nuanced interface effect [30].

To gain further insight into the temperature dependence of spin transport in NiO, we carried out spin pumping measurements for the same samples, in which spin current is injected from YIG through NiO to generate a voltage in Pt via the inverse spin Hall effect (ISHE) [22]. The Pt/NiO/YIG device is placed on a coplanar waveguide that serves as a 5 GHz microwave source at 14 dbm, and the details of the experimental setup can be found elsewhere



FIG. 2. (a) The illustration for the magnetoresistance measurement setup for various magnetic field (*H*) orientations.  $\alpha$ ,  $\beta$ , and  $\gamma$  are the field angles defining the *H* directions when *H* is applied in the *x*-*y*, *x*-*z*, and *y*-*z* planes, respectively. [(b) and (c)] Magnetoresistance measured by *H* sweeping for a Pt/NiO(2.5 nm)/YIG at  $\alpha = 0^{\circ}$  and 90° for various temperatures. (d) Temperature dependence of the SMR ratio  $\Delta \rho_{\text{SMR}} / \rho_{xx}$  for Pt/NiO(2.5 nm)/YIG at  $\alpha = 0^{\circ}$  and 90°. [(e) and (f)] Field angle dependent resistance measured for Pt/NiO(2 nm)/YIG at 260 and 20 K with |H| = 20000 Oe, which shows positive and negative SMR, respectively.



FIG. 3. (a) The SMR ratio measured in  $Pt/NiO(d_{NiO})/YIG$  devices with different NiO thickness  $d_{NiO}$  at various temperatures, which shows that the SMR sign change temperature is lower for a thinner NiO sample. The SMR ratio peak positions are marked by arrows. Negative SMR at low temperatures can be observed for all the NiO thickness except  $d_{NiO} = 30$  nm. The dashed curves are the fitting based on Eq. (2). (b)  $V_{ISHE}$  in Pt/NiO/YIG devices versus temperature from the spin pumping measurement. The peak positions are marked by arrows, which are found to be close to the SMR ratio peak positions marked in Fig. 2(a). The inset shows the normalized  $V_{ISHE}$  temperature dependence.

[24]. The ISHE voltage  $V_{\text{ISHE}}$  from all the samples is plotted against *T* in Fig. 3(b), the behavior of which is very similar to the result we found in Pt/CoO/YIG [24]: spin transmission is nearly 0 for the low temperature limit and increases with temperature to reach the maximum around the Néel point. At room temperature,  $V_{\text{ISHE}}$  shows a nonmonotonic  $d_{\text{NiO}}$ dependence, which is consistent with the previous result. Figure 3(b)'s inset shows the normalized  $V_{\text{ISHE}}$  temperature dependence, in which the data for  $d_{\text{NiO}} = 5.4$ , 7, and 15 nm collapse into a single curve. This confirms that the  $V_{\text{ISHE}}$  is governed by the NiO spin conductivity, which shows the same *T* dependence when NiO is thick enough to exhibit bulk property. For  $d_{\text{NiO}} = 30$  nm,  $V_{\text{ISHE}}$  is below our measurement sensitivity 5 nV.

An important conclusion can be drawn by combining the results from SMR and spin pumping measurements: the negative SMR does not rely on the spin transmission between Pt and YIG, because it reaches the largest magnitude for the lowest temperature at which NiO spin conductivity vanishes. This argument can be further supported by the fact that the negative SMR is present even for  $d_{\text{NiO}} = 15$  nm, where the NiO spin conductivity is nearly 0 throughout the entire temperature range. It indicates that the negative SMR is not caused by the magnetic moment of the YIG layer but that of the NiO layer, which is beyond any model based on spin communication between YIG and Pt [10,31].

Let us next provide an explanation for the negative SMR. The SMR in the trilayer system in this experiment is governed by the spin current through the Pt/NiO interface, which also reflects the effect of the presence of the NiO/YIG interface. The sign change and the thickness dependent behavior can be understood by assuming a spinflop coupling between NiO and YIG [27,32], which means the antiferromagnetic axis (Néel vector unit  $n_{AFM}$ ) in NiO is perpendicular to the YIG magnetization unit vector  $m_{\rm FI}$  as illustrated in Fig. 4(a). Although a perpendicular coupling has not yet been confirmed experimentally for NiO on YIG, spin-flop coupling between NiO and other ferromagnets is quite common and well understood [27,33,34]. For  $d_{\text{NiO}}$ below the domain wall width of NiO (~15 nm) [35], which is the case for nearly all the samples,  $n_{AFM}$  tends to be uniform in NiO, which is strongly coupled with YIG and can be manipulated by magnetic field [36]. Thus,  $n_{AFM}$  is always perpendicular to H below the Néel temperature, because the  $m_{\rm FI}$  is parallel to H. In the low temperature limit, e.g., 10 K, the spin current generated in Pt cannot



FIG. 4. Illustrations for the magnetic structure and spin transport in Pt/NiO/YIG at different temperatures. The red and green arrows represent the phenomenologically described spin currents,  $j_1$  and  $j_2$  in Eq. (1), respectively. The length of the arrow describes the penetration depth of the spin current. (a) *T* close to the low temperature limit. (b) *T* far above the low temperature limit and lower than the Néel temperature. (c) *T* higher than the Néel temperature. (d) Illustration of *T*-dependent SMR in which the temperatures corresponding to the conditions in Figs. 4(a)–4(c) are marked with a red circle.

penetrate through the NiO; thus, the SMR signal is only caused by the NiO layer. The NiO local moments perpendicular to H give rise to a 90-degree phase shift in the SMR field angular dependence with respect to the conventional SMR [9]. Such a 90-degree phase shift in a fourfold SMR field angular dependence is equivalent to a sign reversal in the conventional definition of MR, which explains the negative SMR in Pt/NiO/YIG at low temperatures. For  $d_{\text{NiO}} = 30$  nm, which is beyond the domain wall width,  $n_{\text{AFM}}$  at the Pt/NiO interface decouples with  $m_{\text{FI}}$  and does not respond to H, which explains the vanishing of the negative SMR.

At higher temperatures, but below the Néel point, antiferromagnetic order is maintained but the spin current from Pt has some transmission through NiO, which makes the effect of the YIG more visible as illustrated in Fig. 4(b). The negative SMR contribution from NiO and the positive SMR contribution from YIG compete with each other. With increasing temperature, NiO becomes more transparent to the spin current, so the SMR contribution from YIG is enhanced. The SMR from NiO may also be suppressed because of the attenuation of the antiferromagnetic order at elevated temperatures. As a result, the zero point of the SMR occurs at a temperature where the antiferromagnet is still in the ordered phase. Thinner NiO layers have a lower Néel point due to the finite size effect [37]; hence, the SMR also changes the sign at lower temperatures in thinner-NiO samples, which is in accordance with our observation shown in Fig. 3(a).

Around the Néel point as illustrated in Fig. 4(c), the spin transparency of NiO is maximized [24], where the SMR contribution from YIG reaches its peak value and the SMR contribution from NiO vanishes. As explained above, all the main features of the SMR data in Pt/NiO/YIG, such as negative SMR at low temperatures,  $d_{\rm NiO}$  dependent sign change temperature and peak temperature, can be interpreted by the spin-flop configuration. Figure 4(d) shows an illustration of  $\rho_{\rm SMR}$  temperature dependence, in which the temperature corresponding to these features is marked. We note that negative SMR has also been reported in bilayers of Pt on gadolinium iron garnet and Ar-sputtered YIG, in which the garnet interface moments can align perpendicularly to *H* [30,38].

A simple phenomenological model based on the picture discussed above can also provide a quantitative description of the observed SMR temperature dependence. Let us consider a NM/AFM/FI trilayer system. The key assumption is that we can describe the spin current through the NM/AFM interface by

$$e\mathbf{j}_{s} = G_{\rm AF}\mathbf{n}_{\rm AFM} \times (\mathbf{n}_{\rm AFM} \times \boldsymbol{\mu}_{s}) + t(T)\mathbf{m}_{\rm FI} \times (\mathbf{m}_{\rm FI} \times \boldsymbol{\mu}_{s})$$
$$= e\mathbf{j}_{1} + e\mathbf{j}_{2}; \tag{1}$$

 $G_{\rm AF}$  is the real part of the spin mixing conductance at the NM/AFM interface.  $\mu_s$  is the spin accumulation at the same interface. The first term, which we denote by  $ej_1$ , is

what is expected for NM/AFM bilayer systems as seen in the case studied in Ref. [39]. We have introduced the second term, which is denoted by  $ej_2$ , to phenomenologically capture the effect of the FI layer. t(T) encapsulates the temperature dependent transparency of the AFM to the spin current. In the case that the AFM is completely transparent the NM/FI bilayer result  $m_{\text{FI}} \times (m_{\text{FI}} \times \mu_s)$  is recovered. The linear combination of the NM/AFM and NM/FI terms has been chosen in an attempt to emulate our SMR data in the NM/ AFM/FI system, seen in Fig. 3(a), which seems to indicate a crossover from NM/AFM bilayerlike behavior at low temperatures to NM/FI bilayerlike behavior for higher temperatures.

Once we admit the form of the interfacial spin current in Eq. (1), we can calculate the SMR by employing the diffusion equation and the Onsagar principle, according to Refs. [10]. The SMR contribution to the longitudinal resistivity then is given by

$$\frac{\delta\rho}{\rho_0} = \frac{2\theta_{\text{SHE}}^2 \lambda_N^2}{d_N \sigma} \times \frac{G_{\text{AF}} \cos^2 \phi_n + t(T) \cos^2 \phi_m + \nu t(T) G_{\text{AF}} \sin^2 (\phi_m - \phi_n)}{1 + \nu G_{\text{AF}} + \nu t(T) + \nu^2 t(T) G_{\text{AF}} \sin^2 (\phi_m - \phi_n)} \times \tanh^2 \left(\frac{d_N}{2\lambda_N}\right),$$
(2)

where we defined  $\nu = (2\lambda_N/\sigma) \coth(d_N/\lambda_N)$  with  $\lambda_N$ and  $\theta_{\text{SHE}}$  being the spin diffusion length and the spin Hall angle in NM, respectively, and  $\sigma = \rho_0^{-1}$  the conductivity of the NM layer. Here,  $\phi_{n(m)}$  denotes the angle between  $n_{\text{AFM}}(m_{\text{FI}})$  and the applied current  $j_c$  in NM. Now we set out a hypothesis that the crossover between the negative and positive SMR is of the same origin as the temperature dependence of the spin pumping signal [Fig. 3(b)]. In order to support it, the temperature dependence of t(T) is obtained by fitting to the spin pumping data. The resulting function is then used alongside the other parameters in Eq. (1) to fit the SMR data to test the validity of our model.

Based on the observation that the ISHE signal in Fig. 3(b) is roughly exponential in the intermediate temperature regime, we employ  $V_{\text{ISHE}} \propto t(T) \propto e^{aT} - 1$  to reproduce the temperature dependence of both spin pumping and SMR. The exponential behavior may not apply near the Néel temperature and the data points near and above the Néel temperature have been excluded from the fitting. Under these assumptions, *a* can be determined from the spin pumping data (Table I).

We then fit  $\delta \rho / \rho |_{\phi_m = 0} - \delta \rho / \rho |_{\phi_m = \pi/2}$  based on Eq. (2) to the experimentally obtained SMR ratio  $\Delta \rho_{\text{SMR}} / \rho_{xx}$  in Fig. 3(a) using the fitted value of *a* from the  $V_{\text{ISHE}}$  data. We fix  $\lambda_N = 1.5$  nm,  $d_N = 4.0$  nm,  $\rho_0 = \sigma^{-1} = 860 \Omega$  nm, and  $\theta_{\text{SHE}} = 0.05$ , which are taken to be relevant values to the present experiment, and we further determine  $G_{\text{AF}}$ and  $G_F$  from the data, where the latter two are defined by  $t(T) = G_F(e^{aT} - 1), \ \phi_n - \phi_m = \pi/2$ , respectively.

TABLE I. The results of the fitting with the data from the SMR and spin pumping signals. The parameters are defined in the main text. The units of the last two columns are both  $[\Omega^{-1} \text{ m}^{-2}]$ .

$d_{\rm NiO}$	$a[K^{-1}] \times 10^2$	$G_{ m AF}$	$G_F$
2.0	$1.83\pm0.22$	$3.58 \pm 0.32 \times 10^{12}$	$8.39 \pm 0.57  imes 10^{11}$
2.2	$1.38\pm0.19$	$4.48 \pm 0.17 \times 10^{12}$	$7.78 \pm 0.26  imes 10^{11}$
2.7	$1.42\pm0.10$	$3.67 \pm 0.09 \times 10^{12}$	$3.01 \pm 0.08 \times 10^{11}$
4.0	$1.16\pm0.09$	$2.46 \pm 0.13 \times 10^{12}$	$2.22 \pm 0.14 \times 10^{11}$

The temperature dependence of  $\rho_0$  and  $\theta_{\text{SHE}}$  is ignored since they scale in some powers of *T*, which is wiped out by the exponential change in t(T). The fitting curves can quantitatively reproduce the SMR sign change behavior as shown in Fig. 3(a), and the fitting parameters are summarized in Table I.

Our result highlights the importance of magnetic structure in AFM for spin transport, which suggests an alternative degree of freedom of spin manipulation. The NiO-induced SMR indicates that the spin current response of AFM is anisotropic, which opens the possibility to use AFM insulator as a spin current valve or memory.

The authors thank the valuable discussion with Jia Li and Qian Li, and technical support from Yao Chen. This work was supported by ERATO "Spin Quantum Rectification Project" from JST, Japan and Grant-in-Aid for Scientific Research on Innovative Area "Nano Spin Conversion Science" (No. JP26103005) from JSPS KAKENHI, Japan. J.B. acknowledges support from the Graduate Program in Spintronics, Tohoku University. K.S. was supported by Grants-In-Aid for Scientific Research (Grant No. 15K13531) from the MEXT of Japan. K.Y. is supported by the JSPS Grant-in-Aid for Scientific Research No. 26 · 1204, the Alexander von Humboldt Foundation, the ERC Synergy Grant SC2 (No. 610115), the Transregional Collaborative Research Centre (SFB/ TRR) 173 SPIN+X, Grant Agency of the Czech Republic Grant No. 14-37427G, and DAAD project "MaHoJeRo." The work at CIC nanoGUNE was supported by the Spanish MINECO (Project No. MAT2015-65159-R) and by the Regional Council of Gipuzkoa (Project No. 100/16).

*Note added.*—Recently, we became aware of similar results for the SMR sign change observed in Pt/NiO/YIG by W. Lin *et al.* [31]. The NiO-thickness dependent SMR at room temperature was also reported by Yu-Ming Hung *et al.* [40].

<sup>\*</sup>qiuzy@imr.tohoku.ac.jp

- [1] I. A. Campbell, A. Fert, and O. Jaoul, J. Phys. C **3**, S95 (1970).
- [2] T. McGuire and R. Potter, IEEE Trans. Magn. 11, 1018 (1975).
- [3] G. Binasch, P. Grünberg, F. Saurenbach, and W. Zinn, Phys. Rev. B 39, 4828 (1989).

- [4] M. N. Baibich, J. M. Broto, A. Fert, F. NguyenVanDau, F. Petroff, P. Etienne, G. Creuzet, A. Friederich, and J. Chazelas, Phys. Rev. Lett. 61, 2472 (1988).
- [5] M. Julliere, Phys. Lett. 54A, 225 (1975).
- [6] T. Miyazaki and N. Tezuka, J. Magn. Magn. Mater. 139, L231 (1995).
- [7] S. Yuasa, T. Nagahama, A. Fukushima, Y. Suzuki, and K. Ando, Nat. Mater. 3, 868 (2004).
- [8] S. S. P. Parkin, C. Kaiser, A. Panchula, P. M. Rice, B. Hughes, M. Samant, and S.-H. Yang, Nat. Mater. 3, 862 (2004).
- [9] H. Nakayama, M. Althammer, Y.-T. Chen, K. Uchida, Y. Kajiwara, D. Kikuchi, T. Ohtani, S. Geprägs, M. Opel, S. Takahashi, R. Gross, G. E. W. Bauer, S. T. B. Goennenwein, and E. Saitoh, Phys. Rev. Lett. **110**, 206601 (2013).
- [10] Y.-T. Chen, S. Takahashi, H. Nakayama, M. Althammer, S. T. B. Goennenwein, E. Saitoh, and G. E. W. Bauer, Phys. Rev. B 87, 144411 (2013).
- [11] M. Althammer et al., Phys. Rev. B 87, 224401 (2013).
- [12] M. Weiler, M. Althammer, M. Schreier, J. Lotze, M. Pernpeintner, S. Meyer, H. Huebl, R. Gross, A. Kamra, J. Xiao, Y.-T. Chen, H. J. Jiao, G. E. W. Bauer, and S. T. B. Goennenwein, Phys. Rev. Lett. **111**, 176601 (2013).
- [13] S. Meyer, M. Althammer, S. Geprägs, M. Opel, R. Gross, and S. T. B. Goennenwein, Appl. Phys. Lett. 104, 242411 (2014).
- [14] R. Iguchi, K. Sato, D. Hirobe, S. Daimon, and E. Saitoh, Appl. Phys. Express 7, 013003 (2014).
- [15] M. Schreier, T. Chiba, A. Niedermayr, J. Lotze, H. Huebl, S. Geprägs, S. Takahashi, G. E. W. Bauer, R. Gross, and S. T. B. Goennenwein, Phys. Rev. B 92, 144411 (2015).
- [16] J. Lotze, H. Huebl, R. Gross, and S. T. B. Goennenwein, Phys. Rev. B 90, 174419 (2014).
- [17] S. R. Marmion, M. Ali, M. McLaren, D. A. Williams, and B. J. Hickey, Phys. Rev. B 89, 220404 (2014).
- [18] M. Isasa, A. Bedoya-Pinto, S. Vélez, F. Golmar, F. Sánchez, L. E. Hueso, J. Fontcuberta, and F. Casanova, Appl. Phys. Lett. **105**, 142402 (2014).
- [19] C. O. Avci, K. Garello, A. Ghosh, M. Gabureac, S. F. Alvarado, and P. Gambardella, Nat. Phys. 11, 570 (2015).
- [20] T. Shang, Q. F. Zhan, H. L. Yang, Z. H. Zuo, Y. L. Xie, L. P. Liu, S. L. Zhang, Y. Zhang, H. H. Li, B. M. Wang, Y. H. Wu, S. Zhang, and R.-W. Li, Appl. Phys. Lett. **109**, 032410 (2016).
- [21] C. Hahn, G. de Loubens, V. V. Naletov, J. B. Youssef, O. Klein, and M. Viret, Europhys. Lett. 108, 57005 (2014).
- [22] H. Wang, C. Du, P. C. Hammel, and F. Yang, Phys. Rev. Lett. 113, 097202 (2014).
- [23] T. Moriyama, S. Takei, M. Nagata, Y. Yoshimura, N. Matsuzaki, T. Terashima, Y. Tserkovnyak, and T. Ono, Appl. Phys. Lett. 106, 162406 (2015).
- [24] Z. Qiu, J. Li, D. Hou, E. Arenholz, A. T. N'Diaye, A. Tan, K.-i. Uchida, K. Sato, S. Okamoto, Y. Tserkovnyak, Z. Q. Qiu, and E. Saitoh, Nat. Commun. 7, 12670 (2016).
- [25] L. Frangou, S. Oyarzún, S. Auffret, L. Vila, S. Gambarelli, and V. Baltz, Phys. Rev. Lett. **116**, 077203 (2016).
- [26] W. Lin, K. Chen, S. Zhang, and C. L. Chien, Phys. Rev. Lett. 116, 186601 (2016).
- [27] N.C. Koon, Phys. Rev. Lett. 78, 4865 (1997).
- [28] K. Uchida, Z. Qiu, T. Kikkawa, R. Iguchi, and E. Saitoh, Appl. Phys. Lett. **106**, 052405 (2015).

- [29] X. Zhou, L. Ma, Z. Shi, W. J. Fan, J.-G. Zheng, R. F. L. Evans, and S. M. Zhou, Phys. Rev. B 92, 060402 (2015).
- [30] S. Vélez, A. Bedoya-Pinto, W. Yan, L. E. Hueso, and F. Casanova, Phys. Rev. B 94, 174405 (2016).
- [31] W. Lin and C. L. Chien, Phys. Rev. Lett. **118**, 067202 (2017).
- [32] T. C. Schulthess and W. H. Butler, Phys. Rev. Lett. **81**, 4516 (1998).
- [33] I. P. Krug, F. U. Hillebrecht, M. W. Haverkort, A. Tanaka, L. H. Tjeng, H. Gomonay, A. Fraile-Rodríguez, F. Nolting, S. Cramm, and C. M. Schneider, Phys. Rev. B 78, 064427 (2008).
- [34] J. Li, A. Tan, S. Ma, R. F. Yang, E. Arenholz, C. Hwang, and Z. Q. Qiu, Phys. Rev. Lett. **113**, 147207 (2014).
- [35] M. Fraune, U. Rdiger, G. Gntherodt, S. Cardoso, and P. Freitas, Appl. Phys. Lett. 77, 3815 (2000).

- [36] W. Kim, E. Jin, J. Wu, J. Park, E. Arenholz, A. Scholl, C. Hwang, and Z. Q. Qiu, Phys. Rev. B 81, 174416 (2010).
- [37] D. Alders, L. H. Tjeng, F. C. Voogt, T. Hibma, G. A. Sawatzky, C. T. Chen, J. Vogel, M. Sacchi, and S. Iacobucci, Phys. Rev. B 57, 11623 (1998).
- [38] K. Ganzhorn, J. Barker, R. Schlitz, B. A. Piot, K. Ollefs, F. Guillou, F. Wilhelm, A. Rogalev, M. Opel, M. Althammer, S. Geprägs, H. Huebl, R. Gross, G. E. W. Bauer, and S. T. B. Goennenwein, Phys. Rev. B 94, 094401 (2016).
- [39] S. Takei, B. I. Halperin, A. Yacoby, and Y. Tserkovnyak, Phys. Rev. B 90, 094408 (2014); Y.-T. Chen, S. Takahashi, H. Nakayama, M. Althammer, S. T. B. Goennenwein, E. Saitoh, and G. E. W. Bauer, J. Phys. Condens. Matter 28, 103004 (2016).
- [40] Y.-M. Hung, C. Hahn, H. Chang, M. Wu, H. Ohldag, and A. D. Kent, AIP Adv. 7, 055903 (2017).