Unidirectional Spin Hall Magnetoresistance in Antiferromagnetic Heterostructures

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Unidirectional spin Hall magnetoresistance (USMR) has been widely reported in the heavy metal/ ferromagnet bilayer systems. We observe the USMR in Pt/α -Fe₂O₃ bilayers where the α -Fe₂O₃ is an antiferromagnetic (AFM) insulator. Systematic field and temperature dependent measurements confirm the magnonic origin of the USMR. The appearance of AFM-USMR is driven by the imbalance of creation and annihilation of AFM magnons by spin orbit torque due to the thermal random field. However, unlike its ferromagnetic counterpart, theoretical modeling reveals that the USMR in Pt/ α -Fe₂O₃ is determined by the antiferromagtic magnon number with a non-monotonic field dependence. Our findings extend the generality of the USMR which pave the ways for the highly sensitive detection of AFM spin state.

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The detection of the spin state is one of the central topics in spintronics [1–3]. Spin Hall magnetoresistance (SMR) has been widely used to probe magnetization in heavy metal/ (anti)ferromagnetic [HM/(A)FM] heterostructures [2,4–7]. When the current is applied in a heavy metal layer, the generated spin current is injected to the adjacent magnetic layer. The additional currents induced by inverse spin Hall effects change the resistivity of the heterostructures where its magnitude only depends on the relative angle between magnetization and spin polarization. So far, SMR has been widely reported in both ferromagnetic and antiferromagnetic systems. Recently, a new type of magnetoresistance-unidirectional SMR (USMR) [8-11]-has attracted intense interest. Compared with conventional SMR, the USMR is a non-linear magnetoresistance where the measured voltage depends quadratically on the applied current. What is more, the magnitude unidirectionally depends on the angle between spin polarization and magnetization as its name suggests, which provides a more precise way to probe the spin state in the magnetic layer. USMR has been observed only in HM/FM heterostructures. For metallic FM, the USMR is originated from either spindependent electron scattering (spin-dependent USMR) or electron-magnon scattering (spin-flip USMR) [9,12]. For insulating FM, for example Pt/Y₃Fe₅O₁₂ (YIG) bilayers [10], the observed USMR is attributed to the imbalance of magnon generation and annihilation rate by the spin-orbit torque. Although the SMR has been observed in both FMs and AFMs, the USMR is not expected in the AFMs. Even for YIG which is a ferrimagnet, the observed USMR behaves like a ferromagnet, but is not related to its antiferromagnetic ordering [13]. This is because the order parameter of AFMs. the staggered magnetization (Néel vector), is a pair of two sublattice magnetizations, which makes the spin polarization with a 180° rotational symmetry relative to the Néel vector. A possible way to break this symmetry is by applying an external field to induce a net FM order from sublattice magnetization canting. Here, we report the detection of USMR in Pt/ α -Fe₂O₃ bilayers using the second harmonic measurements. Through systematic field and temperature dependent measurement, and theoretical modeling, we attribute the observed USMR to the imbalance of the magnon generation and annihilation rates, similar to the USMR in FM insulators [10]. However, the antiferromagnetic magnon plays a dominant role whereas the induced magnetization only contributes a small part to the USMR from our simulation.

 α -Fe₂O₃ is an easy-plane antiferromagnet with the Néel temperature around 953 K [14]. The weak anisotropy field of three easy axes in the ab plane (0001) makes the spin flop (or spin reorientation) occur at the critical field of ~ 1 T, where the Néel order is perpendicular to the magnetic field, $n \perp H$ [15]. This makes the field control of the Néel order of α -Fe₂O₃ much easier compared with most of the ypical antiferromagnets, a necessary condition for the extraction of the USMR. We grow a Pt $(5 \text{ nm})/\alpha$ -Fe₂O₃(30 nm) thin film stack on Al₂O₃ (0001) using off-axis magnetron sputtering [15]. Bulk α -Fe₂O₃ experiences the so-called Morin transition, which changes its phase from easy-plane AFM to easy-axis (c axis [0001]) AFM at ~260 K. However, due to the strain induced enhancement of hard-axis anisotropy, our thin film α -Fe₂O₃ does not show such transition down to



FIG. 1. Experimental geometry and first harmonic results. (a) Schematics of a Pt/ α -Fe₂O₃ Hall bar with a 5 µm width and 10 µm length. (b) In plane angular dependence of first harmonic voltage $V_{xx}^{1\omega}$ (blue curve) and $V_{xy}^{1\omega}$ (green curve) for a Pt(5 nm)/ α -Fe₂O₃(30 nm) bilayer at 300 K with 2 T applied field. (c) Field dependence of (transverse) spin Hall magnetoresistance voltage $V_{(T)SMR}$ extracted from the fitting in (b) by Eq. (1) where the inset of (c) shows the ratio of V_{SMR} and V_{TSMR} .

10 K as had been demonstrated in our previous research [15–18]. After the growth of the Pt/α -Fe₂O₃ bilayer, we pattern our sample into a Hall bar device with a length (l) 10 µm and a width (w) 5 µm, as shown in Fig. 1(a).

Since USMR is a nonlinear current effect, an angular dependent harmonic measurement method is commonly used [10,19]. For the harmonic measurement setup, a low frequency 5 Hz ac current $I = I_0 \sin(\omega t)$ is applied to the Hall bar device. The longitudinal (V_{xx}) and transverse (V_{xy}) voltage are measured simultaneously under an in-plane magnetic field. The first harmonic response is the same as the dc measurement, where [20]

$$V_{xx}^{1\omega} = V_{\rm SMR} \sin^2 \varphi_{\rm H},\tag{1}$$

$$V_{xy}^{1\omega} = -V_{\text{TSMR}} \sin \varphi_{\text{H}} \cos \varphi_{\text{H}}.$$
 (2)

Here $\varphi_{\rm H}$ is the in-plane angle between applied field and current direction, as shown in Fig 1(a). $V_{\rm SMR}$ and $V_{\rm TSMR}$ are the longitudinal and transverse spin Hall magnetoresistance where $(V_{\rm SMR}/V_{\rm TSMR}) = (l/w) = 2$, the aspect ratio of our Hall bar [15,21]. Figure 1(b) shows the angular dependent first harmonic measurement at 2 T and 300 K. The current $I_0 = 6$ mA. Figure 1(c) shows the extracted $V_{\rm SMR}$ and $V_{\rm TSMR}$ using Eqs. (1) and (2) as the field increases from 0.3 to 9 T. The magnitude of $V_{\rm SMR}$ and $V_{\rm TSMR}$ saturates near 1 T, which indicates a single domain AFM state at $\mu_0 H > 1$ T. The ratio of $V_{\rm SMR}$ and $V_{\rm TSMR}$ is shown in the inset of Fig. 1(c), which is close to 2, as expected.

As shown in Fig. 2(a), a current induced effect such as spin-orbit torque drives the Néel order away from its equilibrium position where the change of magnetoresistance can be probed via the second harmonic voltage $V_{xx}^{2\omega}$



FIG. 2. (a) Schematic of current induced spin orbit torque in two spin sublattices $m_{A(B)}$. In-plane angular dependence of second harmonic voltage (b) $V_{xx}^{2\omega}$ and (c) $V_{xy}^{2\omega}$ for the Pt(5 nm)/ α -Fe₂O₃(30 nm) bilayer at 300 K with 2 T applied field. The blue, green, and black curves are contributions from the fieldlike torque, spin Seebeck effect, and USMR, respectively. The red curves are the total fit by Eqs. (3) and (4).

and $V_{xy}^{2\omega}$. Based on our previous research [20] (for more details, see Supplemental Material [22]), the second harmonic voltage can be rewritten as

$$V_{xx}^{2\omega} = V_{xx,FL}^{2\omega} + V_{xx,SSE}^{2\omega} + V_{xx,USMR}^{2\omega}$$
$$= -\frac{1}{2} V_{SMR} \frac{H_{FL}}{H} \sin(2\varphi_{\rm H}) \cos\varphi_{\rm H}$$
$$- (V_{xx,SSE} + V_{xx,USMR}) \sin\varphi_{\rm H}, \qquad (3)$$

$$V_{xy}^{2\omega} = V_{xy,FL}^{2\omega} + V_{xy,SSE}^{2\omega}$$

= $\frac{1}{2} V_{TSMR} \frac{H_{FL}}{H} \cos(2\varphi_{H}) \cos \varphi_{H} + V_{xy,SSE} \cos \varphi_{H}.$ (4)

Here, $H_{\rm FL}$ is the fieldlike torque effective field. The longitudinal $V_{xx,{\rm FL}}^{2\omega}$, $V_{xx,{\rm SSE}}^{2\omega}$, and $V_{xx,{\rm USMR}}^{2\omega}$ are the contributions from field-like torque, spin Seebeck effect, and USMR, respectively. The transverse $V_{xy,{\rm FL}}^{2\omega}$ and $V_{xy,{\rm SSE}}^{2\omega}$ are the fieldlike torque and spin Seebeck effect terms. Notice that USMR only shows up in the $V_{xx}^{2\omega}$ term. To extract the USMR contributions $V_{xx,{\rm USMR}}$, we first fit the $V_{xx}^{2\omega}$ and $V_{xy}^{2\omega}$ to get the $V_{xx,{\rm SSE}} + V_{xx,{\rm USMR}}$ and $V_{xy,{\rm SSE}}$; $V_{xx,{\rm USMR}}$ can be separated from $V_{xx,{\rm SSE}}$ given that $(V_{xx,{\rm SSE}}^{2\omega}/V_{xy,{\rm SSE}}^{2\omega}) = (l/w) = 2$. Compared with the fitting to ferromagnets, the transverse second harmonic voltage does not contain the dampinglike (DL) torque term, which makes the results more convincing due to less fitting process [10]. Figures 2(b) and 2(c) show the angular dependence of second harmonic voltage $V_{xx}^{2\omega}$ and $V_{xy}^{2\omega}$ at 2 T and 300 K. We fit the data using Eqs. (3) and (4). Clearly, there is a USMR contribution in $V_{xx,{\rm SSE}}^{2\omega}$ after subtracting the longitudinal spin Seebeck component $V_{xx,{\rm SSE}}^{2\omega}$ with the same angular dependence.



FIG. 3. In-plane angular dependence of second harmonic Hall voltage (a) $V_{xx}^{2\omega}$ and (b) $V_{xy}^{2\omega}$ at different magnetic fields for the Pt(5 nm)/ α -Fe₂O₃(30 nm) bilayer at 300 K. (c) Field dependence of fieldlike torque contribution in $V_{xx}^{2\omega}$ (blue curve) and $V_{xy}^{2\omega}$ (green curve). The solid line is the 1/*H* fit. (d) Field dependence of spin Seebeck effect contribution in $V_{xx}^{2\omega}$ (blue curve) and $V_{xy}^{2\omega}$ (green curve). The solid line is the linear fit. (e) The ratio of $V_{xx,FL}$ and $V_{xy,FL}$ (blue curve) and the ratio of $V_{xx,SSE} + V_{xx,USMR}$ and $V_{xy,SSE}$ (green curve), where the magnitude is calculated from (c) and (d). The ratio of $V_{xx,SSE} + V_{xx,USMR}$ and $V_{xy,SSE}$ is greater than 2, which indicates the presence of USMR. Various contributions in (c) to (e) are obtained by fitting like those in Fig. 2.

Following the same process, we obtain $V_{xx}^{2\omega}$ and $V_{xy}^{2\omega}$ of the Pt(5 nm)/ α -Fe₂O₃(30 nm) bilayer from 1 to 9 T and fitting curves, as shown in Figs. 3(a) and 3(b). Figure 3(c) shows the fitted field-like torque component from Figs. 3(a) and 3(b). The extracted $V_{xx,FL}^{2\omega}$ and $V_{xy,FL}^{2\omega}$ decrease with the increasing field and follow a 1/H dependence. The ratio of $V_{xx,FL}^{2\omega}$ and $V_{xy,FL}^{2\omega}$ is shown in the blue curve of Fig. 3(e), which is around 2 for the entire field range as $(V_{xx,FL}^{2\omega}/V_{xy,FL}^{2\omega}) = (V_{SMR}/V_{TSMR}) = (l/w) = 2.$ At the same time, the sinusoidal and cosinusoidal components in $V_{xx}^{2\omega}$ and $V_{xy}^{2\omega}$ are extracted by fitting the data with Eqs. (3) and (4) where the ratio of them are plotted in the green curve of Fig. 3(e). The sinusoidal term in $V_{xx}^{2\omega}$ is expected to contain both longitudinal spin Seebeck and USMR contributions while the cosinusoidal term in $V_{xy}^{2\omega}$ is only from the transverse spin Seebeck voltage, which is linearly proportional to the field H since $V_{SSE}^{2\omega} \propto m \propto H$ [25]. In the green curve of Fig. 3(e), it is shown that the ratio of $[(V_{xx,SSE}^{2\omega} + V_{xx,USMR}^{2\omega})/V_{xy,SSE}^{2\omega}] > 2$, indicating the existence of USMR.

Figure 4(a) shows the field dependence of extracted USMR at 300 K. Surprisingly, unlike the USMR in a ferromagnet where the magnitude either monotonically decreases or is unchanged as the field increases, the USMR in the antiferromagnetic Pt/α -Fe₂O₃ bilayer shows a nonmonotonic field dependence. The magnitude of USMR increases and reaches maximum at 2 T and then decreases and approaches zero. Since the AFM α -Fe₂O₃ is an insulator, this excludes the possibility of spin-dependent or spin-flip mechanisms that require electron spin carriers. Recently, magnonic USMR has been observed in the insulating ferromagnetic bilayer Pt/YIG [10]. To testify

the role played by magnons in the observed USMR, we perform the temperature dependent measurement as shown in Fig. 4(b). When the applied field is 2 T and the temperature decreases from 325 K, the USMR monotonically drops. At and below 200 K, no USMR is observed. This temperature dependence measurement provides the strong evidence for the magnonic origin of USMR [9].

Following the theory of magnon creation and annihilation imbalance in ferromagnet, we extend it to the antiferromagnetic regime. The coupled Landau-Lifshitz-Gilbert (LLG) equations for two sublattice magnetic moments m_A and m_B are written as [26]

$$\dot{\boldsymbol{m}}_{A} = -\gamma \boldsymbol{m}_{A} \times \boldsymbol{H}_{A}^{eff} - \gamma \boldsymbol{J}_{ex} \boldsymbol{m}_{A} \times \boldsymbol{m}_{B} + \alpha \boldsymbol{m}_{A} \times \dot{\boldsymbol{m}}_{A} + \gamma \boldsymbol{\tau}_{A}^{DL},$$
(5)

$$\dot{m}_{B} = -\gamma m_{B} \times H_{B}^{eff} - \gamma J_{ex} m_{B} \times m_{A} + \alpha m_{B} \times \dot{m}_{B} + \gamma \tau_{B}^{DL},$$
(6)



FIG. 4. (a) The extracted magnetic field dependence of USMR contribution in the $V_{xx}^{2\omega}$. (b) Temperature dependence of USMR in $V_{xx}^{2\omega}$ at 2 T. The error bars are extracted by fitting the data in Fig. 3.



FIG. 5. (a) Magnetic field dependence of antiferromagtic magnon number difference between $\varphi_H = \pm \pi/2$ and (b) between $\varphi_H = 0, \pi$. In the insets, the light color arrows (red and blue) represent the sublattice magnetic moments of AFM after rotating H_0 (green arrow) to the opposite direction. The magnetic fluctuation originates from the thermal random fields in the two orthogonal \hat{e}_{φ} and \hat{e}_r directions. σ_T^2 is the thermal coefficient.

where the effective field $H^{e\!f\!f}_{A(B)}=H_0+h_{{
m A}({
m B})}+H^{DMI}_{A(B)}+$ $H_{A(B)}^{hard}$ contains the external magnetic field H_0 , the thermal random field $h_{A(B)}(T)$, the effective field induced by Dzyaloshinskii–Moriya interaction (DMI) $H_{A(B)}^{DMI} =$ $H_D(\pm m_{B(A)} \times \hat{z})$ [27] and effective field of the hard axis anisotropy $H_{A(B)}^{hard} = 2H_{\perp}m_{A(B)}^{z}\hat{z}$. $J_{ex}(<0)$ is the AFM exchange coupling, γ is the gyromagnetic ratio (positive), and α is the Gilbert damping constant. $\tau_{A(B)}^{DL} = H_{DL}m_{A(B)} \times$ $(\hat{\boldsymbol{\sigma}} \times \boldsymbol{m}_{A(B)})$ is the damping-like (DL) torque that exerts on the unit sublattice magnetization $m_{A(B)}$, as shown in Fig. 2(a). Here, $\hat{\sigma}$ is the unit vector (along \hat{y} axis) of the spin polarization induced by the spin Hall effect in Pt with amplitude H_{DL} being linearly proportional to the charge current density. In our previous work, we have demonstrated that without the thermal random field, the DL torque induced a rotation of the sublattice magnetization (as well as the net magnetization *m* and Néel vector *n*) $\Delta \varphi_{\mathbf{A}(\mathbf{B})} = \Delta \varphi_m =$ $\Delta \varphi_n \propto H_{\rm DL}^2 \propto I^2$ [20]. Therefore, the induced voltage change $\propto I^3$, which cannot be detected in the second harmonic signal $V^{2\omega}$ but rather in third harmonic voltage $V^{3\omega}$, and it is not unidirectional. However, after considering the thermal random field $h_{A(B)}(T)$, the dampinglike torque induces a fluctuation of the sublattice magnetizations, which is now linear to the *I*, and unidirectional. The longitudinal SMR for a Pt/AFM insulator heterostructure can be presented as $\rho_L = \rho_0 - \Delta \rho \langle n_v^2 \rangle$ [5,6,21], where the contribution from net magnetization $-\Delta\rho \langle m_{\nu}^2 \rangle$ is negligible in the AFM regime. When an external magnetic field or applied current is reversed, the asymmetric magnon excitation driven by the thermal random field will result in a USMR in ρ_L [10]. For a current applied along the \hat{x} direction, the USMR signal reaches the maximum (minimum) at $\varphi_H = \pm (\pi/2)(\pm \pi)$ [Fig. 2(a)]. This can be explained by the fact that the induced net FM order is aligned along the external field $H(\varphi_H = \varphi_M)$. Therefore, **H** aligned along $\pm \hat{y}$ are inequivalent for magnon excitation when the spin polarization σ lies at $+\hat{y}$. However, when H is reversed between $\pm \hat{x}$, there is no asymmetry in magnon excitation. The USMR amplitude and antiferromagnetic magnon number difference are both proportional to the difference $\langle n_{y}^{2} \rangle_{+} - \langle n_{y}^{2} \rangle_{-}$ (± sign indicates two opposite directions). In the following, we refer to the term $\langle n_y^2 \rangle_+$ – $\langle n_{y}^{2} \rangle$ as "antiferromagnetic magnon number difference" for convenience since the trivial proportionality does not affect the physical picture [10]. We then numerically calculate the field dependence of antiferromagnetic magnon number difference for $\varphi_H = \pm (\pi/2)$ (see Sec. 1 in the Supplemental Material for more details [22]). We find that the field dependence of the antiferromagnetic magnon number difference qualitatively agrees with the nonmonotonic trend of the USMR signal and the peak around 2 T [Fig. 4(a)] is reproduced [Fig. 5(a)]. We also calculate the antiferromagnetic magnon number difference between $\varphi_H = 0, \pi$ in Fig. 5(b). As expected, we find that $\langle n_v^2(\varphi_H=0)\rangle - \langle n_v^2(\varphi_H=\pi)\rangle = 0$, which is consistent with the results from Fig. 2(a). The decrease of USMR at high fields is due to the suppression of magnon excitations at large H_0 , which is similar to the Pt/FM case. However, unlike FM where the magnetization simply saturates at small fields $(\boldsymbol{m} \| \boldsymbol{H}_0)$, the external field can increase the canting of sublattice magnetization in AFM $(n \perp H_0)$. Therefore, the magnetic fluctuation is governed by H_0 in a more complicated way. Specifically, the canting angle $\Delta \varphi$ for sublattice magnetizations of α -Fe₂O₃ depends on H_D , J_{ex} and H_0 as $\Delta \varphi = -\arcsin[(H_0 + H_D)/2J_{\text{ex}}]$. With the thermal random field **h** acting on the two orthogonal \hat{e}_{φ} and \hat{e}_{r} directions, the dynamical magnetic susceptibility χ that characterizes the fluctuation of Néel vector $\Delta n_y = \chi \cdot h$ contains a highly nontrivial H_0 dependence through $\Delta \varphi$, leading to the nonmonotonic field dependence of USMR in AFM. In Fig. S1(a) (See Supplemental Material [22] for details), we manually eliminate the H_0 dependence of $\Delta \varphi$ and plot the field dependence of magnon number difference. We find that the peak around 2 T disappears and the field dependence returns to be monotonic which is similar to the Pt/FM case. Therefore, the field-assisted canting, which is unique in AFM, increases the imbalance in the creation and annihilation of AFM magnons. However, on the other hand, total magnon number is suppressed with increasing field and the magnon excitation is essentially frozen when the field is large enough. These two competing effects give rise to the nonmonotonic field dependence of magnonic USMR. Finally, we compare the contributions with those of ferromagnetic magnon $\langle m_y^2 \rangle_+ - \langle m_y^2 \rangle_-$ as shown in Fig. S1(b) [22], which is 3 orders of magnitude smaller than that of antiferromagnetic magnons, emphasizing the dominant role of antiferromagnetic magnons in the observed USMR.

In summary, we observe the USMR in the antiferromagnetic heterostructure in Pt/α -Fe₂O₃ bilayers. The magnonic origin of USMR is revealed in the temperature and field dependent measurements. It is shown that the antiferromagnetic magnon plays the dominant role which gives a unique field dependence as compared with that of ferromagnetic materials. This first evidence of USMR in HM/AFI bilayers significantly expands our materials base to include the large family of AFM insulators and paves the way for the highly sensitive detection of AFM spin state in emerging the AFM spintronics through USMR.

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Note added.—Recently we became aware of an independent report [28] which detected UMR in metallic AFM bilayer Pt/FeRh. the UMR observed in Pt/FeRh is originated from the Rashba SOC at metal/AFM interface. Compared with the USMR observed in this work which only has the longitudinal term where Rashba type UMR should also have the corresponding transverse component with the magnitude $(V_{xx,Rashba UMR}^{2\omega}/V_{xy,Rahsba UMR}^{2\omega}) = (l/w)$. Thus, the Rashba SOC induced UMR could be mixed with the spin Seebeck signal in our analysis. Compared with metallic bilayers with almost no thermal effects due to a negligible temperature gradient, the spin Seebeck effect is very likely to overwhelm the Rashba UMR in our insulating α -Fe₂O₃ bilayers. *Corresponding author. cheng991@g.ucla.edu †Corresponding author. wang@ee.ucla.edu

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Correction: A production processing flaw rendered the setting of two diacritics in Eqs. (5) and (6) in the PDF version incorrectly. These diacritics have been fixed and were set without incident in the HTML version.