

Spin Seebeck Imaging of Spin-Torque Switching in Antiferromagnetic Pt/NiO Heterostructures

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As electrical control of Néel order opens the door to reliable antiferromagnetic spintronic devices, understanding the microscopic mechanisms of antiferromagnetic switching is crucial. Spatially resolved studies are necessary to distinguish multiple nonuniform switching mechanisms; however, progress has been hindered by the lack of tabletop techniques to image the Néel order. We demonstrate spin Seebeck microscopy as a sensitive tabletop method for imaging antiferromagnetism in thin films and apply this technique to study spin-torque switching in Pt/NiO and Pt/NiO/Pt heterostructures. We establish the interfacial antiferromagnetic spin Seebeck effect in NiO as a probe of surface Néel order. By imaging before and after applying current-induced spin torque, we resolve spin domain rotation and domain wall motion. We correlate the changes in spin Seebeck images with electrical measurements of the average Néel orientation through the spin Hall magnetoresistance, confirming that we image antiferromagnetic order.

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I. INTRODUCTION

Antiferromagnets (AFs), long relegated to a supporting role as the pinning layers in ferromagnetic spintronic devices [1,2], are emerging as the active element in antiferromagnetic spintronic devices [3–5]. In contrast to ferromagnets (FMs), AFs are insensitive to magnetic fields [6] and exhibit dynamics at the terahertz frequency scale [7,8]. Additionally, AFs have magnetotransport effects that enable electrical readout [9,10]. Taking advantage of these attractive properties, however, requires overcoming the challenge of reliably manipulating Néel order.

Recent breakthroughs in electrical [11,12] and optical [13,14] control provide a path toward reliable devices.

In particular, electrical switching was demonstrated in the metals CuMnAs [11,15] and Mn₂Au [16] using Néel spin-orbit torque, in which the sign of the spin-orbit field from a dc current within the material alternates on each lattice site to coherently rotate the Néel vector [17]. Recently, electrical switching of an AF via spin torque was also demonstrated in insulating NiO [18–20] after several predictions [21,22]. In this mechanism, the dc current passing through an adjacent Pt layer generates a spin current through the spin Hall effect, which then exerts an antidamping torque on the spins at the Pt/NiO interface. Switching by antidamping spin torque does not require that the spin sublattices form inversion partners, which is required for Néel spin-orbit torque; hence, it is a more general approach that could enable all-electrical control over a wider variety of AFs.

Previous experiments have shown that AF switching is nonuniform [15,23,24] and heavily influenced by local magnetoelastic stresses [25]. Nominally identical samples display switching efficiency that varies by almost an order of magnitude at the same current density [18], demonstrating a need for better understanding the switching process at

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the domain level. Systematic spatially resolved studies are necessary to firmly establish the spin rotation mechanisms, the fraction of the domains that switch, and the reproducibility of switching.

A primary challenge when imaging antiferromagnetism is to find an experimental probe that is sensitive to Néel order and also provides the submicrometer resolution necessary to resolve domains. X-ray magnetic linear dichroism photoemission electron microscopy (XMLD-PEEM) has been the most reliable technique [23,26]; however, it requires a coherent x-ray source that is available at only a few facilities. Second-harmonic [27,28] and quadratic magneto-optical techniques [29,30] are available in a tabletop format, but the small signal sizes create a need for background subtraction, which can be a problem because antiferromagnets are difficult to fully saturate. Scanning-probe magnetometers are also available in tabletop format but require stray fields to be produced from the AF [31]. As an alternative, recent demonstrations of the AF anomalous Nernst effect [32] and AF spin Seebeck effect [33,34] open up the possibility of using spin-thermal effects as an imaging probe, because they can be directly sensitive to Néel order [3]. Previous work from both our group and others demonstrates high-sensitivity imaging of ferromagnetic order via the anomalous Nernst and longitudinal spin Seebeck effects [35–39], suggesting that a practical and sensitive magnetothermal microscope for Néel order can also be developed.

In this work, we use antiferromagnetic longitudinal spin Seebeck effect (AF LSSE) microscopy to image spin-orbit torque switching in Pt/NiO(111) bilayers and Pt/NiO(111)/Pt trilayers. We provide the first experimental demonstration of interfacial AF LSSE and use it as a direct probe of the Néel order to resolve 1–10- μm -size antiferromagnetic spin domains. By repeatedly imaging before and after spin-torque switching while varying the

current density and direction in a variety of samples, we reveal effects of antidamping spin torque on the Néel order of NiO that would be difficult to establish either with device-level transport measurements or with limited beam time at an XMLD-PEEM facility. In particular, we show that switching occurs simultaneously by continuous rotation of the Néel orientation within AF domains and by current polarity-dependent AF domain wall motion.

The organization of this paper is as follows: We discuss the antiferromagnetic domain structure in NiO and present initial SSE images. We then establish the interfacial AF LSSE as the source of our signal, and we study spin-torque-induced domain rotation and domain wall motion.

II. IMAGING NÉEL ORDER WITH SPIN SEEBECK MICROSCOPY

A. Resolving antiferromagnetic domains in NiO

NiO is a collinear insulating antiferromagnet with a Néel temperature T_N of 523 K in the bulk [40]. Superexchange between Ni atoms along the $\langle 100 \rangle$ directions aligns the spins in ferromagnetic $\{111\}$ planes, in which spins on one plane are antiparallel to spins on the adjacent plane [26]. In the bulk, the AF domain structure is well known: Magnetostriction along $\langle 111 \rangle$ from the AF ordering causes crystallographic twinning, forming four T (twin) domains [41]. Within each T domain, dipolar next-nearest-neighbor coupling introduces a weak additional in-plane anisotropy along the three equivalent $[11\bar{2}]$ directions, forming three S (spin) domains per T domain [42,43]. In thin films, however, magnetoelastic stresses, the AF equivalent of the demagnetization field in FMs [25], introduce an additional effective anisotropy. This spatially inhomogeneous anisotropy pulls the spins out of well-defined $[11\bar{2}]$ directions, resulting in a disordered in-plane AF domain structure [44].

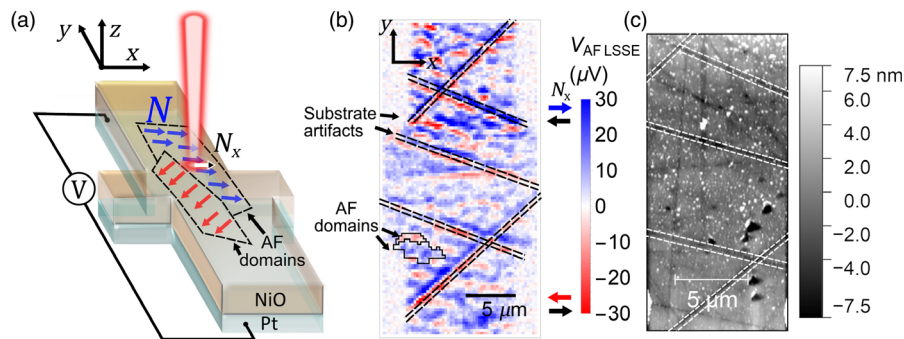


FIG. 1. Demonstration of AF LSSE microscopy. (a) Schematic of the measurement. A Ti:sapphire laser focused to 650 nm spot size thermally generates a local spin current J_s at the Pt/NiO interface, with spin polarization σ parallel to the local Néel orientation N . The sign of σ is determined by the spin direction of the uncompensated monolayer. J_s is transduced into a charge current via the inverse spin Hall effect in the Pt, resulting in a voltage between the contacts. (b) AF LSSE image of a $10 \mu\text{m} \times 50 \mu\text{m}$ Hall cross of MgO/5 nm Pt/7 nm NiO(111). Blue (red) contrast represents interfacial spins pointing right (left). Sharp straight lines highlighted by black dashed line are artifacts from the ordinary Seebeck effect, which may be due to scratches in the MgO substrate. (c) AFM height map of the same sample. Ordinary Seebeck artifacts in the AF LSSE image are accompanied by 1-nm-deep valleys in the height.

We resolve the spin domains in Pt/NiO bilayers and Pt/NiO/Pt trilayers with spin Seebeck effect microscopy [39] using a geometry illustrated in Fig. 1(a). We focus 3-ps-wide, 785 nm wavelength pulses from a Ti:sapphire laser down to a 650 nm spot size, which produces a local out-of-plane thermal gradient. We show in Supplemental Material [45] that the thermal gradient is dominated by an interfacial temperature drop ΔT at the Pt/NiO interfaces. In Sec. II B, we establish that ΔT generates a spin current \mathbf{J}_s with polarization $\boldsymbol{\sigma}_{\text{int}}$ parallel to the orientation \mathbf{s}_{int} of the closest monolayer of spins to the Pt interface. Within the Pt, the spin current is transduced into a charge current $\mathbf{J}_c \propto \mathbf{J}_s \times \boldsymbol{\sigma}_{\text{int}}$ via the inverse spin Hall effect (ISHE), which results in a voltage drop across the sample. By raster scanning the focused laser over the sample, we build a map of the ISHE voltage, which reports the in-plane component of \mathbf{s}_{int} perpendicular to the voltage contacts.

An example AF LSSE image of an epitaxial MgO/5 nm Pt/7 nm NiO(111) device, deposited by sputtering and patterned into a $10 \mu\text{m} \times 50 \mu\text{m}$ Hall cross by optical photolithography, is shown in Fig. 1(b). Blue and red contrast show interfacial spins pointing right and left, respectively. We acquire all images at room temperature and a zero magnetic field using 3.4 mJ/cm^2 laser fluence. By combining finite-element simulations of laser heating with electrical calibration using transient resistivity changes, we measure that this laser fluence locally heats the Pt and NiO layers to 360 and 330 K peak temperature, respectively, from the ambient temperature of 293 K. Note that the voltage we plot—denoted $V_{\text{AF LSSE}}$ —is not the actual spin Seebeck voltage but rather a lock-in voltage after amplification, mixing, and normalization to account for impedance matching, which is also described in Supplemental Material [45].

Sharp straight lines in the AF LSSE image in Fig. 1(b), highlighted with black dashed lines, are also visible in the corresponding atomic force microscopy height map in Fig. 1(c) and may be due to polishing scratches in the MgO(111) substrate. Thermal discontinuities at these scratches cause artifacts in the AF LSSE signal from the ordinary Seebeck effect and are discussed further in Supplemental Material [45]. The rest of the contrast in the AF LSSE image in Fig. 1(b) represents antiferromagnetic spin domains: Examples are highlighted in black enclosures. Since the Pt is beneath the NiO in this sample and the voltage contacts are along y , in Fig. 1(b), we measure the N_x component of the bottom interfacial monolayer $\mathbf{s}_{\text{bottom}}$.

Currently, we cannot saturate the Néel vector along a given direction while imaging, which would require applying an *in situ* magnetic field greater than the spin-flop field (5 T in NiO [46]). Therefore, we cannot calibrate the AF LSSE voltage to an absolute Néel orientation. This limitation is an intrinsic difficulty of detecting antiferromagnetism and is also experienced by other

imaging techniques, including XMLD-PEEM. Instead, the AF LSSE voltage represents the strength of the projection without absolute calibration. The size, shape, and distribution of AF domains are consistent with previous XMLD-PEEM imaging studies of thin-film NiO [44,47]. Note that only domains larger than the 650 nm laser spot size are resolved; some contrast in Fig. 1(b) extends down to the single-pixel limit and may represent incompletely resolved domains.

B. Evidence for interfacial antiferromagnetic LSSE

We attribute the signal in our images to an interfacial AF LSSE. Although the ferromagnetic LSSE is well established both in the bulk [48–50] and at the interface [51–53], the AF LSSE was initially predicted not to exist for a collinear AF [54] and was only recently observed [33,34]. In a collinear AF, the two spin sublattices produce two degenerate magnon modes [55], which produce spin current in opposite directions under a thermal gradient. Therefore, unless the degeneracy is lifted, there is no net spin current [54]. The degeneracy can be lifted in the AF bulk by applying a large magnetic field [33,34] or by exploiting anisotropies that result in additional magnon modes [56,57]. The degeneracy can also be lifted by inversion symmetry breaking at the interface, resulting in an interfacial AF LSSE that has been predicted [58] but has not been previously reported.

The mechanism of the interfacial AF LSSE is schematically illustrated in Figs. 2(a) and 2(b). AF interfaces can be *uncompensated* as in Fig. 2(a), meaning the layer closest to the surface contains an excess of one sublattice, or *compensated* as in Fig. 2(b), in which adjacent spins are antiparallel in each growth plane. In the uncompensated case, an interfacial temperature difference ΔT across the surface normal $\hat{\mathbf{n}}$ induces a local spin current $\mathbf{J}_s \parallel \Delta T \hat{\mathbf{n}}$ whose polarization effectively alternates in sign at each monolayer, following the local spin polarization. The layer closest to the Pt interface— \mathbf{s}_{int} —is more strongly exchange coupled to the spins in the Pt layer than the other layers are [59], and, therefore, a net spin current \mathbf{J}_s diffuses into the Pt with polarization $\boldsymbol{\sigma} \parallel \mathbf{s}_{\text{int}}$ [58]. Note that \mathbf{s}_{int} is \mathbf{s}_{top} if the Pt is above the NiO and $\mathbf{s}_{\text{bottom}}$ if the Pt is beneath the NiO.

At the compensated interface, illustrated in Fig. 2(b), the symmetry between magnon modes is preserved. Therefore, the spin current produced by ΔT is zero on every $\langle 111 \rangle$ plane as long as the in-plane spatial extent of ΔT (650 nm for the laser spot) is much greater than the in-plane lattice spacing [0.4 nm in NiO(001)]. Although the uncompensated interface in the schematic in Fig. 2(a) is atomically flat, the presence of roughness in real samples does not alter the interpretation of the AF LSSE signal as long as the lateral length scale of average height variation is also much smaller than the laser spot diameter (see Supplemental Material [45] for more details).

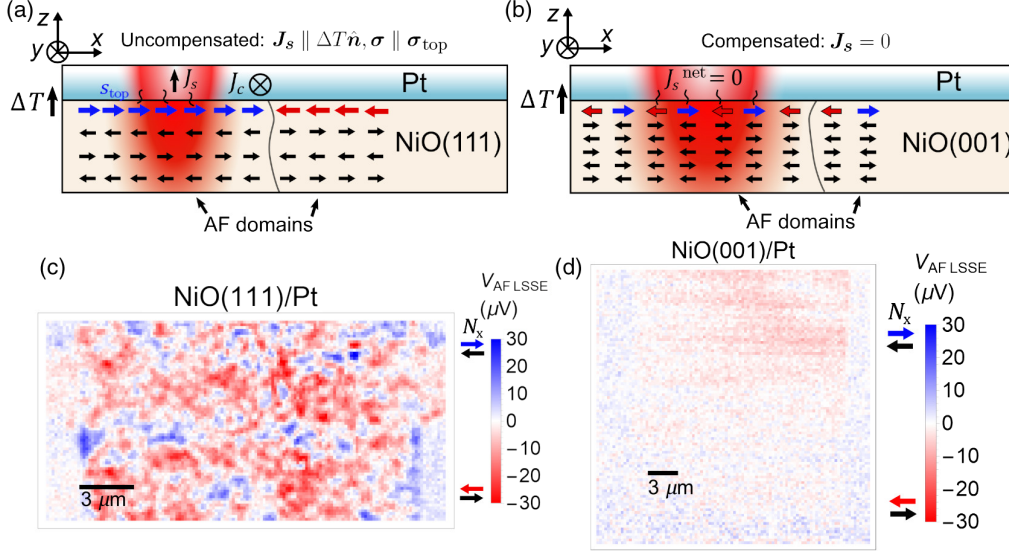


FIG. 2. Mechanism of the interfacial AF LSSE. (a) Schematic illustrating the interfacial AF LSSE at an uncompensated interface. A thermal gradient produces spin current J_s with polarization σ parallel to the spin orientation of the interfacial uncompensated monolayer (s_{top} in the diagram). (b) Schematic illustrating the lack of AF LSSE at a compensated interface. The net spin current produced by every monolayer is zero. (c), (d) AF LSSE images of uncompensated MgAl_2O_4 (111)/165 nm $\text{NiO}(111)$ /6 nm Pt and compensated $\text{MgO}(001)$ /136 nm $\text{NiO}(001)$ /6 nm Pt, respectively. The lack of $V_{\text{AF LSSE}}$ signal from $\text{NiO}(001)$ compared to $\text{NiO}(111)$ indicates that $V_{\text{AF LSSE}}$ originates from the uncompensated interface.

Our experimental test of interfacial AF LSSE is shown in Figs. 2(c) and 2(d). We take AF LSSE images of MBE-grown MgAl_2O_4 /165 nm $\text{NiO}(111)$ /Pt and $\text{MgO}(001)$ /136 nm $\text{NiO}(001)$ /Pt, which have uncompensated and compensated interfaces, respectively. Both samples are patterned into 20- μm -wide Hall bars with similar growth conditions, sample resistivities, and interface qualities as shown in Supplemental Material [45]. Since the Pt is above the NiO, we measure s_{top} here. We find that AF LSSE images of $\text{NiO}(001)$ yield an order of magnitude lower voltage compared with $\text{NiO}(111)$. The residual contrast in $\text{NiO}(001)$ images is nearly uniform and does not resemble domains; it may be due to capacitive coupling or other small experimental artifacts. Although the preferred spin orientation in $\text{MgO}(001)$ / $\text{NiO}(001)$ is not well established [19,20], recent XMLD-PEEM and magneto-optical images of 10-nm-thick $\text{NiO}(001)$ on $\text{MgO}(001)$ indicate approximately 10° out-of-plane tilt [30], which would reduce $V_{\text{AF LSSE}}$ by 2%. If the spins in our 136-nm-thick $\text{NiO}(001)$ are relaxed to the bulk orientation, they would point 35° out of plane (the angle between $\{11\bar{2}\}$ and $\{001\}$). This result would reduce $V_{\text{AF LSSE}}$ by 20%, which is still not enough to account for the order-of-magnitude difference. Therefore, our results are consistent with a model of an uncompensated interface. Our results also indicate that bulk AF LSSE [56,60] does not significantly contribute to our signal. This interpretation is further supported by finite-element simulations, discussed in Supplemental Material [45], which show that the laser-induced thermal profile is dominated by temperature drops at the Pt/NiO interfaces rather than a temperature gradient in the NiO bulk.

We can further distinguish between an AF LSSE at an uncompensated interface and a possible FM LSSE originating from pinned uncompensated moments (UMs) [61]. These UMs could arise from interfacial roughness [62] or defects, both at the interface and in the bulk of the AF [63], and would be detectable by other magnetometry techniques. Therefore, we perform scanning SQUID microscopy at 4 K to search for microscopic moments and polarized neutron reflectometry at room temperature to detect a global moment at 1 T applied field. Both sets of measurements are made on sputtered samples and are described in more detail in Supplemental Material [45]. We find no magnetic moment within sensitivity, which places an upper limit of $8 \times 10^{-4} \mu_B/\text{Ni}$ on the local moment that could be present. From this value, we calculate a maximum bulk magnetization of 110 A/m in our NiO, 3 orders of magnitude less than the bulk magnetization in YIG at room temperature. Based on these measurements, we rule out a FM LSSE and conclude that the AF LSSE signal in NiO arises from the uncompensated interface.

III. IMAGING SPIN-TORQUE SWITCHING IN Pt/NiO/Pt TRILAYERS

Having established a mechanism for the signal contrast, we move onto image current-induced spin-torque switching in Hall crosses, initially following the procedure in Refs. [18,19]. We apply a dc writing current and characterize the Néel state electrically, using the antiferromagnetic analog of spin Hall magnetoresistance (SMR)

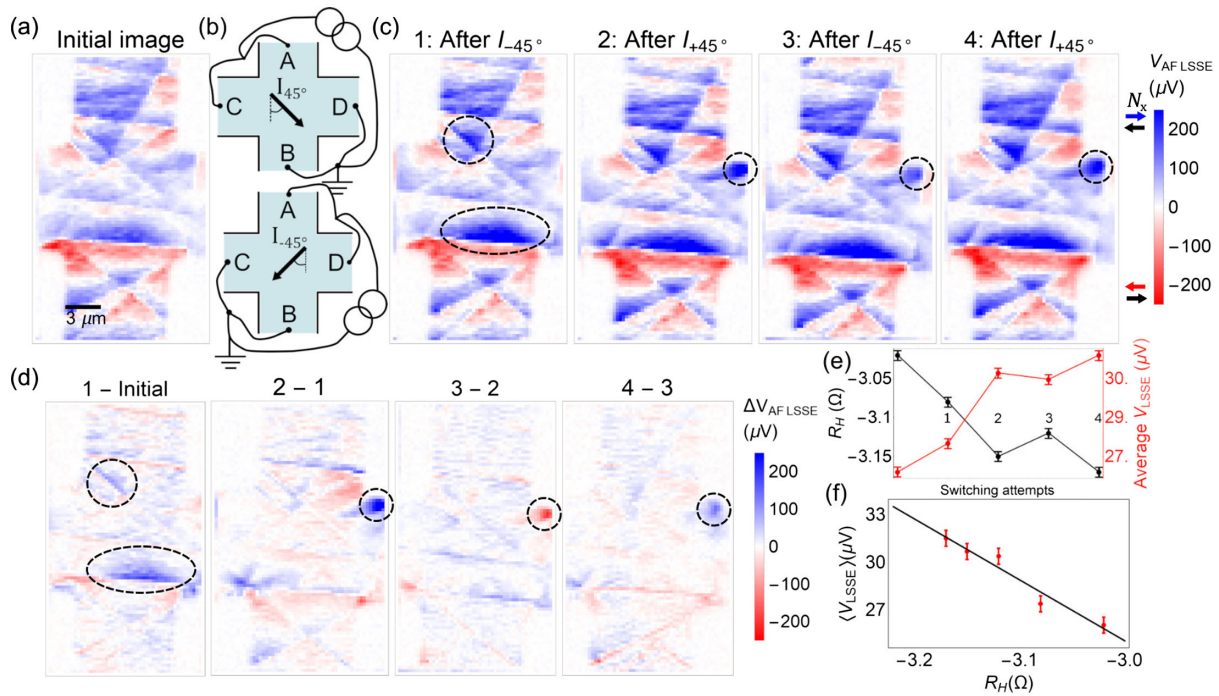


FIG. 3. AF LSSE imaging of spin-torque switching in a 5 nm Pt/13 nm NiO(111)/5 nm Pt trilayer. (a) The initial image, taken before applying the current. (b) Schematic of the writing process. We apply a current to two arms of the Hall cross such that the current density in the center flows along 45° diagonals. (c) Imaging while toggling between I_{45° and I_{-45° . Highlighted in dashed lines are changes in contrast at the top right corner, where the current density is greatest, and near apparent substrate scratches, where the spins may be less strongly exchange coupled. (d) Successive differences between the AF LSSE images in (a), showing the domains that switch more clearly. Both positive and negative contrast in difference images may reflect domains rotating in opposite directions. (e) R_H and the integrated AF LSSE signal $\langle V_{AF\ LSSE} \rangle$, measured while toggling between I_{45° and I_{-45° . Values corresponding to the images shown are labeled. (f) $\langle V_{AF\ LSSE} \rangle$ plotted as a function of SMR, measured through changes in the Hall resistance R_H . The near-linear correlation shows the small-angle correspondence between $\langle V_{AF\ LSSE} \rangle$ and R_H .

[64–66] by measuring the change in the Hall resistance R_H : $R_H = -\Delta R_{SMR} \sin \theta \cos \theta$, where θ is the angle between the spatially averaged Néel vector N and the reading current J_R . To maximize ΔR_H , we apply a writing current to two of the arms of the Hall cross such that the current density in the center of the device flows along $\pm 45^\circ$ [schematically illustrated in Fig. 3(b)]. Using a finite-element simulation described further in Supplemental Material [45], we estimate a writing current density of 8.0×10^7 A/cm² at the corners and 3.1×10^7 A/cm² in the center of the cross. Hereafter, we refer only to the density in the center of the cross, J_W . After each application of writing current, we measure the Hall resistance R_H by applying a reading current density $J_R = 1.5 \times 10^6$ A/cm² from A to B and measuring the voltage from C to D in Fig. 3(b).

We initially employ epitaxially sputtered 5 nm Pt/13 nm NiO(111)/5 nm Pt trilayers, following the argument of Ref. [18] (further demonstrated in Ref. [67] in synthetic antiferromagnets) that spin torque at both the top and bottom interfaces of the NiO leads to more coherent rotation of the Néel orientation throughout the AF thickness. There is a potential complication interpreting the AF LSSE images in

trilayers, because both Pt/NiO interfaces can contribute to the signal; however, we show in Supplemental Material [45] that the presence of two simultaneously contributing interfaces does not alter the interpretation of $V_{AF\ LSSE}$ as long as the AF domains are continuous in thickness, which we expect for our trilayer samples. The AF domains in the trilayer are substantially larger (5–10 μm) than the domains in the bilayer in Fig. 1 (submicrometer to 2 μm), which most likely is due to differing growth conditions but which requires further study.

AF LSSE images of a trilayer before and after four sequential applications of $J_W = 3.1 \times 10^7$ A/cm² are shown in Figs. 3(a) and 3(c), with the writing current direction alternating between $+45^\circ$ and -45° . Although most domains are unaltered, we observe changes in contrast (highlighted inside the black dashed enclosures) at a sample corner, where the current density is highest, and near apparent substrate scratches, where the spins may be less strongly exchange coupled. To quantify these changes to the Néel orientation, we calculate sequential differences between images, shown in Fig. 3(d). We observe both positive and negative changes in contrast in different parts of the sample, which could be due to different AF domains

rotating in opposite directions, as seen in imaging studies of switching in CuMnAs [23]. At the current density used, we estimate that the maximum Oersted field is approximately 10 mT. We show in Supplemental Material [45] that the AF LSSE signal is unaffected by field up to ± 250 mT, which rules out the Oersted field from the writing current as the mechanism responsible for switching.

To compare AF LSSE imaging with electrical measurements of Néel order using R_H , we take the average of all the pixels in and near the cross center in each image (described in Supplemental Material [45]) to obtain the integrated AF LSSE signal $\langle V_{AF\ LSSE} \rangle$. Although $\langle V_{AF\ LSSE} \rangle$ and R_H are both measures of the average Néel orientation in the cross center, they have different symmetries: $R_H \propto \cos \theta \sin \theta$, where θ is the angle between the average Néel vector and the SMR reading current \mathbf{J}_R , while $\langle V_{AF\ LSSE} \rangle \propto \cos \phi$, where ϕ is the angle between the average Néel vector and the voltage contacts. Since we apply \mathbf{J}_R along x in this device, here $\theta = \phi - \pi/2$ and $\langle V_{AF\ LSSE} \rangle \propto -\sin \theta$. In this sample, most of the changes in contrast occur where N appears to be nearly saturated in the $+x$ direction, so that locally $\theta \approx 0$. Near $\theta = 0$, $\sin \theta \cos \theta \approx \sin \theta$. Therefore, $\langle V_{AF\ LSSE} \rangle$ tracks R_H point by point, shown in Fig. 3(e). Plotting one versus the other yields a near-linear correlation with a negative slope, shown in Fig. 3(f) with a linear fit drawn as a guide to the eye. This correspondence indicates that changes in contrast indeed represent antiferromagnetic switching.

We expect that AF switching can occur either by domain rotation, which would manifest in the AF LSSE images as changes of contrast level within domains, or by domain wall motion. The switching in Fig. 3 manifests as changes in contrast within domains while domain walls remain stationary within the resolution limit, which indicates domain rotation. In this sample, we observe changes in color shade but not changes in sign of N_x (blue to red or vice versa), which indicates that N rotates by acute angles. Although we cannot obtain an absolute angle of rotation, we can obtain a lower bound by taking the maximum and minimum $V_{AF\ LSSE}$ to correspond to $\theta = 90^\circ$ and -90° , respectively. In this case, we estimate that the average Néel vector at the corner rotates 22° between images 1 and 2 in Fig. 3(c) and 10° between images 2 and 3.

Previous studies of magnetic-field-induced domain rotation in 120-nm-thick NiO [66] model switching as 120° flopping between $\langle 11\bar{2} \rangle$ in-plane easy axes. In our samples, however, the domains have random in-plane orientation, which is consistent with XMLD-PEEM images of similar Pt/NiO/Pt trilayers [18]. This domain configuration is consistent with an increased role of magnetoelastic stress in our 6-nm-thick samples, which favors a multidomain state with zero average strain. While the effective field from in-plane anisotropy is $H_{Az} = 11$ mT in bulk NiO [57], the destressing field reported in 120-nm-thick NiO in Ref. [66] is 46 mT. We expect the destressing field to be even higher

in 6 nm NiO. Therefore, because the spins are not restricted to the $\langle 11\bar{2} \rangle$ axes in our samples, they can switch by continuous in-plane rotation.

IV. RESOLVING DOMAIN ROTATION AND DOMAIN WALL MOTION IN MgO/Pt/NiO BILAYERS

After correlating the AF LSSE images with an electrical readout of the Néel order through SMR in Pt/NiO/Pt trilayers, we move on to imaging switching in the sputtered NiO(111)/Pt bilayer from Fig. 1 after applying a current along the device channel. The bilayer does not have the potential difficulty of superposing signal from two Pt/NiO interfaces. Furthermore, applying a current along the device channel yields more uniform current density, leading to larger-scale, more easily resolvable changes in image contrast. Figure 4(a) shows AF LSSE images before switching and then after applying progressively greater current densities, from 5.0×10^7 A/cm² at 20 mA to 1.1×10^8 A/cm² at 42 mA, first at positive polarity (flowing down) and then negative polarity (flowing up). Prominent regions of switching are highlighted in the black enclosure as a guide to the eye.

Because the switching in Fig. 4 is spatially distributed and nonuniform, and AF LSSE and SMR have different symmetries, we cannot correlate the AF LSSE signal with SMR like we do in Fig. 3. Therefore, we compare our AF LSSE images to theoretical models of switching. References [18,19] model switching in the high-current limit as coherent rotation of spins within an AF domain (domain rotation), which we observe in Fig. 3. The model in Ref. [20] distinguishes three separate switching mechanisms, with predictions summarized as follows.

- (1) The out-of-plane component of the spin current rotates spins within the easy plane, rotating all AF domains by the same angle.
- (2) The in-plane component of the spin torque creates an additional effective anisotropy, resulting in a translational ponderomotive force \mathbf{F}_{pond} proportional to \mathbf{J}_W^2 on the AF domain wall. \mathbf{F}_{pond} rotates $N \parallel \mathbf{J}_W$ and is current polarity independent.
- (3) The spin torque directly rotates the spins within the domain walls, leading to a chiral domain wall force \mathbf{F}_{DW} that goes as $\mathbf{J}_W \cdot \mathbf{F}_{\text{DW}}$. \mathbf{F}_{DW} can rotate N either towards or perpendicular to \mathbf{J}_W , depending on the domain wall chirality; therefore, it should have no net effect on R_H with randomly oriented domains. \mathbf{F}_{DW} should also reverse direction when \mathbf{J}_W reverses.

To quantitatively characterize the switching, we take sequential image differences in Fig. 4(b). Cumulative difference images as well as AF LSSE images after +20 and -42 mA are given in Supplemental Material [45]. After applying 30 mA, as seen in image 2-1, we see large-scale, nearly uniform positive (blue) contrast in the lower

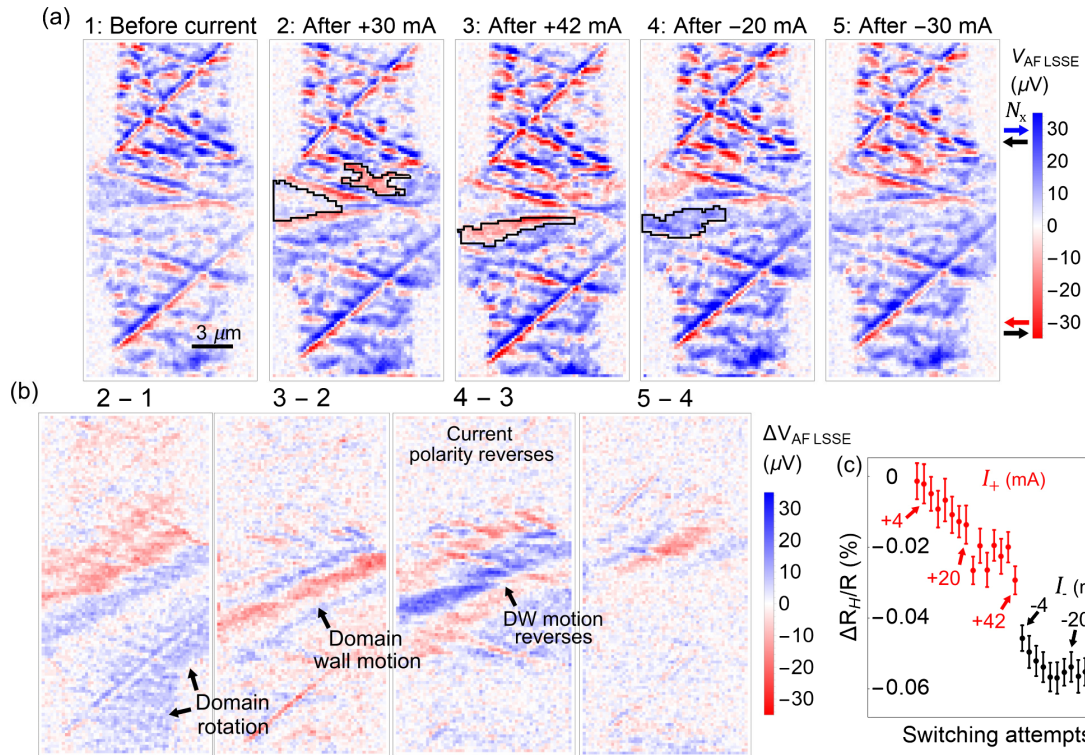


FIG. 4. Switching in the Pt/NiO bilayer from Fig. 1 after applying a current along the channel direction. (a) AF LSSE images before and after applying current. We apply first a positive current, flowing down, and then a negative current, flowing up. Some prominent regions of switching are highlighted in the black line, including domain growth after +42 mA and domain wall motion after -20 mA. (b) Sequential differences from the initial state (1). Large-scale patterns of nearly uniform positive contrast in the lower portion of 2–1 show different AF domains rotating by the same angle. We observe domain wall motion after +42 mA that reverses direction when the current polarity is reversed, consistent with theoretical predictions of a chiral domain wall force F_{DW} . (c) SMR measurements of the average Néel orientation for a similar Pt/NiO bilayer after applying the same currents, first positive and then negative (labeled in mA). R_H does not depend on current polarity, which is consistent with domain wall motion due to F_{DW} .

half of the cross, labeled *domain rotation*. More uniform contrast in the difference images than in the AF LSSE images themselves indicates that different AF domains are rotating by the same angle, consistent with the out-of-plane spin torque in mechanism 1 in Ref. [20]. This domain rotation saturates after +30 mA (7.5×10^7 A/cm²) and does not reverse when the current polarity reverses. We primarily observe $|N_x|$ increasing—blue domains become more blue and red domains more red—which means N rotates $\perp \mathbf{J}_W$.

From images 2 and 3 in the AF LSSE images in Fig. 4(a), we resolve switching by domain wall motion, which appears as a negative (red) horizontal stripe in the sequential 3–2 image in Fig. 4(b). Domain rotation and domain wall motion occur in response to a writing current as low as 4 mA (1.0×10^7 A/cm²). Interestingly, we find that the domains continue to move for 2–3 hr after the current is turned off (shown in Supplemental Material [45]), which may be due to magnetoelastic stresses causing subthreshold domain wall creep after the spin torque rotates the domains out of equilibrium. Although the domain configuration after +42 mA in 3 does not creep in time, the domain wall

motion reverses after applying -20 mA, seen in 4–3, and subsequently almost ceases, seen by weaker contrast in 5–4 after applying -30 mA. DW motion that reverses when the current polarity is reversed points to the chiral force F_{DW} as the origin. In Fig. 4(c), we show SMR measurements of a similar MgO/Pt/NiO cross while applying the same currents, first positive and then negative. We find that R_H does not depend on current polarity, which is consistent with the prediction that the effects of F_{DW} would not be reflected by changes in R_H .

Summarizing our results, we identify both domain rotation and domain wall motion acting simultaneously, which are consistent with the out-of-plane spin torque and chiral domain force, respectively, described in Ref. [20]. At the current densities applied, from 1.0×10^7 to 1.0×10^8 A/cm², we do not observe N rotating towards \mathbf{J}_W from the ponderomotive force \mathbf{F}_{pond} , which is expected to dominate at higher current densities (above $7\text{--}9 \times 10^7$ A/cm², depending on the strain). Further imaging studies on thicker NiO samples with less strain may be required to observe \mathbf{F}_{pond} .

Our results complement the XMLD-PEEM images of switching in Refs. [18,20]. Although Ref. [18] shows domain wall motion and Ref. [20] appears to show domain rotation in response to current, distinguishing several simultaneously acting switching mechanisms requires systematic repeated imaging of multiple samples, which may not be practical with the limited beam time at a synchrotron facility.

V. CONCLUSION

In conclusion, we demonstrate interfacial AF LSSE as the basis for a powerful tabletop technique for imaging in-plane Néel order in an AF insulator. This magnetothermal microscope uses equipment that is readily available in many laboratories, thus enabling in-depth and high-throughput studies of AF spintronics, which was previously limited by the availability of a few coherent x-ray facilities. Using this capability, we probe the microscopic behavior of spin-torque switching of Néel order in Pt/NiO/Pt trilayers and Pt/NiO bilayers. We find that switching occurs by domain rotation and domain wall motion acting simultaneously and that magnetoelastic stresses play an important role in determining both the equilibrium domain structure and the fraction of domains that switch. These insights provide critical understanding of spin-torque switching in NiO and point the way towards systematic optimization of antiferromagnetic spintronic devices. Moreover, we expect AF LSSE microscopy to extend to a wide variety of antiferromagnetic insulators with uncompensated interfaces, which can aid in the development of new device technologies based on different antiferromagnets.

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