

Effect of Interface-Induced Exchange Fields on Cuprate-Manganite Spin Switches

Yaohua Liu,^{1,*} C. Visani,^{2,†} N. M. Nemes,² M. R. Fitzsimmons,³ L. Y. Zhu,¹ J. Tornos,² M. Garcia-Hernandez,⁴ M. Zhernenkov,^{3,‡} A. Hoffmann,¹ C. Leon,² J. Santamaria,² and S. G. E. te Velthuis^{1,§}

¹Materials Science Division, Argonne National Laboratory, Argonne, Illinois 60439, USA

²GFMC, Departamento de Física Aplicada III, Universidad Complutense de Madrid, Campus Moncloa, ES-28040 Madrid, Spain

³Los Alamos National Laboratory, Los Alamos, New Mexico 87545, USA

⁴Instituto de Ciencia de Materiales de Madrid, Consejo Superior de Investigaciones Científicas, ES-28049 Cantoblanco, Spain

(Received 10 January 2012; published 18 May 2012)

We examine the anomalous inverse spin switch behavior in $\text{La}_{0.7}\text{Ca}_{0.3}\text{MnO}_3$ (LCMO)/ $\text{YBa}_2\text{Cu}_3\text{O}_{7-\delta}$ (YBCO)/LCMO trilayers by combined transport studies and polarized neutron reflectometry. Measuring magnetization profiles and magnetoresistance in an in-plane rotating magnetic field, we prove that, contrary to many accepted theoretical scenarios, the relative orientation between the two LCMO's magnetizations is not sufficient to determine the magnetoresistance. Rather the field dependence of magnetoresistance is explained by the interplay between the applied magnetic field and the (exponential tail of the) induced exchange field in YBCO, the latter originating from the electronic reconstruction at the LCMO/YBCO interfaces.

DOI: 10.1103/PhysRevLett.108.207205

PACS numbers: 75.25.-j, 74.78.Fk, 75.47.-m, 75.70.Cn

Interfacial electronic reconstruction offers the possibility to engineer the electronic ground state with unprecedented access to exotic phenomena at epitaxial interfaces of complex oxide heterostructures such as metallicity, superconductivity (SC) and even ferromagnetism (FM) at the interface of two insulating and nonmagnetic oxides [1–3]. Another example is the interface between half-metallic ferromagnet (FM) $\text{La}_{0.7}\text{Ca}_{0.3}\text{MnO}_3$ (LCMO) and high T_C superconductor (SC) $\text{YBa}_2\text{Cu}_3\text{O}_{7-\delta}$ (YBCO), where electronic reconstruction yields an antiferromagnetic coupling between the Cu and Mn's spins [4], which generates an interface induced ferromagnetic exchange field on the Cu ions in YBCO. This induced exchange field in YBCO then gives rise to a net Cu moment, as has been experimentally observed [5–7].

LCMO/YBCO/LCMO (LYL) trilayers are of interest as they are high- T_C superconducting spin switches, yet exhibit the so-called “inverse” superconducting spin switch behavior. It has been shown that, in the superconducting transition region, LYL trilayers have lower resistances when the magnetizations of two ferromagnetic layers are parallel, and higher resistances when they are antiparallel [8]. Consequently, T_C is higher for the parallel state and lower for the antiparallel state, which is opposite to the expectation based on the conventional proximity effect [9,10]. The origin of the inverse spin switch behavior is still controversial. Possible mechanisms include the effect of stray fields [11,12], an imbalance of quasiparticles [13,14], and triplet superconductivity [15]. In these scenarios, the magnetoresistance depends on the relative magnetic alignment between the two ferromagnets, and the applied field direction only plays an indirect role by changing the magnetization configuration [9,10,13,16]. Alternatively, Salafranca and Okamoto have recently proposed a scenario that can explain the inverse superconducting spin switch effect in

LYL trilayers, in which the direction of the applied field plays a direct role. They argue that the superconductivity in the central YBCO is governed by the total field \vec{H}_{tot} that results from the superposition of the applied field \vec{H}_a and (the tail of) the aforementioned exchange field in YBCO, \vec{H}_{ex} , in a way similar to the magnetic field induced superconductivity [17,18]. Accordingly, the alignment between \vec{H}_a and \vec{H}_{ex} influences the superconductivity in YBCO, and consequently a modulation in the alignment between \vec{H}_{ex} and \vec{H}_a should accompany a change of resistance in the superconducting transition region.

In this Letter we examine the angular dependence of the magnetization structures in LYL trilayers in experiments where the magnetic field rotates in-plane. We utilize the polarized neutron reflectometry (PNR) technique, which is capable of resolving the depth profile of the magnetization with subnanometer resolution [19–21], to correlate the angular dependent magnetization structure and magnetoresistance (MR). We show unambiguously that, in the superconducting transition region, MR depends on the alignment between \vec{H}_{ex} and \vec{H}_a , rather than the alignment between the two LCMO's magnetizations. This result strongly supports the Salafranca-Okamoto scenario and settles a longstanding debate.

Samples were grown by sputter deposition in pure oxygen atmosphere on (100) SrTiO_3 substrates [22] with a nominal structure of 40 unit-cells (u. c.) LCMO/8 u.c. YBCO/40 u.c. LCMO. The sample size is $5 \times 10 \text{ mm}^2$. X-ray reflectometry (XRR) experiments were conducted at room temperature using $\text{Cu } K\alpha$ radiation. Polarized neutron reflectometry (PNR) experiments were conducted on the ASTERIX reflectometer at the Lujan Neutron Scattering Center. Magnetic hysteresis loops, magnetotransport data and PNR data were taken at 26 K. The sample's resistance is $\sim 10^{-4}$ of the normal state resistance at 26 K so that the

magnetoresistance is overwhelmed by the modulation of the superconductivity in the YBCO layer.

Our LCMO films have an in-plane cubic anisotropy with the easy axes along the $[110]$ and $[1\bar{1}0]$ axes [7]. Figure 1(a) shows the easy-axis magnetization hysteresis loop. Presumably due to different strain states of the bottom LCMO layer (grown on SrTiO_3) and the top LCMO layer (grown on YBCO), the two layers have different magnetic properties. The well-separated two-step switching with different step sizes indicates different saturation magnetizations, and different anisotropies between the top and bottom LCMO layers. Therefore, the relative magnetization orientation in the top and bottom layers is modulated upon rotating in an in-plane magnetic field, with an amplitude between the two coercivities. Figure 1(b) shows the magnetoresistance in a polar plot for a field of fixed magnitude (150 Oe). The further from the radius origin the larger the resistance. The field direction Φ_H is defined with respect to the $[100]$ direction. The MR shows a quasi-four-fold symmetry with four local R_{\min} 's along the LCMO's magnetic easy-axis directions, i.e., 45° , 135° , 225° , and 315° ; it also shows a hysteresis between clockwise (CW) and counterclockwise (CCW) rotations.

We determine the saturated magnetizations of the top and bottom LCMO layers with complementary studies of XRR and PNR. Figure 2(a) shows the PNR data in saturation with a 5 kOe field applied along the $[100]$ direction. The reflectivities are plotted versus the wavevector transfer along the film's normal direction Q_z . $Q_z = 4\pi \sin(\theta_i)/\lambda$, where θ_i is the incident angle and λ is the neutron's wavelength. R^{++} and R^{--} are the two non-spin-flip reflectivities. With a combined refinement of the XRR and PNR data, we find that the saturation magnetizations of the top and bottom LCMO layers are 380 and 540 emu/cm³, respectively. As also reported previously, the fitting indicates a possible suppression of the magnetization at

the LCMO/YBCO interfaces [23]. However, because of the limited Q_z range, this PNR study is not sufficient to resolve the subtleness of the magnetization profile at the interfaces so that the amplitude of YBCO magnetization cannot be determined accurately. (See Supplemental Material [24] for further details.)

Next we study the response of the top and bottom layer magnetization during rotation of the magnetic field. A 150 Oe field was applied along 0° after having saturated the sample along 180° . The experiments were then conducted at the following field directions sequentially: 0° , 45° , 90° , 135° , 180° , 191° , 202° , 225° ; and then 202° , 180° , and 158° . In contrast to the case for saturation, there the spin-flip reflectivities (R^{SF}) are nonzero. R^{SF} is sensitive to the square of the components of the magnetization perpendicular to the field direction [19–21]. Figures 2(b)–2(d) show some representative data. R^{SF} is high at 90° ; it becomes lower at 135° and slightly increases again at 180° . We determine the direction and magnitude of the magnetizations for the top (\vec{M}_t) and bottom (\vec{M}_b) LCMO layers independently at each field direction by fitting R^{++} , R^{--} , and R^{SF} all together. Figures 3(a)–3(d) shows the \vec{M}_t and \vec{M}_b obtained from the best fit as a function of the field direction. The amplitudes are normalized to their respective saturation magnetizations. θ_t and θ_b are the directions of \vec{M}_t and \vec{M}_b , respectively, with respect to the $[100]$ axis. The magnetic field affects the magnitude of the top layer magnetization, but not its direction. This implies the top layer breaks up into domains. On the other hand, the magnetic field affects the orientation of the bottom layer magnetization but not its magnitude. Thus the bottom layer apparently rotates in response to the field.

Because \vec{M}_b keeps the saturation amplitude during the rotation, we use the coherent rotation model to estimate its expected direction to achieve a more detailed picture of

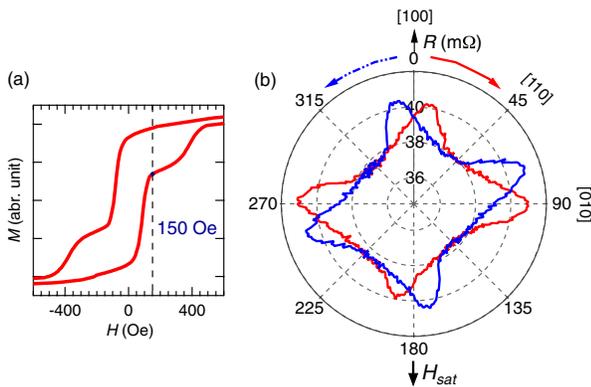


FIG. 1 (color online). (a) Magnetization hysteresis loop along an easy axis ($[110]$). The dashed line shows $H = 150$ Oe. (b) Angular dependence of MR. A 150 Oe in-plane field is applied after having negatively saturated the film along 180° (a hard axis direction). Resistances are recorded when the field rotates either clockwise (CW) or counterclockwise (CCW).

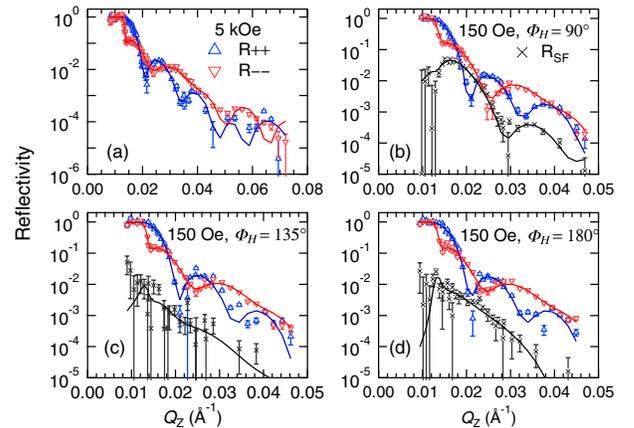


FIG. 2 (color online). (a) PNR data at 5 kOe (the sample is in saturation). (b)–(d) Representative PNR data at 150 Oe with field direction $\Phi_H =$ (b) 90° , (c) 135° and (d) 180° , respectively. Symbols are the experimental data and the lines show the the best fits.

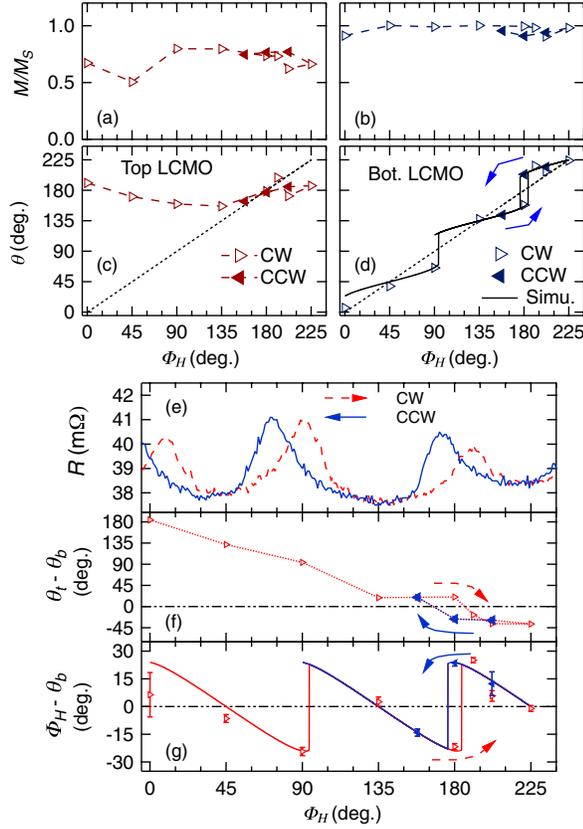


FIG. 3 (color online). Normalized amplitudes (M/M_S) and directions (θ_t and θ_b) for the top [(a), (c)] and bottom [(b), (d)] LCMO magnetizations during rotation, as determined from our PNR experiments. The dotted lines in (c) and (d) show the field direction Φ_H . (e) Angular dependence of MR [same as Fig. 1(b)]. Relative orientations (f) between \vec{M}_t and \vec{M}_b , and (g) between \vec{M}_b and \vec{H}_a , determined from the PNR (triangles), respectively. The solid lines in (d) and (g) show the calculated results based on the energy minimization. Clearly, \vec{M}_b is parallel to \vec{H}_a when the field is along an easy axis.

its magnetization structure during rotation. We consider the Zeeman energy and the anisotropy energy in the free energy, i.e., $F = -\vec{M}_b \cdot \vec{H}_a + K_4 \cos^2(2\theta_b)$, where $M = M_S = 540$ emu/cm³, $H = 150$ Oe, and K_4 is the biaxial magnetocrystalline anisotropy [7]. θ_b is computed via minimizing the free energy. As shown in Fig. 3(d), the calculated values well match the PNR results with $K_4 = 1.6 \times 10^4$ erg/cm³. The only exception is at 0° because of its different magnetic history (field sweeping rather than rotation). Clearly, \vec{M}_b is parallel to \vec{H}_a when \vec{H}_a is along an easy axis. At the same time, the angle between \vec{M}_b and \vec{H}_a reaches a local maximum when the field slightly passes a hard axis, and it shows a hysteresis between clockwise and counterclockwise rotations.

Figure 3(f) shows the relative orientation between \vec{M}_t and \vec{M}_b as the field was rotated. \vec{M}_t and \vec{M}_b are nearly antiparallel when $\Phi_H = 0^\circ$, and parallel when $\Phi_H = 135^\circ$.

More importantly, the difference between the orientations of M_t and M_b decreases monotonically as Φ_H increases from 0° to 135°. Despite a sign change, the amplitude of the relative orientation changes little between $\Phi_H = 135^\circ$ and 225°. If the magnetization alignment governed the MR monotonically, such as for the spin-dependent scattering, then the MR would show no oscillations between 0° and 135° and change little between 135° and 225°. These are obviously in contrast to the MR data shown in Fig. 3(e). Therefore, our results exclude many scenarios that are based on the concept of the mutual magnetization alignment.

On the other hand, the interplay between \vec{H}_{ex} (from the bottom interface) and \vec{H}_a is able to explain the oscillations of the MR with Φ_H . \vec{H}_{ex} is on the order of a few hundred tesla in the first interfacial YBCO unit cell [5]. Salafranca and Okamoto have shown that \vec{H}_{ex} decays exponentially from the interface with an attenuation length less than 1 u.c. and does not quite reach the center of 8 u.c. thick YBCO when $T = T_C$; therefore, \vec{H}_{ex} 's from the top and bottom interfaces influence the superconductivity independently [4]. At the same time, both the coherence length and the mean free path are ≤ 1 u.c. along the c axis in YBCO [25,26]. Therefore, we view the 8 u.c. YBCO layer as a few parallel sublayers for simplicity. This situation is shown in Fig. 4(a). \vec{H}_{ex} in the central YBCO is much weaker than in the interfacial one, so that only the central YBCO becomes superconducting and dominates the resistance of the trilayers when $T = T_C$. The superconductivity in central YBCO is subject to \vec{H}_{tot} that results from the superposition of \vec{H}_a and (the tail of) \vec{H}_{ex} . The relative alignment between \vec{H}_a and \vec{H}_{ex} determines the amplitude of \vec{H}_{tot} during the field rotation [see Fig. 4(b)]. The change of H_{tot} is on the same order of the applied field (150 Oe) during the rotation, slightly shifting the superconducting transition curves and giving rise to a small but observable MR. When \vec{H}_{ex} and \vec{H}_a

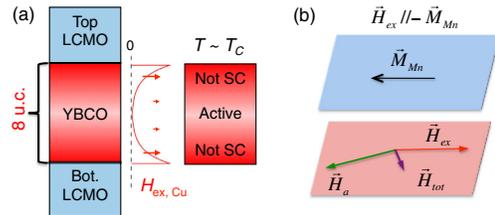


FIG. 4 (color online). (a) A schematic picture of the induced exchange field \vec{H}_{ex} in YBCO. H_{ex} decays exponentially from the interface. When $T \sim T_C$, only the central YBCO undergoes the superconducting transition and therefore dominates the transport properties because the superconductivity in the interfacial YBCO is strongly suppressed. (b) \vec{H}_{ex} is antiparallel to the magnetization of the adjacent LCMO layer (\vec{M}_{Mn}). Meanwhile, the superconductivity in central YBCO is subject to \vec{H}_{tot} that results from the superposition of \vec{H}_a and (the tail of) \vec{H}_{ex} . Therefore, the relative alignment between \vec{H}_a and \vec{M}_{Mn} plays a key role in controlling the superconductivity.

are antiparallel, they effectively cancel each other. Since \vec{H}_{ex} is antiparallel to \vec{M}_{Mn} , H_{tot} is weakest when \vec{H}_a is parallel to \vec{M}_{Mn} , which gives rise to a low resistance state. As shown in Fig. 3(g), \vec{H}_a is parallel to \vec{M}_b when \vec{H}_a is along an easy axis direction with corresponding resistance minima. At intermediate angles, H_{tot} varies and so does MR. This explains the four-fold symmetry of MR. At the same time, the angular hysteresis of \vec{M}_b with respect to the field direction gives rise to the hysteresis in both H_{tot} and MR.

From the Salafranca-Okamoto theory, we also expect a unidirectional offset in MR due to the balance between the external field and the exchange field from the top surface since \vec{M}_t retains the initial saturation direction. The sample used in this PNR study does not show this expected offset and the reason is unclear. One possibility is that as a result of the top LCMO layer breaking down into domains, the effect is compromised. However, such an offset is observed in other samples. Figure 1(b) in Ref. [27] is an example. It clearly shows that, beside the hysteretic four-fold symmetry, there is a unidirectional offset in MR along the initial saturation direction.

A final remark concerns the effect of stray fields created by domain walls of ferromagnetically coupled face-to-face domains in the two FM layers. It has been argued that the magnetic flux closure of the enhanced stray field at domain walls through the SC will cause a large MR [12,28]. This does not occur in our rotation experiment at 150 Oe because the bottom LCMO maintains its saturation magnetization. However, we do observe additional MR features due to the effect of stray fields in other rotation sequences [24].

In summary, we have shown that the interfacial electronic reconstruction controls the inverse spin switch behaviour of half metal-superconductor oxide spin valves. The angular dependence of MR in LYL trilayers along the superconducting transition displays symmetry features that are not correlated with the relative alignment between the two FM's magnetizations, which rules out many MR scenarios proposed so far. Rather the field dependence of the MR is explained by the interplay between the applied field and (the tail of) the induced exchange field on YBCO coming from the electronic reconstruction at the LCMO/YBCO interface. Since the inverse spin switch in LYL is now demonstrated to be governed by interfacial electronic reconstruction and not shape dependent micromagnetic effects, we expect it to survive miniaturization to the nanoscale.

We thank S. Okamoto and J. Salafranca for valuable discussions. Research at Argonne National Laboratory was supported by the U. S. Department of Energy (DOE), Office of Basic Energy Sciences, Division of Materials Sciences and Engineering under Grant No. DE-AC02-06CH11357. Work at UCM was supported by Spanish MICINN Grant No. MAT 2011 27470, Consolider Ingenio CSD2009-00013 (IMAGINE), CAM S2009-MAT 1756 (PHAMA). This work has benefited from the

use of the Lujan Neutron Scattering Center at LANSCE, which is funded by the U. S. Department of Energy's Office of Basic Energy Sciences. Los Alamos National Laboratory is operated by Los Alamos National Security LLC under DOE through Contract No. DE-AC52-06NA25396.

*yhliu@anl.gov

[†]Present address: Unité Mixte de Physique CNRS Thales, Palaiseau, France.

[‡]Present address: Argonne National Laboratory, Argonne, Illinois 60439, USA.

[§]tevelthuis@anl.gov

- [1] A. Ohtomo and H. Hwang, *Nature (London)* **427**, 423 (2004).
- [2] N. Reyren, S. Thiel, A. D. Caviglia, L. F. Kourkoutis, G. Hammerl, C. Richter, C. W. Schneider, T. Kopp, A.-S. Ruetschi, D. Jaccard, M. Gabay, D. A. Müller, J.-M. Triscone, and J. Mannhart, *Science* **317**, 1196 (2007).
- [3] A. Brinkman, M. Huijben, M. Van Zalk, J. Huijben, U. Zeitler, J. C. Maan, W. G. Van der Wiel, G. Rijnders, D. H. A. Blank, and H. Hilgenkamp, *Nature Mater.* **6**, 493 (2007).
- [4] J. Salafranca and S. Okamoto, *Phys. Rev. Lett.* **105**, 256804 (2010).
- [5] J. Chakhalian *et al.*, *Nature Phys.* **2**, 244 (2006).
- [6] R. Werner, C. Raisch, A. Ruosi, B. A. Davidson, P. Nagel, M. Merz, S. Schuppler, M. Glaser, J. Fujii, T. Chassé, R. Kleiner, and D. Koelle, *Phys. Rev. B* **82**, 224509 (2010).
- [7] C. Visani, J. Tornos, N. M. Nemes, M. Rocci, C. Leon, J. Santamaria, S. G. E. te Velthuis, Y. Liu, A. Hoffmann, J. W. Freeland, M. Garcia-Hernandez, M. R. Fitzsimmons, B. J. Kirby, M. Varela, and S. J. Pennycook, *Phys. Rev. B* **84**, 060405 (2011).
- [8] V. Peña, Z. Sefrioui, D. Arias, C. Leon, J. Santamaria, J. L. Martinez, S. G. E. te Velthuis, and A. Hoffmann, *Phys. Rev. Lett.* **94**, 057002 (2005).
- [9] L. R. Tagirov, *Phys. Rev. Lett.* **83**, 2058 (1999).
- [10] A. Buzdin, A. Vedyayev, and N. Ryzhanova, *Europhys. Lett.* **48**, 686 (1999).
- [11] M. van Zalk, M. Veldhorst, A. Brinkman, J. Aarts, and H. Hilgenkamp, *Phys. Rev. B* **79**, 134509 (2009).
- [12] D. Stamopoulos, E. Manios, and M. Pissas, *Supercond. Sci. Technol.* **20**, 1205 (2007).
- [13] S. Takahashi, H. Imamura, and S. Maekawa, *Phys. Rev. Lett.* **82**, 3911 (1999).
- [14] N. M. Nemes, M. García-Hernández, S. G. E. te Velthuis, A. Hoffmann, C. Visani, J. Garcia-Barriocanal, V. Peña, D. Arias, Z. Sefrioui, C. Leon, and J. Santamaría, *Phys. Rev. B* **78**, 094515 (2008).
- [15] K. Dybko, K. Werner-Malento, P. Aleshkevych, M. Wojcik, M. Sawicki, and P. Przyszlupski, *Phys. Rev. B* **80**, 144504 (2009).
- [16] J. Zhu, I. N. Krivorotov, K. Halterman, and O. T. Valls, *Phys. Rev. Lett.* **105**, 207002 (2010). The calculations are based on CuNi/Nb/CuNi trilayers.
- [17] V. Jaccarino and M. Peter, *Phys. Rev. Lett.* **9**, 290 (1962).

- [18] H.W. Meul, C. Rossel, M. Decroux, O. Fischer, G. Remenyi, and A. Briggs, *Phys. Rev. Lett.* **53**, 497 (1984).
- [19] G.P. Felcher, R.O. Hilleke, R.K. Crawford, J. Haumann, R. Kleb, and G. Ostrowski, *Rev. Sci. Instrum.* **58**, 609 (1987).
- [20] C. Majkrzak, *Physica (Amsterdam)* **221B**, 342 (1996).
- [21] M.R. Fitzsimmons and C. Majkrzak, in *Modern Techniques for Characterizing Magnetic Materials*, edited by Y. Zhu (Springer, New York, 2005), pp. 107–155.
- [22] Z. Sefrioui, D. Arias, V. Peña, J.E. Villegas, M. Varela, P. Prieto, C. León, J.L. Martinez, and J. Santamaria, *Phys. Rev. B* **67**, 214511 (2003).
- [23] A. Hoffmann, S.G.E. te Velthuis, Z. Sefrioui, J. Santamaría, M.R. Fitzsimmons, S. Park, and M. Varela, *Phys. Rev. B* **72**, 140407 (2005).
- [24] See Supplemental Material at <http://link.aps.org/supplemental/10.1103/PhysRevLett.108.207205> for additional experimental details for x-ray and polarized neutron reflectometry, and magnetotransport studies.
- [25] T.K. Worthington, W.J. Gallagher, and T.R. Dinger, *Phys. Rev. Lett.* **59**, 1160 (1987).
- [26] Q. Li, X.X. Xi, X.D. Wu, A. Inam, S. Vadlamannati, W.L. McLean, T. Venkatesan, R. Ramesh, D.M. Hwang, J.A. Martinez, and L. Nazar, *Phys. Rev. Lett.* **64**, 3086 (1990).
- [27] N.M. Nemes, C. Visani, C. Leon, M. Garcia-Hernandez, F. Simon, T. Feher, S.G.E. Velthuis, A. Hoffmann, and J. Santamaria, *Appl. Phys. Lett.* **97**, 032501 (2010).
- [28] J. Zhu, X. Cheng, C. Boone, and I.N. Krivorotov, *Phys. Rev. Lett.* **103**, 027004 (2009).