Magnetoresistance studies of La₂^{3Sr}₁³MnO₃-YBa₂Cu₃O₇-La₂^{3Sr}₁³MnO₃ trilayers with ferromagnetic coupling along the nodal direction of $YBa₂Cu₃O₇$

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I have successfully prepared (110) trilayers of $La_{2/3}Sr_{1/3}MnO_3$ -YBa₂Cu₃O₇-La_{2/3}Sr_{1/3}MnO₃. Magnetization measurements on these samples reveal a stronger coupling between the ferromagnetic layers. The coupling is an order of magnitude higher than that seen in the case of (001) trilayers. Magnetoresistance measurements show a first-order transition in the data coinciding with the antiferromagnetic regime deduced from the magnetization measurements. I have also measured the anisotropic magnetoresistance (AMR) of these samples revealing an unusually high AMR (\sim 72 000%). I attribute such a high AMR to the pair-breaking effects in these films.

DOI: [10.1103/PhysRevB.81.014515](http://dx.doi.org/10.1103/PhysRevB.81.014515)

PACS number(s): 74.78.Fk, 75.60. - d, 75.70.Cn

I. INTRODUCTION

The study of hybrid structures composed of oxide superconductor (SC) and oxide ferromagnet (FM) systems have revealed a variety of exotic phenomena in these systems.¹⁻¹⁰ The interest in these systems arises from the fact that both the states, i.e., SC and FM, are mutually exclusive. Proximity of such exclusive states can give rise to phenomena such as Larkin-Ovchinnikov-Fulde-Ferrel state, exchange coupling, Andreev reflection to name a few. Due to recent advances in thin-film fabrication technique it is possible to achieve good quality interface between SC and FM layers and grow these films in any desired orientation by use of suitable substrate. It is to be noted that the same structure in its normal state can act as a well-known spin-valve system where the two mag-netic layers are separated by a nonmagnetic (NM) layer.^{11[,12](#page-6-4)} When the spacer in such structures is a metallic ferromagnet of 3*d* and 4*f* elements the exchange coupling is chiefly driven by the Rudderman-Kittel-Kasuya-Yoshida interaction.¹³

In this paper I am chiefly interested in (110)-oriented trilayers of double-exchange ferromagnet $La_{2/3}Sr_{1/3}MnO_3$ $(LSMO)$ and a non-Fermi-liquid metal $YBa₂Cu₃O₇$ (YBCO). The giant magnetoresistance seen in FM-NM-FM trilayers and multilayers is related to asymmetric scattering of spin-up and spin-down electrons as they crisscross the spacer while diffusing along the plane of the heterostructure.^{14–[16](#page-6-7)} In case the intermediate NM layer becomes superconducting one would expect to see a profound change in the flow of spinpolarized carriers. It is to be noted that most of the earlier studies involving superconducting spacer layers in a spinvalve-type configuration, where the SC is YBCO, the spin injection in SC layer is along the insulating c axis.^{7[,17](#page-6-9)[–19](#page-6-10)} Such structures do not allow the injection of spin-polarized carriers along the fully gapped nodal planes of YBCO. To overcome this problem it is necessary to grow the films in such a way that the $CuO₂$ planes are perpendicular to the plane of the substrate and in direct contact with the ferromagnetic layers. This is possible if the YBCO layer is grown with crystallographic direction $(100)/(010)$ or (110) perpendicular to the plane of the substrate. In an earlier work we have already demonstrated the growth of (110) hybrids.²⁰

The reason for choosing (110) orientation over $(100)/(010)$ is explained as follows. The growth of YBCO films where the *c* axis of the film is in the plane of the substrate involves the use of heterotemplate technique. Now if the template $(PrBa₂Cu₃O₇$ in this case) is grown $(100)/(010)$ and then the first LSMO layer is deposited, then the YBCO layer becomes (001) oriented where the *c* axis is perpendicular to the substrate plane. This is because of the crystallographic symmetry of the LSMO molecule along (001) , (010) , and (100) directions. But in the case of (110) growth the LSMO layer is (110) oriented so the only possible growth directions for YBCO are (110) and (103). The template in this case helps in increasing the (110) volume fraction. In this paper I describe experimental studies of transport and magnetic properties of nodally coupled hybrids. I have also carried out some control experiments to demonstrate the role of ferromagnetic layers on both sides of SC layer.

II. EXPERIMENTS

Thin films of (110) trilayer of LSMO-YBCO-LSMO, bilayer of YBCO-LSMO, and (110) YBCO were deposited on (110) SrTiO₃ substrates. A multitarget pulsed laser deposition technique based on KrF excimer laser $(\lambda = 248 \text{ nm})$ was used to deposit the thin films. The (110) trilayers had 200 and 500 Å YBCO layer sandwiched between 1000 Å of LSMO. The heterostructure was grown using a heterotemplate technique with the template being $PrBa₂Cu₃O₇$. Further details of film deposition giving information about growth rate, deposition temperature, and pressure are given elsewhere. 20 The epitaxial growth in (110) films were established by x-ray diffraction measurements performed in θ -2 θ geometry. The volume fraction of (110) grains in (110) trilayer was determined by the recipe of Westerheim *et al.*[21](#page-6-12) which comes out to be $\geq 65\%$ with the remaining volume of (103) grains. It is true that the trilayers do not have 100% (110) -oriented grains but the presence of (103) grains still allow direct injection of spin-polarized carriers in the $CuO₂$ planes (Fig. 4.4 from Ref. [22](#page-6-13)). For magnetization measurement a commercial magnetometer Quantum Design MPMS XL5 SQUID) was used. For transport measurements, films were patterned in the form of $1000 \times 100 \mu m^2$ bridge with

FIG. 1. Resistivity data for (110) LSMO-YBCO-LSMO trilayer with $d_{YBCO} = 500$ Å (top panel) and 200 Å (bottom panel). Here the films with 200-Å-thick YBCO layer are not superconducting while (001) trilayer films are superconducting even for a YBCO layer thickness of 50 Å.

photolithography and wet etching such that the long axis of the bridge was parallel to $(1\overline{1}0)$ direction for the (110) oriented films. The measurements of resistivity as a function of temperature, magnetic field strength, and the angle (θ) between field and current were performed using a 4.2 K close cycle He refrigerator with a fully automated homemade setup for applying the field at varying angles between 0 and 2π with respect to the direction of current.²³ The sample was mounted in a way to keep the field in the plane of the sample for all values of the angle between \vec{l} and \vec{H} except for the measurements where out of plane contributions were also recorded.

III. RESULTS AND DISCUSSIONS

A. (110) LSMO-YBCO-LSMO

With the optimized growth conditions, trilayers of (110) $La_{2/3}Sr_{1/3}MnO_3-YBa_2Cu_3O_7-La_{2/3}Sr_{1/3}MnO_3$ were synthesized and their various magnetic and electronic properties were measured. Figure [1](#page-1-0) shows the resistivity curves, $\rho(T)$, for two trilayers. The upper panel is the result for a trilayer with a 500 Å YBCO spacer. The $\rho(T)$ curve is characterized by transition to a superconducting state which starts at \sim 80 K and completes when the temperature reaches \sim 60 K. The bottom panel shows the resistivity for a similar structure with a 200 Å YBCO spacer. In this case the trilayer does not go into the superconducting state though it has a metallic behavior. It is interesting to note that while the YBCO of thickness 200 \AA in the (110) trilayer shows no T_c , a superconducting transition can be seen for YBCO thickness of even 50 \AA for the (001) trilayer.¹⁹ This is presumably due to greater T_c suppression in the case of (110) films because

FIG. 2. Panels A and B show the *M*-*H* loop of the trilayer at 40 and 60 K, respectively. Panels C–G show the field dependence of resistivity for the superconducting trilayer at a few representative temperatures across the transition temperature. The MR of the film in the superconducting state is higher than that in the normal state. Panel H shows the comparison of MR% with temperature and sample resistivity. A clear peak seen near the transition temperature can be attributed to the unusual rise in the normal-state properties of the superconductor near the transition temperature.

of direct injection of spin-polarized carriers in the superconducting $CuO₂$ $CuO₂$ $CuO₂$ planes of YBCO. In Fig. 2 (panels A and B), I have shown the *M*-*H* loops of the trilayer with d_{YBCO} = 500 Å, at 40 and 60 K, respectively. A plateau in the *M*-*H* loop near zero magnetization confirms the presence of an antiferromagnetic state. This antiferromagnetic state is present in the normal state of the superconductor as well. Panels C through G in the same figure show the MR of the superconducting trilayer at a few temperatures across the transition. The MR in this case is defined as $R(H)/R(0)$, where $R(H)$ is the resistance of the sample at applied field *H*. The field and current (I) in this case are coplanar but orthogonal to each other. I first discuss panels C and D which present the data for the trilayer in the superconducting state at 40 and 60 K, respectively. Starting from a fully magnetized state of LSMO layers at 800 Oe the MR first increases slowly as the field is decreased. At \sim 400 Oe the rate of increase becomes faster but remains continuous till the zero field. On reversing the field, a small steplike jump is seen around \sim -50 Oe and then the MR keeps rising to a peak value, after which, a local minimum is attained followed by a sudden jump in the MR at \sim 370 Oe to a much lower value. Further increment of the field results in a gradual decrease in MR till a reversed field of 800 Oe is reached. This cycle repeats itself once the field is decreased from −800 Oe and increased to 800 Oe. I have measured MR for the sample at a few more temperatures below T_c . In all those measurements I found that the resistance ratio $(R_{\uparrow\downarrow})_{max}/(R_{\uparrow\uparrow})_{min}$ over the whole range of measurement is \sim 2, where $(R_{\uparrow\downarrow})_{max}$ is the resistance at the peak position in the MR-*H* curve and $(R_{\uparrow\uparrow})_{min}$ is the minimum resistance of the segment of MR-*H* curve where the magnetizations of both the FM layers are parallel to each other. The current flowing through the sample in these measurements is zero field $\sim I_c$ of the sample at that temperature. Panels E and F show the MR vs *H* data for 70 and 80 K, respectively, where the YBCO layer in the trilayer is in the superconducting transition region (top panel of Fig. [1](#page-1-0)). Here one can see that the high-field negative magnetoresistance region, as seen in panels C and D in the field regime \sim 400–800 Oe, is replaced by a positive magnetoresistance which is completely opposite to the negative MR seen on LSMO films.²⁴ The resistance ratio in panel E is \sim 3 which is the highest over the whole range of measurement. This resistance ratio sharply drops once the film starts entering the normal state as is evident from the panels showing the MR at 80 K (panel F) and 100 K (panel G). Panel G shows MR data for 100 K, where negative magnetoresistance is seen in a high field, which is a characteristic feature of LSMO. 24 Even though in panel G the resistance ratio is reduced due to the superconducting spacer entering into the normal state, however the first-order jump in resistance near $H \approx 350$ Oe is still present clearly proving the fact that the resistance ratio is dependent on the spacer-layer properties while the first-order transition is dependent on the FM layer properties. In panel H the MR% plotted is defined as $\Delta R/R(0)$, where $\Delta R = (R_{\uparrow\downarrow})_{max} - (R_{\uparrow\uparrow})_{min}$ and $R(0)$ is the resistance at zero field at that temperature. A distinct behavior of MR% can be seen when the sample becomes superconducting. The sample in the normal state has a very low MR but once the sample starts moving into the SC regime, the MR shoots up rapidly and then comes down to saturate at a constant value at low temperatures. The increase in MR in the vicinity of T_c can be attributed to the abnormal increase in the normal-state properties of the superconductor.²⁵

Another important feature that is quite prominent in Fig. [2](#page-1-1) is the presence of peaks in the MR data. A comparison of MR-*H* and *M*-*H* plots (panels A and C, and panels B and D) shows that the peaks coincide with the region where the *M*-*H* curve has a plateau. The near zero magnetization in the plateau suggests antiferromagnetic coupling between the magnetization vectors of the top and bottom LSMO layers. One can estimate the exchange energy associated with the AF coupling in the following way. The free-energy expression for two magnetic layers of the same thickness coupled via the spacer can be written as^{26}

$$
F = F_c + F_a - \vec{H} \cdot (\vec{M}_1 + \vec{M}_2)t,
$$
 (1)

where M_1 and M_2 are the magnetizations of the top and bottom LSMO layers, F_c is the coupling energy per unit area, and *t* is the thickness of a single LSMO layer. The anisotropy part of the energy (F_a) is primarily dependent on contributions from magnetocrystalline anisotropy as well as the inplane uniaxial anisotropy of the film. Assuming a bilinear coupling, F_c can be written as

$$
F_c = -J_1(\vec{M}_1 \cdot \vec{M}_2),\tag{2}
$$

where \vec{M}_1 and \vec{M}_2 are the unit magnetization vectors, and J_1 < 0 corresponds to antiferromagnetic coupling between FM layers. For a given external field, the minima of Eq. ([1](#page-2-0)) will yield the relative orientation of \vec{M}_1 and \vec{M}_2 . If J_1 is positive, even in zero field $\tilde{M}_1 || \tilde{M}_2$ so increasing the field does not change anything. If J_1 is negative then the minima is achieved when $H=0$ and \vec{M}_1 and \vec{M}_2 are antiparallel or antiferromagnetic in alignment. The anisotropic term in Eq. (1) (1) (1) , F_a can be written as

$$
F_a = KtM_{1,2},\tag{3}
$$

where $M_{1,2}$ is a function of \vec{M}_1 and \vec{M}_2 and K is the anisotropy constant. If $|J_1| \ge K$, then a second-order reorientation transition and a smooth linear *M*-*H* dependence followed by saturation is predicted by the theory. On the other hand if $|J_1| \ll K$ then the magnetization slowly increases in the low field, and then abruptly at some critical field H_s , the system undergoes a first-order transition with an abrupt jump to saturation magnetization. The critical field H_s which is also known as saturation field or switching field can be written in terms of the magnetization density M_s , thickness t of one ferromagnetic layer and coupling energy J_1 as²⁶

$$
H_s = -\left(\frac{J_1}{M_s t}\right). \tag{4}
$$

It is to be noted that this equation is for the case where the two FM layers are of equal thickness *t*. The behavior of the magnetization seen in Fig. [2](#page-1-1) corresponds to this situation. In Fig. [3](#page-3-0)(a), I have shown the variation in J_1 as a function of the temperature calculated using Eq. ([4](#page-2-1)) for $d_{YBCO} = 500$ Å. Panel B of Fig. [3](#page-3-0) shows a comparison between the peak position in the MR-*H* data and the start and end points of the antiferromagnetic phase in *MH* data. Panels C and D show a typical *MH* and MR-*H* data for the sample. The arrows point to positions of the points on the MR-*H* and *MH* data which have been plotted in panel B.

I now discuss the behavior of J_1 as seen in Fig. [3](#page-3-0) (panel A). The temperature dependence of the interlayer exchange coupling in metallic multilayers has been worked out theoretically[.27,](#page-6-18)[28](#page-6-19) The starting point for calculating the coupling is to calculate the energy per unit area in the ferromagnetic and antiferromagnetic configurations. The difference of the two will give the exchange coupling of the system. The energy terms are functions of the reflection coefficients of the electrons in the spacer hitting the spacer-ferromagnet in-

FIG. 3. Panel A shows the variation in coupling energy with temperature between two ferromagnetic layers. The coupling in this case is higher than that seen for (001) layers (details in text). Panel B shows the comparison between the peak position in MR data and the start and end points of the antiferromagnetic phase in the *MH* data. Panels C and D show the position of the points on the *MH* and MR data plotted in panel B. The agreement in the data points clearly shows the dependence of discontinuities in the MR data on the antiferromagnetic phase of the sample.

terface calculated in the light of the free-electron model. Using the above method the dependence of linear exchange coupling J_1 with temperature is given by²⁸

$$
J_1(T) = J_1(0) [(T/T_0)/\sinh(T/T_0)], \tag{5}
$$

where the characteristic temperature T_0 depends on the Fermi wave vector k_F and spacer thickness d_n through the relation $T_0 = \hbar^2 k_F / 2 \pi k_B d_n m$, where *m* is the free-electron mass and \hbar and k_B are the Planck and Boltzmann constants, respectively. In this case, since the transport is along the (110) axis, the relevant wave vector will be $k_{F(110)}$. In Fig. [3](#page-3-0) it is seen that J_1 increases linearly as the temperature decreases. This is dif-ferent from the behavior expected from Eq. ([5](#page-3-1)). In general, functions of the type $x/\sinh(x)$ saturate in the limit $x \to 0$ but in this case I do not see any saturation of J_1 even at a temperature of 2 K. The magnitude of J_1 is almost an order higher than what is seen for (001)-oriented heterostructures.¹⁹ These high values are in line with the predictions of de $Melo²⁹$ where he had pointed out that the coupling along the (110) direction will be higher than that along the (100) or (001) direction. In Fig. [4,](#page-3-2) I have shown the schematic of a (110) trilayer where the spacer is a *d*-wave superconductor. It is clear from the figure that in this case the coupling is mediated by the nodal quasiparticles whose number density remains high even at $T \approx 0$. This explains the large J_1 and the absence of any anomaly in J_1 near T_c . The middle panel of Fig. [3](#page-3-0) shows the comparison between various critical points on the *MH* loop and MR data. Here I have plotted the starting and end points of the plateau in the *MH*

FIG. 4. Schematic showing nodal direction in *dx*2−*y*² orbital. The LSMO layers shown in the schematic signify the position of LSMO layers in a (110) trilayer. This schematic is for a trilayer where the spacer is a *d*-wave superconductor.

loop against temperature. From the MR(*H*) loop, I have taken the points of discontinuity which are indicated in panel D. Panel C shows the position of points H1 and H2 on the *MH* loop. One can clearly see that both the representative points agree well within experimental error with each other clearly demonstrating the fact that the discontinuities in MR data come from the antiferromagnetic regime of the sample.

Figure [5](#page-3-3) presents the AMR data on (110) trilayers. The *x* axis defines the angle θ with respect to the current direction. The left-hand axis shows $R(\theta)$ and the right-hand *y* axis shows MR% defined as $\frac{R(\theta) - R_{min}}{R_{min}}$, where R_{min} is the minimum resistance of the sample over the whole range of measurement. The reason for using this definition will be explained when we discuss the data presented in Fig. [6.](#page-4-0) Before I discuss the data in detail, let me explain the measurement geometry which has been schematically shown in panel D of the diagram. The patterned sample is placed over a solid block of copper as shown in the figure. The current in the

FIG. 5. Angular dependence of MR in the FM-SC-FM trilayer is plotted for $T < T_c$ (panel A), $T > T_c$ (panel B), and a FM-NM-FM trilayer (panel C). The data clearly shows that a superconducting spacer in the superconducting state enhances and modifies the AMR considerably. The AMR of the FM-NM-FM trilayer is mostly dependent on the AMR of the FM layer. Panel D shows the schematic of the measurement geometry.

FIG. 6. Current dependence of AMR measured at three different configurations. The top panel shows the AMR for the configuration when the applied magnetic field stays in the plane of the sample. The middle panel shows the AMR when the current through the sample is always perpendicular to the field. The field in this case moves from a position where it is parallel to $CuO₂$ planes to a position where it is perpendicular to $CuO₂$ planes. The bottom panel shows the AMR when the applied field moves in the plane of the $CuO₂$ planes. The AMR of the sample increases as the current through the sample is decreased, clearly proving the fact that a low resistance spacer enhances MR in these trilayers.

sample is along \hat{y} . The applied field rotates in the *xy* plane. The angle θ is measured with respect to \hat{v} . The sample is patterned along \hat{y} in such a way that the CuO₂ planes of YBCO are in the *yz* plane. In short, \hat{x} , \hat{y} , and \hat{z} are parallel to (001) , $(1\bar{1}0)$, and (110) directions of the sample. Panels A and B show the AMR of the trilayer at $T=40$ K ($) and$ $T = 100 \text{ K}$ $(>T_c)$, , respectively, on a film with d_{YBCO} = 500 Å. Panel C shows the AMR measurement at 20 K on a film with $d_{YBCO} = 200$ Å. From panels A and B, one can see that the trilayer shows a huge MR when the SC layer is in the superconducting state. The same film hardly shows any AMR once the film moves into the normal state. This is also evident from the AMR data in panel C. The trilayer in this case is nonsuperconducting for all temperatures (bottom panel of Fig. [1](#page-1-0)). The angular dependence in panel C is similar to the one seen for plain LSMO films.³⁰ The dependence of AMR in the superconducting state is markedly different from the one in the normal state. For my sample geometry, when the field is perpendicular to the current it is also perpendicular to the $CuO₂$ planes resulting in maximum dissipation in the YBCO layer. So, the logical thing would be that the AMR is higher when the field is perpendicular to the current but what I see here is completely opposite. This can be explained as follows. We know that the dissipation in YBCO when the applied field is perpendicular to the copperoxide planes is due to the formation of vortices and when the field is parallel, the dissipation is mostly due to pair-breaking effects. In my geometry, the effective area of the sample exposed perpendicular and parallel to field is equal to the thickness of the film times the length and breadth, respectively. It is quite possible that the effect of vortex formation in such a small area has lower dissipation than pair-breaking effects. Hence, if I assume that pair breaking is causing larger dissipation in the YBCO layer in these trilayers, I can safely conclude that for fields parallel to the current (or copper oxide planes) the AMR will be higher.

In Fig. [6,](#page-4-0) I have plotted the current dependence of the AMR for three different orientations of the sample as shown in the figure. Before I explain the results in this figure it is important to discuss the reasons behind using the particular definition for AMR. If we look closely $H(0)$ in the figure corresponds to the situation when the angle of the field with respect to the current or sample plane is zero. In the top panel, the measurement geometry is the same as is shown in Fig. $5(d)$ $5(d)$. In the middle panel, the CuO₂ planes and current are perpendicular to field when $\phi = 0$ and in the bottom panel the field is parallel to the $CuO₂$ planes and current for $H(0)$. So, to compare the data in these three panels it is important to find a equivalence point for the calculation of AMR which one cannot find if one considers the field orientation or $CuO₂$ -plane orientation or current as reference point. So the best option would be to consider the point where the resistance of the sample is minimum. Coming back to the top panel, which shows the dependence in the coplanar configuration, one can see that as current through the sample is decreased AMR increases. For the same film, the middle and bottom panels show unusually large AMR. It is to be noted that the AMR in the middle and bottom panels come due to contributions from two different effects. First will be the AMR due to field being parallel or perpendicular to the $CuO₂$ layer of the SC spacer and the second will be due to the out-of-plane field which gives rise to a high resistance state. In the bottom panel, one can see that the applied field always stays parallel to the $CuO₂$ planes. The AMR seen here essentially arises due to the fact that the applied field becomes perpendicular to SC layer. In the middle panel, one can see that the AMR is much higher than that of any other configuration. In case I assume that the contribution is only from the first case, as pointed out earlier, then there should be no angular dependence in the bottom panel and if I assume that all the contribution is coming from the second case, then the middle and bottom panels should show comparable AMR. The fact that the middle panel shows an AMR almost six times that of the bottom panel points to the fact that the AMR is coming from both the contributions which in this case, have same behavior in the configuration shown in middle panel and hence the effect is additive. One can also see that in all the cases, reduction in the current results in an increase in the AMR vindicating the hypothesis that low resistance in the spacer contributes to a higher magnetoresistance. Apart from this, if we try to talk only about the magnitude of change in resistance and not the percentage then we will see that the maximum resistance over the whole range of 2π can increase by as much as \sim 700 times the minimum resistance. The discussion on the trilayers will not be complete without a small discussion on the crystallographic dif-

FIG. 7. Panel (a) shows the resistivity data for a (110) LSMO-YBCO bilayer. Panel (b) shows the current-voltage characteristic for the bilayer film at 76 K. Panels (c) – (e) show the AMR for a LSMO-YBCO bilayer in three different configurations. One can see that the AMR in this case is almost two orders of magnitude smaller than that seen for the trilayer. Panel (f) shows the resistivity data for a (110)-oriented YBCO layer. Panel (g) shows the current-voltage characteristic for the YBCO film at 6 K. Panels (h) - (j) show the AMR for (110) YBCO film in three different configurations. The AMR primarily arises from the orientation of magnetic field with respect to $CuO₂$ planes.

ferences along the two directions in the plane of the film. The directions in the plane of the film are $(1\bar{1}0)$ and (001) with (001) being the easy axis. The easy axis is predominantly determined by the easy axis of the LSMO layers which in this case is along the (001) direction.³⁰ When the field is along the (001) direction it is the easy axis of the LSMO layers but the dissipative state of the YBCO layer and when the field is along the $(1\bar{1}0)$ direction it is parallel to $CuO₂$ layers which is less dissipative but the presence of LSMO layer introduces a strong dissipation resulting in a high resistance state as seen in Figs. [5](#page-3-3) and [6.](#page-4-0)

B. Anisotropic magnetoresistance of (110) LSMO-YBCO and (110) YBCO thin films

To verify my results of unusually high AMR in the trilayers I have done some control experiments involving an LSMO-YBCO bilayer and a YBCO single-layer film. The growth conditions for these films are exactly the same as the trilayer except for the fact that these films were made with $d_{YBCO} = 1000$ Å. Figure $7(a)$ $7(a)$ shows the resistivity data for the bilayer. Panel (b) in the same figure shows a currentvoltage characteristic for the bilayer at 76 K. In Figs. $7(c) - 7(e)$ $7(c) - 7(e)$, I have shown the AMR in this film at 3 kOe for three different configurations. One can see that AMR in this film is much smaller as compared to that of the superconducting trilayer. Most of the contribution comes from the fact that the field moves from parallel to perpendicular to the $CuO₂$ planes. Panel (d) shows the AMR when the field stays in the $CuO₂$ plane. The dependence seen here mostly comes from the fact that the field moves in and out of plane of the sample. In Fig. $7(f)$ $7(f)$, I have plotted the resistivity data for a single layer of (110) YBCO. Panel (g) shows the currentvoltage characteristic for the film at 6 K. Figures $7(h) - 7(j)$ $7(h) - 7(j)$ show the AMR for this film at different configuration. Here one can clearly see that angular dependence comes primarily from the positioning of the field perpendicular or parallel to the $CuO₂$ planes. But a look at panels (e) and (i) of the figure tells us that the AMR for the bilayer is higher than that for the single layer. This is explained by the presence of an FM layer near a superconducting layer. Earlier, people had seen a higher change in the resistivity in a bilayer than on a single layer when the film was pushed into the superconducting state.³¹

IV. CONCLUSION

In conclusion, manganite-cuprate bilayers and trilayers where the $CuO₂$ planes are normal to the plane of the templated (110) SrTiO₃ have been synthesized and their various transport and magnetic properties have been studied. I find that the coupling between the two FM layers is higher in this case than on that of the (001) bilayer as predicted by de Melo.²⁹ I have also observed unusually high $(\sim 72\,000\%)$ angular magnetoresistance in these trilayers. Some control experiments have been done to point out the fact that the unusually high AMR comes from the coupling between the two ferromagnetic layers. The MR% calculated from the MAGNETORESISTANCE STUDIES OF La_{2/3}Sr... PHYSICAL REVIEW B **81**, 014515 (2010)

magnetoresistance measurements shows a peak near the superconducting transition temperature which has been attributed to the unusual increase in the normal-state properties of the superconductor near its transition temperature.

ACKNOWLEDGMENT

The author would like to acknowledge financial support from Indian Institute of Technology Kanpur, Kanpur, India.

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