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PAPER

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Odd-frequency triplet Cooper pairs are believed to be the carriers of long-range superconducting correlations in ferromagnets. Such triplet pairs are generated by an inhomogeneous magnetism at the interface between a superconductor (S) and a ferromagnet (F). So far, reproducible long-range effects were reported only in complex layered structures designed to provide the magnetic inhomogeneity. Here we show that spin triplet pair formation can be found in simple unstructured Nb/permalloy (Py = Ni_{0.8}Fe_{0.2})/Nb trilayers and Nb/Py bilayers, but only when the thickness of the ferromagnetic layer ranges between 140 and 250 nm. The effect is related to the emergence of an intrinsically inhomogeneous magnetic state, which is a precursor of the well-known stripe regime in Py that in our samples sets in at thickness larger than 300 nm.

1. Introduction

Superconductivity and ferromagnetism are competing phases whose coexistence is unlikely to occur. Notable exceptions take place when the electrons responsible for the magnetism are only weakly coupled to those inducing superconductivity, as in some ternary rare-earth compounds [1]. Differently from the case of bulk systems, the coexistence between superconductivity and ferromagnetism may be easily achieved in artificial superconductor–ferromagnet (S/F) hybrids. In these systems the two antagonistic orderings are confined in spatially separated layers interacting via the proximity effect, which arises when a superconductor comes in metallic contact with a ferromagnet [2]. In this case, the spin-singlet Cooper pairs enter the F-layer and magnetic excitations leak into the S-region across the S/F interface. As confirmed by many experiments [3, 4], the penetration depth, ξ_F , of singlets in the F-layer is, in the diffusive limit, basically given by $\xi_F = \sqrt{\hbar D_F / E_{ex}}$ (D_F is the diffusion coefficient), while superconductivity is suppressed in S within a distance ξ_S from the interface (ξ_S is the superconducting coherence length). In addition, the presence of the exchange field, E_{ex} , in F causes an energy shift between the electrons of the pairs entering the F-layer and the Cooper pairs acquire a non-zero center-of-mass momentum. As a consequence, the superconducting order parameter does not decay monotonically in F, as it would happen in the case of a normal metal, but it shows oscillatory decay in the direction perpendicular to the interface over a length scale given (again) by ξ_F [5, 6]. In strong ferromagnets, since $E_{ex} \sim 1$ eV, ξ_F is only few nanometers.

However, at the interface between a superconductor and a ferromagnet, conventional singlet Cooper pairs can be converted into equal-spin triplet ones. Since the triplets have their spins equally aligned, they are much less affected by the pair breaking caused by E_{ex} in F. Thus, once injected in the F-layer, at low T they can survive over distances of the order of hundreds of nanometers [7], contrary to what happens for the singlets. Such spin-triplet correlations are predicted to have even symmetry in space (s-wave), which makes them robust against

scattering, but have odd symmetry with respect to time (hence named odd-frequency). The key factor to achieve singlet-to-triplet conversion is the presence of a certain degree of magnetic inhomogeneity at the S/F interface [7]. There are different ways of providing such inhomogeneity. In the original theoretical proposal, the magnetic inhomogeneity was described in terms of a rotating vector with the angle of the magnetization direction rotating in the plane of the S/F interface when moving away from it. This scenario could be realized, for example, in a domain wall within which the magnetization gradually rotates. So far, however, almost all the experiments which gave evidence of a long-range proximity effect relied on different ways to provide the required magnetic inhomogeneity [8–11]. In most of the experimental works, an extra ferromagnetic layer F_1 is inserted in between S/F and the inhomogeneity is controlled by varying the collinearity between the magnetization of F and F_1 [12–17]. By using holmium (Ho) as F_1 -layer and Co as F-layer, Robinson *et al* [12] more closely reproduced the original theoretical model. Ho, indeed, is a rare-earth ferromagnet with conical magnetic ordering, whose magnetization vector rotates around the c -axis, if one moves along it. In this case, the inhomogeneity is expected to be intrinsically present in the Ho layer, however the multilayer geometry $F_1/F/F_1$ is still needed. For these heterostructures a theoretical explanation is also available [18, 19].

In this framework the properties of permalloy can be particularly useful since it is well known that, if grown under certain specific conditions, it can form stripe-domains [20–22]. This is realized when its thickness exceeds a critical value, d_{cr} , which depends on the growth parameters such as, among others, the deposition rate and the substrate temperature [21, 23, 24]. In this configuration the magnetization vector lies mainly in-plane, parallel to the stripe direction in all domains, but it develops an out-of-plane component which goes alternately upward and downward. The out-of-plane component of the different stripes is therefore aligned antiparallel with Néel-domain-walls in between, in which the magnetization rotates coherently. Recently, we extensively described the magnetic properties of Py and we characterized the stripe-domain (SD) regime [25]. We also showed that below d_{cr} exists a broad regime, approximately between $0.5 d_{cr}$ and d_{cr} , where the magnetization can easily become inhomogeneous without being arranged in stripes. We called this state emerging SD (ESD) regime. The intrinsic magnetic inhomogeneity of Py implied in the occurrence of a SD phase led us to investigate the possibility of using it as possible generator for triplet correlations. The question is whether it is possible to have S/F/S (or S/F) structures where the conversion is intrinsically provided by the F-layer itself, due to its magnetic configuration, as proposed in the [7].

In this article we investigate the temperature dependence of the parallel upper critical field, $H_{c2||}(T)$, of simple Nb/Py/Nb trilayers and Nb/Py bilayers. The thickness of the Nb layers, d_{Nb} , is kept constant at 25 nm while the thickness of the Py layer, d_{Py} , is varied across the different thickness regimes: homogeneous (H), ESD and SD. For the trilayer with d_{Py} in the ESD regime, namely for $125 \text{ nm} \lesssim d_{Py} \lesssim 300 \text{ nm}$, a 2D–3D dimensional crossover (DCO) was observed at $T \simeq 0.9 T_c$, where T_c is the superconducting critical temperature of the system. Moreover, a clear kink is present in the $H_{c2||}(T)$ curves of Nb/Py bilayers when $d_{Py} = 200 \text{ nm}$. These observations, which we attribute to an increased effective thickness of the superconducting layer, cannot be explained within the spin-singlet proximity effect, because of the short coherence length of Py, estimated to be about 1.9 nm [6, 26]. The results are rather compatible with a long-range spin-triplet proximity effect, induced by the inhomogeneous magnetic configuration of the ESD regime.

2. Experimental methods

Nb/Py/Nb trilayers and Nb/Py bilayers were grown on Si(100) substrates by ultrahigh vacuum dc diode magnetron sputtering at an Ar pressure of 2.25×10^{-3} Torr after obtaining a base pressure of 1.5×10^{-8} Torr. The substrates were nominally kept at room temperature during the deposition process. The typical deposition rates were 0.25 nm s^{-1} for Nb and 0.30 nm s^{-1} for Py measured by a quartz crystal monitor previously calibrated by low-angle x-ray reflectivity measurements on deliberately deposited thin films of each material. The prepared samples are unstructured and the typical in-plane dimensions are $5 \times 10 \text{ mm}^2$. Samples have constant Nb thickness, $d_{Nb} = 25 \text{ nm}$, and variable Py thickness with d_{Py} in the range 20–430 nm. Thanks to the presence of a movable shutter in the deposition chamber, which selectively covers the substrates, three different samples can be grown in the same deposition run. A single 25 nm-thick Nb film and several single Py films, having the same thickness as the Py layers in the corresponding hybrids, were also deposited and characterized for comparison.

Single layers of Py were magnetically characterized in-depth using magnetic force microscopy (MFM), ferromagnetic resonance (FMR), SQUID magnetometry and magnetoresistance measurements (MR). Details of the used techniques can be found in the [25].

The resistive transitions of unstructured multilayers were performed using an in-line four-terminal geometry with a constant bias current of $500 \mu\text{A}$. The distance between the current (voltage) pads was about 8 mm (3 mm). The samples, mounted on a copper block and placed at the center of a NbTi superconducting

solenoid, were immersed in a ^4He cryostat. During measurements the temperature stabilization was around 1 mK.

3. Results

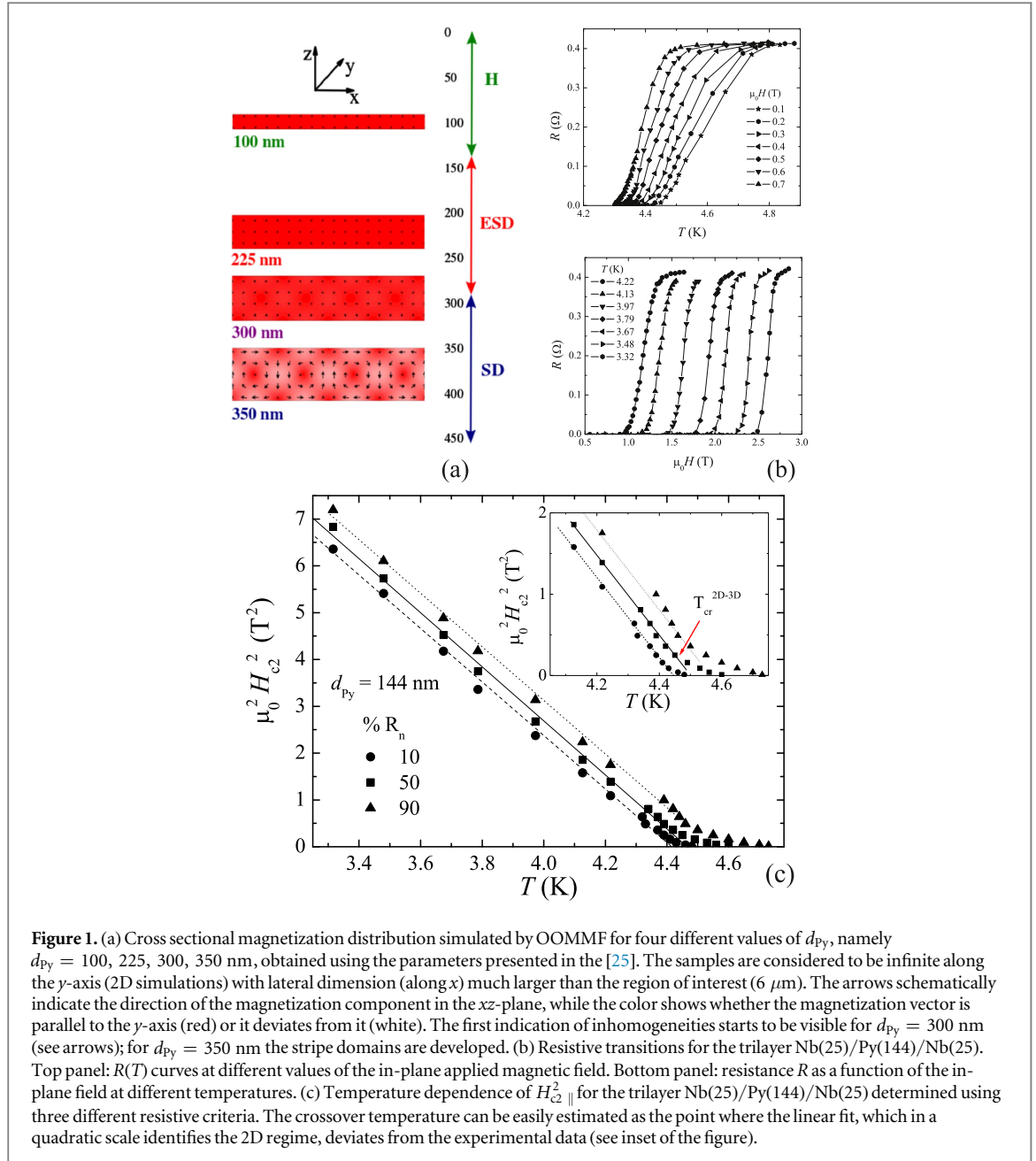
3.1. Magnetic characterization

A detailed magnetic characterization of Py as a function of the layer thickness was recently presented in the [25]. MFM measurements on unstructured Py single layers [25] showed that magnetic SDs are visible only above a thickness $d_{\text{py}} \simeq 300$ nm, which we defined to be the critical thickness value d_{cr} , i. e. the lower boundary of the SD regime. For $d < d_{\text{cr}}$, two different regimes could be recognized: a homogeneous regime (H), for $d \lesssim 125$ nm, and a so-called ESD regime, for $125 \text{ nm} \lesssim d \lesssim 300$ nm. For the samples in the ESD regime, the MFM measurements did not provide evidence of an inhomogeneous magnetic configuration, except for thicknesses close to the transitions to the SD regime. This suggests that either the magnetization is fully in-plane or the out-of-plane component is too weak to be detected by the MFM technique [25]. However, signatures of inhomogeneity emerged with other measurement techniques. Magnetic hysteresis loops $M(H)$ for samples in the ESD regime showed hints of a linear dependence before the saturation was reached. This linear behavior in general is a well-known feature which signals the presence of SDs and it is attributed to the coherent rotation of the stripes before the saturation. The observation of such dependence in the ESD regime, although less pronounced, is an indication of a certain degree of inhomogeneity, even if not necessarily in the stripe-form. The tendency of the samples in the ESD regime to have an inhomogeneous configuration emerged more clearly by looking at structured samples [25], in particular in the domain-wall configuration and in the outcome of MR measurements. The domain wall configuration of the structured samples was reproduced by simulations realized with the object oriented micromagnetic framework (OOMMF) software for all the three different regimes [25].

Since the devices we studied here are unstructured, we performed simulations on semi-infinite samples, for different thicknesses. The results are shown in figure 1(a). The magnetic parameters used (exchange stiffness constant, A , saturation magnetization, M_s , and out-of-plane anisotropy, K_{\perp}) are the same as in the [25], namely $A = 13 \times 10^{-12} \text{ J m}^{-1}$, $M_s = 8.59 \times 10^5 \text{ A m}^{-1}$, and $K_{\perp} = 5.6 \times 10^3 \text{ J m}^{-3}$. The sample is considered to be infinite along the y - and x -axes while it spans the thickness of the sample along the z -axis. The cross section shown in the figure is thus taken in the xz -plane, while the y -axis points inside this plane and represents the direction along which the sample is initially magnetized, prior to the magnetic measurements. The color code indicates the direction of the magnetization with respect to the y -axis, namely red and white mean the magnetization is respectively parallel or perpendicular to the initial direction. Black arrows indicate the orientation of the components of the magnetization in the xz -plane. The magnetization in the thinner samples ($d_{\text{py}} = 100, 225$ nm topmost sketches) stays parallel to the initial state (full dark color, indicating an homogeneous magnetic state), while deviations from such state begin to occur at $d_{\text{py}} = 300$ nm, for which areas with magnetization perpendicular to the y -axis start to appear. As d_{py} increases, the size and density of such areas increase and the magnitude of the component of the magnetization which deviates from the y -axis also grows. At $d_{\text{py}} = 350$ nm (bottom sketch), the domains with an out-of-plane magnetization component become wider, with thinner domain walls in between. This is the SD regime. Thus, from these simulations no clear evidence of inhomogeneity appear for the ESD regime. However, it is possible that the ‘semi-infinite’ approximation only partially reproduces the real physics of our devices. Furthermore, in the simulations the role of the proximity with superconducting layer is not taken into account. In the study below, the magnetic field is applied parallel to the film plane, and consequently the magnetic flux perpendicular to interface is minimum, however the diamagnetic nature of Nb below T_c could influence the magnetic configuration at the interface.

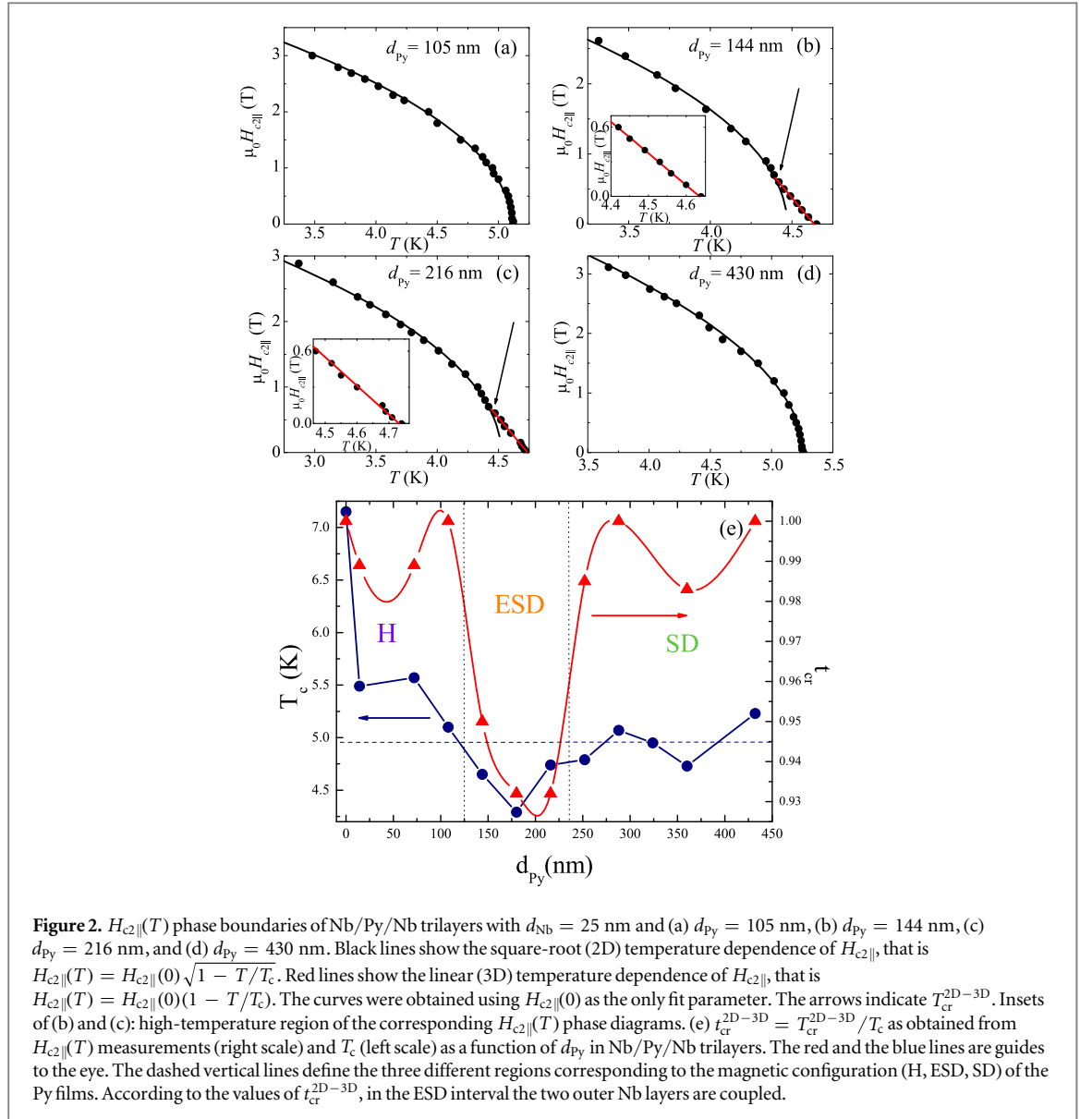
3.2. Superconducting transport properties of Nb/Py/Nb trilayers

In order to determine the $H_{c2}(T)$ phase diagrams of the Nb/Py/Nb trilayers, the resistance R was measured either as a function of the temperature T (at a fixed applied magnetic field H) or as a function of H (at a fixed T). The field was applied in-plane, perpendicular to the direction of the bias current. In figure 1(b) a selection of $R(T)$ and $R(H)$ curves for the trilayer Nb(25)/Py(144)/Nb(25) is presented (numbers in parentheses indicate the thickness expressed in nanometers). T_c was defined at $T_{50\%}$, namely at the temperature at which the resistance value is 50% of the normal state resistance R_N , measured at $T = 10$ K. Before measuring, a strong magnetic field (approximately 1 T) was applied in the plane of the substrate at low temperatures and then removed. This was done in order to ‘induce’ the stripes in the SD regime, and for consistency in the other two regimes. The width of the transitions at zero field, defined by $T_{90\%} - T_{10\%}$, is about 200 mK for all the samples and does not increase when a field is applied. The single Nb film 25 nm-thick has a critical temperature (at zero field) $T_c = 6.5$ K and shows a two-dimensional (2D) behavior [27] ($H_{c2\parallel}(T) \propto \sqrt{1 - T/T_c}$) in the whole investigated temperature



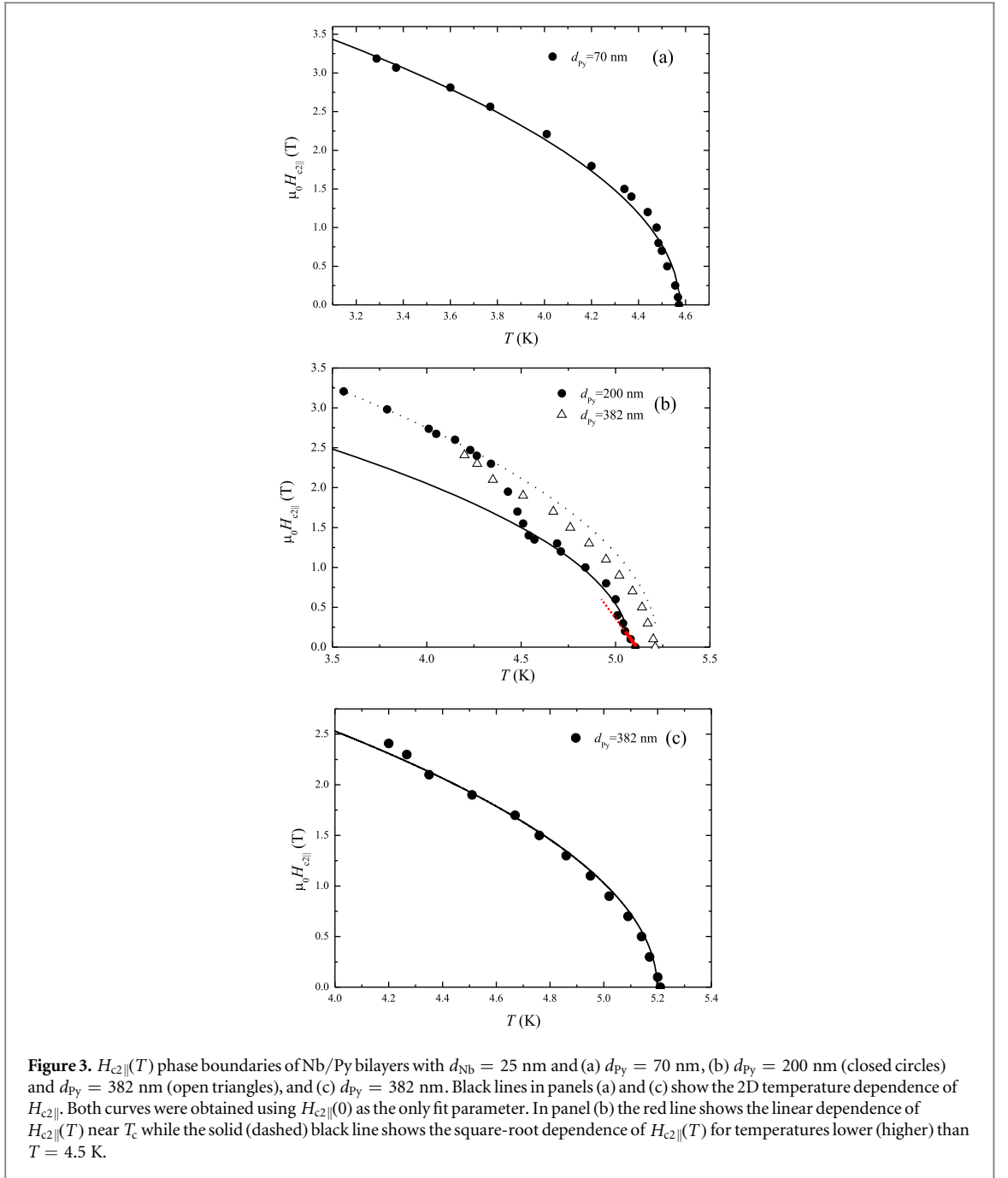
range. The Ginzburg–Landau (GL) coherence length at zero temperature, $\xi_{\text{GL}}(0)$, was extracted from the linear temperature dependence of the perpendicular upper critical field $H_{c2\perp}(T) = (\phi_0/2\pi\xi_{\text{GL}}^2(0))(1 - T/T_c)$ [28]. The obtained value is $\xi_{\text{GL}}(0) \simeq 10$ nm, implying a superconducting coherence length $\xi_s(0) = (2/\pi)\xi_{\text{GL}}(0) \simeq 7$ nm.

The $H_{c2\parallel}(T)$ phase diagrams for a representative set of the trilayers are presented in figure 2. The samples Nb(25)/Py(105)/Nb(25) (figure 2(a)) and Nb(25)/Py(430)/Nb(25) (figure 2(d)) show a 2D like behavior in the whole temperature range. The black line is the square-root temperature dependence of $H_{c2\parallel}(T)$ obtained leaving $H_{c2\parallel}(0)$ as the only fitting parameter. The first Py thickness is in the H regime, the latter in the SD regime. In both cases, thus, the Nb layers behave as the isolated single layer. Like in any conventional S/F interface, the singlet component cannot penetrate the Py layer more than 1–2 nm (on both sides), so the Nb layers result isolated. In the ESD regime, instead, the behavior is quite different. For both Nb(25)/Py(144)/Nb(25) (figure 2(b)) and Nb(25)/Py(216)/Nb(25) (figure 2(c)) there is a 2D–3D dimensional crossover (DCO) at a temperature $T_{\text{cr}}^{2\text{D}-3\text{D}} \simeq 0.9 T_c$. Close to T_c , $H_{c2\parallel}(T)$ is linear, as for a three-dimensional (3D) system, while at $T_{\text{cr}}^{2\text{D}-3\text{D}}$ it presents a square-root behavior of a 2D system. The insets in these panels show an enlargement of the data for temperatures close to T_c where the linear behavior of $H_{c2\parallel}(T)$ is much more evident. The DCO indicates a change in the dimensionality of the superconducting layers (in relation to ξ_{GL}) and can be explained within the framework of a long-range proximity effect. The peculiar inhomogeneous magnetic configuration of the ESD layer generates equal-spin triplet Cooper pairs which can ‘leak’ into Py, therefore extending the effective



thickness of the superconducting layers. One possible explanation is that the effect, as a consequence of the presence of magnetic inhomogeneities in Py, is due to a coupling between the top and bottom Nb layer across the Py. However, given the length scales involved, this seems to be very unlikely. The penetration length of the triplet component, indeed, is expected to be limited by the spin diffusion length, ℓ_{sf} , which for Py is relatively small, about 5 nm [26, 30], much shorter than the Py thickness. The reason why the 3D-behavior is observed only above $T_{\text{cr}}^{2\text{D}-3\text{D}}$ has to do with the temperature dependence of the coherence length

$\xi_s(T) = \xi_s(0)/\sqrt{1 - T/T_c}$ and it will be more extensively discussed later. The previous analysis of the $H_{c2||}(T)$ phase diagrams is summarized in figure 2(e) (right hand scale) where the reduced crossover temperature $t_{\text{cr}}^{2\text{D}-3\text{D}} = T_{\text{cr}}^{2\text{D}-3\text{D}}/T_c$ is reported for all trilayers. The crossover temperature can be more easily estimated by plotting $H_{c2||}^2$ versus T , as shown in figure 1(c) for the trilayer with $d_{\text{Py}} = 144$ nm: $t_{\text{cr}}^{2\text{D}-3\text{D}}$ is the value at which the dependence deviates from the linear fit (see inset of the figure). For values of d_{Py} up to 125 nm (region H) and larger than 300 nm (region SD) $t_{\text{cr}}^{2\text{D}-3\text{D}}$ is essentially equal to 1 (namely there is no crossover), while in the ESD region it is $t_{\text{cr}}^{2\text{D}-3\text{D}} = 0.93-0.95$. As a remark, as we show in figure 1(c), the choice of the 50% of R_N as a criterion to obtain the $H_{c2||}(T)$ -phase boundaries of the different heterostructures does not alter the main results presented above, since the position of the 2D–3D crossover in the $H_{c2||}(T)$ -plane, if present, is confirmed also if the 10% or 90% criteria for the determination of T_c are adopted. The dependence of T_c (at zero field) on the thickness of Py, plotted in figure 2(e) (left scale), also shows a clear variation, with a dip in the middle of the ESD regime. Notably, T_c for the trilayer with $d_{\text{Py}} = 170$ nm is at least 0.5 K lower than T_c of the trilayers with d_{Py} above 300 nm. This is fully consistent with the explanation for the 2D–3D crossover. When triplets are formed, the leakage of Cooper pairs into the Py layer depletes the superconducting order parameter on the S side



of the S/F interface. The length scale of the effect is determined by $\xi_S(0)$ and since the thickness of the superconducting layer is only a few times $\xi_S(0)$, the result is a suppression of T_c of the whole layer. This observation was predicted theoretically [31] and recently demonstrated in different systems [13, 17].

3.3. Superconducting transport properties of Nb/Py bilayers

Since the observed behavior is expected to be due to two separate S-layers at top and bottom, decoupled by the thick Py layer, a similar effect should be observed for a Nb/Py bilayer. In figure 3 we show the phase diagrams $H_{c2||}(T)$ for the bilayers Nb/Py, with $d_{\text{Nb}} = 25$ nm and $d_{\text{Py}} = 70, 200, 382$ nm, in order to explore again the H, ESD, and SD regions, respectively.

As expected, for $d_{\text{Py}} = 70$ and 382 nm (figures 3(a) and (c), respectively; H and SD regime) the dependence is 2D in the whole temperature range, as for the trilayers. The black lines are the fits of the 2D relation, obtained using $H_{c2||}(0)$ as the only fitting parameter. The bilayer with $d_{\text{Py}} = 200$ nm (ESD regime), which in figure 3(b) is compared with the Nb(25)/Py(382) one, also shows a DCO but in this case the transition is 2D–2D, at about $T = 4.5$ K ($t_{\text{cr}}^{2\text{D}-2\text{D}} \simeq 0.9$). Very close to T_c there is a hint of 2D–3D dimensional crossover, but the small range makes it difficult to judge whether it is a real feature or an artifact. A possible explanation for the observations is given later.

4. Discussion

From the analysis of the phase diagrams of the S/F/S trilayers it emerges that a 2D–3D DCO is present only when the thickness of the Py layer is in the ESD regime. Such a DCO is typically observed in S/N multilayers [27] (with N a normal metal), and it is ascribed to a change in the dimensionality of the superconducting layer with respect to the coherence length $\xi_{GL}(T)$. This occurs when the thickness, d_S , of a single layer is lower than (or of the order of) the coherence length but the total thickness of two (or more) layers is larger. When ξ_{GL} , which is temperature dependent, becomes of the order of the spacer length d_N , the S-layers are coupled and the $H_{c2\parallel}(T)$ dependence becomes linear (3D-behavior), provided that $d_N < 2\xi_N$. The latter condition makes a DCO unexpected in a S/F multilayer unless d_F is very small [32] or the ferromagnet is diluted [28]. In our case $d_F \gg \xi_F$. For Py, indeed, $E_{ex} \simeq 200$ meV [6] and the diffusion coefficient can be obtained via the relation $D_F = (1/3)v_F\ell_F$, where $v_F = 2.2 \times 10^5$ m s⁻¹ [6] is the Fermi velocity and ℓ_F is the mean free path. ℓ_F can be obtained from $\rho\ell_F = 31.5 \times 10^{-6}$ $\mu\Omega$ cm² [29], knowing that $\rho = 20$ $\mu\Omega$ cm (value measured at low temperatures in our samples). Thus, from $\xi_F = \sqrt{\hbar D_F/E_{ex}}$, we obtain $\xi_F \simeq 1.9$ nm. The scenario is different if long-range equal-spin triplet Cooper pairs are induced, because of the inhomogeneity in the ESD regime. In this case the coherence length is given by $\xi_F^T(T) = \sqrt{\hbar D_F/2\pi k_B T}$ [7]. By using the value of D_F extrapolated above, at $T = 4.2$ K, $\xi_F^T(T = 4.2 \text{ K}) \simeq 20$ nm $\gg \xi_F$. While estimating the length scale of the proximity effect, also the spin diffusion length ℓ_{sf} , typically the main limiting factor, has to be taken into account. For strong ferromagnets such as Co, ℓ_{sf} is about 60 nm [30, 33], while for Py is expected to be much shorter, $\ell_{sf} \simeq 5$ nm [26, 30]. Therefore, the possibility that the two Nb layers are coupled across a 200 nm-thick Py layer is extremely unlikely even though one can consider that for $d_{Py} = 144$ nm the order parameter is attenuated at the center of the ferromagnetic layer by the factor $\exp(-d_{Py}/2\ell_{sf}) = 5.5 \times 10^{-7}$, value that is much larger than $\exp(-d_{Py}/2\xi_F) = 3.5 \times 10^{-17}$, as expected in the spin-singlet scenario. The DCO in this case seems to be the result of the extended effective S-thickness, with only the proximity of the top S/F interface contributing. A similar DCO, indeed, was observed for a simple Nb/Cu bilayer [34]. Whether the estimated short penetration length, due to ℓ_{sf} , is enough to explain a transition from a 2D to a 3D regime, is unclear. The spin diffusion length for Py was evaluated with a two-current model [35], considering a homogeneously magnetized layer of Py. In the ESD regime, however, there is a certain degree of inhomogeneity which could be responsible for mixing the two channels, resulting in an enhanced effective spin-diffusion length [36, 37]. Moreover, the observation of a dip in the T_c versus d_{Py} in correspondence of the ESD regime is a further strong indication of the origin of the effects being the leakage triplet Cooper pairs into Py. If the suppression of T_c in the trilayer was only due to the leakage of spin-singlet Cooper pairs, the critical temperature values in the studied Py thickness range should be d_{Py} independent, because in this case $d_{Py} \gg \xi_F$. The reason why the triplet generation occurs only in the ESD regime has to do with the particular inhomogeneity of the magnetization in this thickness region. One could expect that the SDs in the SD regime, and in particular the rotating domain-walls between stripes, could be a source of singlet-to-triplet conversion. However, this is not what the outcome of the measurements suggests. This can be explained by looking at the length scale of the inhomogeneities in this case. The typical width w of the stripes is of the same order of magnitude of the thickness d_{Py} [38], thus $w > 300$ nm, with the domain-wall width δ_w of a similar order of magnitude ($\delta_w \approx w/3$). This length scale is large, if compared to ξ_F . As a consequence the short-ranged Cooper pairs injected into Py ‘feel’ a homogeneous magnetization and therefore there is no conversion. In the ESD regime, instead, the magnetization is not arranged in stripes. Because of the weak perpendicular anisotropy the magnetic moments have a ‘tendency’ to rotate out-of-plane, and the result is the presence of more localized inhomogeneities [38]. The 3D–2D transition which occurs by lowering the temperature and increasing the magnetic field, is probably due to the alignment of the magnetization which is no longer inhomogeneous. Surprisingly, the transition field ($\simeq 0.5$ T) is an order of magnitude higher than the field of complete saturation [25]. A possible explanation could be that the uniaxial anisotropy and the interface roughness lead to a residue of unsaturated and misaligned moments close to the interface. Triplet correlations can actually be induced by quite small amounts of misaligned moments, as already noticed in early work involving CrO₂: in sandwiches of NbTiN/CrO₂ [8] and MoGe/CrO₂ [11] triplet correlations were found in the absence of engineered magnetic inhomogeneities and they were sustained in fields well above the nominal saturation field of CrO₂.

The measurements performed on the bilayers confirm the picture described above, with a DCO observed only for the ESD regime, and at an even higher magnetic field ($\simeq 1.5$ T). Why the transition in this case is 2D–2D is not entirely trivial. In the [39] a 2D–2D DCO was observed in a Pb/Ge multilayer because the total thickness of the coupled Pb layers is still in the 2D regime. Here, if we fit the $H_{c2\parallel}(T)$ curve of Nb(25)/Py(200) at low temperatures, which coincides with the curve of Nb(25)/Py(382) (figure 3) using the expression $\mu_0 H_{c2\parallel}(0) = (\sqrt{12} \phi_0)/(2\pi\xi_{GL}(0)d_{S_{eff}})$ [40], we obtain an effective superconducting thickness $d_{S_{eff}}$ of about 20 nm. This is also the thickness inferred from the Nb(25)/Py(382) data. A lower thickness than the actual layer thickness is to be expected for an S/F bilayer where strong pair breaking on the F-side of the interface lowers the Cooper pair density. From the extrapolation close to T_c , instead, the estimated $d_{S_{eff}}$ value is approximately

25 nm. On the one hand, this is significantly larger than the low-temperature thickness; on the other hand, it is thin enough to be in the 2D regime. Maybe even more importantly, the crossover in the $H_{c2\parallel}(T)$ curve of Nb(25)/Py(200) suggests that not only the effective thickness decreases, but that also T_c increases, which would be in full agreement with the triplet leakage scenario. An effective thickness of 25 nm could even be expected to arise from the limits set by the small spin diffusion length. Unfortunately, this is difficult to reconcile with the coupling between the S-layers observed in the trilayers. Both experiments point to the presence of triplets in the ESD regime, but with different length scales. The key here still lies in a better understanding of the inhomogeneous ESD regime, which is not yet available. Just summing the thicknesses of the superconducting and proximized layer can give a qualitative idea but it is a too simplistic analysis. First of all, the extrapolation of the $d_{\text{S,eff}}$ from $H_{c2\parallel}(T)$ is not completely reliable for a proximized system, where T_c and therefore $H_{c2}(0)$ are modified. In general, the whole picture can be better described by saying that in these systems the order parameter adjusts itself, such that it can sustain the highest critical field [41]. The outcome, therefore, is the result of the interplay between parameters such as $\xi_S(T)$ and ξ_F (or $\xi_F^T(T)$), with the magnetic field playing a crucial role in determining the magnetic configuration and thus the (possible) triplet generation.

5. Conclusions

To summarize, we showed that equal-spin triplet proximity can be induced in simple Nb/Py/Nb trilayers and Nb/Py bilayers by exploiting the intrinsic magnetic inhomogeneities of Py. The conclusion is indirectly inferred from the DCO observed in the phase diagram $H_{c2\parallel}(T)$. Indeed, the DCO, observed for both tri- and bilayers (2D–3D transition in the first case, 2D–2D in the latter), cannot be described by a short-range singlet proximity. The crossover appears only for Py thicknesses in the ESD regime, which seems to provide the optimal degree of inhomogeneous magnetization for the singlet-to-triplet conversion. The interpretation is confirmed by the dependence T_c versus d_{Py} for the trilayers, which shows a strong suppression of T_c in the ESD regime where the leakage of long-range Cooper pairs is maximum.

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References

- [1] Fertig W A, Johnston D C, DeLong L E, McCallum R W, Maple M B and Matthias B T 1977 *Phys. Rev. Lett.* **38** 987–90
- [2] Buzdin A I 2005 *Rev. Mod. Phys.* **77** 935–76
- [3] Lazar L, Westerholt K, Zabel H, Tagirov L R, Goryunov Y R, Garif'yanov N N and Garifullin I A 2000 *Phys. Rev. B* **61** 3711–22
- [4] Cirillo C, Prischepa S L, Salvato M, Attanasio C, Hesselberth M and Aarts J 2000 *Phys. Rev. B* **72** 144511
- [5] Ryazanov V V, Oboznov V A, Rusanov A Y, Veretennikov A V, Golubov A A and Aarts J 2001 *Phys. Rev. Lett.* **86** 2427–30
- [6] Robinson J W A, Piano S, Burnell G, Bell C and Blamire M G 2006 *Phys. Rev. Lett.* **97** 177003
- [7] Bergeret F S, Volkov A F and Efetov K B 2001 *Phys. Rev. Lett.* **86** 4096–9
- [8] Keizer R S, Gönnerwein S T B, Klapwijk T M, Miao G, Xiao G and Gupta A 2006 *Nature* **439** 825–7
- [9] Khaire T S, Khasawneh M A, Pratt W P and Birge N O 2010 *Phys. Rev. Lett.* **104** 137002
- [10] Sprungmann D, Westerholt K, Zabel H, Weides M and Kohlstedt H 2010 *Phys. Rev. B* **82** 060505
- [11] Anwar M S, Veldhorst M, Brinkman A and Aarts J 2012 *Appl. Phys. Lett.* **100** 052602
- [12] Robinson J W A, Witt J D S and Blamire M G 2010 *Science* **329** 59–61
- [13] Leksin P V, Garif'yanov N N, Garifullin I A, Fominov Y V, Schumann J, Krupskaya Y, Kataev V, Schmidt O G and Büchner B 2012 *Phys. Rev. Lett.* **109** 057005
- [14] Zdravkov V I et al 2013 *Phys. Rev. B* **87** 144507
- [15] Banerjee N, Smiet C B, Smits R G J, Ozaeta A, Bergeret F S, Blamire M G and Robinson J W A 2014 *Nat. Commun.* **5** 3048
- [16] Flokstra M G et al 2015 *Phys. Rev. B* **91** 060501
- [17] Singh A, Voltan S, Lahabi K and Aarts J 2015 *Phys. Rev. X* **5** 021019
- [18] Fritsch D and Annett J F 2014 *New J. Phys.* **16** 055005
- [19] Fritsch D and Annett J F 2015 *Phil. Mag.* **95** 441–50
- [20] Saito N, Fujiwara H and Sugita Y 1964 *J. Phys. Soc. Japan* **19** 421–2
- [21] Amos N, Fernandez R, Ikkawi R, Lee B, Lavrenov A, Krichevsky A, Litvinov D and Khizroev S 2008 *J. Appl. Phys.* **103** 07E732
- [22] Belkin A, Novosad V, Iavarone M, Fedor J, Pearson J E, Petrean-Troncilli A and Karapetrov G 2008 *Appl. Phys. Lett.* **93** 072510
- [23] Ben Youssef J, Vukadinovic N, Billet D and Labrune M 2004 *Phys. Rev. B* **69** 174402
- [24] Dastagir T, Xu W, Sinha S, Wu H, Cao Y and Yu H 2010 *Appl. Phys. Lett.* **97** 162506
- [25] Voltan S, Cirillo C, Snijders H J, Lahabi K, García-Santiago A, Hernández J M, Attanasio C and Aarts J 2016 *Phys. Rev. B* **94** 094406

- [26] Moraru I C, Pratt W P and Birge N O 2006 *Phys. Rev. B* **74** 220507
- [27] Chun C S L, Zheng G G, Vicent J L and Schuller I K 1984 *Phys. Rev. B* **29** 4915–20
- [28] Schöck M, Sürgers C and von Löhneysen H 2000 *Eur. Phys. J. B* **14** 1–10
- [29] Mayadas A F, Janak J F and Gangulee A 1974 *J. Appl. Phys.* **45** 2780–1
- [30] Bass J and Pratt W P Jr 2007 *J. Phys.: Condens. Matter* **19** 183201
- [31] Fominov Y V, Golubov A A, Karminskaya T Y, Kupriyanov M Y, Deminov R G and Tagirov L R 2010 *JETP Lett.* **91** 308–13
- [32] Koorevaar P, Suzuki Y, Coehoorn R and Aarts J 1994 *Phys. Rev. B* **49** 441–9
- [33] Piraux L, Dubois S, Fert A and Belliard L 1998 *Eur. Phys. J. B* **4** 413–20
- [34] Otop A, Boogaard G R, Hendriks R W A, Hesselberth M B S, Ciuhu C, Lodder A and Aarts J 2003 *Europhys. Lett.* **64** 91–7
- [35] Dubois S, Piraux L, George J M, Ounadjela K, Duvail J L and Fert A 1999 *Phys. Rev. B* **60** 477–84
- [36] Rudiger U, Yu J, Zhang S, Kent A D and Parkin S S P 1998 *Phys. Rev. Lett.* **80** 5639–42
- [37] Kent A D, Rudiger U, Yu J, Thomas L and Parkin S S P 1999 *J. Appl. Phys.* **85** 5243–8
- [38] Hubert A and Schäfer R 1998 *Magnetic domains The Analysis of Magnetic Microstructure* (Berlin: Springer)
- [39] Neerincx D, Temst K, van Haesendonck C, Bruynseraede Y, Gilbert A and Schuller I K 1991 *Phys. Rev. B* **43** 8676–8
- [40] Jin B Y and Ketterson J B 1989 *Adv. Phys.* **38** 189–366
- [41] Nojima T, Kinoshita M, Nakano S and Kuwasawa Y 1993 *Physica C* **206** 387–92