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Second order anisotropy contribution in perpendicular magnetic tunnel junctions

A. A. Timopheev^{1,2,3}, R. Sousa^{1,2,3}, M. Chshiev^{1,2,3}, H. T. Nguyen^{1,2,3} & B. Dieny^{1,2,3}

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Hard-axis magnetoresistance loops were measured on perpendicular magnetic tunnel junction pillars of diameter ranging from 50 to 150 nm. By fitting these loops to an analytical model, the effective anisotropy fields in both free and reference layers were derived and their variations in temperature range between 340 K and 5 K were determined. It is found that a second-order anisotropy term of the form $-K_2 \cos^4 \theta$ must be added to the conventional uniaxial $-K_1 \cos^2 \theta$ term to explain the experimental data. This higher order contribution exists both in the free and reference layers. At $T = 300$ K, the estimated $-K_2/K_1$ ratios are 0.1 and 0.24 for the free and reference layers, respectively. The ratio is more than doubled at low temperatures changing the ground state of the reference layer from “easy-axis” to “easy-cone” regime. The easy-cone regime has clear signatures in the shape of the hard-axis magnetoresistance loops. The existence of this higher order anisotropy was also confirmed by ferromagnetic resonance experiments on FeCoB/MgO sheet films. It is of interfacial nature and is believed to be due to spatial fluctuations at the nanoscale of the first order anisotropy parameter at the FeCoB/MgO interface.

Magnetic anisotropy is a key feature of a ferromagnetic material playing a crucial role in technical applications of these materials. Generally, this phenomenon takes its origin from magnetic dipole-dipole, exchange and/or spin-orbit interactions. These interactions provide respectively shape, exchange and magnetocrystalline (magnetoelastic) anisotropies. One can also divide the magnetic anisotropy as arising from the bulk and/or from the surface or interface of the layer.

Concept of interfacial anisotropy was proposed in the pioneering work of L. Neel¹ predicting the perpendicular interfacial anisotropy as a result of the lowered symmetry at the surface/interface. This work was followed by experiments carried out on ultrathin NiFe films grown on Cu(111)² which confirmed the interfacial nature of the perpendicular magnetic anisotropy (PMA) observed in this system. Within the last fifty years, a lot of work has been carried out on interfacial anisotropy both from theoretical and experimental points of view^{3–8}. Nowadays, perpendicular interfacial anisotropy has become one of the main ingredients of novel magnetic memory elements employing out-of-plane magnetized (perpendicular) magnetic tunnel junctions (pMTJ) stacks^{9–11}. In such structures, perpendicular anisotropy of the free layer is provided by the interface between FeCoB and MgO layers while in the reference layer, it is additionally enhanced by exchange coupling with Co/Pt or Co/Pd multilayers¹² with PMA of interfacial nature as well.

Taking into account the system symmetry, the PMA energy density originating from the interface can be written as:

$$E_{PMA} = -\frac{1}{t}(K_{1s} \cos^2 \theta + K_{2s} \cos^4 \theta + \dots), \quad (1)$$

where θ is the angle between magnetization and normal to the plane of the layers, K_{1s} , K_{2s} , ... are constants of the first and second order surface anisotropy energy per unit area and t is the thickness of the FM layer. One can then define effective bulk anisotropy constants which also include the demagnetizing energy for a thin film (CGS units): $K_1 = \left(\frac{K_{1s}}{t} - 2\pi M_S^2\right)$, $K_2 = \frac{K_{2s}}{t}$. In case of very thin Fe films magnetization saturation parameter M_S is typically reduced in comparison with its bulk value¹³. If $K_1 > 0$, $K_2 < 0$ and $0.5 < -K_2/K_1 < 1$, the ground state of

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the system will correspond to so-called “easy-cone” regime, or canted state. In the easy-cone regime, the magnetization is tilted away from the symmetry axis by angle θ_c given by $\cos^2 \theta_c = -K_1/2K_2$. Due to the axial symmetry, the system energy remains invariant around a cone with opening angle θ_c yielding a so-called “easy-cone” anisotropy. Quite frequently, in systems where interfacial anisotropy is present, the first order term proportional to K_{1s} dominates the higher order term proportional to K_{2s} . However, the influence of this second order term has been clearly observed experimentally around the spin-reorientation transition region where the demagnetizing energy partially or fully balances the $K_{1s} \cos^2 \theta$ term (i.e. effective anisotropy K_i is close to zero)^{14–19}. $K_{2s} \cos^4 \theta$ term can arise due to peculiarities of atomic structure at the interface or as a result of non-uniform mechanical stresses existing at interfaces presenting a large crystallographic mismatch. Also, B. Dieny and A. Vedyayev have shown analytically that spatial fluctuations of the film thickness under $K_{1s} = \text{const}$ term can lead to a higher order $K_{2s} \cos^4 \theta$ term if the period of the fluctuations is lower than the exchange length of FM material²⁰. Recently, J. Sun has reported similar results²¹.

Experimental determination and understanding of magnetic anisotropy in FM layers and multilayers is very important towards the pMTJ stack optimization for future use in STT-MRAM applications. Experiments conducted on sheet films combining magnetometry (VSM, SQUID etc.) with ferromagnetic resonance (FMR) allow the determination of magnetic anisotropy constants at sheet film level. However, in the context of STT-MRAM development, it is also important to know how these anisotropy parameters are affected by the patterning process and how they are distributed from dot to dot in an array of magnetic tunnel junctions. Magnetotransport measurements with field applied in the plane of the layers provide a convenient way to determine the anisotropy characteristics in pMTJ. Magnetic field applied along hard-axis tilts the magnetic moments of both layers away from the normal to the plane direction which produces a change in the tunneling conductance of the system. The curvature of the obtained MR(H) dependences and their different shapes for initially parallel and antiparallel magnetic configurations allow direct extraction of the effective anisotropy fields in both magnetic electrodes assuming that micromagnetic distortions are not developing much under the applied field (macrospin approximation). Such analysis can be performed on an automated wafer prober equipped with an electromagnet allowing large-scale analysis of pMTJ pillar arrays with good statistics. For deeper analysis on a limited number of pMTJs, experiments can also be carried out on experimental setups such as Physical Property Measuring System (PPMS) allowing measurements in a wide range of temperatures and magnetic fields.

In this study, we investigated the anisotropy in pMTJs via hard-axis magnetoresistance loops analysis and derived the effective anisotropy fields of these pMTJ pillars of various nominal diameters ranging from 50 nm to 150 nm. The 1st and 2nd order magnetic anisotropies in both layers were derived as well as their temperature dependences. It was found that a significant $K_{2s} \cos^4 \theta$ term is present in both free and polarizing layers. In this term, K_{2s} has a negative sign which can result in an easy-cone magnetic state with canted remanence of the magnetic layers.

Experimental Details

pMTJ pillars array with nominal diameters ranging between 50 nm and 500 nm were fabricated from an MTJ stack grown by DC and RF magnetron sputtering on thermally oxidized Si substrate. The stack is a bottom pinned pMTJ with the composition close to ref. 22. Enumerating from the substrate, the stack is Ta(5)/Pt(5)/[Co(0.4)/Pt(0.4)]₆/Co(0.4)/Ru(0.42)/[Co(0.4)/Pt(0.4)]₂/Co(0.4)/Ta(0.3)/Fe₆₀Co₂₀B₂₀(0.9)/MgO(1.2)/Fe₆₀Co₂₀B₂₀(0.9)/Ta(0.3)/Fe₆₀Co₂₀B₂₀(0.8)/MgO(0.4)/Ta(1.2)/Ru(5). The layers nominal thicknesses are in nm.

Saturation magnetization parameter of the free layer was measured to be 1030 emu/cm³. Current in-plane magnetotransport measurements yielded $R_{xA} = 5.7 \Omega \mu\text{m}^2$ and TMR = 126%. The second MgO barrier was introduced to increase the perpendicular anisotropy of the free layer. It has a negligible resistance-area (RA) product compared to the main tunnel barrier. Additional information and experiments on these samples can be found in ref. 23.

Statistical measurements of coercivity, coupling field and TMR values were performed using an automated wafer prober setup equipped with an electromagnet. Temperature-dependent measurements on single pMTJ pillars were carried out using PPMS system. Magnetoresistance loops were measured by applying a magnetic field along the easy and hard axis directions and passing a constant current through the pillars which amplitude was set not to exceed 30 mV across the tunneling barrier in the antiparallel configuration in order to minimize any spin-transfer-torque influence during the measurement. At each field point, the voltage drop was measured and the resistance determined. Magnetic field was swept from –6 kOe to +6 kOe and then back to –6 kOe with a constant sweep rate.

Analysis of Hard-Axis Magnetoresistance Loops

Assuming $K_1 \neq 0$, $K_2 = 0$, macrospin behavior and linear dependence of the tunneling conductance versus cosine of the relative angle between magnetization vectors in the two magnetic electrodes²⁴, one can analytically derive the hard-axis magnetoresistance as a function of applied field H for initially (at H = 0) parallel and antiparallel states:

$$MR(H) = 2 R_p R_{Ap} \frac{(R_p + R_{Ap} + (R_{Ap} - R_p))}{(\pm \sqrt{(1 - H^2/H_{\perp 1}^2)(1 - H^2/H_{\perp 2}^2)} + H^2/(H_{\perp 1} H_{\perp 2}))^{-1}}, \quad (2)$$

where $H_{\perp 1}$, $H_{\perp 2}$ are the effective perpendicular anisotropy fields in the two electrodes, R_p , R_{Ap} are the resistance values in parallel and antiparallel states, plus/minus sign of the square root corresponds to MR curve for the initially parallel/antiparallel state.

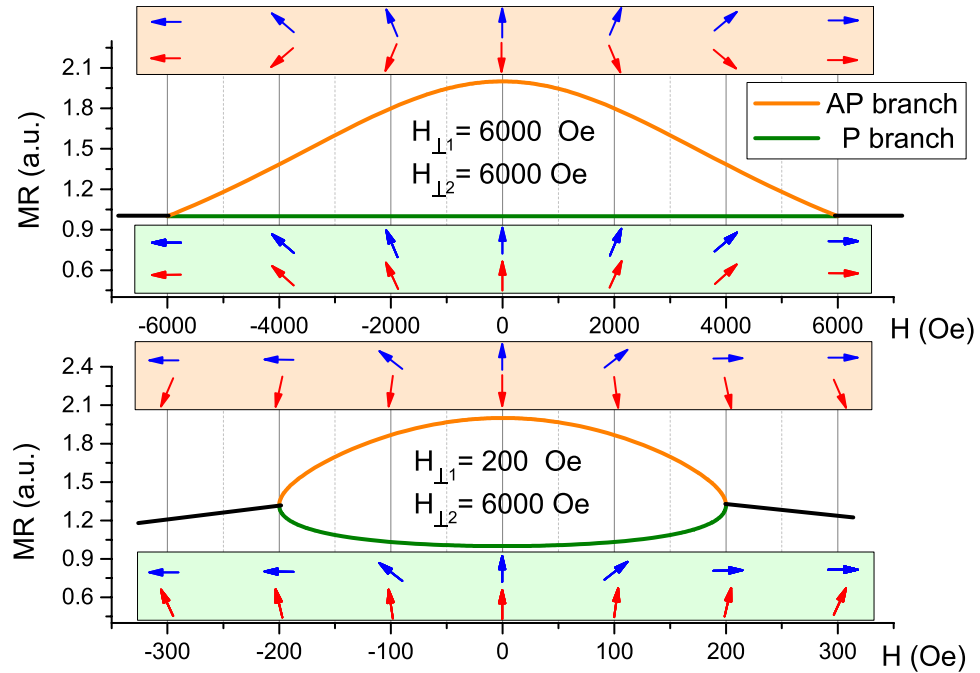


Figure 1. AP and P branches of hard-axis magnetoresistance loops calculated using model (2) assuming constant anisotropy field $H_{\perp 2}$ in the reference layer and two different anisotropy fields $H_{\perp 1}$ in the free layer. The arrows sets (blue for free layer and red for reference layer) below and atop of each graph show configuration of magnetic moments for AP and P branches respectively at several points of the in-plane magnetic field.

Figure 1 shows the variation of MR curves defined by Eq. (2) starting from the P or AP states with respect to $H_{\perp 1}/H_{\perp 2}$ ratio. When $H_{\perp 1}/H_{\perp 2} \ll 1$ (bottom graph), both curves starting from P or AP states have parabolic shape with similar curvatures. This behavior corresponds to the limit of strictly fixed reference layer. On the contrary, if both layers have the same anisotropy fields, $H_{\perp 1} = H_{\perp 2}$ (top graph), the resistance starting from P state will remain unchanged whatever the field (both magnetization rotates together), while the MR curve starting from the AP state will vary from R_{Ap} to R_p value. The variation of the curvatures with respect to $H_{\perp 1}/H_{\perp 2}$ ratio allows one to estimate $H_{\perp 1}$ and $H_{\perp 2}$ directly from the experiments by fitting the experimental hard-axis MR curve starting from P and AP states with expression (2). Knowing the magnetization saturation parameter and ferromagnetic film thickness, one can derive the surface anisotropy constant K_s from the relation: $\frac{H_{\perp 1} M_s}{2} = \left(\frac{K_s}{t} - 2\pi M_s^2 \right)$. If higher order anisotropy contributions have to be taken into account in (1) in order to improve the fits, then no analytical expression similar to Eq. (2) is available but the fitting of the MR(H) curves is still numerically feasible.

One should notice, however, that micromagnetic distortions, very strong interlayer coupling and superparamagnetic thermal fluctuations can play a role in the MR(H) dependences and worsen significantly the fitting quality. Analysis of easy-axis MR(H) loops can help to identify possible contributions of these effects.

Room-Temperature Easy-Axis Magnetoresistance Loops

Using the automated wafer prober setup, about 90 pillars of each diameter were measured to obtain statistically reliable information and to guide the choice of the samples for a further more detailed investigation of the anisotropy properties. 15-loop magnetoresistance hysteresis loops were measured on each device. The magnitudes of the TMR, coercive field and coupling field were extracted from the averaged loops. Few devices showing TMR < 90% were excluded from the statistical analysis. Figure 2(a) shows these three parameters as a function of pillar diameter. In average, all samples have a coercive field ~ 1.1 kOe, a coupling field ~ 90 Oe and TMR $\sim 113\%$. For most devices, the interlayer exchange coupling is ferromagnetic with a positive sign. It is hard to track its diameter dependence since the standard deviation is of the same order of magnitude as the same measured value. It is believed that fluctuations of the coupling field are mainly caused by damages of the pillar edges. No correlations are observed between H_p , H_c and TMR values. The coercivity and TMR are observed to weakly decrease versus pillar diameter, which can be ascribed to the appearance of micromagnetic distortions at pillar edges as the diameter increases (e.g. flower state). Individual easy-axis magnetoresistance loops of some selected devices at RT are shown in Fig. 2(b). The measurements were performed on a PPMS-based setup at room temperature. All measured devices have similar TMR amplitude and perfect rectangular shape with no evidence of any intermediate states between full P and AP configurations.

Temperature Dependent Measurements

In a hard-axis measurement of the MR(H) loops, the magnetization of the storage layer only rotates by 90° between the remanent state and the saturated state. The system has to be prepared either in the initial P

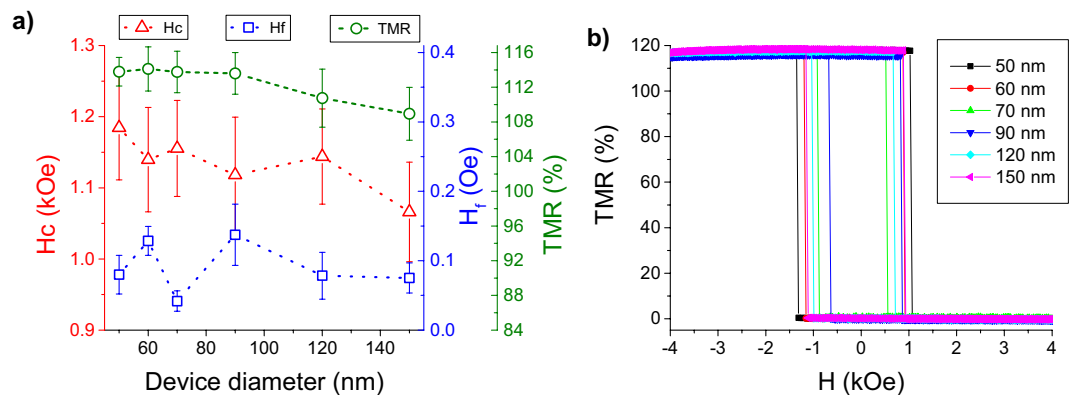


Figure 2. (a) Statistically averaged coercivity (H_c), coupling field (H_f) and TMR as a function of pillar diameter. The error bar heights represent the standard deviation over ~ 90 samples. (b) easy-axis magnetoresistance loops for the selected devices at room temperature (RT).

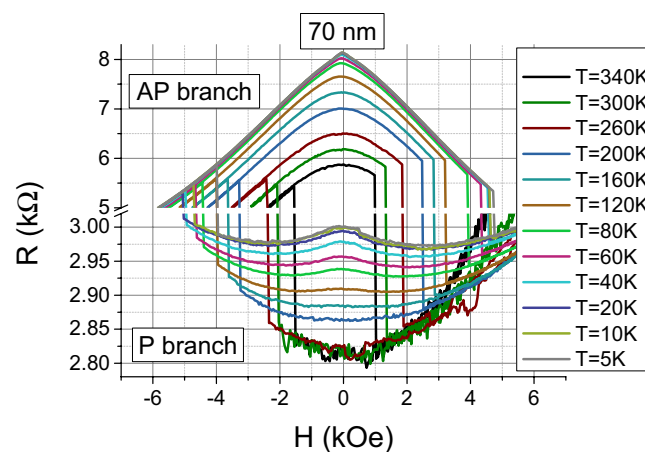


Figure 3. MR(H) loops at different temperatures measured on 70 nm device.

configuration or in the initial AP configuration giving two hard-axis hysteresis branches. If the field is strictly applied in the plane of the sample during the hard-axis measurement, the two MR(H) branches can be obtained according to the following protocol with 8 steps using a PPMS setup with rotating sample holder: 1) switch the pMTJ pillar in P state by applying the magnetic field along the easy axis, set $H = 0$ and rotate the pillar into hard-axis configuration; 2) make a MR(H) measurement from $H = 0$ to $H = H_{max}$; 3) repeat step 1; 4) make a MR(H) measurement from $H = 0$ to $H = -H_{max}$; 5) rotate the sample back to the position with field applied parallel to the normal to the plane (i.e. along easy axis) and set the pillar in AP state, set $H = 0$ and rotate the pillar into hard-axis configuration; 6) repeat step 2; 7) repeat step 5; 8) repeat step 4. By putting together MR(H) dependences obtained for negative and positive magnetic field sweeps, one finally obtains the two MR(H) branches corresponding to initially P and AP states, i.e. a full hard-axis MR loop.

We have implemented a simplified method for hard-axis MR(H) measurements. If the magnetic field is slightly tilted away from the hard axis by a few degrees, then the small out-of-plane remaining component of the applied field will allow the switching of the magnetization from the P hard-axis branch to the AP hard axis branch. Thermal fluctuations and interlayer exchange coupling across the tunnel barrier determine the minimal angle of misalignment necessary to observe these jumps between the two branches. In our samples where the coupling field is one order of magnitude lower than the switching field, it is enough to tilt the magnetic field by 3–4 degrees out-of-plane which slightly distorts the MR(H) curves, making them slightly asymmetric around the vertical axis. But at the same time, it allows obtaining the full hysteresis loop containing both AP and P branches in a much easier way than in the case where the field is applied strictly in-plane. Here and further, we will call AP/P branches those corresponding to the reversible parts of a MR(H) hysteresis loop with respective AP/P state at $H = 0$. As an example, let us describe a MR(H) loop measured on 70 nm pMTJ pillar at $T = 340$ K (the most inner loop in Fig. 3). The MR(H) loop contains both AP and P branches both having a parabolic shape before the switching occurs. The AP (resp. P) branch has a maximum (resp. minimum) at $H = 0$ with $R = 5.8$ kΩ (resp. 2.81 kΩ). The switching fields between the branches (which are seen as vertical lines) are -1.7 kOe and $+1.5$ kOe for $P \rightarrow AP$ and $AP \rightarrow P$ branch transitions, respectively. The resistance range corresponding to a discontinuous change in magnetoresistance (the switching) is cut out from the graph in order to focus the reader attention on the reversible parts of MR(H) dependence situated in-between the switching fields and which is only discussed

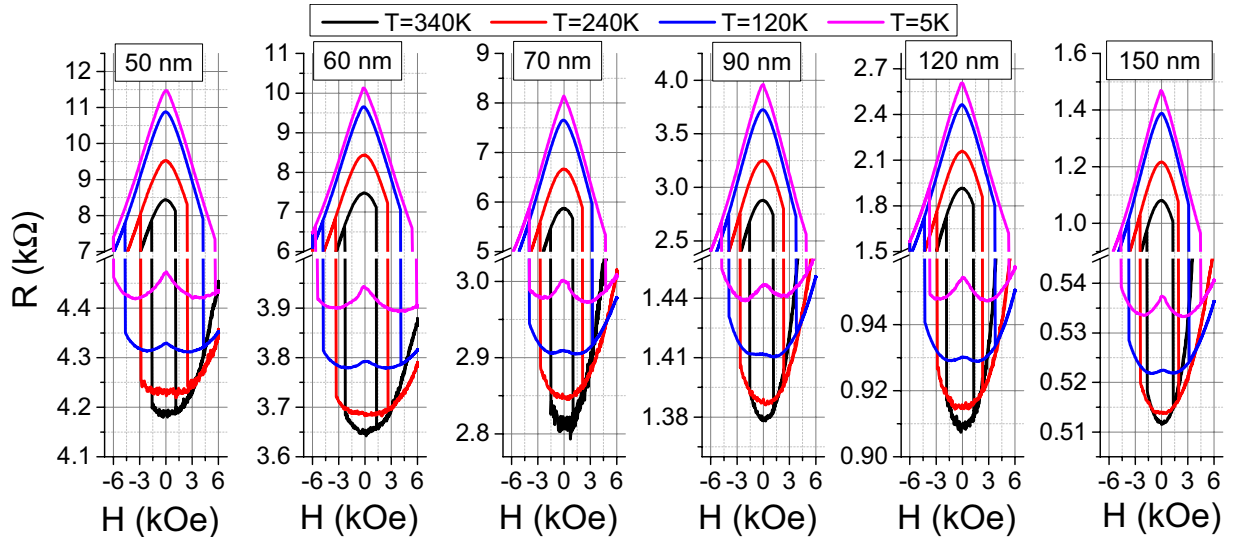


Figure 4. The same as in Fig. 3 for the selected devices of various diameters; only several temperatures are shown.

in the following of the text. Thus, the graph has a brake hiding a range between 3 and 5 kΩ and it has a different vertical scale before and after the brake due to noticeable difference in MR(H) curvature for P and AP branches. The same is applied below in Fig. 4.

Figure 3 shows MR(H) loops behavior as a function of temperature ranging between 5 K and 340 K for a 70 nm diameter pMTJ pillar. For $T > 140$ –120 K, it qualitatively reproduces the situation described in Section 3, i.e. both AP and P branches have a characteristic parabolic shape. In the AP state, the curvature is more pronounced; the resistance variation for the AP branch is one order of magnitude larger than for P branch, which can be ascribed to the finite PMA of the reference layer and correlatively to a rotation of its magnetization. The fitting according to Eq. (2), however, is not ideal even at high temperatures and it is getting worse at decreasing temperature. For $T < 120$ K, MR(H) loops starts showing qualitatively different features and it becomes impossible to reproduce the shape of AP and P branches using Eq. (2). Indeed, at $T = 5$ K, the AP branch exhibits a triangular shape while the P branch shows a local maximum of resistance at $H = 0$ and two respective minima located at ± 2.5 –2.7 kOe. The same behavior is observed for all device diameters, as shown in Fig. 4.

To reproduce experimentally the obtained results in a wide range of temperatures, the model giving Eq. (2) needs to be improved by introducing a second-order uniaxial anisotropy term both in the free and reference layers. The total magnetic energy density (normalized by magnetization saturation parameter M_S) in each layer can be then written as follows:

$$\frac{E_{tot}}{M_S} = -\frac{K_1}{M_S} \cos^2 \theta - \frac{K_2}{M_S} \cos^4 \theta - H \sin \theta, \quad (3)$$

Each layer is assumed to behave as a macrospin. Considering that the uniaxial anisotropy has an interfacial origin, the effective anisotropy constants can be written as $K_1 = \left(\frac{K_{s1}}{t} - 2\pi M_S^2 \right)$, $K_2 = \frac{K_{s2}}{t}$, where K_{s1} , K_{s2} are the interfacial perpendicular anisotropy constants, t is the thickness of a layer.

Unfortunately, no analytical expression of the R(H) variation can be derived in this case. However, the fitting can be carried out numerically. In this case, we folded up the AP and P branches around the $H = 0$ horizontal axis in order to have a more accurate fitting and to cancel, at least partially, the asymmetry of the left and right wings of the MR(H) dependences appearing due to tilted orientation of the external magnetic field. We also increased the relative magnitude of the P branch to give equal weight in the fitting procedure of the P and AP branches. Both AP and P branches are actually fitted simultaneously, so that each fitting result gives $\frac{K_1}{M_S}$, $\frac{K_2}{M_S}$ values in both free and reference layers. Typical results of the fit are shown in Fig. 5. The higher magnitudes set of K_1 , K_2 corresponds obviously to the reference layer. We will use “F” and “R” sub-indexes in the constants to specify to which layer these constants are associated. Accuracy of the fitting is very high in the temperature range between 340 and 160 K. At lower temperatures, the fitting is less accurate but still good enough to reproduce both the triangular shape of the AP branch and characteristic double-well shape of the P branch. It is also important to illustrate how the fitting with K_{2F} , $K_{2R} = 0$ (model Eq. (2)) looks like for the same experimental data (see Fig. 5, dotted lines). At $T = 300$ K, the fitting according to Eq. (2) becomes acceptable. However even in this case, a deviation from the experimental curves is clearly observed: the obtained R(H) curvature is not as accurately reproduced as in the case where the fitting includes the second order anisotropy term. At low temperatures, the fitting without including the second order anisotropy terms does not work at all because of the impossibility to reproduce the double-well shape of the P branch.

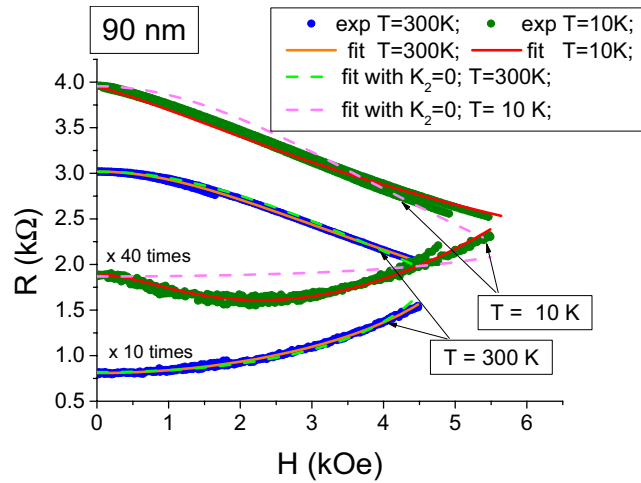


Figure 5. Fitting of the MR(H) loop at T = 300 K and T = 10 K. Extracted values for model (3): T = 300 K: $\frac{K_{1F}}{M_S} = 2815$ Oe, $\frac{K_{2F}}{M_S} = -294$ Oe; $\frac{K_{1R}}{M_S} = 11084$ Oe, $\frac{K_{2R}}{M_S} = -2935$ Oe; T = 10 K: $\frac{K_{1F}}{M_S} = 5184$ Oe, $\frac{K_{2F}}{M_S} = -1034$ Oe; $\frac{K_{1R}}{M_S} = 37285$ Oe, $\frac{K_{2R}}{M_S} = -19178$ Oe. The sub-indexes F and R specify free and reference layers' constants respectively.

The origin of the appearance of a triangular shape in the AP R(H) branch as well as double well shape in the P R(H) branch at low-temperature can be understood from the values of the extracted K_1 and K_2 parameters for T = 10 K. In both layers K_{2F} , K_{2R} is negative. In the free layer $-K_{2F}/K_{1F} = 0.2$ at 10 K while $-K_{2F}/K_{1F} = 0.104$ at T = 300 K. The increase of $-K_{2F}/K_{1F}$ ratio at low temperatures results mainly in a decrease of the free layer switching field and deformation of hard-axis M(H) and R(H) dependences. In the case of the reference layer at T = 300 K, $-K_{2R}/K_{1R} = 0.265$ while at T = 10 K $-K_{2R}/K_{1R} = 0.514$, i.e. higher than 0.5 which yields the onset of an “easy-cone” ground state of the reference layer magnetization instead of “easy-axis” at high temperatures. In the easy-cone regime, the reference layer magnetization is tilted out from the symmetry axis by the angle θ_c , $\cos^2 \theta_c = -K_1/2K_2$; From the fitting parameters, the easy-cone angle at 10 K is $\theta_{cR} \sim 10^\circ$. In this regime, an infinitely small reversal of the in-plane applied field yields a 180° rotation of the in-plane component of the reference layer magnetization around its easy cone thereby skipping the parabolic part of the R(H) curve, thus resulting in the observed triangular shape of the R(H) response at low temperature.

The double-well shape of the P branch can also be explained by the easy-cone regime in the reference layer. The free layer is in the “easy-axis” state (i.e. at $H = 0, \theta = 0$) since $-K_{2F}/K_{1F} < 0.5$. Its anisotropy is about 4 times lower than that of the reference layer. For this reason, the in-plane magnetic field tilts the free layer magnetization away from the normal to the plane direction faster than the reference layer magnetization. Starting from zero field, for $0 < H < 2.5$ kOe, the in-plane magnetic field first yields a decrease in the relative angle between the magnetic moments in the two electrodes. Indeed, because the reference layer is initially oriented in a canted direction, $\theta_{cR} \sim 10^\circ$, the field-induced rotation of the free layer magnetization towards the applied field brings it closer to the reference layer magnetization. The minimum of resistance at $H \sim 2.5$ kOe therefore corresponds to the parallel orientation of both magnetic moments. Further increasing the magnetic field gives rise to an increase of the relative angle between the two moments so that correlatively the resistance starts increasing again. It is expected that at larger fields, the resistance would decrease again since the system would evolve towards the parallel configuration if full saturation could be reached at very large fields. But full saturation of the reference layer magnetization would require overcoming both the anisotropy energy and the antiferromagnetic RKKY coupling across the ruthenium layer. Field of the order of 2 T would be needed to observe this behavior which is out of our range of measurements.

A summarized view of the temperature dependences of K_{1F}/M_S , $-K_{2F}/K_{1F}$, K_{1R}/M_S , $-K_{2R}/K_{1R}$ extracted from the fitting for all measured devices is shown in Fig. 6. All samples exhibit the same trends and similar magnitudes of the anisotropy constants extracted from the fitting. Scattering of the extracted values gives an idea of the dispersion of the fitting parameters. The temperature dependences of average values of these parameters over all measured devices are also shown. For the free layer, in average, K_{1F}/M_S increases almost linearly as the temperature decreases in the range 120–300 K. The corresponding $-K_{2F}/K_{1F}$ ratio also increases in this range of temperature, not exceeding 20% at low T and therefore never reaching the easy cone regime. Concerning the reference layer, the situation is generally similar, but $-K_{2R}/K_{1R}$ ratio is much larger at all temperatures. Below 120–160 K, $-K_{2R}/K_{1R}$ ratio is above 0.5 so that the reference layer magnetization enters the “easy-cone” regime as pointed out above. Among the measured devices, the fitting for the 60 nm pillar demonstrates a rather strange behavior of $-K_{2F}/K_{1F}$ ratio for T < 100 K. It is hard to give a definite explanation for this observation without additional statistical measurements on other devices of the same diameter. One of the possible scenarios is the development of

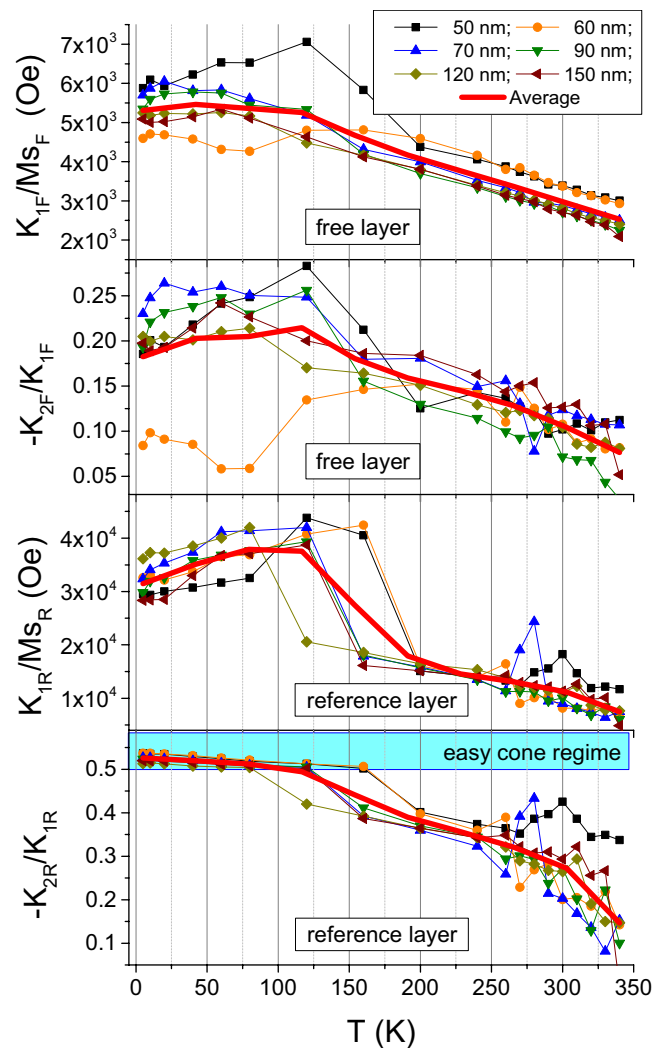


Figure 6. Temperature dependences of the anisotropy fields and $-K_2/K_1$ ratios extracted from the fitting.

a micromagnetic distortion near the pillar edges due to magnetic imperfections introduced by nanofabrication process.

We also recalculated the easy-cone angle θ_{cR} of the reference layer magnetization versus temperature. As shown in Fig. 7, the easy-cone angle increases almost linearly as temperature decreases below ~ 180 K. Furthermore, θ_{cR} is observed to increase with decreasing sample diameter. As will be shown further in section 7, the K_2 contribution is interpreted in terms of spatial fluctuations of the uniaxial K_1 first order term. In this case, for smaller diameter, edge defects may increase K_2 due to increased spatial fluctuations of K_1 . This could explain the larger $-K_2/K_1$ ratio and correlatively the large easy cone angle observed at small pillar diameters.

Easy-Cone Regime in Sheet FeCoB/MgO Films

Generally, one cannot rule out *a priori* that certain types of micromagnetic distortions in the ferromagnetic electrodes could be responsible for the observed hard-axis MR(H) curve deformations in the studied pillars at low temperatures. To exclude this possibility, experiments were conducted at sheet film level in order to check whether the second order anisotropy is also evidenced in this case. In thin films, the demagnetizing (magneto-static) energy and first order perpendicular interfacial anisotropy K_{s1} have the same symmetry. They can be combined in one effective anisotropy density constant $K_1 = \left(\frac{K_{s1}}{t} - 2\pi M_S^2 \right)$. Consequently, an easy way to tune the K_2/K_1 ratio simply consists in changing the thickness t of the FM film. For any $K_2 \neq 0$ amplitude, a range of FM thickness around the anisotropy reorientation transition from out-of-plane to in-plane direction should always exist, wherein the easy-cone regime should be observable.

To check this, several samples were grown, consisting of an $\text{Fe}_{72}\text{Co}_8\text{B}_{20}$ layer in contact with MgO with nominal thickness of FM material $t = 17.4 \text{ \AA}$, 16.9 \AA and 15.8 \AA . Room-temperature magnetization measurements with magnetic field applied parallel to the films plane clearly exhibit three different M(H) loop shapes as shown in Fig. 8. The thickest and thinnest samples demonstrate M(H) loops respectively typical of XY-easy-plane and Z-easy-axis anisotropies (the field is applied in the XY-plane, which is the sample plane). The sample with

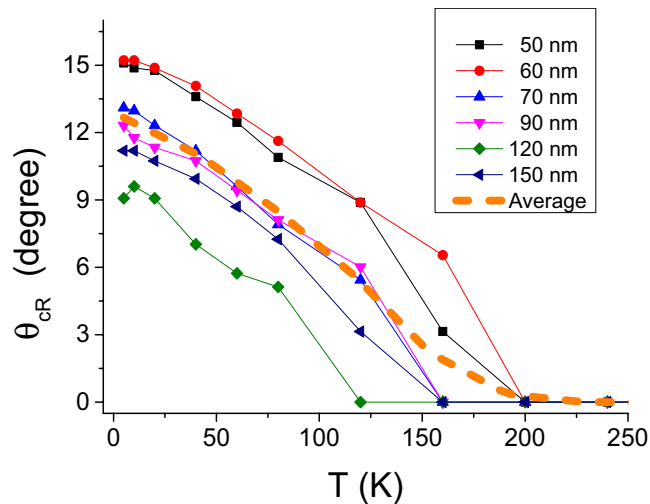


Figure 7. Angle between the symmetry axis and the easy cone direction as a function of the temperature.

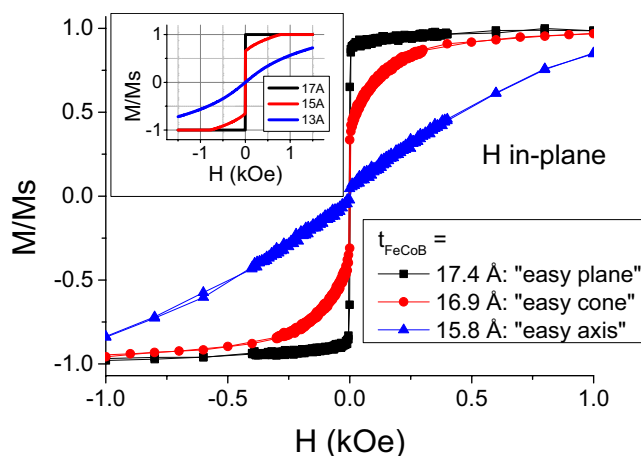


Figure 8. $M(H)$ curves for three samples around the anisotropy reorientation transition. Z-axis is out-of-plane. The field is applied in-plane. The inset shows simulated $M(H)$ dependences for three different thicknesses using the model (3) with $M_S = 1000 \text{ emu/cm}^3$, $K_{1S} = 1 \text{ erg/cm}^2$, $K_{2S} = -0.05 K_{1S}$.

intermediate FM layer thickness shows features of a two-step magnetization process. Firstly, an abrupt switching of magnetization, as in the thickest sample, followed by a slower non-linear $M(H)$ magnetization increase. Such features are exactly expected in presence of easy-cone anisotropy. As discussed in the previous section, when the magnetic field is applied perpendicularly to the easy cone symmetry axis, it is initially very easy to rotate the in-plane component of magnetization around the easy-cone. This corresponds to the low-field part of the $M(H)$ curve with an abrupt variation of the magnetization. Following this rapid rotation, at larger fields, the magnetization has to depart from the easy cone to gradually align with the in-plane applied field. This yields a more gradual increase of magnetization since the easy cone anisotropy has to be gradually overcome by the Zeeman energy. Corresponding macrospin simulations using model of Eq. (3) are shown in the inset of Fig. 8 and reproduce the qualitative modifications of the in-plane $M(H)$ loop shapes with the film thickness variation. Knowing that the “real” films are in multidomain state, we did not try to match the macrospin simulations and experiments exactly.

Discussion

Regarding the origin of the second order anisotropy term which gives rise to the easy cone regime, at least two explanations *a priori* can be provided and discussed. The first one is based on the possible existence of a bulk magnetocrystalline cubic anisotropy in the centered cubic Fe rich alloy constituting the magnetic electrodes of the MTJ. Our samples are polycrystalline so that the in-plane mosaicity of the FeCoB grains can average out the in-plane anisotropy. In contrast, due to the (100) texture of the film, the out-of-plane component of this anisotropy can be conserved with easy axis of anisotropy along the $\langle 001 \rangle$, $\langle 010 \rangle$, and $\langle 100 \rangle$ directions²⁵. The four-fold bulk cubic anisotropy combined with the two-fold uniaxial anisotropies for the out-of-plane direction can yield the observed behavior for the $M(H)$ dependence²⁶.

We have employed X-band magnetic resonance technique (9.45 GHz) in order to investigate this possible source of 2nd order anisotropy in the sample with $t = 16.9 \text{ \AA}$. Room temperature ferromagnetic resonance (FMR)

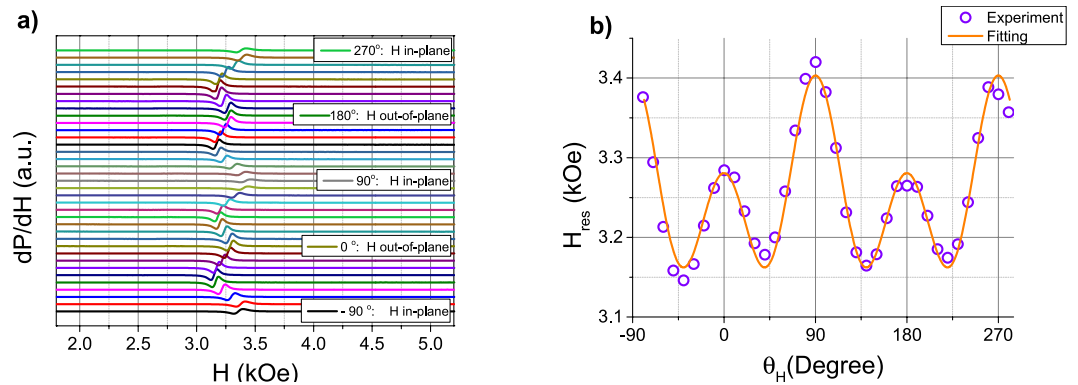


Figure 9. Angular dependent out-of-plane FMR measurements on the sample with $t = 16.9$ Å. (a) 10-degree step FMR spectra with magnetic field angle counted from the film normal. (b) extracted angular dependence of FMR resonance field and respective fitting using Smit-Beljers formalism.

spectra were measured for different angles of magnetic field with respect to the sample normal. The results are shown in Fig. 9. FMR signal, as seen by comparing Figs 8 and 9 is observed in a magnetic field range wherein the sample is completely saturated. The angular dependence is composed of four-fold and two-fold angular contributions of comparable amplitudes. These two contributions exhibit energy maxima when the field is oriented in-plane. Conversely, the four-fold anisotropy also reaches maxima when the field is out-of-plane while the two-fold anisotropy reaches minima for this field orientation. Without entering into the details of magnetic resonance, one can therefore definitely state that the hard axis directions of the four-fold anisotropy correspond to the normal to the film or to the in-plane direction. For the out-of-plane angular dependence of FMR, the expected behavior is qualitatively similar for the case of uniaxial + cubic anisotropy and for the case of uniaxial with the second order uniaxial term. The extracted constants in the case of uniaxial + cubic anisotropies are $K_1 = 6.2 \cdot 10^6$ erg/cm³ and $K_{1c} = -7.7 \cdot 10^4$ erg/cm³ (assuming H rotating in (010) plane). In comparison with the bulk values of bcc iron ($\sim 5 \cdot 10^5$ erg/cm³), the obtained cubic anisotropy constant K_{1c} is six times lower and has the opposite sign. It is well known that by adding cobalt into iron, the anisotropy constant K_{1c} is expected to gradually change from positive to negative with $K_{1c} \sim 0$ for Fe₄₅Co₅₅ composition²⁵. In our case, the layer is iron-rich so that the lower value of K_{1c} could be explained by the Co content of the alloy in this 1.7 nm thick layer. However, the opposite sign of K_{1c} is not expected. Moreover, for positive K_{1c} , the easy direction of uniaxial and cubic anisotropies would coincide along <001> direction not allowing therefore the formation of the canted state in contradiction with its experimental observation. We can therefore conclude that bulk cubic anisotropy of iron rich alloy does not play a significant role in these samples and other explanations have to be found for the second order anisotropy term.

Several experimental studies reported anisotropy reorientation phenomena about which the role of a second-order uniaxial anisotropy term could be evidenced. Easy-cone regime was observed experimentally near the magnetic reorientation transition in Co films grown on Pt(111) and Pd(111) substrates¹⁶ as well as on Co/Pt multilayers¹⁴. Recently, J. Shaw *et al.* have reported FMR measurements on Ta/Co₆₀Fe₂₀B₂₀/MgO films¹⁹. The authors obtained an angular dependence of FMR with maxima of FMR field corresponding to in-plane and out-of-plane magnetic field orientation with four-fold and two-fold angular dependencies, as in our case. The authors, however, used a cobalt-rich crystallized alloy, which in bulk has negative constants of cubic anisotropy²⁵. They put forward a possible $\sim \sin^4\theta$ contribution without too much explanation on its possible origin.

Besides, several studies have pointed out the possible influence of strain on second order anisotropy in thin magnetic films^{27,28}. For instance in ref. 28, the authors observed an anomaly in the tunneling conductance in Ta/CoFeB/MgO based MTJ at low temperatures ($T = 160$ K), that they interpreted as a structural-magnetic phase transition of a magnetic oxide formed at the interface between MgO and CoFeB. Magnetic phase transition alters both spintronic and magnetic properties of the MTJ stack. While the proposed interpretation definitely needs a more detailed study, we should accept the fact that in our experiment we also observe a fast increase of K_1 for the reference layer in the temperature range 120–160 K. It can be speculated that this observation might be associated with a low-temperature structural transition in one of the stack layers, not necessarily a magnetic one. Along the same line, mismatch of thermal expansion coefficients of the different materials in the stack and substrate can also play an important role. Considering the large magnetostriction of Fe rich FeCo alloys, stresses in the pillar can change the magnetic anisotropy in the magnetic layers through magnetoelastic coupling. Even the crystallization of MgO during the post-deposition annealing can produce some residual stresses in the neighboring ferromagnetic electrodes. Therefore one cannot rule out that magnetoelastic effects play a role in the second order anisotropy term that we observe in our samples. Further structural characterization and stress analysis would be required to clarify that.

Another possible origin of the second-order uniaxial term was proposed theoretically by B. Dieny and A. Vedyayev²⁰. They have shown analytically that spatial fluctuations in the magnitude of first order surface anisotropy can give rise to a second order anisotropy contribution provided the characteristic wavelength of these fluctuations is much smaller than the exchange length. The sought second-order contribution has a negative sign

with respect to the main first-order term, thus allowing the onset of easy cone anisotropy. Topology of the interface in their model determines the relative strength of the second-order contribution. In case of Fe/MgO systems, spatial fluctuations of the effective perpendicular anisotropy can be responsible for the second order anisotropy term. These fluctuations can be due to local variations in the ferromagnetic layer thickness associated with film roughness. Due to the competition between interfacial anisotropy and bulk demagnetizing energy, around the anisotropy reorientation transition, a monolayer variation in the thickness of the FeCoB layer due to interfacial roughness is sufficient to yield spatial variations of effective anisotropy from in-plane to out-of-plane. Following the model of ref. 20, using an average film thickness of 15 Å and variations in FM layer thickness ± 2 Å, one can expect spatial modulation of the surface anisotropy parameter of the order ~ 0.2 erg/cm². Considering an exchange constant $\sim 1.5 \cdot 10^{-6}$ erg/cm, $K_{1S} \sim 1$ erg/cm², period of spatial fluctuations ~ 15 nm, one should expect $K_{2S} \sim -0.0024$ erg/cm² which magnitude is much lower than that estimated from the aforementioned fits (Figs 5 and 6). Alternatively, one may think about the possible presence of nanometric “dead” spots where contribution to the net interfacial anisotropy could be locally strongly reduced. In Ta/CoFeB/MgO, a possible explanation for the existence of dead spots could be the preferential diffusion of Ta along the grain boundaries of the CoFe(B) layer to the MgO barrier upon post-deposition annealing. The presence of Ta next to the barrier can locally alter the interfacial perpendicular anisotropy yielding strong local variations of interfacial anisotropy between the inner part of the grains and the grain boundaries. Assuming a grain size ~ 16 nm and a spatial modulation of the interfacial anisotropy ~ 1 erg/cm² yields $K_{2S} \sim 0.07 K_{1S}$ which is the right order of magnitude.

The observed temperature dependence can be explained by the model of ref. 20. According to it, K_2 scales as square of K_1 . This is generally what is observed in the temperature range 160–340 K: K_1 decreases with temperature but $-K_2/K_1$ also decreases, meaning that K_2 drops with temperature faster than K_1 . However, in the temperature range between 5 and 120 K, the behaviors of the reference and free layer are different. Indeed, the free layer keeps following the above described tendency while the reference layer shows abrupt changes in K_1 and further decrease of its magnitude versus decreasing temperature. This may indicate that for the reference layer which has a complex structure (SyAF), the single macrospin description may not be sufficient. Different temperature dependences of perpendicular anisotropies arising from MgO/FM interface and from the synthetic Co/Pt multilayer as well as temperature-dependent coupling field through the NM layer may complicate the overall picture.

From practical point of view, the easy cone anisotropy can be used to significantly improve the writing performances of pMTJ-based STT-MRAM elements^{29,30}. In a standard pMTJ system, the magnetic moments of both free and reference layers are aligned parallel or antiparallel in standby regime. Upon writing, when the write current starts flowing through the MTJ, the initial STT-torque is zero and only thermal fluctuations or micromagnetic distortions provide the non-collinearity required to trigger the reversal of the storage layer magnetization. Both effects are generally undesirable in STT-MRAM technology. Indeed, thermal fluctuations are stochastic by nature and therefore the write pulse duration and intensity must be overdesigned to reach the specified write error rate. As for micromagnetic distortions, the latter induce non-uniform switching process which can result in the need for higher switching current and variability in the switching process. An easy cone regime in the free layer and the easy axis configuration in the reference one would be the optimal configuration for a STT-MRAM memory element. Unique features of easy cone regime is that it allows for a canted state and at the same time conserves the axial symmetry so essential for effective transfer of the STT torque into the angular motion.

We point out that because the magnetostatic term reduces the effective K_1 but keeps K_2 unchanged, the ratio K_2/K_1 is at least twice larger than the surface constants ratio K_{2S}/K_{1S} . Thus, keeping $K_2/K_1 \sim 10\%$ as it is in the free layer, the expected K_{2S}/K_{1S} ratio should not exceed 5% at room temperature which is quite easy to overlook both in experiments and theories.

Conclusion

While easy-axis magnetoresistance loops allow for determination of switching current and coupling fields, hard axis magnetoresistance loops provide additional information about the magnetic anisotropy in pMTJ pillars. Reversible parts of the hard-axis magnetoresistance loops starting from parallel or antiparallel configuration can be simultaneously fitted providing quantitative estimation of the effective anisotropy fields both in the free and reference layers.

In this work magnetoresistance loops of pMTJ pillars with radius 50–150 nm were measured in a wide range of temperatures. The anisotropy fields in both free and reference layers were derived in the temperature range between 340 and 5 K. At temperatures below 160–120 K, the shape of the hard-axis magnetoresistance loops changes qualitatively from parabolic to triangular which cannot be described by a model taking into account only first order magnetic anisotropy. By adding a higher order anisotropy term, the magnetoresistance loops could be fitted to the model over the whole temperature range and for all measured devices. The extracted anisotropy constants have shown that the second order term is noticeable and it has a negative sign with respect to the first order anisotropy term in both layers. At room temperature, the magnitude of the second order term is about 10% of the first order one in the free layer and about 20% in the reference layer. With decreasing temperature, the second order term contribution increases faster than the first order one and exceeds 50% of the first order term in the reference layer below 120 K. This results in a change of the reference layer net anisotropy from easy-axis along the normal to the plane to easy cone. In this state, hard-axis magnetoresistance loops acquire a triangular shape for the antiparallel branch and a double well shape with a maximum at $H = 0$ for parallel branch. The free layer remains with a net easy axis anisotropy at all temperatures. Extracted temperature dependences of the anisotropy in both layers are quantitatively and qualitatively similar for all measured devices whatever their diameter. Therefore, the anisotropy transition from easy-axis to easy cone regime seems to be diameter-independent.

We have evidenced the existence of the higher-order term in simple FeCoB/MgO sheet films and it is experimentally accessible for the thicknesses corresponding to the magnetization reorientation transition. The Dieny-Vedyayev model proposed in ref. 20 explains the second order magnetic uniaxial anisotropy contribution,

$-K_{2s} \cos^4 \theta$ with $K_{2s} < 0$, as a result of spatial fluctuations of the first order anisotropy parameter, $-K_{1s} \cos^2 \theta$ with $K_{1s} > 0$. The preferred diffusion of Ta through the CoFe(B) layers towards the MgO interface upon post-deposition annealing and CoFeB crystallisation was proposed as a possible mechanism at the origin of these spatial fluctuations of the CoFeB/MgO interfacial anisotropy.

The canted (easy cone) state of the free layer could be advantageously used to improve STT writing performance in pMTJ pillars in STT-MRAM applications. In this state, the system conserves its axial symmetry allowing STT to work efficiently over the whole precession orbit. At the same time, it provides an initial noncollinearity between free layer and polarizer which considerably reduces the threshold switching current and stochasticity in switching time at finite temperatures. Reduction of the thermal stability is a negative effect which is also expected in the easy cone state. However, simple macrospin simulations show that the threshold current is reduced faster than the stability factor meaning that the overall performances of pMTJ pillar with the free layer in the easy cone regime should be improved. Thus further research aiming at engineering high $-K_2/K_1$ ratio while keeping K_1 large enough to achieve sufficient thermal stability of the storage layer is highly desirable.

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Author Contributions

R.S. and H.T.N. grew the samples; A.A.T. carried out the magnetotransport and magnetoresonance experiments; H.T.N. made magnetostatic measurements; A.A.T. and B.D. wrote the manuscript; M.C. proofread the manuscript; all the authors participated in the discussions of the experiment, model and manuscript.

Additional Information

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