Splitting of a moving transverse domain wall in a magnetic nanostripe in a transverse field

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The field-driven domain-wall dynamics in magnetic nanostripes under the influence of transverse fields was examined experimentally and by numerical simulations. For a small range of transverse fields in a regime below the Walker field an unexpected instantaneous change in the domain-wall velocity was found. Micromagnetic simulations have shown that this peculiar behavior is caused by a transformation of the moving asymmetric transverse wall: the wall is stretched along the nanostripe due to different velocities of the upper and lower half antivortices constituting the domain wall.

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Domain walls (DWs) in nanostripes are the key elements for different spintronic applications. The control of the domain-wall position in magnetic nanostripes by means of magnetic fields and electric currents was addressed recently in many contributions. These were motivated by possible DW applications for data storage logic devices and an existing application based on moving DWs in nanostripes. The control of DWs, moved through a system of magnetic nanostripes by rotating magnetic fields. Understanding of domain-wall dynamics under these conditions is crucial for the application of these devices. Here, besides the longitudinal field component \( H_{\text{long}} \) (driving the DW through a nanostripe), an additional transverse field \( H_{\text{tr}} \) directed in-plane perpendicular to the nanostripe long axis is present during the DW motion. The effect of such a field on the DW dynamics was recently studied in simulations and experiments. It was shown that a transverse field strongly influences the DW dynamics both below and above the critical Walker field \( H_{\text{tr}} \).

In this Rapid Communication we report a dynamic phenomenon appearing at relatively large transverse fields and manifesting itself in an abrupt change in the DW velocity outside the Walker breakdown regime. Micromagnetic simulation has revealed a peculiar DW motion type responsible for the observed behavior.

We have examined the DW motion in \( \text{Ni}_{81}\text{Fe}_{19} \) nanostripes of width \( 300 \) nm and thickness \( 10 \) nm. These nanostripes were the free layer of giant magnetoresistance (GMR) stacks with the composition \( \text{Ta}(5 \text{ nm})/\text{NiFe}(10 \text{ nm})/\text{CoFe}(0.8 \text{ nm})/\text{Cu}(2 \text{ nm})/\text{CoFe}(3 \text{ nm})/\text{Ru}(0.8 \text{ nm})/\text{CoFe}(3 \text{ nm})/\text{IrMn}(7 \text{ nm})/\text{Ta}(5 \text{ nm}) \). The stack was deposited on a sapphire substrate using physical vapor dc magnetron sputtering (base pressure \( <2 \times 10^{-8} \) mbar) and patterned by means of e-beam lithography and Ar ion etching. Electrical Al contacts in coplanar waveguide geometry were deposited using a lift-off process.

A sample design is shown in Fig. 1(a). The reference magnetization of the 12-\( \mu \text{m}-\)long nanostripe was oriented parallel to its long axis (chosen as \( x \) axis). Using the experimental setup sketched in Fig. 1(a), we measured the resistance time dependence in single shot measurements. The GMR value is a direct measure of the sense layer magnetization component \( m_{\text{long}} \) parallel to the nanostripe long axis so that we could track the DW during its movement. For the present Rapid Communication we have used the following procedure. After saturating the NiFe layer in the stripe direction by using a longitudinal field \( H_{\text{long}} = -40 \text{ kA/m} \), we applied first a constant transverse field \( H_{\text{tr}} \) and then a linearly increasing longitudinal field \( \frac{dH_{\text{long}}}{dt} = 160 \text{ (kA/m)/s} \). Note that during DW motion (\( \approx 1 \) \( \mu \text{s} \)) the longitudinal field is practically constant. Both the critical longitudinal field \( H_{\text{tr}}^{c} \) required to nucleate a DW and the DW velocity \( v_{\text{DW}} \) (measured in this field \( H_{\text{tr}}^{c} \)) exhibited a strong dependence on the transverse field \( H_{\text{long}} \) as shown in Fig. 1(b).

Figure 2(a) shows single shot measurements at transverse fields marked as A and B in Fig. 1(b). For large \( H_{\text{long}} \) (case A, \( H_{\text{tr}}^{c} = 9.8 \text{ kA/m} \) and \( H_{\text{tr}} = 6 \text{ kA/m} \)) we can clearly resolve the stop-and-go motion typical for the Walker breakdown process. When the transverse field is increased to \( H_{\text{tr}} = 8.6 \text{ kA/m} \), then the longitudinal field necessary for

![FIG. 1.](Color online) (a) Measurement setup and a micrograph of the GMR device. The inset shows the GMR nanostripe. (b) The DW mean velocity (squares) and longitudinal field (circles) \( H_{\text{tr}}^{c} \) as a function of \( H_{\text{tr}} \) (averaged over 25 repetitive measurements at each \( H_{\text{tr}} \)). Error bars for \( H_{\text{tr}}^{c} \) are smaller than the symbol size.)
To understand the reason for the decrease in the DW velocity and increase in its dispersion, we show a typical single shot measurement of the DW movement for \( H_{\text{tr}} = 11.3 \) kA/m. In some cases the DW moves through the nanostripe with constant (high) velocity. These strong differences in the DW behavior for different measurements at the same \( H_{\text{tr}} \) causes the large dispersion of the DW velocity.

In order to clarify this peculiar feature of the DW motion, we have performed micromagnetic simulations for the system similar to that used experimentally using the MICROMAGUS package.\(^{21}\) We have simulated the DW dynamics in a \( 10 \times 200 \) (and 300, respectively) \( \times 6000 \) nm\(^3\) nanostripe using discretization cells of \( 10 \times 5 \times 5 \) nm\(^3\). The material parameters are chosen as for Permalloy \((M_s = 750 \) kA/m, \( A = 1.3 \times 10^{-11} \) J/m). In the starting configuration of our simulations the DW is pinned to an artificial notch in the nanostripe, positioned 1 \( \mu \)m apart from the left stripe edge; for this reason the magnetization at \( t=0 \) is \( m_{\text{out}} = -0.6 \) (and not \(-1.0\)).

We note that simulations were performed for the perfect nanostripe shape, i.e., not taking into account the nanostripe edge roughness, present in real systems. An edge roughness decreases the effective nanostripe width and influences the DW dynamics significantly, causing, e.g., a shift of the Walker field.\(^{22}\) Therefore, the quantitative agreement between the DW dynamics found experimentally and in simulations cannot be expected even for nanostripes having the same dimensions. As mentioned above, we performed the simulations for two different nanostripe widths 200 and 300 nm. In both cases we have found this type of DW behavior (of course, for different field values). However, in the 300-nm-wide nanostripes the DW dynamics demonstrates a much wider variety than for the 200 nm stripes (which will be published elsewhere). Thus, in order to simplify the explanation, we concentrate in the following on the simulations of the 200-nm-wide nanostripes.

In previous micromagnetic studies where small transverse fields were used,\(^{19,20}\) the type of the DW dynamics found here has not been reported so that the transverse field strength seems to play a key role. Hence we have studied the DW motion in dependence on \( H_{\text{tr}} \). Detailed results of this study will be reported elsewhere. Here we present a typical example for a transverse field \( H_{\text{tr}} = 23.5 \) kA/m (and \( H_{\text{long}} = 4.8 \) kA/m). The time dependence \( m_{\text{long}}(t) \) for the parameters specified above is shown in Fig. 2(c). This simulated dependence is remarkably similar to that found experimentally [shown in Fig. 2(b)]. The velocity also changes instantaneously, dropping from the large initial value of \( v_{1,\text{sim}} = 830 \) m/s to \( v_{2,\text{sim}} = 240 \) m/s.

Visualization of the simulated magnetization configurations (see Fig. 3) immediately reveals the reason for this highly nontrivial magnetization time dependence and the abrupt change in the effective DW velocity. Namely, it can be clearly recognized that after the DW starts to move, it quickly splits into two parts, and the upper part moves much faster than the lower one. As long as both parts remain within the nanostripe, the total longitudinal magnetization component changes very fast, resulting in the large apparent DW velocity, calculated as \( \dot{v}_{\text{DW}} \approx dm_{\text{long}}/dt \). When the upper DW part, moving much faster, reaches the right nanostripe end, the magnetization component \( m_{\text{long}} \) continues to change but due to the motion of the lower DW part only. Hence the
change rate $dm_{\text{long}}/dt$ becomes much smaller, resulting in the abrupt decrease in the calculated $v_{\text{DW}}$ [Figs. 2(b) and 2(c)].

In order to understand this behavior, we note first that for magnetic and geometric parameters used in simulations, the DW formed at $t=0$ (when only $H_{tr}$ is applied) is the so-called asymmetric transverse wall. This wall is formed by two half antivortices (HAVs) with opposite winding numbers $n$: the upper HAV with $n=+1/2$, and a lower HAV with $n=-1/2$. For the asymmetric transverse wall these two HAVs are shifted with respect to each other in the longitudinal direction.

Due to the different magnetization rotation senses within the upper and lower HAVs, they have very different stray field energies. This leads to the different widths of the two HAVs (clearly visible already in the equilibrium state). This difference also leads to different mobilities of the two HAVs in the longitudinal applied field so that the upper HAV moves initially faster than the lower one.

The last statement is qualitatively true both in the presence and absence of a transverse field. However, when $H_{tr}$ is absent, the initially fast motion of the upper HAV slows down when the DW continues to move, and then the motion direction of the upper HAV is even reversed (see Fig. 4). This process, combined with the formation of the full antivortex and its periodic motion across the nanostripe, leads to the well-known periodic oscillations of the magnetization components, observed in simulations of the transverse wall dynamics in a nanostripe (see, e.g., Refs. 23 and 25).

In contrast, our simulations show, that in a sufficiently large transverse field the upper DW part always moves faster than the lower one. We note also that in presence of $H_{tr}$ the full antivortex (AV) is also formed at the nanostripe edge where the HAV with the lower mobility is located (as for $H_{tr}=0$). However, this AV never reaches the opposite nanostripe edge—it is pushed back to its initial position, where it annihilates, emitting a burst of spin wave (see last picture row in Fig. 3).

The analytical theory, which would allow us to understand why the increase in the transverse field value qualitatively changes the DW dynamics is, up to our knowledge, not available. In particular, we cannot calculate the mobility of the two HAVs mentioned above, especially for the case, when an additionally transverse field is applied. For this reason, we can present only qualitative arguments explaining why the application of $H_{tr}$ is crucially important.

From the “energetic” point of view the explanation for the DW splitting is relatively straightforward. When the upper HAV at the initial stage of the DW motion moves forward faster than the lower HAV, the wall is tilted and stretched along the nanostripe direction as it can be seen at the first three image rows in Fig. 4. The magnetization in the tilted part of the domain wall is directed in plane and nearly perpendicular to the $x$ axis. When the transverse field $H_{tr}$ is applied, its direction coincides with the magnetization direction in this tilted DW part, thus leading to a large gain in the Zeeman energy. When the magnitude $H_{tr}$ reaches a critical value, this energy gain compensates the exchange energy increase due the wall elongation. Hence the formation and growth of the wall region parallel to the long strip side becomes energetically favorable, resulting in the DW splitting as shown in Fig. 3.

However, for the small damping typical for real NiFe layers studied in our experiments (in simulations we have used $\alpha=0.01$) the DW motion is essentially a dynamic process governed by the magnetization precession, so that the “energetic” argument presented above is not enough to explain the qualitative change in the DW dynamics when $H_{tr}$ is switched on.

To better understand the transverse field influence on the DW dynamics, we have to recapitulate in more details the behavior of the upper (initially faster) HAV during the DW
motion without a transverse field. The important point is that when this half antivortex moves (initially) to the right, its velocity decreases, then the HAV stops and starts to move in the opposite direction. This process can be clearly seen on the color maps of the in-plane magnetization projection (first column in Fig. 4). The reason for this change in the movement direction is the gradual change in the HAV polarity during its motion. Namely, initially the HAV-core magnetization points along the positive out-of-plane direction (bright spots showing the HAV position at the first two maps in Fig. 4), then the out-of-plane component of the HAV magnetization reduces and finally changes its sign as it can be seen from the dark spots marking the HAV on the last two maps in Fig. 4. The detailed analysis (which will be reported elsewhere) shows that $T_z$ component of the torque responsible for the longitudinal HAV motion is dominated by demagnetizing field torque $T_{\text{dem}}$. Hence, when the $m_\perp$ component of the HAV core changes sign, $T_z$ component changes its sign also, thus causing the backward HAV motion (see the third column in Fig. 4).

The decrease and the sign change in the initially positive $m_\perp$ component is caused, in turn, by the out-of-plane torque projection $T_\perp$; dark spots indicated by arrows on the first two maps in Fig. 4 clearly show that this component is negative at the HAV location. The detailed analysis shows that the (negative) out-of-plane torque projection is a result of the nearly cancellation of the demagnetizing $T_{\text{dem}}$ and exchange torques $T_{\text{exch}}$, whereby the magnitude of $T_{\text{exch}}$ is slightly larger.

When, in addition, a transverse field is applied, it makes a magnetization configuration more homogeneous. The first important point is that this has a larger impact on the exchange torque $T_{\text{exch}}$ (compared to $T_{\text{dem}}$) because $T_{\text{exch}}$ is propotional to the spatial derivatives of the magnetization components. Second, for the same reason, the magnitude $T_{\text{exch}}$ reduces with increasing $H_x$ [because $M(r)$ becomes more homogeneous]. Hence, starting from some value of $H_x$, the exchange torque component $T_{\text{exch}}$ at the HAV location is not more able to overcompensate the positive demagnetizing torque $T_{\text{dem}}$. Thus the $m_\perp$ component of the HAV core remains positive as it can be seen on the maps in Fig. 3, where the HAV positions are marked by the arrows. For this reason the upper HAV velocity is always positive and exceeds the velocity of the lower HAV causing the DW splitting.

In conclusion, we have found a peculiar type of the DW dynamics in a nanostripe under the influence of an in-plane transverse field. Due to this field the upper and lower half antivortices of the transverse wall can behave as independent quasiparticles. The different mobilities of these two half-antivortices results in different velocities of the two corresponding DW parts in the given longitudinal field. This leads to an abrupt change in the effective DW velocity measured from the time dependence of the total longitudinal magnetization component.

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