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# Characterization of Spin-Wave Eigenmodes in Normally Magnetized Nano-Pillars

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**Abstract** The eigen-modes of spin waves in a circular spin-valve nano-pillar that is perpendicularly magnetised along z are studied spectroscopically and reported here. Magnetic Resonance Force Microscopy (MRFM) is used to conduct spectroscopy. Depending on whether an RF current running perpendicularly through the layers or a homogeneous in-plane RF magnetic field excites the nano-pillar, different spectra are observed. These findings are consistent with micromagnetic simulations of the local magnetization's time decay response to excitations with various azimuthal symmetries, (x iy)e-i4. Because of the orthoradial symmetry of the induced RF Oersted field, only four 0 modes are stimulated by the RF magnetic field and only four 1 modes are excited by the RF current, indicating that the azimuthal 4-index is the discriminating parameter for the selection criteria.

#### 1.1 Introduction

A new science focused on harnessing spin dependent effects in the electrical transport characteristics has emerged as a result of technological advancements in the creation of hybrid nanostructures employing magnetic metals [1]. Two magnetic layers are separated by a normal layer in a basic spintronics device. It displays both the opposite spin transfer phenomenon [4, 5] and the well-known giant magneto-resistance (GMR) effect [2, 3].

From an experimental perspective, it is still necessary to precisely identify the

spin-wave (SW) eigen-modes in hybrid magnetic nanostructures [6–11]. The precise kind of modes induced by a polarised current in spin transfer nano-oscillators (STNOs) is of special interest. Finding the associated symmetry underlying each mode is crucial in this situation. It provides a basic understanding of their selection criteria and potential mutual connection processes.

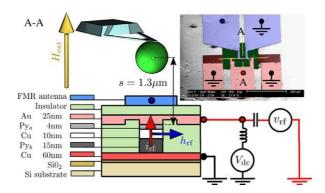


Fig. 1.1Schematic representation of the experimental setup. Two independent excitation circuits are used: in red, the circuit allowing the injection of an RF perpendicular-to-planethrough nano-pillar current (irf. redarrow);inblue,thecircuit allowing the generation of an RF in-plane magnetic field blue (hrf.intraorinterSTNOs. Italsodetermines the optimum strategy to couple to the autooscillatingmodeobservedwhenthespintransfertorquecompensatesthedamping, vital knowledge to achieve phase synchronization in arrays of nano-pillars [12]. These SW modes also have a fundamental influence on the high frequency propertiesofspin-valvedevices, and in particular on the noise of magneto-resistive sensors [13, 14].

This work concentrates on a comprehensive identification of the SW eigenmodes in a normally magnetized circular spin-valve nano-pillar, where the axial symmetry is preserved. We shall perform a comparative spectroscopic study of the SW modes excited either by an RF current flowing perpendicularly throughthe nano-pillar, as used in spin-torque driven ferromagnetic resonance (ST-FMR) [15,16],orbyahomogeneousRFin-planemagneticfield,asusedinconventional

FMR. The experimental spectra are compared to micromagnetic simulations of the time decay response of the local magnetization to a small perturbation of the magnetic configuration. This allows identifying and labeling the observed SW eigen-modes, according to their symmetry and to the dipolar coupling between the two magnetic layers.

Thespin-valvestructureusedinthisstudyisastandardPermalloy (Ni<sub>80</sub>Fe<sub>20</sub>Py) bi-layer structure sandwiching a 10 nm copper (Cu) spacer: the thicknesses of the thin Py<sub>a</sub>and the thick Py<sub>b</sub>layers are respectively  $t_{\overline{a}}$ 4 nm and  $t_b$ 15nm.Theextendedfilmispatternedbystandarde-beamlithographyandion-milling techniques to a nano-pillar of radius 125 nm. The magnetic parameters of this sample have been measured in [17], and are reported in Table 1.1. The top Cu andbottomAucontactelectrodesareshowninredinFig.1.1.Theyareimpedance matched to allow for high frequency characterization by injecting an RF current  $i_{rf}$  through the device. Hereafter, spectra associated to SW excitations by this part of the microwave circuit will be displayed in a red tone.

The originality of our design is the addition of an independent top microwave antennawhosepurposeistoproduceanin-planeRFmagneticfield  $h_{\rm rf}$  atthenano- pillar location. In Fig. 1.1, this part of the microwave circuit is shown in blue. In- jecting a microwave current from a synthesizer inside the top antenna produces a homogeneous inplanelinearly polarized microwave magnetic field, oriented perpendicularly to thest ripedirection. Hereafter, spectra associated to SW excitations by this part of the microwave circuit will be displayed in a blue tone.

We shall use in this study a method independent of transport to detect the magnetic resonance in neticresonance in side as pinvalvenanostructure: a Magnetic Resonance Force Microscope (MRFM) [18–20], here afternamed mechanical-FMR. A first decisive advantage of the mechanical-FMR technique is that the detections cheme does not rely on the SW spatial symmetry because it measures the change in the longitudinal component of the magnetization. It thus probes all the excited SW modes, independently of their phase [21, 22]. A second decisive advantage is that mechanical-FMR is a very sensitive technique that can measure the magnetization dynamics in nanostructures buried under metallic electrodes [23]. Indeed, the probe is a magnetic particle attached at the endofasoft can tile verand is coupled to the sample through dipolar interaction.

Themechanical detector consists of a 800 nm diameter sphere of soft amorphous Fe (with 3 % Si) glued to the apex of an Olympus Bio-Lever having a spring

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con- stantk5 mN/m. In our setup, the separation between the center of the spherical probe and the nano-pillar is set to 1.3  $\mu$ m (see Fig. 1.1), which is a large distance considering the lateral size of the sample. The external magnetic field produced by an electromagnet is oriented out-of-plane, exactly along the nano-pillar axis z. In our study, the strength of the applied magnetic field shall exceed the saturation field (8 kOe), so that the nano-pillar is studied in the saturated regime. The mechanical sensor measures then the spatial average of the longitudinal component  $M_z$  of magnetization:

$$(M) = \frac{1}{M} (\mathbf{r}) d^3 \mathbf{r}, \tag{1.1}$$

#### 1.2 IdentificationoftheSpin-WaveModes

In this section, we discuss the boundary-value problem for SW propagation inside normallymagnetizeddisks, where the confinement leads to a discrete SW spectrum.

Neglecting the thickness dependence, only three indices are required to label the resonance peaks: the usual azimuthal and radial indices for a single disk (4,m), plus an additional index referring to the anti-binding or binding (A or B) coupling between the two magnetic layers in mutual dipolar interaction.

#### 1.2.1 SingleMagneticDisk

SWeigen-modesarethesolutionsofthelinearized equation of motion of the magnetization, obtained by decomposing the instantaneous magnetization vector M(t) into a static and dynamic component [25]:

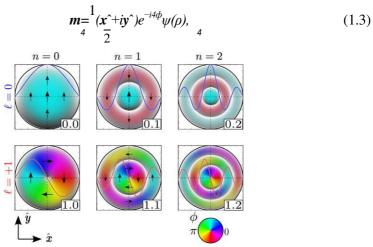
$$\underline{\underline{M(t)}} = \underline{u}^+ + \underline{m(t)} + O^{\dagger} \underline{m}^{2}^{\,c}, \tag{1.2}$$

where the transverse component m(t) is the small dimensionless deviation (m

of the magnetization from the local equilibrium direction, u. Inferromagnets,  $|M| = M_s$  is a constant of the motion, so that the local orthogonality condition  $\hat{u}m = 0$  is required.

We restrict our study to the case of thin layers so that one can assume that the magnetizationdynamicsisuniformalongthethickness. For an ormally magnetized

disk,where uz, the SW modes can be classified according to their behavior under rotations in the x-yplane (2D vector equations with polar coordinates  $\rho$  and  $\phi$ ):



**Fig.1.2**Colorrepresentation of the Bessel spatial patterns for different values of the azimuthal mode index 4 (by row)andradialmodeindex n(bycolumn). The arrows area snapshot of the transverse magnetization  $m_V$ , labeled by the index v=4, n

The top left panel shows the  $\Rightarrow 0.0$  (40 = n0) mode, also called the uniform mode. It usually corresponds to the lowest energy mode since all the vectors are pointing in the same direction at all time. Below is the 4 = n0 mode. It corresponds to SWs that are rotating around the disk in the same direction as the

Larmorprecession. The corresponding phase is in quadrature between two orthogonal positions and this mode has an ode at the center of the disk. The variation upon the n0,1,2 index (4 being fixed) shows higher order modes with an increasing

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number of nodal rings. Each ring separates regions of opposite phase along the dialdirection. Allthese spatial patterns preserve the rotation invariance symmetry.

The coupling to an external coherent sourced epends primarily on the 4-index [27]astheexcitationefficiencyisproportionaltotheoverlapintegral

$$h_{\nu} \propto (\boldsymbol{m}_{\nu} \cdot \boldsymbol{h}_{1}), \tag{1.4}$$

where  $h_1(r)$  is the spatial profile of the external excitation field. It can be easily shown that a uniform RF magnetic field,  $h_1h_{rt}x$ , can only excite 40 SWmodes. Obviously, the largest overlap is obtained with the socalleduniformmode (n0). Higher radial index modes (n0) still couple to the uniform excitation butwithastrength that decreases as n increases [19,28]. The 40 normalmodes, however, are hidden because they have strictly no overlap with excitation. contrast, the RF current-created Oersted field,  $h_1 = h_{Oe}(\rho)$  (—  $\sin \phi x^+ \cos \phi y^-$ )has

anorthoradialsymmetryandcanonlyexcite4=+1SWmodes.

#### 1.2.2 DoubleMagneticDisks

The interaction between two identical magnetic layers leads to the splitting of uniformmodeineachlayerintotwocollectivemodes:thebindingandantibinding modes. This has been observed in interlayer-exchange-coupled thin films [29] intrilayeredwireswherethetwomagneticstripesaredipolarlycoupled[30].Intheca se where the two magnetic layers are not identical (different geometry or mag- netic parameters), this general picture continues to subsist. Although both isolated layershaveeigen-modeswithdifferenteigenfrequencies, the collective magnetization dynamics in each layer are still symmetrically or anti-symmetrically coupled. But here, the precession of magnetization can be more intense in one of the two layers and the spectral respect isolated coupled SWmodes with SWmodesisreduced, asitwas observed in both the dipolarly-[30] and exchangecoupled cases [31].

Here, we assume that the dominant coupling mechanism between the two mag- netic disks in the nano-pillar is the dipolar interaction. We neglect any exchange coupling between the magnetic layers mediated through the normal couplingassociatedtopurespincurrents[7,32]inourallor metallicspin-valvestructure.Ournano-

pillarconsistsoftwodislikemagneticlayers,havingdifferentthicknesses, with  $t_a < t_b$ , and different saturation magnetizations, with  $M_a < M_b$ . In the normally saturated state which is considered here, this difference of magnetization leads to a difference in the internal demagnetizing field of each layer (self dipolarcoupling)andinthestrayfieldofonelayerontheotherone(crossdipolarcoupling). Asaresult,theeigen-

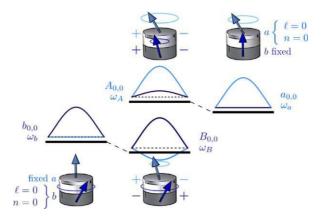
frequencyoftheuniformmodeofthethinlayerislargerthan the one of the thicklayer,  $\omega_a > \omega_b$ . This is represented in Fig.1.3(a), onboth sides of the bilayer diagram, where we have reported on an energy scale their relative positions.

In the perpendicular geometry, the strength of the dynamical dipolar coupling  $\Omega$  between the layers depends on the cross tensor element  $jj^r$ , associated to the in-plane component of the dipolar magnetic field produced inside layer j by the in-plane component of the magnetization of layer  $j^r$  (see [17, 33]). It is attractive (lower in energy) when both layers precessinantiphase because the dynamical

dipolar charges in each layer are alternate (anti-ferromagnetic coupling). Thus the binding state corresponds to a collective motion where the two layers vibrate anti- symmetrically (B) and the anti-binding state to a collective motion where they vi- brate symmetrically (A). In this case, the larger of the frequencies  $(\omega_a)$  shifts upby

$$\Delta \omega = \frac{\Omega^2}{a - \omega_b}, \tag{1.5}$$

while the smaller one  $(\omega_b)$  shifts down by the same amount. This effect is summa-rized in Fig. 1.3.



**Fig.1.3**Schematicrepresentationofthecoupleddynamicsbetweentwodifferent magnetic disks. When the two disks are dynamically coupled through the dipolar interaction, the binding state B corresponds to the two layers oscillating in antiphase at  $\omega B$ , with the precession occurring mostly in the thick layer, whereas the anti-binding state A corresponds to the layer so scillating in phase at  $\omega A$ , with the precession mostly in the thin layer

amplitudesinthetwolayersisgivenby

$$|c_b/c_a|_{\omega A} = \Delta \omega/(\gamma h_{a,b})' \omega \frac{\Omega}{\omega}$$
 (1.6)

Fortheexperimental parameters,  $\Omega/(\omega_a \omega_b)$  0.1, i.e., the precession amplitude in the disk b is about 10% of that in the disk a. Thus, although the dipolar coupling induces a small spectral shift (second order in the coupling parameter (1.5)), its influence in the relative precession amplitude is significant (first order in the coupling parameter (1.6)).

## 1.2.3 MicromagneticSimulationsvs.Mechanical-FMR Experiments

Althoughtheanalyticalformalismpresented above allows estimating the spectrum, several approximations have been made. In particular, we have assumed total pin-ning at the disks boundary for the SW modes and no variation of the precession profile along the disks thicknesses (2D model), and we have neglected the dependence on the mode index vof the dynamic dipolar coupling. Instead of developing a more complex analytical

formalism, we have performed a complete calculation of the SW spectra inside our nano-pillar sample using the open source micromag- netic simulation package Nmag [34]. Nmag is a finite element solver based on the generalpurposemulti-physicslibraryNsim.ItisdevelopedbythegroupofH.Fan-gohrandT.FischbacherintheSchoolofEngineeringSciencesattheUniversityof Southampton.

**Table 1.1**Magnetic parameters of the thin Pya and thick Pyblayers measured by mechanical- FMR in the nano-pillar

measurea e	y inicentamean	I will the ha	no pinai	
$4\pi Ma(G)$	$\alpha a$	$4\pi Mb(G)$	ab	γ(rads—
				$^{1}G^{-1}$ )
$8.0 \times 10^3$	1.4×10 <sup>-2</sup>	$9.6 \times 10^{3}$	0.85 ×10	-2 1.87×10 <sup>7</sup>

Inthissimulation,thefullthree-dimensional(3D)dynamicsofthebi-layersystem is calculated. The thin layer is discretized with a mesh of 4 nm (equal to its thickness), and the thick layer with a mesh of 3 nm. The numbers of nodes

are

re-

spectively6135inthethinlayerand37598inthethicklayer. Themagnetic parameters introduced in the code are the ones reported in Table 1.1. The magnetization

vectorisassumedtobeuniforminsideeachcell,whichisvalidbecausethecellsize issmallerthantheexchangelength  $\Delta 5$ nm inPermalloy.Weemphasizethatthe simulationsincorporatetheperturbingpresenceofthemagneticsphereattached on

theMRFMcantilever.Moreover,the10nmthickCuspacerisreplacedbyvacuum , so that the layers are only coupled trough the dipolar interaction (spin diffusion effects are absent).

The first step is to calculate the equilibrium configuration of the normally mag- netized nano-pillarat  $H_{\rm fix}$  10 kOe. The external magnetic field is applied exactly =  $\times$ 

along z and the spherical probe with a magnetic moment  $m210^{-10}$  emu is placedontheaxialsymmetryaxisatadistances 1.3  $\mu$ mabove the upper sur-

face of the nano-pillar (see Fig. 1.1). The convergence criterion introduced in the code is  $1/M_s dM/dt < 1^\circ/ns$ . The result reveals that the equilibrium configuration is almost uniformly saturated along z. Still, a small radial flaring ( $<5^\circ$ ) of the mag- netization from z is observed at the periphery of the thick and thin layers.

InordertocalculatetheSW spectrum corresponding to a given excitation of this equilibrium state at  $H_{\rm ffx}10$  kOe, we record the time decay response of the local

magnetizationtoasmallperturbationofthemagneticconfiguration, andweFouri er transform the ringdown of the transverse magnetization [35]. Two different initial conditions are simulated. For the conventionalFMR spectrum, in which a uniform P.EfieldieusedtoevoiteSW modes weusethefollowing perturbation vector field:

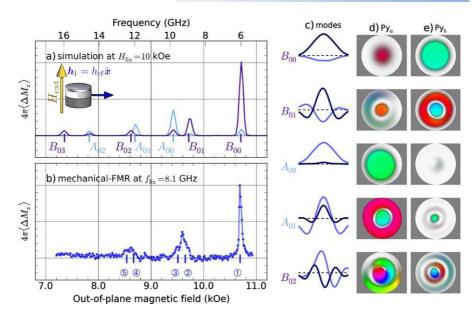
RFfieldisusedtoexciteSWmodes, weusethefollowing perturbation vector field: m = 9  $J_0(k_{0n}\rho)x^{\hat{}}$ . (1.7)

Thisformcorrespondstoanexcitation whichputsthesameenergyinthelowest 6

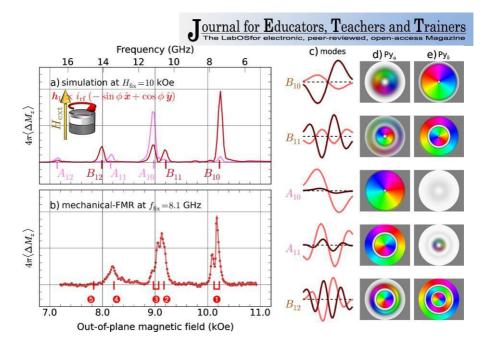
*n*-index modes with azimuthal index 40. For the spectrum excited by an RF current flowing through the nano-pillar, which creates an orthoradial RF Oersted field in the magnetic volume, we use:

$$m=9$$
  $J_1(k_{1n}\rho)(-\sin\phi x^+\cos\phi y^-).$  (1.8)  $n < 6$ 

Attime t0, we add at every mesh the perturbation vector field defined in (1.7) or in (1.8) to the local unit vector along the equilibrium magnetization. The first one



**Fig.1.4**Panel(**a**)isthenumericallycalculatedspectralresponsetoauniformexcit ati®nfield  $h_1x$ , from a 3D micromagnetic simulation performed at  $H_{\text{fix}}10$  kOe. The peaks are labeled according to their precession profiles shown in (**c**), (**d**), and (**e**). A *light* (*dark*) *color* is used to indicate the energy stored in the thin (thick) layer. Panel (**b**) shows the experimental spectrum measuredbymechanical-FMRwhenexcitingthenanopillarbyahomogeneousRFmagneticfield at  $f_{\text{fix}} = 8.1 \text{ GHz}$ 



 $\begin{tabular}{ll} Fig. 1.5 Panel (a) is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to an orthogonal excitation field $d^{-\infty}$ is the simulated spectral response to a simulated spectral$ 

 $h_1 \sin \phi x \cos \phi y$ . Panel(**b**) is the experimental spectrum measured by mechanical FMR for an RF current excitation. Panels (**c**)—(**e**) show the simulated precession patterns of the eigen-vectors

#### 1.3 Conclusion

In conclusion, we investigated the SW eigen-modes in the prototype of a STNO, a normally magnetisednano-pillar made of two magnetic layers in dipolar contact, using the MRFM method [20]. The SW spectra of this passive STNO stimulated by an RF current flowing perpendicularly across the layers and by a homogeneous in-plane RF magnetic field were compared. We discovered that the two aforementioned excitation techniques ignite rather distinct SW modes, each with an azimuthal index of 4 0 and 4 1, respectively. Additionally, we created a straightforward analytical theory that offers a thorough labelling of every SW eigen-mode of a magnetic nano-pillar in the axially symmetric scenario under study. The standard azimuthal and radial indices 4 and n, which are used for the SW modes of a single magnetic disc, as well as an extra index that refers to the binding or anti-binding (B or A) dipolar coupling between the two magnetic discs are needed for this labelling. The outcomes of 3D micromagnetic simulations using the Nmag software were also contrasted with the experimental and analytical findings that were achieved. We discovered that the discriminating parameter for the selection rules of the SW mode excitation is the 4-index, which is associated with the azimuthal symmetry of the SW modes. Our findings, in our opinion, are

crucial for both the experimental determination of the STNO parameters and the optimisation of the properties of nano-spintronic devices, and specifically STNOs.

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