



Angle-dependent spin wave spectra of permalloy ring arrays

Shuxuan Wu(吴书旋), Zengtai Zhu(朱增泰), Yunxu Ma(马云旭), Jinwu Wei(魏晋武), Senfu Zhang(张森富), Jianbo Wang(王建波), and Qingfang Liu(刘青芳)

Citation: Chin. Phys. B, 2022, 31 (11): 117505. DOI: 10.1088/1674-1056/ac8af9

What follows is a list of articles you may be interested in

Experimental observation of interlayer perpendicular standing spin wave mode with low damping in skyrmion-hosting [Pt/Co/Ta]₁₀ multilayer

Zhen-Dong Chen(陈振东), Mei-Yang Ma(马眉扬), Sen-Fu Zhang(张森富), Mang-Yuan Ma(马莽原), Zi-Zhao Pan(潘咨兆), Xi-Xiang Zhang(张西祥), Xue-Zhong Ruan(阮学忠), Yong-Bing Xu(徐永兵), and Fu-Sheng Ma(马付胜)

Chin. Phys. B, 2022, 31 (11): 117501. DOI: 10.1088/1674-1056/ac7bf9

Interaction region of magnon-mediated spin torques and novel magnetic states

Zai-Dong Li(李再东), Qi-Qi Guo(郭奇奇), Yong Guo(郭永), Peng-Bin He(贺鹏斌), and Wu-Ming Liu(刘伍明)

Chin. Phys. B, 2021, 30 (10): 107506. DOI: 10.1088/1674-1056/ac04a8

Spin waves and transverse domain walls driven by spin waves: Role of damping

Zi-Xiang Zhao(赵梓翔), Peng-Bin He(贺鹏斌), Meng-Qiu Cai(蔡孟秋), Zai-Dong Li(李再东)

Chin. Phys. B, 2020, 29 (7): 077502. DOI: 10.1088/1674-1056/ab90e5

Voltage control of ferromagnetic resonance and spin waves

Xinger Zhao(赵星儿), Zhongqiang Hu(胡忠强), Qu Yang(杨曲), Bin Peng(彭斌), Ziyao Zhou(周子尧), Ming Liu(刘明)

Chin. Phys. B, 2018, 27 (9): 097505. DOI: 10.1088/1674-1056/27/9/097505

Angle-dependent spin waves in antidot bilayers

Hu Chun-Lian (胡春莲), Liao Leng (廖棱), Stamps R

Chin. Phys. B, 2014, 23 (12): 127501. DOI: 10.1088/1674-1056/23/12/127501

Angle-dependent spin wave spectra of permalloy ring arrays

Shuxuan Wu(吴书旋)¹, Zengtai Zhu(朱增泰)², Yunxu Ma(马云旭)¹, Jinwu Wei(魏晋武)¹,
Senfu Zhang(张森富)¹, Jianbo Wang(王建波)^{1,3}, and Qingfang Liu(刘青芳)^{1,†}

¹Key Laboratory for Magnetism and Magnetic Materials of the Ministry of Education, Lanzhou University, Lanzhou 730000, China

²Songshan Lake Materials Laboratory, Dongguan 523808, China

³Key Laboratory for Special Function Materials and Structural Design of the Ministry of the Education, Lanzhou University, Lanzhou 730000, China

(Received 30 June 2022; revised manuscript received 4 August 2022; accepted manuscript online 19 August 2022)

We investigated the angle-dependent spin wave spectra of permalloy ring arrays with the fixed outer diameter and various inner diameters by ferromagnetic resonance spectroscopy and micromagnetic simulation. When the field is obliquely applied to the ring, local resonance mode can be observed in different parts of the rings. And the resonance mode will change to perpendicular spin standing waves if the magnetic field is applied along the perpendicular direction. The simulation results demonstrated this evolution and implied more resonance modes that maybe exist. And the mathematical fitting results based on the Kittel equation further proved the existence of local resonance mode.

Keywords: spin waves, magnetic resonance spectra, spintronic devices

PACS: 75.30.Ds, 32.30.Dx, 85.75.-d

DOI: 10.1088/1674-1056/ac8af9

1. Introduction

Periodically magnetic patterned structures have been extensively studied in the last decades.^[1–8] As the progress of nanofabrication techniques, it is possible to prepare the nanoscale or submicron-scale pattern arrays on the magnetic films. To date, the arrays of dots, rings, and wires have been investigated by experimental observation or micromagnetic simulation.^[4,9–13] Magnetic patterned films have opened the possibility of studying and designing the new magnetic storage, magnetic logic and spintronic devices. For example, there have been many studies on nanoring magnetic tunnel junctions.^[14–17] Their static and dynamic magnetic properties strongly depend on the size and topological structures. The patterned magnetic films are also known as the magnonic crystals. As the propagation of light in the photonic crystals, the propagation properties of spin waves in the magnonic crystals can be modified by the well-designed patterns.^[18] Therefore, magnonic crystals are considered to offer new functionalities that are unavailable in regular electronic devices. Depending on the scale of magnetic pattern structures, exchange interaction and dipolar interaction play significant roles, respectively. In fact, the patterns introduce a kind of size confine, which makes the magnetization motion become very different from that in the continuous films. Demagnetizing field, edge pinning and anisotropy of patterns are all supposed to be important for the magnetic properties of patterned films.^[5,11,19]

Although the ferromagnetic resonance (FMR) spectra of patterned films have been wisely researched, recent studies showed rich spin wave spectra of magnetic disks and rings

at different magnetized states.^[13,20,21] For example, when the magnetic field was applied in the film plane, the magnetic disks at remanence showed a magnetization transition from onion to vortex state depending on the magnitude of the applied field.^[22–24] On the other hand, splitting of excitation spectra was observed if the magnetic rings magnetized along the direction of out-of-plane.^[21] Furthermore, the resonance spectra of ring arrays are more complex than those of disks because of the unique topological structure of rings. In other words, disks are internally connected but rings are not. The geometric ring will introduce the radial demagnetizing field, which provides more dimensions to design the spintronic devices or storage the information. The more complex structure represents more difficulties in manipulating the performance of rings. The previous research focused on the FMR spectra and dynamic magnetic properties of rings. Multi-peak resonance was always observed at both vertical and horizontal magnetization states. However, the evolution of resonance modes is still not clear, which is important to understand and adjust the dynamic magnetic properties of rings.

In this paper, we prepared a series of permalloy ring arrays with the fixed outer diameter and various inner diameters and presented the observation of FMR spectra when the applied magnetic field rotated from in-plane to out-of-plane direction. Several absorption modes were observed at different magnetized angles. Micromagnetic simulations qualitatively agree with the experimental results and indicate the evolution of resonance modes. When the field is applied obliquely, the local resonance modes depend on the magnetic field magni-

[†]Corresponding author. E-mail: liuqf@lzu.edu.cn

tude, while the perpendicular spin standing wave (PSSW) can be observed if the magnetic field is applied vertically. Mathematical fitting results further prove the evolution pattern of resonance modes.

2. Experimental and simulation methods

Isotropic permalloy (Py, $\text{Fe}_{20}\text{Ni}_{80}$) film was deposited on Si (100) wafer by radio frequency magnetron sputtering technique. The base pressure of the sputtering chamber was better than 5×10^{-5} Pa. Based on the sectional images of scanning electron microscopy (SEM), the thickness of the permalloy film was about 220 nm. The as-deposited film was divided into several $5 \text{ mm} \times 5 \text{ mm}$ square chips. Periodic ring arrays were fabricated on the prepared samples by using a laser direct write lithography (DWL, Heidelberg 66FS) and lift-out processes. The outer diameters (D) of rings were fixed at $30 \mu\text{m}$. The inner diameters (d) were $10 \mu\text{m}$, $15 \mu\text{m}$, $20 \mu\text{m}$, $25 \mu\text{m}$, respectively. The space between centers of rings was kept at $40 \mu\text{m}$. For convenience, the samples with d of $10 \mu\text{m}$, $15 \mu\text{m}$, $20 \mu\text{m}$, $25 \mu\text{m}$ are represented as sample 1–4, respectively. The morphology of the prepared ring arrays was characterized by field emission scanning electron microscopy (FESEM, Tescan Mira 3 xmu, Czech Republic). Dynamic properties were characterized by electron spin resonance (ESR, JES-FA300) commercial equipment. All measurements were conducted at room temperature.

To validate the experimental results, micromagnetic simulation is performed by the MuMax3 code.^[25] The magnetization dynamics of a ring is governed by the Landau–Lifshitz–Gilbert (LLG) equation^[26]

$$\frac{d\mathbf{M}}{dt} = -\gamma(\mathbf{M} \times \mathbf{H}_{\text{eff}}) + \frac{\alpha}{M_s} \mathbf{M} \times \frac{d\mathbf{M}}{dt}, \quad (1)$$

where \mathbf{M} represents the magnetization, M_s is the saturation magnetization, \mathbf{H}_{eff} is the effective magnetic field, γ is the gyromagnetic ratio and α is the dimensionless Gilbert damping constant. The magnetic parameters of the Py film are chosen as $M_s = 8.0 \times 10^5$ A/m, exchange constant $A = 1.3 \times 10^{-11}$ J/m, uniaxial anisotropy constant $K_u = 0$ J/m³, $\alpha = 0.02$ in the simulation. The unit cell size of a single ring is set as $5 \text{ nm} \times 5 \text{ nm} \times 4 \text{ nm}$. To indicate the spatial profiles of resonance amplitude for dynamic magnetization at different fields, the excitation field is applied with sinc function type, $H = H_0 \text{sinc}(2\pi f_0(t - t_0))$, where the amplitude $H_0 = 5$ mT, frequency $f_0 = 9$ GHz, and $t_0 = 1$ ns. Then, the magnetization components as a function of time and space are computed by the fast Fourier transform. Therefore, the resonance frequencies of different spin excitation modes can be acquired, as well as the spatial distributions of the fast Fourier transformation amplitude at specific oscillation frequency.

3. Results and discussion

Figure 1 shows the SEM image of sample 4. It can be seen that uniform edges were achieved in most of the rings. The measured diameters of rings are the same as the designed one. For convenient description, we will use the designed size to represent the samples in the following text.

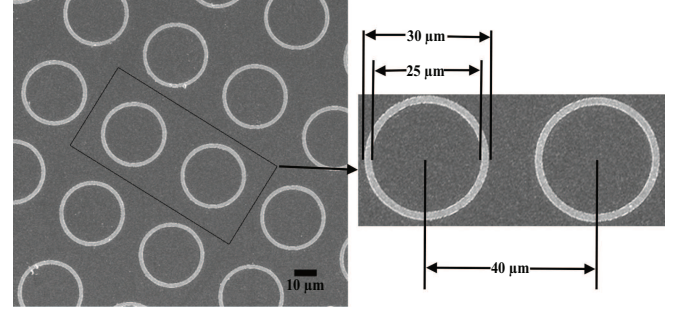


Fig. 1. Scanning electron micrograph of permalloy ring arrays with the inner diameter of $25 \mu\text{m}$.

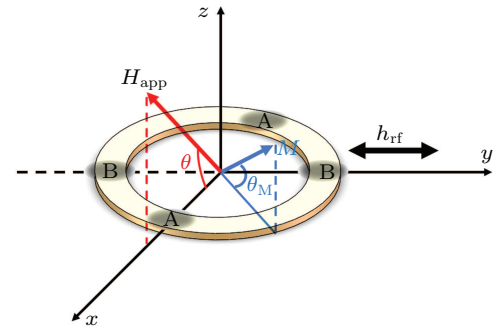


Fig. 2. The coordinate system used for the measurement and analysis of FMR, and the spatial relationship between \mathbf{H}_{app} , \mathbf{h}_{rf} , and \mathbf{M} . The sample lies in the xy -plane.

Figure 2 shows the coordinate system used to analyze FMR results. \mathbf{H}_{app} and \mathbf{h}_{rf} are the applied DC and microwave magnetic fields, respectively. θ is the angle between \mathbf{H}_{app} and film plane. θ_M is the angle between \mathbf{M} and film plane. Figure 3 shows the in-plane ($\theta = 0$) field-swept FMR differential spectra of samples at 9 GHz. As can be seen, high field (HF) mode and low field (LF) mode are observed for sample 1. Similar results were also found in nano-ring magnetic tunnel junction samples, where HF and LF modes were identified as acoustic-like and optical-like fundamental resonance modes, respectively.^[16,17] The HF mode exhibits larger intensity. When the inner diameter increased, the HF mode weakened while the LF mode enhanced. Meanwhile, a transition resonance mode is clearly visible between the two modes when $d = 20 \mu\text{m}$. Furthermore, resonance peaks shift slightly with the change of d . Since all the resonance modes should follow the Kittel equation, this can be explained as the result of spatial variations of demagnetizing fields.

In order to understand how the magnetization state affects resonance spectra, we measured the out-of-plane FMR spectra of samples. Figure 4 shows the FMR spectra of different samples as a function of θ , which is the angle between

film plane and applied magnetic field. The resonance fields would increase when the applied field rotates from the easy axis (in-plane) to the hard axis (out-of-plane). Let us focus on the resonance spectra of sample 1 as shown in Fig. 4(a). When θ changed from 2° to 72° , noticeable mutation was not observed no matter in relative intensity or the number of resonance peaks. When θ changed from 72° to 81° , two resonance peaks tended to merge into one peak with a relatively larger linewidth. If the magnetic field was close to the perpendicular direction ($\theta = 90^\circ$), the resonance peak split into 3 modes in which the intensity increased with magnetic field. For samples 2–4, the similar trend was observed as shown in Figs. 4(b)–4(d). That is, resonance peaks gradually merged at lower angle and re-split at higher angle, which happened near $\theta = 70^\circ, 80^\circ$. But for sample 3, the transition mode is always

visible when $\theta \leq 80^\circ$. To conclude, the resonance modes are divided into three stages with the rise of θ , and $70^\circ, 80^\circ$ are two characteristic points of mode evolution.

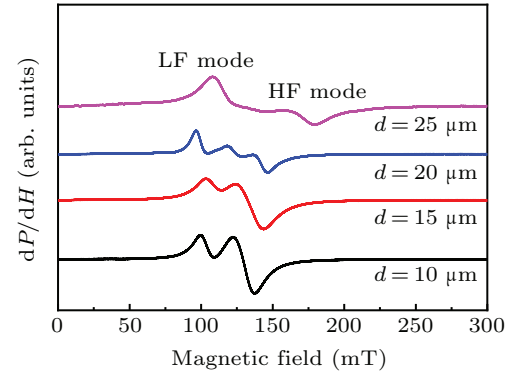


Fig. 3. In-plane FMR spectra of samples with different d at 9 GHz.

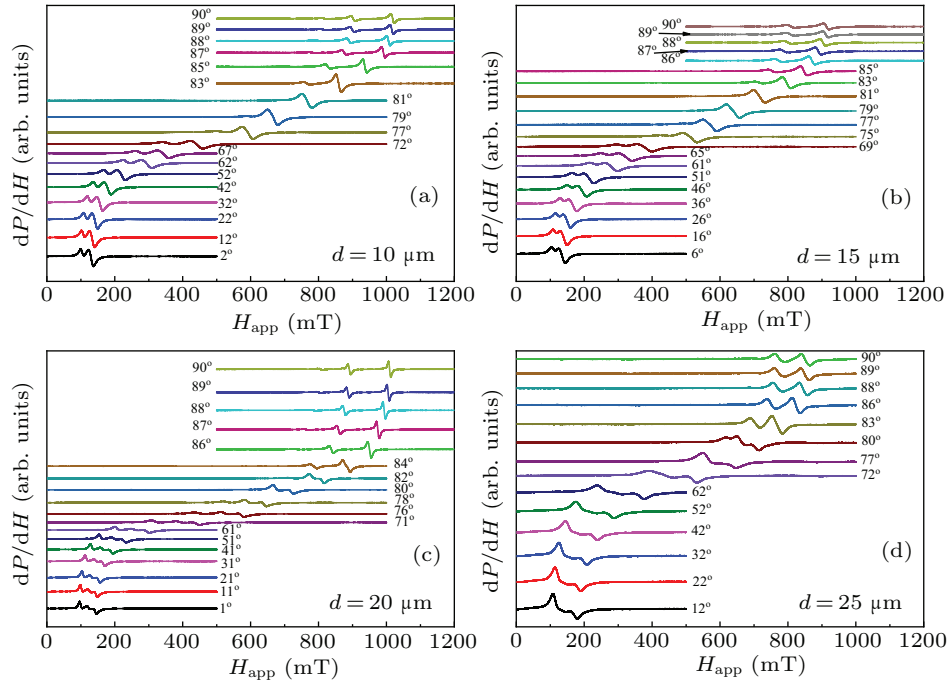


Fig. 4. Angle-dependent out-of-plane FMR spectra of samples with different d at 9 GHz. The angle indicated in the figure is the angle between the external magnetic field and the film plane.

Micromagnetic simulation was used to further understand the evolution of resonance modes. Figure 5 shows the simulation results of angle-dependent FMR differential spectra and resonance mode profiles for a Py ring with the size of $D = 1200$ nm and $d = 1000$ nm. The thickness of the ring is 4 nm. The ratio of D and d for simulation item is the same as that of sample 4. DC magnetic field is along the x axis, and excitation field is along the y axis. We have selected four characteristic points of $0^\circ, 70^\circ, 80^\circ$, and 90° to simulate the actual resonance evolution in the samples. Simulation results show two distinctly different resonance modes when the field is obliquely or horizontally applied on the Py ring. These two resonance modes correspond to LF and HF modes in the experimental results. For LF mode, the magnetization precession

mainly occurs in area B, as the gray part shown in Fig. 2. For HF modes, multiple resonance peaks are observed. The magnetization precession mainly occurs in area A. The upper parts of Fig. 5 show the resonance profiles under different DC magnetic fields, which shows that the multiple HF modes are spin standing waves (SSW) of different orders. For area A, the DC magnetic field is applied along the radial direction of the Py ring. The limited size of radial direction promotes the formation of SSW, which makes multi-order HF modes easier to be observed than the LF mode. High-order HF modes were not observed in the experimental FMR spectra. This may be due to the first-order HF mode with a strong intensity, which makes other high-order signals difficult to be measured. When θ is 70° and 80° [Figs. 5(b) and 5(c)], the main resonance modes

are still LF mode and multi-order HF mode, and the resonance fields of each modes rise slightly. When $\theta = 90^\circ$ [Fig. 5(d)], the resonance mode transforms to the PSSW mode.^[27] The resonance profiles of the Py ring show that the precession of magnetization occurs throughout the sample instead of local resonance mode.

A distinct difference between experimental and simulation results is the critical angle of mode evolution. For experimental results, as we have pointed out, $\theta = 70^\circ, 80^\circ$ are two characteristic points. However, when $\theta \leq 80^\circ$, simulation results maintain the similar local resonance mode. When $\theta = 90^\circ$, the resonance mode of the Py ring changes to PSSW mode of multiple orders, as shown in Fig. 5(d). the evolution of resonance modes from local resonance to PSSW probably takes place between 80° and 90° . Because of the different sizes between experimental and simulation samples, the positions of peaks in simulation results would not agree well with the experimental data. Despite this, the simulation results give a reliable evolutionary tendency of the resonance spectra.

To further verify the reliability of simulation results, the Kittel equation is used to analyze resonance field (H_r) as a function of θ . Considering the geometric structure of Py rings, the demagnetizing fields are important parameters. Based on the coordinate system shown in Fig. 2, the demagnetization factors of x , y and z directions are represented as N_x , N_y , and N_z , respectively. For the HF mode existing in area A, the Kittel equation can be written as^[28]

$$(\omega/\gamma)^2 = (H_r + (N_z - N_x)M_s)(H_r + (N_y - N_x)M_s), \quad (2)$$

where ω is the angular frequency of the AC excitation field. Also, for the LF mode,

$$(\omega/\gamma)^2 = (H_r + (N_z - N_y)M_s)(H_r + (N_x - N_y)M_s). \quad (3)$$

For the thin films, $N_z \geq N_y, N_x$, equations (2) and (3) can be rewritten as a unified equation

$$(\omega/\gamma)^2 = (H_r \mp H_d^R)(H_r \mp H_d^R + M_{\text{eff}}), \quad (4)$$

here, $(N_x - N_y)M_s = N_d^R M_s = H_d^R$ and $N_z M_s = M_{\text{eff}}$. H_d^R represents the effective radial demagnetizing field, and M_{eff} is the effective perpendicular demagnetizing field. If θ is considered, equation (4) can be further written as^[29]

$$(\omega/\gamma)^2 = (H_r \cos(\theta - \theta_M) - (M_{\text{eff}} \mp H_d^R) \sin^2 \theta_M \mp H_d^R) \times (H_r \cos(\theta - \theta_M) + (M_{\text{eff}} \mp H_d^R) \cos 2\theta_M), \quad (5)$$

here, θ_M is the angle between magnetization and film plane. We use Eq. (5) to fit the FMR data. Figure 6 shows the experimental results (dots) and fitting results (lines) of H_r as a function of θ . Obviously, the fitting lines agree well with the experimental results, which strongly verifies our discussion about the evolution of resonance modes. The fitting parameters of all the samples are listed in Table 1. With the rise of d , M_{eff} gradually decreases, which is due to the reduction of N_z . At the same time, H_d^R grows with the rise of d . In other words, the effective radial demagnetizing field will increase if the ring becomes narrower.

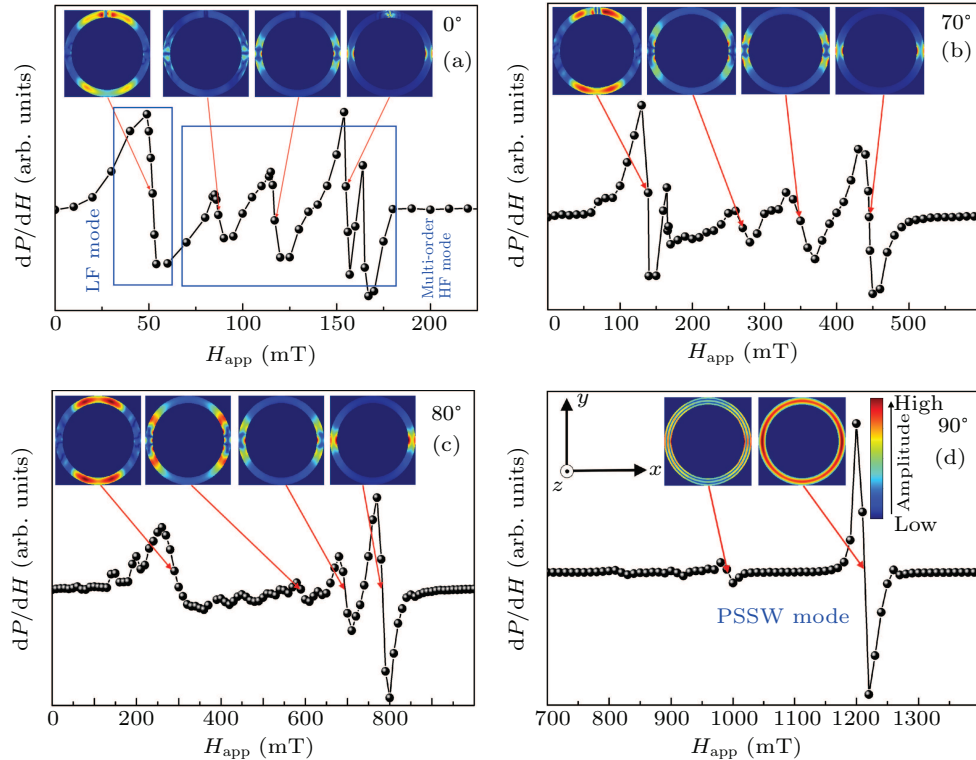


Fig. 5. Simulation FMR spectra and the profiles of resonance modes at different resonance fields. The angle between film plane and applied magnetic field is (a) 0° , (b) 70° , (c) 80° , and (d) 90° , respectively. The coordinate system shown in (d) is applicable for all the simulation results, which is the same as Fig. 2.

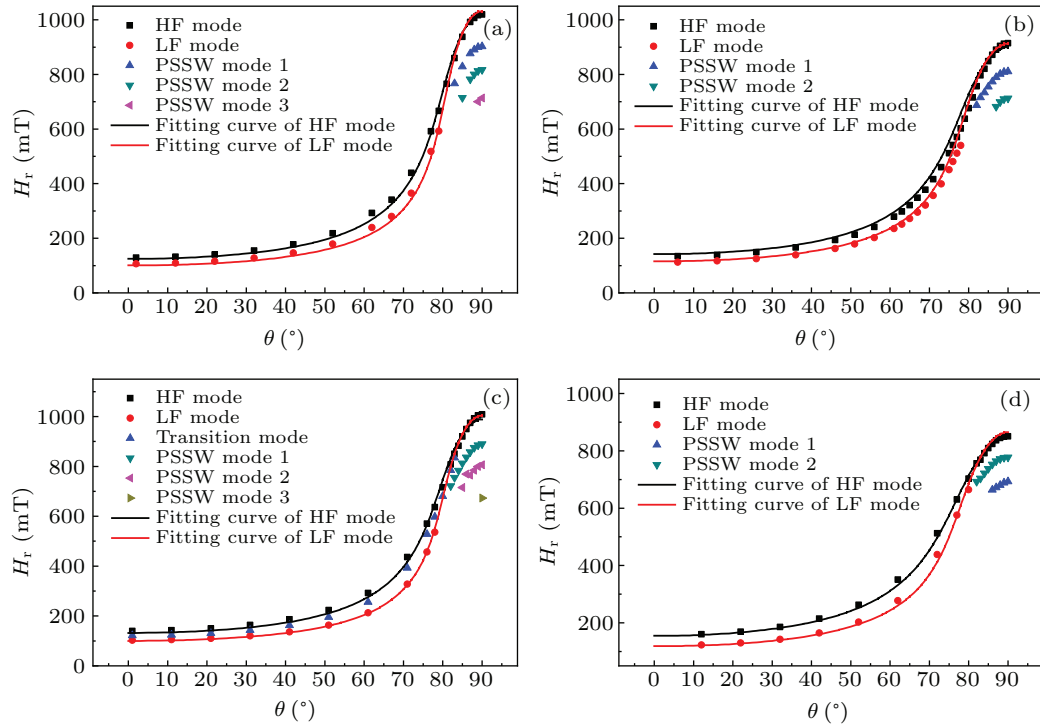


Fig. 6. Experimental (dots) and fitting (lines) results of FMR resonance field as a function of θ . (a) $d = 0 \mu\text{m}$; (b) $d = 15 \mu\text{m}$; (c) $d = 20 \mu\text{m}$; (d) $d = 25 \mu\text{m}$.

Table 1. The magnetic parameters obtained from FMR theoretical fitting for all the samples.

d (μm)	10	15	20	25
$4\pi M_{\text{eff}}$ (10^2 G)	72	60	69	55
H_d^R (10^2 Oe)	1.2	1.3	1.6	1.8

4. Conclusions

In summary, the experimental, micromagnetic simulation and mathematical fitting results drew a picture of resonance modes evolution for the Py rings. When H_{app} was applied on the film plane, it would break the original symmetry of Py rings at remanence. Local resonance modes consisting of the high and low field modes occurred at different parts of the rings. The simulation results revealed the existence of high-order high field modes, which were the SSW resonances derived from size limitation. As the magnetic field angle increased, the asymmetry of Py rings gradually disappeared. The high and low field modes merged to a single resonance peak with a relatively larger linewidth. When the magnetic field was applied vertically to the film plane, the multiple resonance peaks shown in the FMR spectra were actually PSSW resonance mode, which was due to the size limitation in the thickness direction of the thin films. Even though multiple peaks were usually measured in the FMR spectra of the magnetic patterned films, which probably possess completely different resonance modes. Our results revealed the evolution of resonance modes at different magnetization state for the Py ring arrays, which is meaningful in designing the microwave and spintronic devices.

Acknowledgement

Project supported by the National Natural Science Foundation of China (Grant Nos. 12074158, 12174166, and 12104197).

References

- [1] Neusser S and Grundler D 2009 *Adv. Mater.* **21** 2927
- [2] Podbielski J, Giesen F and Grundler F 2006 *Phys. Rev. Lett.* **96** 167207
- [3] Ding J, Singh N, Kostylev M and Adeyeye A O 2013 *Phys. Rev. B* **88** 014301
- [4] Zhou X and Adeyeye A O 2016 *Phys. Rev. B* **94** 054410
- [5] Zhu Z, Feng Z, Cheng X, Xie H, Liu Q and Wang J 2018 *J. Phys. D: Appl. Phys.* **51** 045004
- [6] Jorjick J, Demokritov S O, Hillebrands B, Bailleul M, Fermon C, Guslienko K Y, Slavin A N, Berkov D V and Gorn N L 2002 *Phys. Rev. Lett.* **88** 047204
- [7] Breitling A, Bublat T and Goll D 2009 *Phys. Status Solidi - R* **3** 130
- [8] Adeyeye A O and Singh N 2008 *J. Phys. D: Appl. Phys.* **41** 153001
- [9] Shaw J M, Silva T J, Schneider M L and McMichael R D 2009 *Phys. Rev. B* **79** 184404
- [10] Tse D H Y, Steinmuller S J, Trypiniotis T, Anderson D, Jones G A C, Bland J A C and Barnes C H W 2009 *Phys. Rev. B* **79** 054426
- [11] Wang C C, Adeyeye A O and Singh N 2006 *Nanotechnology* **17** 1629
- [12] Martyanov O N, Yudanov V F, Lee R N, Nepijko S A, Elmers H J, Hertel R, Schneider C M and Schönhense G 2007 *Phys. Rev. B* **75** 174429
- [13] Dutra R, Gonzalez-Chavez D E, Marcondes T L, Sommer R L, Parreiras S O and Martins M D 2019 *Phys. Rev. B* **99** 014413
- [14] Wen Z C, Wei H X, Han X F 2007 *Appl. Phys. Lett.* **91** 122511
- [15] Wei H X, He J X, Wen Z C, Han X F, Zhan W S and Zhang S F 2008 *Phys. Rev. B* **77** 134432
- [16] Qin J Y, Chen X, Yu T, Wang X, Guo C Y, Wan C H, Feng J F, Wei H X, Liu Y W and Han X F 2018 *Phys. Rev. Appl.* **10** 044067
- [17] Chen X, Qin J Y, Yu T, Han X F and Liu Y W 2018 *Appl. Phys. Lett.* **113** 142406
- [18] Herring C and Kittel C 1951 *Phys. Rev.* **81** 869
- [19] Shimon G, Adeyeye A O and Ross C A 2014 *Phys. Rev. B* **89** 024302

- [20] Zhou X, Tartakovskaya E V, Kakazei G N and Adeyeye A O 2017 *Phys. Rev. B* **96** 024446
- [21] Zhou X, Ding J, Kostylev M and Adeyeye A O 2015 *Appl. Phys. Lett.* **106** 112403
- [22] Taurel B, Valet T, Naletov V V, Vukadinovic N, de Loubens G and Klein O 2016 *Phys. Rev. B* **93** 184427
- [23] R Zarzuela, E M Chudnovsky and Tejada J 2013 *Phys. Rev. B* **87** 014413
- [24] Castel V, Ben Youssef J, Boust F, Weil R, Pigeau B, de Loubens G, Naletov V V, Klein O and Vukadinovic N 2012 *Phys. Rev. B* **85** 184419
- [25] Vansteenkiste A, Leliaert J, Dvornik M, Helsen M, Garcia-Sanchez F and Van Waeyenberge B 2014 *AIP Advances* **4** 107133
- [26] Gilbert T L 2004 *IEEE Trans. Magn.* **40** 3443
- [27] Rameev B, Yildiz F, Kazan S, Aktas B, Gupta A, Tagirov L R, Rata D, Buegler D, Gruenberg P, Schneider C M, Kämmerer S, Reiss G and Hütten A 2006 *Phys. Status Solidi A* **203** 1503
- [28] Kittel C 1948 *Phys. Rev.* **73** 155
- [29] Mizukami S, Ando Y and Miyazaki T 2001 *Jpn. J. Appl. Phys.* **40** 580