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Order/disorder phase composite Fe-Pt exchange-spring film magnets

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Introduction

As for thick film magnets, Aoyama *et al.* reported a 129- μ m-thick Fe-Pt film magnet with coercivity of 580 kA/m and (BH)_{max} of 104 kJ/m³, respectively.¹ Recently, W. F. Liu *et al.* have reported an approximately 7 μ m-thick Fe-Pt film magnet with H_c = 446 kA/m and (BH)_{max} = 124 kJ/m³, respectively.² On the other hand, a couple of researchers also reported Fe-Pt exchange-spring film magnets with a large (BH)_{max} value. For example, J. P. Liu *et al.* reported a 58 nm-thick Fe-Pt film magnet with (BH)_{max} = 420 kJ/m³. However the thickness of almost all the exchange-spring films reported previously is less than 1 μ m. In order to improve the magnetic properties of Fe-Pt film magnets, this contribution reports a fabrication of Fe-Pt exchange spring film magnets with the thickness of several microns by taking advantage of order and disorder phase transformation.

Experimental

Fe-Pt film magnets were prepared with a RF-sputtering apparatus. A target was composed of Pt chips placed on a 100-mm-diameter Fe disk. The composition of the films, which was evaluated with a SEM-EDX (Scanning Electron Microscope-Energy Dispersive X-ray spectrometer), was controlled by the number of Pt chips and set to Fe₅₀Pt₅₀. As-deposited films were composed of only disorder phase, and then a post-annealing was carried out in order to bring out the order and disorder to obtain order/disorder phase composite Fe-Pt transformation. In order exchange-spring thick film magnets, the samples were heated up to about 773 K under the heating rate of 323 K/min, in the vacuum of 10^{-4} Pa with an infrared furnace and then were cooled down to room temperature without a holding time. A sample composed of only order phase could be obtained under the post annealing at 1023 K. Magnetic and thermomagnetic properties were evaluated with a VSM (Vibrating Sample Magnetometer) and microstructure observation was carried out with a TEM (Transmission Electron Microscope). The analysis of crystal structure was carried out with an X-ray diffractometer (XRD).

Results and Discussions

Figure 1 shows bright-field TEM images of a 1.7 μ m-thick Fe-Pt film magnet (Sample A) annealed at 773 K. Nano-beam diffraction patterns of Figs .1 (a) and (b) indicated that the Spot Points X and Y shown in the figure are the disordered (A1) and the ordered (L1₀) phases, respectively, which means that the sample were composed of the both order and disorder phases.

As it was found that the annealing at 773 K enables us to synthesize exchange-spring magnets, we prepared a film annealed at 773K with 7 μ m- thicknesses. The M-H loop of the sample showed large coercivity of approximately 500 kA/m. In addition, the remanence value of approximately 0.9 T which was higher than the theoretical value of 0.7 T in the isotropic Fe-Pt alloy. In-plan and perpendicular values of remanence were almost the same, and it was clarified that the high value of 0.9 T is attributed to remanence enhancement in the sample. The above-mentioned 7 μ m-thick film had a superior recoil rate compared to that of Fe-Pt film with the thickness of 7 μ m composed of only order phase. This result indicates Fe-Pt exchange-spring thick films could be obtained by taking advantage of order and disorder composite phases.

References

- 1. H. Aoyama, and Y. Honkura, J. Magn. Soc. Jpn. 20, 237-240 (1996). (in Japanese).
- 2. W. F. Liu et al., J. Magn. Magn. Mater. 302, 201-205 (2006).
- 3. J. P. Liu, C. P. Luo, Y. Liu, and D. J. Sellmyer, Appl. Phys. Lett. 72, 483-485 (1998).



Fig. 1 (a) A1 phase.

Fig. 1 (b) $L1_0$ phase.

Fig. 1 Bright-field TEM images of a Fe-Pt film magnet annealed at 773 K (Sample A). NBD (Nano Beam Diffraction) indicates a spot point of the selected area diffraction analysis and the insets are the corresponding [001] diffraction patterns.