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Switching Field and Thermal Stability of CoPt/Ru Dot Arrays With Various Thicknesses

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The switching fields and thermal stability of CoPt/Ru dot arrays with various dot thickness δ (5–20 nm) were experimentally investigated as a function of the dot diameter, D , (130–300 nm). All dot arrays showed a single domain state, even after removal of an applied field equal to the remanence coercivity H_r . The angular dependence of H_r for the dot arrays indicated coherent rotation of the magnetization during nucleation. We estimated the values of the “intrinsic” remanence coercivity H_0 obtained by subtracting the effect of thermal agitation on the magnetization and the stabilizing energy barrier to nucleation $E_0/(k_B T)$. The variation in H_0 as a function of δ and D was qualitatively in good agreement with that of the effective anisotropy field at the dot center $H_k^{\text{eff}}(r=0)$, calculated taking account of the demagnetizing field in the dots. The ratio of H_0 to $H_k^{\text{eff}}(r=0)$ for the dot arrays with $\delta = 10$ nm increased from 0.53 to 0.70 as D decreased from 300 to 140 nm, and no significant difference in the $H_0/H_k^{\text{eff}}(r=0)$ ratio due to the difference in δ was observed. On the other hand, $E_0/(k_B T)$ decreased as δ decreased. $E_0/(k_B T)$ increased slightly as D decreased, but, was not so sensitive to D over the present D range.

Index Terms—CoPt perpendicular films, dot arrays, remanence coercivity, single domain, thermal stability, patterned media.

I. INTRODUCTION

AN introduction of a new recording system using patterned media is one possibility to overcome the tradeoff between thermal stability and recording writability. A large uniaxial magnetic anisotropy K_u of $1 \sim 2 \times 10^7$ erg/cm³ was obtained in Co-20at%Pt perpendicular films deposited on Ru seed layers [1]. Our studies for patterned films of CoPt(20 nm)/Ru revealed [2], [3] that dot arrays with dot diameter $D = 80$ –245 nm have a high thermal stability. Magnetic force microscopy (MFM) images revealed that all dot arrays showed a single domain state after removal of an applied field equal to the coercivity. It is generally assumed that the reversal of the single domain dot arrays begins with a nucleation, immediately followed by domain wall propagation [3]–[7]. The nucleation point is assumed to be located at the dot edges due to the damage during the dot fabrication [4], or some point within the dots due to structural inhomogeneities [5]–[7]. In this paper, the magnetic properties of CoPt/Ru dot arrays with various dot thicknesses δ were experimentally investigated as a function of D . We discuss the switching field and thermal stability of CoPt/Ru dot arrays in relation to the distribution of the effective anisotropy field in the dots due to the demagnetizing field.

II. EXPERIMENTAL PROCEDURE

Co₈₀Pt₂₀ films with film thicknesses of 5, 10, and 20 nm were deposited with Ru seed layers on 4-in SiO_x/Si substrates using a dc-magnetron sputtering system (ANELVA, E8002) [1]. No substrate heating was carried out during the deposition process. Dot arrays having D of 130–300 nm were made using laser interference lithography (LIL), with a constant periodicity of 600 nm [2]. Inter-dot magnetostatic coupling was negligibly

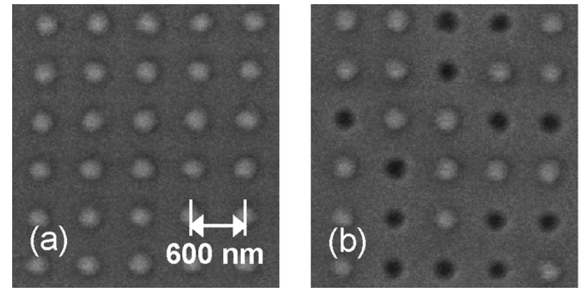


Fig. 1. MFM images of CoPt dot films with $D = 245$ nm. (a) Saturation remanent state. (b) After removal of an applied field equal to H_r .

small in these dot arrays. The dot patterns were formed in the magnetic layers by means of ion beam etching (IBE) with Ar⁺ ions. The magnetization curves of the dot arrays were examined by anomalous Hall effect (AHE) measurements [8]. The averaged AHE signals of about 3000 dots were detected. The remanence magnetization curves were measured at applied field sweep rates of ~ 10 Oe/s (using an electromagnet) and $\sim 10^8$ Oe/s (using a pulse field). Remanence coercivities at ~ 10 Oe/s and $\sim 10^8$ Oe/s were defined as H_r and H_r^p .

III. RESULTS AND DISCUSSIONS

A. Magnetic and Structural Properties of CoPt Dot Arrays

X-ray diffraction patterns indicated that CoPt films had a hcp structure with the c -axis perpendicular to the film plane and a c -axis distribution $\Delta\theta_{50}$ of about 2.8° , independent of film thickness [1]. Fig. 1 shows typical MFM images of dot arrays of $D = 245$ nm with $\delta = 20$ nm. The images were observed in (a) the saturation remanent and (b) after removal of a perpendicular applied field equal to H_r . MFM analysis revealed that all the dots showed a single domain state during the magnetization reversal in the present D (130–245 nm) and δ (5–20 nm) regions, although the exchange length of the magnetization of these films should be much smaller than D .

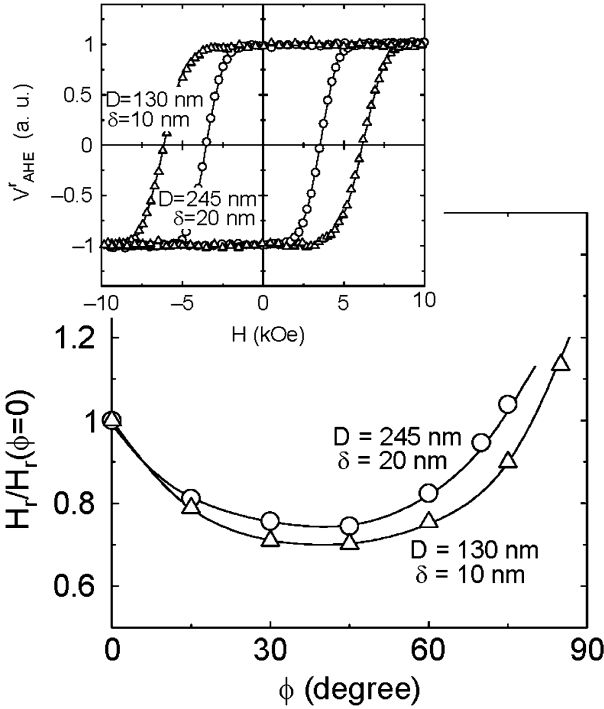


Fig. 2. Values of $H_r/H_r(\phi = 0)$ as a function of the angle ϕ between the applied field direction and the direction normal to the film plane. The inset shows magnetization curves and remanence magnetization curves along the perpendicular direction.

The inset to Fig. 2 shows magnetization curves (solid lines) and remanence magnetization curves (open symbols) measured along the perpendicular direction for dot arrays with different D and δ . The remanence magnetization curves were identical to the normal magnetization curves for all the dot arrays, indicating the magnetization reversal processes in the dots were irreversible. All the dot arrays showed similarly shaped magnetization curves, but the remanence coercivity H_r changed as a function of D and δ as we described later. It should be noted that the angular dependence of H_r for all dot arrays showed a minimum value at around $\phi = 45^\circ$ (ϕ is the angle between the applied field direction and the direction normal to the film plane) as shown in Fig. 2, indicating coherent rotation of the magnetization during nucleation in a small nucleation volume. No significant difference in the angular dependence of magnetization was observed due to the differences in D and δ .

B. Effective Anisotropy Field in Dots

We calculated the demagnetizing field in the dots along the perpendicular direction H_d as a function of the distance from the dot center r [3] and evaluated the distribution of the effective anisotropy field in the dots H_k^{eff} using

$$H_k^{\text{eff}}(r) = H_k - H_d(r) \quad (1).$$

where H_k is the anisotropy field ($H_k = 2K_u/M_s$, M_s is the saturation magnetization.) Fig. 3 shows the distribution of H_k^{eff} in dots having various D and δ . H_k^{eff} shows a minimum at the dot center $r = 0$. Dots with a large D/δ ratio have a broad minimum of H_k^{eff} but a decrease in the D/δ ratio results in a relatively pronounced minimum of H_k^{eff} at the dot center. It should

TABLE I
VALUES OF THE RATIO OF $H_k^{\text{eff}}(r = 0)$ TO $H_k^{\text{eff}}(r = 70)$ FOR DOT ARRAYS MADE OF VARIOUS MATERIALS ($D = 140$ nm AND $\delta = 20$ nm)

	CoPt/Ru	Co/Pd superlattice	FePt
M_s (emu/cm ³)	1180	500	1100
K_u (x10 ⁶ erg/cm ³)	12.4	5	50
H_k (kOe)	21	20	91
$H_k^{\text{eff}}(r=0)/H_k^{\text{eff}}(r=70)$	0.56	0.84	0.93

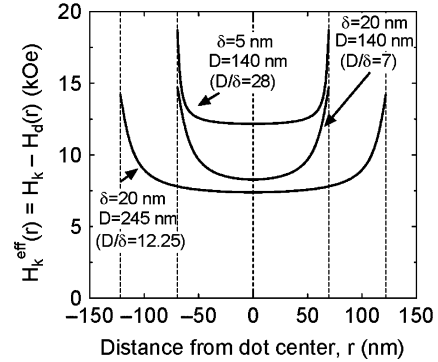


Fig. 3. Values of H_k^{eff} in dots as a function of the distance from the dot center, r , for various CoPt dot having different D and δ . ($M_s = 1180$ emu/cm³)

be noted that the value of H_k^{eff} at the dot center $H_k^{\text{eff}}(r = 0)$ for the dot arrays with $\delta = 5$ nm was larger than the others, in spite of having the largest D/δ ratio (the largest H_d), because of a significant increase in K_u on reducing δ in CoPt/Ru films [1].

Table I shows the value of the ratio of $H_k^{\text{eff}}(r = 0)$ to $H_k^{\text{eff}}(r = 70)$ for CoPt/Ru dot arrays with $D = 140$ nm and $\delta = 20$ nm ($D/\delta = 7$). The values of $H_k^{\text{eff}}(r = 0)/H_k^{\text{eff}}(r = 70)$ calculated for dot arrays made of Co/Pd superlattice film and FePt ordered alloy film are also shown in the table, together with the K_u and M_s values used for the calculation. The value of $H_k^{\text{eff}}(r = 0)/H_k^{\text{eff}}(r = 70)$ for CoPt dot arrays was remarkably smaller than the others, indicating a significant reduction in H_k^{eff} at the dot center, due to a large M_s value (large H_d) relative to H_k .

C. H_r and Thermal Stability as Functions of D and δ

We assumed that the nucleation point for CoPt dot arrays was located around the center of the dot because of the significant reduction in H_k^{eff} at the dot center. Assuming coherent rotation of the magnetization during nucleation, we estimated the values of H_0 and $E_0/k_B T$ of all dot arrays by fitting the values of H_r and H_r^P to the Sharrock equation [9]. H_0 corresponds to the “intrinsic” remanence coercivity obtained by subtracting the effect of thermal agitation on the magnetization, and E_0 is the stabilizing energy barrier to nucleation.

Fig. 4 shows the values of H_r , H_0 and $H_k^{\text{eff}}(r = 0)$ as a function of D for dot arrays with various δ . $H_k^{\text{eff}}(r = 0)$ for thin dot arrays was larger than the others, in spite of having the largest D/δ ratio, as mentioned above. $H_k^{\text{eff}}(r = 0)$ increased slightly as D decreased in all series of dot arrays due to a decrease in H_d . It should be noted that the variation in H_0 was qualitatively

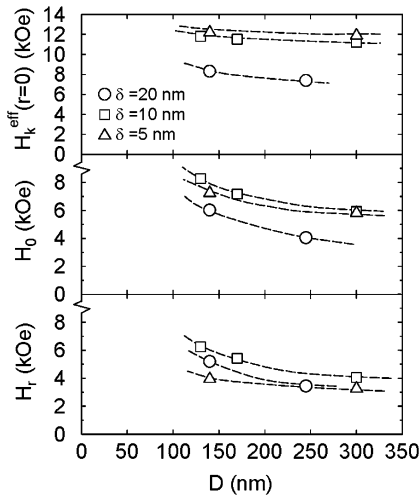


Fig. 4. Values of H_r , H_0 , and $H_k^{\text{eff}}(r = 0)$ as a function of D for the dot arrays with various δ .

in good agreement with that in $H_k^{\text{eff}}(r = 0)$; H_0 for the dot arrays with $\delta = 20$ nm was smaller than that for the thinner dot arrays; moreover, H_0 increased as D decreased in all series of dot arrays. The ratio of H_0 to $H_k^{\text{eff}}(r = 0)$ for the dot arrays with $\delta = 10$ nm increased from 0.53 to 0.70 as D decreased from 300 to 130 nm, and no significant difference in the $H_0/H_k^{\text{eff}}(r = 0)$ ratio due to the difference in δ was observed. A numerical calculation assuming Stoner–Wohlfarth type magnetization reversal [3] revealed that the c -axis distribution ($\Delta\theta_{50} = 2.8^\circ$) should reduce H_0 to be about 70%–78% of $H_k^{\text{eff}}(r = 0)$ in the present D/δ range. The calculated $H_0/H_k^{\text{eff}}(r = 0)$ values were in relatively good agreement with those obtained experimentally for dot arrays with a small D/δ ratio. On the other hand, H_r was slightly smaller than H_0 for the dot arrays with $\delta = 20$ nm, but H_r was reduced to nearly half of H_0 for the dot arrays with $\delta = 5$ nm due to the thermal agitation of magnetization.

Fig. 5 shows the values of $E_0/(k_B T)$ of these dot arrays. (k_B is the Boltzmann constant and T is the absolute temperature.) The values of $E_0/(k_B T)$ of the dot arrays with $\delta = 20$ nm were more than 600, indicating a high thermal stability, but $E_0/(k_B T)$ decreased significantly as δ decreased, resulting in the reduction in the H_r/H_0 ratio mentioned above. $E_0/(k_B T)$ increased as D decreased in thick dot arrays, but $E_0/(k_B T)$ of the dot array with $\delta = 5$ nm was not sensitive to the values of D over the present D range. The value of E_0 is described as the product of the effective anisotropy for nucleation K_u^{eff} and the activated switching (nucleation) volume V_{act} . K_u^{eff} should increase as D decreases because of a reduction in H_d . On the other hand, V_{act} should decrease as δ decreases, since the exchange length of magnetization, $L_{\text{ex}} = 4(A/K_u^{\text{eff}})^{1/2}$, of the dot arrays (about 15 nm) is nearly the same as the δ value. It is likely

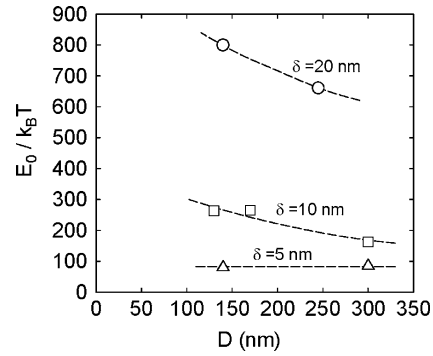


Fig. 5. Values of $E_0/k_B T$ as a function of D for the dot arrays with various δ .

that the reduction in V_{act} on reducing δ is significant, compared to the increase in K_u^{eff} on reducing D , and that $E_0/(k_B T)$ was mainly determined by δ in the present D range.

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