

# Fabrication of $L1_2$ -CrPt<sub>3</sub> alloy films using rapid thermal annealing for planar bit patterned media

T. Kato<sup>1</sup>, *Member, IEEE*, D. Oshima<sup>1</sup>, Y. Yamauchi<sup>2</sup>, S. Iwata<sup>1</sup>, *Member, IEEE*, S. Tsunashima<sup>2</sup>, *Member, IEEE*

<sup>1</sup>Department of Quantum Engineering, Nagoya University, Nagoya, 464-8603, Japan

<sup>2</sup>Department of Electrical Engineering and Computer Science, Nagoya University, Nagoya, 464-8603, Japan

SiO<sub>2</sub> (2 nm) / Cr<sub>25</sub>Pt<sub>75</sub> (15 nm) and SiO<sub>2</sub> (2 nm) / Cr<sub>25</sub>Pt<sub>75</sub> (15 nm) / SiO<sub>2</sub> (10, 20 nm) / Co<sub>80</sub>Zr<sub>10</sub>Nb<sub>10</sub> (10 nm) were prepared by magnetron sputtering method and post-annealed by rapid thermal annealing (RTA) at temperatures of 600 – 1000 °C for 1 – 60 sec. The saturation magnetization  $M_s$  and coercivity  $H_c$  measured by applying a maximum field of 18 kOe were 150 emu/cc and 12 kOe, respectively, for the sample after RTA at 1000 °C for 30 sec. This means that  $L1_2$ -CrPt<sub>3</sub> phase was obtained by RTA process. The RTA process was applied to fabricate the multilayered structure having  $L1_2$ -CrPt<sub>3</sub> and CoZrNb soft magnetic underlayer (SUL). The polar Kerr loop of CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm) after RTA at 1000 °C for 30 sec exhibited a large coercivity  $H_c > 9$  kOe corresponding to that of the CrPt<sub>3</sub> single layer. From the compositional depth profile of CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm) processed by RTA at 1000 °C for 30 sec, sharp layered structure was confirmed despite the high temperature heat treatment. When the SiO<sub>2</sub> thickness was reduced to 10 nm, such a layered structure was completely destroyed due to the interdiffusion between CrPt<sub>3</sub> and CoZrNb layers. Thus it was concluded that the RTA process and the interlayer of SiO<sub>2</sub> (20 nm) were effective to fabricate layered structure having  $L1_2$ -CrPt<sub>3</sub> and SUL.

**Index Terms**—Rapid thermal annealing, Magnetic layered films, Planar bit patterned media, CrPt<sub>3</sub>

## I. INTRODUCTION

BIT patterned media have attracted considerable interest as future high density magnetic recording media, since they provide a promising technology to postpone the problem of superparamagnetic limit, i.e., thermal instability of the recorded bits in the media. One of the problems for the practical use of the bit patterned media is topography of discrete magnetic bits defined by lithographical fabrication, because the rough surface of the disk disturbs stable flying of the hard disk drive (HDD) head. Ion beam irradiation has been proposed as a new approach to pattern magnetic materials locally without etching magnetic materials, i.e., without altering the surface topography [1, 2], and ion irradiation into Co/Pt [1 - 4] and Co/Pd [5, 6] multilayers (MLs) has been reported for the modification of their perpendicular anisotropies. However, in the Co/Pt and Co/Pd MLs patterned by ion irradiation, the adjacent magnetic bits are not magnetically isolated due to the exchange coupling through in-plane magnetized spacing, which will limit the ultimate recording density of the media.

Previously, we have reported the ion-beam patterned medium using a CrPt<sub>3</sub> alloy film [7]. The CrPt<sub>3</sub> shows ferrimagnetism when it has an ordered  $L1_2$  phase, while paramagnetism when disordered fcc phase, and the ferrimagnetic  $L1_2$ -CrPt<sub>3</sub> was found to be transformed to paramagnetic Cr<sub>25</sub>Pt<sub>75</sub> by a quite low ion dose of  $2 \times 10^{14}$  ions/cm<sup>2</sup> with 30 keV Kr<sup>+</sup> ions [8]. The amount of ions  $2 \times 10^{14}$  ions/cm<sup>2</sup> corresponds roughly to number of atoms within 0.1 atomic layer of CrPt<sub>3</sub>, and such a low ion dose has no influence on the surface topology and the thickness of the CrPt<sub>3</sub> layer [9]. The  $L1_2$ -CrPt<sub>3</sub> film onto SiO<sub>2</sub> substrate exhibits a magnetization of 250 emu/cc and a large

perpendicular anisotropy of  $5 \times 10^6$  erg/cc [7, 10 - 13], which is suitable for the high-density bit patterned media. Thus we consider that Kr<sup>+</sup> ion irradiation onto the  $L1_2$ -CrPt<sub>3</sub> film could be a practical approach to achieve planar bit patterned media beyond 1 Tb/in<sup>2</sup> recording density. One of the problems to use CrPt<sub>3</sub> for the practical application is high temperature heat treatment to obtain  $L1_2$  phase CrPt<sub>3</sub>, which makes difficult to fabricate layered structure for perpendicular recording media. In this study, we report rapid thermal annealing (RTA) processing of CrPt<sub>3</sub> and discuss magnetic properties of RTA processed CrPt<sub>3</sub>. Furthermore we report the application of RTA to the fabrication of multilayered structure having  $L1_2$ -CrPt<sub>3</sub> and soft magnetic underlayer (SUL).

## II. EXPERIMENTAL METHOD

[Cr (0.4 nm) / Pt (1.5 – 1.7 nm)]<sub>10</sub> MLs / substrate, SiO<sub>2</sub> (2 nm) / Cr<sub>25</sub>Pt<sub>75</sub> (15 nm) / substrate, and SiO<sub>2</sub> (2 nm) / Cr<sub>25</sub>Pt<sub>75</sub> (15 nm) / SiO<sub>2</sub> (10, 20 nm) / Co<sub>80</sub>Zr<sub>10</sub>Nb<sub>10</sub> (10 nm) / substrate were prepared by magnetron sputtering method. Fused quartz or thermally oxidized Si was used as a substrate. The Cr<sub>25</sub>Pt<sub>75</sub> alloy layer was fabricated by the co-sputtering of Cr and Pt sources. The Co<sub>80</sub>Zr<sub>10</sub>Nb<sub>10</sub> was deposited using an alloy target.  $L1_2$  phase CrPt<sub>3</sub> were obtained by rapid thermal annealing (RTA) at temperatures of 600 – 1000 °C for 1 – 60 sec in N<sub>2</sub> atmosphere. The ramp rate was set to 70 °C/sec. For comparison, some of the samples were annealed in vacuum at a temperature of 850 °C for 15 min. In the vacuum annealing, the cooling rate to room temperature was set at 10 °C/min.

Magnetic properties were measured by using alternating gradient field magnetometer (AGM) and polar Kerr loop tracer using polarized angle modulation method. Crystal structure and surface morphology were checked by X-ray diffraction (XRD) with Cu K $\alpha$  radiation and atomic force microscopy (AFM), respectively. Magnetic domain structure was observed by magnetic force microscope (MFM).

Compositional depth profile was characterized by scanning Auger electron spectrometer (AES) equipped with  $\text{Ar}^+$  ion beam milling system. The energy of the primary electron for AES measurements was 10 keV, and  $\text{Ar}^+$  ion beam energy was 3 kV. The primary electron beam was scanned within the region of  $200 \times 200 \mu\text{m}^2$ , which is adjusted around the center of the ion-etching region of  $1 \text{ mm}\phi$ .

### III. RESULTS AND DISCUSSIONS

Figure 1 shows the RTA process temperature dependence of (a) saturation magnetization  $M_s$  and (b) coercivity  $H_c$  of  $\text{CrPt}_3$  films on  $\text{SiO}_2$ . The annealing time was set at 30 sec. The open and closed circles were obtained by annealing the multilayer sample  $[\text{Cr} (0.4 \text{ nm}) / \text{Pt} (1.51 \text{ nm})]_{10}$  and alloy sample  $\text{Cr}_{25}\text{Pt}_{75}$  (15 nm). Both samples have almost the same chemical composition of Cr 25 at.% and Pt 75 at.%. Upper two figures in Fig. 1 show  $M$ - $H$  loops of  $\text{Cr}_{25}\text{Pt}_{75}$  (15 nm) annealed at  $850^\circ\text{C}$  and  $1000^\circ\text{C}$ . The saturation magnetization  $M_s$  and coercivity  $H_c$  were estimated from  $M$ - $H$  loops taken at a maximum field of 18 kOe. As shown in Fig. 1, the  $M_s$  and  $H_c$  increased with increasing the process temperature. Furthermore, large perpendicular anisotropy just as the vacuum annealed  $\text{CrPt}_3$  in the previous reports [7, 8] is confirmed. The disordered  $\text{Cr}_{25}\text{Pt}_{75}$  alloy and Cr / Pt MLs do not show ferro- or ferri-magnetism, and thus the increase of the magnetization and coercivity are understood by the formation of the ferrimagnetic  $L1_2$  ordered  $\text{CrPt}_3$  phase. The formation of the  $L1_2$ - $\text{CrPt}_3$  was also confirmed by the appearance of the superlattice line of  $L1_2$  phase in X-ray diffraction profiles after RTA (not shown here). The RTA made  $\text{CrPt}_3$  has strong 111 orientation with  $\Delta\theta_{50} = 6$  deg (FWHM of rocking curve for 111 peak). Thus the order parameter  $S$  of RTA made  $\text{CrPt}_3$  at  $950^\circ\text{C}$  was calculated from the in-plane XRD profile as in the same manner as Ref. [8], and estimated to be  $S > 0.9$ .

The saturation magnetization of bulk  $\text{CrPt}_3$  is around 250 emu/cc [10], and the  $\text{CrPt}_3$  (20 nm) on  $\text{SiO}_2$  substrate annealed in vacuum at  $850^\circ\text{C}$  for 15 min exhibits a large coercivity about 12 kOe [7]. The present sample has almost the same coercivity when the sample was processed at around  $1000^\circ\text{C}$ . The  $M_s$  of the sample after RTA at  $1000^\circ\text{C}$  was around 150 emu/cc, which is smaller than those of the bulk  $\text{CrPt}_3$  and the vacuum annealed  $\text{CrPt}_3$  film. The reason of smaller magnetization of the RTA made  $\text{CrPt}_3$  than that of vacuum annealed  $\text{CrPt}_3$  is considered to be larger saturation field of RTA made  $\text{CrPt}_3$  than that of the vacuum annealed one. As for the annealing time dependence, the  $M_s$  of  $\text{CrPt}_3$  processed by RTA  $950^\circ\text{C}$  for 1 sec was only 30 % of the samples processed for 30 sec (not shown in the figure), which means  $950^\circ\text{C}$  and 1 sec RTA is considered to be insufficient for the formation of  $L1_2$ - $\text{CrPt}_3$ . The  $M_s$  of  $\text{CrPt}_3$  obtained by 1 min RTA was almost the same as that by 30 sec RTA.

#### FIG. 1 HERE

Figure 2 shows AFM and MFM images of (a) as-sputtered

$[\text{Cr} (0.4 \text{ nm}) / \text{Pt} (1.51 \text{ nm})]_{10}$  ML, (b)  $\text{CrPt}_3$  (20 nm) made by RTA at  $950^\circ\text{C}$  for 30 sec, and  $\text{CrPt}_3$  (20 nm) annealed in vacuum at  $850^\circ\text{C}$  for 15 min. The  $\text{CrPt}_3$  films are made out of the Cr / Pt MLs. For the as-sputtered ML, quite flat surface having a surface roughness of  $R_a = 0.14 \text{ nm}$  was confirmed, and no contrast in MFM images indicates that the as-deposited ML is non-magnetic state. When the sample was vacuum annealed, large grain with the diameter of  $\sim 100 \text{ nm}$  was confirmed in the AFM image (Fig. 2 (c)). Accompanied by the growth of  $L1_2$ - $\text{CrPt}_3$  grains, the surface roughness was significantly increased to  $R_a = 0.70 \text{ nm}$ . In the MFM image of Fig. 3(c), clear magnetic contrast resulting from the formation of the ferrimagnetic  $L1_2$ - $\text{CrPt}_3$  was seen. The size of the magnetic domain (or magnetic cluster) was almost the same as the crystal grain size. In the case of RTA made  $\text{CrPt}_3$ , small grain size of  $\sim 30 \text{ nm}$  was confirmed as shown in Fig. 2 (b). The surface roughness was  $R_a = 0.53 \text{ nm}$ , which is smooth compared to the vacuum annealed  $\text{CrPt}_3$ , even though it is still rough compared to the as-sputtered ML. Just as the vacuum annealed sample, the RTA made  $\text{CrPt}_3$  exhibited clear magnetic contrast in the MFM image (Fig. 2(b)). The domain size or magnetic cluster size was around 30 nm, which is much smaller than that of vacuum annealed  $\text{CrPt}_3$  and roughly corresponds to crystal grain size.

#### FIG. 2 HERE

Compared to the vacuum annealing, RTA is quite effective to fabricate the recording medium with flat surface. Moreover, it is considered to have a great advantage to make a layered structure having  $L1_2$   $\text{CrPt}_3$  layer for the magnetic recording, since it can control the interdiffusion between  $\text{CrPt}_3$  recording layer and soft under layer (SUL). Figure 3 shows polar Kerr loops of  $\text{CrPt}_3$  (15 nm) /  $\text{SiO}_2$  ( $t$  nm) /  $\text{CoZrNb}$  (10 nm) annealed at  $1000^\circ\text{C}$  for 30 sec. The loops were measured by applying a maximum field of 16 kOe and taken at a wavelength  $\lambda = 800 \text{ nm}$  from the film side ( $\text{CrPt}_3$  side), which means that the loop mainly reflects the hysteresis of the  $\text{CrPt}_3$  layer. As shown in Fig. 3 (a), for the  $\text{SiO}_2$  thickness of  $t = 20 \text{ nm}$ , the sample exhibits a large coercivity  $H_c > 9 \text{ kOe}$  indicating large perpendicular magnetic anisotropy. The  $H_c$  was comparable to the RTA made  $\text{CrPt}_3$  single layer (see Fig. 1 (b)), and the loop shape was consistent with that of the RTA made single layer (not shown here) and that of the vacuum annealed  $\text{CrPt}_3$  [7]. When the  $\text{SiO}_2$  layer was decreased to  $t = 10 \text{ nm}$ , the coercivity of  $\text{CrPt}_3$  was found to reduce significantly as shown in Fig. 3 (b). This is considered to be due to the interdiffusion between the  $\text{CrPt}_3$  layer and  $\text{CoZrNb}$  SUL.

#### FIG. 3 HERE

In order to clarify the interdiffusion between the  $\text{CrPt}_3$  and SUL, AES depth profiles were measured for  $\text{CrPt}_3$  /  $\text{SiO}_2$  / SUL before and after the RTA process. Si(KLL), O(KLL), Cr(LMM), Co(LMM), Pt(MNN) AES were monitored during the interval of  $\text{Ar}^+$  ion beam etching. Figure 4 shows the

Auger peak intensities of the above elements as a function of the etching time. The Auger peak intensity was evaluated from the derivative spectra  $d(E \cdot N(E)) / dE$ . The AES depth profile taken for the as-sputtered CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm) is shown in Fig. 4 (a). The sharp layered structure of CrPt<sub>3</sub> / SiO<sub>2</sub> / SUL was confirmed. After the sample was annealed at 1000 °C for 30 sec, it is confirmed the layered structure still exists as shown in Fig. 4 (b) even though the sharpness of the compositional change at the interface deteriorates slightly. When the SiO<sub>2</sub> thickness was reduced to 10 nm (see Fig. 4 (c)), the layered structure was completely destroyed after RTA of 1000 °C for 30 sec due to the interdiffusion between CrPt<sub>3</sub> layer and CoZrNb SUL. These results are consistent with the Kerr loops shown in Fig. 3. Thus it was concluded that processing by RTA and interlayer of SiO<sub>2</sub> (20 nm) were effective to fabricate layered structure having L1<sub>2</sub>-CrPt<sub>3</sub> layer and SUL.

Finally we have to briefly comment on the magnetic properties of SUL after the RTA. When the as-sputtered CoZrNb (20 nm) was annealed at 950°C for 30 sec, it was confirmed that the  $H_c$  significantly increased from 2 Oe to 25 Oe. In the X-ray diffraction profile, no diffraction peaks indicating the formation of the crystalline phase of CoZrNb even after RTA were detected, however, the increase of  $H_c$  after RTA may indicate a structural difference between as-sputtered and RTA processed SUL. The increase of  $H_c$  will be a problem to achieve a practical recording media having L1<sub>2</sub>-CrPt<sub>3</sub>, and the development of the SUL tolerating RTA process and/or the reduction of the process temperature for making L1<sub>2</sub>-CrPt<sub>3</sub> will be necessary.

#### FIG. 4 HERE

#### IV. CONCLUSION

Rapid thermal annealing process was applied to obtain L1<sub>2</sub> ordered CrPt<sub>3</sub> film with high perpendicular anisotropy, since the RTA process is considered to have a great advantage to make a layered structure having L1<sub>2</sub>-CrPt<sub>3</sub> and soft under layers for the magnetic recording. We confirmed that the L1<sub>2</sub>-CrPt<sub>3</sub> phase was formed by RTA at 950 – 1000 °C for 30 sec. From AFM observations, the RTA processed CrPt<sub>3</sub> was found to have smooth surface compared to the vacuum annealed CrPt<sub>3</sub> due to the suppression of the grain growth in the RTA process. The polar Kerr loop of CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm) annealed at 1000 °C for 30 sec exhibits a large coercivity  $H_c > 9$  kOe. This indicates a large perpendicular magnetic anisotropy of the CrPt<sub>3</sub> layer even after RTA at 1000 °C. From the AES depth profiles of CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (10 and 20 nm) / CoZrNb (10 nm) annealed at 1000 °C for 30 sec, sharp layered structure was confirmed for the SiO<sub>2</sub> (20 nm) sample despite high temperature heat treatment of 1000 °C. When the SiO<sub>2</sub> thickness was reduced to 10 nm, such a layered structure was completely destroyed due to the interdiffusion between CrPt<sub>3</sub> and CoZrNb layers. Thus it was concluded that application of the RTA process and the interlayer of SiO<sub>2</sub> (20 nm) are effective to fabricate layered structure having L1<sub>2</sub>-CrPt<sub>3</sub> and SUL.

#### ACKNOWLEDGMENT

The authors are grateful to Dr. K. Matsumoto, Dr. T. Morikawa, and Dr. K. Ozaki of Fujitsu Laboratories Ltd. for helpful discussions, and would like to thank Mr. M. Kumazawa, and Mr. Y. Adachi of Nagoya University for assistance in the experiments and film composition analysis, respectively.

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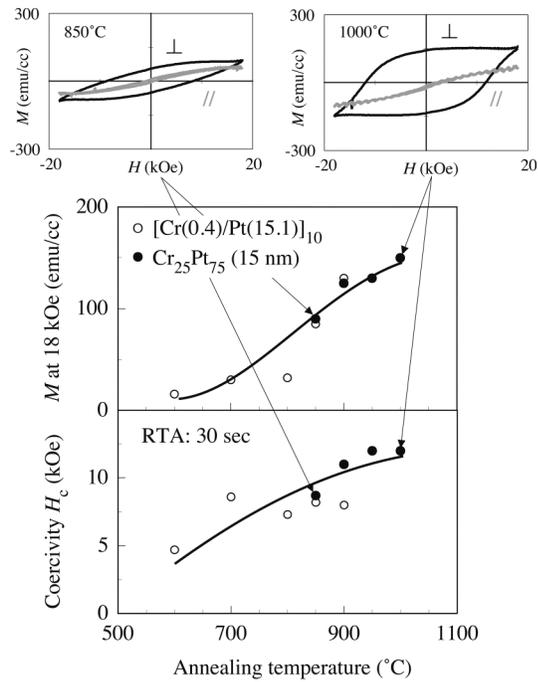


Fig. 1. RTA process temperature dependence of (a) magnetization at 18 kOe and (b) coercivity  $H_c$  of CrPt<sub>3</sub> films. The annealing time was set at 30 sec. The open and closed circles were obtained by annealing the multilayer sample [Cr(0.4 nm) / Pt(15.1 nm)]<sub>10</sub> and alloy sample Cr<sub>25</sub>Pt<sub>75</sub> (15 nm), respectively. Upper figures are  $M$ - $H$  loops of Cr<sub>25</sub>Pt<sub>75</sub> (15 nm) annealed at 850°C and 1000°C.

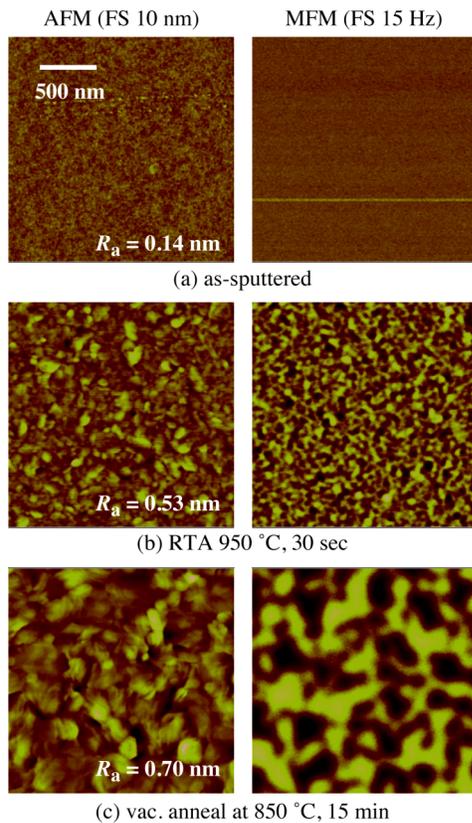


Fig. 2. AFM and MFM images of (a) as-sputtered [Cr(0.4 nm) / Pt(15.1 nm)]<sub>10</sub> ML, (b) CrPt<sub>3</sub> (20 nm) made by RTA 950 °C for 30 sec, and (c) CrPt<sub>3</sub> (20 nm) annealed in vacuum at 850 °C for 15 min. Left are AFM images with a height full scale (FS) of 10 nm, and right are MFM images with a contrast FS of 15 Hz.

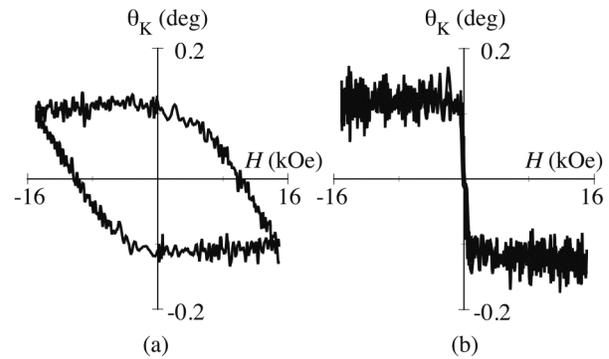


Fig. 3. Polar Kerr loops of CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> ( $t$  nm) / CoZrNb (10 nm) annealed at 1000 °C for 30 sec. The loops were measured by applying a maximum field of 16 kOe, and taken from film side (CrPt<sub>3</sub> side) at  $\lambda = 800$  nm. The SiO<sub>2</sub> thickness was  $t = 20$  nm for (a) and  $t = 10$  nm for (b).

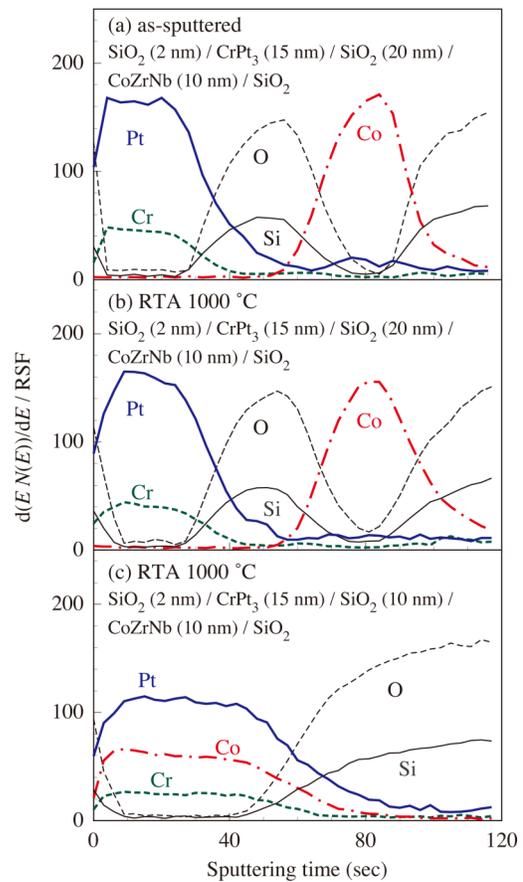


Fig. 4. AES depth profile taken for the (a) as-sputtered CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm), (b) CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (20 nm) / CoZrNb (10 nm) after RTA at 1000 °C for 30 sec, and (c) CrPt<sub>3</sub> (15 nm) / SiO<sub>2</sub> (10 nm) / CoZrNb (10 nm) after RTA at 1000 °C for 30 sec. The Auger peak intensity was evaluated from the derivative spectra  $d(E \cdot N(E)) / dE$  taking into account the relative sensitivity factor (RSF) of each element.