

Supplementary material

Large voltage-induced coercivity change in Pt/Co/CoO/amorphous TiO_x structure and heavy metal insertion effect

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S1 I-V characteristics of heavy metal inserted samples

Figure S1 (a)-(f) shows I-V characteristics of the sample without and with heavy metal insertion. Except W inserted sample, asymmetric I-V characteristics was observed, while the degree differ for each sample. By post annealing, the asymmetry was weakened and the applicable voltage was increased. In contrast, the W inserted sample exhibits a symmetric I-V characteristics with lower resistivity. Fig. S1 (g) summarize the post anneal temperature dependence of the resistivity @ 1 V of samples without and with heavy metal insertion (nominally $t_{\text{Co}} = 1.7$ nm samples). The resistivity increased by post annealing up to ~ 350 °C. After post annealing at higher temperature, the resistivity degraded. A degradation of insulating layers (SiO_2) between bottom and top electrode, which deposited during micro-fabrication process, may partly contribute to the degradation of the resistivity at high temperature.

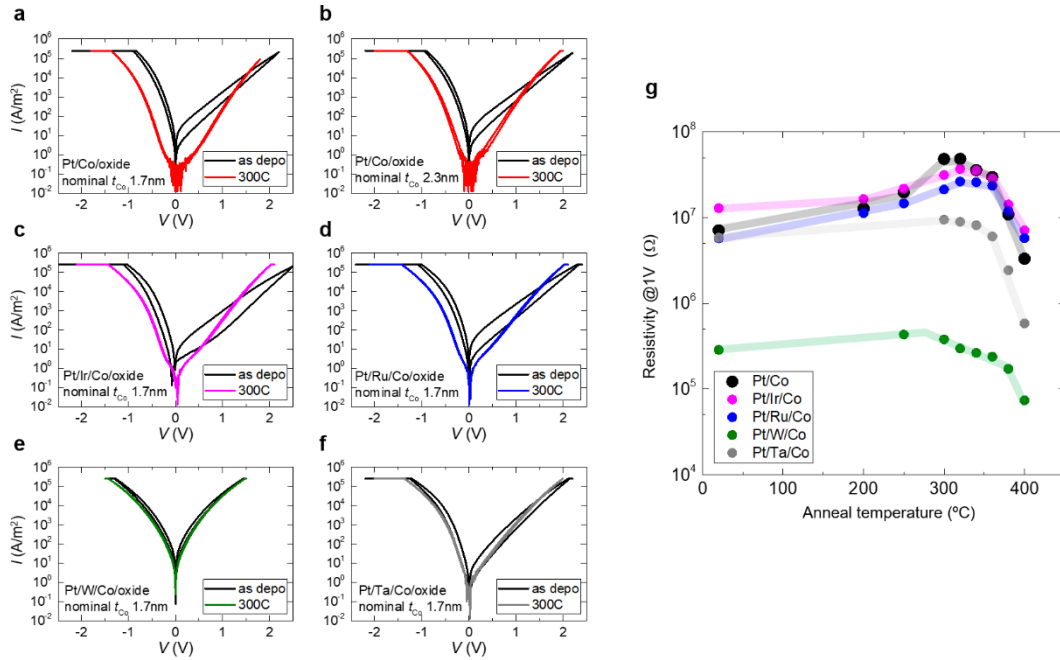


Fig. S1 (a)-(f) I-V characteristics of the sample without and with heavy metal insertion. The black lines and colored lines corresponding to those of as-deposited and post annealed samples. (g) Annealing temperature dependence of the resistivity @ 1 V of samples without and with heavy metal insertion. The nominal film structures are SiO_x sub./Ta (5nm)/Ru (10 nm)/Ta (5 nm)/Pt (10 nm)/heavy metal insertion layer (0.2 nm)/Co (1.7 or 2.3 nm)/ TiO_x (5 nm)/Pt (5 nm).

S2 RHEED patterns of Co in heavy metal inserted samples

Figure S2 shows the RHEED patterns of Co surface (before TiO_x deposition) in heavy metal 0.2 or 1 nm inserted samples. The RHEED patterns of Co in heavy metal 0.2 nm samples looks identical to that in Fig. 1 (b), confirming (111) texture orientation of Co for all samples. On the other hand, an obvious degrade of the streaky RHEED patterns was observed for Ta 1 nm inserted samples. These results indicate that the effect of 0.2 nm-thick insertion layers to the crystal orientation of Co were small, regardless the preferable crystal structure of the inserted heavy metals.

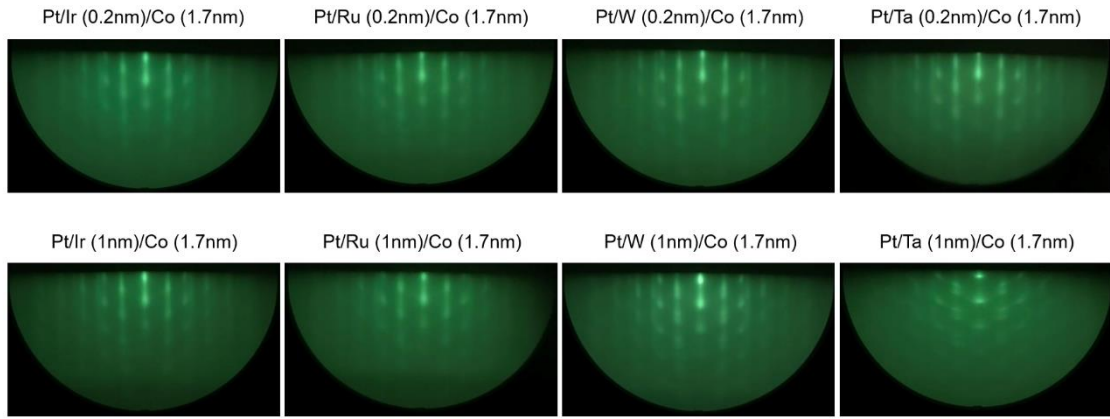


Fig. S2 RHEED patterns of Co surface in heavy metal 0.2 nm inserted structures (upper images) and 1 nm inserted structures (lower images). The nominal film structures are SiO_x sub./Ta (5nm)/Ru (10 nm)/Ta (5 nm)/Pt (10 nm)/heavy metal insertion layer (0.2 or 1 nm)/Co (1.7 nm).

S3 dielectric constant of bilayer dielectric layers

An averaged dielectric constant of bilayer dielectric layers are represented by the following equation;

$$\text{averaged } \varepsilon_r = \left(\frac{1}{\varepsilon_{r1}} \frac{d_1}{d_1 + d_2} + \frac{1}{\varepsilon_{r2}} \frac{d_2}{d_1 + d_2} \right)^{-1} \quad (\text{S1})$$

Here, $\varepsilon_{r1}(\varepsilon_{r2})$ and $d_1(d_2)$ represent relative dielectric constant and thickness of dielectric layer 1 (2), respectively. Fig. S3 shows d_2 dependence of averaged ε_r of bilayer dielectric layers, assuming $\varepsilon_{r1} = 50$, $\varepsilon_{r2} = 10$, $d_1 = 5$ nm. We suppose dielectric layer 1 and 2 as a high-k TiO_x barrier and interfacial CoO layer, respectively. Fig. 3 shows a large influence of thin interfacial CoO layer on the average ε_r . Assuming 1.2 nm CoO layer at the interface, the ε_r of TiO_x layer is estimated to be ~ 50 .

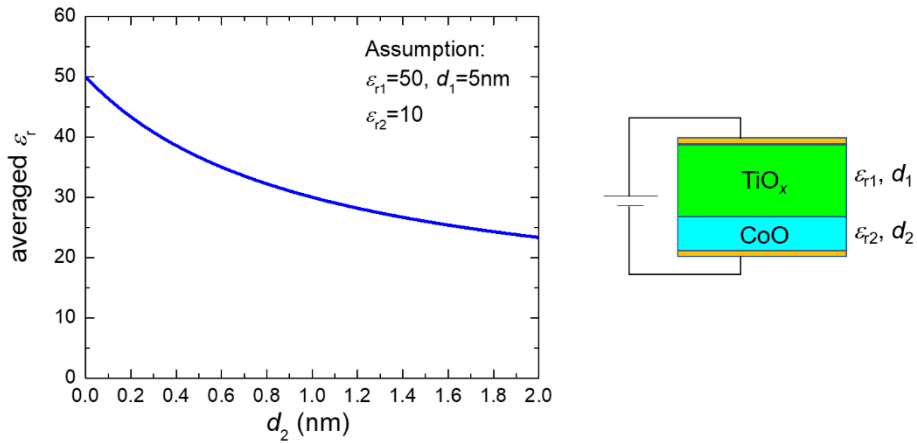


Fig. S3 d_2 dependence of averaged ε_r of bilayer dielectrics calculated by eq. (S1), assuming $\varepsilon_{r1} = 50$, $\varepsilon_{r2} = 10$, $d_1 = 5$ nm.