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Quantitative determination of anisotropic magnetoelectric coupling in BiFeO$_3$–CoFe$_2$O$_4$ nanostructures


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The transverse and longitudinal magnetoelectric susceptibilities (MES) were quantitatively determined for (001) heteroepitaxial BiFeO$_3$–CoFe$_2$O$_4$ nanostructures. Both of these MES values were sharply enhanced at magnetic fields below 6 kOe and revealed asymmetric line shapes with respect to the dc magnetic field, demonstrating the strain-induced magnetoelectric effect. The maximum transverse MES, which reached as high as $-60$ mV/cm Oe, was about five times larger than the longitudinal MES. This observation signifies that transverse magnetostriction of the CoFe$_2$O$_4$ nanopillars is enhanced more than the bulk value due to preferred magnetic domain alignment along the [001] direction coming from compressive, heteroepitaxial strain. © 2010 American Institute of Physics. [doi:10.1063/1.3475420]

The magnetoelectric (ME) effect is a physical phenomenon in which the electric polarization $P$ (magnetization $M$) is modulated by the magnetic field $H$ (electric field $E$). There is a growing interest in the application of ME effects toward various devices, including magnetic sensors and energy harvesters. As such, numerous efforts have been made to obtain strong ME couplings in ME composites made of ferroelectric (or piezoelectric) and ferromagnetic materials. In these types of ME composites, $P$ is varied with $M$ via the strain ($u$)-coupling at the interface between the piezoelectric and magnetostrictive phases. Thus, the configuration of this interface is a significant control parameter for determining the extent of ME coupling. In this respect, a layered sandwich structure [i.e., (2–2) structures] produces a larger ME coupling than for particulates that are dispersed in a matrix [i.e., (0–3) structures] because the former has a larger interface area. In contrast, this common practice cannot be applied to multilayered thin films, where clamping of the nonmagnetic substrate can prevent strain-coupling between the layers.

In 2004, an epitaxial thin film composed of CoFe$_2$O$_4$ (CFO) nanopillars embedded in a BaTiO$_3$ (BTO) matrix [i.e., (1–3) structures] was grown and suggested to be an alternative to circumvent the substrate clamping effect. On the other hand, it has been quite difficult to determine quantitatively the ME coupling of such nanostructures and, more generally, numerous ME films, except observing the existence of non-trivial ME coupling. One dominant reason for this difficulty is that the ME voltage signal is proportional to the film thickness; it typically becomes smaller than 1 $\mu$V for film thicknesses less than 1 $\mu$m. To overcome this difficulty, a large ac magnetic field ($H_{ac}$) of up to $\sim 1$ kOe was recently used to obtain the effective MES as a function of $H_{ac}$. However, direct measurements of the MES based on a conventional scheme that employs small $H_{ac}$ values with variations in the dc magnetic field ($H_{dc}$) would be useful for understanding the ME coupling of numerous multiferroic films or nanostructures at the quantitative level. In this work, by use of the conventional scheme, we provide experimental evidences that a 300 nm thick BiFeO$_3$–CoFe$_2$O$_4$ (BFO–CFO) nanostructure has a peculiar MES anisotropy that is not expected in bulk forms of those materials.

The self-assembled epitaxial BFO–CFO film with thickness of 300 nm was grown on a (001) SrRuO$_3$/SrTiO$_3$ substrate by pulsed laser deposition. The film had the CFO nanopillars embedded in a BFO matrix with a volume fraction of 1:1. Top electrodes Pt/SrRuO$_3$ were deposited on the film surface [Fig. 1(a)]. For magnetic hysteresis measurements, a vibrating sample magnetometer was utilized. For ferroelectric hysteresis loops, the displacement current was measured using a fast digitizer and a high voltage amplifier.

To investigate the MES ($\alpha = \delta P/\delta H_{ac}$), especially for thin films, we developed a highly sensitive ME susceptometer that operates inside the PPMS (Quantum Design). A pair of solenoids was designed to induce a $H_{ac}$ of $\sim 4$ Oe inside the solenoid pair and a voltage pick-up coil was used to determine the phase of $H_{ac}$. In particular, modulated charges, instead of voltages, were measured using a high-impedance

![FIG. 1. (Color online) (a) Schematic picture, (b) AFM image, and (c) XRD pattern of the BFO–CFO film with nanopillar structures.](http://www.aip.org/about/aip/pressroom/press.released_images/2010/052902-1.png)
charge amplifier with a gain factor of $10^{12}$ V/C. This makes the amplified signal independent of film thickness but proportional to electrode area so that the signal-to-noise ratio was improved. Based on this scheme, we were able to detect small ME changes in a thin film with a thickness $\sim 40$ nm and a circular electrode of diameter $\sim 100$ µm. The lowest charge noise ($\Delta Q$) was $10^{-17}$ C, which corresponded to a voltage noise of $\Delta V \sim 10^{-9}$ V.

Moreover, there existed a large uniaxial magnetic anisotropy field of $25$ kOe. Therefore, even in the BFO–CFO film studied here, the magnetic anisotropy field of $25$ kOe seems to originate from a compressive, heteroepitaxial strain of $0.33\%$ inside the CFO nanopillars. This experimental result reflects that the presence of a compressive, heteroepitaxial strain of $0.33\%$ inside the CFO nanopillars was estimated as $2.0867$ Å from the crystal, the transverse magnetostriction $\lambda_{31}$ decreases under $H_{dc} = 0$. This is in good agreement with the reported CFO nanopillars. The saturated magnetic anisotropy field of $25$ kOe was obtained for $H_{dc} = 0$. When the $H_{dc} = 0$ was applied to the CFO crystal, the transverse magnetostriction $\lambda_{31}$ was positive [dashed lines in Fig. 3(c)]. Thus, the CFO nanopillars and the BFO matrix, via strain-coupling, were expected to be elongated along the $[001]$ direction. It is known from an earlier study that $P_3$ increases due to the rotation of $P$ with increasing length along the $[001]$ direction. Therefore, the observation of $\alpha_{31} > 0$ for $H_{dc} > 0$ can be qualitatively understood.

As shown in Fig. 3, $\alpha_{31} < 0$ for $H_{dc} > 0$ while $\alpha_{31} > 0$ for $H_{dc} > 0$. This experimental result reflects that $P_3$ increases under $H_{dc} = 0$, while $P_3$ decreases under $H_{dc} = 0$. When the $H_{dc} = 0$ was applied to the CFO crystal, the transverse magnetostriction $\lambda_{31}$ was positive [dashed lines in Fig. 3(c)]. Thus, the CFO nanopillars and the BFO matrix, via strain-coupling, were expected to be elongated along the $[001]$ direction. It is known from an earlier study that $P_3$ increases due to the rotation of $P$ with increasing length along the $[001]$ direction. Therefore, the observation of $\alpha_{31} > 0$ for $H_{dc} > 0$ can be qualitatively understood.
stood as the result of an elongation of CFO/BFO along the [001] direction under \( H_{dc} \parallel [100] \) and the subsequent increase in \( P_3 \). The case of decreasing \( P_3 \) under \( H_{dc} \parallel [001] \) can also be understood in a similar way because the longitudinal magnetostriction (\( \alpha_{33} \)) in the CFO crystal was negative.

However, our MES data are seemingly inconsistent with the magnetostriction behavior of a bulk CFO at the quantitative level. The maximum to minimum value of \( \alpha_{33}(\Delta \alpha_{33}) \) in Fig. 3(a), amounting to \( \sim 260 \) ps/m (\( \sim 120 \) mV/cm Oe), is about five times larger than that of \( \alpha_{33}(\Delta \alpha_{33}) \). According to the magnetostriction of a \( \text{Co}_{0.8}\text{Fe}_{2.2}\text{O}_4 \) crystal with demagnetized domains [Fig. 3(e)], a slope of the \( \lambda_{33} \) versus \( H \) curve is at least twice that of the \( \lambda_{33} \) versus \( H \) curve. Upon assuming that the magnetostriction of CFO nanopillars follows a bulk behavior, these magnetostriction data predict that \( \Delta \alpha_{33} \) should be at least twice of \( \Delta \alpha_{31} \), which is in sharp contrast with the results in Figs. 3(a) and 3(b).

Although there might exist several mechanisms to induce enhanced \( \alpha_{31} \) as discussed in a recent anisotropic MES study using large \( H_{dc} \), one decisive factor could be the preferred magnetic domains existing in the CFO nanopillars. The solid lines in Fig. 3(c) reproduce published \( \lambda_{31} \) and \( \lambda_{33} \) for a \( \text{Co}_{0.8}\text{Fe}_{2.2}\text{O}_4 \) crystal with a single magnetic domain along the [001] direction. The single magnetic domain was obtained through the magnetic annealing process, i.e., cooling under \( H \) from high to room temperature. In this situation, applied \( H \parallel [001] \) gave rise to the 180° domain wall motion, which resulted in negligible \( \lambda_{31} \). In contrast, \( H \parallel [100] \) resulted in a 90° domain wall motion so that it produced quite large \( \lambda_{31} \). As we discussed above, the compressive, heteroepitaxial strain was a main source of enhanced magnetic anisotropy along the [001] direction in the CFO nanopillars. It is thus likely that the magnetic domains inside the CFO nanopillars have preferred alignment along the [001] direction, as illustrated in Fig. 3(f). If so, similar to the case of single magnetic domains, the CFO nanopillars are expected to have enhanced \( \alpha_{31} \) and suppressed \( \alpha_{33} \). As a result, as observed in Figs. 3(a) and 3(b), the BFO–CFO nanostructure will give rise to a bigger \( \alpha_{31} \) (smaller \( \alpha_{33} \)) than that expected based on the behavior of bulk CFO magnetostriction.

These results point to the possibility that the strain-induced ME coupling in the nanostructured film can be quite different from the macroscopic bulk composite. Application of a compressive, heteroepitaxial strain to the CFO nanopillars enables to achieve increased \( \alpha_{31} \) as high as \( \sim 130 \) ps/m (\( \sim 60 \) mV/cm Oe) at 6 kOe. Upon increasing \( \mu_{001} \), as done in the BTO–CFO nanostructures with growth temperatures, \( \alpha_{31} \) can be further optimized. In comparison, our previous study on a thin film made of \( \text{NiFe}_2\text{O}_4 \) nanoparticulates embedded in a \( \text{PbZr}_{0.2}\text{Ti}_{0.8}\text{O}_3 \) matrix, [i.e., the (0–3) structure] showed maximum \( \alpha_{31} \) \( = 4 \) mV/cm Oe (\( \sim 4 \) ps/m) and \( \alpha_{33} \) \( = 16 \) mV/cm Oe (\( \sim 14 \) ps/m), which was clearly smaller than the maximum \( \alpha_{31} \) \( \sim 130 \) ps/m found here. Therefore, our results strongly support that thin films with the (1–3) nanostructure have larger ME couplings than the other nanostructures [e.g., (0–3) structure].

In conclusion, we have determined the anisotropic MES of a 300 nm thick \( \text{BiFeO}_3\)–\( \text{CoFe}_2\text{O}_4 \) nanostructure. An enhancement was observed in the transverse configuration, which can be explained by the preferred alignment of magnetic domain resulting from the heteropitaxial strain that is unique to the present (1–3) nanostructure. This investigation offers quantitative evidences that the nanoscale engineering of strain coupling is useful for the design of ME devices.

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