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Citation: Journal of Applied Physics 117, 17D711 (2015); doi: 10.1063/1.4913943
View online: http://dx.doi.org/10.1063/1.4913943
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Magnetoelectric polymer nanocomposite for flexible electronics

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(Received 5 November 2014; accepted 30 October 2014; published online 6 March 2015)

This paper reports the fabrication and characterization of a new type of magnetoelectric polymer nanocomposite that exhibits excellent ferromagnetism and ferroelectricity simultaneously at room temperature. The multiferroic nanocomposite consists of high aspect ratio ferromagnetic iron nanowires embedded inside a ferroelectric co-polymer poly(vinylidene fluoride-trifluoroethylene), P(VDF-TrFE). The nanocomposite has been fabricated via a simple low temperature spin coating technique. Structural, ferromagnetic, ferroelectric, and magnetoelectric properties of the developed nanocomposite have been characterized. The nanocomposite films showed isotropic magnetic properties due to the random orientation of the iron nanowires inside the film. In addition, the embedded nanowires did not hinder the ferroelectric phase development of the nanocomposite. The developed nanocomposite showed a high magnetoelectric coupling response of 156 mV/cmOe measured at 3.1 kOe DC bias field. This value is among the highest reported magnetoelectric coupling in two phase particulate polymer nanocomposites. © 2015 AIP Publishing LLC.

[http://dx.doi.org/10.1063/1.4913943]

I. INTRODUCTION

Multiferroic nanocomposites are of great interest for fundamental and technological aspects as they enable new functionalities and concepts for applications in sensors, actuators, data storage, or micro energy harvesting. The coupling between the constituent ferroic orders of these nanocomposite structures offers multifunctional properties and enhanced functionalities that are absent in each of the individual phases. Specifically, the coupling between ferroelectricity and ferromagnetism, the magnetoelectric effect (ME), brings about new potential applications in magnetoelectric sensors, energy harvesters, novel information storage, or smart ME devices. Composite ME materials show a spontaneous electrical polarization when subjected to a magnetic field, and an induced magnetization when subjected to an electrical field. Unlike ceramic-based ME composites, polymer-based multiferroic ME nanocomposites provide the advantages of being flexible, versatile, lightweight, low cost, able to conform to complicated shapes, and they typically involve a low-temperature fabrication process. These unique characteristics make polymer nanocomposites attractive for applications like flexible or wearable devices. Depending on the interface between the ferromagnetic and the polymer matrix, three main types of multiferroic polymer nanocomposites can be found in literature, particulate nanocomposite, laminate composites, and polymer as a binder composite. The first type received the least attention among researchers.1–4 Martins et al. reported incorporation of ferrite nanoparticles (35–55 nm) of CoFe2O4 or Ni0.5Zn0.5Fe2O4 into P(VDF-TrFE) ferroelectric polymer matrix. In both cases, the nanocomposites exhibited improved ferroelectric, ferromagnetic, and ME properties, which were strongly dependent on the ferrite filler concentrations. The highest ME response of Martins’ reported nanocomposites was 41 mV/cmOe with 72 wt. % of CoFe2O4 at 2.5 kOe DC bias magnetic field.3,4 In this kind of multiferroic nanocomposites, the effective coupling properties are strongly dependent on the characteristics of the interface between the polymer and the ferromagnetic fillers. Surface interfacial area to volume ratio is a key factor and therefore high aspect ratio ferromagnetic nanostructures are preferred over spherical nanoparticles. This paper describes the development of a new type of multiferroic nanocomposite consisting of a ferroelectric copolymer poly(vinylidene fluoride-trifluoroethylene), P(VDF-TrFE), and high aspect ratio ferromagnetic iron (Fe) nanowires (NWs) embedded inside the co-polymer. When subjected to a magnetic field, the strain transfer from the Fe NWs to the P(VDF-TrFE) matrix is maximized, owing to the large surface interfacial area of NWs with the polymer. This phenomena in the developed nanocomposite promises a strongly improved ME coupling when compared to low aspect ratio nanoparticles-based nanocomposites.

II. EXPERIMENTAL PROCEDURES

A. Fabrication of ferromagnetic Fe NWs

Anodic aluminum oxide (AAO) membranes offer self-assembled highly ordered hexagonal nanopores, which are ideal templates for homogenous nanoscale materials fabrication. Owing to their easy preparation, controllable pore size and spacing, as well as their low cost fabrication, nanowires, and nanotubes of polymers, metals, semiconductors, and composite materials have been fabricated successfully.5–8 In this work, AAO membranes were prepared by a two-step anodization process using oxalic acid.9 First, highly pure Al discs with 500 μm in thickness were electro polished for

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B. Fabrication of multiferroic nanocomposite

P(VDF-TrFE), with 70/30% in molar ratio of VDF to TrFE, was dissolved in dimethylformamide (DMF) to form an 8 wt.% solution. Fe NWs were dispersed in the same DMF solvent. The NWs were then added to the P(VDF-TrFE) solution to get the polymer nanocomposite solution. The formed nanocomposite mixture had 1.5 vol.% Fe content. The solution was mixed thoroughly using a shaker followed by ultra-sonication for 10 min for dispersion and minimum NWs agglomeration. Thin film samples of the nanocomposite were obtained by spin coating at 1500 rpm for 30 s on Si substrates with patterned Pt bottom electrodes obtained by standard lithography and lift off processes. After spinning, the nanocomposite film was baked at 80°C for 30 min for solvent evaporation. The ferroelectric β phase of the polymer was facilitated by subsequent annealing in vacuum at 135°C for 2 h. Then the films were allowed to gradually cool down to room temperature under vacuum. The thickness of the films was 950 nm measured using a stylus profiler (Dektak). For electrical and ME characterization, a 100 nm thick Au top electrode was thermally evaporated inside the AAO template measured parallel, and perpendicular to the NW’s axis. Clearly, the curve indicates an easy magnetization direction along the length of the wires, which is caused by the shape anisotropy of the high aspect ratio Fe wires. On the other side, the magnetization curve for the nanocomposite does not show field direction dependence (Fig. 2(b)), indicating that the Fe NWs are randomly oriented and well dispersed inside the ferroelectric polymer, and therefore the film is magnetically isotropic.

B. Ferroelectric and magnetoelectric characterization

To measure the ferroelectric polarization versus the electric field (P-E) of the developed nanocomposite, a precision multiferroic tester (Radiant Technologies) was used. The hysteresis loops were measured at room temperature with a maximum electric field of 125 MV/m. By applying triangular waves oscillating at 10 Hz of frequency, well-defined ferroelectric loops were obtained for the fabricated nanocomposite sample (Fig. 3). When compared to pristine P(VDF-TrFE) films, the measured ferroelectric loops indicate that the development of the ferroelectric β phase is not hindered by the addition of the ferromagnetic Fe NWs.

Due to the dimensions of the NWs compared to the thickness of the film, all NWs are lying in-plane with random orientations as also confirmed by the magnetization curve (Fig. 2(b)). Therefore, the ME effect was characterized by measuring the longitudinal ME voltage coefficient $\varepsilon_{ME}$ using the dynamic lock-in amplifier method. A pair of Helmholtz coils was used to generate an AC magnetic field with amplitude of 3 Oe and frequency of 300 Hz that is superimposed to a DC bias field driven by an electromagnet. Both fields are applied in-plane of the nanocomposite film and the generated small signal voltage across the sample thickness is measured.
using a digital Lock-in amplifier (Stanford Research SR850). The ME voltage coefficient \( \alpha_{ME} \) is calculated from the measured voltage using the equation \( \alpha_{ME} \text{ (mV/cmOe)} = \frac{V_{out}}{h_m t_p} \), where \( V_{out} \) is the measured output voltage, \( h_m \) is the amplitude of the AC magnetic field, and \( t_p \) is the thickness of the nanocomposite film.\(^6\)

Prior to ME characterization, the nanocomposite was electrically poled by applying a maximum electric field of 125 MV/m. The ME voltage coefficient of the nanocomposite film, shown in Fig. 4, was measured by sweeping the DC field from 0 Oe to 3.5 kOe. A high ME response of 156 mV/cmOe was measured under 3.1 kOe DC bias field. This value is considered among the highest reported for two phase particulate polymer nanocomposites,\(^1,3\) and it is attributed to the large interfacial area between the NWs and the polymer. The ME response curve also shows that the ME coupling is driven by a strain mediated effect and is mainly due to strain transfer caused by the non-linear magnetostriction effect of the Fe NWs. The drop in the ME curve afterwards is attributed to the saturation of the Fe NWs, where no further elongation or stress is present. The initial fluctuations in the ME voltage coefficient response is mainly due to the nature of the nanocomposite in which the Fe NWs are randomly aligned inside the ferroelectric polymer. Therefore, NWs, which are not in the same direction as the applied DC magnetic field cause torques and stresses depending on their distributions and angles with respect to the applied field. We expect this kind of initial fluctuations to be mitigated, when the NWs are aligned in a preferred direction while fabricating the nanocomposite.

IV. CONCLUSION

A multiferroic polymer nanocomposite was successfully fabricated on a platinized Si substrate using P(VDF-TrFE) as the ferroelectric phase, and high aspect ratio Fe NWs as the ferromagnetic phase. Results showed that the addition of NWs to the P(VDF-TrFE) does not have a significant influence on the ferroelectric properties of the developed nanocomposite, and the nanocomposite exhibits both ferroelectric

FIG. 2. Magnetic moment (m) as a function of applied field (H) of the (a) Fe NWs inside the AAO membrane and (b) the P(VDF-TrFE)/Fe NWs nanocomposite.

FIG. 3. Ferroelectric polarization (P) over electric field (E) measurements for the developed nanocomposite compared to a pristine P(VDF-TrFE).

FIG. 4. ME voltage coefficient versus DC magnetic field bias of the nanocomposite.
and ferromagnetic properties at room temperature. The use of high aspect ratio Fe NWs as ferromagnetic filler resulted in a ME coefficient up to 156 mV/cmOe, which is higher than what has been achieved using ferrite nanoparticles. The enhancement is attributed to a higher interfacial area between the constituent phases. The developed nanocomposite is a promising candidate for flexible and smart ME applications.

ACKNOWLEDGMENTS

Research reported in this publication was supported by King Abdullah University of Science and Technology (KAUST). The authors also would like to acknowledge the financial support provided by SABIC.