Temperature induced Spin Switching in SmFeO₃ Single Crystal
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The prospect of controlling the magnetization (M) of a material is of great importance from the viewpoints of fundamental physics and future applications of emerging spintronics. A class of rare-earth orthoferrites RFeO₃ (R is rare-earth element) materials exhibit striking physical properties of spin switching and magnetization reversal induced by temperature and/or applied magnetic field. Furthermore, due to the novel magnetic, magneto-optic and multiferroic properties etc., RFeO₃ materials are attracting more and more interests in recent years. We have prepared and investigated a prototype of RFeO₃ materials, namely SmFeO₃ single-crystal. And we report magnetic measurements upon both field cooling (FC) and zero-field cooling (ZFC) of the sample, as a function of temperature and applied magnetic field. The central findings of this study include that the magnetization of single-crystal SmFeO₃ can be switched by temperature, and tuning the magnitude of applied magnetic field allows us to realize such spin switching even at room temperature.

The emergent spintronics calls for novel materials with unprecedented controls of magnetism. Rare-earth orthoferrites RFeO₃ (R is rare-earth element) materials are excellent examples to show temperature and/or magnetic field induced spin switching and magnetization reversal, intriguing magnetic and multiferroic properties. RFeO₃ possesses two magnetic sublattices, from 4f-electrons of the rare-earth ions (R-sublattice) and 3d-electrons of the iron ions (Fe-sublattice), respectively. At relatively higher temperatures, a weak ferromagnetism (FM) originates from the canted antiferromagnetism (AFM) in the Fe-sublattice, while the R-sublattice only orders antiferromagnetically at much lower temperatures. Moreover, there exists a delicate exchange interplay between R 4f- and Fe 3d-electrons in these complex oxides. As a result, interesting magnetic behaviors are strongly dependent on several external stimulations, such as applied magnetic field, temperature, and pressure etc.

Among many family members of RFeO₃, SmFeO₃ has been shown to have excellent device characteristics such as the fast magnetic switching, and an easy axis rotation transition (also known as spin reorientation transition) from c-axis to a-axis which occurs at quite high temperature between $T_{SR1}=450$ K and $T_{SR2}=480$ K (see Figure 1). This is the highest spin reorientation transition temperature of the whole RFeO₃ family, which indeed deserves special attention for practical uses. At the same time, the interplay between Sm 4f- and Fe 3d-electrons is also intriguing. For example, below 4 K, SmFeO₃ exhibits an interesting phenomenon of spontaneous magnetization reversal. This reversal is attributed to the activation of the long range ordering of Sm$^{3+}$ spins whose total magnetic moment is antiparallel to the weak ferromagnetic moment of canted AFM ordering of Fe-sublattice. Moreover, the onset of strong competition interplay between Sm 4f- and Fe 3d-electrons can be observed to occur at a relatively high temperature of 140 K, below which the total magnetization is monotonically suppressed and dramatically reversed sign at low temperature.

We first look at the notable feature of single crystal SmFeO₃ well above room temperature. The magnetic anisotropy dependence of temperature can be obtained by applying external magnetic fields along a- and c-axes, i.e. $H/a$ and $H/c$ to measure magnetizations $M_a$ and $M_c$, as illustrated in Figure 1. The magnitude of the applied magnetic field is $H=300$ Oe. Below its Néel temperature $T_N=680$ K, SmFeO₃ becomes a canted antiferromagnet with an FM vector from Fe-sublattice along c-axis as indicated by the red curve. The marked region in a dashed rectangle near 450–480 K exhibits the spin reorientation transition of the Fe-sublattice, where $M_c-T$ and $M_a-T$ show crossover behavior with exchanged magnetization magnitude. This continuous transition (so-called $F_{G}(G_x,F_y)$ to $G_y(F_x,F_z)$ magnetic configuration, i.e. canted G-type AFM and resultant weak FM along a, c directions) is in good agreement with that reported in Refs. [14,18,20] for SmFeO₃ single crystal.

We further discover here, below a certain critical point near room temperature, $M_c-T$ curves after FC and ZFC may exhibit interesting magnetic behaviors of opposite signs (green and black curves in Figure 1). At such critical temperature ($T_{SR1}=278.5$ K for $H=300$ Oe), we observe a very sharp magnetization jump that clearly demon-
strates to a spin switching transition. For both FC and ZFC, a compensation temperature ($T_{\text{comp}} = 3.9$ K for $H = 300$ Oe) corresponding to zero magnetization is obtained. Note that near $T_{\text{comp}}$, $M_s-T$ dramatically decreases in magnitude and changes its sign, whereas $M_s$ curve remains essentially a zero magnetization below $T_{\text{SR1}}$. In the FC case, $M_s-T$ shows a less temperature-dependent form in a large temperature range of 100–350 K, and two strongly temperature-sensitive regimes at the compensation and spin-reorientation temperatures. This measured behavior of $M_s-T$ after the FC is also consistent with Refs. [7,14,20] when $H = 100$ or 500 Oe, but the more interesting $M_s-T$ curve measured after the ZFC has never been reported.

Let us now focus on the $M_s-T$ measured after the ZFC along a-axis of single-crystal SmFeO$_3$ as shown in Figure 2, under an applied magnetic field of $H = 300$ Oe. When we increase the temperature from 3 K to 350 K, the total magnetization $M_s$ falls rapidly and changes sign by crossing a full annihilation (zero magnetization) at 3.9 K, and then achieves a relatively saturate negative magnetization above 100 K. Note that a similar compensation behavior was also reported in other RFeO$_3$ (R= Nd, Er)$_{22–24}$ and in RMnO$_3$ (R= Nd, Sm)$_{22–24}$ systems. As consistent with what is advocated in the literature, we interpret such feature as indicative that the resulting magnetization associated with the FM vector of Sm-sublattice considerably decreases in size (with respect to that of Fe-sublattice) when the temperature increases (see the red and blue arrows illustrating the relative directions and magnitudes of FM vectors of Sm- and Fe-sublattices). At $T_{\text{comp}} = 3.9$ K, the FM vector of Sm-sublattice is equal in magnitude to that of Fe-sublattice but reversed in direction (AFM-type coupled with each other), producing in a vanishing total magnetization. For temperatures above $T_{\text{comp}} = 3.9$ K and below $T_{\text{sw}} = 278.5$ K, the FM vector of Sm-sublattice becomes smaller than that of Fe-sublattice (and still reversed in direction), therefore explaining why the total net magnetization is negative. After a fast decrease, the total magnetization reaches its relative saturation in the temperature region of 100–278.5 K, since the magnetic ordering of the Sm-sublattice is weakened by temperature much faster than that of the Fe-sublattice.

Figure 2 also highlights the main finding of this work, namely a remarkable first-order transition at $T_{\text{sw}} = 278.5$ K. The magnetization undergoes a sudden jump with another sign change from negative to positive value, showing a mirror-like symmetry with respect to zero magnetization at transition point. We note that the observed switching with temperature is not reversible, since our ZFC curve is obtained in a warming up process. This distinctive spin switching effect should thus be associated with a spontaneous spin-flip transition of Sm- and Fe-sublattices at the same time, accompanied by an exchange of their FM vector directions. This is in analogy with what we observed in NdFeO$_3$ near 29 K when $H = 100$ Oe. However, the negative $M_s$ exists in a much wider temperature range from $T_{\text{comp}} = 3.9$ K to near room temperature of $T_{\text{sw}} = 278.5$ K in SmFeO$_3$ system, whereas in NdFeO$_3$ it only happens in a narrow temperature region from 7.6 K to 29 K. This indicates that the unyielding feature of Sm-sublattice that keeps its magnetization parallel to the field $H$ and antiparallel to magnetization of Fe-sublattice up to a high temperature (even at $T_{\text{sw}} = 356$ K for $H = 250$ Oe as shown below). This striking result reveals an evidence for the existence of strong interaction between Sm-4f and Fe-3d electrons in SmFeO$_3$ system. And we speculate that the Sm-sublattice may have a long-range ordering to some extent even near room temperature.

In order to study the relationship between the spin switching temperature $T_{\text{sw}}$ and the applied magnetic field $H$ with a possible modulation, we have measured the $M_s-T$ curves (in ZFC regime) under different $H$. We find that $T_{\text{sw}}$ can be readily controlled in a wide temperature range by varying the magnitude of the applied magnetic field, as shown in Figure 3. Thus, $T_{\text{sw}}$ is observed to be very field-sensitive, but the magnitudes of the positive and negative $M_s$ around $T_{\text{sw}}$ is less sensitive to the initial increase of field (Figure 3a, Figure 3b). As the applied magnetic field is strengthened from 250 Oe to 20000 Oe, $T_{\text{sw}}$ changes dramatically from 356 K to 4 K (Figure 3c). Note that for a large $H > 2000$ Oe, the switched $M$ values from negative to positive will become nonsymmetrical and a smaller spin jump to positive $M$ will be observed. It is clear that, under low magnetic field of 250–600 Oe, $T_{\text{sw}}$ covers the most useful
temperature regime from 356 K to about 20 K and possible spin switching or magnetic sensor devices can be easily designed. For higher fields of 700–1000 Oe, the \( T_{\text{sw}}-H \) curve undergoes a knee point transition. And for the further high field of \( H>2000 \) Oe, the \( T_{\text{sw}}-H \) curve starts to reach a saturation. When \( H \) is 10000 Oe or higher, the negative magnetization and \( T_{\text{comp}} \) totally vanishes, whereas the residual spin switching transition \( T_{\text{sw}} \) can still be observed with a very small magnetization jump, and the total magnetization \( M_a \) is aligned parallel to the direction of field \( H \).

According to the previously reported work, the rare earth Sm ions in SmFeO\(_3\) system seem to establish long-range ordering below 140 K\(^\circ\)\(^{19-20}\), which is the highest temperature in the rare earth orthoferrites RFeO\(_3\) compounds. For other RFeO\(_3\) systems like R\( ^{5}\)Nd, Er, Ho, etc., their onset temperatures of long-range ordering of rare earth sublattices are all below 100 K. Our first-principles calculation study\(^{20}\) for NdFeO\(_3\) system indicates that the spin reorientation transition of Fe-sublattice can be ascribed to the exchange interaction between Nd-4f and Fe-3d electrons, which are mediated by O-2p state. As the temperature decreases, the superexchange angle of Fe-O-Fe gets larger, the Fe-O and Nd-O bonds become more covalent, and the exchange interactions become stronger. This study reveals that 4f-electrons of rare earth ions play the main role in triggering the spin reorientation transition of Fe-sublattice. The magnetic properties (magnetization) of RFeO\(_3\) below spin reorientation temperature should be dominated by the rare-earth sublattice, which is FM-coupled or AFM-coupled with the net FM vector of Fe-sublattice. For our case of SmFeO\(_3\) system, the delicate interactions among R-R, R-Fe, and Fe-Fe ions, and the role of rare earth Sm-sublattice are similar with other RFeO\(_3\) family members. Nonetheless, the uniquely high \( T_{SR} \) window (450–480 K) well above room temperature makes SmFeO\(_3\) system an exceptional material with spin switching \( T_{\text{sw}} \) near room temperature. Above \( T_{\text{sw}} \) for both ZFC and FC, the total magnetization stays parallel to the \( a \)-axis and slightly decreases with increasing temperature until the spin reorientation transition region of Fe-sublattice (Figure 1). This result suggests the gradual decrease of FM vector from Fe-sublattice and the progressive disappearance of the long-range ordering moments of Sm-sublattice, as consistent with literature of RFeO\(_3\) materials\(^{18}\).

Figure 4 shows the magnetic field dependence of the magnetization at the spin switching temperature of SmFeO\(_3\) single crystal, where the “lower” and “upper” refer to the magnitudes of magnetization before and after the spin switching transition, respectively; the “average” refers to the midpoint magnetization of the “lower” and “upper”; the upper inset shows zoom-in region marked by the dashed black rectangle for the field \( H<2000 \) Oe; the lower inset shows the magnetization jump \( (\Delta M) \) before and after the spin switching transition as a function of magnetic field. When \( H \) is higher than 1000 Oe, the transition temperature \( T_{\text{sw}} \) is suppressed to be...
likely that the key role of rare earth Sm ions might be found in many spin switching devices by using very low field. Furthermore, it is indeed the case since enhancing the magnetic field from 250 to 2000 Oe, the magnetization becomes always positive in the whole range of temperature. The slight rise of \( T_{\text{comp}} \) with the increasing field is because that the magnetic field enhances the value of the positive magnetization at the lower temperatures. Note that interestingly \( T_{\text{comp}} \) shows a small dip when \( H = 700-900 \) Oe, which is corresponding to the inflection point in the \( T_{\text{sw}} \)-\( H \) curve in Figure 3c and the sharp decrease of \( \Delta M \) in Figure 4. These details again demonstrate that the strong exchange interaction between the Sm-4f and Fe-3d electrons renders SmFeO\(_3\) possessing extremely alterable spin configurations by small perturbations like weak magnetic field, especially near room temperature.

Let us now come back and pay more attention to the first-order transition occurring at \( T_{\text{comp}} \) since such a sharp spin-reversal transition may be put in use for designing novel spin switching or magnetic sensor device. One may wonder what the origin of that first-order transition is. It is important to realize that, in addition to the exchange interaction between Sm-4f and Fe-3d electrons that result in opposite, temperature-dependent magnetizations in the Sm- and Fe-sublattices, the investigated rare-earth orthoferrites has also another energetic preference: it desires to have a total ferromagnetic moment being aligned along the field’s direction. Such desire should become more and more pronounced, with respect to the interactions between Fe-3d and Sm-4f electrons, when increasing the magnitude of the applied field. Figure 3 indicates that this is indeed the case since enhancing the magnetic field from 250 to 2000 Oe significantly shifts \( T_{\text{comp}} \) towards lower temperature, therefore leading to a maximum negative magnetization that considerably reduces. In fact, for the field stronger than 2000 Oe, the interaction between the total magnetization and the applied field prevails over the intrinsic exchange interactions between Sm-4f electrons and Fe-3d electrons, since the magnetization is always positive for any studied temperature – therefore annihilating the existence of a compensation temperature, but the temperature-induced spin flip is still observable through a very small jump (the so called spin switching effect in the above, see Figure 3). It is noticeable that, for the case of SmFeO\(_3\), the \( T_{\text{sw}} \) (under low field, such as 250 Oe) is much higher than that of other RFeO\(_3\) compounds (such as 29 K for NdFeO\(_3\), 65 K for ErFeO\(_3\) under 100 Oe), and the transition temperature can be modulated to a temperature as high as 356 K. This outstanding feature exhibits an evidence of strong exchange interaction between Sm-4f and Fe-3d electrons in the SmFeO\(_3\) system, since such an interaction coexists and lasts even up to room temperature. It is the strong Sm-4f and Fe-3d electrons interaction that makes SmFeO\(_3\) single crystal unique and may be put in use for designing novel spin switching devices in the near future. For device technology where small magnetic field of mT is applied by current pulses, we might propose substitutional doping Sm by other rare earth, e.g. Nd, to further lower the switching field at room temperature.

In conclusion, we have studied the temperature-induced multiple magnetic transition properties of single-crystal SmFeO\(_3\), by demonstrating spin reorientation, compensation and switching phenomena that originated from the Sm-4f and Fe-3d electrons and their interaction. In particular, SmFeO\(_3\) possesses an extremely useful magnetization that is very sensitive to small field perturbations even at room temperature. This makes SmFeO\(_3\) an easily manipulable candidate for practical application in spin switching devices by using very low field. Furthermore, this study indicates that the significant role of rare earth Sm ions in RFeO\(_3\) or other perovskite ABO\(_3\) compounds should be focused both in experimental and theoretical study. Sm-4f electrons may trigger more novel effects of spin state transitions near room temperature, which is desirable for the use of spin in future spintronics.

**Methods**

Single crystal of SmFeO\(_3\) was grown in a four-mirror optical-floating-zone furnace (FZ-T-10000-H-VI-P-SH, Crystal Systems Corp.) using 1.5 kW halogen lamps as the infrared radiation source with flowing air. The temperature of the molten zone was precisely controlled by adjusting the power of the lamps. During the growth process, the molten zone moved upwards at a rate of 3 mm h\(^{-1}\) with the seed rod (lower shaft) and the feed rod (upper shaft) counter rotating at 30 rpm in air flow of 5 L min\(^{-1}\). The compositional homogeneity and crystal morphology were confirmed by X-ray diffraction (XRD), and scanning electron microscopy (SEM) with energy-dispersive X-ray spectroscopy (EDX). All results confirmed the high homogeneity of the crystals studied.

Measurements of magnetization as a function of temperature (\( M-T \)) and magnetic field (\( M-H \)) were performed using the Quantum Design Physical Property Measurement System (type PPMMS-9) for temperatures below 400 K and the Lakeshore Vibrating Sample Magnetometer (VSM, type 7407) for temperatures from 300 K to 750 K. For the zero-field-cooling (ZFC) measurement, the sample was progressively cooled down under zero magnetic field until the temperature of 3 K is achieved. A magnetic field was then applied, and the sample was heated under this field to measure the magnetization as the temperature increases. For the field-cooling (FC) measurement, the sample was progressively cooled down to the temperature of 3 K under an applied magnetic field. Then, the sample was heated under the same field to measure the magnetization as the temperature increases.

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Acknowledgments

This work is supported by the National Natural Science Foundation of China (NSFC, Nos. 51372149, 50932003, 11274221, 11274222), QiMingXing Project (14QA1402000) of Shanghai Municipal Science and Technology Commission, Eastern Scholar Program and Shuguang Program (Grant No. 12SG34) from Shanghai Municipal Education Commission.

Author contributions

S.C., H.Z. and W.R. conceived the idea for this project and wrote the manuscript. S.C., H.Z. and B.K. did all the experiments and prepared the figures 1–5. S.C., H.Z., J.Z. and W.R. reviewed the manuscript. All authors contributed to the discussion of the results and commented on the manuscript.

Additional information

Competing financial interests: The authors declare no competing financial interests.

How to cite this article: Cao, S., Zhao, H., Kang, B., Zhang, J. & Ren, W. Temperature induced Spin Switching in SmFeO₃ Single Crystal. Sci. Rep. 4, 5960; DOI:10.1038/srep05960 (2014).

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