Electric field manipulation of magnetic and transport properties in SrRuO₃/Pb(Mg₁/₃Nb₂/₃)O₃-PbTiO₃ heterostructure

W. P. Zhou, Q. Li, Y. Q. Xiong, Q. M. Zhang, D. H. Wang, Q. Q. Cao, L. Y. Lv & Y. W. Du

Jiangsu Key Laboratory for Nano Technology and National Laboratory of Solid State Microstructures, Nanjing University, Nanjing 210093, People’s Republic of China.

The electric field manipulation of magnetic properties is currently of great interest for the opportunities provided in low-energy-consuming spintronics devices. Here, we report the effect of electric field on magnetic and transport properties of the ferromagnetic SrRuO₃ film which is epitaxially grown on Pb(Mg₁/₃Nb₂/₃)O₃-PbTiO₃ ferroelectric substrate. With the application of electric field on the substrate, the magnetization, Curie temperature and resistivity of SrRuO₃ are effectively modified. The mechanism of the electric field manipulation of these properties is ascribed to the rotations of RuO₆ oxygen octahedra caused by the electric-field-induced strain, which changes the overlap and hybridization between the Ru 4d orbitals and O 2p orbitals, resulting in the modification of the magnetic and electronic properties.

Magnetoelastic (ME) heterostructures consisting of ferromagnetic and ferroelectric elements have aroused great interest in recent years for their rich coupling mechanisms and promising applications in next-generation electronic devices¹⁻⁶. The ME effects can be classified as direct ME (DME) and converse ME (CME), which are characterized as magnetic-field-induced polarization and electric-field-induced magnetization, respectively. The CME effect, i.e. electric field control of magnetism, is now a research focus, since it can provide a low-consuming method to manipulate the magnetization¹⁻⁴. Up to now, the most used CME mode is the strain-mediated coupling¹⁻⁷. By depositing the ferromagnetic component on ferroelectric/piezoelectric substrate, the electric-field-induced strain can remarkably manipulate magnetization, giving rise to large ME effects⁸⁻¹⁰. Pb(Mg₁/₃Nb₂/₃)O₃-PbTiO₃ (PMN-PT) and PbZrₓTi₁₋ₓO₃ (PZT) are frequently selected as the ferroelectric/piezoelectric substrates due to their excellent piezoelectric and ferroelectric performance. As for the ferromagnetic components, magnetic perovskites are ideal candidates because they exhibit abundant physical properties and are easy to grow epitaxially on the ferroelectric substrates¹⁰⁻¹³. In magnetic-perovskite-based ME heterostructures, the magnetism can be manipulated by means of altering the magnetic anisotropy¹⁰⁻¹², affecting the phase separation¹¹⁻¹², or adjusting the spin state of magnetic ions¹³ through the electric-field-induced strain. In addition to strain-mediated mechanism, ferroelectric field-effect¹⁴ may also play a role in tuning magnetic and transport properties of magnetic perovskites in those ME heterostructures due to the accumulation or depletion of charge carries at interface through ferroelectric polarization reversal¹⁵⁻¹⁷. It is known that networks of corner-sharing oxygen octahedra play an important role in determining the magnetic and electronic properties of the perovskite through its deformations, rotations or tilts¹⁸⁻²⁰. Recently, theoretical results based on the first-principles calculation point out that the magnetization can be controlled via octahedral rotations in a perovskite related oxide²¹. Since the oxygen octahedral rotations in perovskites can be manipulated by applying stress¹⁶, it may provide an opportunity to achieve CME effect in magnetic perovskites.

SrRuO₃ (SRO) is a moderately correlated material and the only known 4d transition metal oxide with ferromagnetic order and metallic behavior, which has attracted an enormous amount of investigation for its intriguing magnetic and electronic properties²². The perovskite structure of SRO crystallizes in orthorhombic symmetry, exhibiting a GdFeO₃-kind of distortion associated with rotation and tilt of RuO₆ octahedra²². The manipulation of resistivity by ferroelectric field-effect has been reported in a SRO/PZT heterostructure, in which a 9% change of the resistivity upon reversing the ferroelectric polarization is observed²³. However, there are few reports about the electric-field control of magnetization or resistivity in SRO film using strain-mediated...
mechanism. As we know, density-functional theory calculations have predicted that the magnetic moment is suppressed under compressive strain and enhanced with tensile strain for (001)-epitaxial SRO films due to the strain-induced oxygen octahedral rotations. As for the experimental results, Gan et al reported that, compared with the strain relaxed film, the saturation magnetic moment can be suppressed in the compressive SRO film. Moreover, Terai et al found that the Curie temperature \(T_C\) of SRO is enhanced under tensile strain. Up to now, most of the studies about the strain dependent of magnetic properties for SRO are focused on the films deposited on various substrates, in which the different strain states are attributed to the mismatch of lattice parameters. Since the deposited SRO film on the substrates are observed almost one type of correlation between magnetic properties and structural and morphological features. Therefore, the structural and morphological features of the film exhibit the strain field causes a significant structural and magnetic effects by comparing with each other. However, depositing SRO film on the piezoelectric substrate can neglect these intricacies and provide an electric-field-controlled strain state in-situ, which is helpful to gain more insight on the strain impact on the magnetic properties of the film. In this paper, we deposit SRO film on PMN-PT substrate to investigate the electric field control of magnetization and resistivity by applying an electric field on the piezoelectric single crystal. The experimental results show that not only the saturation magnetization but also \(T_C\) can be affected by the electric field, showing an intrinsic CME effect. The effect of electric field on the resistivity further demonstrates the correlation between magnetic and electric properties of SRO film.

### Results

The XRD \(0\)-2\(\theta\) scan of SRO/PMN-PT heterostructure is shown in Fig. 1(a). Obviously, only the diffraction peaks of substrate and film can be observed, suggesting the single phase and highly (001)-oriented nature of the film. Although the structure of PMN-PT crystal and bulk SRO is rhombohedral and orthorhombic respectively, they can be regarded as the pseudocubic structures with lattice parameters of \(a_{PMN-PT}=4.02\) Å and \(a_{SRO}=3.93\) Å. Due to the lattice mismatch of 2.3% between SRO and PMN-PT, the SRO film is under the substrate-induced tensile strain along the in-plane direction. Besides the epitaxial growth, the surface morphology of the film is also important for the efficiency of strain transferred from the substrate to the film in strain-mediated ME heterostructures. The AFM image exhibited in Fig. 1(b) with an area of 2 \(\times\) 2 \(\mu\)m\(^2\) shows a rather smooth surface with the root mean square roughness about 0.4 nm. Therefore, the structural and morphological features of SRO film demonstrate a favorable condition for the following CME coupling measurements.

For investigating the effect of electric field on the magnetism, we measure the thermomagnetic curves for SRO film with or without electric field, where the applying magnetic field and electric field are 500 Oe and 8 kV/cm, respectively. As shown in Fig. 2, SRO film shows a typical ferromagnetic transition and the value of \(T_C\) under zero electric field is around 117 K. With the application of electric field, an obvious suppression of the magnetization and a shift of \(T_C\) are observed simultaneously. The upper inset of Fig. 2 shows the temperature dependence of the relative change of magnetization \(\Delta M/M = (M(E) - M(0))/M(0)\), where \(M(E)\) and \(M(0)\) are the magnetization with and without electric field, respectively. A negative CME effect with a peak value of \(-26.8\)% near 117 K is observed in SRO film by applying an electric field on the PMN-PT substrate. To further determine the variation of \(T_C\) caused by the electric field, we extract \(T_C\) from extrapolating the linear part of \(M^2-T\) curve to \(M=0\) in a temperature range below \(T_C\). As shown in the bottom inset of Fig. 2, \(T_C\) shifts from 117 K to 115 K with an electric field of 8 kV/cm. According to the earlier reports, the magnetization of the strain-mediated ME system is usually modified by altering the magnetic anisotropy, giving rise to an extrinsic CME effect. However, in the case of SRO/PMN-PT heterostructure, not only the magnetization but also \(T_C\) varies with the electric field, suggesting that the CME effect in SRO film may have other coupling mechanism.

Figure 3 shows the magnetic hysteresis loops \((M-H)\) at 115 K under the electric fields of zero and 12 kV/cm with the magnetic field parallel to the film surface, and the schematic diagram of laminate structure and magnetic measurement is shown in the inset. It is obvious that both \(M-H\) curves show typical ferromagnetic character and are almost saturated under the magnetic field of 5 kOe. With the application of electric field, a noticeable separation of magnetization can be observed. The value of \(\Delta M/M\) under the magnetic field of 5 kOe is about \(-8.6\)% As we know, the modification of magnetic anisotropy can affect the magnetization, but cannot change the saturation magnetization. Therefore, the electric field would have an intrinsic effect on the magnetism of SRO film due to the change of saturation magnetization.

To further explore the origin of electric field control of magnetism in SRO/PMN-PT heterostructure, the relative change of magnetization \(\Delta M/M\) as a function of bipolar electric field with a bias magnetic field of 500 Oe is measured at 115 K. As shown in Fig. 4(a), a hysteresis loop with a rather symmetric butterfly shape is observed, which is consistent with the shape of strain vs electric field hysteresis loop and different from that of polarization vs electric field hysteresis loop of the PMN-PT substrate. Thus the observed electric field manipulation of magnetization in SRO/PMN-PT heterostructure

![Figure 1](image-url) | (a) XRD pattern for SRO/PMN-PT heterostructure at room temperature. (b) Surface morphology of SRO film with an area of 2 \(\times\) 2 \(\mu\)m\(^2\).
would be ascribed to the electric-field-induced strain. When an electric field is applied on PMN-PT with the direction parallel to the polarization, the substrate would contract along the in-plane direction which causes the reduction of magnetization of the upper SRO layer about 9.2% under an electric field of 20 kV/cm. As the electric field reverse its direction, a tensile strain is produced, which leads to the enhancement of magnetization with a maximum value of about 6.7% at the coercivity field of 10.8 kV/cm. Compared with the earlier report about depositing SRO film on different substrates, this method of controlling magnetism in SRO is adjustable and scalable, since the applying electric field can be continuously tuned and the value of generated stress is also countable. After the substrate is poled with a positive electric field, the time dependence of magnetization and electric field is measured with a bias magnetic field of 500 Oe. As shown in Fig. 4(b), when an electric field of 12 kV/cm is applied, SRO film exhibits a low magnetization state of 0.139 μB/Ru. While the electric field turns off, it subsequently transforms into a high magnetization state of 0.149 μB/Ru. Furthermore, the magnetization of SRO film shows almost stable change after several cycles with the electric field switching on and off alternatively, suggesting that the magnetization of this heterostructure can be reversibly and reproducibly controlled by the electric field.

As mentioned above, SRO is a kind of moderately correlated material, in which the transport and magnetism are related with the conduction $d$ band, leading to an intrinsic coupling between the magnetic and transport properties. Therefore, the electric field would have influence on the transport properties of SRO film as well. Figure 5(a) shows the temperature dependence of resistivity for SRO film with and without electric fields, and the schematic image of electric measurement is shown in the inset. Under zero electric field, the resistivity decreases almost linearly with the temperature cooling down from room temperature, exhibiting a typical behavior of “bad metal”. When the temperature further decreases, a kink is observed around $T_C$ and the resistivity decreases rapidly below this temperature due to the loss of spin disorder scattering. It is obvious that the resistivity of SRO film decreases in the measured temperature regions with the application of an electric field on PMN-PT substrate. We also measure the hysteresis loop of the relative change of resistivity $\Delta R/R$ ($\Delta R/R = |R(E) - R(0)|/R(0)$) as a function of bipolar electric field at 115 K. As demonstrated in Fig. 5(b), $\Delta R/R$ exhibits a nearly symmetric butterfly shape with the variation of electric field, which is similar with the $\Delta M/M - E$ hysteresis loop, further suggesting that the electric field modulation of the resistivity stems from the electric-field-induced strain as well. Under the compressive strain, the resistivity of SRO film decreases about 1.3% at an electric field of 20 kV/cm. While the tensile strain is found to increase the resistivity with a maximum value about 1.1% at the coercivity field of 10 kV/cm. The resistivity response to the variation...
of electric field as a function of time for SRO film is also measured at 115 K with the substrate poles under a positive electric field. As shown in Fig. 5 (c), the resistivity reduces with the electric field switching on and restores its initial state immediately with the electric field switching off, showing a same behavior with that of magnetization. The reversible modulation of the resistivity $\Delta R/R$ is about $-0.5\%$ with an applied field of 6 kV/cm.

**Discussion**

It is well accepted that there is a rather strong hybridization between the Ru 4$d$ orbitals and O 2$p$ orbitals of SRO, leading to a ferromagnetic ground state$^{35-38}$. However, in the case of CaRuO$_3$, it possesses the same structure as that of SrRuO$_3$, but has larger rotations of the RO$_6$ oxygen octahedra, which results in a paramagnetic ground state$^{32}$. This fact suggests that the octahedral rotations and tilts play a key role in determining the magnetic state of ARuO$_3$ (A = Ca, Sr) system and the magnetic properties of them can be adjusted through modulation of the hybridization between the Ru 4$d$ and O 2$p$ states by structural distortions$^{38}$. In SRO/PMN-PT heterostructure, when an electric field is applied with the direction parallel to the polarization, the PMN-PT substrate would contract along in-plane direction and weaken the as-grown tensile strain. As illustrated in Fig. 6, in order to adapt this contraction, the RuO$_6$ octahedral rotation angle of SRO film would enlarge correspondingly, giving rise to the enhancement of the overlap and hybridization between the Ru 4$d$ and O 2$p$ orbitals. Therefore, the magnetic interaction would reduce$^{29,39}$, which leads to the decrease of magnetization and $T_C$, showing an intrinsic CME effect.

Since the strain, $\Delta M/M$ and $\Delta R/R$ have the similar electric-field dependence, we can assume that the magnetization and resistivity change linearly with the strain. When an electric field of 8 kV/cm is applied, an in-plane compression of about 0.1% is generated$^{40}$, leading to the decrease of $T_C$ of SRO film for about 2 K, as shown in Fig. 2. This result is comparable with that reported by Terai $et$ $al$$^{25}$, where the strain of 0.25% results in the variation of $T_C$ for about 4 K for SRO film grown on Ba$_{1-x}$Sr$_x$TiO$_3$/BaTiO$_3$ buffer layers. As for the magnetization, the saturation magnetization of SRO film decreases about 8.6% near $T_C$ under an electric field of 12 kV/cm (about 0.12% compressive strain$^{39}$), corresponding to about 71.7% reduction of magnetization under 1% compressive strain. This result is dramatically larger than that reported by Gan $et$ $al$$^{39}$, in which the saturation magnetization of SRO decreases about 20% under a 0.67% compressive strain at 10 K. The larger effect of elastic strain on saturation magnetization in our sample reflects that the magnetism of SRO film responds more sensitive to the strain near $T_C$, as revealed in the upper inset of Fig. 2.

The electric field manipulation of resistivity can also be attributed to the oxygen octahedral rotations caused by the strain. In order to accommodate the electric-field-induced strain, the Ru-O-Ru bond angles, which is determined by the rotation of the oxygen octahedra, would change and the overlap between the Ru 4$d$ states and O 2$p$ states would enhance correspondingly$^{15,40}$. As a result, the variation of the Ru-O-Ru bond angles affects the electron hopping matrix elements while the enhanced orbital overlap broadens the electron bandwidth and reduces the correlation effect, leading to the decrease of resistivity$^{15,40,41}$.

In conclusion, the electric field control of magnetic and transport properties of SRO/PMN-PT heterostructure have been investigated. We demonstrate that the saturation magnetization, $T_C$ and resistivity of SRO film can be manipulated with the application of electric field. Moreover, the magnetizaton and resistivity can be controlled reversibly and reproducibly with in situ electric field. The observed CME effect in SRO/PMN-PT heterostructure can be understood in the scenario of electric-field- induce strain modulates the hybridization between the Ru 4$d$ orbitals and O 2$p$ orbitals through altering the octahedral rotations. This result suggests an alternative mechanism for intrinsic CME effect in perovskite films through controlling octahedral rotations. The electric field control of magnetic and transport properties in SRO/PMN-PT heterostructure provide a potential application in low-energy-consuming spintronics devices.
Methods

SRO film with thickness about 300 nm was deposited on polished (001)-oriented PMN-PT substrate with dimensions of 5 × 0.5 mm² by pulsed laser deposition method with a 248 nm KrF excimer laser. During the growth, the substrate temperature was 700°C and the background oxygen pressure was 13 Pa. After deposition, the film was cooled down in oxygen of 2 × 10⁻⁶ Pa. X-ray diffraction (XRD) and atomic force microscopy (AFM) measurements were performed to characterize the structure and surface topography of the film. To conduct the investigation of electric field on magnetic and transport properties in situ, Au layer was sputtered on the backside of the PMN-PT substrate as bottom electrode while the conducting SRO film was deposited on polished (001)-oriented substrate symmetry. To conduct the investigation of electric field on magnetic and transport properties, in situ electric voltage was applied. The transport properties were measured with a magnetic property measurement system (MPMS, Quantum Design) and the transport properties were measured with a magnetic property measurement system (MPMS, Quantum Design).

Electric voltage was applied in situ by a source meter (Keithley, model 2410).

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Author contributions

W.P.Z. and D.H.W. designed the experiments. W.P.Z., Q.L., Y.Q.X., Q.M.Z. and L.Y.L. conducted the experiments. W.P.Z. and D.H.W. wrote the paper. W.P.Z. and D.H.W. analysed the data. Results were analyzed and interpreted by W.P.Z., D.H.W., Q.Q.C. and Y.W.D. The author contributions have been reviewed by the authors. W.P.Z. and D.H.W. designed the experiments. W.P.Z., Q.L., Y.Q.X., Q.M.Z. and L.Y.L. conducted the experiments. W.P.Z. and D.H.W. wrote the paper. W.P.Z. and D.H.W. analysed the data. Results were analyzed and interpreted by W.P.Z., D.H.W., Q.Q.C. and Y.W.D. The author contributions have been reviewed by the authors.

Additional information

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