Resistive switching and optical properties of strontium ferrate titanate thin film prepared via chemical solution deposition

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Abstract: The polycrystalline strontium ferrate titanate (SrFe_{0.1}Ti_{0.9}O_{3}, SFTO) thin films have been successfully prepared by chemical solution method. By analyzing the current–voltage (I–V) characteristics, we discuss the conduction mechanism of SFTO. It is found that the number of oxygen vacancy defects is increased by Fe ion doping, making SFTO be with better resistive switching property. Fe ion doping can also enhance the absorption of strontium titanate to be exposed to visible light, which is associated with the change of energy band. The band gap width (2.84 eV) of SFTO films is figured out, which is less than that of pure strontium titanate. Due to more oxygen vacancy defects caused by Fe ion doping, the band gap width of strontium titanate was reduced slightly. The defect types of SFTO thin films can be determined by electron paramagnetic resonance spectroscopy. In addition, we analyzed the energy band and state density of SFTO by first-principles calculation based on density functional theory, and found that Fe ion doping can reduce the band gap width of strontium titanate with micro-regulation on the band structure. A chemical state of SFTO was analyzed by X-ray photo electron spectroscopy. At the same time, the structure and morphology of SFTO were characterized by X-ray diffraction and scanning electron microscope. This study deepens further understanding of the influence of Fe ion doping on the structure and properties of strontium ferrate titanate, which is expected to be a functional thin film material for memristor devices.

Keywords: SrTiO_{3}; thin films; resistive switching; oxygen vacancy; first principles

1 Introduction

During the past few decades, a large number of experimental and theoretical studies on ABO_{3} perovskite compounds have been carried out, especially the typical material SrTiO_{3}. One motive force to study SrTiO_{3} perovskite is the opportunity to regulate and control its electronic and ionic defect structures, so as to gain a wide range of applications (energy harvesting, memory device, oxygen sensor, catalysis, fuel cell cathodes, etc.) [1–5]. There are a number of researches by first-principles studies of defective perovskite to investigate electronic structure and performance [6–11]. As for SrTiO_{3}, it can be changed into a p-type material by substituting Fe to Ti ions [7]. Some previous work revealed the interesting properties such as ferroelectric [8], photochromic [9], oxygen sensitive properties [10,11], as well as Jahn–Teller distortion [12]. In addition, dielectric properties, defect state, and conduction mechanism have been studied in strontium ferrate titanate system [13–16].

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In recent years, strontium ferrate titanate has attracted more and more attention in studying defect chemistry, carrier transport properties, and ferroelectric [17,18]. Compared with the pure SrTiO$_3$, defect state like oxygen vacancies in strontium ferrate titanate sustains charge balance due to the diverse valence states between Fe and Ti ions. The formation of vacancies and defect state affects both electrical and magnetic behaviors [19]. As the previous report said, the potential barrier height is influenced as a result of oxygen vacancy concentration [20]. Fe ion doping of 10% content improves the ferroelectric of SrTiO$_3$ exactly. Moreover, ferromagnetic properties in perovskites depend on not only oxygen vacancies and defects but also the annealing atmosphere and temperature [21]. Afterward, the annealing atmosphere effects on device characteristics have been investigated [22,23].

Up to now, iron-substituted SrTiO$_3$ system has been studied by multifarious experimental and theoretical researches. Nevertheless, there is still not any thorough elucidation of the relation to the concentration of point defects. An intensive understanding of the energy band regulation–property relation is essential. In this work, we presented the preparation and electric properties including resistive switching characteristic, ferroelectric, and optoelectronic of SrFe$_{0.1}$Ti$_{0.9}$O$_{3–x}$ thin film deposited on fluorine doped tin oxide (FTO) coated glass substrate via chemical solution deposition method. In this study, it is the first time to utilize the first-principles theoretical calculation based on density functional theory (DFT) for investigating the electronic structure and state density of perovskite material SrFe$_{0.1}$Ti$_{0.9}$O$_{3–x}$ with 10% Fe ions to elaborate the experimental results, which can give us enlightenment to explore and verify the effect of doping in band gap regulation.

2 Experimental

The method we used in our work to prepare the SrFe$_{0.1}$Ti$_{0.9}$O$_{3–x}$ (SFTO) thin film on the FTO (the thickness is 330 nm and the resistance is 7 Ω) coated glass substrate (purchased from the School Experimental Raw Materials Purchasing Platform, Guangdong University of Technology) was chemical solution deposition (CSD) [24]. The raw materials we selected were strontium acetate (0.859 g), iron 9-hydrate nitrate (0.161 g), and butyl titanate (1.225 mL). In addition, methanol, and 36% acetic acid and acetylacetone were selected as solvent and stabilizer, respectively. First, strontium acetate was dissolved in methanol and iron nitrate was dissolved in 36% acetic acid. In order to dissolve the strontium acetate and iron 9-hydrate nitrate completely, we put the beaker on the heating table and stirred it for 2 h to be dissolved at a temperature of 50 °C. Secondly, appropriate amount of butyl titanate was taken from the measuring cylinder, and 2 drops (about 0.04 mL) of acetylacetonate were added to stabilize it. Finally, we mixed the three solutions together and bottled the mixed solution after filtering. The concentration of the final mixed solution was 0.2 mol/L. In order to ensure the quality of the prepared thin film, we put the prepared solution stand at room temperature for one day and observed that there was no precipitate before casting the film. FTO-coated glass substrates were cleaned completely before the preparation. We put the FTO-coated glass substrates into ultrasonic cleaner for 15 min and dried the substrates on the heating platform. The above operation needs to be repeated three times to ensure that the substrates are thoroughly cleaned. The solution was dropped onto the FTO substrate and spun at a speed of 4000 rpm for 30 s. Two high speed spins were performed according to the thickness of the required film. After that, thin film samples were baked on a heating table at 400 °C for 30 min. Hydrofluoric acid with low concentration will not corrode the FTO layer. Therefore, we used the hydrofluoric acid (0.1 mol/L) to wipe one corner on the SFTO layer of the thin film samples—in order to plate the electrode on the surface after annealing. At the end, all the thin film samples (SFTO and STO) were annealed at 650 °C for 15 min in the oxygen atmosphere by the rapid thermal annealing furnace RTP-1000D4 facility.

X-ray diffraction, X-ray photoelectron spectroscopy (XPS), and electron paramagnetic resonance spectroscopy were used to characterize the structure, chemical states, and defect center of the thin film samples. The current–voltage ($I–V$) characteristic of the Au/SFTO/FTO/Glass device was measured with the two-probe method by the Keithley 2400 programmable electrometer under room temperature. Additionally, first-principles calculations based on density functional theory were used to analyze the band structure and electronic states of the SFTO samples.

3 Results and discussion

3.1 Structure, morphology, and chemical states

To prove the crystalline state of the strontium ferrate...
titanate thin film we prepared, X-ray diffraction was applied to characterize the structure of the samples. As we can see from Fig. 1, the red and blue curves represent the XRD results of SFTO and pure strontium titanate (STO was prepared with SFTO at the same time and same conditions), respectively. It can be seen from the periodic table of chemical elements that the ionic radius of iron is smaller than that of titanium. According to the Bragg diffraction equation (2d sinθ = nλ), the XRD diffraction peaks of SFTO shift slightly to the direction of small angle. Peaks of 23.3° (100), 32.5° (110), 46.6° (200), and 58.2° (211) crystal planes of SFTO were observed obviously which are corresponding to JCPDS No. 35-0734 from the Joint Committee on Power Diffraction Standards (JCPDS) database, indicating that SFTO thin film is crystallized in cubic structure. No second phase was observed in the X-ray diffraction result. Therefore, Fe ions successfully replace Ti ions in strontium titanate matrix and the same perovskite structure of strontium ferrite as strontium titanate is formed [25]. The scanning electron microscopy (SEM) result of cross section is shown in Fig. 2(a). As can be seen from the results of the cross section image (Fig. 2(a)), the stratification is obvious. The top layer is the strontium ferrate titanate thin film, while the middle layer is FTO conductive layer, the bottom layer is the glass, the thickness of SFTO film is about 120 nm, and the particles are relatively dense [26]. Figure 2(b) shows the morphology observed by the atomic force microscopy (AFM). From Fig. 2(b), we can see the slight sharp grain morphology with the typical surface roughness which is closely related to the annealing temperature (SFTO was treated at the high temperature of 650 °C).

In order to analyze the chemical composition, proportion, and chemical valence of SFTO thin film sample, XPS results are shown in Fig. 3. Narrow spectrum scanning was acquired from the XPS spectra for the Sr 3d, Ti 2p, O 1s, and Fe 2p levels of SrFe₀.₁Ti₀.₉O₃₋ₓ thin film sample. Two peaks of Sr 3d locating at 132.10 eV (Sr 3d₅/₂) and 133.78 eV (Sr 3d₃/₂) are observed in Fig. 3(a), which indicates Sr ions with chemical state of 2⁺ [27]. Two peaks of Ti 2p₁/₂ and Ti 2p₃/₂ are located at 457.52 and 463.24 eV in Fig. 3(b), which indicates Ti ions with chemical state of 4⁺. The O 1s spectra are shown in Fig. 3(c), and the binding energy of O 1s spectra possesses two peaks. The stronger peak appears at 528.59 eV and it is closely related to O²⁻ ions which is associated with O element in SFTO lattice [4]. The lower peak at 530.54 eV is owing to intermediate oxidation state for O element. It may be related to the chemical-adsorbed oxygen on the thin film surface, which is associated with defects such as oxygen vacancies. As shown in Fig. 3(d), Fe 2p XPS spectra contain a doublet of Fe 2p₁/₂ and 2p₃/₂. The positions of the peaks are located at 709.60 and 723.25 eV, respectively. It indicates Fe element with chemical state of 3⁺ [28]. Due to spin–orbit coupling, Fe 2p₁/₂ peak is stronger than Fe 2p₃/₂ peak. Fe²⁺ electronic configuration is 3d⁶, while Fe³⁺ is 3d⁵. It means that the Fe²⁺ would have a longer life-time in the comparison to Fe³⁺, and therefore the selected full width at half maximum (FWHM) of Fe 2p₁/₂ peak is expected to be smaller than the Fe 2p₃/₂ peak.

![Fig. 1](https://www.springer.com/journal/40145)

**Fig. 1** XRD patterns of the SFTO film on FTO/Glass substrates.

![Fig. 2](https://www.springer.com/journal/40145)

**Fig. 2** (a) Cross section SEM image and (b) AFM image of the SFTO/FTO/Glass device.
In addition, the bimodal spectra of Fe 2p indicate Fe element with 4+ valence. Therefore, both Fe$^{3+}$ and Fe$^{4+}$ ions exist in SFTO thin film [29]. It has been reported in Ref. [30] that the increase of charge at the Fe site leads to the shifting of Fe 2p spectrum toward higher binding energy. Thus, it can be concluded that SrTiO$_3$ with 10% Fe ion doping can increase Fe$^{3+}$ and Fe$^{4+}$ ions effectively, resulting in the formation of oxygen vacancy defects [30]. The process can be briefly written into the defect reaction equations to deepen the understanding of Fe ion doping effect. The reactions are as following [31]:

$$\begin{align*}
2O_2^{-} & \rightarrow 2V_0^{-} + 4e^\prime + O_2^{-} \quad (1) \\
2Fe^{4+} + 2e^\prime & \rightarrow 2Fe^{3+} \quad (2) \\
2Fe^{3+} + 2e^\prime & \rightarrow 2Fe^{2+} \quad (3)
\end{align*}$$

where $e^\prime$ is the free electron, $O_2^{-}$ represents the oxidation of lattice oxygen, and $V_0^{-}$ is the oxygen vacancy.

### 3.2 \(I-V\) characteristics

Resistive switching behavior of Au/SFTO/FTO/Glass device is shown in Fig. 4. Voltage sweeping ($0 \rightarrow -3 \rightarrow 0 \rightarrow +3 \rightarrow 0$ V) is employed for \(I-V\) measurement. It can be seen from Fig. 4(b) that the current of the Au/SFTO/FTO/Glass device increases with the gradual increase of negative voltage at the beginning, and the device is in a low resistance state (LRS), as shown in path 1 in the Fig. 4(b). When the negative bias exceeds the threshold voltage of 2.8 V, the current rapidly decreases from $1.7 \times 10^{-2}$ to $6 \times 10^{-4}$ mA, and the state of the device switches from LRS to high resistance state (HRS). As the negative bias decreases, the current approaches 0, as shown in path 2 in Fig. 4(b). When the scanning voltage is from 0 to 3 V, the current of the thin film device gradually increases from 0. When the voltage exceeds the forward threshold voltage of 2.95 V, the current drops rapidly, and the device switches from LRS to HRS, as shown in path 3 in Fig. 4(b). When the forward bias gradually decreases from 3 to 0 V, the current intensity also decreases to 0, returning to the original state. At this point, a complete cycle test is completed. It can be concluded that the SFTO thin film device has the resistance switching behavior and the performance of memristors. Under the same test conditions, the resistive switching characteristics of SrTiO$_3$ cannot be found. We will discuss the reasons in the following sections. Figure 4(c) shows the \(I-V\) curve under semi-logarithmic coordinates. The 100-cycle \(I-V\) curves are shown in Fig. S1 in the Electronic Supplementary Material.
We can calculate the switch ratio from Fig. 4(c). Switch ratio ($\frac{R_{\text{High}}}{R_{\text{Low}}}$) at $-2.8$ V is about 30. We also tested the switch retention characteristics to ensure the resistive switching stability of the strontium ferrate titanate. The cycle endurance results are shown in Figs. 5(a) and 5(b). Resistances of HRS and LRS are read at $-2.8$ V and $+2.8$ V. Average ratio of HRS/LRS are about 30 and 10, respectively. From the results we can conclude that the resistive switching property of strontium ferrate titanate has not only good stability but also higher switching ratio compared with the pure strontium titanate.

In order to figure out the resistive switching mechanism of Au/SFTO/FTO/Glass device, $I$–$V$ characteristics under negative and positive applied bias are replotted as log $I$ vs. log $V$ and shown in Figs. 6(a) and 6(b), respectively. The slopes of fitting curve in LRS under negative and positive applied bias are 1.05 and 1.09, respectively, corresponding to the ohmic conduction mechanism. For HRS, ohmic conduction mechanism is dominant in low voltage region under both negative and positive applied bias, which is confirmed by the slopes of fitting curves of 1.05 and 1.07, respectively. However, conduction mechanism switches to space-charge limited current (SCLC) mechanism in high voltage region, which is confirmed by the slopes of fitting curves of 1.93 and 2.13. In general, SCLC mechanism is mainly related to the presence of defect centers. The current density of space-charge limited current $J_{\text{SCLC}}$ can be explained as following [32]:

$$J_{\text{SCLC}} = \frac{9\mu^2}{8d^3} \varepsilon \mu$$

where $V$ is the bias voltage, $d$ is the thickness of the thin film, $\varepsilon$ is the dielectric constant of the thin film, and $\mu$ is the electron mobility. Under the application of bias voltage, the defect centers can capture the free carriers excited by the electric field. In order to understand the conduction mechanism of strontium ferrate titanate thin film devices more intuitively, a reasonable model is proposed to explain the role of oxygen vacancy in the conduction process shown in Fig. 7.

![Fig. 4](image-url) **Fig. 4** Resistance properties of the SFTO/FTO/Glass device. (a) Measurement schematic diagram of the Au/SFTO/FTO/Glass device. (b) $I$–$V$ characteristic curve in Cartesian coordinates at the bias voltage of ±3 V and $I$–$V$ characteristics measuring structure is shown in the inset. The inset is the $I$–$V$ characteristic of pure SrTiO$_3$ in Cartesian coordinates. (c) $I$–$V$ characteristic curve in logarithmic coordinates and the direction of measurement is shown in the order of arrows from 1 to 4. The inset is the $I$–$V$ characteristic of pure SrTiO$_3$ in logarithmic coordinates.

![Fig. 5](image-url) **Fig. 5** Resistance retention characteristics at: (a) $-2.8$ V and (b) $+2.8$ V of the SFTO/FTO/Glass device.
Oxygen vacancies are inevitable in the preparation of thin films and the redistribution of electric charge on the surface of the devices is generated when the electrodes are plated to form the structure of the strontium ferrate titanate thin film devices. Therefore, before the device structure is formed without bias voltage, due to different work functions of the metal electrode and the strontium ferrate titanate thin film, the two different materials contact at the interface and form the Schottky junction. The electrons cannot freely cross the Schottky barrier, and the Fermi level of the gold electrode is at the same level with the Fermi level of the strontium ferrate titanate thin film which is shown in Fig. 7(a). $E_C$ is the conduction band in the SFTO band structure, $E_V$ is the valence band, and $E_{FS}$ is the Fermi level. $W_D$ represents the barrier width formed by the contact between gold electrode and SFTO film. When the forward bias voltage is added, the potential barrier at the interface will decrease, the Fermi energy level of the strontium ferrate titanate thin film will rise, and the electrons will fill into the oxygen vacancy and be captured by the oxygen vacancy. A large number of electrons tend to move toward the direction of the gold electrode, forming an LRS, as shown in Fig. 7(b). When the reverse bias voltage is applied, the barrier at the interface will increase, the oxygen vacancy will release the captured electrons, and the Fermi energy level of the film will decrease. However, due to the increase of the barrier, only a small part of the electrons on the surface of the gold electrode can cross the barrier and enter the conduction band of the semiconductor thin film, thus forming an HRS which is shown in Fig. 7(c). When the direction of electric field is constantly changed, the strontium ferrate titanate thin film device can switch between high and low resistance states. Additionally, the more oxygen vacancies the thin film has, the higher ratio of high and low resistance states the thin film would have. By doping Fe ions, the defect concentration of strontium ferrate titanate thin film can be changed to produce more oxygen vacancies which affect the migration of the free electrons. Thus, by contrasting the $I-V$ characteristics between the strontium ferrate titanate thin film and the pure strontium titanate thin film (Figs. 4(b) and 4(c)) and the conduction mechanism simulation diagram analysis (Fig. 7), we can draw the conclusion that strontium ferrate titanate thin film has relatively favorable resistive switching performance.
3. 3 Optical properties

UV–Vis absorption spectra of SFTO thin film are shown in Fig. 8(a). We utilize UV–Vis spectrophotometer (ThermoFisher Evolution 220) to measure absorption spectrum of SFTO and STO. Before the measurement, we use blank sample (which provided from the measurement system) for calibration. After that, we choose the measurement mode of reflectance. The wavelength range is from 330 to 600 nm. In order to ensure the accuracy of reflection spectra, we repeat the measurement three times for each sample. We also utilize the same UV–Vis spectrophotometer to measure the transmittance. The test wavelength range keeps the same with the reflection measurement. Transmittance of SFTO and STO thin film rises with the increase of test wavelength. It is noted that the reflection and transmittance measurement on different thin film samples must be calibrated respectively. We repeat the transmittance measurement three times for each sample as well for attaining accuracy results. At last, we use the reflection spectra and transmission spectrum of SFTO and STO thin film samples to calculate the absorption spectrum respectively. From Fig. 8(a), it can be seen that SFTO thin film absorbs more UV light with a wavelength lower than 365 nm, and the UV response is relatively sensitive. According to Tauc’s law, the relationship between absorption coefficient (α) and photon energy (hν) can be expressed as

\[ \alpha = \frac{1}{d} \times \ln \left( \frac{1}{T} \right) \]  \hspace{1cm} (5)

\[ h\nu = \frac{1240}{\lambda} \]  \hspace{1cm} (6)

where \( T \) and \( \lambda \) are the transmittance and wavelength, respectively. The optical band gap (\( E_g \)) can be derived by the tangent line of the known curve. \( E_g \) of 2.84 eV is obtained for SFTO thin film as shown in Fig. 8(b), being lower than the \( E_g \) of pure SrTiO\(_3\) (3.24 eV). The doping of Fe ions can lower the \( E_g \) of strontium titanate. Thus, SFTO thin films have better electric properties which are expected for the application of photodetector with faster response [33].

3. 4 Electron paramagnetic resonance spectroscopy of SFTO and STO

In order to determine the defect type of the thin film of strontium titanate with 10% Fe iron doping, we studied its electron paramagnetic resonance (EPR) spectra. EPR is a magnetic resonance technique used to characterize the unpaired electrons or single electron states of materials. The relative intensity of SFTO is three times larger than that of the pure SrTiO\(_3\). In fact, there are intrinsic defects (such as oxygen vacancies and line defect) in pure SrTiO\(_3\) crystal structure and the defect states would produce defect energy level in the band gap which can trap the electrons during the electron transition process. From the EPR spectra, g factor of the sample can also be calculated and used to determine whether there is an electron defect center or a whole defect center by comparing it with the g factor of the free electron. Some researchers used EPR technology to analyze O\(_2^–\) ions and Ti\(^{3+}\) ions on the surface of polycrystalline TiO\(_2\) and SrTiO\(_3\), respectively [34,35]. Figure 9 shows the EPR results of SFTO thin film sample. It is found that the EPR peak occurs at a position of 3800 Gs magnetic field intensity. The g factor of strontium titanate SFTO thin film sample can be calculated by the following equation:

\[ g = \frac{0.7144773 \times \nu}{B_0} \]  \hspace{1cm} (7)

where \( \nu \) is the frequency and \( B_0 \) is the magnetic field. g factor of SFTO thin film sample is 1.9237, which is
Fig. 9 EPR spectroscopy of strontium titanate films with iron content of 10%. The defect type can be determined by calculating $g$ from the EPR spectroscopy.

less than that of free electron (2.0023), indicating that there is an electron defect center in the thin film sample. To some extent, it can be concluded that Fe ions have successfully replaced the Ti ions in strontium titanate matrix. Fe ions can attract more electron density distribution than Ti ions and lead to the redistribution of electron cloud density near the Fe-doped impurity, which is closely related to the formation of oxygen vacancies.

3.5 First-principles calculations

It has been reported that a phenomenon of Jahn–Teller distortion in strontium ferrate titanate was found [17]. It is closely related to the Fe–O bond length. Fe–O bond length will change when Fe ions substitute to Ti ions, producing relative displacement of oxygen and related oxygen defects [36]. The experiments show that Jahn–Teller distortion decreases with the increase of Fe ion concentration. Further analysis on electronic state density, band structure, and absorption spectrum can be studied via first-principles calculation based on density functional theory for SFTO. In order to better illustrate the band structure and electron state density of SrFe$_{0.1}$Ti$_{0.9}$O$_3$, SrTiO$_3$, and SrFeO$_3$ are also studied. First-principles calculations are carried out via the Vienna ab initio simulation package (VASP) utilizing the projector augmented waves (PAW) technique [37]. Before the calculation based upon the density functional theory (DFT), the $k$-point grid size and the advolution are set up with respect to the cut of energy. We ensure that each structure (referring to SrFe$_{0.1}$Ti$_{0.9}$O$_3$, SrTiO$_3$, and SrFeO$_3$) has a appropriate number of $k$ points for the integration of Brillouin-zone. The electronic energies are converged to at least $10^{-5}$ eV/atom. As for the modeling of SFTO (10% Fe atoms on the Ti site), it is important to build an appropriate crystalline structure which is shown in Fig. S1 in the ESM. Calculation results are shown in Fig. 10. Among them, Figs. 10(a) and 10(b) show the band structure and electron state density of SrTiO$_3$, respectively. The top of the valence band and the bottom...
of conduction band are mainly made up of O 2p states and Ti 3d states, respectively. Figures 10(e) and 10(f) illustrate the band structure and electron state density of SrFeO$_3$–. It is obviously seen from the comparison of the electron state density for spinning up and spinning down of the electron densities, where the effect of spin polarization is derived from the Fe 2p states, a little bit from Fe 3d states. The results in Figs. 10(e) and 10(f) also indicate that there is a strong superposition of Fe 3d orbital electron states and O 2p orbital electron states near the Fermi level. In addition, the contribution of the main electron states near the Fermi level is derived from the Eg states of the Fe ions and O 2p electron states. Figures 10(c) and 10(d) show the band structure and electron state density of SrFe$_{0.1}$Ti$_{0.9}$O$_3$–. The top of the valence band consists of O 2p orbital electron states, while the bottom of virtual bands mainly originates from Ti 3d electron states [38]. It can be concluded that the band gap of strontium titanate with 10% Fe-doping is smaller than that of the pure strontium titanate, which indicates that Fe ions can regulate the energy band of strontium titanate and form the defect state effectively.

4 Conclusions

In this study, the resistance characteristics and band structure of strontium ferrate titanate/FTO devices are studied. Polycrystalline strontium ferrate titanate thin films were successfully prepared on FTO/Glass substrate by chemical solution deposition. XRD results showed that the characteristic peaks of strontium ferrite titanate were consistent with that of polycrystalline strontium titanate. It can be seen from SEM that the grains of strontium ferrate titanate films are relatively dense and have good crystallinity. The analysis results of EPR and XPS show that Fe ions can produce oxygen vacancy and other defects, and form electron trap center, which has important influence on the resistance characteristics of strontium ferrate titanate. Typical bipolar resistance is easy to be found in the $I$–$V$ characteristic test. By the means of log–log fitting, we found that ohmic conduction is the conduction mechanism in the LRS, and ohmic conduction and space charge limited current conduction are the two mechanisms in the HRS. The first-principles calculation shows that the band structure of strontium titanate has changed after the doping of Fe ions. In addition, through the absorption spectrum test of strontium ferrate titanate, it was found that the absorption of partial visible light was enhanced. By absorption spectrum calculation, the band gap width $E_g$ of the sample was 2.84 eV, slightly smaller than the band gap width 3.24 eV of strontium titanate. The experimental results were consistent with the theoretical calculation.

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Electronic Supplementary Material

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