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RESEARCH



Thermionic Emission Induced Current-Voltage behaviour of GdMnO₃/Al:ZnO/STO and GdMnO₃/ZnO/STO Thin Film Heterostructures

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ABSTRACT

We report the results of studies on the rectifying behaviour and tunnelling conduction in GdMnO3/ZnO/STO and GdMnO3/Al:ZnO/STO thin film heterostructures comprising of p-n junctions fabricated using the Pulsed Laser Deposition (PLD) technique. A structural study using grazing angle mode XRD depicts polycrystalline growth and confirms the phase purity. The AFM micrographs reveals the different grain growth and grain sizes of the prepared thin film heterostructures. Room temperature Raman spectra shows the presence of various vibrational modes in both the thin film heterostructures, the transport studies using I–V measurements at room temperature is explained using various models. The temperature dependent transport studies using I-V measurements at various temperatures reveal the rectifying behaviour and the difference in the I-V behaviour at various temperature can be understood on the basis of interface effect at the junction, which has been attributed to the presence of the various conduction phenomena through the junctions and the change in barrier height with the temperature for both presently studied thin film heterostructures.

KEYWORDS

Heterostructures, pulsed laser deposition, space charge limited conduction, thermionic emission, barrier height.

INTRODUCTION

Multiferroics are the important class of materials with various ferroic properties like ferromagnetism, ferroelectricity and ferro-elasticity in single phase which convey temenos interest in research. From discovery of perovskite structure especially rare earth manganites with $AMnO_3$ (A= rare earth elements) have obtained extensive attention, as they possess interesting properties like magnetoresistance, Jahn-Teller distortion, Zener Double Exchange, multiferroic properties [1-7]. All these properties can be modified under different external conditions like Heat treatment, pressure, applied fields, irradiation effect, A-site/B-site doping effect etc. [8-14] are induced modification in properties of manganite materials. Manganites can enhance the properties/efficiency of capacitors, transistors, FETs, sensors, memory storage devices etc. [15-19]. Manganites attribute the strong relations between spins, lattice, orbitals etc. like fundamental parameters which are responsible for various behaviors such as paramagnetic insulator, ferromagnetic insulator, antiferromagnetic insulator, paramagnetic metallic, ferromagnetic metallic [15,16]. To explore this kind of behavior, manganite have been studies in form of polycrystalline bulk materials, composites, thin films, corecells etc. [10,15].

Comparison of polycrystalline bulk and thin films shows a large variation in microstructural as well as different physical properties. Properties of thin films depends on different conditions like choice of substrate, deposition techniques, deposition parameters etc. [17,18]. There are several deposition techniques to fabricate thin films such as Spin Coating, Deep Coating, Atomic Layer Deposition (ALD), RF & DC Sputtering, Chemical Vapor Deposition (CVD), Pulse Laser Deposition (PLD) etc.[18-20]. Pulsed Laser Deposition (PLD) is most used thin film fabrication technique because as compared to other techniques used in research it provides less stoichiometric distortion. In PLD technique one can control several parameters like substrate temperature, oxygen pressure etc.

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There are several reports available for I-V properties of manganite thin films. Khacher et al. [21] have investigated the temperature dependent I-V behaviour of manganite based thin films, reported that positive magnetoresistance shows strong dependency on the temperature as well as magnetic field. Kataria et al. [14] have reported the temperature and magnetic field sensitive resistivity behaviour with SHI irradiation. Temperature plays important role in modification of physical properties of manganite-based structures. Ravalia et al. [13] investigate the variation in barrier height of manganite based thin films with temperature. Solanki et al. [22] reported the light dependency of I-V behaviour of LCMO/Si p-n junction and the thickness effect on charge conduction. Various responsible charge conduction mechanism for temperature dependent I-V behaviour have studied by Gadani et al. [23].

Among the all-manganite structures, GdMnO₃ based structures are not far studied as compared to other manganites. There are few reports are available for single layer GdMnO₃ thin films. Andreev *et al.* [24] have fabricated GdMnO₃/NdGaO₃ thin film to investigate the magnetic phase transition. Romaguera *et al.* [25] have studied the Raman and magnetic properties of GdMnO₃ thin film grown on Pt substrate. Negi *et al.* [26] reported the weal ferromagnetic phase at lower temperature in PLD grown GdMnO₃/Si thin film. Hitherto no reports available for GdMnO₃ based multilayers/heterostructures.

A noticeable amount of work has carrier out to study the I-V characteristic and resistive switching behavior of rhombohedral BiFeO₃/ZnO based heterostructures [**27-30**], GdMnO₃/Al:ZnO/SNTO heterostructure exhibits a good resistive switching behaviour [**31**], it will be very interesting to study the I-V behaviour and the effect of temperature on the electrical transport characteristics of the GdMnO₃/ZnO (Al:ZnO) bilayers fabricated on SrTiO₃ substrate.

In present communication, the results of the studies on temperature effect on transport behaviour have been reported for $GdMnO_3/Al:ZnO/SrTiO_3$ and $GdMnO_3/ZnO/SrTiO_3$ manganite-based bilayer thin film heterostructures. Herein, manganite-semiconductor structures were fabricated using Pulsed Laser Deposition (PLD) system. We investigated the structural, topographical, vibrational and transport properties of $GdMnO_3/Al:ZnO/SrTiO_3$ and $GdMnO_3/Al:ZnO/SrTiO_3$ bilayer thin films, especially focused on the transport behaviour of both films. To understand transport mechanism some theoretical models have been used.

EXPERIMENTAL DETAILS

Firstly, all the required targets (GdMnO₃, ZnO and 2% Al doped ZnO) were prepared using high temperature SSR route. Then, Al:ZnO/ZnO semiconductor layers were fabricated on (*100*) single crystalline SrTiO₃ substrate, then after masking GdMnO₃ manganite layer was fabricated on



Al:ZnO/SrTiO₃ and ZnO/SrTiO₃ films using Pulsed Laser Deposition system (parameters: laser energy: -~ 225 mJ and ~240 mJ, Pulse rate: - 5 Hz, substrate temperature: - 800°C, partial oxygen pressure: - ~ 250 mTarr and ~75 mTarr for GdMnO₃ and ZnO/Al-ZnO layers respectively). The X-ray Diffraction pattern were taken using Cu Ka radiation for structural analysis of prepared thin films. For better understanding phase identification, X-ray Diffraction Pattern were taken with grazing angle mode. For topographical analysis, Atomic Force Micrographs were taken using Nanoscop V-3100 (Bruker). For vibrational mode identification, the Raman spectra were obtained over a range from 100 cm⁻¹ to 1200 cm⁻¹ with a spectral resolution of 1 cm⁻¹ at an excitation wavelength of 785 nm, using A Renishaw confocal micro-Raman spectrometer with a liquid-nitrogen-cooled charge-coupled device was used to analyze the backscattered light. The laser spot power incident on the sample was kept below 5mW to avoid any sample heating. The temperature dependent Current-Voltage (I-V) measurements were taken to investigate the transport behaviour of all fabricated films using the Keithley Source Metter (Model- 4200SCS). Current-Voltage (I-V) measurements were taken using the DC two probe method with Current Perpendicular to plane (CPP) mode in voltage range of -5 V to +5 V for both heterostructures for this Ag (silver) patches formed on the surface of both the GMO manganite layer and the AZO/ZO layer.



Fig. 1. X-ray diffraction patterns for polycrystalline (a) $GdMnO_3/Al:ZnO/SrTiO_3$ and (b) $GdMnO_3/ZnO/SrTiO_3$ thin film heterostructures taken at 1.5° grazing angle.

RESULTS AND DISCUSSIONS

X-ray Diffraction (XRD) Measurements

Fig. 1 displays the X-ray diffraction patterns for GdMnO₃/Al:ZnO/SrTiO₃ (GMO/AZ/STO) and GdMnO₃/ZnO/SrTiO₃(GMO/Z/STO). X-ray Diffraction

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pattern taken at 1.5° grazing angle to confirm the polycrystalline growth of all proposed layers. The X-ray Diffraction patterns confirm the orthorhombic (*Pnma*) growth of GdMnO₃ and wurtzite hexagonal phase growth of Al:ZnO and ZnO layers. The tensile strain (+0.07%) and compressive strain (-0.15%) were present at interface of GdMnO₃/Al:ZnO and GdMnO₃/ZnO bilayers respectively. The lattice mismatch/strain calculated using following relation,

$$\delta(\%) = \left[\frac{(d_{substrate} - d_{film})}{d_{film}}\right] \times 100 \tag{1}$$

Atomic Force Microscopy (AFM) measurements

Atomic Force Microscopy (AFM) is the technique which helps to study the surface topography of thin films. **Fig. 2 (a-h)** reveals the surface topographic images with 3-dimentional (Fig. 2(a) & Fig. 2(e)) and 2-dimentional (Fig. 2(b) & Fig. 2(f)) view as well as surface profile (Fig. 2(c) & Fig. 2(g)) and RMS Roughness (Fig. 2(d) & Fig. 2(h)) studies for prepared GMO/AZ/STO and GMO/Z/STO thin films respectively.

Island type growth for both the strictures is confirmed from the 3D view of surface, GMO/AZ/STO possesses large islands as compared to GMO/Z/STO film. Also, Zaxis scale bar reveals the large difference in grain height for GMO/AZ/STO (~236 nm) and GMO/Z/STO (~70.8 nm) thin films. Surface profile suggests homogeneous grain growth with ~1.93 µm and ~172 nm average grain size for both films respectively as well as RMS roughness found to be ~ 36.7075 (GMO/AZ/STO) and ~ 7.1297 (GMO/Z/STO) respectively. The difference in microstructure of both the layered films helps to understand the difference in Current-Voltage relation of both the structures.



Fig. 2. AFM Images for GMO/AZ/SNTO and GMO/Z/SNTO heterostructures; (a & e) 2D view surface topography, (b & f) 3D view of surface topography with Z- axis scale bar, (c & g) granular (surface) profile for randomly selected grains, (d & h) average RMS roughness analysis respectively.

Raman measurements

Raman spectroscopy studies related vibrations of atomic bonds present in structure, all vibrations are associate with significant phonon modes. For the orthorhombic structure, 60 phonon vibrational modes are associated with Γ -point among them 24 modes (7A g+5B 1g+7B 2g+5B 3g) Raman active, 25 modes (9B 1u+7B 2u+9B 3u) infrared active, 8 modes (8A u) silent mode, and 3 are acoustic modes (3B 1u+B 2u+B 3u) respectively. Two MnO₆ octahedra were found in the GdMnO₃ crystal lattice, as well as two axial O2 and four equatorial O1 ions on the y-axis. The xy-plane and y-axis are parallel to the Mn-O1 and Mn-O2 bonds, respectively. **Fig. 3(a)** and **Fig. 3(b)** represents the typical room temperature Raman spectra for GMO/AZ/STO and GMO/Z/STO thin films respectively. The Raman shifts were identified using the Lorentz function fitted on experimentally observed spectrum. As shown in **Fig. 3(a)** and **Fig. 3(b)**, Raman spectra reveals strong Raman Shifts 245.48 cm⁻¹, 273.98 cm⁻¹, 374.98 cm⁻¹,415.33 cm⁻¹, 486.56 cm⁻¹, 504.41 cm⁻¹, 616.06 cm⁻¹, 679.01 cm⁻¹ and 728.89 cm⁻¹ as well as 243.10 cm⁻¹, 275.17 cm⁻¹, 374.98 cm⁻¹, 411.76 cm⁻¹, 486.59 cm⁻¹, 505.60 cm⁻¹, 618.43 cm⁻¹, 677.82 cm⁻¹ and 726.52 cm⁻¹ for GMO/AZ/STO and GMO/Z/STO thin films respectively, which shows significance with previously reported data [**32-34**]. The majority Raman shifts corresponds with

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GdMnO₃, only 245.48 cm⁻¹, 679.01 cm⁻¹ (for GMO/AZ/STO) and 243.10 cm⁻¹, 677.82 cm⁻¹ (for GMO/Z/STO) Raman Shifts are related with lattice-substrate interaction. From the close observation of the Raman spectra, one finds the five Raman active modes especially four A_g and one B_{2g} which is associated with MnO₆ octahedra present in GdMnO₃.



Fig. 3. Typical Micro-Raman Spectra for (a) GMO/AZ/STO and (b) GMO/Z/STO thin film heterostructures taken at room temperature.

The A_g symmetry which corresponds to tilting of MnO₆ octahedra reveals by 273.98 cm⁻¹ and 275.17 cm⁻¹. The Raman shift at 374.98 cm⁻¹ reveals the A_g mode associated with the out of phase *x* rotations of MnO₆. The A_g mode having MnO₆ bending and O2 anti-stretching were located at 486.56 cm⁻¹ and 504.41 cm⁻¹ as well as 486.59 cm⁻¹ and 505.60 cm⁻¹ respectively. The B_{2g} mode with the symmetric stretching mode involving the equatorial oxygen atoms is associated with the prominent band at 616.06 cm⁻¹ and 618.43 cm⁻¹ respectively.

The other modes, at 245.48 cm⁻¹, 679.01 cm⁻¹ (for GMO/AZ/STO) and 243.10 cm⁻¹, 677.82 cm⁻¹ (for GMO/Z/STO), possibly represent interactions between the GdMnO₃ films and the SrTiO₃ substrate [**25,34-36**]. Moreover, the Raman shift at 728.89 cm⁻¹ as well as 726.52 cm⁻¹ can be attributed to longitudinal optical (LO) branches of SrTiO₃ substrate [**37**]. It is observed that most Raman spectral features are approximately identical, although the intensity decreases. The Raman spectra of GdMnO₃/ZnO bilayers exhibit a broad band in the 380-480 cm⁻¹ region, which may attributed to the emergence of the E_2^{high} mode of Wurtzite ZnO structure [**38**].

Current-Voltage (I-V) measurements

Before delving into the transport studies, it is worth noting that Ag (silver) patches formed on the surface of both the GMO manganite layer and the AZO/ZO layers. In this study, these Ag patches served as electrodes for the Current-Voltage (I-V) behavior. During the experiment, these patches were made with the same size and thickness. The effects of the ohmic contact between the Ag electrode and all layers (for both films) on the transport properties of both films will be the same when analyzing the transport measurement results. In this case, changes in the results of transport characterizations are regarded as film behavior rather than any effective ohmic contact.



Fig. 4. Room temperature Current-Voltage (I-V) characteristics for GMO/AZ/STO and GMO/Z/STO thin film. (Insets: CPP geometry and Semi-logarithmic scale view)

To understand the transport behaviour across the Ag-GMO/AZ_Z-Ag interfaces of prepared GMO/AZ_Z/STO thin films, Current-Voltage (I-V) measurements were taken using the DC two probe method with Current Perpendicular to Plane (CPP) mode at room temperature. Room temperature I-V behaviour in range of -5 V to +5 V for both films are shown in Figure 4. The insets of Figure 4 represent schematic diagram of CPP mode as well as logarithmic plot of I-V behaviour of prepared films.

Fig. 4 reveals that both Thin film heterostructures possess rectifying behaviour with rectification ratio (I_F/I_R) of ~1.12 and ~0.98 at 5 V respectively, indicating majority charge carriers flow from semiconductor layer to manganite layer [39]. Herein one can consider that there is no major role of GdMnO₃ in charge transport behaviour in particular study deciding the rectification. Also, the filamentary paths created at the interface, which creates tunneling effect at the interface, are responsible for the increment in current and the rectification [31]. From Fig. 4, it is observed that GMO/AZ/STO film contains higher value of current compared to that in GMO/Z/STO, because Al:ZnO contains higher carrier concentration as compared to ZnO and for present case the charge carries are flowing from Al:ZnO and ZnO layer to GMO layer of GMO/AZ/STO and GMO/Z/STO films respectively which may be responsible for the high value of current for GMO/AZ/STO thin film heterostructures. Another factor affecting the charge on rectifying ratio is the microstructure presented in Fig. 2, in which the GMO layer in GMO/Z/STO films is having larger grains and hence smaller number of grain boundaries offering the small resistance to the current, whereas the smaller grains and larger number of grain boundaries offer comparatively large resistance for the case of GMO/Z/STO layered thin films.



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In context to understand the chare transport mechanism responsible for the observed room temperature Current-Voltage (I-V) behavior across the GMO/AZ and GMO/Z interface, the experimental data were fitted to various theoretical models, namely,

(i) Fowler-Nordheim (FN) Tunneling [40]

$$J_{FN} = AE^2 exp\left(\frac{-B^2\sqrt{\emptyset^3}}{E}\right)$$
(2)

where, A and B are constant; E is the applied electric field and \emptyset is potential barrier height.

(ii) Poole-Frenkel (PF) Emission [41]

$$J_{PF} = BE \ exp\left(\frac{-E_1 - \sqrt{q^3 \frac{E}{4\pi\epsilon_0 k}}}{K_B T}\right) \tag{3}$$

where *B* is a constant, *E* is the electric field applied to interfaces, E_1 is the trap ionization energy, *q* is electron charge K_B is Boltzmann's Constant, T is absolute temperature, ε_0 is the permittivity of free space and *k* is the optical dielectric constant of heterostructure.

(iii) Schottky Emission/Barrier [42],

$$J_{Schottky} = AT^2 exp\left(\frac{\emptyset - \sqrt{q^3 \frac{E}{4\pi\epsilon_0 k}}}{K_B T}\right)$$
(4)

where, *A* is a constant; *T* is absolute temperature; *E* is the electric field; k_B is Boltzmann's constant; \emptyset is the barrier height; ε_0 is the permittivity of free space and *k* is the optical dielectric permittivity of thin film

(iv) Space Charge Limited (SCL) Conduction [42],

$$J_{SCLC} = \frac{9\mu\varepsilon_0 k V^2 \theta}{8d_s^3} \exp\left[0.891\beta \left(\frac{V}{d_s}\right)^{1/2}\right]$$
(5)

where, μ is the mobility of electron, *k* is the permittivity, ε_0 is permittivity of free space, θ is free to total carrier density ratio, d_s is thickness of film, (v) Simmons Model [42] etc.

$$I = (\sigma \times V) + (k \times V^n) \tag{6}$$

where, σ represents the conductivity, k represents the constant which depending upon the barrier at interface.

Our obtained data best fits to two different theoretical models namely (i) Space Charge Limited (SCL) conduction and (ii) Simmons model. **Fig. 5(a)** & **Fig. 5(c)** show the Space Charge Limited Conduction (SCLC) fittings for Forward and Reverse bias of GMO/AZ/STO and GMO/Z/STO Thin film heterostructures respectively. Space Charge Limited Conduction (SCLC) model fitted for full voltage range which reveals that GMO/AZ and GMO/Z interfaces shows the ohmic behavior. To support the Space Charge Limited Conduction mechanism, linear fit for dV/d (*ln I*) vs I were performed. **Fig. 5(b)** & **Fig. 5(d)** provide the value of series resistance present at the interface of the film. The series resistance is dominant factor for the ohmic behavior of any junction [**43**]. The fitting parameters are tabulated in **Table 1**.



Fig. 5(e) and **Fig. 5(f)** show the Simmons model fitting for the GMO/AZ/STO and GMO/Z/STO thin filmns respectively. Fitting the Simmons moded on nonlinear curve, provides value of n which is associate with the different posible charge transport mechanism. If the value of $n \ge 0.6$ then charge conduction can be considerad tunneling through the disorderad metalic oxides. For the n = 1.33, quasiparticle tunneling via the pairs localized states. The stron spin-flip scattering at the insulating barries attirbute for chare conduction if n > 1.4. When n=2, direct tunneling [**42**]. In present cases, obtained value of n (0.72 \le n \le 1.33) indicate that the charge conduction is controlled by disordered metalic oxide (at interface due to lattice mismatch). Value of n is tabulated in **Table 1**.



Fig. 5. (a) & (c) Space Charge Limited (SCL) conduction, (b) & (d) Liner fits of dV/d(lnI) vs I linear, (e) & (f) Simmons model fittings to room temperature Current-Voltage (I-V) characteristics for GMO/AZ/STO and GMO/Z/STO thin film heterostructures respectively.

Table 1. Fitting parameters obtained from dV/d(lnI) vs I linear fitting and Simmons model.

		<i>dV/d(lnI) vs I</i> graph		Simmons Model Fitting
		Ideality Factor (η)	Series Resistance (Ω)	n
GMO/AZ/STO	FB	0.16	1.43	1.31
	RB	1.10	1.68	1.33
GMO/Z/STO	FB	0.02	7.36×10^{7}	0.72
	RB	0.03	7.18×10^{7}	0.84

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To understand the temperature effect on charge conduction at insulating- semiconductor interface, the temperature dependent Current-Voltage (I-V) measurements are being carried out. Figure 6 (a) and (b) show the I-V behavior obtained at various temperature (50°C, 100°C & 150°C) for GMO/AZ/STO and GMO/Z/STO thin film heterostructures respectively and insets of Fig. 6(a) & Fig. 6(b) shows current variation at lower voltage. As shown in Fig. 6(a) & Fig. 6(b), as temperature increases the value of current is increases for both bilayers. The increment in current with temperature can be describes in two different ways, (i) The thermal injection charge carries [39] and (ii) internal annealing [44] at the interface respectively.



Fig. 6. Temperature dependent Current-Voltage (I-V) characteristics for (a) GMO/AZ/STO (b) GMO/Z/STO thin film heterostructures (at 50°C, 100°C & 150°C). (Inset: enlarged view of lower voltage region).

The thermal injection of charge carrier can be predicted because of the partial substitution of Zn⁺² by Mn⁺³ in GMO/AZ and GMO/Z lattices of the thin film heterostructures which increase the carrier concentration at the interface as temperature increases [45]. Khushal et al. [46] also reported the partial substitution or effective diffusion of Zn⁺² in LSMO manganite at interface that results in La_{0.7}Sr_{0.3}Mn_{1-y}Zn_yO₃ form of stoichiometry. In thin films, the electrical resistance strongly obeys the annealing temperature in context of grain boundaries. The increment in temperature causes the partial annealing at the interfaces which causes decrement in grain boundaries, resulting observed increment in current due to increment in crystallinity which decrease the electrical resistance of thin films at interfaces. To convey the fact of temperature effect on charge conduction, the modified Thermionic Emission model were fitted for I-V data collected at different temperature (50°C, 100°C & 150°C) using the relation,

$$I = AT^{2} exp\left(-\left[\frac{\left\{\Phi_{B} - q\left(\frac{qV}{4\pi\varepsilon d}\right)^{1/2}\right\}}{\kappa T}\right]\right)$$
(7)

where Φ_B is the barrier height, q is the charge of carrier, ε is the permittivity of material, d is the barrier width and K is Boltzmann constant. Fig. 7(a) and Fig. 7(b) represent the theoretically fitted experimental I-V data at different temperature (50°C, 100°C & 150°C). Form the fitting, there is a decrement in Φ_B value (tabulated in Table 2) with increment in temperature for both thin film

Adv. Mater. Lett., | Issue (July-September) 2023, 23031730

heterostructures, suggesting that with temperature crystallinity increases (decreasing lattice mismatch) at the interface which in turn increase the charge conduction due to the thermal excitation effect on the charge carriers [47]. In other words, the interface of heterostructures is modified with temperature and the discontinuity at manganite-semiconductor interface decreases [48].



Fig. 7. Thermionic emission model fitting for Temperature Dependent Current-Voltage (I-V) characteristics for (a) GMO/AZ/STO and (b) GMO/Z/STO thin film heterostructures (at 50°C, 100°C & 150°C). Variation of barrier height with temperature for (c) GMO/AZ/STO and (d) GMO/Z/STO thin film heterostructures.

Table 2. Fitting parameters from Thermionic emission model.

	Temperature		Thermionic Model
			$\Phi_B(eV)$
GMO/AZ/STO	50 °C	FB	8.85×10^{-4}
	30 C	RB	5.50×10^{-4}
	100°C	FB	2.59×10^{-5}
	150°C	RB	1.70×10^{-5}
		FB	1.16×10^{-5}
		RB	5.85×10^{-6}
GMO/Z/STO	50 °C	FB	1.23×10^{-4}
		RB	1.43×10^{-4}
	100°C	FB	1.01×10^{-5}
		RB	8.10×10^{-5}
	150°C	FB	8.35×10^{-6}
		RB	5.48×10^{-5}

CONCLUSION

In conclusion, we successfully fabricated the polycrystalline GMO/AZ/STO and GMO/Z/STO thin film heterostructures using Pulsed Laser Deposition System. The growth of orthorhombic GMO layer and Wurtzite hexagonal Al:ZnO/ZnO structure was confirmed by X-ray diffraction patterns. The AFM images reveals the topographical behaviour of prepared films. From AFM analysis, found that GMO/AZ/STO film possess larger grain size as compared to GMO/Z/STO film. The Raman spectra confirms the bending and stretching of MnO₆ octahedra presence in GdMnO3 layer. The transport

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behaviour of prepared films understood with Current-Voltage (I-V) behaviour. Both films show rectifying behaviour at room temperature. From the I-V measurements one found that GMO/AZ/STO shows higher value of current because it contains more charge carriers as compared to GMO/Z/STO films. For better understanding of room temperature conduction mechanism, Space Charge Limited Conduction (SCLC) and Simmons model were fitted. SCLC fitting reveals the ohmic behaviour of prepared films which rather supported by dV/d(lnI) vs I graphs and provides value of series resistance for both films. Simmons model suggested that charge conduction attributed with tunneling though disordered metallic oxides. Temperature dependent I-V behaviour shows that increment in temperature increases current and shows rectifying nature at high temperature. The increment in current at comparatively high temperatures attributed as partial annealing effect at the interface. Thermionic emission model was employed to understand the temperature effect at interface, reveals that with increasing temperature barrier height decreases, promoting the charge conduction. The obtained results show that interfacial coupling between manganite and semiconductor materials could be controlled by applied voltages and temperatures that immensely improves the promising applications of manganite/semiconductor heterostructures in multifunctional electronic devices.

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DATA AVAILABILITY

The datasets generated during and/or analysed during the current study are available from the corresponding author on reasonable request.

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