


Cite this: *RSC Adv.*, 2024, 14, 21692

Study of the correlation between the magnetic and electrical properties of the $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ compound†

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In this work, we investigated the relationship between the electrical and magnetic properties of the superparamagnetic (SPM) $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0) compound prepared by the sol–gel method. The (S1C0) sample displayed a ferromagnetic metallic (FMM) behavior at low temperatures and a paramagnetic semiconductor (PMSC) behavior at high temperatures. The FMM behavior was described by the Zener Double Exchange (ZDE) polynomial law containing the contributions of the electron–electron (e–e) interactions and the electron–magnon (e–m) scattering. The PMSC behavior was described by the Mott Variable Range Hopping (Mott–VRH) transport model. The semiconductor/metallic transition temperature has been approximated at the blocking temperature. The Thermal Coefficient of Resistivity (TCR), which exhibits a linear variation around ambient temperature, can be used as a calibration curve for thermometry. Thus, our sample can be considered as a good candidate for the detection of infrared radiation used in night vision bolometer technologies.

Received 13th May 2024
Accepted 25th June 2024
DOI: 10.1039/d4ra03528c
rsc.li/rsc-advances

1. Introduction

The manganites of perovskite structure, having the general formula $\text{RE}_{1-x}\text{AE}_x\text{MnO}_3$ (RE = trivalent Rare-Earth, AE = bivalent Alkaline Earth Ca, Sr and Ba), offer a high degree of chemical flexibility with a complex interplay between structure and the electronic and magnetic properties. This leads to a very rich phase diagram involving various metallic, insulating and magnetic phases.^{1–7} At high temperature, the paramagnetic (PM) phase has been detected. However, at low temperatures, both antiferromagnetic (AFM) and ferromagnetic (FM) phases have been observed. The FM phase can be either insulating for $x < 0.2$, or metallic for $0.2 < x < 0.45$. For the higher concentration values ($0.45 < x < 1$), the ground state was found to be AFM and nonmetallic.⁸ In this type of compounds, manganese presents the $\text{Mn}^{3+}/\text{Mn}^{4+}$ mixed valence, which is controlled by the rate of substitution x . This valence is considered as a crucial parameter controlling the structural, electrical and magnetic properties of manganites. Jonker and Santen⁹ showed that, in this type of

compounds, the FM properties can be understood as the result of a strong positive $\text{Mn}^{3+}/\text{Mn}^{4+}$ exchange interaction combined with a weak $\text{Mn}^{3+}/\text{Mn}^{3+}$ interaction and a negative $\text{Mn}^{4+}/\text{Mn}^{4+}$ interaction. These mixed valence manganites have attracted a lot of attention, since the middle of the twentieth century, due to their potential magnetic applications using their high magnetoresistance effect (MR) defined as a decrease of the resistivity when a magnetic field is applied.^{10,11} Also, these manganites are considered of great interest due to their specific features, such as metal–insulator transition, ferromagnetic–(FM–PM) phase transition, charge and orbital ordering and lattice degree of freedom.^{12,13} These materials are found to exhibit a ferromagnetic-metal (FMM) to paramagnetic-insulator (PMI) transition near the Curie temperature T_C .

From the last decade of the 20th century, numerous studies have been published on the magnetic and magnetoresistive properties of lanthanum manganites. The $\text{La}_{1-x}\text{Sr}_x\text{MnO}_3$ compounds are among those which have been the subject of the greatest number of publications concerning the properties of substituted lanthanum manganites.^{14–18} In fact, the phase diagram of $\text{La}_{1-x}\text{Sr}_x\text{MnO}_3$ (ref. 16) show that these compounds can exist in different magnetic and electric phases according to the value of the concentration (x) of the AE bivalent element.

Magnetic nanoparticles display physical and chemical properties different to those found in their corresponding bulk materials. These properties make them attractive in widespread applications, such as energy, electronics, sensor designs of all kinds, catalysts, magnetic refrigeration, optics and in various biomedical applications.^{17–19}

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† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d4ra03528c>



For the reasons previously mentioned, we will study in this paper, the correlation between the electrical and magnetic properties of super-paramagnetic (SPM) nanoparticles $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0), who exhibits the highest FM/PM temperature transition of the $\text{La}_{1-x}\text{Sr}_x\text{MnO}_3$ series.

2. Experiment

2.1. Sample preparation

The polycrystalline $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0) sample was prepared by the citric-gel method using $\text{La}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$, $\text{Sr}(\text{NO}_3)_2$ and $\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}$ as precursors. These precursors were dissolved in deionized water in a separate beaker and mixed with the solution of citric acid and ethylene glycol, at constant stirring, in the ratio of $(1[\text{Mn}(\text{NO}_3)_2 \cdot 4\text{H}_2\text{O}]/1[\text{citric acid}]/\frac{2}{3}[\text{ethylene glycol}])$ to get a transparent stable solution. The obtained solution was then heated at 80 °C to eliminate excess water. The resulting solution was converted into viscous glassy gels and dried at 120 °C to get dark grayish products. In order to ensure the decomposition of nitrate/citrate complexes, the organic components and the residual organic compounds of the precursors, the obtained products were ground, calcined at 700 °C for 12 h and converted into pellets. Finally, these pellets were annealed at 900 °C for 18 h to get the desired nanoparticles manganite.²⁰

2.2. Characterizations

The morphology of the surface and fracture surfaces of the sample as well as an elemental analysis by X-ray microanalysis (Energy Dispersion Spectroscopy EDS) were carried out by Transmission Electron Microscopy (TEM) at the Beni-Mellal Materials Physics Laboratory of the Faculty of Science and Technology; Sultan Moulay Slimane University (Morocco). A small amount of the sample was dissolved in ethanol and then exposed to ultrasound to ensure good dispersion. A drop is then deposited on a copper grid, which serves as a sample holder. Finally, the drop is dried by a lamp in order to evaporate the ethanol. Surfaces were covered with carbon before microscopic observation.

The crystalline structure and phase purity of the prepared sample were examined by powder X-ray Diffraction (XRD) with Cu K α radiation ($\lambda = 1.5406 \text{ \AA}$), at room temperature, by a step scanning of 0.016° in the range of $20^\circ \leq 2\theta \leq 90^\circ$. The data were analyzed by the Rietveld method using the Fullproof SOFTWARE.^{21,22}

Magnetic measurements of magnetization (M) as a function of temperature (T), under different values of an applied magnetic field ($\mu_0 H$), have been carried out with the BS1 and BS2 developed at the Louis Néel Laboratory in Grenoble. The isothermal magnetization curves $M(\mu_0 H)$, at a temperature of 5 K, were carried out under a magnetic field between 0 and 10 T. The hysteresis cycles, at different temperatures below the Curie temperature and at magnetic fields between (-10 T) and (10 T), were performed.

The temperature dependence of electrical resistance was measured with standard four-probe method in a commercial Physical Property Measurement System (Quantum Design PPMS) under different magnetic field.

3. Results and discussion

3.1. Purity, homogeneity and structure

The elemental composition and the homogeneity of our (S1C0) sample were studied by X-ray microanalysis (Energy Dispersion Spectroscopy: EDS). Fig. 1(a) shows the presence of the characteristic peaks of the La, Sr, Mn and O chemical elements introduced during the elaboration. These elements have practically the same proportions at different locations,⁴ which confirms the homogeneity of the sample. The characteristic copper (Cu) peaks correspond to the sample holders, while the carbon is used to prepare the sample for observation.

In order to study the grain size distribution, we have used the images provided by (TEM). Fig. 1(b) shows an example of the top view of the surface morphologies of (S1C0) sample. A manual statistical count of the grain size was performed on this image using the Image-J software. The results, showed in Fig. 1(c) as grain number (counts) *versus* grain size (nm), reveal a dispersion in the grain size following a Lorentzian law with a most common size around 78 nm and an average value of 96 nm. The goodness of fit was assessed by the square of the linear correlation coefficient (R^2). The obtained value of (R^2) reaches 99.9%.

The refinement of the structure as well as the lattice parameters were carried out by the Rietveld method,¹⁷ using the “Full Prof” program.¹⁸ As mentioned in our previous work,²⁰ the Rietveld refinements of the XRD data revealed that our sample is indexed in the orthorhombic structure with the $Pbnm$ space group. The Rietveld refinement of the XRD Pattern is presented in Fig. 1(d) showing the most intense reflections (hkl) peaks. The related refined structural parameters are given in Table S1 (seeESI).†

The crystallite size (D_{SC}) and the micro strain (ε) were determined, using the following relationships:^{23–26}

$$D_{\text{SC}} = \frac{0.9 \times \lambda}{\beta \times \cos(\theta)} \quad (1)$$

$$\varepsilon = \frac{\beta}{4 \times \tan(\theta)} \quad (2)$$

where λ is the used wavelength ($\lambda = 1.5406 \text{ \AA}$), $2\theta = 32.704^\circ$ is the position of the most intense peak (112) reflection, and $\beta = 0.30232^\circ$ is the full width at half maximum (FWHM) of the same peak.

The obtained values are then $D_{\text{SC}} = 27 \text{ nm}$ and $\varepsilon = 4.50 \times 10^{-3}$.^{20,27}

The agglomeration rate, defined as the ratio of the average grain size to that of the crystallites, is about 3.59. This can explain the relatively high value of micro strain compared to those obtained with thin films, in TiSiN, having micro strain varying from 1.6×10^{-3} to 2.58×10^{-3} , as mentioned by Bushroa *et al.*²⁸

3.2. Magnetic properties

Fig. 2(a) shows the variation of magnetization as a function of temperature in Zero-Field-Cooled (ZFC) and Field-Cooled (FC) modes, under an applied magnetic field of 0.05 T. From the FC



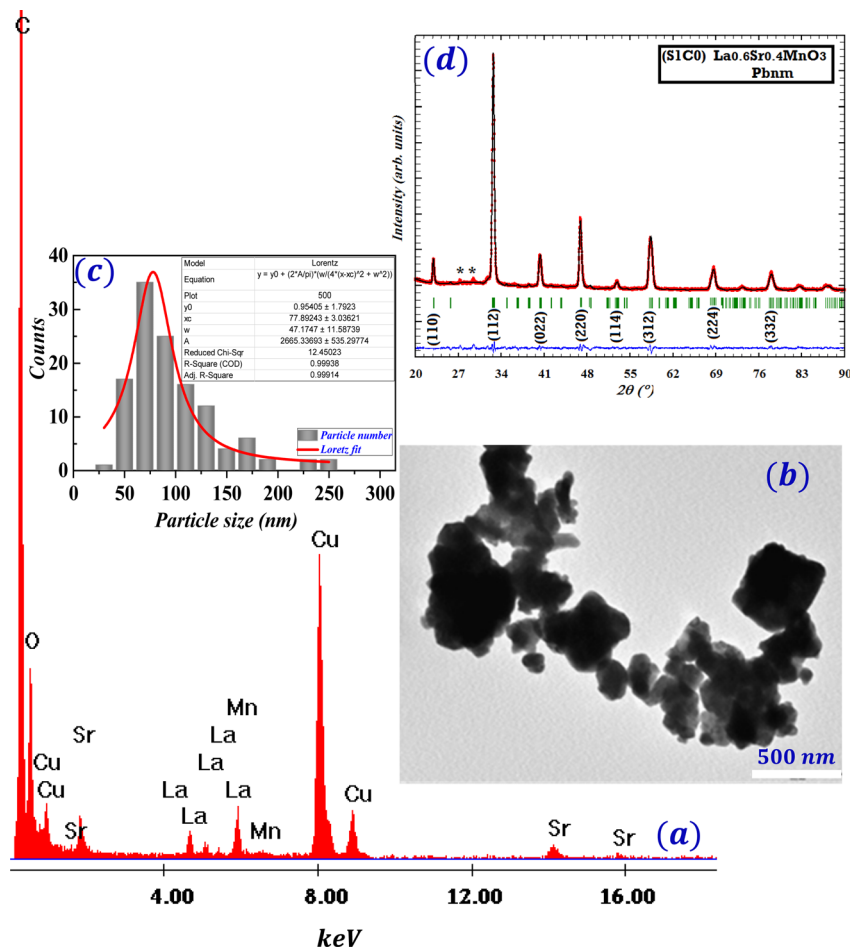


Fig. 1 (a) X-ray microanalysis (Energy Dispersion Spectroscopy: EDS). (b) TEM micrograph of particle size. (c) The dispersion in the number of grains (counts) according to their size (nm). (d) X-ray diffraction pattern: The observed (red circle), calculated (continuous black line) and difference patterns (blue line at the bottom) of X-ray diffraction data. Asterisks show the presence of the minority secondary phase Mn₂O₃.

mode, it is clear that (S1C0) sample exhibits a PM behavior at high temperatures and a FM order at low temperatures. This FM behavior is confirmed by the evolution, at 5 K, of the magnetization as a function of the applied magnetic field, in the inset of Fig. 2(a). In fact, the magnetization increases abruptly by application of weak fields and saturates for fields greater than 1 T. The transition temperature T_C , called Curie temperature, corresponding to the minimum of the first derivative dM_{FC}/dT was found to be equal to $T_C = 365$ K.

The irreversibility between the $M(T)$ curves in FC and ZFC modes, below a so-called irreversibility temperature $T_i \approx 350$ K, indicates the presence of a magnetic anisotropy within (S1C0) sample.^{27,29} The ZFC curve describes a broad peak centered on the blocking temperature $T_B = 195$ K. This magnetic anisotropy is observed with nanoparticles, having the stable monodomain configuration, where the magnetization has a preferential direction.³⁰ This magnetization can reverse spontaneously under the effect of thermal agitation or magnetic field. This regime of magnetic fluctuations is quite simply the paramagnetism of the macro spin of the monodomain and it is thus called superparamagnetism (SPM). For temperatures below the blocking temperature (T_B), the magnetic anisotropy energy

outweighs the thermal agitation energy, which explain the increase of the magnetization when the temperature rises. For temperatures above the blocking temperature, the thermal agitation energy predominates leading to the usual decrease in magnetization when increasing the temperature, characteristic of the SPM phenomenon. These observations are confirmed by the variation of the inverse of the magnetic susceptibility (χ^{-1}) as a function of the temperature (T), represented in Fig. S1 (see ESI).[†] The $\chi^{-1}(T)$ curve presents two linear branches. The first one is fitted with the Curie–Weiss law,³¹ indicating the PM–FM transition at $\theta_{WC} = 368$ K value quite equal to the Curie temperature $T_C \approx 365$ K. On the other hand, the linear adjustment of the second branch, shows the existence of another temperature characteristic of the magnetic properties $T_i \approx 350$ K. This indicates the existence of (SPM) clusters within the (FM) phase in the temperature interval between T_B and T_C . So, the irreversibility temperature T_i can be defined as the temperature of a SPM/PM transition.

In order to confirm the single-domain configuration and the existence of (SPM) clusters in the (FM) phase in the temperature range between T_B and T_C , hysteresis cycles, for magnetic fields ranging from -10 T to 10 T at different temperatures, were



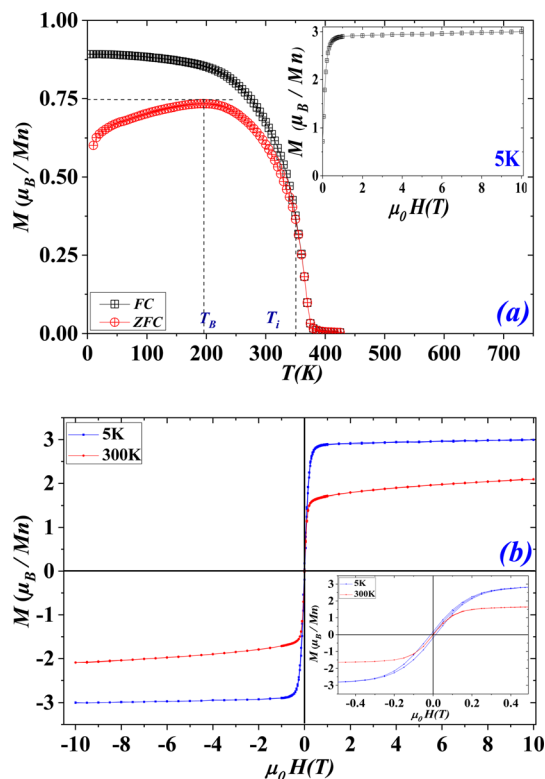


Fig. 2 (a) The temperature dependence of the magnetization measured under a magnetic field of 0.05 T, in the ZFC and FC modes. In the insert, the applied magnetic field-dependence of the magnetization measured at temperature of 5 K. (b) The complete $M(\mu_0H)$ hysteresis loop for S1C0 at 5 and 300 K. The inset of (b) a zoom of the hysteresis cycles for weak magnetic fields.

performed in Fig. 2(b). These curves clearly indicate very low hysteresis losses for very low temperatures, which justifies the monodomain configuration of our sample. The inset of Fig. 2(b) shows that, when the temperature increases between T_B and T_C , the remanent magnetization and the coercive field decrease and become zero, confirming the presence of a SPM behavior. These results are similar to those obtained by Manh *et al.*,³⁰ revealing that nanometric particles, of $\text{La}_{0.7}\text{Sr}_{0.3}\text{MnO}_3$ solution sintered at 700 °C, are the site of a superparamagnetism phenomenon.

3.3. Magneto-transport study

The temperature dependence of electrical resistance $R(T)$ (proportional to the resistivity $\rho(T)$) of the (S1C0) sample is shown in Fig. 3(a), for the temperature range of 100 to 500 K and under different applied magnetic fields of 0, 2 and 5 T. By decreasing temperature, (S1C0) sample exhibits a single insulator (or semiconductor)–metallic (I–M) transition at T_R . This transition temperature is found to increase from 200 K in the absence of the magnetic field, to 235 K for 5 T. We can also note the decrease in resistance by the increase of the magnetic field (Table 1) revealing that a magnetic field promotes the electrical conduction. This transition is the result of the competition between the double exchange (DE) mechanism, favoring the FM interaction, and the super exchange (SE) one, approving the AFM interaction.

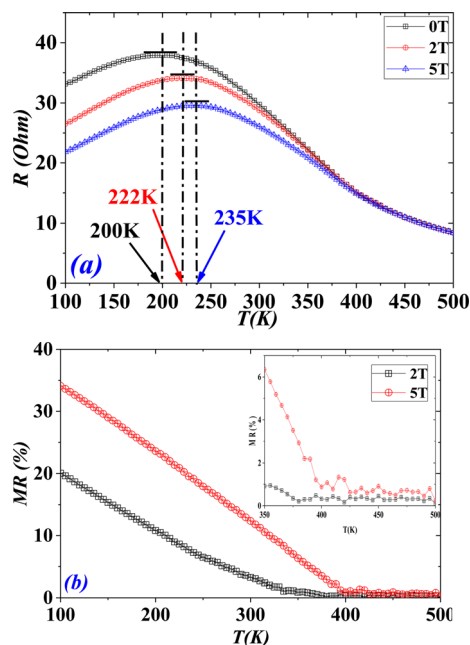


Fig. 3 (a) Temperature dependence of resistance $R(T)$ for (S1C0) under different applied magnetic fields of 0, 2 and 5 T. (b) The variation of $\text{MR}\% (T, \mu_0H)$, as a function of temperature for 2 T and 5 T.

Table 1 Summary of the magneto transport properties of $\text{La}_{0.6}\text{Sr}_{0.4}\text{-MnO}_3$ (S1C0), for different values of the magnetic field 0, 2 and 5 T

μ_0H (T)	0	2	5
T_R (K)	200	222	235
R_{max} (Ω)	37.907	34.266	29.632
$R_{\text{max}}(\mu_0H)$ (%)	100	90.39	78.17
$R_{\text{max}}(0 \text{ T})$			
MR_{max} (%)		20.1	34.1
$T(\text{MR}_{\text{max}})$ (K)		100	100
$100 \times \text{TCR} (\text{K}^{-1}) _{\text{max}}$	0.88	0.84	0.75
$T(\text{TCR} _{\text{max}})$ (K)	370	370	370

The (DE) model introduced by Zener^{32,33} and taken up by Anderson and De Gennes^{34,35} aims to explain the FMM behavior of certain doped compounds.³⁶ This coupling considers the hop of electrons e_g between $\text{Mn}^{3+} (3t_{2g}1e_g)$ and $\text{Mn}^{4+} (3t_{2g})$ ions, using oxygen as the jump intermediate. Thus, an electron e_g from Mn^{3+} jumps to the O^{2-} ion, at the same time as an electron from the O^{2-} ion jumps to the Mn^{4+} cation. The hop probability of the electrons e_g being governed by the scalar product between the core spins of the electrons t_{2g} of the Mn^{3+} and Mn^{4+} ions in order to satisfy Hund's rule. As a result, the hop of an electron e_g will be all the easier if the spins of the two considered Mn^{3+} and Mn^{4+} ions are parallels. This leads to a very strong parallel between magnetism and transport properties, where ferromagnetism favoring electronic conduction and *vice versa*.

The (SE) theory was developed by several physicists as early as 1934 by Kramers,³⁷ Anderson,^{38–40} Van Vleck⁴¹ and Goodenough.^{42,43} The super exchange, explained by the Goodenough–

Kanamori rules for a Mn–O–Mn bond,⁴⁴ makes it possible to explain, among other things, the AFM coupling between the manganese ions $\text{Mn}^{3+}\text{--O}^{2-}\text{--Mn}^{3+}$. In fact, the e_g orbitals of Mn^{3+} cations each have one electron. Electrons placed on the 2p orbitals of oxygen can occupy the e_g orbitals, provided that each of their spins is in an antiparallel configuration with respect to the e_g electrons, and this in accordance with the Pauli principle. We can therefore predict AFM and strong $\text{Mn}^{3+}\text{--O}^{2-}\text{--Mn}^{3+}$ interactions due to the strong hybridization of the e_g orbitals. SE theory assumes localized electrons. Therefore, a compound in which the magnetic structure is dominated by SE shows, at the level of electrical conductivity, the behavior of an insulator. These explanations are perfectly illustrated in Fig. S2(a)–(c) (see ESI),[†] where we have represented, on the same graph, the variations according to the temperature of the magnetization and the electrical resistance of (S1C0) sample. At low temperatures, the metallic behavior ($d\rho(T)/dT > 0$) is associated with a FM comportment. While the insulator behavior (or semiconductor) ($d\rho(T)/dT < 0$), at high temperatures, is associated with a magnetic disorder. It remains to be noted that the metal/insulator (M/I) transition temperature is significantly lower than that of the magnetic (FM/PM) one. The same comportment has been observed for many studies.^{19,45,46} In our case, $T_C \approx 365$ K for a magnetic field of 0.05 T and $T_R \approx 200$ K for a zero magnetic field. This last temperature is substantially equal to the Blocking temperature $T_B \approx 195$ K. This seems to be logical, since this temperature T_B is the limit of dominance of the magnetic anisotropy energy (for T lower than T_B) which favors the orientation of the macro spins along a preferential direction. Under these conditions, electrical conduction is favored. Hence the metallic behavior for T lower than T_B . For temperatures higher than T_B , it is the energy of thermal agitation which prevails, hence the fluctuation of the macro spins and the SPM behavior of the material for such temperatures. This explains the insulating (or semi-conducting) dependence of the material at high temperatures and hence the increase of the T_R temperature value with increasing the magnetic field ($\mu_0 H$).

The coexistence of the FM order and the metallic behavior, at low temperatures, causes a relative decrease of the resistance under the effect of a magnetic field known under the name of the magnetoresistance (MR) effect (Fig. 3(a)). This effect can be evaluated by the following relationship:^{1,3,45,46}

$$\text{MR}_{\%}(T, \mu_0 H) = 100 \times \frac{R(0, T) - R(\mu_0 H, T)}{R(0, T)} \quad (3)$$

where $R(0, T)$ and $R(\mu_0 H, T)$ are the resistances at zero magnetic field and at an applied magnetic field $\mu_0 H$, respectively. The variation of $\text{MR}_{\%}(T, \mu_0 H)$, as a function of temperature for 2 and 5 T, is shown in Fig. 3(b). It is clear that $\text{MR}_{\%}(T, \mu_0 H)$ does not exceed a few percent in the PM phase where our (S1C0) sample behaves like an insulator or a semiconductor. By decreasing the temperature, $\text{MR}_{\%}(T, \mu_0 H)$ increases to reach its maximum, which is about 20.1% at 100 K for a magnetic field of 2 T and 34.1% at 100 K for a magnetic field of 5 T (Table 1).

These values and this behavior are similar to those obtained by Fu Yonglai and C. K. Ong,⁴⁷ in the bulk polycrystalline samples of $\text{La}_{0.7}\text{Ca}_{0.3-x}\text{Sr}_x\text{MnO}_3$ for $x = 0.05, 0.15$ and 0.2 . Also,

these characteristics are similar to those obtained by Y. K. Lakshmi and P. V. Reddy,⁴⁸ in the case of nanocrystalline $\text{La}_{0.7}\text{Ca}_{0.3-x}\text{Ag}_x\text{MnO}_3$, for $x = 0$ over the entire temperature range of the measurements, but different for $x = 0.1, 0.2$ and 0.3 , where we note the existence of a peak around the (M/I) transition temperature.

The MR observed at low temperatures, qualified as extrinsic MR (EMR), is due to the spin tunneling effect through the grain boundaries.^{49,50} However, the MR observed near the transition temperature T_R , qualified as intrinsic MR (IMR), takes place in the volume of the grain and is governed by the DE mechanism.^{32,51–53} A comparative study of the electrical properties of (S1C0) with those of the compound mentioned above as well as other compounds has been summarized in Table S2 (see ESI[†]).^{45–48,52–55}

To probe the sensitivity of infrared detector to temperature, in oxide-based manganites, a key requirement is to have materials with high thermal coefficient of resistivity (TCR) value, expressed by equation:

$$\text{TCR}(T) = \frac{1}{R(T)} \frac{dR(T)}{dT} \quad (4)$$

The (TCR) represents an important parameter for device applications concerning the infrared detectors (IR) for night vision technologies (bolometer).^{56–60} The temperature dependence of (TCR), for 0, 2 and 5 T, have been plotted in Fig. 4(a).

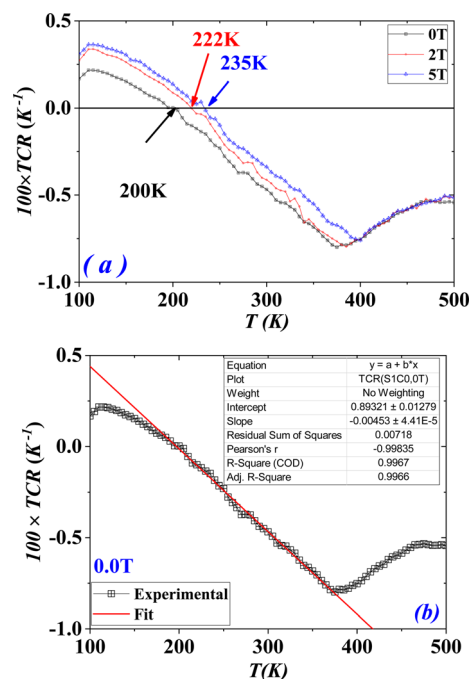


Fig. 4 (a) The temperature dependence of (TCR), for 0, 2 and 5 T. (b) The adjustment of the TCR, in the absence of a magnetic field, according to the temperature shows a linear evolution between 195 and 375 K. This curve can therefore be used to calibrate a measuring thermometer in the temperature range indicated above. Thus, our sample can be considered as a good candidate for the detection of infrared radiation used in night vision bolometer technologies.



Our (S1C0) sample is characterized by a positive (TCR) value for low temperatures (related to metallic behavior) and a negative (TCR) one for high temperatures (related to insulating or semiconductor behavior). We can remark that the (TCR) values, for our compound, are not very sensitive to the application of an external magnetic field (Fig. 4(a)). In fact, it's clear that the application of an external magnetic field leads to a slightly increase of the (TCR) value in the (FM) phase without any effect in the (PM) one. Over the entire measurement temperatures range, |TCR| does not exceed 1%. For a zero magnetic field, the maximums of |TCR| is obtained around the FM/PM transition Curie temperature: $|\text{TCR}|_{\text{peak}}$ (0 T, 375 K) = 0.0080 K⁻¹.

In the presence of a magnetic field, we obtain practically the same maximum value of $|\text{TCR}|_{\text{peak}}$, for FM/PM transition temperatures ($|\text{TCR}|_{\text{peak}}$ (2 T, 385 K) = 0.0080 K⁻¹ and $|\text{TCR}|_{\text{peak}}$ (5 T, 400 K) = 0.0076 K⁻¹) (Table 2) slightly higher than that obtained in the absence of the magnetic field. This is predictable, since the application of a magnetic field favors magnetic order in favor of magnetic disorder. We also note that the variation of the TCR, as a function of temperature, is practically linear over a wide temperature range around room temperature, whatever the magnetic field applied. For example, the adjustment of the TCR, in the absence of a magnetic field, according to the temperature shows a linear evolution between 195 and 375 K (Fig. 4(b)). This curve can therefore be used to calibrate a measuring thermometer in the temperature range indicated above. Thus, our sample can be considered as a good candidate for the detection of infrared radiation used in night vision bolometer technologies. A comparative study of the sensitivity, of our (S1C0) compound with other samples, to infrared radiation has been summarized in Table S2 (see ESI).†

3.4. Transport mechanism

To better understand the transport mechanism in the (S1C0) compound, we tried to test the validity of some theoretical model.

At low temperatures, the behavior is metallic and the resistivity is therefore interpreted in terms of elastic and inelastic scattering. The different diffusion mechanisms are generally temperature dependent with particular power laws. Thus, the resistivity is often described by a polynomial in T , of the form $\rho(T) = \rho_0 + \sum \rho_n T^n$, where ρ_0 is the residual resistivity, at $T = 0$ K, caused by grain boundaries.^{61,62} Based on previous work,^{63–69} we choose to work with the models described by the eqn (5) and (6) based on the Zener Double Exchange (ZDE) polynomial law and eqn (7) based on the Small Polaron Conduction Mechanism (SPCM):⁷⁰

$$\rho(T) = \rho_0 + \rho_2 T^2 + \rho_{4.5} T^{4.5} \quad (5)$$

$$\rho(T) = \rho_0 + \rho_2 T^2 + \rho_5 T^5 \quad (6)$$

$$\rho(T) = \rho_0 + \frac{E\omega_s}{\sinh^2\left(\frac{\hbar\omega_s}{2k_B T}\right)} + \rho_n T^n \quad (7)$$

where ρ_0 , ρ_2 , $\rho_{4.5}$ and ρ_5 are parameters, function of the applied magnetic field, which are expressed in Ω m, Ω m K⁻², Ω m K^{-9/2} and Ω m K⁻⁵, respectively, while $k_B = 1.38 \times 10^{-23}$ J s⁻¹ is one of the fundamental constants: it is the Boltzmann constant.

The variation in T^2 has been attributed to the electron–electron (e–e) interactions.^{63,64} The term $\rho_{4.5} T^{4.5}$ is caused by the contribution of the electron–magnon (e–m) scattering.^{65,66} However, the $\rho_5 T^5$ term is due to the electron–phonon (e–ph) interactions.^{67–69} Finally, in the (SPCM) model proposed by Alexandrov and Bratkovsky,⁷⁰ where E is a constant proportional to the effective mass of polarons; *i.e.*, the (e–ph) coupling strength; ω_s is the average frequency of the softest optical mode and ρ_n is the fitting coefficient.

The fittings of the experimental curves of resistance $R(T)$, with the eqn (5)–(7), show, in Fig. S3 (see ESI),† a good agreement between the experiment and the chosen model. The quality of these fittings was evaluated by the squared linear

Table 2 The best fit parameters, for La_{0.6}Sr_{0.4}MnO₃ (S1C0), obtained from the experimental resistance data of the metallic behavior

$\mu_0 H$ (T)	0	2	5
Model ①	$R(T) = R_0 + R_2 T^2 + R_{4.5} T^{4.5}$		
R_0 (Ω)	29.09	21.66	17.57
R_2 (Ω K ⁻²)	4.44×10^{-4}	5.22×10^{-4}	4.57×10^{-4}
$R_{4.5}$ (Ω K ^{-4.5})	-0.41×10^{-9}	-0.40×10^{-9}	-0.31×10^{-9}
Model ②	$R(T) = R_0 + R_2 T^2 + R_5 T^5$		
R_0 (Ω)	29.30	21.86	17.73
R_2 (Ω K ⁻²)	4.09×10^{-4}	4.88×10^{-4}	4.30×10^{-4}
R_5 (Ω K ⁻⁵)	-0.26×10^{-10}	-0.25×10^{-10}	-0.20×10^{-10}
Model ③	$R(T) = R_0 + \frac{E\omega_s}{\sinh^2\left(\frac{\hbar\omega_s}{2k_B T}\right)} + R_n T^n = R_0 + \frac{R_2}{\sinh^2\left(\frac{T_2}{T}\right)} + R_n T^n$		
R_0 (Ω)	16.66	8.53	6.39
R_2 (Ω K ⁻²)	93.78×10^4	30.72×10^4	3.05×10^4
T_2 (K)	25 182	8226	10 309
R_n (Ω K ⁻ⁿ)	2.17	1.48	1.02
n	0.44	0.54	0.54



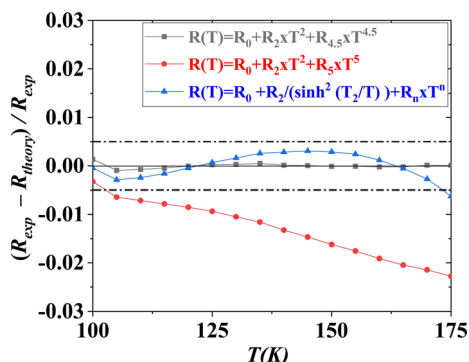


Fig. 5 The relative difference in resistance $\left(\frac{R_{\text{exp}} - R_{\text{theory}}}{R_{\text{exp}}}\right)$ vs. T plots for the three models under the zero applied magnetic field.

correlation coefficients (R^2). The obtained values of (R^2), for all field, were as high as 99.9%. Plots of the best fits to the data are shown by red solid lines in the main panels of Fig. 5 and the corresponding parameters are listed in Table 2. At first sight, the three models are suitably adapted to describe the

mechanism responsible for electrical conduction in the metallic region of our sample. To select the most appropriate model, the relative difference in resistance $\left(\frac{R_{\text{exp}} - R_{\text{theory}}}{R_{\text{exp}}}\right)$, in the absence of the magnetic field, was plotted in Fig. 5, using the adjustment parameters listed in Table 2. According to Fig. 5, the relative error for the (ZDE) polynomial, with (e-m) interactions, is the smallest. So, we can suggest that this model is the better one to describe the charge transport behavior in the metallic region.

The coefficients R_0 , R_2 and $R_{4.5}$, relating to this model, have different signs (Table 2). In fact, coefficients R_0 and R_2 have a positive contribution in the electrical resistance of the studied material while the $R_{4.5}$ one has a negative contribution. Fig. 6(a)–(c) show that the R_0 residual term presents the main contribution in the electrical resistance. The term $R_2 T^2$, relating to (e-e) interactions, comes second while the lowest contribution is that of the term $R_{4.5} T^{4.5}$, relating to (e-m) interactions. Fig. 6(d) shows that the residual term R_0 decreases with both temperature and the applied magnetic field. This decrease is for

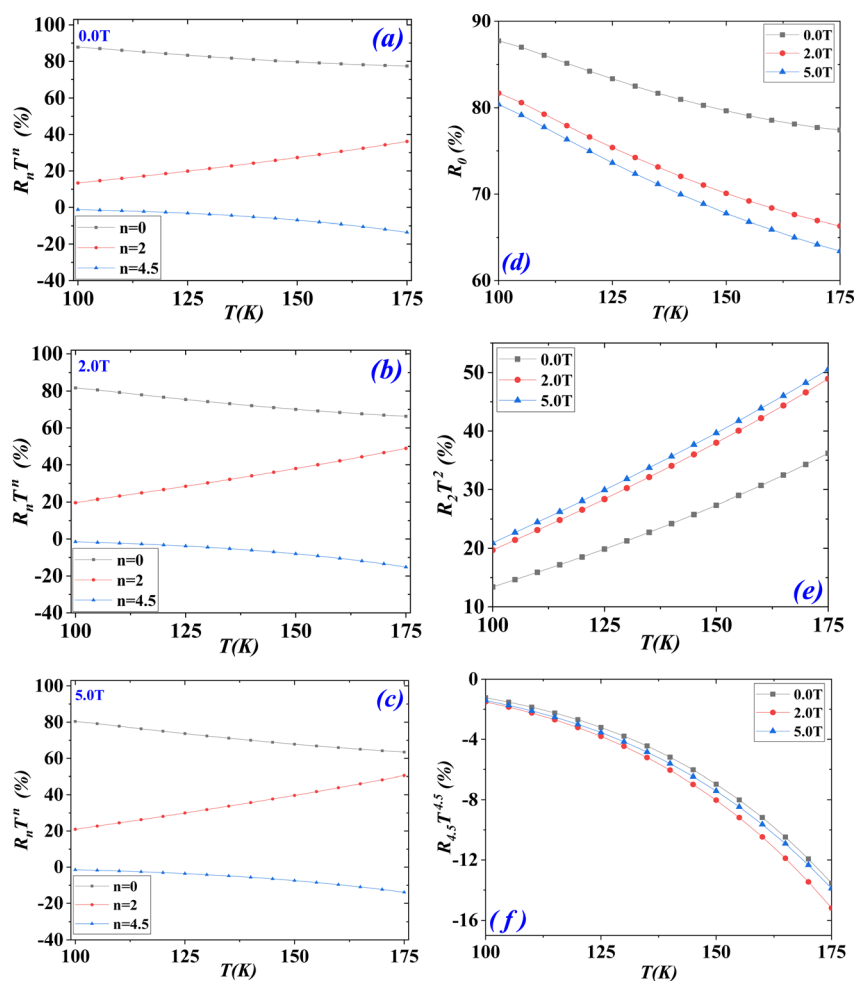


Fig. 6 Evolution, as a function of temperature, of the different terms and, of the (ZDE) polynomial model with (e-m) interactions, under different applied magnetic fields of 0, 2 and 5 T, respectively. (a)–(c) Applied magnetic field was maintained constant for different n values. (d)–(f) n value was maintained constant for different applied magnetic field values.



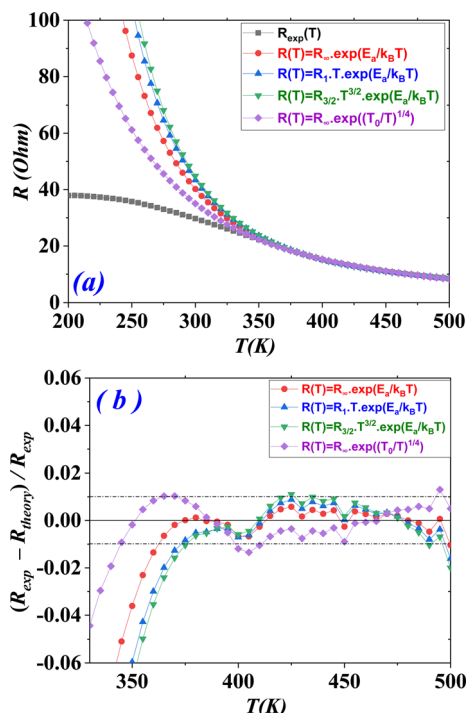


Fig. 7 (a) The experimental and the theoretical $R(T)$ curves for the four models. (b) The $\left(\frac{R_{\text{exp}} - R_{\text{theory}}}{R_{\text{exp}}}\right)$ vs. T plots for the four models under $\mu_0 H = 0$ T.

the benefit of the (e-e) interactions which increase with both temperature and the applied magnetic field (Fig. 6(d)). Fig. 6(e) shows that the negative contribution of (e-m) interactions is negligible at low temperatures, whether in the absence of the applied field or under a field of 2 or 5 T. By increasing the

temperature, this term promotes electrical conduction. The influence of the applied magnetic field is random.

At high temperatures, the behavior is that of an insulator or a semiconductor, where the electrical conductivity is governed by electron jumps. Three major mechanisms are possible: the thermally activated hopping (TAH) model,^{71,72} the model of small polaron hopping (SPH) between nearest neighbors⁶² and the Mott Variable Range hopping (Mott-VRH) transport model.⁷³

- The (TAH) model, governed by the Boltzmann factor, is written by the relation:

$$\rho(T) = \rho_{\infty} \exp\left(\frac{E_a}{k_B T}\right) \quad (8)$$

- The model, of (SPH) between nearest neighbors in an adiabatic limit, is governed by the equation:

$$\rho(T) = \rho_1 T \exp\left(\frac{E_a}{k_B T}\right) \quad (9)$$

The strong electron-polaron coupling, in a non-adiabatic limit, causes a slow motion of the charge carriers and leads to the following law:

$$\rho(T) = \rho_{3/2} T^{3/2} \exp\left(\frac{E_a}{k_B T}\right) \quad (10)$$

- In the (Mott-VRH) transport model, the resistivity is governed by the law:

$$\rho(T) = \rho_{\infty} \exp\left(\left(\frac{T_0}{T}\right)^{1/4}\right) \quad (11)$$

Table 3 The best fit parameters, for $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0), obtained from the experimental resistance data of the semiconductor behavior

$\mu_0 H$ (T)	0	2	5
Model ①	$\ln(R(T)) = \ln(R_{\infty}) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T}$		
$\ln(R_{\infty})$	-0.19203	-0.18252	-0.13683
E_a (meV)	100.50	100.07	98.04
Model ②	$\ln\left(\frac{R(T)}{T}\right) = \ln(R_1) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T}$		
$\ln(\rho_1)$	-7.27236	-7.21314	-7.15460
E_a (meV)	138.06	135.70	133.16
Model ③	$\ln\left(\frac{R(T)}{T^{3/2}}\right) = \ln(R_{3/2}) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T}$		
$\ln(R_{3/2})$	-10.8023	-10.67394	-10.49935
E_a (meV)	156.44	151.43	144.43
Model ④	$\ln(R(T)) = \ln(R_{\infty}) + \left(\frac{T_0}{T}\right)^{1/4}$		
$\ln(R_{\infty})$	-8.39416	-8.14518	-7.93170
T_0 (10^6 K)	6.12013	5.57338	5.11946
N_{EF} (10^{21} (eV) ⁻¹ cm ⁻³)	4.48064	4.92019	5.35644



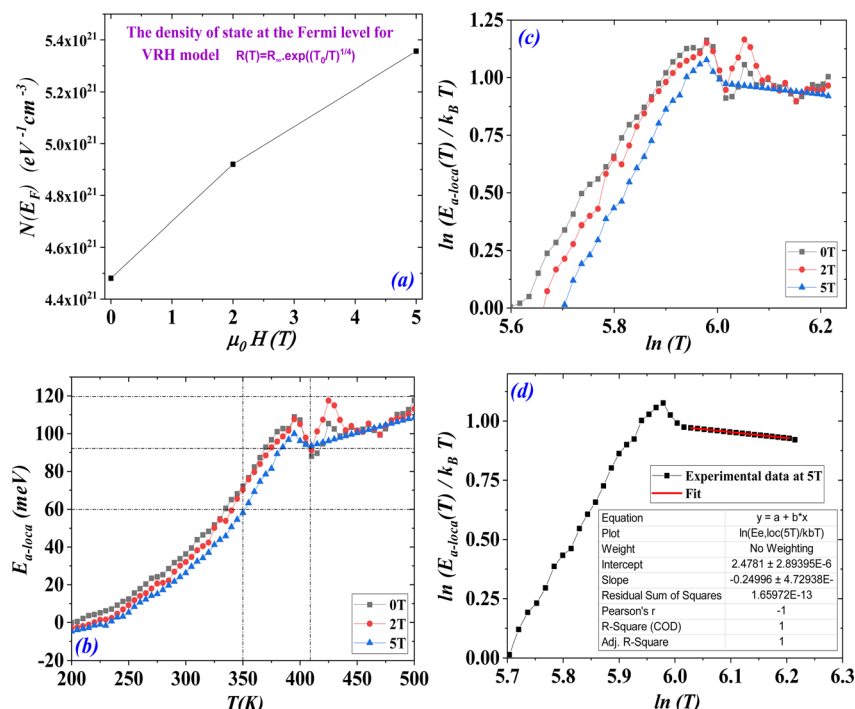


Fig. 8 (a) The density of state at the Fermi level, for (VRH) model, under different magnetic fields of 0, 2 and 5 T. (b) The temperature dependence of the local activation energy, under different magnetic fields of 0, 2 and 5 T. (c) The evolution of $\ln(E_{a-local}(T)/k_B T)$ as a function of $\ln(T)$ under different magnetic fields of 0, 2 and 5 T.

where, E_a is the activation or gap energy (width of the forbidden band), ρ_∞ is the value of the residual resistivity at an infinite temperature, and $T_0 = \frac{18}{k_B \times \mathcal{N}(E_F) \times \xi^3}$ denotes the characteristic temperature of Mott. $\mathcal{N}(E_F)$ is the density of state at the Fermi level and ξ represents the localization length d_{Mn-O} .

On the different components of Fig. S4 (see ESI),[†] we tried to model the experimental data of the resistance, at high temperatures, by the four eqn (8)–(11).

In order to search for the model giving the best fit of the experimental data, a fit with the three relations (12), (13), (14) and (15) was performed on Fig. 7(a), (b), (c) and (d), respectively.

$$\ln(R(T)) = \ln(R_\infty) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T} \quad (12)$$

$$\ln\left(\frac{R(T)}{T}\right) = \ln(R_1) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T} \quad (13)$$

$$\ln\left(\frac{R(T)}{T^{3/2}}\right) = \ln(R_{3/2}) + \frac{E_a(\text{eV}) \times 1.16 \times 10^4}{T} \quad (14)$$

$$\ln(R(T)) = \ln(R_\infty) + (T_0)^{1/4} \left(\frac{1}{T}\right)^{1/4} \quad (15)$$

The fit of the experimental curves, shows a good agreement between the experiment and the models mentioned above. The obtained values of the squared linear correlation coefficients (R^2), for all field, were as high as 99.9% and are listed in Table 3.

Using the adjustment parameters, the curves of eqn (8)–(11), relative to the resistance $R(T)$ were plotted on the same graph with the experimental curve (Fig. 7(a)). We note a good agreement between the experiment and the models studied for high temperatures. The divergence of the theoretical models is obtained at temperatures 355, 365, 367 and 340 K for the laws described by eqn (8)–(11), respectively. It is therefore clear that the (Mott-VRH) model is the most appropriate to describe the electrical resistance in the (PM) phase. This is confirmed by the curve giving the evolution of $\left(\frac{R_{\text{exp}} - R_{\text{theory}}}{R_{\text{exp}}}\right)$ as a function of temperature in Fig. 7(b). In fact, $\left(\frac{R_{\text{exp}} - R_{\text{theory}}}{R_{\text{exp}}}\right)$ of the (Mott-VRH) model is the smallest and does not exceed 1% over the extended temperature range between 345 and 500 K. The resistance in the (PM) phase is so characterized by a semiconductor behavior, which is best described by the (Mott-VRH) mechanism. This model describes low-temperature conduction in strongly disordered systems with localized charge-carrier states.^{74,75} From values of the characteristic temperature

$T_0 = \frac{18}{k_B \times \mathcal{N}(E_F) \times \xi^3}$ of Mott, listed in Table 3, we are able to determine the state density $\mathcal{N}(E_F)$ at the Fermi level, for different values of the magnetic field listed in Table 3. Fig. 8(a) shows that $\mathcal{N}(E_F)$, which is of the order of $10^{21} (\text{eV})^{-1} \text{cm}^{-3}$, increases slightly with the magnetic field. This evolution, confirms the previous findings. In fact, even in the PM phase, the application of an external magnetic field increases the magnetization, due to the suppression of magnetic domain

scattering,⁷⁶ which promotes conduction and gives a positive MR even if it is weak. These results are similar to those obtained by Venkataiah and Venugopal Reddy⁷⁷ in the orthorhombic $\text{Nd}_{0.67}\text{A}_{0.33}\text{MnO}_3$, ($\text{A} = \text{Ca}, \text{Sr}, \text{Pb}$ and Ba), and those obtained by N. Hasanli *et al.*⁷⁸ in the orthorhombic $\text{La}_{0.5}\text{Sr}_{0.5}\text{A}_{0.5}\text{Rh}_{0.5}\text{O}_3$, ($\text{A} = \text{Mn}$ and Fe). In the (Mott-VRH) model, the activation energy is a function of temperature.⁷⁹ We have examined the temperature variation of the local activation energy, using the following low:

$$E_{\text{a-local}}(T) = \frac{d(\ln(R(T)))}{d(1/k_{\text{B}}T)} \quad (16)$$

In Fig. 8(b), we show the plot of $E_{\text{a-local}}(T)$ vs. temperature. This curve reveals that this energy varies between 60 and 120 meV, for temperatures above 350 K, for all magnetic field values. Moreover, we can note that for a magnetic field of 5 T and for temperatures above 400 K this energy gradually increases in a quasi-linear manner. In the absence of the magnetic field ($\mu_0 H$) and even for a $\mu_0 H = 2$ T, the local activation energy performs fluctuations, around $E_{\text{a-local}}(T, 5 \text{ T})$. These fluctuations can be explained by the cation disorder σ^2 in A-site,[†] occupied by La^{3+} and Sr^{2+} cations with respective radii of 1.216 and 1.310 Å.⁸⁰ In fact, the size difference between different cations located in the A-site^{81–83} induces local distortions distributed randomly in the sample volume and, consequently, causes considerable variations of Mn–O–Mn bond lengths and angles.^{84–88} Thus, we have both the variance in the distribution of Mn–O and A–O distances $\sigma^2(\text{Mn–O})$ and $\sigma^2(\text{A–O})$, respectively.⁸⁹ The $\sigma^2(\text{Mn–O})$ quantifies the orthorhombic distortion and $\sigma^2(\text{A–O})$ quantifies the local one. An orthorhombic distortion induces the localization of the carriers, and therefore leads to the reduction of the FM behavior leading to the increase of the super-exchange AFM interactions. The application of an enough strong magnetic field is able to overcome these AFM interactions and suppress these fluctuations.

In order to confirm these results, we have plotted on Fig. 8(c), the evolution of $\ln(E_{\text{a-local}}(T)/k_{\text{B}}T)$ as a function of $\ln(T)$. These curves show, for high temperatures and for zero and 2 T magnetic field, the presence of fluctuations around $\ln\left(\frac{E_{\text{a-local}}(T, 5 \text{ T})}{k_{\text{B}}T}\right)$. The fit of the linear part Fig. 8 (d), for a 5 T magnetic field, shows a negative slope of $(-1/4)$. This confirms the validity of the (Mott-VRH) model to describe the conduction mechanism in the studied compound at high temperatures⁹⁰

Thus, there is a FMM at low temperatures and a semi-conducting-paramagnetic one (PMSC) at high ones.

In the intermediate temperatures range, around the transition, there is coexistence of (FM), (PM) or (SPM) clusters (macro spin paramagnetism). This can be justified by the second-order

† The cation disorder σ^2 is given by $\sigma^2 = \sum x_i r_i^2 - r_{\text{A}}^2$ where x_i and r_i are the atomic fraction and ionic radii of i -type ions at A-site, respectively.

$$\sigma^2 = (0.6 \times r^2(\text{La}^{3+}) + 0.4 \times r^2(\text{Sr}^{2+})) - (0.6 \times r(\text{La}^{3+}) + 0.4 \times r(\text{Sr}^{2+}))^2 = 21.20 \times 10^{-4} \text{ \AA}^2$$

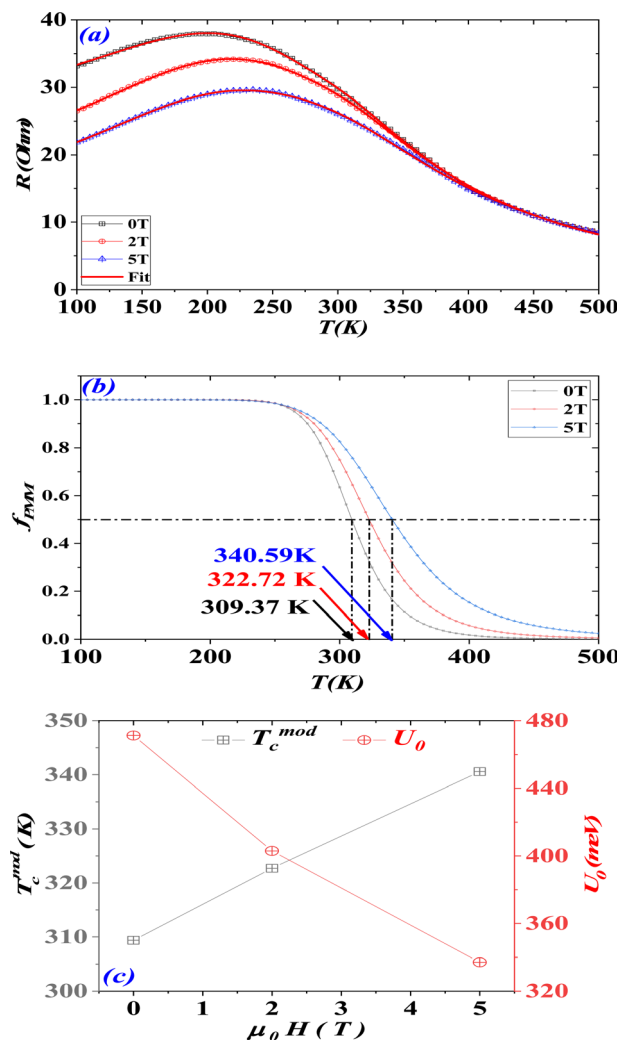


Fig. 9 (a) Fit of the temperature dependence of resistance, over the whole temperature range, under different applied magnetic fields of 0, 2 and 5 T, with the theoretical models: of percolation. (b) The temperature dependence on the volume fraction of the ferromagnetic phase f_{FMM} under different magnetic fields of 0, 2 and 5 T, (c) the evolution of $T_{\text{c}}^{\text{mod}}$ and of U_0 according to applied magnetic field. (d) The fit of the linear part at high temperatures, for a 5 T field, of $\ln\left(\frac{E_{\text{a-local}}(T)}{k_{\text{B}}T}\right)$ as a function of $\ln(T)$.

transition width, which is manifested by a dispersion in the temperature transition. To describe the behavior of the resistivity between the (FMM) and (PMSC) transition, we will adopt the percolation model, based on the phase segregation mechanism. This model, proposed by Li *et al.*,⁹¹ assumes that materials are composed by FMM and PMSC regions, of volume

Table 4 The best fit parameters, for $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0) sample, obtained from the experimental resistance data and the percolation model, over the entire temperature range

$\mu_0 H$ (T)	0	2	5
U_0 (meV)	471.20	402.97	337.08
$T_{\text{c}}^{\text{mod}}$ (K)	309.37	322.72	340.59

fractions (f) and $(1 - f)$, respectively. Thus, the electrical resistance of the system at any temperature is determined by the following expression:

$$R(T) = f \times R_{\text{FMM}}(T) + (1 - f)R_{\text{PMSC}}(T) \quad (17)$$

where:

$$R_{\text{FMM}}(T) = R_0 + R_2 T^2 + R_{4.5} T^{4.5} \quad (18)$$

$$R_{\text{PMSC}}(T) = R_\infty \exp\left(\left(\frac{T_0}{T}\right)^{1/4}\right) \quad (19)$$

The f function satisfies well the Fermi-Dirac distribution given by:

$$f = \frac{1}{1 + \exp\left(\frac{U_0}{k_B T} \left(\frac{T}{T_C^{\text{mod}}} - 1\right)\right)} \quad (20)$$

where U_0 is the energy difference between the FM and PM states at $T = 0$ K and T_C^{mod} is the PM/FM transition temperature used in the model.

So, we have:

$$\begin{cases} f(T \ll T_C^{\text{mod}}) = 1 \\ f(T = T_C^{\text{mod}}) = \frac{1}{2} \\ f(T \gg T_C^{\text{mod}}) = 0 \end{cases}$$

which give:

$$\begin{cases} R(T \ll T_C^{\text{mod}}) = R_{\text{FMM}}(T) \\ R(T = T_C^{\text{mod}}) = \frac{1}{2} \times (R_{\text{FMM}}(T) + R_{\text{PMSC}}(T)) \\ R(T \gg T_C^{\text{mod}}) = R_{\text{PMSC}}(T) \end{cases}$$

The adjustment of the experimental resistance curves over the entire temperature range, using the eqn (17)–(20), allows us to determine the U_0 and T_C^{mod} parameters. After, we can go back up to the f function given by eqn (20) and afterwards determine the volume fractions of each phase. The R_0 , R_2 , $R_{4.5}$, T_0 and R_∞ parameters are initiated with those listed in Tables 2 and 3.

Fig. 9(a) presents the experimental results of $R(T)$ as well as the simulated curves (red line) obtained at magnetic field of 0, 2 and 5 T. The good agreement between the experimental data and the simulated curves affirm that the percolation model describes well the behavior of the resistivity in a wide temperature range including the phase transition region whatever the applied magnetic field. The best fit of the U_0 and T_C^{mod} parameters are given in Table 4. With these two parameters only, we have drawn using relation 20, in Fig. 9(b), the curves giving, the temperature dependence on the volume fraction of the FM phase f_{FMM} under different magnetic fields. The application of an external magnetic field has practically no effect on the volume fractions of the FMM and PMSC phase, at low and at high temperatures. In the transition temperature

range, a magnetic field favors the volume fraction of the FMM phase over the PMSC phase, leading to the increase in the conductivity and, as consequence, the appearance of the colossal magnetoresistance around the transition. These interpretations are in agreement with the evolution of T_C^{mod} according to applied magnetic field, in the inset of Fig. 9(b). The PMSC/FMM transition temperature T_C^{mod} , increases with the application of the magnetic field (Fig. 9(b) and (c)), which favors the FMM phase. These interpretations are in good consistency with the evolution of U_0 according to applied magnetic field, traced in Fig. 9(c). The energy difference U_0 between the FM and PM states at $T = 0$ K, proved to decrease with the application of the magnetic field, which promotes electrical conduction. However, the PMSC/FMM transition temperature T_C^{mod} , increases with the application of the magnetic field, which favors the FMM phase. All these results prove the strong correlation between magnetic and electrical properties.

4. Conclusion

The correlation between the electrical and the magnetic properties of nanoparticles $\text{La}_{0.6}\text{Sr}_{0.4}\text{MnO}_3$ (S1C0) compound has been studied. By decreasing the temperature, the (S1C0) compound was found to undergo a transition, of a second order, from a paramagnetic (PM) phase to a ferromagnetic (FM) one. The images, obtained using a transmission electron microscope, have showed a dispersion in grain size. The dispersion in the transition (PM/FM) temperature has been related to the presence of an irreversibility between the magnetization in (ZFC) and (FC) modes. This irreversibility, at the temperature $T_i \approx 350$ K below the critical temperature of the PM/FM transition $T_C \approx 365$ K, indicated a magnetic anisotropy around this temperature. So, we have deduced that our compound is the seat of a superparamagnetism (SPM) phenomenon justified by the absence of hysteresis losses.

The study of the electrical properties showed a semi-conductor behavior (SC) at high temperatures and a metallic one (M) at low temperatures. The SC/M transition temperature, which was found to increase under the effect of a magnetic field, has been approximated at the Blocking temperature. The conduction mechanisms have been tested by three models at low temperatures as well as at high ones. The Zener double exchange (ZDE) polynomial model, with the electron-magnon interaction, was found to be the best one to describe the electrical conduction in the metallic region, verifying a deviation relative to the experimental data closest to zero.

At high temperatures, the Mott variable range hopping (Mott-VRH) transport model was considered as the most appropriate one to describe the electrical conduction. Finally, we have adopted the percolation model $R(T) = f \times R_{\text{FMM}}(T) + (1 - f)R_{\text{PMSC}}(T)$, based on the phase segregation mechanism, to describe the conduction mechanism over the whole temperature range, where the function (f), giving the volume fraction of the ferromagnetic metallic (FMM) regions, satisfies the Fermi-Dirac distribution.



Data availability

The data are not publicly available.

Conflicts of interest

There are no conflicts to declare.

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