


 Cite this: *RSC Adv.*, 2024, 14, 23592

# Tailoring of structural, morphological, electrical, and magnetic properties of $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$ ceramics

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This study undertakes a comparative analysis of the structural, morphological, electrical, and magnetic characteristics of Fe-doped  $\text{LaMnO}_3$  ceramics. The solid-state reaction method was used to prepare Fe-doped  $\text{LaMnO}_3$  at different concentrations ( $0.00 \leq x \leq 1.00$ ) and has been characterized using X-ray diffraction (XRD), Fourier transforms infrared spectroscopy (FTIR), Field emission scanning electron microscopy (FE-SEM), energy-dispersive spectroscopy (EDS), and vibrating sample magnetometry (VSM). The structural transformation from rhombohedral to orthorhombic with Fe-doping is demonstrated by Rietveld's refined XRD patterns. The positive slope in Williamsons–Hall's (W–H) plots confirms the presence of tensile strain with increasing average crystallite size. Quasi-spherical morphology of all the compositions with similar uniformity was confirmed by FESEM images. The chemical distribution of all the elements has been identified by EDS mapping images. Normal dielectric dispersion behaviour of all the samples with NTCR response is confirmed by dielectric and impedance analysis respectively. Increasing lattice volume with Fe-concentration results in increasing  $E_a$ . The presence of antiferromagnetic ordering, in addition to weak ferromagnetic ordering, is indicated by the unsaturated magnetization even up to a high external field. The decrease in  $M_s$  and increase in  $H_C$  values due to Fe-doping reflect the influence of particle size on various magnetic parameters.

 Received 8th July 2024  
 Accepted 17th July 2024

DOI: 10.1039/d4ra04931d

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## Introduction

Perovskite materials based on lanthanum have been widely used in numerous applications because of their apparent structural, electrical, optical, catalytic, and magnetic capabilities.<sup>1–4</sup> Their structures' substitution of metals, varying oxidation states, variable oxygen stoichiometry, and stability of these materials show their wide applicability in solid oxide fuel cell electrodes,<sup>5</sup> chemical sensors,<sup>6</sup> optoelectronic devices,<sup>7</sup> spintronics, and solar cells,<sup>8</sup> because of their low power consumptions. The features of lanthanum-based perovskite

materials' microwave absorption have been extensively researched to solve issues connected to radiation pollution and electromagnetic interference (EMI), which pose a threat to information security as well as human health and device operation.<sup>9–11</sup>

Lanthanum-based perovskite materials such as  $\text{LaMnO}_3$ ,  $\text{LaFeO}_3$ ,  $\text{LaCrO}_3$ ,  $\text{LaNiO}_3$ , and  $\text{LaCoO}_3$ , have been widely studied due to their ease of manufacture, flexibility, and low production cost.  $\text{LaMnO}_3$  and  $\text{LaFeO}_3$  are perovskite-type materials generally having orthorhombic crystal structures at room temperature.  $\text{LaMnO}_3$  (LMO) is an A-type antiferromagnetic perovskite material that shows orthorhombic crystal symmetry only around its stoichiometric composition.<sup>12–14</sup> The magneto-resistance properties of LMO make it an important multiferroic material that can be exploited for the use of LMO as electrode material in supercapacitors.<sup>15</sup> On the other hand,  $\text{LaFeO}_3$  (LFO) is an orthoferrite, has a Néel temperature ( $T_N$ ) of roughly 738 K, and a G-type antiferromagnetic structure.<sup>9,16</sup> Orthorhombically distorted LFO-perovskite was found to have multiferroic properties and has potential applications as a spin filter in spintronics.<sup>17,18</sup>

It has been observed that 3d cations on the B-site (*i.e.*,  $\text{Mn}^{4+}$  or  $\text{Fe}^{3+}$ ) of the crystal lattice drive the electrical and magnetic properties of LMO and LFO. The double exchange mechanism (DE) and the Jahn–Teller effect are the two main reasons for the

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co-existence of two ferroic orders in LMO and LFO ceramics. DE mechanism is the hopping of  $e_g$  electrons between  $Mn^{3+}$  and  $Mn^{4+}$  mediated by oxygen anions, while the Jahn–Teller effect is strong electron-phonon coupling arises due to the deformation of  $[MnO_6]$  or  $[FeO_6]$  octahedral. These two machines were observed in lanthanum-based ceramics by appropriate doping or by introducing oxygen vacancies in the crystal structure.<sup>19–21</sup> Lanthanum is a non-magnetic rare earth metal with zero unpaired electrons in its  $La^{3+}$  oxidation state which does not influence magnetic properties directly.<sup>22</sup> Hence, due to the presence of unpaired electrons and high magnetic moment values of B-site ions (Fe and Mn ions), a change in magnetic properties has been observed in lanthanum-based perovskite materials. Thus, it is important to study the Mn-site substitution in  $LaMnO_3$  as the Mn-site doped ion will consequently result in the distortion of the Mn–O plane, which will cause a considerable change in structural and electrical and magnetic properties.<sup>23</sup> In recent years, extensive research has been conducted on the effects of substituting Mn sites with various cations ( $M = Fe, Cr, Co, Ni$ ) in both charge-delocalized and localized manganite's.<sup>24–27</sup> Among these, Fe ions, due to their close ionic radii to  $Mn^{3+}$ , are particularly interesting as dopants for  $LaMnO_3$ , as they can occupy Mn sites as  $Fe^{3+}$  without causing significant lattice distortion.<sup>28</sup>

Different groups have reported on a number of investigations including Fe doping at the Mn location.<sup>29–32</sup> With decreasing temperature, the perovskite system  $La_{0.67}Ca_{0.33}Mn_{1-x}Fe_xO_3$  ( $x = 0.00, 0.01, 0.03, \text{ and } 0.07$ ) exhibits an essentially paramagnetic (PM) to ferromagnetic (FM) transition, and as Fe concentration increases, the PM to FM transition temperature is decreased.<sup>32</sup> Magnetization measurements of the system  $La_{0.6}Ca_{0.4}Mn_{1-x}Fe_xO_3$  ( $x = 0, 0.05, 0.1, 0.15, \text{ and } 0.2$ ) revealed that the coexistence of ferromagnetic and antiferromagnetic interactions for critical composition  $x \sim 0.1$  and that the Curie temperature ( $T_C$ ) decreases upon Fe-doping from 275 K to 75 K for  $x = 0$  and  $x = 0.2$ , respectively.<sup>31</sup> In their studied system  $La_{0.7}Ca_{0.3}Mn_{1-x}Fe_xO_3$  ( $x = 0.08, 0.1, \text{ and } 0.12$ ), Sahasrabudhe *et al.*<sup>33</sup> found only one ferromagnetic phase exist below the transition temperature without any spin glass behaviour. Kundaliya *et al.*<sup>34</sup> also noted comparable outcomes for the combination  $La_{0.67}Ca_{0.33}Mn_{0.9}Fe_{0.1}O_3$ . Fe-doping of  $La_{1-x}Ca_xMnO_3$  results in the suppression of ferromagnetism and conduction in both the ferromagnetic ( $x = 0.37$ ) and antiferromagnetic ( $x = 0.53$ ) phases.<sup>35</sup> This is because double-exchange interactions are reduced and the number of hopping electrons is reduced. In the  $La_{0.67}Ca_{0.33}Mn_{0.9}Fe_{0.1}O_3$  perovskite, the random substitution of  $Fe^{3+}$  with  $Mn^{3+}$  lowers the number of locations where the Mn  $e_g$  (up) electron can hop, which lowers ferromagnetic exchange. At low temperatures, the system is driven into a randomly canted ferromagnetic state by the competition between the co-existing antiferromagnetic super-exchange interactions and the ferromagnetic double-exchange interactions.<sup>36</sup>

$Fe^{3+}$  can readily take the place of  $Mn^{3+}$  in the  $MnO_6$  octahedron. Reduced Jahn–Teller distortion from  $Mn^{3+}$  ions results from partial replacement of  $Fe^{3+}$  for  $Mn^{3+}$ . In the meantime, resistivity rises as a result of Fe doping, which lessens the

double exchange contact between  $Mn^{3+}$  and  $Mn^{4+}$ . The degree of the double exchange effect's destruction deepens, the degree of the crystal structure's distortion is lessened, the symmetry of the crystal structure is strengthened, the magnetoresistance (MR) progressively rises, and the temperature coefficient of resistance (TCR) initially rises and then falls with an increase in Fe doping.<sup>37–39</sup> Kholil *et al.*,<sup>40</sup> reported in their report that Fe-doping can be used to reduce bandgap in perovskites and also shift the optical conductivity in the visible region. Thus, Fe-doped perovskites provide promising candidates for applications in optoelectronics, solar cells, solid oxide fuel cells and in fabrication of new magnetic devices.<sup>41–43</sup>

Numerous studies have investigated Fe-doped  $LaMnO_3$ ; however, none have explored the complete substitution of Mn ions with Fe-ions. Thus, this study offers an extensive examination of the structural, morphological, magnetic, and electrical characteristics of  $LaMn_{1-x}Fe_xO_3$  ceramics. In addition to examining how magnetic behaviour changes with different Fe doping concentrations in  $LaMnO_3$ , we also analysed dielectric and impedance characteristics at various temperatures and frequencies.

## Results and discussion

FullProf software was utilized to process the Rietveld refinement of XRD data for  $LaMn_{1-x}Fe_xO_3$  ( $x = 0.00, 0.25, 0.50, 0.75, \text{ and } 1.00$ ), as shown in Fig. 1 (a–e). The refinement confirms the phase transition from rhombohedral to orthorhombic as the concentration of Fe-ions is increased in  $LaMnO_3$ . Rhombohedral crystal structure with  $R\bar{3}C$  space group was confirmed for  $0.00 \leq x \leq 0.50$  concentration of Fe-doping in  $LaMnO_3$ , while the orthorhombic crystal structure with the  $Pbnm$  space group was obtained for  $0.75 \leq x \leq 1.00$ . Different refined parameters along with their residual factors ( $R_p, R_{wp}, R_e$ ) are summarized in Table 1. The lattice parameters  $a = b = 5.5227 \text{ \AA}$ ,  $c = 13.3646 \text{ \AA}$  and  $a = 5.5540 \text{ \AA}$ ,  $b = 5.5637 \text{ \AA}$ ,  $c = 7.8519 \text{ \AA}$  for LMO and LFO respectively, were found well-matched with the literature.<sup>44,45</sup> Phase transition can be attributed to the ionic radii difference between  $Mn^{4+}$  (0.053  $\text{\AA}$ ) and  $Fe^{3+}$  (0.064  $\text{\AA}$ ) ions. Due to larger ionic radii of  $Fe^{3+}$  ions, as the  $Mn^{4+}$  ions are replaced by  $Fe^{3+}$  ions, it generates high pressure on the grain boundaries and results in structural transformation from rhombohedral to orthorhombic.

To investigate the kind of micro-strain seen in the crystal structures of  $LaMn_{1-x}Fe_xO_3$ , the Williamsons–Hall (W–H) method was employed. To determine the crystallite size and micro-strain in the rhombohedral and orthorhombic crystal structures, the linear fitted line (as indicated in Fig. 2) was employed. For all samples except for compositions  $x = 0.75$ , which display a negative slope as illustrated in Fig. 2(d), the positive slope of the linear fit validates the tensile strain. The negative slope line confirmed the compressive strain. Because Fe-ions have large ionic radii, it is observed that when Fe-ions are doped in  $LaMnO_3$ , the average crystallite size increases as the number of Fe-ions increases. The two intrinsic quantities that provide the dislocation network's measure are the lattice strain and the dislocation density. In Fig. 2(c) for  $x = 0.50$  an

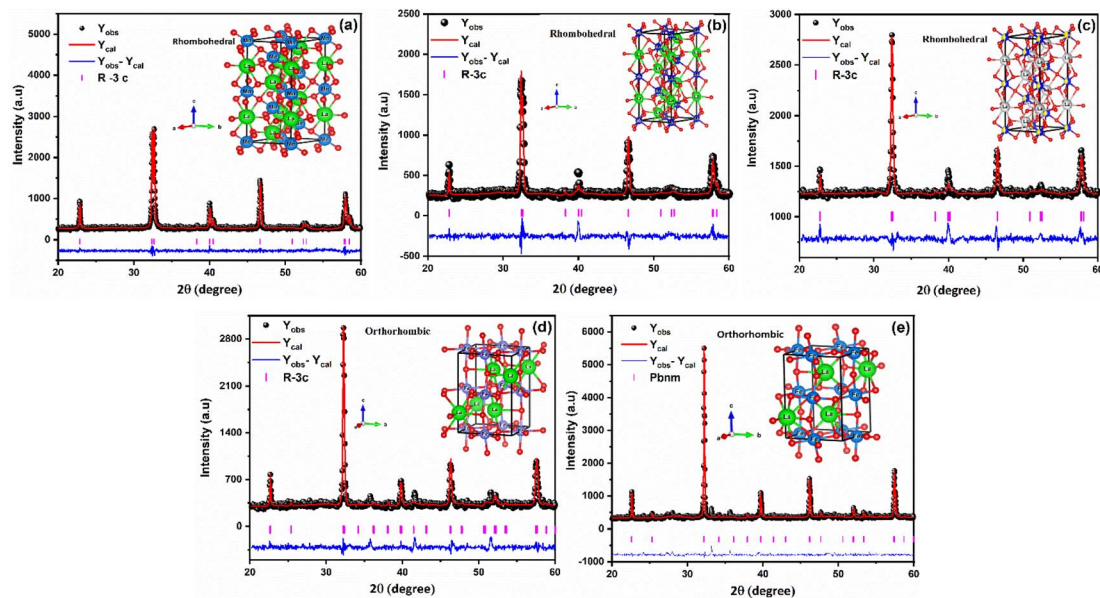


Fig. 1 Rietveld refined XRD pattern of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics (where (a)  $x = 0.00$ , (b)  $x = 0.25$ , (c)  $x = 0.50$ , (d)  $x = 0.75$ , (e)  $x = 1.00$ ).

Table 1 Rietveld refined and calculated the structural parameters of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics

		$x = 0.00$	$x = 0.25$	$x = 0.50$	$x = 0.75$	$x = 1.00$	
Simulated parameters		(Rhombohedral) ( $R\bar{3}c$ )			(Orthorhombic) ( $Pbnm$ )		
Lattice parameters	$a$ (Å)	5.5227	5.524194	5.52837	5.527039	5.5539	
	$b$ (Å)	5.5227	5.524194	5.52837	5.527039	5.5635	
	$c$ (Å)	13.3646	13.3869	13.4476	7.8535	7.8517	
Volume of unit cell	$V$ (Å <sup>3</sup> )	353.0115	353.7920	355.9479	241.033	242.6303	
Atomic positions	La	$x$	0.0000	0.0000	0.0000	0.99211	0.99694
		$y$	0.0000	0.0000	0.0000	0.01920	0.02651
		$z$	0.2500	0.2500	0.2500	0.2500	0.2500
	Mn/Fe	$x$	0.0000	0.0000	0.0000	0.0000	0.0000
		$y$	0.0000	0.0000	0.0000	0.5000	0.5000
		$z$	0.0000	0.0000	0.0000	0.0000	0.0000
	$O_1$	$x$	0.44736	0.45901	0.45901	0.75178	0.70561
		$y$	0.0000	0.0000	0.0000	0.70500	0.29975
		$z$	0.2500	0.2500	0.2500	0.0159	0.03019
	$O_2$	$x$	—	—	—	0.09087	0.07964
		$y$	—	—	—	0.48723	0.47727
		$z$	—	—	—	0.2500	0.2500
$R$ -factors (%)	$R_p$	25.80	27.5	41.2	26.6	41.70	
	$R_{exp}$	16.67	12.6	19.7	12.7	18.34	
	$R_{wp}$	16.60	24.3	24.0	20.9	22.70	
	$R_f$	4.28	16.2	18.2	10.9	4.50	
	$R_{Bragg}$	1.930	6.28	2.43	4.88	2.664	
GoF		1.00	1.87	0.45	1.97	1.53	
Crystallite size ( $D$ (nm))		81.25	119.24	120.83	128.52	132.13	
Lattice strain ( $\epsilon$ )		$5.58 \times 10^{-4}$	$2.94 \times 10^{-4}$	$3.24 \times 10^{-5}$	$-2.15 \times 10^{-4}$	$1.79 \times 10^{-4}$	
Dislocation density $\delta$ (nm <sup>-1</sup> )		$1.52 \times 10^{-3}$	$7.03 \times 10^{-5}$	$6.84 \times 10^{-5}$	$6.05 \times 10^{-5}$	$5.73 \times 10^{-5}$	
X-ray density $\rho_x$ (g cm <sup>-3</sup> )		6.83	6.99	6.89	6.74	6.65	
Specific surface area $S$ (m <sup>2</sup> g <sup>-1</sup> )		10.81	7.20	7.19	6.93	6.83	

exceptionally high value is observed. This may be due to the reason of equal concentration of  $\text{Fe}^{3+}$  and  $\text{Mn}^{4+}$ -ions, which results into inhomogeneity at B-site. This inhomogeneity leads to lattice distortion due to the ionic radii of  $\text{Fe}^{3+}$  and  $\text{Mn}^{4+}$ -ions.

Therefore, maximum strain in the ceramic at  $x = 0.50$  is observed. Fig. 2(e) a small deviations in the data point is seen, which may be due to the reason B-site is fully occupied by  $\text{Fe}^{3+}$ -ions. Table 1 shows that a smaller lattice strain value

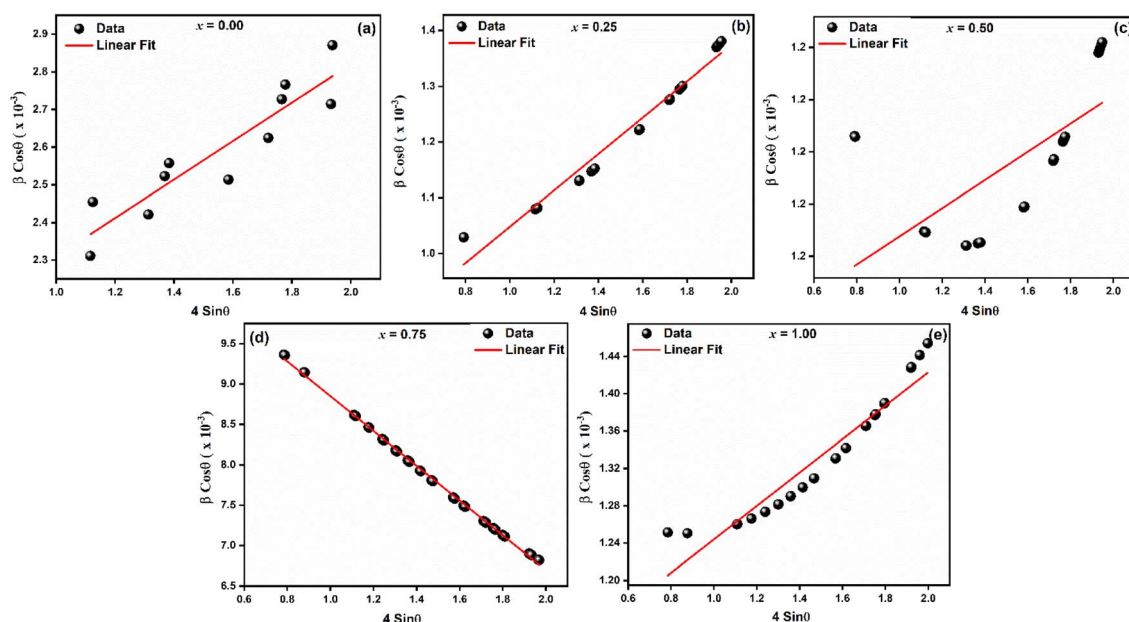


Fig. 2 William-Hall plots of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

corresponds to a lower concentration of lattice defects, whereas a reduced dislocation density corresponds to the production of a higher-quality sample. Eqn (1) and (2) were used to determine the synthesized NPs' X-ray densities ( $\rho_x$ ) and specific surface area ( $S$ ).

$$\rho_x = \left( \frac{ZM}{NV} \right) \quad (1)$$

$$S = 6l(D\rho_x) \quad (2)$$

where  $M$  is the samples' molecular weight,  $N$  is Avogadro's number,  $V$  is the unit cell's volume, and  $D$  is the average

crystallite size,  $Z$  is the number of atoms in the rhombohedral/orthorhombic phase's unit cell. The obtained values of X-ray densities for these ceramics are listed in Table 1. The specific surface area ( $S$ ) was found to be decreasing with increasing crystallite size which can be attributed to increasing surface-to-volume ratio.

To well understand the microstructural features of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ), FESEM was performed as shown in Fig. 3. All the samples appear to contain quasi-spherical particles with shapes of similar uniformity. The microstructure consisted of sub-micron-sized particles with very fine morphology. All ceramics exhibit distinct, well-defined grains and boundaries.

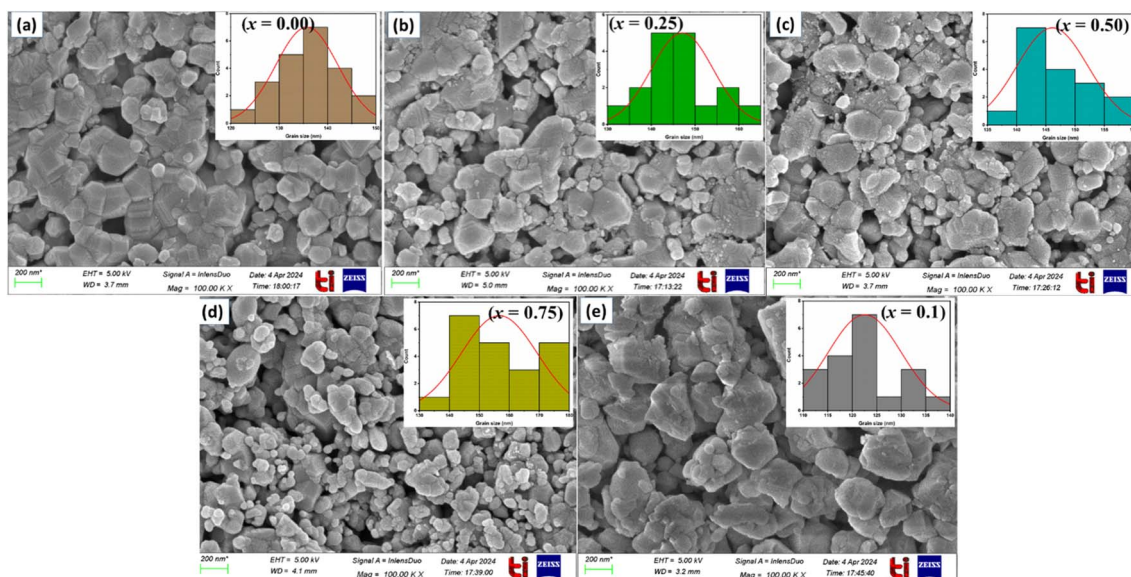


Fig. 3 FESEM images of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

Table 2 Elemental details and average grain size of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics

Composition	Elements	O	Mn	La	Fe	Grain size (nm)
$x = 0.00$	Wt (%)	18.68	23.14	58.18	—	135
	At (%)	58.15	20.98	20.98	—	
$x = 0.25$	Wt (%)	18.76	14.40	57.65	9.19	145
	At (%)	58.21	13.01	20.60	8.17	
$x = 0.50$	Wt (%)	17.61	4.28	57.63	20.48	146
	At (%)	56.14	3.98	21.17	18.71	
$x = 0.75$	Wt (%)	19.78	8.40	56.81	15.00	156
	At (%)	59.82	7.40	19.79	13.00	
$x = 1.00$	Wt (%)	18.92	—	55.50	25.58	122
	At (%)	57.97	—	19.59	22.45	

Plotting histograms (inset) from SEM images using Image-J software allowed us to determine the average grain size, which ranged from 122 to 156 nm (see Table 2). The existence of secondary particles created by agglomeration accounts for the smaller average crystallite size found in SEM pictures compared to that found in X-ray diffraction.

EDS spectra confirm the homogeneous composition of the synthesized samples (as shown in Fig. 4), with elemental distribution mapped in  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ). La, Mn, Fe, and O elements are detected at their expected energy levels, with atomic and weight percentages listed in Table 2, consistent with the nominal composition. No additional elements are detected, affirming the purity of the synthesized materials.

The thermal variations of dielectric constants ( $\epsilon$ ) and dielectric loss ( $\tan \delta$ ) in the temperature range 30–250 °C of the samples observed at frequencies 1 kHz, 10 kHz, 100 kHz, and 1 MHz, respectively are shown in Fig. 5(a–e). Typical dielectric dispersion behavior like ferroelectric materials has been observed in all the samples that is with increasing temperature the dielectric constants ( $\epsilon$ ) first increase and then attain a peak at Curie temperature ( $T_C$ ), and above  $T_C$ , further increase in the temperature causes a rapid decrease in the dielectric constant

( $\epsilon$ ). It is noted that at low temperatures,  $\epsilon$  is temperature independent and frequency independent for all samples. Then, for  $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00,$  respectively, it increases progressively with increasing temperature to its maximum value ( $\epsilon_{\text{max}}$ ), which corresponds to the change from a ferroelectric to a paraelectric phase, at approximately 60, 110, 150, 230, and 222 °C.

Moreover, dielectric constant decreases with increasing frequency as shown in Fig. 5(a–e). Change in  $\epsilon$  values with frequency depends on the extrinsic as well as intrinsic contribution. For every sample, the contribution (extrinsic) from the grain boundary is greater than the bulk grain, as indicated by the huge values of  $\epsilon$  in the low frequency. The Maxwell–Wagner model may provide an explanation for this, as the charges that build at the grain borders cause frequency-dependent behaviour that suggests the conducting grains and insulating grain boundaries that separate the polycrystalline perovskites. According to findings in the literature, the grain borders (extrinsic) contribute more at a low frequency than the grain contribution. The dielectric loss variation with temperature exhibits the same characteristics as the dielectric constant temperature variation, and it might be described using the same methodology as the dielectric constant discussion. It is discovered that the dielectric loss rises as the temperature climbs. Temperature-related increases in charge carrier mobility cause polarization to rise and significant dielectric loss. Charge accumulating at grain boundaries is the cause of the increased dielectric loss value that has been found at high temperatures.

Frequency-dependent of real part of impedance ( $Z'$ ) is shown in Fig. 6 for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ) ceramics at high temperatures 200, 210, 220, 230, 240 and 250 °C, respectively. As shown in the figures, the magnitude of  $Z'$  is higher at lower temperatures and decreases with increasing frequency for all compositions, confirming the NTCR behaviors of these ceramics. This decrease in  $Z'$  corresponds to an increase in electrical conductivity. As frequency increases,  $Z'$  values converge, leading to decreased barrier

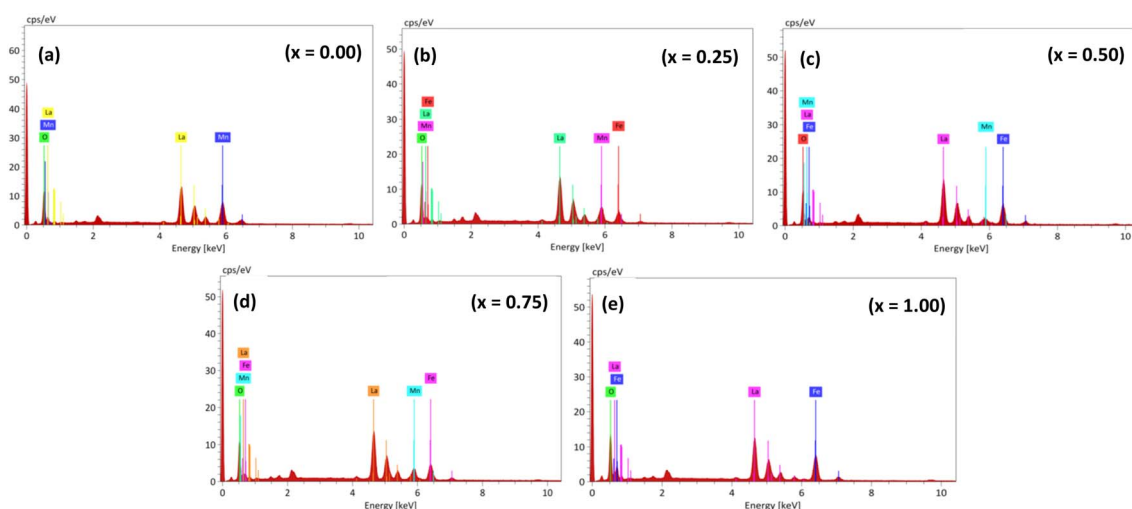


Fig. 4 EDS spectra of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

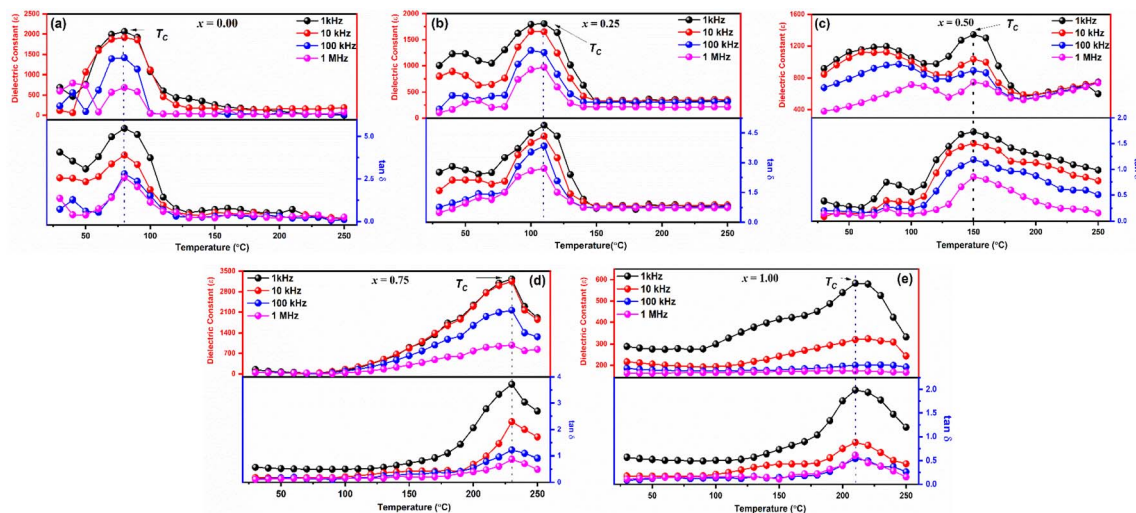


Fig. 5 Temperature dependent dielectric constant ( $\epsilon'$ ) and dielectric loss ( $\tan \delta$ ) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

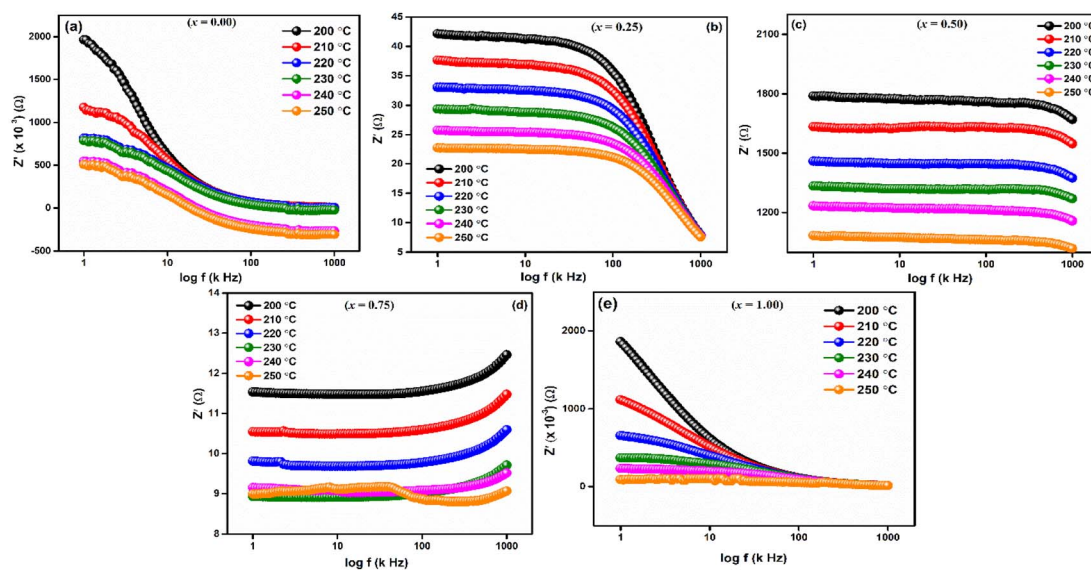


Fig. 6 Frequency dependent real part of impedance ( $Z'$ ) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

properties and the disappearance of polarization caused by space charge.<sup>46</sup>

Fig. 7(a–e) displays the imaginary portion of impedance ( $Z''$ ) frequency response curves for a wide range of temperatures (200–250 °C) for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ). Type and strength of electrical relaxation events in the system can be determined by looking for temperature-dependent peaks ( $Z''_{\text{max}}$ ) at a specific frequency. The system's space charge is evident from this behaviour.<sup>25,26</sup> As frequency and temperature rise, we observe that the value of ( $Z''$ ) first rises until peaking at  $Z''_{\text{max}}$ , after which it falls. The peaks shift towards higher frequencies as the temperature rises, which is a significant observation. It is significant to observe that as temperature rises, the peaks shift towards higher frequencies. The change in the peaks shows how the system's relaxation

period is spreading, and the fact that this relaxation phenomenon evolved as the temperature rose supports the theory that temperature-dependent dielectric relaxation exists. The relaxation process, which causes electrical conduction in materials, may be caused by flaws created at higher temperatures. The fact that all compositions' height peaks or  $Z''_{\text{max}}$  values fall as temperature rises further supports the development of thermally activated charge carriers, which power the materials' conduction mechanism.<sup>47,48</sup> The confirmation of a non-Debye-type relaxation process in the materials is provided by the asymmetric peak broadening, which reflects the relaxation time distribution. Furthermore, it was discovered that in the higher frequency area, all the curves combine at a particular frequency. This is thought to be caused by the space charge polarization decreasing at higher frequencies.<sup>25–28</sup>

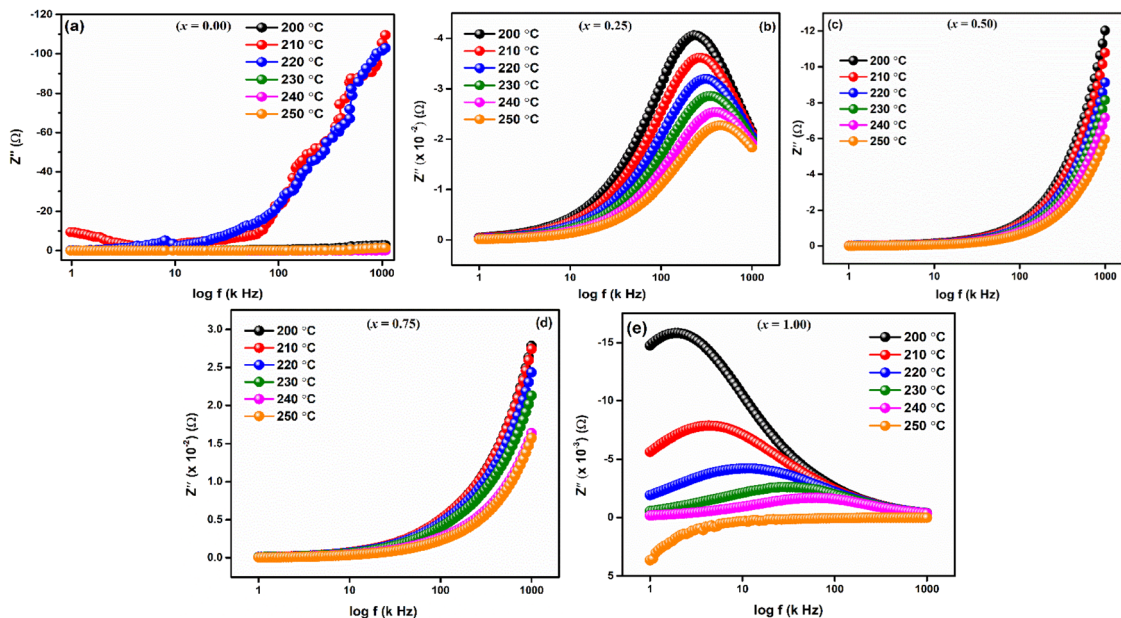


Fig. 7 Frequency dependent imaginary part of impedance ( $Z''$ ) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

The Nyquist representation, which plots the imaginary part  $Z''$  vs. the real part  $Z'$  in a Cartesian orthonormal reference frame, is one of the most used graphical representations in complex impedance spectroscopy. The literature<sup>23</sup> states that the contribution of the grain boundary is represented by the semicircle in lower frequencies, while the semicircle in higher frequencies symbolizes the phenomena of intrinsic conduction, the response of the grains, and gives rise to the resistances of the grains. Fig. 8(a–e) shows our compound's Nyquist diagrams obtained at various temperatures. Each composition has a single, depressed

semi-circular arc, its centre located below the real axis, signifying that it is semiconducting and has an NTCR. Every semicircle's diameter shrinks as temperature rises, suggesting a thermally triggered conduction process.<sup>49</sup> The Nyquist plots are fitted with an equivalent circuit made up of two parallel resistances ( $R_g$  and  $R_{gb}$ ) and a constant phase element ( $Q$ ) in order to verify the distinct contributions of grains and grain borders. Table 3 lists the fitted values resistance for grain and grain borders, whereas the black, red and blue lines show the fitted lines and dots shows the experimental data, respectively.

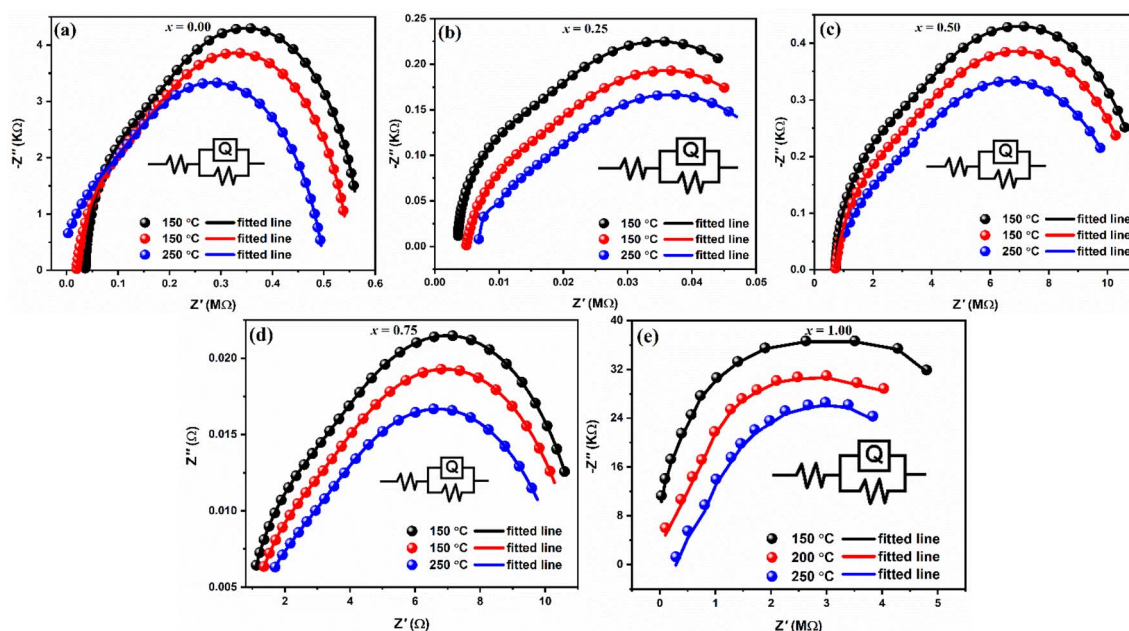


Fig. 8 Cole–Cole plot for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

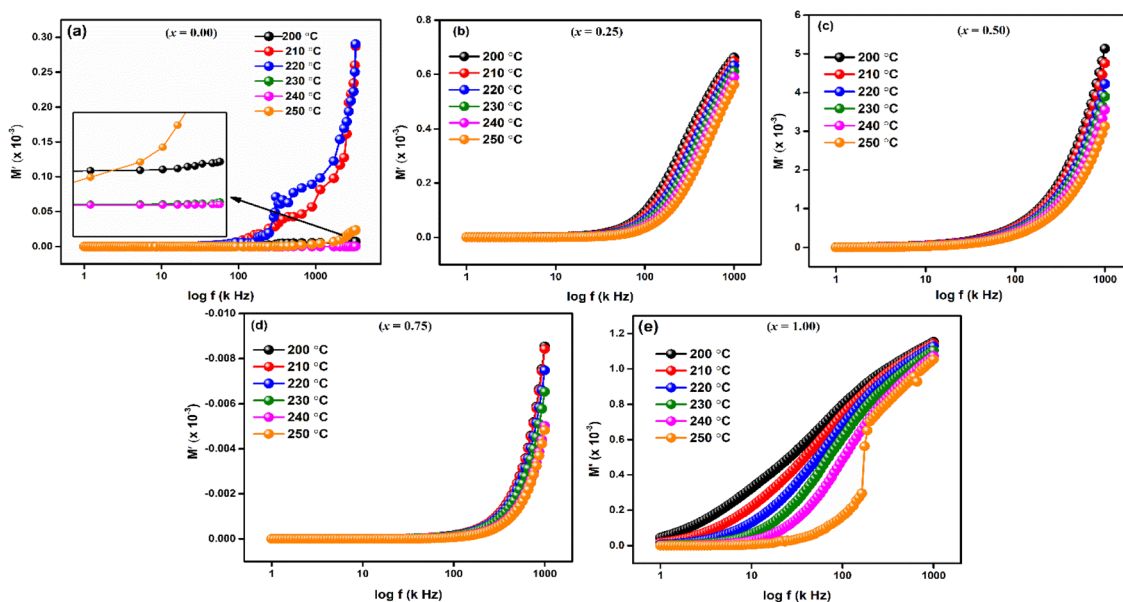
Table 3 Parameters calculated from impedance fitted data of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics

Compositions ( $x$ )	Temperature ( $^{\circ}\text{C}$ )	Resistance		$n$	$Q$
		$R_g$ ( $\Omega\text{cm}^2$ )	$R_{gb}$ ( $\Omega\text{cm}^2$ )		
0.00	150	$2.0 \times 10^{-9}$	$2.1 \times 10^4$	$6.0 \times 10^{-1}$	$9.6 \times 10^{-10}$
	200	$1.5 \times 10^{-9}$	$2.3 \times 10^3$	$6.0 \times 10^{-1}$	$5.7 \times 10^{-10}$
	250	$1.0 \times 10^{-9}$	$2.3 \times 10^3$	$6.0 \times 10^{-1}$	$3.8 \times 10^{-10}$
0.25	150	$3.6 \times 10^1$	$3.3 \times 10^3$	$6.0 \times 10^{-1}$	$3.5 \times 10^{-9}$
	200	$4.7 \times 10^2$	$4.3 \times 10^3$	$6.0 \times 10^{-1}$	$1.1 \times 10^{-9}$
	250	$1.1 \times 10^1$	$2.9 \times 10^2$	$6.0 \times 10^{-1}$	$3.2 \times 10^{-9}$
0.50	150	$3.0 \times 10^{-7}$	$1.0 \times 10^2$	$6.0 \times 10^{-1}$	$7.8 \times 10^{-8}$
	200	$2.0 \times 10^{-7}$	$2.5 \times 10^5$	$6.0 \times 10^{-1}$	$2.8 \times 10^{-9}$
	250	$1.3 \times 10^{-7}$	$5.3 \times 10^5$	$6.0 \times 10^{-1}$	$5.9 \times 10^{-6}$
0.75	150	$2.5 \times 10^{-7}$	$1.2 \times 10^9$	$6.0 \times 10^{-1}$	$5.3 \times 10^{-7}$
	200	$3.0 \times 10^{-7}$	$1.6 \times 10^4$	$6.0 \times 10^{-1}$	$7.7 \times 10^{-7}$
	250	$1.6 \times 10^{-7}$	$2.3 \times 10^6$	$6.0 \times 10^{-1}$	$2.9 \times 10^{-7}$
1.00	150	$4.0 \times 10^{-7}$	$6.3 \times 10^8$	$6.0 \times 10^{-1}$	$8.6 \times 10^{-10}$
	200	$3.0 \times 10^{-7}$	$3.4 \times 10^6$	$6.0 \times 10^{-1}$	$2.8 \times 10^{-10}$
	250	$2.0 \times 10^{-7}$	$4.5 \times 10^4$	$6.0 \times 10^{-1}$	$1.9 \times 10^{-10}$

For  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ) at temperature range  $200$ – $250$   $^{\circ}\text{C}$ , Fig. 9(a–e) shows the frequency response of the real component of modulus ( $M'$ ). The Figures show that for all compositions, the value of  $M'$  is extremely low (near zero) in the low-frequency zone, and that it continuously increases as frequency increases, suggesting a tendency to saturate at a maximum temperature. The idea that the conduction process is caused by the charge carriers' short-range mobility is supported by this saturation characteristic.<sup>30</sup> These explanations may relate to the idea that the mobility of charge carriers is brought about by an induced electric field when a restoring force isn't present. Because of the long-range mobility of charge carriers and the minimal impact of electrode polarization on the material, the low-frequency side's tiny value of  $M'$  supports the conduction phenomenon.<sup>51</sup>

The imaginary component of modulus ( $M''$ ) for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ) in a temperature range of  $200$ – $250$   $^{\circ}\text{C}$  is displayed in Fig. 10(a–e). The relaxation peak shifts towards a higher frequency as the temperature rises. The observed asymmetry in peak broadening, which illustrates the spread of relaxation time with different time constants, supports the non-Debye type of relaxation in all the compositions.<sup>52</sup>

Fig. 11(a–e) displays the complex electric modulus spectrum ( $M''$  vs.  $M'$ ) for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ( $x = 0.00, 0.25, 0.50, 0.75,$  and  $1.00$ ) at various temperatures. Electrical transport characteristics, including hopping rate and conductivity relaxation time, are interpreted by the electric modulus graph. By examining even, the tiniest changes in the materials' capacitance, it provides information on electrical processes. When it comes to

Fig. 9 Frequency dependent real part of modulus ( $M'$ ) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.



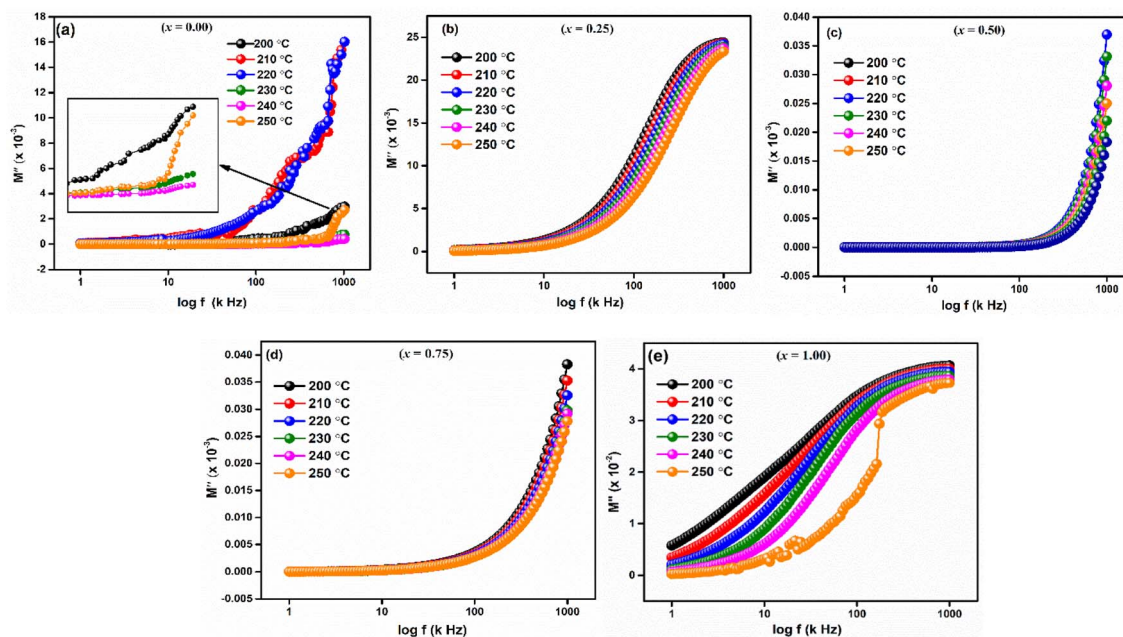


Fig. 10 Frequency dependent real part of modulus ( $M''$ ) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

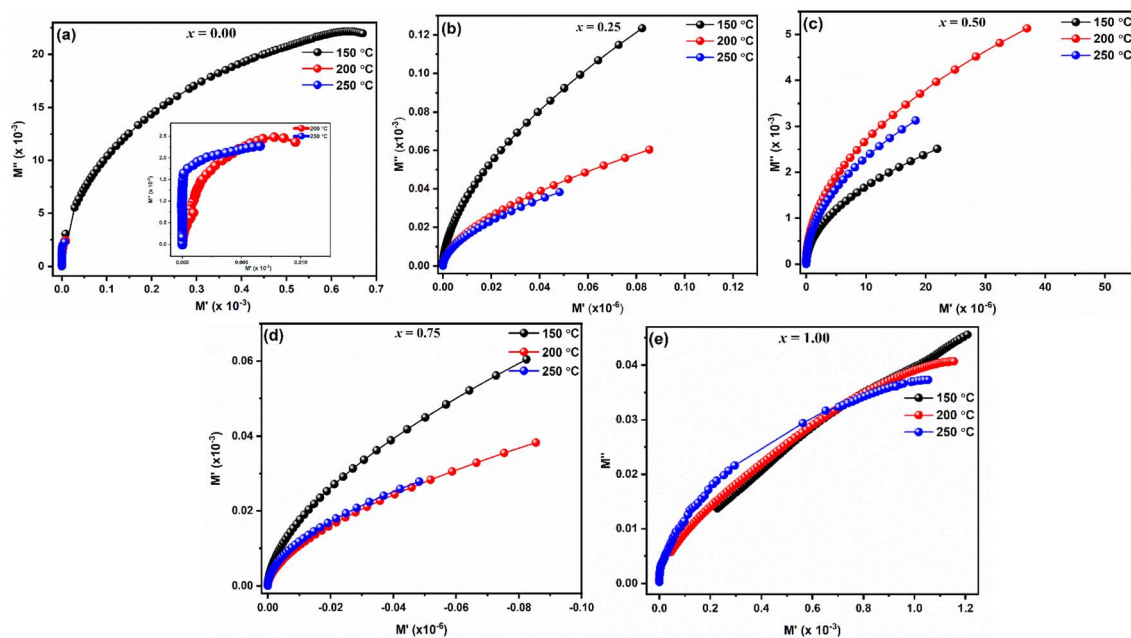


Fig. 11 Modulus plots of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

differentiating the relaxation effects from grains (conducting regions) and grain borders (resistive plates) in materials, the  $M''$  vs.  $M'$  plot performs better than the Nyquist plot of impedance ( $Z''$  vs.  $Z'$ ). The phenomena with the smallest capacitance are identified by the modulus plot, while those with the most resistance is revealed by the impedance plot. Complex modulus analysis is suitable when two materials have similar resistances but differing capacitances.<sup>53</sup> As seen in Fig. 11(a–e), the presence of a single semicircle for every composition whose centres

are located below the real axis suggests that the grain effect predominates in the conduction mechanism over the grain boundaries effect.

For every composition, a well-resolved semicircle at the higher frequency side is visible. This semicircle indicates the capacitive grain effect and shows that grains actively participate in the conduction mechanism. Furthermore, when temperature rises, a change in the grain semicircle's intercept on the  $M'_x$  axis towards lower values of  $M'$  is seen, suggesting an increase in

capacitance that supports NTCR-type behavior. For  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramic compositions, the normalized plot of  $Z'/Z'_{\text{max}}$  vs.  $\log f/f_{\text{max}}$  at various temperatures nearly overlapped on a single master curve at various temperatures (Fig. 12(a–e)). The  $Z''$  peak frequencies exhibited a minor variation in full width at half maximum (FWHM) with an increase in temperature, but they still exhibited the same form and pattern at the peak position. The temperature-independent conduction transfer mechanism was confirmed by the master curve. Additionally, the non-

symmetry of the curves was noted, suggesting that the conductivity relaxation did not behave in an exponential manner.  $Z'/Z'_{\text{max}}$  FWHM as a function of  $\log f/f_{\text{max}}$  was wider than a Debye peak's breadth, suggesting the existence of a relaxation process that is not Debye-type.

The fluctuation of  $\ln(\sigma_{\text{ac}})$  with reciprocal temperature ( $10^3/T$ ) for compounds  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  at certain frequencies (1 and 10 kHz) is displayed in Fig. 13(a–e). The NTCR behaviour of every sample is confirmed by the observation that the ac conductivity

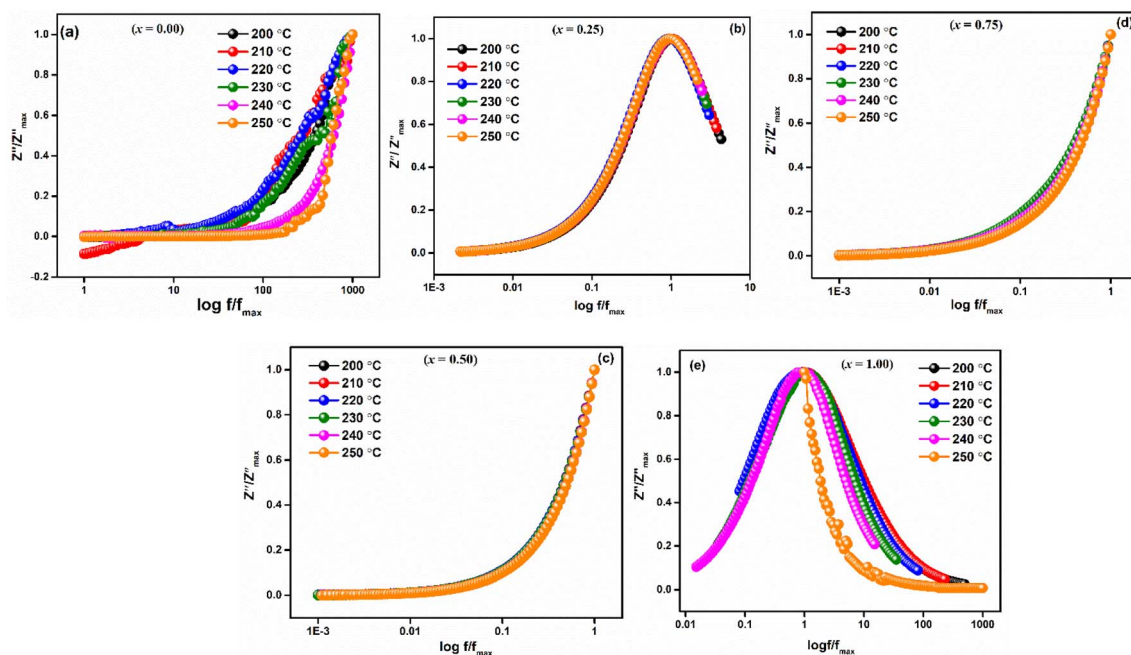


Fig. 12 Master's plots of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

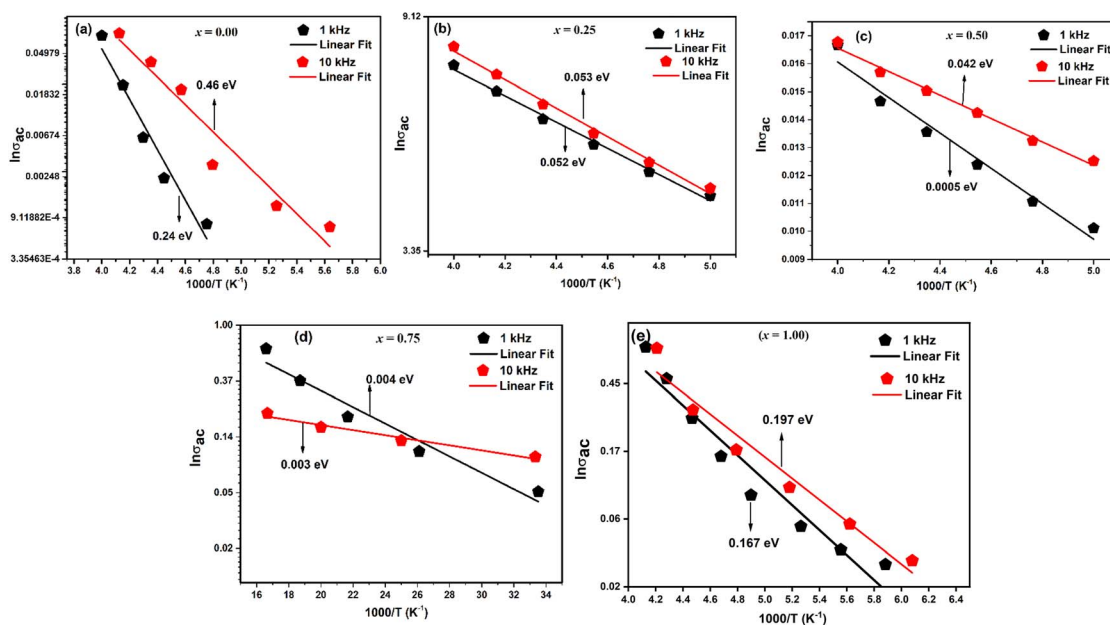


Fig. 13 AC-conductivity as a function of  $10^3/T$  for  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics (1 kHz & 10 kHz).

Table 4 Comparison of the activation energy of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics at different frequencies with the selected literature

Ceramics	Composition ( $x$ )	Activation energy ( $(E_a)$ eV)				References
		1 kHz	10 kHz	100 kHz	1 MHz	
$\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$	0.00	0.24	0.46	0.39	0.22	This work
	0.25	0.052	0.053	0.052	0.166	
	0.50	0.0005	0.042	0.018	0.044	
	0.75	0.004	0.003	0.004	0.004	
	1.00	0.167	0.197	0.149	0.028	
$\text{La}_{0.8}\text{Ca}_{0.15}\text{Na}_{0.05}\text{Mn}_{1-x}\text{Fe}_x\text{O}_3$	0.00	0.173				54
	0.25	0.248				
	0.50	0.258				
	0.75	0.289				
$\text{La}_{0.7}\text{Sr}_{0.3}\text{Mn}_{1-x}\text{Fe}_x\text{O}_3$	0.2	0.143				46
	0.3	0.108				
$\text{La}_{0.7}\text{Sr}_{0.3}\text{Mn}_{1-x}\text{Fe}_x\text{O}_3$	0.08	0.135				33
	0.10	0.138				
	0.12	0.140				
$\text{La}_{0.7}\text{Sr}_{0.3}\text{Mn}_{1-x}\text{Fe}_x\text{O}_3$	0.0	0.20				55
	0.10	0.19				
	0.15	0.17				
	0.20	0.16				
$\text{La}_{0.67}\text{Sr}_{0.33}\text{Mn}_{1-x}\text{Fe}_x\text{O}_3$	0.05	0.073				56
	0.1	0.152				

increases with temperature. Table 4 lists the activation energy of the relaxation mechanism for each sample. As the Fe content grew, the activation energy decreased, suggesting that the charge carrier concentration was hopping between the adjacent lattice sites more frequently. The obtained activation energies fell as the ceramic's Fe concentration increased, ranging from 0.24 to 0.167 eV. Because of the Fe substituent's larger lattice volume, the conduction-related specimens were released more readily and needed less energy to move.

The room temperature hysteresis loop (HL) of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  is presented in Fig. 14. While  $\text{La}^{3+}$  is typically nonmagnetic due to paired electrons,<sup>57</sup> the magnetic moments of Mn and Fe are responsible for magnetic ordering in this compound. Although  $\text{LaMnO}_3$  and  $\text{LaFeO}_3$  are known as antiferromagnetic materials, the HL of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  indicates weak ferromagnetism (FM) in all samples. The unsaturation of magnetization, even at high magnetic fields (20 kOe), suggests antiferromagnetic ordering. This weak ferromagnetism is attributed to antiferromagnetic

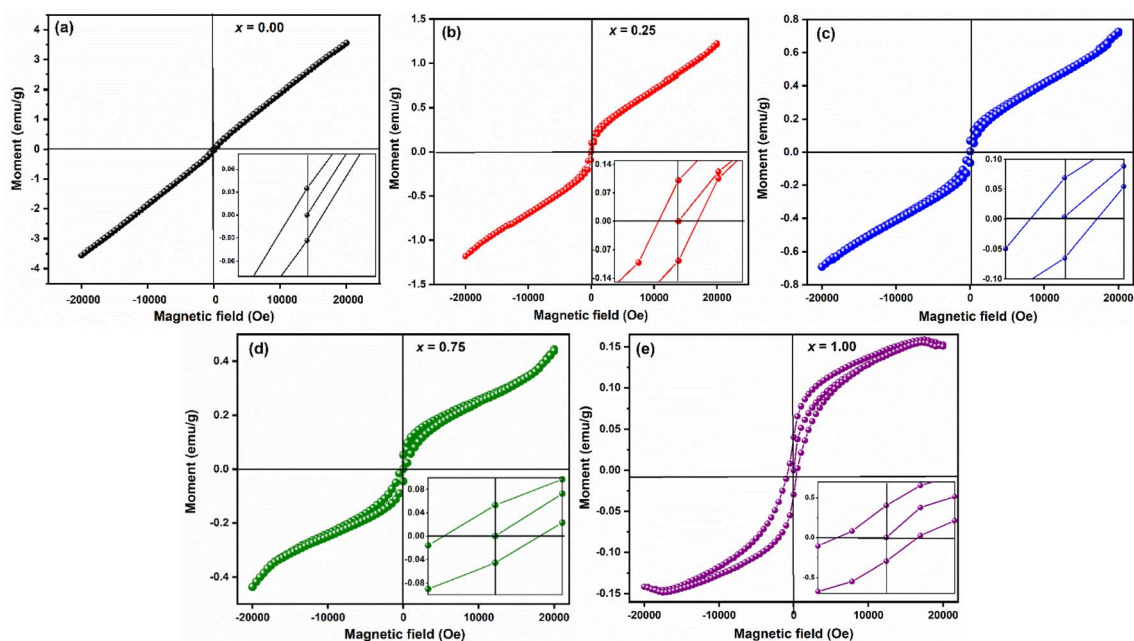
Fig. 14 Room-temperature M – H loop of  $\text{LaMn}_{1-x}\text{Fe}_x\text{O}_3$  ceramics.

Table 5 Various magnetic properties of LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ceramics

Compositions	$M_r$ (emu g <sup>-1</sup> )	$H_{C(i)}$ (Oe)	$H_{C(d)}$ (Oe)	$M_s$ (emu g <sup>-1</sup> )	SQR = $M_r/M_s$	$\mu_B$
$x = 0.00$	0.035	84.23	159.39	3.55	0.01	0.15
$x = 0.25$	0.086	230.73	268.66	1.25	0.07	0.05
$x = 0.50$	0.069	293.38	295.21	0.73	0.09	0.03
$x = 0.75$	0.054	333.97	360.93	0.45	0.12	0.019
$x = 1.00$	0.033	491.50	696.08	0.14	0.24	0.006

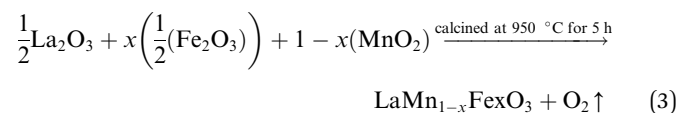
order with canted spins, induced by the presence of a Dzyaloshinskii–Moriya (DM) interaction,<sup>58</sup> which leads to a small magnetic moment in LMO and LFO NPs due to the spin canting of Mn and Fe ions.

Table 5 lists the estimated values of the various magnetic characteristics, including squareness ratio (SQR), coercive field ( $H_C$ ), remanent magnetization ( $M_r$ ), and saturation magnetization ( $M_s$ ). The  $M_s$  value was found to be decreased from 3.55 emu g<sup>-1</sup> to 0.14 emu g<sup>-1</sup> with increasing Fe-concentration. The difference in magnetization values reflects the impact of small particle size resulting in a large surface-to-volume ratio for LaMnO<sub>3</sub>.

On the other hand, coercive field ( $H_C$ ) was found to be increasing with increasing Fe-concentration, as bigger particles give a higher coercivity ( $H_c \propto D^6$ ). This is in good agreement with the law of the nano-magnetic particles. The coercivity value calculated at increasing and decreasing field (inset of Fig. 14, Table 5) indicated a shift in the HL around the origin, which confirms the presence of ferromagnetic/antiferromagnetic interfaces.<sup>59</sup> The squareness ratio ( $S$ ) of LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> was found to be nearly equal to zero, while  $H_C \neq 0$ , and  $M_r \neq 0$  which indicates that the prepared samples have particles with multiple domain sizes.

## Experimental

Polycrystalline LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ( $x = 0.00, 0.25, 0.50, 0.75, 1.00$ ) were synthesized *via* high-temperature solid-state reaction method using high purity La<sub>2</sub>O<sub>3</sub> (Sigma-Aldrich, 99.98%), MnO<sub>2</sub> (Sigma Aldrich, 99%), and Fe<sub>2</sub>O<sub>3</sub> (Merck, 99%) as the starting materials. The stoichiometric proportion of these oxides was weighted and mixed thoroughly (5–6 h) using mortar and pestle in the presence of hot distilled water as media. Following that, all of the sample reactant powders were calcined for five hours at 950 °C. Eqn (3) provides the following description of the suggested chemical reaction that would produce LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub>.



After the phase was confirmed by X-ray diffraction, a hydraulic press was used to form the powder into pallets with the appropriate dimensions. For improved densification, these green pellets were sintered for five hours at 1000 °C. Using CuK

radiation ( $\lambda = 1.5406 \text{ \AA}$ ) and a PANalytical X'Pert Pro Diffractometer at room temperature, the phase purity and crystal structure were once more examined. The surfaces of the sample were examined using Oxford Analytical instruments in combination with a Zeiss Sipra 55 field emission scanning electron microscope to determine micrographs, purity, composition, and chemical compositions. Using an LCR meter, dielectric characteristics were measured. The magnetic characteristics were measured at room temperature using a vibrating sample magnetometer (Microsense, Model: EZ9).

## Conclusions

A standard solid-state reaction technique was utilized to effectively synthesize LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> (where  $x = 0.00, 0.25, 0.50, 0.75$ , and  $1.00$ ). Rietveld refinement of XRD patterns confirms a phase transition from pure rhombohedral to orthorhombic as Fe concentration increases in the LaMnO<sub>3</sub> lattice. FESEM images reveal quasi-spherical grain morphology across all compositions. EDS spectra demonstrate elements present in stoichiometric ratios. Temperature-dependent dielectric spectra exhibit typical dispersion behavior, with high  $\epsilon$  values at low frequencies attributed to grain boundaries, supported by the Maxwell–Wagner model. Complex impedance and modulus spectroscopy confirm thermally activated conduction mechanisms. Increasing AC conductivity with temperature confirms negative temperature coefficient resistance (NTCR) behavior. Activation energy, calculated from AC conductivity, decreases with higher Fe content due to increased lattice volume, easing specimen mobility with lower energy requirements. The magnetic hysteresis loop (HL) reveals weak ferromagnetism with antiferromagnetic (AFM) ordering, possibly due to Dzyaloshinskii–Moriya (DM) interactions inducing spin-lattice canting. Squareness ratios nearing zero suggest multiple domain sizes, correlating with increasing coercive field values. Based on the above-received properties these ceramics are suitable for the spintronic applications.

## Future research scope

This study focused on the experimental tailoring of structural, morphological, electrical, and magnetic properties of LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ceramics, future research could greatly benefit from the integration of Density Functional Theory (DFT) calculations. DFT could provide detailed insights into the electronic structure, predict various material properties, and

assist in understanding the behavior of dopants at the atomic level. For instance, DFT calculations could help elucidate the electronic band structure and density of states, providing a deeper understanding of the electrical conductivity mechanisms in these ceramics. Additionally, it could offer predictions on the magnetic interactions within the material, guiding further experimental investigations.

A recent study by Wang *et al.* has demonstrated the utility of DFT in understanding competitive adsorption behaviors in environmental contexts, specifically on the facets of Goethite.<sup>69</sup> This approach could be similarly beneficial in studying LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ceramics, where DFT could help in understanding the coordination structure affinity of different dopants and their influence on the material's overall properties.

The tailored properties of LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ceramics open up a wide range of potential applications. Due to their tunable electrical and magnetic properties, these materials could be used in electronic devices such as sensors, actuators, and memory devices. Their unique magnetic properties also make them suitable for applications in magnetic storage media and spintronics. Additionally, the structural and morphological versatility of these ceramics could be exploited in catalysis and other industrial applications where specific surface characteristics are crucial. In conclusion, integrating DFT calculations with experimental research could significantly advance the understanding and development of LaMn<sub>1-x</sub>Fe<sub>x</sub>O<sub>3</sub> ceramics, enabling the optimization of their properties for various high-performance applications.

## Data availability

Data will be made available on reasonable request by corresponding author.

## Author contributions

P. T.: methodology, writing-original draft, software, visualization. K. I. N.: writing-review & editing. D. K.: writing-review & editing. P. K.: writing-review & editing. P. S.: resources. V. T.: writing-review & editing. A. S. A.: writing-review & editing. A. A.: writing-review & editing. M. E.: writing-review & editing. M. L.: conceptualization, software, visualization, data curation, supervision, writing-review & editing.

## Conflicts of interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

## Acknowledgements

The authors extend their appreciation to the Deanship of Scientific Research at King Khalid University Abha 61421, Asir, Kingdom of Saudi Arabia for funding this work through the Large Groups Project under the grant number RGP.2/545/44.

## References

- 1 X. Zhu, H. Wang, R. Hou, J. Li, Y. Li, Q. Chen and H. Zhang, *Ceram. Int.*, 2024, **50**, 13258–13265.
- 2 X. Qi, P. Ren, X. Tong, X. Wang and Y. Wan, *J. Alloys Compd.*, 2024, **989**, 174342.
- 3 X. Jin, Z. Fu, D. He, Y. Liu, X. Huang, P. Zhao, Q. Zhao and A. Chang, *Ceram. Int.*, 2024, **50**, 24743–24753.
- 4 P. Thakur, N. Sharma, D. Pathak, P. Sharma, K. Kishore, S. Dhar and M. Lal, *Emergent Mater.*, 2024, **7**, 1–28.
- 5 Q. Zheng, Y. Zhang, C. Wang, C. Zhang and Y. Guo, *Energy Fuels*, 2021, **36**, 1091–1099.
- 6 F. Li, Z. Wang, A. Wang, S. Wu and L. Zhang, *J. Alloys Compd.*, 2020, **816**, 152647.
- 7 K. Sebayang, D. Aryanto, S. Simbolon, C. Kurniawan, S. F. Hulu, T. Sudiro, M. Ginting and P. Sebayang, 2018.
- 8 W. Azouzi, W. Sigle, H. Labrim and M. Benaissa, *Mater. Sci. Semicond. Process.*, 2019, **104**, 104682.
- 9 L. Huang, L. Cheng, S. Pan, Y. He, C. Tian, J. Yu and H. Zhou, *Ceram. Int.*, 2020, **46**, 27352–27361.
- 10 S. Zhang, Z. Jia, B. Cheng, Z. Zhao, F. Lu and G. Wu, *Adv. Compos. Hybrid Mater.*, 2022, **5**, 2440–2460.
- 11 J. Su, H. Zhao, R. Yang, B. Wang, Y. Xu, X. Lin, Y. Xie and C. Wang, *Ceram. Int.*, 2023, **49**, 32049–32057.
- 12 S. Belhamra, R. Masrour, A. Jabar and E. Hlil, *Polyhedron*, 2021, **193**, 114891.
- 13 A. Ashok, A. Kumar, R. R. Bhosale, F. Almomani, S. S. Malik, S. Suslov and F. Tarlochan, *J. Electroanal. Chem.*, 2018, **809**, 22–30.
- 14 G. Takalkar, R. R. Bhosale, F. Almomani, A. Kumar, A. Banu, A. Ashok, S. Rashid, M. Khraisheh, A. Shakoor and A. al Ashraf, *Appl. Surf. Sci.*, 2020, **509**, 144908.
- 15 R. Dhahri, M. Bejar, M. Hajlaoui, N. Sdiri, M. Valente and E. Dhahri, *J. Magn. Magn. Mater.*, 2009, **321**, 1735–1738.
- 16 A. Mitra, A. Mahapatra, A. Mallick and P. Chakrabarti, *J. Magn. Magn. Mater.*, 2017, **435**, 117–125.
- 17 A. Scholl, J. Stohr, J. Luning, J. W. Seo, J. Fompeyrine, H. Siegwart, J.-P. Locquet, F. Nolting, S. Anders and E. Fullerton, *Science*, 2000, **287**, 1014–1016.
- 18 A. Mubarak, S. Tariq, B. Alsobhi, F. Hamioud and A. O. Alrashdi, *Int. J. Quantum Chem.*, 2022, **122**, e26850.
- 19 A. Mahmood, M. F. Warsi, M. N. Ashiq and M. Sher, *Mater. Res. Bull.*, 2012, **47**, 4197–4202.
- 20 S. Ye, W. Song, J. Dai, K. Wang, S. Wang, C. Zhang, J. Du, Y. Sun and J. Fang, *J. Magn. Magn. Mater.*, 2002, **248**, 26–33.
- 21 S. Nandi, J. X. Wu, P. Simon, N. Nuns, M. Trentesaux, A. Tougeriti, E. Fonda, J.-S. Girardon, J.-F. Paul and A.-S. Mamede, *Appl. Catal., B*, 2021, **296**, 120330.
- 22 J. A. Chavarría-Rubio, D. A. Cortés-Hernández, A. M. Garay-Tapia and G. F. Hurtado-López, *J. Magn. Magn. Mater.*, 2022, **553**, 169253.
- 23 W. Khan, A. H. Naqvi, M. Gupta, S. Husain and R. Kumar, *J. Chem. Phys.*, 2011, **135**, 054501.
- 24 A. Barnabé, A. Maignan, M. Hervieu and B. Raveau, *Eur. Phys. J. B*, 1998, **1**, 145–150.

- 25 R. Mahendiran, B. Raveau, M. Hervieu, C. Michel and A. Maignan, *Phys. Rev. B*, 2001, **64**, 064424.
- 26 K. De, R. Ray, R. N. Panda, S. Giri, H. Nakamura and T. Kohara, *J. Magn. Magn. Mater.*, 2005, **288**, 339–346.
- 27 P. Thakur, S. Kumari, S. Chaudhary, N. Sharma and M. Lal, *Emergent Mater.*, 2024, 1–20.
- 28 K. Ahn, X. Wu, K. Liu and C. Chien, *Phys. Rev. B*, 1996, **54**, 15299.
- 29 G. Huo, D. Song, Q. Yang and F. Dong, *Ceram. Int.*, 2008, **34**, 497–503.
- 30 X. Liu, Z. Li, P. Wu, H. Bai and E. Jiang, *Solid State Commun.*, 2007, **142**, 525–530.
- 31 S. Othmani, M. Balli and A. Cheikhrouhou, *Solid State Commun.*, 2014, **192**, 51–55.
- 32 G. Zhang and J. Lin, *J. Alloys Compd.*, 2010, **507**, 47–52.
- 33 M. Sahasrabudhe, S. Patil, S. Date, K. Adhi, S. Kulkarni, P. Joy and R. Bathe, *Solid State Commun.*, 2006, **137**, 595–600.
- 34 D. C. Kundaliya, R. Vij, R. Kulkarni, A. Tulapurkar, R. Pinto, S. Malik and W. Yelon, *J. Magn. Magn. Mater.*, 2003, **264**, 62–69.
- 35 K. Ahn, X. Wu, K. Liu and C. Chien, *J. Appl. Phys.*, 1997, **81**, 5505–5507.
- 36 S. Yusuf, M. Sahana, K. Dörr, U. Rößler and K.-H. Müller, *Phys. Rev. B*, 2002, **66**, 064414.
- 37 X. Jin, Z. Fu, D. He, Y. Liu, X. Huang, P. Zhao, Q. Zhao and A. Chang, *Ceram. Int.*, 2024, **50**, 24743–24753.
- 38 Q. Ni, A. Tang, T. Cai, Q. Zhang, Z. Zhang and L. Xiao, *Chem. Eng. J.*, 2024, **490**, 151715.
- 39 A. Vazhayil, R. Vinaykumar, J. Thomas, A. A. Jeffery, I. Hasan, N. Thomas, A. K. Yadav and Y.-H. Ahn, *J. Energy Storage*, 2024, **86**, 111145.
- 40 M. Kholil, M. Bhuiyan, M. A. Rahman, M. Ali and M. Aftabuzzaman, *AIP Adv.*, 2021, **11**, 035229.
- 41 X. Zhang, Y. Zhang, S. Wang, S. Deng, S. Liu, Q. Lin, H. Zhou, J. Zhao and G. Rao, *J. Magn. Magn. Mater.*, 2022, **563**, 169881.
- 42 J. Wang, F. Wang, L. Zhang, Y. Wei, X. Li, X. Xu, Y. Luo and Y. Jia, *J. Phys. Chem. C*, 2023, **127**, 12795–12801.
- 43 N. A. Nik-Jaafar, R. Abd-Shukor and M. A. Kamarudin, *Solid State Phenom.*, 2021, **317**, 10–16.
- 44 W. Wenwei, C. Jinchao, W. Xuehang, L. Sen, W. Kaituo and T. Lin, *Adv. Powder Technol.*, 2013, **24**, 154–159.
- 45 M. Gabal, F. Al-Solami, Y. Al Angari, A. Ali, A. Al-Juaid, K.-w. Huang and M. Alsabban, *Ceram. Int.*, 2019, **45**, 16530–16539.
- 46 S. Paul, A. K. Manokamna, D. Sharma and A. Singh, *Rasayan J. Chem*, 2022, **15**, 210–216.
- 47 S. Selvasekarapandian and M. Vijayakumar, *Mater. Chem. Phys.*, 2003, **80**, 29–33.
- 48 K. Iben Nassar and N. Rammeh, *Indian J. Phys.*, 2023, **97**, 1749–1757.
- 49 K. I. Nassar, F. Tayari, M. Benamara, S. S. Teixeira and M. P. F. Graça, *RSC Adv.*, 2023, **13**, 24023–24030.
- 50 K. Padmasree, D. Kanchan and A. Kulkarni, *Solid State Ionics*, 2006, **177**, 475–482.
- 51 M. Belal Hossen and A. Akther Hossain, *J. Adv. Ceram.*, 2015, **4**, 217–225.
- 52 B. Kaur, L. Singh, V. A. Reddy, D.-Y. Jeong, N. Dabra and J. S. Hundal, *Int. J. Electrochem. Sci.*, 2016, **11**, 4120–4135.
- 53 R. Ranjan, R. Kumar, N. Kumar, B. Behera and R. Choudhary, *J. Alloys Compd.*, 2011, **509**, 6388–6394.
- 54 S. Singh and D. Singh, *J. Alloys Compd.*, 2017, **702**, 249–257.
- 55 A. A. S. Hassan, W. Khan, S. Husain, P. Dhiman and M. Singh, *Int. J. Appl. Ceram. Technol.*, 2020, **17**, 2430–2438.
- 56 M. Baazaoui, S. Zemni, M. Boudard, H. Rahmouni, M. Oumezzine and A. Selmi, *Phys. B*, 2010, **405**, 1470–1474.
- 57 P. Coutinho, F. Cunha and P. Barrozo, *Solid State Commun.*, 2017, **252**, 59–63.
- 58 R. Saha, A. Sundaresan and C. Rao, *Mater. Horiz.*, 2014, **1**, 20–31.
- 59 D. Xue, G. Chai, X. Li and X. Fan, *J. Magn. Magn. Mater.*, 2008, **320**, 1541–1543.
- 60 X. Wang, Y. Zhang, C. Song, Z. Shen, T. Wang, K. Yang, H. Miao, J. Yang, J. Wang and X. Xu, *Chem. Eng. J.*, 2024, **479**, 147677.