


 Cite this: *RSC Adv.*, 2023, **13**, 9260

Research on the physical properties of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ spinel ferrites by the combination of optical, magnetic, and dielectric behaviors

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Ferrite compounds have recently attracted significant interest because of their multifunctional properties. This work investigates the optical, magnetic, and dielectric properties of a $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ ferrite prepared by a solid-state reaction. Raman spectroscopy analysis substantiated the presence of the 5 active modes representing the vibration of the oxygen anion at both tetrahedral and octahedral sites. The direct optical band gap was estimated to be 3.51 eV, which indicates the semiconductor behavior of the compound. A theoretical modulation of the hysteresis loop was done to confirm the dominance of the ferromagnetic contribution over the antiferromagnetic one. Furthermore, the dielectric permittivity result indicated a colossal value of the order of 10^3 . The dielectric losses are characterized by the Giuntini law to extract the relaxation process, which is hidden by the DC-conductivity process. Our results indicate the potential of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ for applications in multifunctional devices.

Received 15th February 2023

Accepted 8th March 2023

DOI: 10.1039/d3ra00985h

rsc.li/rsc-advances

1 Introduction

Research in nanotechnology is a fast-growing field at the forefront of physics and is attracting special interest from the scientific community.¹ Multifunctional materials, in particular, are investigated for their advantageous properties that are combined in one smart compound.^{2,3} A broad range of intelligent materials have been proposed and among them, ferrites have drawn significant attention and are deemed to be promoter hosts due to their extremely prominent features.^{4,5}

Significant findings in this category of ceramics generally reveal excellent physicochemical properties, such as environmental benignity,⁶ chemical stability,⁷ high electrical resistivity,⁸ and availability at low cost.⁹ It is noteworthy to mention that a promising path has been paved for spinel ferrites to be useful for potential technological applications. Some outstanding review articles have indicated that using spinel ferrites for biomedical applications may become inevitable due to their high biocompatibility¹⁰ and low toxicity.⁶ It has also

been shown that they have special properties for magnetic hyperthermia, and magnetic resonance imaging.^{11,12} Similarly, many discoveries have confirmed the development of this type of material as a suitable approach for ferrofluid technology,¹³ wastewater treatment,¹⁴ catalysts,¹⁴ selective sun absorbers,¹⁵ and medical tools.¹⁶

Some achievements and review reports relating to different spinel ferrites are proposed and investigated in detail for their properties, including lithium-based ferrites Li-Co,¹⁷ Li-Zn,¹⁸ Li-Mg,¹⁹ and Ni-Li.²⁰ Lithium ferrites are among the striking chief ferrite materials on account of their low eddy current losses, high spontaneous magnetization, excellent ferromagnetic squareness, high Curie temperature, and catalytic behavior.²¹⁻²⁴ Various reports have indicated that Li-based ferrites are usually electrically highly resistive.²⁵ However, under an applied external electric field in the microwave and radio frequency ranges, they provoke high dielectric losses.

Several strategies and synthesis pathways are used for the formation of Li-based spinel ferrites, including solid-state reactions, co-precipitation,²⁶ hydrothermal,²⁷ microemulsion,²⁸ and sol-gel methods,²⁹ and the cation distribution and crystal structure change from one ferrite to another. All these factors prevent the materials from keeping the usual properties of a typical spinel ferrite structure¹⁰ but different preparation processes and thermal treatment allow such difficulties to be overcome.

Lithium manganese ferrite is one of the most interesting ferrite materials. The XRD diffraction peaks correspond to those of the cubic $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ spinel.³⁰ Our previous results³⁰ revealed that the compound could be a multifunctional

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material with the potential to drive technological innovation forward. The electrical properties of the compound have proven that it can be used in various applications such as optoelectronics, photodetectors, photovoltaics, and full-cell low temperature-co-fired ceramics (LTCCs).³⁰ Moreover, our sample can be suitable for device performance ameliorating the sensitivity of uncooled infrared bolometers after comparing its (TCR) with other previous works. It also represents anomalous behaviors because it combines magnetic and non-magnetic elements. In this kind of ferrite, the magnetic properties are governed by the spin coupling of the 3d electrons.³¹

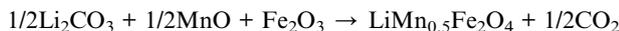
Our previous electrical search³⁰ is insufficient to evaluate, describe, and clarify all possible fields where it is suitable for use. Therefore, in this scientific report, an attempt has been made to analyze the magnetic, and optical properties. Besides, we have undertaken further studies about dielectric relaxation and modulus formalism depending on temperature and we present an exhaustive idea about the studied sample to make it useful in diverse applications.

2 Experimental details

2.1 Preparation method

The conventional solid-state method was used to prepare the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ spinel ferrite ceramic. The protocol for preparation is depicted in detail in our previous paper.³⁰

All precursors (Li_2CO_3 , MnO , Fe_2O_3) were purchased from Sigma-Aldrich (purity of 99.99%) and utilized without further purification. The pure chemical reactants were at first dried and intimately mixed in stoichiometric quantities according to the reaction as follows:



The obtained mixture was well-ground and pressed into a pellet (disk-shaped) of about 1 mm in thickness, which is essential for a homogenous mixture of reactants and to facilitate element diffusion. The pellet was calcined at 850 °C in an electrical muffle furnace for 7 h. In the final step, it was subjected to many sintering heating cycles up to 1100 °C for 2 h, accompanied by grinding and pelletizing yields to obtain a homogenous, pure, and compact crystallized sample.

2.2 Characterization techniques

Raman scattering measurements were recorded at room temperature using a SENTERRA Spectrometer (Bruker) with green laser excitation of 540 nm. Fourier-transform infrared (FTIR) spectra were obtained on a Bruker Tensor 27 spectrophotometer.

To determine the energy bandgap (E_g) and confirm the semiconductor nature of the sintered sample from the absorption spectra, UV-vis spectroscopy analysis was employed. The hysteresis cycle of the ferrite was carried out using a SQUID magnetometer at 5 K and the ambient temperature (300 K) under a magnetic field up to 10 T.

To prepare the sample for dielectric measurement, a thin layer of silver was coated on both faces of the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ pellet by a vacuum thermal evaporator. The measurements were deduced from impedance spectroscopy with a Janis VPF800 cryostat and an Agilent 4294A analyzer in the explored range of frequency [100–10 MHz]. We measured the real and imaginary parts of the complex impedance Z^* versus the frequency.

3 Results and discussion

3.1 X-ray diffraction analysis

The X-ray diffractogram patterns of the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ sample were recorded at room temperature, as shown in Fig. 1. All the reflection peaks are well indexed in the cubic system with $Fd\bar{3}m$ space group. The absence of impurity peaks proves the pure and high quality of the produced compound. All the structural parameters deduced from the Rietveld analysis were mentioned in our previous work.³⁰

3.2 Scanning electron microscopy analysis

The morphology of the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ sample, carried out by scanning electron microscopy (SEM), is depicted in Fig. 2. The grains are polygonal-shaped and polyhedron-shaped. The average estimated diameter of the grain was relatively large, of the order of 2.48 μm , as detailed in our preceding paper.³⁰

3.3 Raman spectroscopy

Raman spectroscopy is broadly used as an effective microscopic technique to determine more structural properties and complementary details of the prepared compounds. It is also used to explore the short-range cationic disorder due to its high sensitivity and its lofty spatial resolution to investigate the vibrational levels in spinel ferrites.³²

The factor group process foretells the subsequent vibrational modes in spinel by the following:

$$\Gamma = A_{1g}(\text{R}) + E_g(\text{R}) + 3F_{2g}(\text{R}) \quad (1)$$

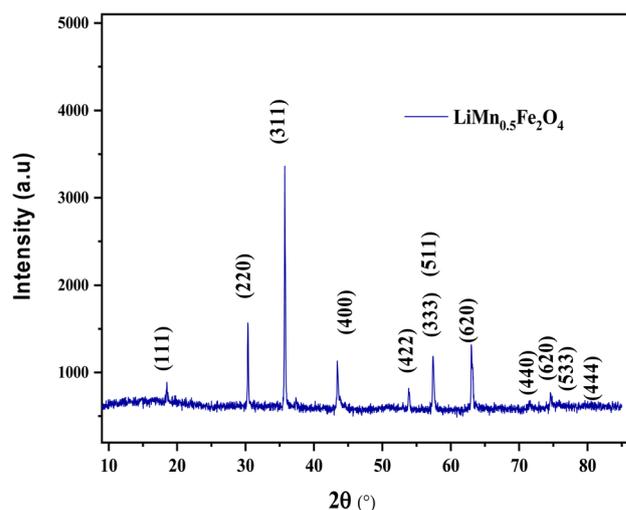


Fig. 1 XRD pattern of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$.

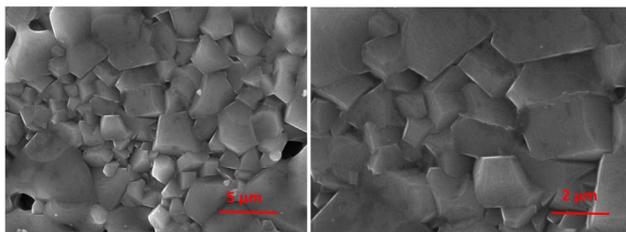


Fig. 2 SEM images of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ at scales of $5\ \mu\text{m}$ and $2\ \mu\text{m}$.

where (R) represents Raman active vibrational modes. $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ has a cubic structure belonging to the $Fd\bar{3}m$ space group³⁰ and according to group theory, 5 Raman active modes ($A_{1g} + E_g + 3F_{2g}$), acquired under ambient conditions, are recognized for this structure space group. They are dependent on the motion of both A and B site cations and oxygen ions.³³

The Raman spectrum was acquired at room temperature in the spectral range varying from 150 to $800\ \text{cm}^{-1}$ and illustrated in Fig. 3. It was properly analyzed and deconvoluted *via* fitting by the Lorentzian peak shape function to evaluate the full width at half maximum (FWHM), and the positions of obtained bands. The different peaks were assigned pursuant to the theoretical vibrational mode. The detected Raman modes match well with the literature reports. They were centered at 192 , 259 , 360 , 490 , and $612\ \text{cm}^{-1}$ and allocated to $F_{2g}(1)$, E_g , $F_{2g}(2)$, $F_{2g}(3)$, and $A_{1g}(1)$ Raman vibrational modes, respectively, as shown in Table 1.

The A_{1g} mode attributed to the uppermost frequency is represented as though a symmetric stretching exists between the oxygen atom and the iron metal ion located in tetrahedral sites.³⁴ The E_g Raman mode is assigned to the symmetric bending motion of the oxygen anion with respect to the iron situated at the octahedral site. The second highest frequency $F_{2g}(3)$ mode is the anti-symmetric bending motion of the oxygen anion within the octahedral site. $F_{2g}(2)$ is the mode antisymmetric stretching vibration mode of the oxygen anion with respect to the iron at the octahedral site (it is displaced in

Table 1 Raman vibrational modes of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ with the assigned polyhedron

Position (cm^{-1})	Raman mode	FWHM (cm^{-1})	Assigned polyhedron
192	$F_{2g}(1)$	118	FeO_6 Octahedron
259	E_g	120	FeO_6 Octahedron
360	$F_{2g}(2)$	120	FeO_6 Octahedron
490	$F_{2g}(3)$	121	FeO_6 Octahedron
611	A_{1g}	120	FeO_4 Tetrahedron

differing directions) and the fifth active Raman mode $F_{2g}(1)$ represents a complete translation development of the whole at the octahedral unit in the spinel lattice.

Raman peak frequencies higher than $600\ \text{cm}^{-1}$ were assigned to the Raman active mode A_{1g} , related to the motion of O^{2-} ions within the tetrahedral sublattice (site A), while the rest of the peaks are due to the vibration of the oxygen atom and octahedral group (site B).³⁵ The absence of peaks attributed to any foreign oxide phase (iron oxide phase) is probing the chemical purity of the sample.

The Raman spectrum detects the attendance of extra peaks at $611\ \text{cm}^{-1}$ and $705\ \text{cm}^{-1}$ attributed to $A_{1g}(1)$ and $A_{1g}(2)$. Biplab K. Chatterjee *et al.*^{36,37} suggested that they are probably linked to a breakdown of the momentum conservation rule because the wavelength of exciting radiation is higher than the diameter of the particle. Other authors estimate that this behavior is a typical feature of a mixed or inverse spinel structure.³⁸

3.4 FTIR studies

Fourier transform infrared (FTIR) measurements aid in determining and monitoring the degree of crystallization in the studied crystalline compound. The FTIR results of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ after absorption in the wave number range of 400 – $2500\ \text{cm}^{-1}$ can be viewed in Fig. 4. The presence of two infrared absorption bands in the high-energy region (at low frequency) proves that our sample is well-crystallized and confirms the presence of the spinel structure, as reported in Waldron's classification.³⁹ The absorption band (ν_B) observed at $425\ \text{cm}^{-1}$ is attributed to metal–oxygen stretching in the octahedral site while the second band (ν_A) at $555\ \text{cm}^{-1}$ is related to the stretching of metal–oxygen at the tetrahedral site. Given that the bond length in the octahedral is longer than that in the tetrahedral site, this can explain the fact that the band frequency of the stretching vibration in the tetrahedral site is higher than that of the octahedral site.⁴⁰

3.5 Absorption study

Our target in this section is to estimate the light efficiency and the quality of our prepared spinel ferrite compound by

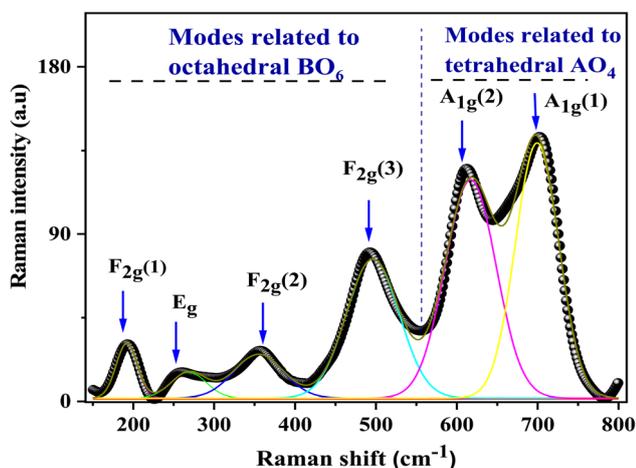


Fig. 3 Raman spectra of the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$.



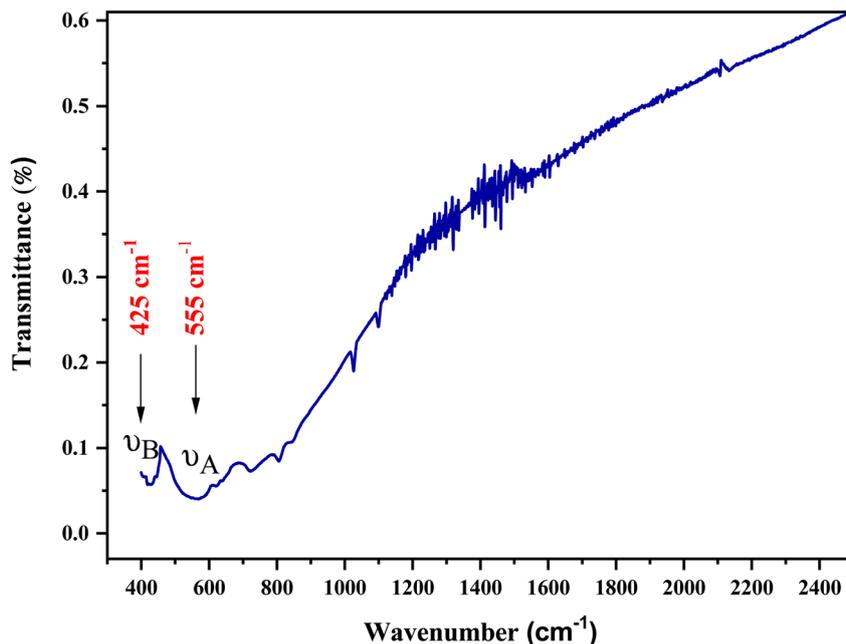


Fig. 4 Fourier transformation infrared spectrum of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$.

presenting and discussing its fundamental optical properties. The ultraviolet-visible absorbance spectrum was registered in the range extending from 200 to 800 nm at ambient temperature (Fig. 5). According to previous reports on spinel ferrites, the absorption can be related to the Fe^{3+} transitions (from $3d^5$ to $3d^4 4s^1$) or the photoexcitation process, which appears where the electron leaves the valence band (VB) (O_{2p} level) to go to the conduction band (CB) (Fe_{3d} level) after excitation by visible light.⁴¹

The optical bandgap (E'_g) denotes the photon energy that should be absorbed by the electron to transit from the (VB) to

(CB). This value is obtained by using both the Kubelka-Munk function and the Tauc plot.⁴²

On the one hand, the Kubelka-Munk function is as follows:

$$F(R) = \frac{(1 - R)^2}{2R} = \frac{\alpha}{s} \quad (2)$$

where R is the reflection factor, α is the absorption coefficient, and s is a scattering factor.

On the other hand, $F(R)h\nu = A_1[h\nu - E'_g]^n$, where A_1 is a steady parameter that reveals the degree of disorder in the structure of our compound, $h\nu$ is the incident photon's energy

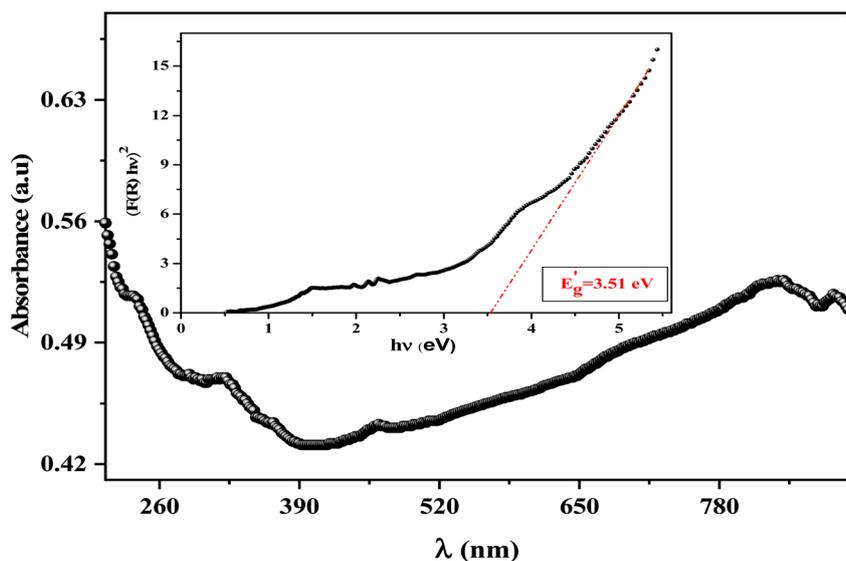


Fig. 5 UV-visible absorbance spectra of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ at room temperature; the inset shows the Tauc plot for the determination of the bandgap value for the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ compound.



and the exponent n has 2 values (1/2, and 2) according to the type of allowed optical transition. n is equal to 1/2 in the direct transition and equal to 2 for the indirect transition. $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ shows a direct band gap like other compounds belonging to the spinel ferrite family.^{41,43,44} The inset in Fig. 5 exhibits the variation of $(F(R)h\nu)^2$ as a function of $h\nu$, the forbidden band value (E_g) is obtained by the extrapolation line of the curve to the energy axis (at the origin of the y-axis). The optical bandgap of the sample is 3.51 eV which falls in the range of a semiconductor bandgap. The semiconducting nature of this material makes it a promising candidate for photodetectors and photovoltaics applications.⁴⁴

To ensure the value of the energy gap, the curve of $dR/d\lambda$ versus wavelength was plotted in Fig. 6. It gives a shred of evidence for the presence of an optical bandgap that lies at 3.57 eV, close to what is obtained by Kubelka–Munk. Thereby, we confirmed the direct type of transition and the semiconducting nature of the synthesized ceramic.

The energy gap value is significantly higher than the energy prerequisite for water splitting ($E > 1.23$ eV). This allows the use of the material for photocatalysis under visible light.⁴¹ The refractive index (n_0) has an essential role in the control of the optical and electric properties of the studied material. Also, the knowledge of the index n_0 indicates the compactness of our sample and aids in gaining insight into its good quality.⁴⁵

In this paper, the refractive index of ferrite is determined according to Moss's empirical relation as reported:³²

$$E'_g n_0^4 = 104 \text{ eV} \quad (3)$$

The obtained value of the refractive index was 2.37. This reflects the good quality of our compound. Likewise, the high-frequency optical dielectric constant (ϵ_∞) of our prepared ferrite is related to the refractive index (n_0) as stated in the following relation:²²

$$\epsilon_\infty = n_0^2 \quad (4)$$

The optical dielectric is equal to 5.62. The refractive index and high-frequency dielectric constant are similar to those

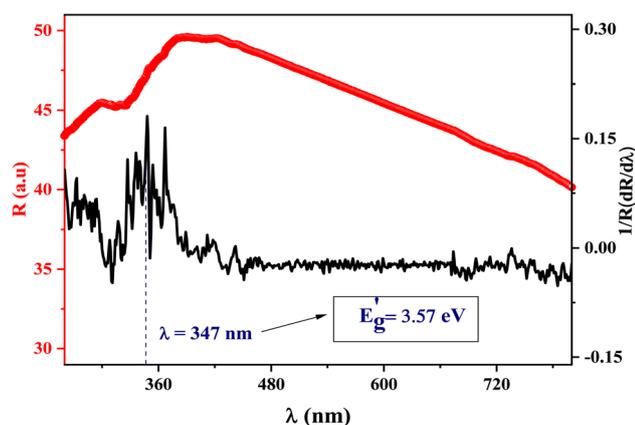


Fig. 6 The smoothed reflectance spectra of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ with its first derivative ($1/R$) ($dR/d\lambda$) curve.

obtained for the material, which is considered a good candidate in the modeling of heterostructure solar cells as well as in laser optoelectronic devices.⁴⁶ Ultimately, the prepared spinel ferrite $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ is full of promise in the optical domain.

3.6 Magnetic study

The magnetic hysteresis loop $M(H)$ of the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ powder at 5 K and 300 K are depicted in Fig. 7a and b, respectively, with the applied field reaching 10 T. At room temperature, the hysteresis loop shows a small surface area that highlights the low energy loss over the magnetization–demagnetization procedure.⁴⁷ Hence, the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ is suitable for magnetic refrigeration application.

The hysteresis cycle at 5 K showed a significant remanence which became negligible at 300 K (Fig. 7(b)). The coercivity is low, which may be related to the superparamagnetic behavior of this material. The large size of particles D_{SEM} deduced from scanning emission microscopy (D_{SEM} is about 2.48 μm (ref. 30)) indicates great potential for developing and improving superparamagnetic nanoparticles from metal oxide material, especially spinel ferrite.⁴⁸

This makes the studied sample suitable for the biomedical domain as a contrast improvement instrument for magnetic resonance imaging (MRI) and hypothermia, as well as for designing polyvalent tools to change magnetization with a low outer magnetic field.⁴⁷

We found that the magnetization reached saturation at weak fields below 0.5 T. This behavior is explained by the alignment of the spins of the fine nanoparticles in the direction of the applied field. Two types of interactions can exist in the sample, namely, antiferromagnetism (AFM) and ferromagnetism (FM). In the next step, we aspired to better understand the origin of magnetization and segregate the antiferromagnetic (AFM) and ferromagnetic (FM) contributions in the synthesized sample. A theoretical modulation of hysteresis loops was done according to the following equation:⁴⁹

$$M(H) = \chi H + 2 \frac{M_{\text{FM}}^{\text{S}}}{\pi} \tan^{-1} \left\{ \left(\frac{H + H_{\text{CI}}}{H_{\text{CI}}} \right) \tan \left(\frac{\pi M_{\text{FM}}^{\text{R}}}{2 M_{\text{FM}}^{\text{S}}} \right) \right\} \quad (5)$$

where the first part put forward the linear antiferromagnetic contributions (AFM) and the second represents the ferromagnetic quote-part (FM).⁴⁹ M_{FM}^{S} and M_{FM}^{R} represent the FM saturation and remanant magnetization, respectively. H_{CI} is an intrinsic coercive field. χ is the AFM susceptibility. All adjusted parameters are reported in Table 2. It shows that the magnetization of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ is due to the FM contribution, which represents $\sim 90\%$ vs. the AFM contribution and does not exceed $\sim 10\%$ of the total magnetization at both temperatures 5 K and 300 K.

In the framework of investigation in the relation between the saturation magnetization (M_{S}) and coercive field (H_{C}), Stoner–Wohlfarth estimates the value of constant anisotropy related to the coercivity using the below relation:⁵⁰

$$K_1 = \frac{0.98 H_{\text{C}}}{M_{\text{S}}} \quad (6)$$



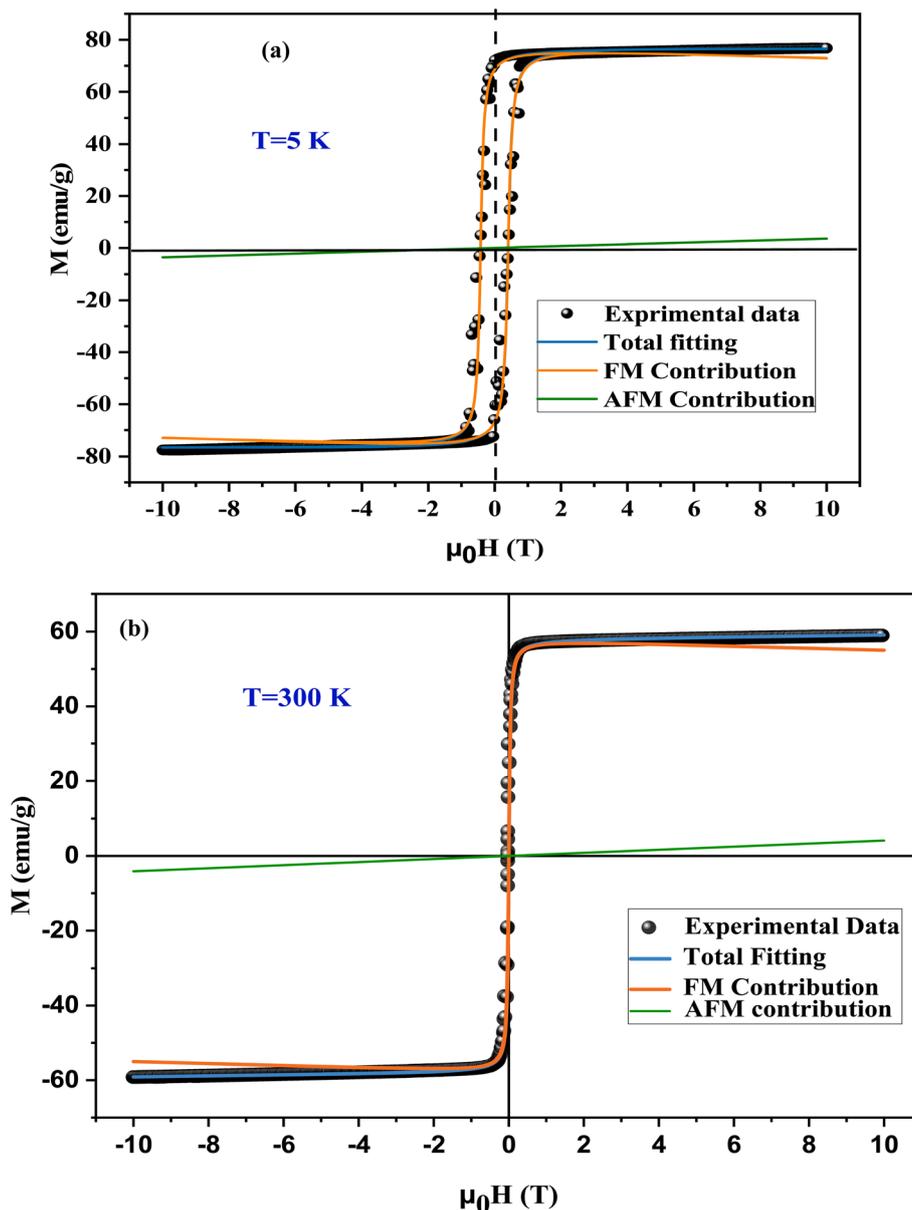


Fig. 7 The magnetization curves versus magnetic field at 5 K (a) and 300 K (b) for $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$.

This relation shows that constant anisotropy K_1 is proportional to H_C and inversely proportional to M_S which is confirmed by other studies.³²

To go further, we ascertained whether our sample is useful for high-frequency microwave devices. Here, we determined the microwave frequency (ω_m) utilizing the given relation:³²

$$\omega_m = \delta_1 8\pi^2 M_S \quad (7)$$

where δ_1 designates the gyromagnetic ratio ($\delta_1 = 2.8 \text{ MHz Oe}^{-1}$). The obtained value is 11.6 GHz, which is in the same range as those found in other materials used in this application.^{51,52}

Table 2 Theoretical parameters obtained from fitting the hysteresis loops of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$

	AFM contribution		FM contribution			%	
	χ ($10^{-5} \text{ emu} \times \text{Oe g}^{-1}$)	H_{Cl} (Oe g^{-1})	M (10 T)	M_{FM}^S (emu g^{-1})	M_{FM}^R (emu g^{-1})	FM	AFM
5 (K)	2.776	4241	76.62	73.02	68.51	95.3	4.7
300 (K)	3.1267	40	59.10	54.97	1.01	93	7



$\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ can, therefore, be considered an aspiring candidate for the domain of microwave frequency bands.

3.7 Dielectric study

The dielectric permittivity affords important insight into the prepared material. It represents the ability of the material to polarize under the effects of applied electric fields.⁵³ Spinel ferrites are usually good dielectric materials,⁵⁴ depending firmly on some variables such as the cation distribution among the interstitial sites, grain size, stoichiometries, and the synthesis procedure. The complex permittivity of the studied material was expressed by the following relation:⁵³

$$\varepsilon^* = \varepsilon' - j\varepsilon'' = \frac{1}{j\omega C_0 Z^*} \quad (8)$$

It is well known that ε' is the real part of the dielectric permittivity that gives an idea of the stored energy and ε'' is the imaginary part that describes dissipation energy in the material. C_0 denotes the vacuum capacitance and ω is the angular frequency ($=2\pi f$).

The recorded dielectric data of the real permittivity ε' is plotted in Fig. 8. A high dielectric permittivity value is presented in the low-frequency region, which is promising in many technological applications.⁵⁵ The dielectric constant diminishes continuously with increasing frequency. This decreasing trend is observed in many investigations of different spinel materials.^{56,57} Exploring the structure of ferrites, which are generally composed of grains separated by grain boundaries, aids in understanding the behavior of ε' . Pubby *et al.*⁵⁸ suggested that grain boundaries impose dielectric properties at low frequencies.⁵⁹ Mostly, the grains are good conductors and the grain boundaries are poor conductors. This conductivity difference induces an accumulation of load carriers at boundaries, which creates space charge polarization. The authors explain the effect

of the behavior of the real part of the permittivity with the variation of the temperature and the frequency by the reduction of the polarization.⁶⁰ There exist 4 types of polarization, namely, interfacial, electronic, ionic, and dipolar polarization, which have an apparent effect on the dielectric constant evolution but their contribution declines with frequency.⁶¹

On our measurement frequency range, we can only identify dipolar or interfacial polarization. These types of polarization are highly temperature-dependent.

Fig. 8 (inset) depicts the temperature dependence of the dielectric constant of our compound at selected frequencies. In the range [200–250 K], ε' increases gradually with temperature but its variation is less marked and almost invariant with the increase in frequency. This result may be due to the orientation of polarization, which is related to the molecule's thermal motion.⁶² It is explained by the fact that the dipoles remain frozen and cannot orient themselves at low temperatures.⁶³ In the temperature range (>250 K), the dipoles can rotate loosely and the molecular interaction energy effect becomes lower than the thermal energy. The bound load carriers gradually obtained thermal excitation energy. This in turn makes them able to reply to the change in the externally applied field more easily.⁶² Therefore, it enhances their contribution to the polarization, resulting in an increase in the dielectric constant with temperature.

Fig. 9 shows the variation of ε'' as a function of frequency in a temperature range of [200–380 K]. It is clear that ε'' also decreases with the frequency until it becomes stable in the high-frequency range (above 10^4 Hz). This behavior can be explained by the migration of electrons between Fe^{2+} and Fe^{3+} and the ion polarization loss, which can be considered as the cause of the very high value of dielectric loss ε'' at low frequency. Above 10^4 Hz, only the exchange ions can go along with the applied electric field frequency guide to the decrease in ε'' across the studied temperature range.

Fig. 10 shows the evolution of ε'' as a function of frequency at several temperatures on a logarithmic scale. In ε'' can be analyzed according to Guitini's theory, as given by the following:⁶⁴

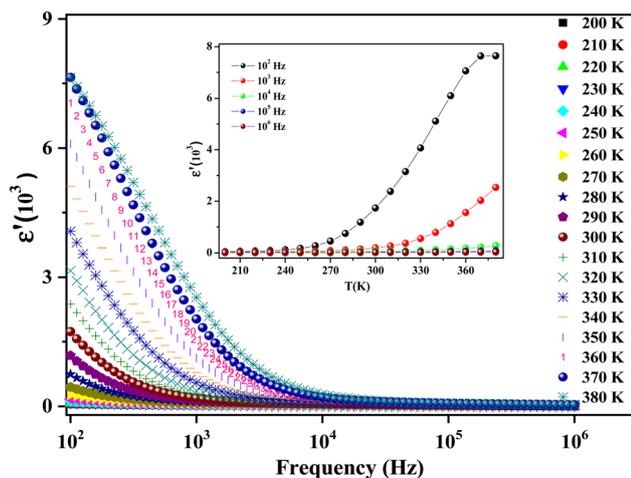


Fig. 8 The dielectric data of the real permittivity (ε') at several temperatures [200–380 K] for the synthesized ferrite. The inset shows the temperature dependence of the real part ε' of the permittivity at selected frequencies.

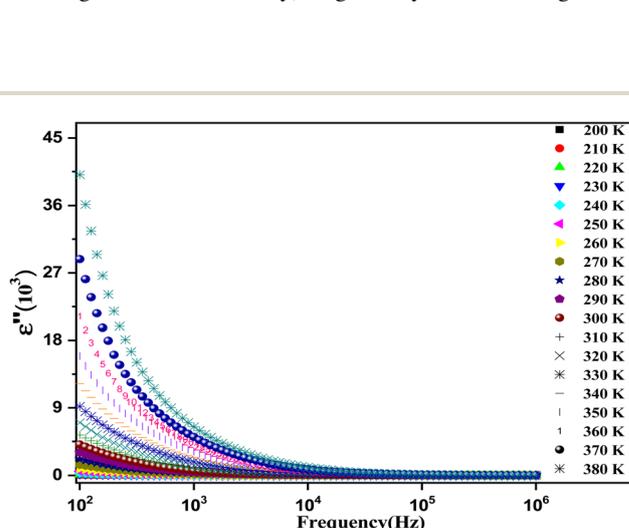


Fig. 9 The recorded dielectric data of the imaginary permittivity (ε'') at several temperatures [200–380 K] for the synthesized ferrite.



$$\varepsilon'' = (\varepsilon_s - \varepsilon_\infty) 2\pi^2 N \left(\frac{nq^2}{\varepsilon_0} \right)^3 K T \omega^m \tau_0 (W_m)^4 \quad (9)$$

where ε_s is the dielectric constant at the low frequencies, ε_∞ is the dielectric constant at the high frequencies, N is the state concentration, n is the hopping electron number, W_m is the barrier height, q is the elementary electron charge and τ_0 signifies the relaxation time.

This law can be simplified in another form:⁶⁵

$$\varepsilon'' = C(T)\omega^m \quad (10)$$

where $C(T)$ is a constant that depends only on temperature and m is an exponent describing the interaction between electric dipoles, which is expressed as follows:⁶⁵

$$m = -\frac{4k_B T}{W_m} \quad (11)$$

In this expression, k_B is the Boltzmann constant and W_m is the energy required for charge carriers to leap above the potential barrier. The obtained values of parameter m versus temperature are shown in the inset in Fig. 10. We observed that the variation of m was proportional to the temperature. The exponent m decreased with increasing temperature in the intervals [200–270 K] and [340–380 K], which matches well with the correlated barrier hopping (CBH) model. Nevertheless, m increases with temperature in the intermediary region, so the nonoverlapping small polaron tunneling (NSPT) mechanism is predominant. These results are well correlated with our previous electrical findings for this material.³⁰

Studying the temperature effect on the evolution of the imaginary part of the dielectric permittivity can give us a clear insight into the dielectric relaxation number within a system. From Fig. 10, we observed the absence of the relaxation phenomenon in the frequency range, which was concealed by

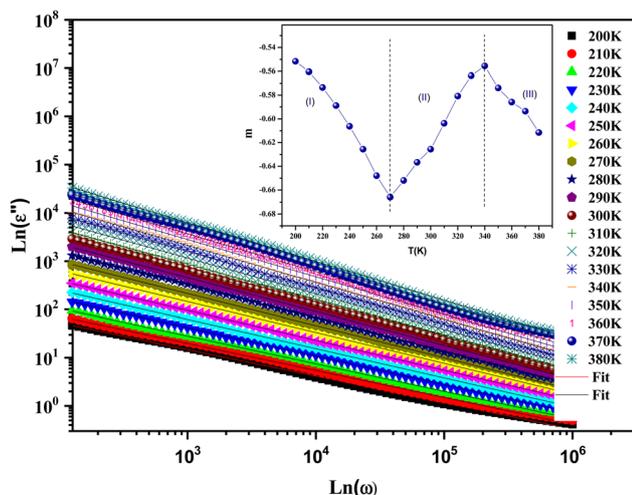


Fig. 10 The frequency dependence of $\ln(\varepsilon'')$ at several temperatures [200–380 K] for the synthesized ferrite. The black solid line depicts the fit of the data with relation (11). The temperature evolution of the exponent (m) is shown in the inset of the figure.

the DC-conductivity mechanism. It can be elicited using the following relation:⁶⁶

$$\varepsilon''_1 = \varepsilon''_{\text{measured}} - \varepsilon''_{\text{DC}} \quad (12)$$

where $\varepsilon''_{\text{DC}}$ is the contribution of DC-conductivity to the imaginary part of the dielectric permittivity, it can be calculated by the following equation:⁶⁶

$$\varepsilon''_{\text{DC}} = \frac{\sigma_{\text{DC}}}{\varepsilon_0 \omega} \quad (13)$$

where σ_{DC} is the static component of the conductivity, and ε_0 is the vacuum permittivity. As expected, Fig. 11 illustrates peaks that are shifting toward high frequencies with the increase in temperature. The obtained result asserts that this approach is an advisable tool for studying the dielectric relaxation phenomenon in the material.⁶⁷

3.8 Polarizability study

A new method was suggested by Scaife⁶⁸ to investigate the dielectric response of the sample. It is based on the substitution of the complex permittivity by the polarizability according to the relation:⁶⁹

$$\alpha^* = \alpha' - i\alpha'' \quad (14)$$

where the imaginary part α'' can be easily obtained as follows:

$$\alpha'' = \frac{3\varepsilon''}{(\varepsilon' + 2)^2 + (\varepsilon'')^2} \quad (15)$$

Complex polarizability usually refers to the tendency of the studied material to polarize and ease the analysis of the dielectric results of various samples. It unveils the new relaxations and clarifies the previous ones. The dependence temperature of α'' is plotted in Fig. 12(a). It exhibits well-separated relaxation peaks that are centered at angular frequency ω_α (where $\omega_\alpha = 2\pi f_\alpha$) and move towards the high frequency with the increase in temperature.

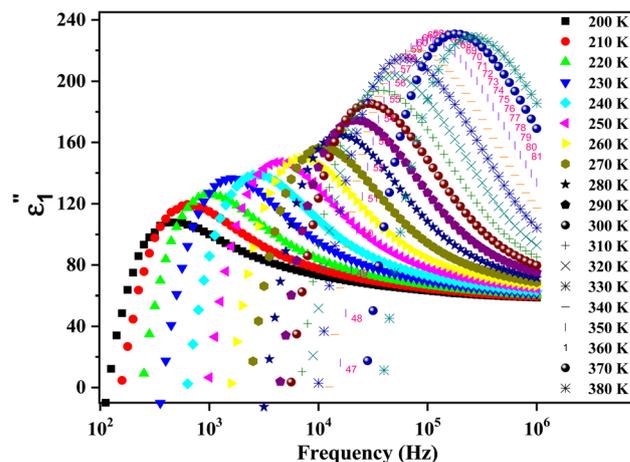


Fig. 11 The variation of ε''_1 at various temperatures for the prepared compound.



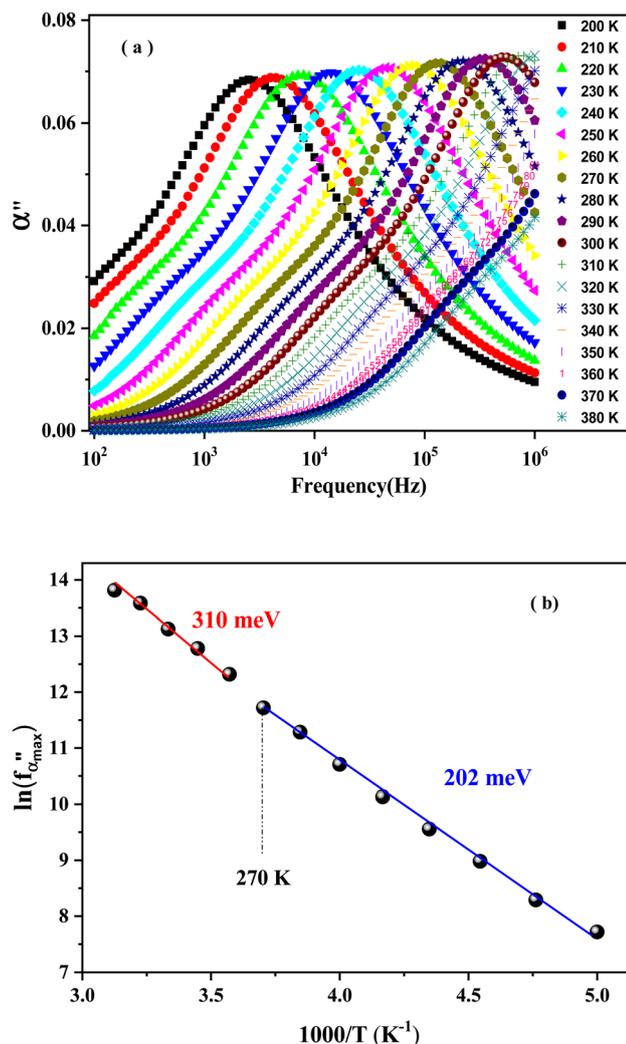


Fig. 12 The frequency dependence of polarizability at different temperatures (a). The variation of $\ln(f''_{\alpha_{\max}})$ vs. $(1000/T)$ (b).

The logarithmic variation of ω_{α} versus the inverse of the temperature is illustrated in Fig. 12(b). This curve can be described according to the Arrhenius relation:⁶⁴

$$\omega_{\alpha} = \omega_0 \exp\left(-\frac{E_a}{k_B T}\right) \quad (16)$$

where ω_0 is a constant and E_a is the activation energy. The evaluated activation energies are 202 meV for [200–270 K] and 310 meV for [280–310 K].

The quantitative loss of energy (the dielectric loss tangent) in an assorted dielectric material is generally caused by different physical processes, for example, electrical conduction, or dielectric relaxation.⁶⁹ Fig. 13 exhibits the temperature-dependent loss tangent ($\tan \theta$). $\tan \theta$ increases with the increase in temperature. The peaking behavior may be related to the arising relaxation effect.⁷⁰ The applied electric field period fits well with those of the relaxation process. The peaks move towards the higher frequency side ($>10^4$ Hz) with increasing

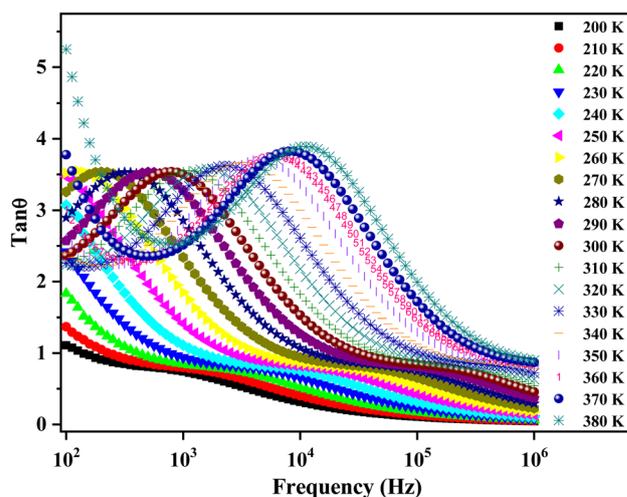


Fig. 13 The variation of dielectric loss as a function of frequencies at various temperatures.

temperatures, showing a pronounced thermally activated relaxation phenomenon. The latter is associated with the curtailment of resistance with the increasing frequencies.⁶⁴

3.9 Modulus study

In this section, we pay attention to the utility of modulus formalism. Its main role is to supply us with an idea about the electrical process implicated in our investigated material and to study the same as the function of temperature over a broad range of frequencies.

The total complex electric modulus follows the equation⁷¹

$$M^* = M' + M'' = i\omega C_0 Z^* \quad (17)$$

where M' and M'' are the real and imaginary parts of the modulus, respectively. C_0 denotes the geometrical capacitance and Z^* is the complex impedance. The variation of the real part of the electrical modulus as a function of frequency with the rising of temperature is depicted in Fig. 14. As shown, the real part of the modulus exhibits a weak value (bring closer to zero) at the low frequencies for all temperatures. This corroborates that the contribution of the electrode polarization is almost absent in the structure.⁷¹ M' increased with the increase in frequency, showing a dispersive region assigned to the short-range mobility of load carriers in our compound that is attributed to the conduction process.⁷² Then, it achieved a maximum at a high frequency of around 10^6 Hz.

Fig. 14 depicts the frequency dependence of the imaginary part of the modulus at different temperatures. The curves of M'' illustrate a broad peak that represents the assembly of the two peaks at temperatures below 310 K. For the rest, we assume the presence of a broad peak in the high frequencies exceeding the scale of our available impedance analyzer. The peak in the lower frequency range is related to the grain boundaries effect.⁷³ The pronounced peaks in the higher frequency range are associated with the bulk effect (the grain's effect).⁶⁷ The more the



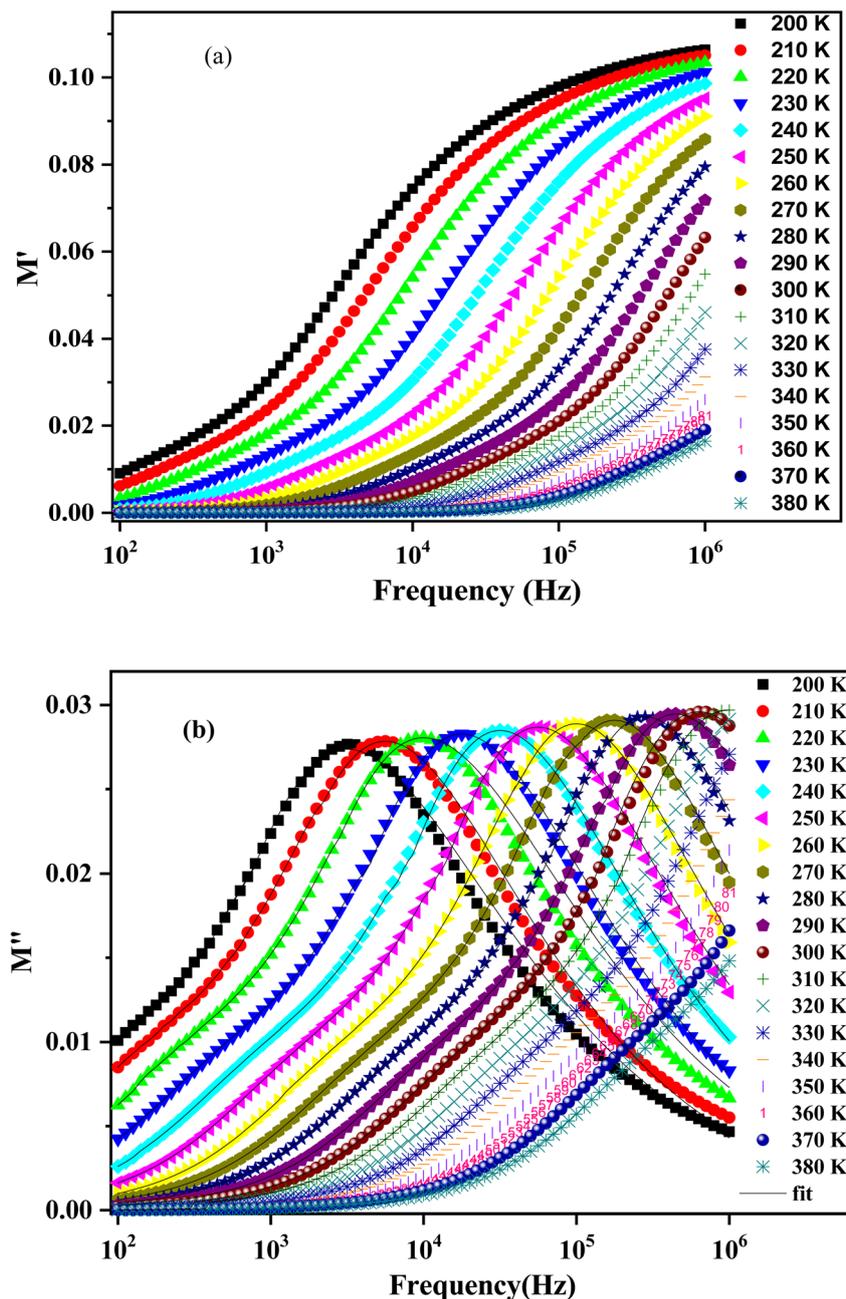


Fig. 14 The real part of the modulus plot as a function of frequency at different temperatures, (a) and the imaginary part of the modulus plot as a function of frequency at different temperatures (b), with black solid lines showing the fit of the recording data using relation (22).

temperatures increase, the peaks shift towards the higher region of frequencies, indicating the existence of a thermally activated relaxation process. Thus, we can differentiate two frequency regions according to the frequency corresponding to the modulus peak maxima positions. At the frequency region below the peak, there is the pure conduction process,⁷⁴ where the load carrier drifts loosely across long distances. However, in the second region above the peak, the load carriers are spatially restricted and can move only over short distances.⁷⁵

The experimental data of the imaginary part of the modulus plot can be interpreted by the modified function

Kohlraush–Williams–Watts (KWW). This fitting approach (KWW) is proposed by Bergman according to the following formula:⁷⁶

$$M'' = \frac{M''_{g_{\max}}}{(1 - \beta_g) + \frac{\beta_g}{1 + \beta_g} \left[\beta_g \left(\frac{f_{g_{\max}}}{f} \right) + \left(\frac{f}{f_{g_{\max}}} \right)^{\beta_g} \right]} + \frac{M''_{g_{b_{\max}}}}{(1 - \beta_{gb}) + \frac{\beta_{gb}}{1 + \beta_{gb}} \left[\beta_{gb} \left(\frac{f_{g_{b_{\max}}}}{f} \right) + \left(\frac{f}{f_{g_{b_{\max}}}} \right)^{\beta_{gb}} \right]} \quad (18)$$



Table 3 Parameters extracted from the fitted data of the imaginary part of the modulus *versus* frequency at several temperatures

T (K)	M''_{\max} (10^{-2})	β	f_{\max} (Hz)
200	2.767	0.502	3162
210	2.785	0.534	5623
220	2.806	0.501	10 000
230	2.828	0.537	17 783
240	2.849	0.550	31 623
250	2.869	0.542	56 234
260	2.889	0.560	100 000
270	2.910	0.578	177 828
280	2.930	0.570	281 838
290	2.947	0.559	446 684
300	2.960	0.530	630 957
310	2.970	0.490	972 117

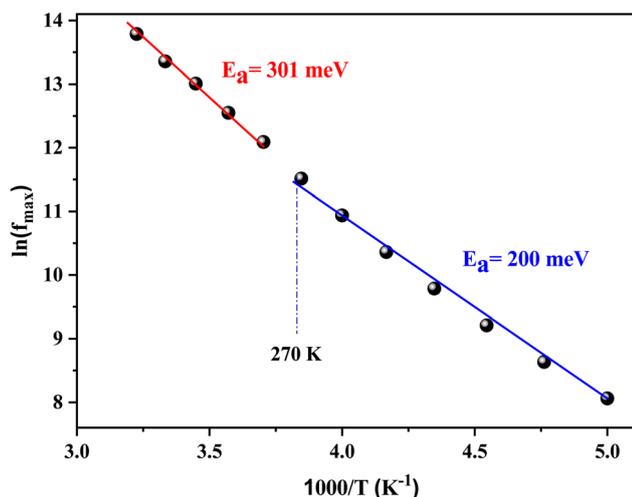


Fig. 15 The evolution of $\ln(f_{\max})$ with $1000/T$ to evaluate the activation energies for $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$.

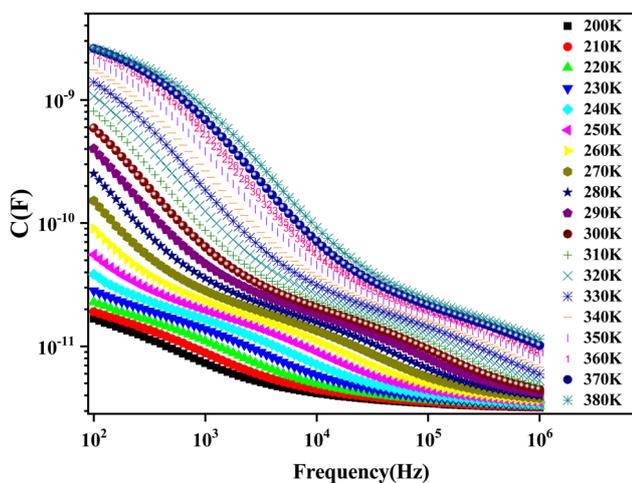


Fig. 16 The variation of capacity (C) as a function of frequency at different temperatures.

where $M''_{g_{\max}}$ and $M''_{gb_{\max}}$ are the maximum peaks value of the imaginary part of modulus match to grain and grain boundary, respectively. β_g and β_{gb} designate the deviation of the line from the linear exponential, known as the stretching factor corresponding to the grain and grain boundary, respectively. It is broadly placed in the range $0 < \beta < 1$.

The theoretical parameters deduced from the fitting of modulus data are summarized in Table 3. The value of the stretching factor does not exceed unity, and this kind of information bears out the non-Debye behavior of the examined compound. Fig. 15 illustrates the variation of $\ln(f_{\max})$ *versus* the temperature, which obeys the Arrhenius relation. The values of activation energy estimated from the slopes of lines in the plot are 288 meV and 192 meV for [200–270 K] and [280–310 K], respectively. These values are almost the same as those obtained from the imaginary part of the polarizability. This suggests that transport in the studied sample is decided by the electron-hopping mechanism.⁴²

The variation of capacitance *versus* frequency at various temperatures is shown in Fig. 16. The high value of capacitance can be obtained in the region of lower frequencies that can be related to the existence of interface states that can follow the alternating current (AC) signal.⁷⁷ The capacitance decreases with frequency. This behavior is typically similar to that of a material with traps. This can be explained by the interface states' inability to follow the applied electric field.⁶⁷

Finally, these results prove that the investigated sample could be suitable for many technological applications (Li-ion batteries, electronic devices, microwaves, fuel cells, and low-temperature cofired ceramics applications).⁴²

4 Conclusion

This study has examined the optical, magnetic, and dielectric properties of $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ prepared by the solid-state reaction. Vibrational modes were further analyzed by Raman spectra confirming the bonding of oxygen with cations in the octahedral and tetrahedral sites. The optical band gap value shows semiconductor behavior that was estimated at 3.51 eV, which was deduced from the absorption spectrum. Hence, the sample can be considered good candidates for various applications such as optoelectronics, photodetectors, photovoltaics, and gas sensors. The magnetic hysteresis loops were well investigated by the ferromagnetic (FM) and antiferromagnetic (AFM) contributions to understanding the origin of our studied sample. It was found that the ferromagnetic contribution is the most dominant. It depicts the presence of superparamagnetic behavior and a high value of microwave frequency. Therefore, the $\text{LiMn}_{0.5}\text{Fe}_2\text{O}_4$ ferrite material is a promising compound for the domain of microwave frequency bands and biomedical applications. The dielectric results represent a colossal value of dielectric permittivity, which proves the importance of ferrites as promising materials for electronic devices. The dielectric relaxations are hidden by the direct current conductivity (σ_{dc}) process and can be extracted from dielectric measurements.

This study is just a step that can be taken further. It gives interesting perspectives and makes way for new horizons to be



explored, exploited, and for practical applications of this family of ferrites compounds in multiple technological fields.

Conflicts of interest

The authors declare that there are no competing interests regarding the publication of this paper.

References

- M. D. Hossain, A. T. M. K. Jamil, M. S. Hossain, S. J. Ahmed, H. N. Das, R. Rashid and M. N. I. Khan, *RSC Adv.*, 2022, **12**, 4656.
- J. M. Gonçalves, M. N. Silva, K. K. Naik, P. R. Martins, D. P. Rocha, E. Nossol and C. S. Rout, *J. Mater. Chem.*, 2021, **9**, 3095.
- C. Yao, M. Ismail, A. Hao, S. K. Thatikonda, W. Huang, N. Qin and D. Bao, *RSC Adv.*, 2019, **9**, 12615.
- M. Chandel, D. Moitra, P. Makkar, H. Sinha, H. S. Hora and N. N. Ghosh, *RSC Adv.*, 2018, **8**, 27725.
- Z. Li, Z. L. Wang and Z. Wang, *RSC Adv.*, 2018, **8**, 23522.
- N. M. Malima, M. D. Khan, J. Choi, R. K. Gupta, P. Mashazi, T. Nyokong and N. Revaprasadu, *RSC Adv.*, 2021, **11**, 31002.
- K. Jangam, K. Patil, S. Balgude, S. Patange and P. More, *RSC Adv.*, 2020, **10**, 42766.
- M. Bastianello, S. Gross and M. T. Elm, *RSC Adv.*, 2019, **9**, 33282.
- N. Kobylinska, D. Klymchuk, A. Shakhovskiy, O. Khainakova, Y. Ratushnyak, V. Duplij and N. Matvieieva, *RSC Adv.*, 2021, **11**, 26974.
- T. N. Pham, T. Q. Huy and A. T. Le, *RSC Adv.*, 2020, **10**, 31622.
- A. Manohar, K. Chintagumpala and K. H. Kim, *Ceram. Int.*, 2021, **47**, 7052.
- K. K. Kefeni, T. A. M. Msagati, T. T. I. Nkambule and B. B. Mamba, *Mater. Sci. Eng.*, 2020, **107**, 110314.
- P. B. Kharat, M. V. Shisode, S. D. Birajdar, D. N. Bhojar and K. M. Jadhav, *AIP Conf. Proc.*, 2017, **1832**, 050122.
- S. Taneja, P. Thakur, B. Ravelo and A. Thakur, *Mater. Res. Bull.*, 2022, **154**, 111937.
- J. H. Kim, H. E. Kim, J. H. Kim and J. S. Lee, *J. Mater. Chem. A*, 2020, **8**, 9447.
- S. B. Somvanshi, S. A. Jadhav, M. V. Khedkar, P. B. Kharat, S. D. More and K. M. Jadhav, *Ceram. Int.*, 2020, **46**, 13170.
- M. Junaid, M. A. Khan, Z. M. Hashmi, G. Nasar, N. A. Kattan and A. Laref, *J. Mol. Struct.*, 2020, **1221**, 128859.
- N. Singh, A. Agarwal and S. Sanghi, *Curr. Appl. Phys.*, 2011, **11**, 783.
- F. Gandomi, S. M. Peymani-Motlagh, M. Rostami, A. Sobhani-Nasab, M. Fasihi-Ramandi, M. Eghbali-Arani and M. R. Ganjali, *J. Mater. Sci.: Mater. Electron.*, 2019, **30**, 19691.
- J. Jing, L. Liangchao and X. Feng, *J. Rare Earths*, 2007, **25**, 79.
- M. M. Hessien, *J. Magn. Magn. Mater.*, 2008, **320**, 2800.
- D. Ravinder and P. Vijaya Bhasker Reddy, *Mater. Lett.*, 2003, **57**, 4344.
- A. M. Shaikh, S. C. Watawe, S. S. Bellad, S. A. Jadhav and B. K. Chougule, *Mater. Chem. Phys.*, 2000, **65**, 46.
- T. Nakamura, M. Naoe and Y. Yamada, *J. Magn. Magn. Mater.*, 2006, **305**, 120.
- P. Hernández-Gómez, J. M. Muñoz, M. A. Valente and M. P. F. Graça, *Mater. Res. Bull.*, 2019, **112**, 432.
- M. Anis-ur-Rehman, W. Mahmood, H. Ghazanfar, M. A. A. Khan and A. Haq, *J. Supercond. Nov. Magn.*, 2019, **32**, 2787.
- E. K. Al-Shakarchi, S. H. Lafta, A. M. Musa, M. Farle and R. Salikov, *J. Supercond. Nov. Magn.*, 2016, **29**, 923.
- M. K. Abbas, M. A. Khan, F. Mushtaq, M. F. Warsi, M. Sher, I. Shakir and M. F. A. Aboud, *Ceram. Int.*, 2017, **43**, 5524.
- I. H. Gul and E. Pervaiz, *Mater. Res. Bull.*, 2012, **47**, 1353.
- I. Soudani, K. B. Brahim, A. Oueslati, H. Slimi, A. Aydi and K. Khirouni, *RSC Adv.*, 2022, **12**, 18697.
- M. A. Maksoud, A. El-Ghandour, A. H. Ashour, M. M. Atta, S. Abdelhaleem, A. H. El-Hanbaly and A. S. Awed, *J. Rare Earths*, 2021, **39**, 75.
- A. Kumar, S. K. Sharma, V. Grover, Y. Singh, V. Kumar, V. K. Shukla and P. K. Kulriya, *J. Nucl. Mater.*, 2022, **564**, 153682.
- V. D'Ippolito, G. B. Andreozzi, D. Bersani and P. P. Lottici, *J. Raman Spectrosc.*, 2015, **46**, 1255.
- G. Datt, M. S. Bishwas, M. M. Raja and A. C. Abhyankar, *Nanoscale*, 2016, **8**, 5200.
- D. K. Pradhan, S. Kumari, V. S. Puli, P. T. Das, D. K. Pradhan, A. Kumar and R. S. Katiyar, *Phys. Chem. Chem. Phys.*, 2017, **19**, 210.
- B. K. Chatterjee, K. Bhattacharjee, A. Dey, C. K. Ghosh and K. K. Chattopadhyay, *Dalton Trans.*, 2014, **43**, 7930.
- M. Virumbrales-del Olmo, A. Delgado-Cabello, A. Andrada-Chacón, J. Sánchez-Benítez, E. Urones-Garrote, V. Blanco-Gutiérrez and R. Sáez-Puche, *Phys. Chem. Chem. Phys.*, 2017, **19**, 8363.
- D. Bouokkeze, J. Massoudi, W. Hzez, M. Smari, A. Bougoffa, K. Khirouni, E. Dhahri and L. Bessais, *RSC Adv.*, 2019, **9**, 40940.
- R. D. Waldron, *Phys. Rev.*, 1955, **99**, 1727.
- I. El Heda, J. Massoudi, R. Dhahri, E. Dhahri, F. Bahri and B. F. O. Costa, *J. Alloys Compd.*, 2023, **931**, 167479.
- J. Massoudi, D. Bouekkeze, A. Bougoffa, K. Khirouni, E. Dhahri and L. Bessais, *Adv. Powder Technol.*, 2020, **31**, 4714.
- P. Kubelka and F. Munk, *Z. Phys. D*, 1931, **12**, 259.
- A. Ghoudi, K. B. Brahim, A. Oueslati, K. Khirouni and A. Aydi, *J. Mater. Sci.: Mater. Electron.*, 2022, **33**, 1.
- A. Hadded, J. Massoudi, E. Dhahri, K. Khirouni and B. F. O. Costa, *RSC Adv.*, 2020, **10**, 42542.
- R. Mguedla, A. B. J. Kharrat, O. Taktak, H. Souissi, S. Kammoun, K. Khirouni and W. Boujelben, *Opt. Mater.*, 2020, **101**, 109742.
- P. Chand, S. Vaish and P. Kumar, *Phys. B*, 2017, **524**, 53.
- N. Amri, J. Massoudi, K. Nouri, M. Triki, E. Dhahri and L. Bessais, *RSC Adv.*, 2021, **11**, 13256.
- Q. Chen and Z. J. Zhang, *Appl. Phys. Lett.*, 1998, **73**, 3156.
- L. R. Shah, H. Zhu, W. G. Wang, B. Ali, T. Zhu, X. Fan and J. Q. Xiao, *J. Phys. D*, 2010, **43**, 035002.



- 50 S. E. Shirsath, S. S. Jadhav, B. G. Toksha, S. M. Patange and K. M. Jadhav, *J. Appl. Phys.*, 2011, **110**, 013914.
- 51 M. N. Akhtar, M. Saleem and M. A. Khan, *J. Phys. Chem. Solids*, 2018, **123**, 260.
- 52 M. Akhtar, M. Babar, S. Qamar, Z. Rehman and M. A. Khan, *Ceram. Int.*, 2019, **45**, 10187.
- 53 G. Sahu, M. Das, M. Yadav, B. P. Sahoo and J. Tripathy, *Polymers*, 2020, **12**, 374.
- 54 M. M. L. Sonia, S. Anand, V. M. Vinose, M. A. Janifer, S. Pauline and A. Manikandan, *J. Magn. Magn. Mater.*, 2018, **466**, 238.
- 55 M. H. Dhaou, S. Hcini, A. Mallah, M. L. Bouazizi and A. Jemni, *Appl. Phys. A*, 2016, **123**, 1.
- 56 M. Saha, S. Mukherjee, P. Bera, M. M. Seikh and A. Gayen, *Ceram. Int.*, 2022, **48**, 35719.
- 57 A. Thakur, P. Mathur and M. Singh, *J. Phys. Chem. Solids*, 2007, **68**, 378.
- 58 K. Pubby, K. Vijay Babu and S. Bindra Narang, *Mater. Sci. Eng. B*, 2020, **255**, 114513.
- 59 A. M. Shaikh, S. S. Bellad and B. K. Chougule, *J. Magn. Magn. Mater.*, 1999, **195**, 384.
- 60 M. C. Dimri, A. Verma, S. C. Kashyap, D. C. Dube, O. P. Thakur and C. Prakash, *Mater. Sci. Eng. B*, 2006, **133**, 42.
- 61 Y. B. Taher, A. Oueslati, K. Khirouni and M. Gargouri, *J. Clust. Sci.*, 2015, **5**, 1655.
- 62 I. S. Yahia, N. A. Hegab, A. M. Shakra and A. M. AL-Ribaty, *Phys. B*, 2012, **13**, 2476.
- 63 N. J. Tharayil, S. Sagar, R. Raveendran and A. V. Vaidyan, *Phys. B*, 2007, **399**, 1.
- 64 J. C. Giuntini, J. V. Zanchetta, D. Jullien, R. Eholie and P. Houenou, *J. Non-Cryst. Solids*, 1981, **45**, 57.
- 65 K. M. Batoo, S. Kumar, C. G. Lee and A. Alimuddin, *Curr. Appl. Phys.*, 2009, **9**, 1397.
- 66 A. Ben Jazia Kharrat, M. Bourouina, N. Moutia, K. Khirouni and W. Boujelben, *J. Alloys Compd.*, 2018, **741**, 723.
- 67 R. Mguedla, A. Ben Jazia Kharrat, N. Moutia, K. Khirouni, N. Chniba-Boudjada and W. Boujelben, *J. Alloys Compd.*, 2020, **836**, 155186.
- 68 B. K. P. Scaife, *Proc. Phys. Soc.*, 1963, **81**, 124.
- 69 M. Amghar, A. Bougoffa, A. Trabelsi, A. Oueslati and E. Dhahri, *RSC Adv.*, 2022, **12**, 20348.
- 70 S. Saha and T. P. Sinha, *Phys. Rev.*, 2002, **65**, 134103.
- 71 P. Norouzzadeh, K. Mabhouti, M. M. Golzan and R. Naderali, *J. Mater. Sci.*, 2020, **31**, 7335.
- 72 N. K. Mohanty, S. K. Satpathy, B. Behera, P. Nayak and R. N. P. Choudhary, *J. Adv. Ceram.*, 2012, **1**, 221.
- 73 P. Oruc, P. Turan, Y. Demiroglu, A. Seckin, S. Cavdar, H. Korlav and N. Tugluoglu, *J. Mater. Sci.*, 2021, **32**, 15837.
- 74 C. Bharti and T. P. Sinha, *Solid State Sci.*, 2010, **4**, 498.
- 75 K. Holderna-Natkaniec, M. Ouled Mohamed Sghaier, P. Ławniczak, M. Zdanowska-Frączek, A. Wozniak-Braszak and S. Chaabouni, *Polyhedron*, 2015, **85**, 131.
- 76 R. Bergman, *J. Appl. Phys.*, 2000, **88**, 1356.
- 77 D. M. Taylor and H. L. Gomes, *J. Phys. D: Appl. Phys.*, 1995, **28**, 2554.

