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Novel HPMC/PEDOT:PSS nanocomposite for optoelectronic and energy storage applications

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This study investigates a class of materials known as polymer nanodielectrics, which are formed by incorporating ceramic fillers into polymers. These materials offer the unique advantage of tunable electrical and optical properties. The research focuses on the incorporation of high-purity stannic oxide nanoparticles (SnO₂ NPs) into a ternary blend matrix of hydroxypropyl methylcellulose (HPMC) and poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT:PSS) using a solution casting method. Characterization techniques like X-ray diffraction (XRD) and Fourier-transform infrared spectroscopy (FT-IR) revealed alterations in the amorphous nature of the HPMC/PEDOT:PSS blend upon the introduction of SnO₂ NPs. These analyses also suggest the formation of interactions between the polymer and nanoparticles. Scanning electron microscopy (SEM) images confirmed the successful dispersion of SnO₂ NPs on the surface of the polymer blend, particularly at lower concentrations. The optical properties of the nanocomposite films were investigated using UV-vis spectrophotometry. This analysis allowed for the calculation of optical constants like the bandgap and refractive index. The results showed a dual-bandgap structure, with the direct and indirect bandgaps ranging from 4.92 eV to 4.26 eV and 3.52 eV to 1.68 eV, respectively. Electrical characterization using AC conductivity and dielectric permittivity measurements revealed a dependence on the SnO₂ NPs concentration within the frequency range of 0.1 Hz to 10 MHz. The relaxation processes and interfacial polarization effects within these nanocomposites are further discussed in the study. At a frequency of 10 Hz, the AC conductivity exhibited a significant increase, rising from $1.85 \times 10^{-12} \text{ S m}^{-1}$ to $1.04 \times 10^{-9} \text{ S m}^{-1}$ upon the addition of 0.7 wt% SnO₂ NPs. These findings highlight the multifunctional nature of the developed nanocomposites. They hold promise for various applications, including UV blockers, optical bandgap tuners, and optical coatings in advanced optoelectronic devices. Additionally, their tunable high permittivity suggests potential use as dielectric substrates for next-generation, high-performance energy storage devices.

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1. Introduction

In the past few decades, there has been notable progress in technology, accompanied by a rise in digital products and services. Moreover, emerging technologies such as artificial intelligence (AI), robotics, and nanotechnology appear poised to accelerate the pace of advancement.¹ Nanotechnology's versatility offers a wealth of opportunities to improve our lives. Notably, materials science, nanomedicine, and device engineering hold immense promise for groundbreaking advancements. At the nanoscale, materials exhibit properties distinct from their bulk counterparts. These properties can encompass significant changes in mechanical strength, electrical conductivity, magnetism, thermal behavior, and light interaction. This

has attracted significant research interest, with transition metal oxides like stannic oxide nanoparticles (SnO₂ NPs) gaining particular attention for their desirable properties. SnO₂ NPs offer several advantages: they are low-cost, highly stable, and possess valuable optoelectronic characteristics.²

One promising near-term strategy to leverage the benefits of SnO₂ NPs is through the creation of composites with polymers or polymer blends. Hydroxypropyl methylcellulose (HPMC) and PEDOT:PSS are particularly attractive candidates because of optical, and electrical properties, making them highly relevant for industrial applications. HPMC, a cellulose derivative, is a water-soluble anionic polymer known for its film and gel-forming capabilities at high temperatures.³ The polymer's structure incorporates OH group and COO[−] functionalities, promoting interaction between the nanofillers and enhancing the overall performance of the composite system.⁴ HPMC's versatility stems from its unique combination of properties. It is transparent, possesses high viscosity, and exhibits

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biocompatibility and biodegradability. These characteristics make HPMC valuable across diverse applications, ranging from paper manufacturing and textile printing to controlled drug delivery in medications.⁵

PEDOT:PSS is a remarkable material with a unique combination of properties that make it highly sought after in various fields. It boasts impressive conductivity, allowing it to efficiently carry electrical current. This characteristic makes it a perfect candidate for applications like transparent electrodes in solar cells and OLEDs.⁶ Furthermore, PEDOT:PSS demonstrates good biocompatibility, minimizing negative interactions with living tissues. This biocompatible nature opens doors for exciting biomedical applications such as neural interfaces, implantable biosensors, and tissue engineering scaffolds. Additionally, PEDOT:PSS offers partial transparency, permitting some light to pass through. This transparency is crucial for the development of organic solar cells and transparent displays.⁷ Finally, PEDOT:PSS exhibits good thermal stability, meaning it can withstand relatively high temperatures without significant degradation. This property makes it suitable for processing and device fabrication at elevated temperatures. While PEDOT:PSS offers significant advantages, research continues to refine its properties and explore even more applications for this versatile material.

Recent studies have utilized the solution casting method to create lightweight and flexible polymer nanocomposites containing HPMC, PEDOT:PSS, and SnO₂ NPs, aiming to enhance their electrical properties for electronic applications. The research focuses on using an HPMC/PEDOT:PSS blend as the main polymeric matrix, incorporating SnO₂ NPs as conductive additives. This work delves into the functional properties of the nanocomposites through a battery of characterization techniques, including structural, optical, and electrical analyses. By unveiling the interplay between these properties at the nano-scale, the investigation aims to shed light on the underlying mechanisms responsible for the improved performance of these materials.

2. Experimental work

2.1. Chemicals

Stannic oxide nanoparticles (SnO₂ NPs) with a particle size less than 40 nm and a purity of 99.7% trace metal basis were acquired from Sigma-Aldrich. Hydroxypropyl methylcellulose (HPMC) with a molecular weight of about 133 400 g mol⁻¹ and poly(3,4-ethylenedioxythiophene) polystyrene sulfonate (PEDOT:PSS) as a 1.3 wt% dispersion in water with a conductivity of 1 S cm⁻¹ were also obtained from the same supplier.

2.2. Preparation of HPMC/PEDOT:PSS-SnO₂ nanocomposite

Nanocomposite films were prepared using a solution casting method. First, 0.25 g of HPMC was dissolved in 80 ml of deionized water at 50 °C with the aid of magnetic stirring to ensure complete dissolution. After allowing the HPMC solution to cool, 20 ml of the mixture was combined with 0.5 ml of PEDOT:PSS. This combined solution was then equally divided

into four parts and stirred for an additional 2 hours. In parallel, varying amounts of SnO₂ NPs (0, 0.003, 0.005, and 0.007 g) were each dispersed in 2 ml of deionized water. Each SnO₂ NP dispersion was then incorporated into one of the four HPMC/PEDOT:PSS mixtures and stirred for another 2 hours. Finally, the resulting mixtures were cast into plastic Petri dishes with a diameter of 55 mm and dried at 45 °C. Following the casting process, the films underwent thorough drying to eliminate residual moisture. This was achieved by placing them in a desiccator containing silica gel desiccants. To facilitate future reference, the dried films were labeled S1, S2, S3, and S4, with the labels corresponding to the increasing amount of incorporated SnO₂ NPs (0 g for S1 and progressively more for subsequent samples).

2.3. Characterizations

The structural and morphological properties of the nanocomposite films were characterized using various techniques. Fourier transform infrared (FT-IR) spectroscopy (JASCO 430, Japan) identified functional groups within the 400–4000 cm⁻¹ wavenumber range. XRD using CuK_α radiation (DIANO, USA) revealed the crystalline structure of the samples by analyzing the Bragg angle (2θ) between 5° and 70°. Optical properties were investigated using a UV-visible spectrophotometer (JASCO V-630, Japan) to measure light absorption across the spectrum. Scanning electron microscopy (SEM) with a 20 kV electron beam (Quanta Inspect F50, FEI Company, Netherlands) was employed to visualize the surface morphology. To characterize the electrical properties of the films at room temperature (303 K), broadband dielectric spectroscopy was employed.

3. Result and discussion

3.1. XRD studies

Fig. 1 presents the X-ray diffraction pattern of the SnO₂ nanoparticles. The detected diffraction peaks were associated with the standard JCPDS data (# 41-1445) to identify the corresponding crystalline planes. This analysis confirms the formation of a pure, polycrystalline SnO₂ phase with a tetragonal crystal system. The results are consistent with previous reports on SnO₂ nanoparticles.⁸ The XRD patterns of the HPMC/PEDOT:PSS nanocomposites with varying SnO₂ NPs concentrations are presented in Fig. 2. The pattern for pristine PEDOT:PSS exhibits broad, low-intensity diffraction peaks around 2θ = 12.27°. These peaks can be attributed to either the intermolecular spacing of the polymer backbone or the (020) reflection in the crystal structure.⁹ The XRD patterns of the nanocomposites exhibit several new diffraction peaks compared to pristine HPMC/PEDOT:PSS (Fig. 2). These peaks appear at 2θ values of approximately 26.6°, 33.9°, and so on, which is consistent with the reported diffraction pattern of tetragonal SnO₂. This observation strongly suggests the successful incorporation of SnO₂ nanoparticles into the HPMC/PEDOT:PSS matrix. While the XRD patterns of the nanocomposites confirmed the presence of SnO₂ (Fig. 1), the peaks appeared broader and less intense compared to pure SnO₂. This



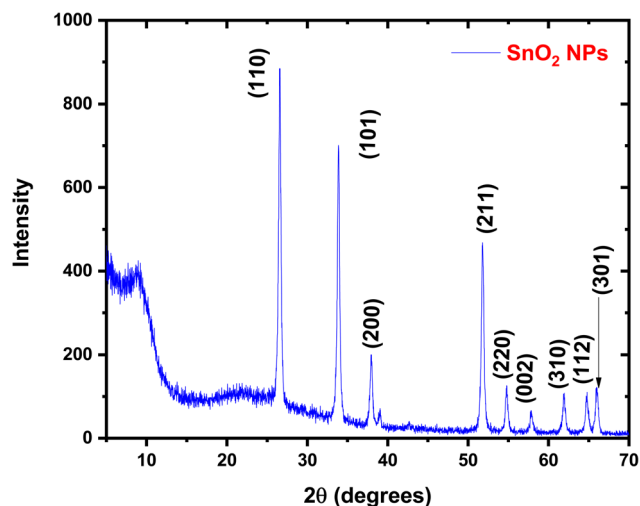


Fig. 1 XRD pattern of pure SnO₂ nanoparticles.

suggests a less well-crystallized SnO₂ phase within the composite matrix. The Debye–Scherrer equation,¹⁰ was employed to estimate the average crystallite size (*D*) of SnO₂ nanoparticles using the most prominent peak (likely corresponding to the (110) plane). The calculated crystallite sizes ranged from approximately 43 nm to 26 nm across the different nanocomposite films.

$$D = \frac{0.9\lambda}{\beta \cos \theta} \quad (1)$$

The Debye–Scherrer equation, a well-established technique for estimating crystallite size from XRD data, was employed in this study. This method considers factors like the peak broadening (full width at half maximum, FWHM) and the Bragg angle (*θ*) to calculate the average crystallite size (*D*). Notably, the

analysis revealed a decrease in the average crystallite size of SnO₂ nanoparticles from 43.12 nm to 25.81 nm as the concentration of SnO₂ increased within the HPMC/PEDOT:PSS nanocomposites.

3.2. FT-IR analysis

Fourier Transform Infrared (FT-IR) spectroscopy was employed to investigate the formation of the HPMC/PEDOT:PSS blend and the incorporation of SnO₂ nanoparticles within the composite (Fig. 3). The FT-IR spectra of the HPMC/PEDOT:PSS blend doped with varying SnO₂ concentrations reveal distinct peaks corresponding to different functional groups. The pristine HPMC/PEDOT:PSS spectrum exhibits characteristic transmission bands at 1222 cm⁻¹, 1116 cm⁻¹, and 1090 cm⁻¹, indicative of CH₂ bending, CH₂ symmetrical stretching, and CO stretching vibrations, respectively.^{1,3,5,10,13,14} The FT-IR spectra (Fig. 3) further confirm the presence of HPMC and PEDOT:PSS within the composites. Characteristic peaks for HPMC were observed at 1403 cm⁻¹ (CH₂ scissoring),¹¹ while PEDOT:PSS displayed bands at 1655 cm⁻¹ (C=C stretching) and 3471 cm⁻¹ (OH stretching).¹ A more detailed peak assignment is provided in Table 1. Interestingly, the incorporation of SnO₂ NPs resulted in subtle changes to the overall FT-IR profile. Notably, a new peak emerged at 468 cm⁻¹, attributed to Sn–O vibrational modes, and its intensity increased with higher SnO₂ concentration. This observation suggests successful interaction between the HPMC/PEDOT:PSS blend and the SnO₂ nanoparticles. Further analysis of these spectral shifts, particularly for characteristic bands of the polymer components, can provide valuable insights into how the SnO₂ filler influences the structural composition of the interpolymer matrix. It's important to note that the presence of double bond segments within the composite structure might be advantageous. These segments could potentially serve as preferential sites for

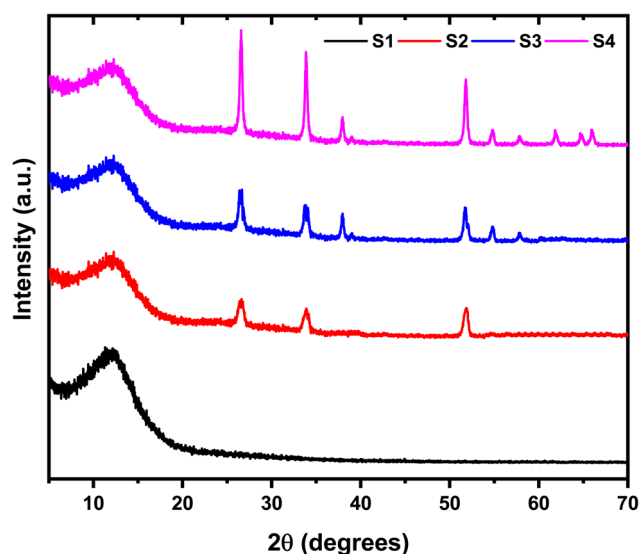


Fig. 2 XRD pattern of HPMC/PEDOT:PSS ternary blend and HPMC/PEDOT:PSS doped with SnO₂ nanoparticles.

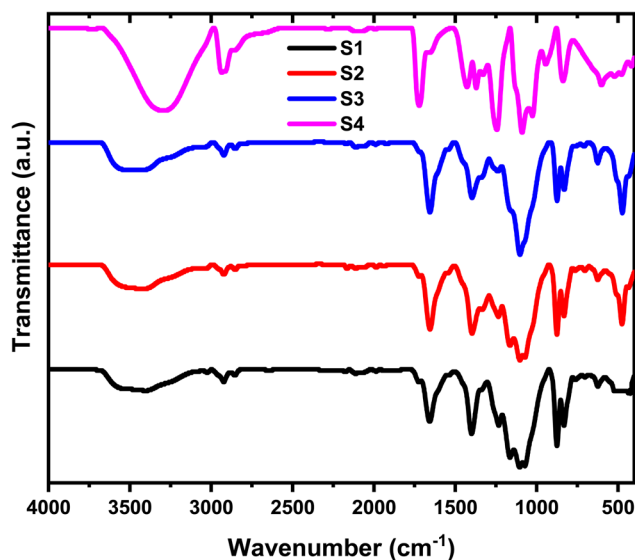
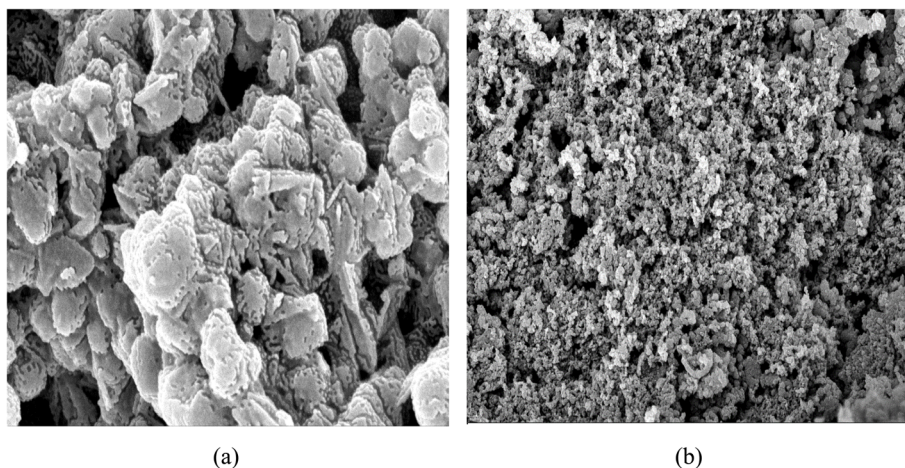


Fig. 3 FT-IR spectra of HPMC/PEDOT:PSS ternary blend doped with SnO₂ nanoparticles.



Table 1 FT-IR Peak assignments for the synthesized nanocomposites

Wavenumber (cm ⁻¹)	Bands assignment	Sources	References
3471	OH stretching	HPMC + PEDOT:PSS	1, 5 and 13
2926	CH ₂ asymmetric stretching	HPMC	3 and 14
2850	CH ₂ symmetrical stretching	HPMC	3
1655	C=C stretching	PEDOT:PSS	1
1402	CH ₂ scissoring	HPMC	5 and 14
1222	CH ₂ bending	HPMC	3 and 5
1116	CH ₂ bending	HPMC	3 and 5
1090	CO stretching	HPMC	5
872	CH ₂ bending	HPMC	5
468	Sn–O vibration	SnO ₂ NPs	10

Fig. 4 FESEM images (a) HPMC/PEDOT:PSS, (b) HPMC/PEDOT:PSS/0.007 g SnO₂ nanocomposite.

polarons and bipolarons, which can act as charge carriers and contribute to the electrical properties of the nanocomposite.¹²

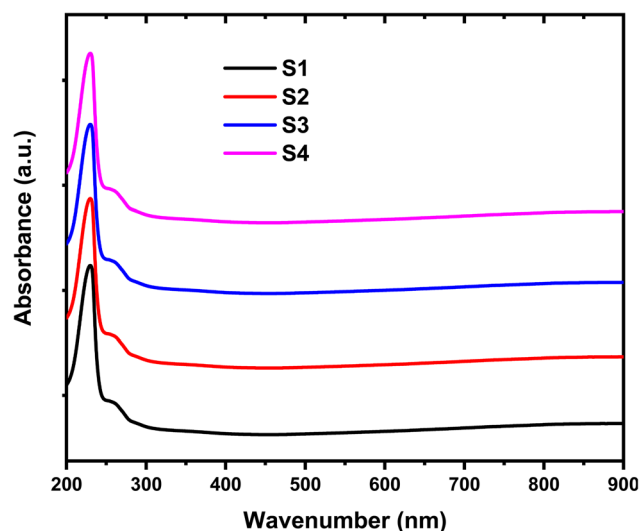
3.3. Morphology studies

Field emission scanning electron microscopy (FESEM) was employed to investigate the surface morphology of the HPMC/PEDOT:PSS composites with varying SnO₂ nanoparticle concentrations (Fig. 4). The FESEM image of pristine HPMC/PEDOT:PSS (Fig. 4a) reveals a smooth surface with minimal porosity, suggesting a relatively low surface area. In contrast, the FESEM image of the composite containing the highest SnO₂ loading (0.007 g) exhibits a markedly different morphology (Fig. 4b). This image shows a highly porous structure with the presence of well-dispersed SnO₂ nanoparticles. This observation suggests that SnO₂ nanoparticle incorporation effectively disrupts the formation of large HPMC/PEDOT:PSS agglomerates, leading to an increased surface area. Furthermore, the introduction of SnO₂ nanoparticles has been shown to enhance the electrical conductivity of the HPMC/PEDOT:PSS composites.

3.4. UV-visible spectroscopy

Fig. 5 illustrates the relationship between optical absorbance and wavelength (λ) for the HPMC/PEDOT:PSS composite and the HPMC/PEDOT:PSS-SnO₂ nanocomposites. The data reveals

that the absorbance of the nanocomposites is generally higher than that of the pristine HPMC/PEDOT:PSS blend across most of the investigated wavelength range. Furthermore, the

Fig. 5 UV-Vis absorption spectra of the HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS nanocomposites containing different SnO₂ nanoparticle loadings.

absorbance tends to increase with increasing SnO₂ concentration.¹⁵ The observed variations in optical properties suggest the formation of homogeneous polymer nanocomposites. This characteristic, combined with the increased light absorption due to SnO₂ incorporation, makes these nanocomposites promising candidates for applications in optical coatings and UV shielding materials.¹⁶ The UV-Vis spectrum reveals an absorption band around 230 nm. Interestingly, the incorporation of SnO₂ nanoparticles leads to a red-shift of this peak and the overall absorption edge in the UV region. This observation suggests a decrease in the optical band gap (E_g) of the nanocomposite compared to pristine HPMC/PEDOT:PSS. The red-shifted absorption edge might be attributed to the creation of new localized states and energy levels within the band gap. These new states, potentially induced by increased disorder and defects due to SnO₂ incorporation, could facilitate the movement of electrons from the valence band (VB) to the conduction band (CB). This ultimately leads to a narrower band gap and potentially allows for more efficient electronic transitions within the nanocomposite.¹⁷ The incorporation of SnO₂ nanoparticles within the HPMC/PEDOT:PSS matrix likely influences

the electronic interactions and creates more defect states within the composite.¹⁸ These defect states and potential disorder regions may enhance the mobility of polymer segments and facilitate the movement of electrons between VB and CB of the nanocomposite. This can lead to a reduction in the overall band gap (E_g) and a consequent red-shift of the absorption edge observed in the UV-Vis spectrum.

The Tauc equation is employed to estimate the optical band gap (E_g) of the prepared films for both allowed indirect (E_{ig}) and direct (E_{dg}) transitions:¹⁹

$$(\alpha h\nu)^2 = B(h\nu - E_{dg}) \quad (2)$$

$$(\alpha h\nu)^{1/2} = B(h\nu - E_{dg}) \quad (3)$$

This approach relates the absorption coefficient (α) to the incident photon energy ($h\nu$) through a material-specific constant (B). Fig. 6 and 7 figures depict plots of $(\alpha h\nu)^2$ and $(\alpha h\nu)^{1/2}$ for indirect and direct transitions, respectively, as functions of $h\nu$. The band gap values (E_{dg} and E_{ig}) can be determined by extrapolating the linear portions of these curves

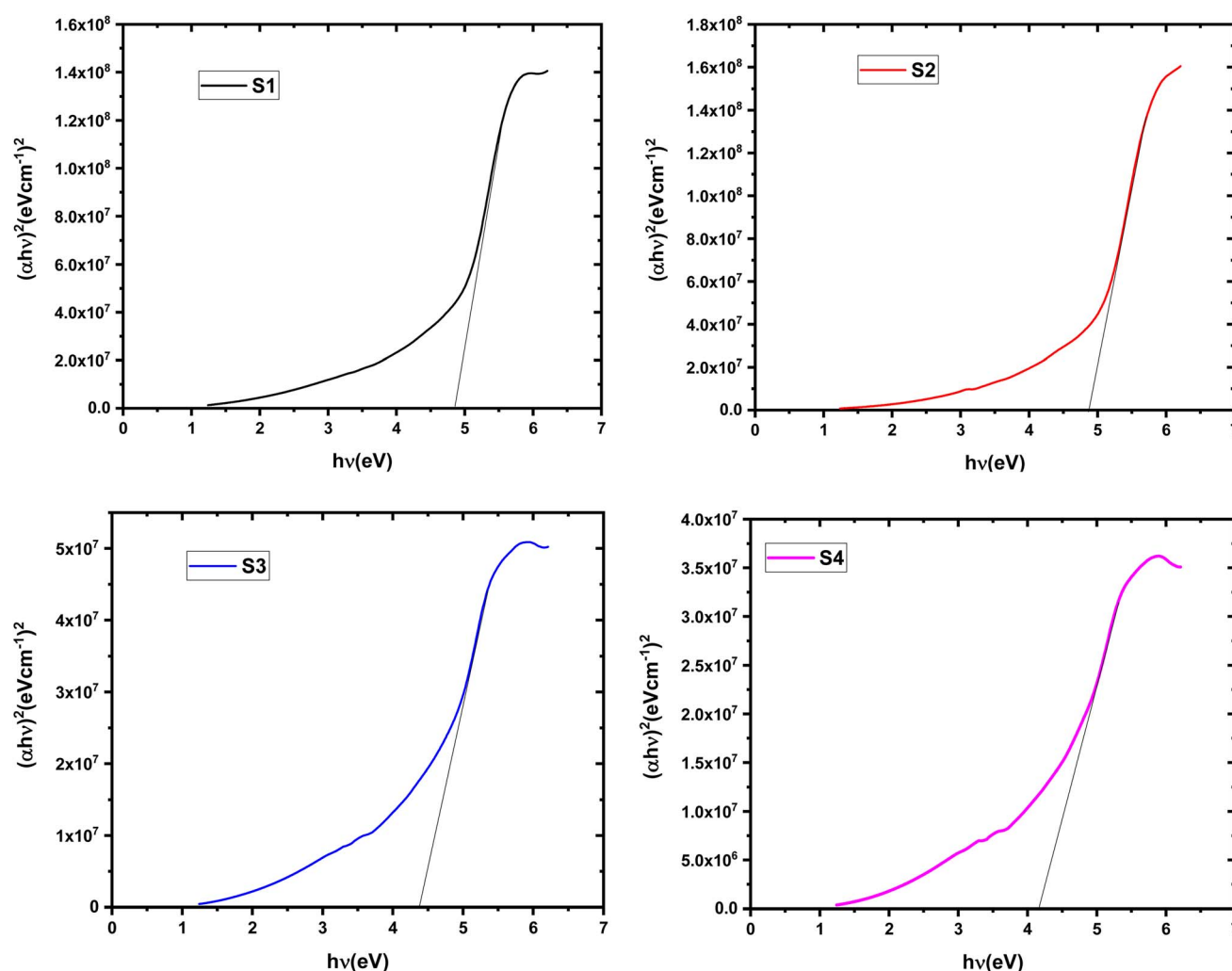


Fig. 6 Tauc plot for determining the direct band gap (E_{dg}) of HPMC/PEDOT:PSS composites with varying SnO₂ nanoparticle loadings.



to the point where the absorption coefficient (α) intercepts zero. The obtained E_g values for the nanocomposite films are presented in Table 2. The E_g value of the pristine HPMC/PEDOT:PSS composite is consistent with previously reported values.²⁰ Notably, the incorporation of SnO₂ nanoparticles leads to a decrease in the E_g compared to the pure polymer composite. This observation aligns with the potential formation of charge transfer complexes and defect states within the band gap. These defect states, possibly arising from increased disorder due to SnO₂ inclusion, could facilitate electron movement between VB and CB. Additionally, the lower E_g of

SnO₂ nanoparticles compared to HPMC/PEDOT:PSS might contribute to the overall reduction in the nanocomposite's band gap.⁵ The observed decrease in E_g suggests that the nanocomposite films exhibit enhanced semiconducting behavior upon SnO₂ nanoparticle incorporation. This could potentially translate to the formation of more efficient 3D conducting networks within the material.¹⁵

The linear refractive index (n_0) of the nanocomposite films was estimated using an empirical relationship established by Dimitrov and Sakka,²¹ and based on the indirectly derived band gap values (E_{gi}) (Table 2).

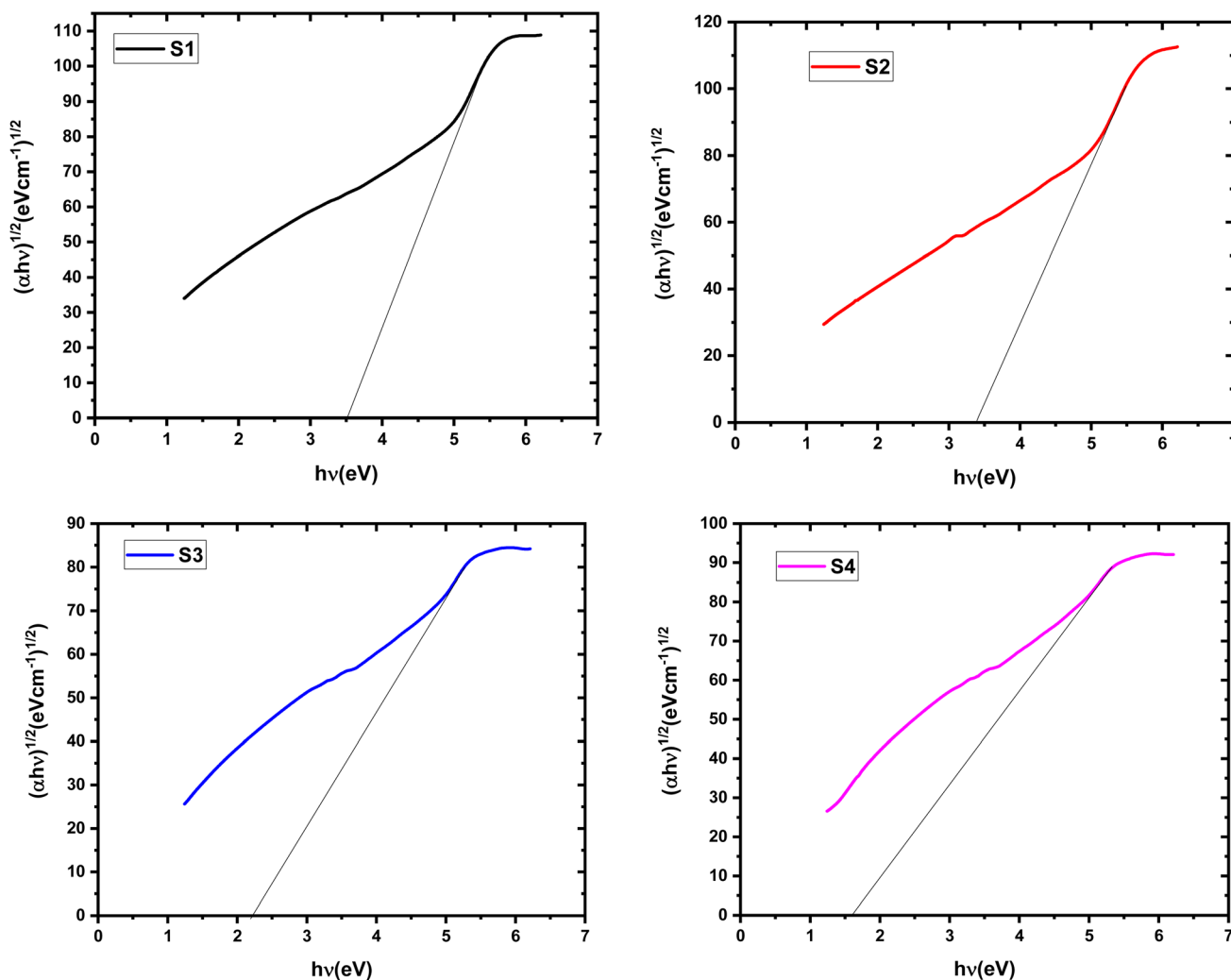


Fig. 7 Tauc plot for determining the indirect band gap (E_{ig}) of HPMC/PEDOT:PSS composites with varying SnO₂ nanoparticle loadings.

Table 2 Energy gap, refractive index, and AC conductivity for the prepared samples

Samples	Description	E_{dg} (eV)	E_{ig} (eV)	n_0	AC conductivity ($S\ cm^{-1}$) at 10 Hz	S
S1	HPMC/PEDOT:PSS	4.92	3.52	2.07	1.85×10^{-12}	0.85
S2	HPMC/PEDOT:PSS-0.003 SnO ₂ NPs	4.85	3.43	2.81	1.32×10^{-11}	0.78
S3	HPMC/PEDOT:PSS-0.005 SnO ₂ NPs	4.41	2.25	3.25	7.46×10^{-11}	0.66
S4	HPMC/PEDOT:PSS-0.007 SnO ₂ NPs	4.26	1.68	3.54	1.04×10^{-9}	0.52

$$\frac{n_0^2 - 1}{n_0^2 + 1} = 1 - \sqrt{\frac{E_{gi}}{20}} \quad (4)$$

This equation suggests an inverse proportionality between n_0 and E_{gi} , which is consistent with the observed trends. Our data reveals an improvement in n_0 upon incorporation of SnO₂ nanoparticles. The HPMC/PEDOT:PSS/0.007 g SnO₂ NPs sample exhibits the highest n_0 value. These variations in the optical properties can be attributed to the internal structure of the material, including factors like molecular weight distribution and packing density. The introduction of SnO₂ nanoparticles as a nanofiller might have increased the degree of disorder within the composite. This, in turn, could potentially enhance the mobility of free radicals within the material. As free radicals are inherently dynamic, their increased movement might lead to a rise in excited states, ultimately contributing to the observed improvement in n_0 for the doped samples.²² The enhanced n_0 values make these nanocomposite films promising candidates for applications requiring high refractive index materials, such as high-performance lenses or antireflection coatings.²³

3.5. Electrical conductivity

The frequency dependence of electrical conductivity is a well-known characteristic of amorphous polymer composites. This phenomenon arises from the interplay between the polymeric matrix and the charge carriers within it. The mobility of these charge carriers can be influenced by the frequency of the applied electric field. To assess the AC conductivity of the samples, a sinusoidal voltage with varying frequencies (f) is employed. The response of the material to this alternating voltage is measured by monitoring the current that flows through it. This current measurement includes both its amplitude and the phase shift relative to the applied voltage. By analyzing this current response, the AC conductivity ($\sigma(f)$) can be calculated using the following equation:²⁴

$$\sigma_{ac} = 2\pi f \epsilon' \epsilon_0 \tan \delta \quad (5)$$

where, f represents the frequency, ϵ' denotes the measured dielectric constant, and ϵ_0 is the permittivity of free space. Fig. 8 depicts the frequency dependence of σ for the HPMC/PEDOT:PSS composites containing varying weight percentages of SnO₂ nanoparticles, measured at room temperature. The data reveals a clear trend of increasing conductivity with increasing SnO₂ content. This observation suggests that the well-dispersed SnO₂ nanoparticles contribute to the formation of additional conductive pathways within the composite. The overall conductivity of these nanocomposites likely depends on several factors, including the type, morphology (shape and size), and spatial distribution of the nanofillers within the polymer matrix.²⁵ The observed conductivity enhancement might be attributed to a combination of two mechanisms. First, the SnO₂ nanoparticles themselves may introduce additional charge carriers into the system. Second, the incorporation of SnO₂ could potentially increase the amorphous character of the polymer matrix. This, in turn, could facilitate the movement of

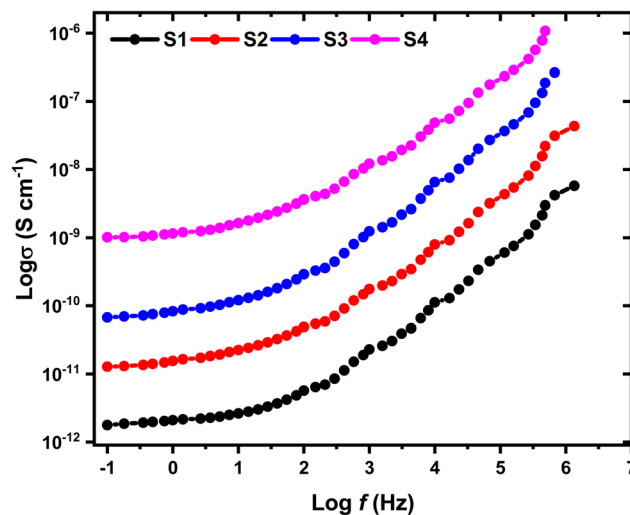


Fig. 8 The relationship between $\log \sigma$ and $\log f$ for both HPMC/PEDOT:PSS composite films and HPMC/PEDOT:PSS films incorporating SnO₂ nanoparticles.

ions by lowering the energy barriers associated with their transport.²⁶ Fig. 8 illustrates the dependence of electrical conductivity on frequency for the prepared HPMC/PEDOT:PSS nanocomposite films. The data reveals a clear trend of increasing conductivity with increasing frequency for all samples. This observation suggests a frequency-dependent behavior, particularly at higher frequencies. The observed behavior aligns with the power-law relationship proposed by Jonscher for electrical conductivity in many materials:²⁷

$$\sigma_{ac}(\omega) = \sigma_{dc} + A\omega^s \quad (6)$$

In this context, σ_{dc} represents the direct current conductivity (specifically at $\omega = 0$), A denotes a frequency-dependent coefficient, and s varies with frequency. The findings presented in Table 2 align with the correlated barrier hopping (CBH) model for electrical conduction in the films. This interpretation stems from the consistent observation that all “ s ” values are below 1.^{28,29} Additionally, the σ values for HPMC/PEDOT:PSS-SnO₂ films show an increase of approximately four orders of magnitude compared to pure HPMC/PEDOT:PSS. With higher frequencies of the applied field, the conductivity of charge carriers demonstrates a linear escalation, indicating enhanced mobility at high frequencies compared to low frequencies.³⁰ Moreover, increased dominance of the loss factor at higher frequencies contributes to a proportional enhancement in electrical conductivity within this frequency range. Similar trends have been noted in diverse polymer nanocomposites.^{31,32}

3.6. Dielectric properties

The dielectric constant (ϵ') and dielectric loss (ϵ'') of a material are crucial parameters that reflect its ability to store and dissipate electrical charge. Materials with high dielectric constants can store more charge for a given electric field, making them valuable for capacitors where smaller dimensions are desired.



Conversely, materials with low dielectric constants exhibit minimal energy loss at high frequencies or under high power conditions, minimizing power loss in these applications.³³ These contrasting properties highlight the importance of understanding dielectric behavior when selecting materials for specific functionalities. The following equation outlines the calculation for both dielectric constant and loss:³⁴

$$\varepsilon' = \frac{Cd}{\varepsilon_0 A} \quad (7)$$

$$\varepsilon'' = \frac{\sigma}{\omega \varepsilon_0} \quad (8)$$

In the investigation of HPMC/PEDOT nanocomposites, Fig. 9 and 10 illustrate the variation of ε' and ε'' with frequency at room temperature, where A and d denote the sample's area and thickness, and C represents the sample capacitance. The findings indicate distinct behaviors based on the concentration of SnO₂ NPs relative to their percolation threshold. Samples containing SnO₂ NPs above the percolation threshold show a significant decrease in both ε' and ε'' as frequency increases, ultimately stabilizing at lower values. In contrast, samples with SnO₂ NP concentrations below this threshold exhibit relatively constant ε' and ε'' values. This trend suggests that electrode and interface effects, predominant in significant parts of the sample, contribute to the observed high dielectric constants at low frequencies. Additionally, for polar materials, initial dielectric values tend to be high but diminish as the frequency of the electric field increases due to limitations in dipole alignment with field fluctuations.³⁵ Moreover, the incorporation of SnO₂ NPs enhances ε' and ε'' values across all frequency ranges due to the introduction of polar groups. SnO₂ NPs act as nanocapacitors within the insulating matrix, and as their concentration in the composites increases, these nanocapacitors become more prominent, thereby augmenting the overall capacitance (*i.e.*, dielectric constant).³⁶ Furthermore,

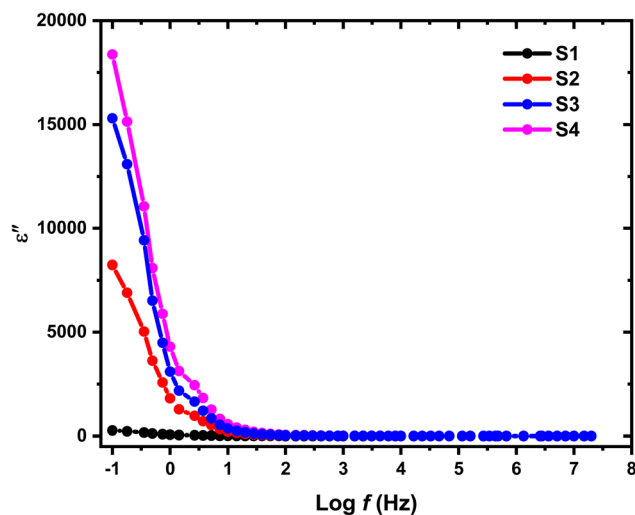


Fig. 10 Variation in dielectric loss (ε'') with $\log(f)$ for HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS loaded with SnO₂ NPs.

nanofillers are known to alleviate restrictions on dipole responsiveness to applied electric fields, thereby further enhancing the dielectric constant.³⁷

3.7. Complex electric modulus

The study of dielectric spectra often involves the use of advanced mathematical functions like complex electric modulus (M^*) and complex impedance (Z^*).³⁸ These functions provide crucial insights into how materials behave electrically at different frequencies. Complex impedance (Z^*) is particularly effective for studying systems dominated by long-range charge transport, typical of resistive or capacitive materials. On the other hand, complex electric modulus (M^*) is preferred when the material primarily exhibits localized relaxation processes as its main electrical response. The equation below illustrates the determination of complex electric modulus (M^*):³⁹

$$M^* = \frac{1}{\varepsilon^*} = \frac{\varepsilon'}{(\varepsilon')^2 + (\varepsilon'')^2} + \frac{i\varepsilon''}{(\varepsilon')^2 + (\varepsilon'')^2} = M' + iM'' \quad (9)$$

Fig. 11 and 12 depict the variations of M' and M'' with the applied electric field frequency (presented on a logarithmic scale) for the investigated films. The observed changes in M' exhibit a sigmoidal-shaped curve. This behavior suggests that capacitive behavior dominates the electrical response at low frequencies. In other words, the films exhibit a high initial ability to store electrical energy. Interestingly, the M' values decrease with increasing SnO₂ nanoparticle concentration. This trend implies a potential improvement in the real component (storage capability) of the dielectric constant at higher nanoparticle loadings. Fig. 12 presents the variations in the imaginary component of the electric modulus (M'') with frequency for the investigated films. A prominent relaxation peak is observed in these curves. Interestingly, the magnitude of this peak decreases with increasing SnO₂ nanoparticle concentration. The presence of relaxation peaks in the M'' spectra is often

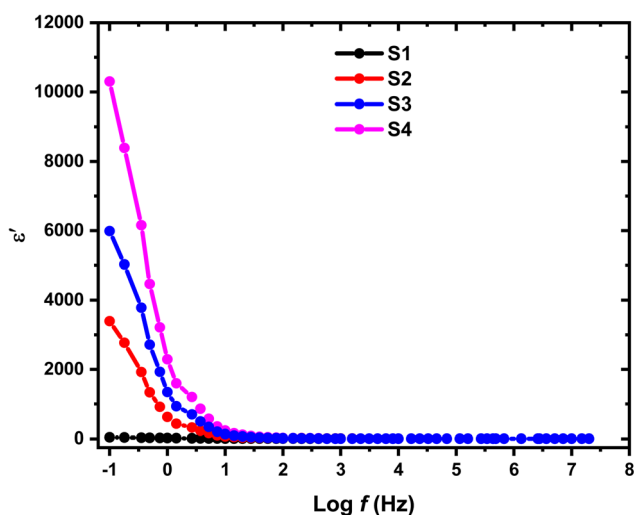


Fig. 9 Variation in dielectric constant (ε') with $\log(f)$ for HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS loaded with SnO₂ NPs.

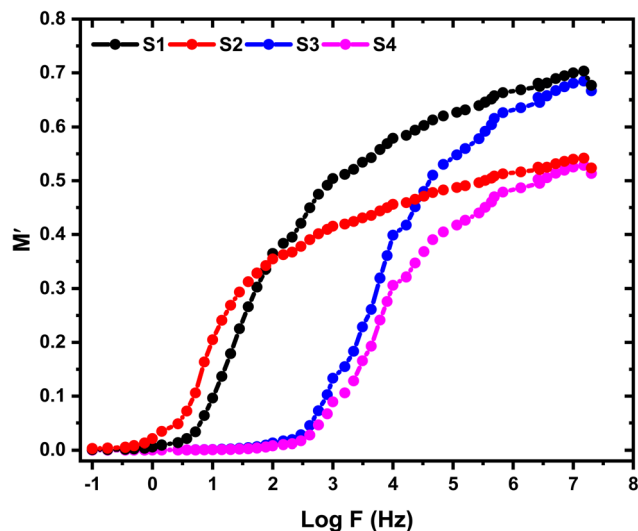


Fig. 11 Frequency dependence of M' for the pristine HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS nanocomposites containing various loadings of SnO_2 nanoparticles.

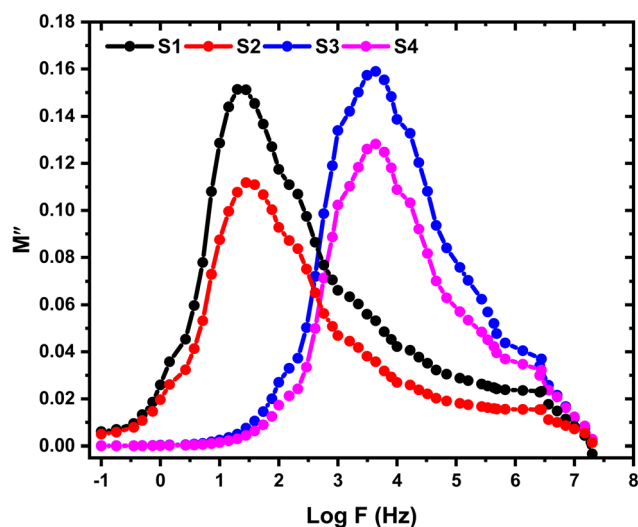


Fig. 12 Frequency dependence of M'' for the pristine HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS nanocomposites containing various loadings of SnO_2 nanoparticles.

associated with the existence of ionic conductivity within the material. The observed peak's significant height and asymmetry suggest a distribution of relaxation times within the films. This behavior deviates from an ideal Debye-type relaxation process, which assumes a single, well-defined relaxation time.⁴⁰ At frequencies lower than the peak frequency observed in M'' , the M'' response suggests a dominance of DC conductivity. This likely involves the long-range movement of charge carriers through hopping mechanisms.⁴¹ In contrast, at frequencies exceeding the peak, the M'' behavior indicates a shift towards AC conductivity. Here, ions become confined within potential wells, and their motion becomes restricted to short distances within these confined regions. Consequently, the peak

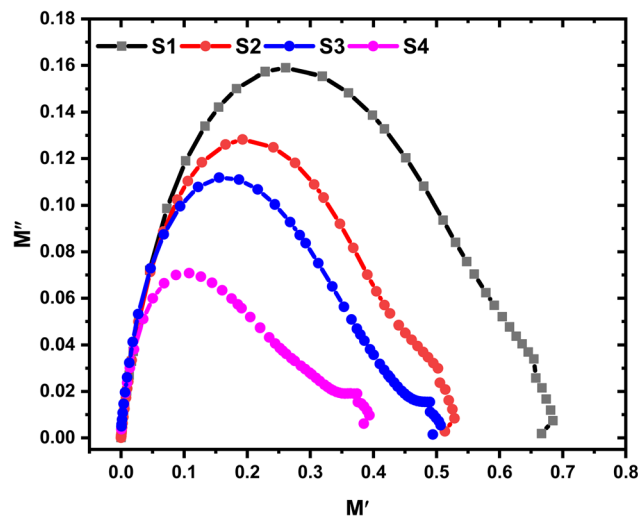


Fig. 13 Argand plot for HPMC/PEDOT:PSS ternary blend and HPMC/PEDOT:PSS loaded with SnO_2 NPs.

frequency in the M'' spectra can be interpreted as a transition point. As the applied field frequency increases, the mobility of charge carriers changes from long-range (at lower frequencies) to short-range (at higher frequencies).

3.8. Argand plot

Fig. 13 presents the Argand plots for the HPMC/PEDOT:PSS composite and HPMC/PEDOT:PSS nanocomposites containing varying SnO_2 nanoparticle concentrations. An Argand plot visualizes the relationship between the real (M') and imaginary (M'') components of the electric modulus. Ideal Debye-type relaxations, characterized by a single relaxation time, appear as perfect semicircles in the Argand plot. However, polymers often exhibit a distribution of relaxation times, leading to a depressed semicircular arc.⁴² The center of this arc, located on the M' axis, represents a characteristic relaxation time for the material. The length of this depressed semicircle has been associated with the material's conductivity. Shorter arc lengths tend to indicate higher conductivity.⁴³ In Fig. 13, we observe a subtle decrease in the arc length with increasing SnO_2 concentrations, suggesting a potential improvement in conductivity for these nanocomposites. Furthermore, the movement of the Argand plots towards the origin at higher filler loadings further supports this trend. The observed deviation from ideal Debye behavior likely arises from the presence of multiple polarization mechanisms, diverse relaxation processes within the material, and various interactions between ions and dipoles within the nanocomposite.

4. Conclusion

A solution casting method was utilized to create nanocomposite films. These films incorporated varying concentrations of SnO_2 nanoparticles dispersed within a matrix composed of HPMC and PEDOT:PSS. X-ray diffraction (XRD) analysis confirmed the presence of tetragonal SnO_2 NPs with sizes ranging from 26 to



43 nm. Interestingly, the XRD results also revealed a decrease in the crystallinity of the SnO₂ NPs upon incorporation into the HPMC/PEDOT:PSS composite. Fourier-transform infrared spectroscopy (FTIR) analysis of the nanocomposites indicated changes in the vibrational peak positions following the introduction of SnO₂ NPs. This suggests potential interactions between the polymer matrix and the nanoparticles. UV-visible spectroscopy measurements revealed a decrease in the optical bandgap of the nanocomposites with increasing SnO₂ NP content. This finding suggests a potential for bandgap engineering through the controlled incorporation of nanoparticles. The combined analysis using XRD, UV-visible spectroscopy, and FTIR techniques highlights the complex interactions occurring within the nanocomposites upon SnO₂ NP incorporation. Furthermore, the research demonstrates that the nanocomposites exhibit improved dielectric properties and AC electrical conductivity at room temperature with increasing SnO₂ NP concentration. These findings, coupled with the observed tunable optical properties, suggest the potential of these nanocomposites for applications in optoelectronic devices.

Data availability

The data that support the findings of this study are available from the corresponding author upon reasonable request.

Author contributions

H. M. Ragab: methodology, formal analysis, investigation, writing – review & editing. N. S. Diab: investigation, writing – review & editing. Shimaa Mohammed Aboelnaga: methodology, formal analysis, investigation. Jumana N. Dawoud: methodology, formal analysis, investigation, writing – review & editing. Mahmoud AlElaimi: investigation, writing – review & editing. M. O. Farea: investigation, writing – review & editing.

Conflicts of interest

There are no conflicts to declare.

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