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Structural, dielectric, and antimicrobial evaluation of PMMA/CeO₂ for optoelectronic devices

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In the current report, we have successfully synthesized nanocomposites of PMMA incorporating different doping of CeO₂ through a chemical approach. XRD results reflects decent matching for CeO₂ nanoparticles with 29 nm crystallite size. FTIR spectroscopy demonstrates the characteristic functional groups validating the successful formation of the composite. The optical study of PMMA and the nanocomposites has proven that the optical properties such as band gap, refractive index, optical permittivity, and loss tangent factor are affected by adding CeO₂ to the PMMA matrix. The peak residing around 420 nm by UV measurements is allocated to occurring electrons photoexcitation from the valence to conduction band inherent in CeO₂. The dielectric measurements were achieved using broadband dielectric spectroscopy upon a wide span of frequencies (10⁻¹–10⁷ Hz) and within temperatures from – 10 to 80 °C with a step of 10 °C. The permittivity decreases by adding CeO₂ and the dielectric parameters are thermally enhanced, however, the temperature influence is based on CeO₂ content, the higher the CeO₂ amount, the higher the influence of temperature. The results of the nanocomposites revealed antibacterial activity counter to gram-positive bacteria strain (*S. aureus*, and *B. subtilis*), and gram-negative bacteria (*E. coli*, and *K. pneumoniae*), yeast (*C. albicans*, as well as fungi (*A. niger*). Inherently, the change in CeO₂ concentration from 0.01 to 0.1 wt% delivers maximum influence against gram-negative bacteria. These PMMA CeO₂-doped composites are beneficial for optoelectronic areas and devices.

The research is evolving rapidly to provide enhanced life conditions regarding the industrial and biomedical areas for numerous aspects. Nanotechnology and related nanoscale composites promote versatile characteristics that could not be accomplished for their counterpart coarse grains. The conjugation of polymers and inorganic 0D–3D nanoparticles in the nanocomposite structure offers a desirable set of physical and chemical merits based on their confinement size. Likewise, this conjugation paves the route for varied optimization in the optical, electrical, thermal, and biomedical specifications¹.

Nanocomposites explicitly engineered for several biomedical applications are regularly stated as "biomedical nanocomposites". There exists a multitude of probable biomedical nanocomposites, which can be characterized into many interconnected classifications. The foremost biological practices for these biomedical nanocomposites encompass the delivery of drugs, wound dressings, antibacterial assets, tissue engineering, stem cell remedy, cancer treatment, cardiac prosthesis, peripheral organs, biosensors, artificial blood vessels, as well as enzyme immobilization².

Polymethyl methacrylate (PMMA) is an amorphous polymer material that provides significant specifications due to its high transparency, increased mechanical properties, environmental safety, reduced cost, and feasible

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formation at limited temperatures accompanied by reduced thermal conductivity (around 0.0012 cal/s cm K)^{3,4}. These merits nominate it as a prominent contender for the polymer matrix^{1,5}. Furthermore, it is familiar with the ability to transmit light within around 300–1000 nm wavelength without significant loss which highlights its eminent optical performance^{3,6}. PMMA is naturally insulating, however upon certain doping with the preceding metal oxides its electrical properties could be modulated⁴.

Metal oxides are a major abundant material on Earth⁷, composed of more than one element thus they might be modulated regarding their electrical, optical, and morphological specifications for a certain application. These metal oxides are familiar with their elevated bandgap value that delivers multiple properties beneficial for sensing appliances^{8,9}, catalysis instruments¹⁰, storage devices, and optoelectronic devices^{11,12}.

Cerium oxide nanoparticles, regularly known as nanoceria (CeNPs), are broadly recognized as highly auspicious metal oxide nanomaterials. The options available are either a pristine (bare) state or a state stabilized by ligands. Cerium-based nanoparticles (CeNPs) have grown in recognition as effective therapeutic agents in the fields of regenerative medicine and tissue engineering. In vitro studies have exploited their capability to promote cell proliferation, while in vivo experiments have exposed their role in expediting the healing process of lesions. These findings have significantly influenced the outlook on wound therapy, offering new possibilities for treatment^{13–16}.

CeO₂ nanoparticles are reported for their stability as well as biocompatibility where they demonstrate transparency within the visible light area and a 2.2 refractive index magnitude around 630 nm wavelength value^{17,18}, besides a raised bandgap of 3.2 eV¹⁹. These advantages are favored for optoelectronic applications²⁰. The CeO₂ emerges as a significant metal oxide in the photocatalysis field²¹. Broker et al., studied the performance of photocatalytic activity for CeO₂ when exposed to the sunlight directed towards the organic dye degradation²². Other reported research efforts revealed the impacts of the additives on CeO₂ and their performance in catalytic²³. TiO₂ and CeO₂ are usually combined for the collective features of their catalytic activity in general²⁴. Former research works were directed to the impact of CeO₂ on PMMA¹ or TiO₂/PMMA²⁵ and CeO₂/TiO₂²⁶.

The specifications provided by the polymeric network structure reinforced with a cerium oxide comprising elevated surface area, wide bandgap, conductivity, visible light transparency, provided electronic mobility, and antibacterial tendency offers a wide spectrum of photocatalysis, solar energy, and optoelectronic applications^{18,27,28}.

In this report, we introduce the simple synthesis route to acquire PMMA/CeO₂ with a limited doping molar ratio of the metal oxides to reveal their impact on the optical, electrical, and antimicrobial specifications. XRD, FTIR, optical, dielectric, and antimicrobial measurements were assessed via their conventional devices. This combined composite might shed light on the next stage of nanotechnological and industrial applications. The purpose of the intended effort is to improve the optical, electrical, and antimicrobial characteristics of the PMMA upon incorporating small portions of dopants. These specifications can be applied in a wide set of biomedical and optoelectronic applications.

Experimental work

Materials

Poly methyl methacrylate (PMMA), Sigma Aldrich, Toluene anhydrous (C₆H₅CH₃), 99.8%, Merck, Cerium Oxide (CeO₂), Sigma Aldrich.

Preparation of PMMA/CeO₂ nanocomposites

To prepare the samples, a certain amount of PMMA was dissolved in 50 ml of toluene, then CeO₂ was added gradually. After that, they were homogeneously blended by high-speed stirring for 2 h at 50 °C to acquire a consistent dispersion of the CeO₂ in the PMMA matrix. Lastly, the constituted nanocomposite mixtures were decanted using petri dishes, and then they were aged for two weeks. The dry powders of PMMA and CeO₂ were precisely weighted to acquire quantified weight percentages: zero, 0.01, 0.05, and 0.1 wt.%, which were marked as PMMA, PC1, PC2, and PC3, respectively.

Measurements

The XRD measurements were conducted using Cu K α -X-ray powder diffraction (XRD) by Bruker D-8 diffractometer instrument functioned at 35 kV and 30 mA, using a (0.05°) step size within a scanning range (5–70°). For measuring the IR spectra, we have employed the device Bruker optics VERTEX 7000 Fourier Transform Infrared Spectrometer. The acquired spectra were plotted in a spectral span of 4000–400 cm^{−1} having a 2 cm^{−1} resolution and 2 mm/s scanning speed. The optical measurements were attained by a Jasco V-570 spectrophotometer throughout the wavelength range (0.2–2.5 μ m). The dielectric studies were achieved using broadband dielectric spectroscopy upon a wide range of frequencies (10^{−1} to 10⁷ Hz), and at temperatures between (−10 to 80 °C) with a step of 10 °C. The technique employs an increased-resolution ALPHA analyzer having an active sample holder head (Model: Novocontrol, Montabaur, Germany), accompanied by an active sample head. A Quatro temperature controller system was utilized for the stabilization of temperatures less than 0.2 K, and pure nitrogen was used as a heating agent. Gold-plated stainless-steel electrodes with a 20 mm diameter having a configuration of parallel plate capacitor were used for the measuring process.

Antimicrobial activity

To evaluate the CeO₂ antibacterial efficiency. Test organisms comprised yeast (*Candida albicans* ATCC 10231), fungi (*Aspergillus niger* NRRLA-326), and gram-positive and gram-negative bacteria (*Staphylococcus aureus* NRRLB-767 and *Bacillus subtilis* ATCC 6633, *Escherichia coli* ATCC 25922, *Klebsiella pneumoniae* ATCC 10145) were employed. Flat polystyrene plates with 96 wells were functioned for performing the tests. Succeeding the

addition of 10 μL of the test extracts (to obtain a final concentration of 500 g/ml) to 80 μL of lysogeny broth (LB broth), and 10 μL of the bacterial culture suspension media (log phase), further, the plates were incubated at 37 °C overnight. The absorbance amount was counted after around 20 h at OD600 in a Spectrostar Nano Microplate Reader (BMG LABTECH GmbH, Allmendgrun, Germany)^{29,30}.

Antibiofilm activity

The biofilm inhibitory efficiency of the pigment extract was assessed using the microtiter plate assay (MTP) approach. The experiment elaborated 96 wells of flat-bottom polystyrene plates, each encompassing two clinical pathogens (*E. coli* ATCC 25922 and *S. aureus* NRRLB-767). Every well was occupied by 180 μL of lysogeny broth (LB) and 10 μL of overnight bacterial growth sample. The plate was then incubated for 24 h at 37 °C with 10 μL of CeO_2 at dissimilar concentrations, accompanied by a negative control entailing filtrate without the sample. To eradicate the presence of suspended bacteria, the liquid within each well was detached, and subsequently, each well was subjected to a washing procedure including the use of 200 μL of phosphate-buffered saline solution with a pH value of 7.2. The plate underwent a staining process for one hour, during which a solution of crystal violet with a concentration of 0.1% (w/v) was introduced to every well. Subsequently, rinsed by distilled water (200 μL), after which the plate was left to dry inside a laminar flow environment. The dry plate was treated with ethanol (95%), and the optical density (OD)-at 570 nm-was monitored using a SPECTROSTAR nano absorbance plate reader (BMG LABTECH) for quantification purposes^{31,32}.

Results and discussion

XRD

The PMMA/ CeO_2 synthesized nanocomposites constructed with CeO_2 weight portions of 0, 0.01, 0.05, and 0.1 wt.%, were marked as PMMA, PC1, PC2, and PC3, respectively. The chemical structure was evaluated through XRD examination, as demonstrated in Fig. 1. Figure 1 (inset) illustrates the XRD spectrum of the CeO_2 nanoparticles. The pattern displays that the distinctive peaks are found at $2\theta = 28.9, 33.4, 47.7, 56.8, 59.1, 69.6, 77.1, 79.4$, and 88.5° which are owing to (111), (200), (220), (311), (222), (400), (331), (420), and (422) lattice planes, correspondingly. The acquired lattice parameters of CeO_2 nanoparticles coincide with former research data (JCPDS 34-0394)^{33,34}. The XRD spectrum of cerium dioxide nanoparticles possesses wide peaks, which validate the construction of mini-sized nanoparticles. The familiar lattice parameters values attained from the XRD information were observed as ($a = b = c = 5.411 \text{ \AA}$), where ($\alpha = \beta = \gamma = 90^\circ$). The average crystallite dimension of CeO_2 nanoparticles was identified using Scherrer's equation^{35–37}.

$$D = \frac{0.94\lambda}{\beta \cos \theta} \quad (1)$$

Inherently, λ is incident X-ray source wavelength (1.54060 \AA), θ is the Bragg's diffraction angle, and β is the full width at mid-maximum (FWHM) stated in radians³⁸. The nominal crystallite size was calculated and found about 29 nm.

The PMMA sample displays three principal, wide XRD peaks, localized at 13.8° (the high-intensity band), besides two reduced-intensity peaks, noticed at 29.7° and 41.9° , which designate the amorphous character of the PMMA polymeric matrix³⁹.

Additionally, this behavior was noticed in all the PMMA/ CeO_2 nanocomposites samples with low intensity. The PMMA/ CeO_2 nanocomposites samples' XRD diffraction patterns disclosed the amorphous nature of PMMA, signifying that CeO_2 integration had no impact on the structural features of the PMMA and that no chemical interaction had occurred between the PMMA and CeO_2 .

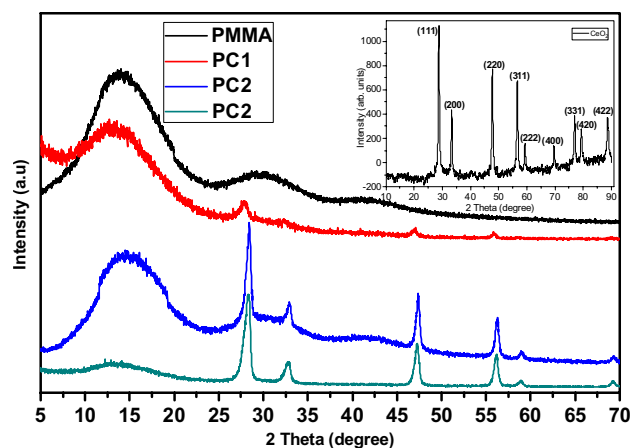


Figure 1. XRD pattern for PMMA, CeO_2 nanoparticles, and PMMA/ CeO_2 nanocomposites. The right inset shows the indexed diffraction planes and their corresponding angles.

Optical analysis

The investigation of the optical aspects of the prepared samples provides powerful details about their electronic properties and design. When examining materials using spectroscopy, the electronic vibration states are applicable. The electronic excitation is represented by the UV–Vis diffuse reflectance (DRS) part of the electromagnetic (EM) spectrum, and the energy levels are determined by the chemical bonds existent throughout the composite structure. To determine the optical properties of PMMA doped with CeO₂ nanoparticles, the UV–Vis spectra were reported. Figure 2 reveals the UV spectra for all samples. The UV–Vis spectrum of PMMA displays very weak reflectance peaks at 300 nm, while the nanocomposites exhibit reflectance peaks and their intensity increases with CeO₂ content in PMMA matrix. The peaks of nanocomposites were at 424, 456, and 480 nm for PC1, PC2, and PC3 respectively. Moreover, the reflectance of the nanocomposites is altered by different quantities of CeO₂, resulting in a corresponding change in absorbance. The Kubelka–Munk relation was employed to measure the degree of change in the band gap energy (E_g),^{40,41}

$$F(R) = \left(\frac{\alpha}{S} \right) = \left[\frac{(1 - R)^2}{2R} \right] \quad (2)$$

$F(R)$, α , and S are all present, and denoted as the Kubelka–Munk function, the absorption coefficient, and the scattering factor, correspondingly.

The optical energy bandgap (E_g) was evaluated by the proposed method introduced by Tauc equation⁴²,

$$F(R)h\nu = A(h\nu - E_g)^r \quad (3)$$

where $h\nu$ is the incident radiation's photon energy, A expresses a constant depending on the electronic transition probability, E_g denotes the optical band gap. The power factor, expressed by r , varies based on the type of transition. Additionally, the electronic transition's characteristics can be specified by defining the value of r . For direct transitions, r can be either $1/2$ or $3/2$, while for indirect transitions, r is equal to 2 or 3, depending on whether they are allowed or disallowed, respectively⁴³. Normally there are two categories of materials for insulators and semiconductors: direct band gaps and indirect band gaps. The valence band maximum (VBM) and conduction band minimum (CBM) in direct band gap materials concur at the identical zero-crystal momentum position (i.e., wave vector $k=0$)⁴⁴. In this occasion, r receives the magnitude of $1/2$. When the quantum selection rule prohibits a direct transition amongst the valence band maximum (VBM) and conduction band minimum (CBM) in certain materials, this transition is referred to as a forbidden direct transition. In this case, the value of r is equal to $3/2$. An indirect electron transition arises when the VBM and the CBM are not located on the identical wave vector. In this instance, the VB to CB electronic transition with the appropriate crystal momentum magnitude will continually be associated with the absorption/emission of phonon energy⁴⁴.

The optical energy gap is calculated from the relation $(F(R)h\nu)^2$ versus $h\nu$ through extrapolating the linear part of the curve to zero as illustrated in Fig. 2b. The band gap variation is shown in the inset figure, which is an amplification of Fig. 2b. The energy band gap of pure PMMA was determined to be 4.15 eV and it is consistent with numerous earlier reports^{45,46}. The energy band gap of the nanocomposite samples was found 4.15, 3.34, 3.30, and 3.31 eV for pure PMMA, PC1, PC2, and PC3 respectively. Evidently, as the CeO₂ content elevates in PMMA matrix, the optical energy band gap magnitudes drop. The addition of CeO₂ nanoparticles in the polymeric host matrix leads to the emergence of extra absorption bands in the UV and visible portions of the optical spectra⁴⁷. The decrease in the energy band gap could potentially be explicated by the intermolecular interactions that occur between the polymer and CeO₂ particles. The gradual accumulation of CeO₂ in the polymer matrix promotes significant modifications in the composites' structure, such as the enhancement of the conjugated chain lengths,

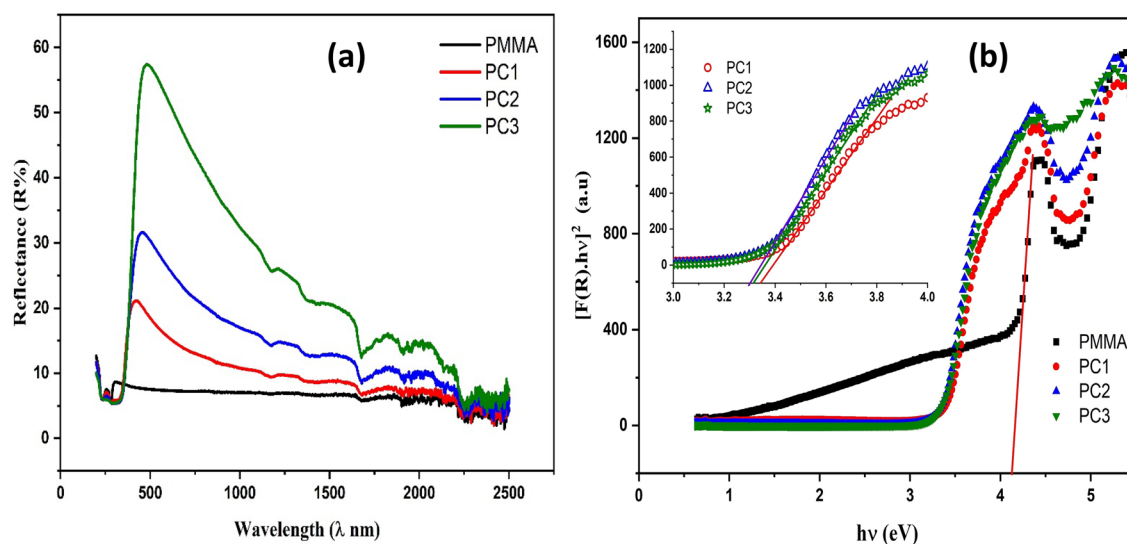


Figure 2. (a) Diffuse reflectance (R), (b) Tauc's plot of PMMA and PMMA/CeO₂ nanocomposites.

resulting in a reduction of the band gap⁴⁷. When assessing the electronic and optoelectronic qualities of a material, the composite's refractive index and optical dielectric characteristics are crucial. The refractive index of the PMMA/CeO₂ nanocomposites can be considered using the following equation⁴⁸,

$$\left\{ n = \left[\frac{(1+R)}{(1-R)} \right] + \sqrt{\frac{4R}{(1-R)^2} - k^2} \right\} \quad (4)$$

Herein, k represents the extinction coefficient and has been determined via the relation ($k = \alpha\lambda/4\pi$), where λ is the wavelength. Figure 3a demonstrates the relationship between the refractive index (n) and wavelength for both PMMA and the nanocomposites that were developed. The refractive index upsurges with raising the wavelength to reach the maximum value then decreases for all samples. Figure 3a shows that as CeO₂ content enlarged, the refractive index of the nanocomposites also increased. The rise in the refractive index may be accredited to the change in PMMA structure due to CeO₂ addition, which leads to the enhancement of the conjugated chain lengths.

The optical permittivity is connected with the refractive index (n), and extinction coefficient (k) via the following equation⁴⁸,

$$\varepsilon' + i\varepsilon'' = (n^*)^2 = (n + ik)^2 = n^2 - k^2 + i2nk \quad (5)$$

Figure 3b depicts the relationship between the real part of optical permittivity (ε') and wavelength. The real part of optical permittivity (ε') shows the same trend as the refractive index where it increases with frequency to a maximum value and decreases and increases with CeO₂ content. The increased optical permittivity with CeO₂ content might be ascribed to the generation of additional free charges triggered by the presence of CeO₂, acting as a polarization center, hence enhancing the polarizability of the nanocomposites. The imaginary part of optical permittivity (ε'') versus wavelength in Fig. 3c demonstrated the same behavior of the real part of optical permittivity with wavelength, but the behavior of the imaginary part of dielectric (ε'') with CeO₂ concentration is opposite to real part of optical permittivity (ε'), which decreases with increasing CeO₂ concentration, as

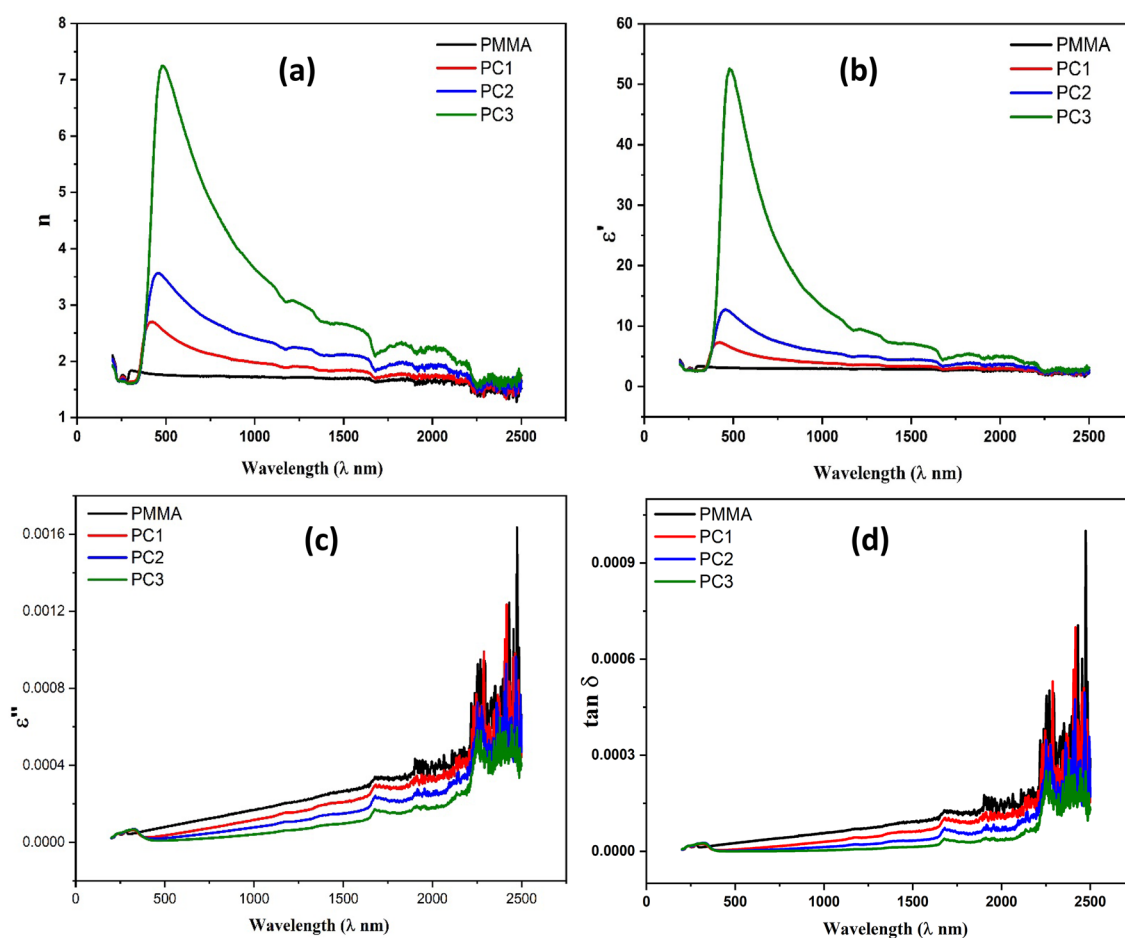


Figure 3. (a) The propagation of the refractive index (n), (b) Real section of the optical permittivity (ε'), (c) Imaginary part of the optical permittivity (ε'') and (d) Optical dielectric loss tangent ($\tan\delta$) of PMMA and PMMA/CeO₂ nanocomposites.

illustrated in Fig. 3c. The loss factor (dielectric loss tangent) was either calculated through the subsequent relation⁴⁹:

$$\left[\tan \delta = \frac{\varepsilon''}{\varepsilon'} \right] \quad (6)$$

The relationship ruling wavelength and optical dielectric loss, for pure PMMA and PMMA/CeO₂ nanocomposites is displayed in Fig. 3d. The optical dielectric loss ($\tan \delta$) was found to decrease as the concentration of CeO₂ in the nanocomposites increases and it increases with frequency.

FTIR analysis

(FTIR) spectroscopy is an advanced tool that might be employed to determine the handy functional groups of the prepared nanocomposite samples. As infrared radiation overrides via the sample, a specific ratio of the radiation intensity would be captivated and that will be revealed in the FITR spectrum. According to occurring stretching and bending vibrations within the samples, the IR radiation absorption appears at specific frequency values depending on the fabric of the sample. The chemical formula, sample's functional groups can be identified by following the absorption bands or peaks in the wavenumber range 400 cm⁻¹ to 4000 cm⁻¹ of the FTIR spectrum. The FTIR spectrum of PMMA displays two peaks at 299 and 2950 cm⁻¹, as presented in Fig. 4, which are assigned to C-H stretching vibration. The intense band positioned at 1724 cm⁻¹ was attributed to C=O stretching vibration, whereas the two peaks that appeared at 1487 and 1442 cm⁻¹ were assigned to -CH₂ and -CH₃ asymmetric stretching or deformation of PMMA. The peaks presented at 1383, 1203, and 1140 cm⁻¹ are credited to O-CH₃ deformation, C-O stretching, and -O-CH₃ stretching of PMMA, respectively. CH₂ twisting, wagging, and rocking vibrations of PMMA are located at 1195, 990, and 745 cm⁻¹. The FTIR spectra for PMMA/CeO₂ nanocomposites samples demonstrate the main characteristic peaks of PMMA as it denoted the spectra, except for the emergence of small peaks located at 430 cm⁻¹, which is accredited to the metal-oxygen bond of cerium dioxide nanoparticles.

Dielectric study

The permittivity is a complex function given by⁵⁰

$$\varepsilon^* = \varepsilon' - i\varepsilon'' \quad (7)$$

where ε' is the real part which is a degree of the material's capability to store charges, and ε'' is the imaginary part which reflects the energy loss in a material. The variation of permittivity of all samples vs frequency at different temperatures is manifested in Figs. 5 and 6. The permittivity of PMMA has higher values at lower frequencies and then falls gradually with increasing frequency to be almost frequency-independent. The increased magnitude of permittivity in the low-frequency regime is believed to arise from the dipoles' alignment and obeying the applied field. The field variation becomes faster at a higher frequency and the dipoles lag and can't follow up the changes in the applied field which causes a drop in the polarization, hence the permittivity decreases and is no longer affected by frequency⁵⁰. The addition of small amounts of CeO₂ to PMMA altered its dielectric properties.

Adding CeO₂ had two major effects. First, it hinders the polymeric chain mobility, which reduces its submission to the field and hence decreases the permittivity⁵¹. The second, it produces an interface inside the polymer matrix at which charges accumulate, which boosts the interfacial polarization and increases the permittivity. The permittivity is found to decrease with increasing CeO₂ content, which means that the first factor prevails. The permittivity is thermally activated for all samples and the rate of increment of permittivity with temperature is dependent on CeO₂ content, the highest content has the greatest rate. For the pure sample PMMA, by increasing temperature the permittivity values increase, the temperature increases the mobility of

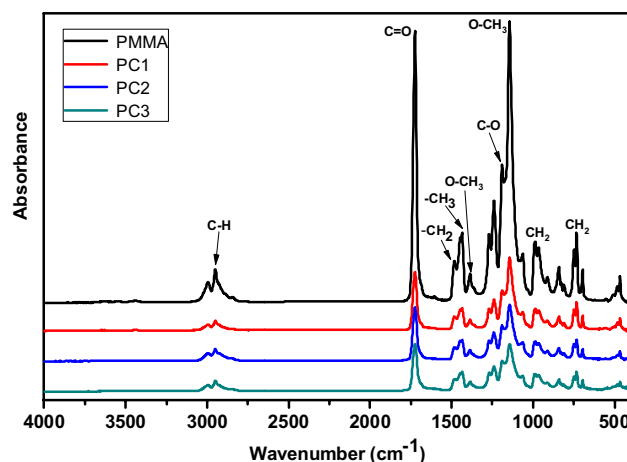


Figure 4. FTIR spectrum for PMMA, and PMMA/CeO₂ nanocomposites.

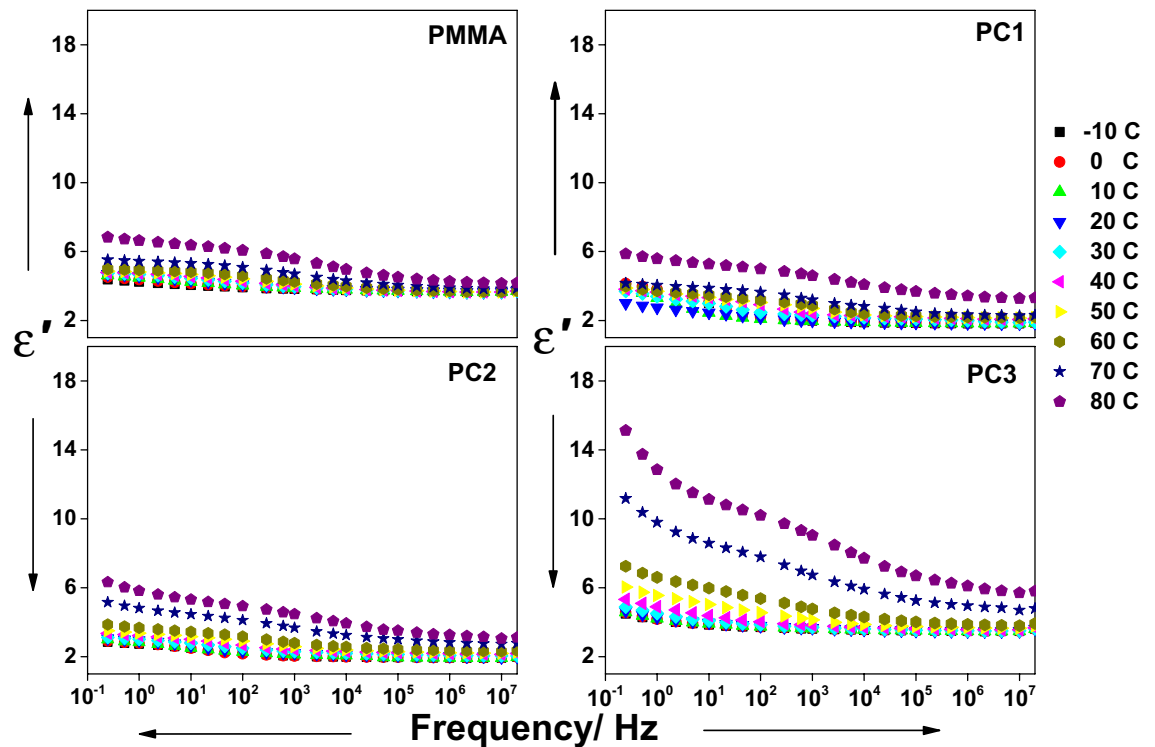


Figure 5. The real section of permittivity (ϵ') vs frequency for all composites at temperatures between -10 and 80 °C, as indicated.

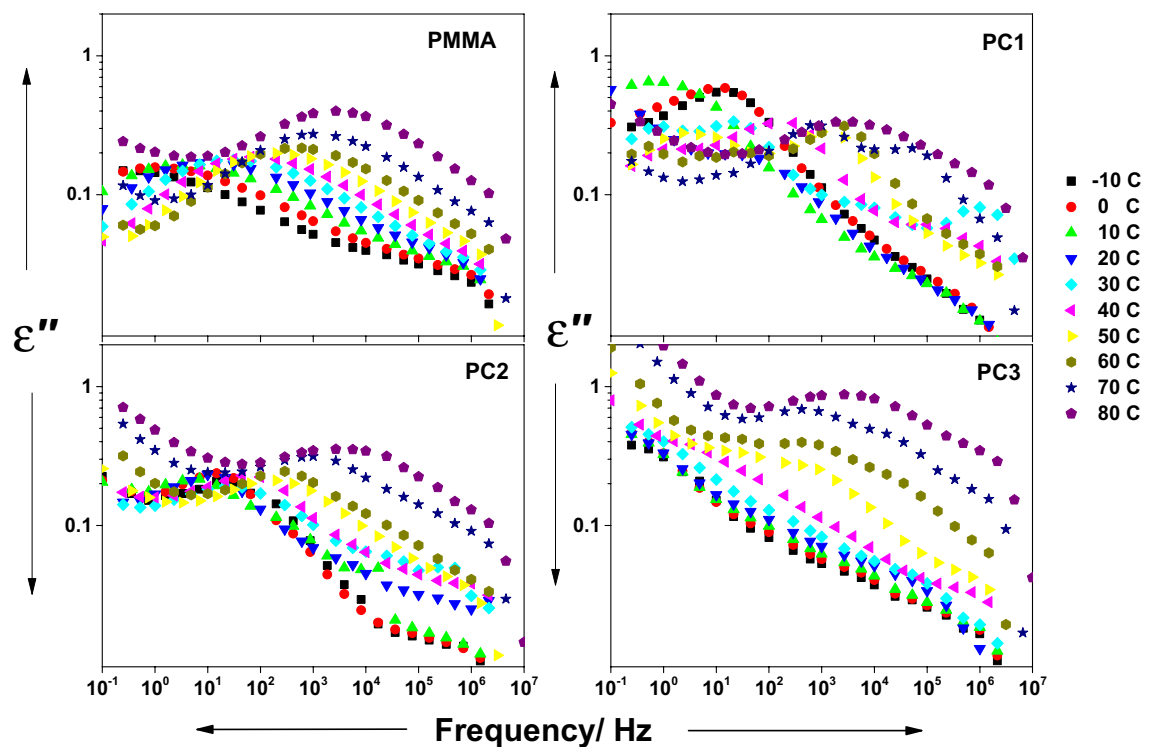


Figure 6. The imaginary part of permittivity (ϵ'') vs frequency for the whole composites, at temperatures between -10 and 80 °C, as indicated.

the dipoles and upsurges the conductivity, which increases the permittivity. By increasing the temperature for the samples containing small amounts of CeO_2 , besides the effects on PMMA dipoles, more charge carriers are liberated to accumulate at the interfaces between CeO_2 and PMMA, which increases the permittivity.

The imaginary part of permittivity is presented in Fig. 6. The (ϵ'') of pure PMMA has three dielectric relaxations, interfacial polarization, conductivity contribution at low frequency, and β -relaxation at high frequency, which is ascribed to ester group with other accompanied methylene group⁵².

The conductivity contribution, which appeared in the curve as a linear decrease in the dielectric loss, is only noticed when the temperature is raised. The interfacial polarization and β -relaxation become faster with increasing temperature, and their corresponding peaks merge into a larger one. The conductivity contribution in the dielectric loss for samples PC1, PC2, and PC3 shows up at lower temperatures compared with the neat sample, and PC3 shows the highest conductivity contribution which suggests it is more conductive than other samples. The relaxation peaks are shifted to elevated frequencies with increasing CeO_2 content which means their relaxation time decreases.

Electrical conductivity in the materials is a thermally activated procedure that occurs due to the motion of relatively loose charge carriers influenced by an electric field. The conductivity in numerous materials is frequency-dependent and obeys Jonscher's power law⁵³,

$$\sigma'(\nu) = \sigma_{dc} [1 + (\nu/\nu_c)^s] \quad (8)$$

where ν_c the hopping frequency separating DC and AC regimes, σ_{dc} represents DC conductivity and the exponent s represents a temperature-dependent parameter that holds $0 < s \leq 1$, depending on the conduction mechanism⁵⁴.

The AC conductivity vs frequency for all samples is shown in Fig. 7. The conductivity appears to be frequency dependent and no plateau or DC conductance is observed even when the temperature is raised or when CeO_2 is added, which is the prime trait of insulators. The conductivity increases with frequency and it increases by increasing temperature or CeO_2 content. The heat causes liberation to some charge carriers. The addition of CeO_2 increases the crystallinity of the samples which provides facile routes for charge carriers' transport. The sample PC3 has relatively higher conductivity and is slightly affected more than other samples by increasing temperature, which appears as semi-linear decrease in the dielectric loss at lower frequency.

Antimicrobial and antibiofilm results

In our work, the antibacterial activity of CeO_2 was evaluated using six distinct microorganisms (*B. subtilis* ATCC6633, *S. aureus* NRRLB-767, *K. pneumoniae* ATCC10145, *E. coli* ATCC25922, *C. albicans* ATCC 10231 and *A. niger* NRRLA-326). The outcomes in the documentation demonstrated antibacterial activity against the whole bacterial strains, mainly with increasing the concentration of CeO_2 from 0.01 to 0.1, especially against

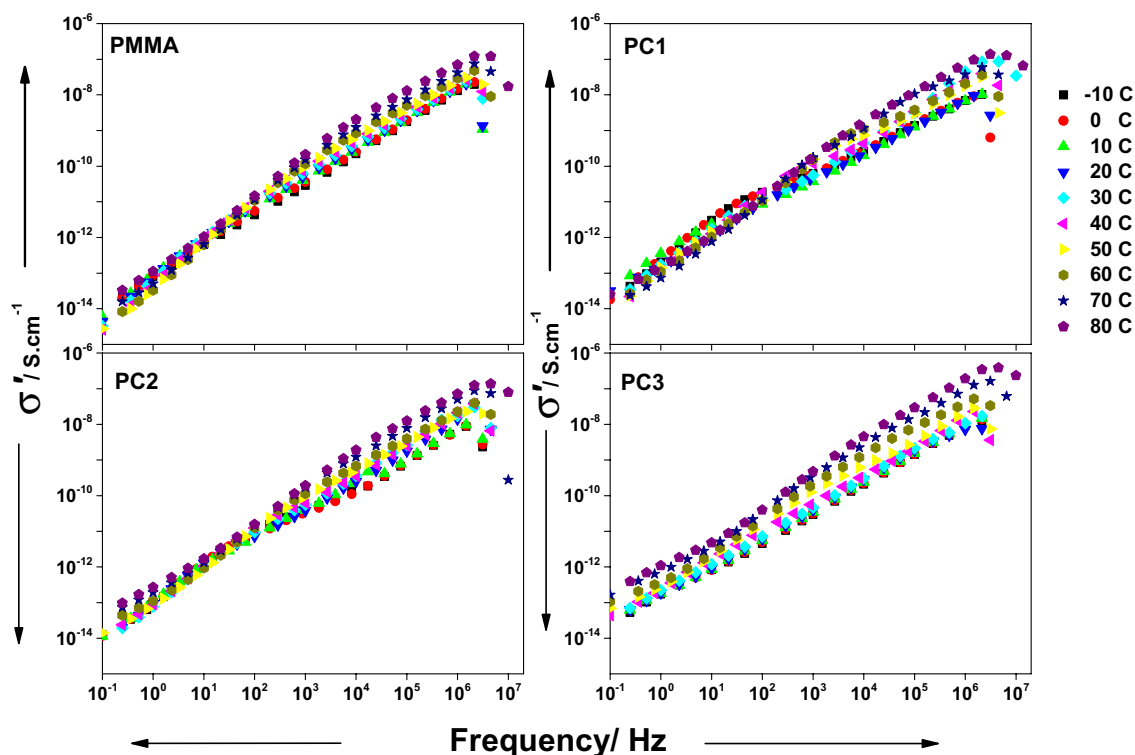


Figure 7. AC conductivity (σ') vs frequency for all samples at temperatures ranging from -10 to 80 °C, as indicated.

gram-negative bacteria (*K. pneumoniae* ATCC10145 and *E. coli* ATCC25922). In contrast, the antifungal activity of the CeO₂ against *C. albicans* ATCC 10231 and *A. niger* NRRLA-326 was insignificant. While the antibiofilm activity of CeO₂ displayed a significant result for *S. aureus* NRRLB-767 with increasing the concentration of CeO₂ (Tables 1, 2) The findings of this study align with those of former research⁵⁵, which demonstrated the synthesis of a nanocomposite consisting of cerium oxide (CeO₂) and graphene oxide. The nanocomposite exhibited promising antibacterial activity against various wound pathogens, including *E. coli*, *P. aeruginosa*, *S. aureus*, and *S. typhi*. The concentrations employed in the study were 25 µg/ml, 50 µg/ml, 75 µg/ml, and 100 µg/ml.

The membranes of both gram-positive and gram-negative bacteria were exposed to adsorption by the positively charged nanoparticles due to the electrostatic interaction, which can be credited to several mechanisms. The prolonged existence of nanoparticles (NPs) on the bacterial surface can be accredited to the electrostatic contact and the obstruction of the bacterial membrane, which suppress their penetration into the membrane. Subsequently, the introduction of nanoparticles (NPs) has the potential to vary the cellular membrane's viscosity, hinder the functionality of particular ionic pumps, and ultimately disrupt the transport processes involved in the interchange of substances between the bacterial cell and its surrounding solution, thereby perturbing bacterial evolution⁵⁶. After adsorption onto the exterior membrane of the bacterial cell, CeO₂ has the potential to interact with and disrupt proteins. According to preceding studies⁵⁷, the incidence of cerium ions has the potential to interrupt electron flow and respiratory processes in bacteria. Additionally, these ions can interact with thiol groups (–SH) or bind to transporters and/or porins, thereby impeding the delivery of vital nutrients. Further, the irregular forms and rough edges of CeO₂ itself are responsible for causing physical impairment to bacterial membranes, principally in gram-positive bacteria case⁵⁸.

Conclusion

The feasible synthesis approach of PMMA/CeO₂ nanocomposites is delivered in this work with varying amounts of CeO₂. XRD crystallite size is moderately small as 29 nm. The investigation of the optical properties by UV–Vis showed that the band gap decreased from 4.15 eV for pure PMMA to 3.30 eV for PC2. The refractive index and optical dielectric constant increase with CeO₂, while the imaginary part of the dielectric constant and optical dielectric loss tangent decrease with CeO₂ concentration. The UV findings clarify electron photoexcitation at around 420 nm arising from the valence band and directing to the conduction band. The addition of small amounts of CeO₂ to PMMA decreased the permittivity but it made the influence of temperature higher. The permittivity is thermally activated for all samples and the rate of increment of permittivity with temperature is dependent on CeO₂ content. The dielectric spectra show three dielectric relaxations, interfacial polarization, conductivity contribution at low frequency, and β-relaxation at high frequency. The relaxation peaks are shifted to higher frequencies with increasing temperature or CeO₂ content, which means their relaxation time decreases and they move faster. The conductivity is frequency dependent which is the prime trait of insulators and it increases by increasing temperature or CeO₂ content. The antimicrobial specifications of the nanocomposites depict an antimicrobial activity towards gram-positive and gram-negative bacteria, further, yeast and fungi. These results might be beneficial and applied in the optoelectronic and biomedical application areas.

		{Antimicrobial activity (%)}					
		Gram-positive		Gram-negative		Yeast	Fungi
Compounds		(<i>S. aureus</i>) NRRLB-767	(<i>B. subtilis</i>) ATCC 6633	(<i>E. coli</i>) ATCC 25922	(<i>K. pneumoniae</i>) ATCC 10145	(<i>C. albicans</i>) ATCC 10231	(<i>A. niger</i>) NRRLA-326
PC 1	0.01 CeO ₂	13.08 ± 0.59	15.43 ± 0.59	22.39 ± 0.66	25.74 ± 0.49	0.0	0.0
PC 2	0.05	26.73 ± 0.58	28.54 ± 0.71	45.95 ± 0.30	49.60 ± 0.47	19.56 ± 0.68	17.62 ± 0.53
PC 3	0.1	38.85 ± 0.73	43.60 ± 0.89	59.46 ± 0.71	62.04 ± 0.51	34.83 ± 0.73	28.04 ± 0.48
Ciprofloxacin		96.01 ± 0.43	97.24 ± 0.18	98.07 ± 0.35	98.10 ± 0.27	–	–
Nystatine	–	–	–	–	–	97.16 ± 0.90	98.23 ± 0.16

Table 1. Antimicrobial activity (%) of the PMMA/CeO₂ nanocomposites.

		Biofilm inhibition ratio (%)	
Samples		(<i>E. coli</i>) ATCC 25922	(<i>S. aureus</i>) NRRLB-767
PC 1	0.01 CeO ₂	11.48 ± 0.73	17.40 ± 0.90
PC 2	0.05	22.06 ± 0.61	31.21 ± 0.47
PC 3	0.1	36.92 ± 0.80	42.75 ± 0.55

Table 2. Biofilm inhibition ratio (%) of the PMMA/CeO₂ nanocomposites.

Data availability

The data related to or connected with the work are all encompassed in the manuscript and its supporting information.

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Author contributions

A.M.B. conceived the offered idea, developed the experimental, and contributed to the first draft writing as well as data analysis. A.D. contributed to the experimental, writing of the manuscript content, data analysis, and interpretation. A.A.A. revised the manuscript, approved procedure validity, and contributed to the manuscript writing. M.E.E. and A.A.H. conducted the experimental, data collection, and results interpretation. A.E. contributed to the manuscript writing, data interpretation, and coordinated and supervised the results of this work. All authors discussed the results, agreed to the manuscript content, and contributed to the revised version of the manuscript.

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Competing interests

The authors declare no competing interests.

Additional information

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