







Dual nitrogen-sulfur-doping induce microwave absorption and EMI shielding in nanocomposites based on graphene

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Graphene-oxide (GO) is one of the most commonly used carbon nanomaterials in advanced applications such as microwave absorption and EMI shielding, due to various advantages such as ease of synthesis and exfoliation, effective doping capability, and superior composite compatibility. In this study, we used the modified Hummer's method to synthesize GO by exfoliating graphite powder, and a simple hydrothermal approach was employed for elemental doping and GO reduction. As nitrogen-sulfur (N, S) dual-doping precursors, thiourea and L-cysteine amino acids were utilized. The structural features and microporous network structure of GO aerogel foams were investigated. The microwave absorption capabilities of polyethersulfone-based nanocomposite films incorporating the as-produced nitrogen-sulfur enriched reduced GO (NS-rGO) are also explored. According to the physico-chemical characterization, the existence of remarkable structural defects with a porous three-dimensional (3D) network was discovered due to heteroatom insertion and hydrothermal doping. Additionally, the dual-doped sample exhibited high Nitrogen and sulfur content of 8.93% and 13.19%, respectively. While NS-rGO possesses a higher conductivity of 174.7 μS compared to 12.65 μS for GO. The nanocomposites filled with NS-rGO foams demonstrated a high shielding efficiency (SE) of 45 dB in the X-band with a filler loading of 0.5 wt.%. This high SE arises from dopant heteroatoms and the heterogeneous interface, which induce interface polarization, thereby increasing microwave absorption and dielectric constant. It also results from multi-level reflections caused by the 3D porous structures. These findings offer valuable insights into the functionalization of carbon nanostructures and the development of 3D networks in GO-based functional materials, providing further guidance for engineering high-performance electromagnetic interference shielding materials.

Keywords: Shielding performance; multi-level reflection; hydrothermal synthesis; porous structure.

1. Introduction

In response to the escalating problem of electromagnetic (EM) pollution, a considerable amount of attention has been directed toward the field of advanced functional materials¹ for the design and development of microwave absorption materials.^{2,3} Among these materials, polymer composites could be considered a promising choice for addressing the electromagnetic interference (EMI) problem.^{4,5} Their anti-corrosion properties, light weight, flexibility, and ease of processing make them beneficial for this purpose.⁶

In EMI shielding applications, nanocomposites based on generic polymers filled with carbonaceous materials such as carbon nanotubes (CNTs),⁷ carbon black,⁸ carbon fiber,⁹

and two-dimensional (2D) transition metal nitride/ carbide /carbonitrides (MXene),¹⁰ graphene,¹¹ and graphene oxide (GO)¹² have an advantage over rigid, metal-based shielding materials.

These conductive/carbonaceous fillers mitigate EMI by reflecting and absorbing EM waves through dielectric relaxation and conduction losses.¹³ Carbonaceous nanostructured materials, in general, are used as fillers in polymeric matrices due to their low density, high conductivity, thermal stability, high permittivity, and controllable aspect ratio.¹⁴ The polymer matrix plays two key roles in EMI shielding; it provides flexibility and mechanical strength to the EMI shielding layer and allows incoming EM waves to

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pass through the shielding layer because polymers often exhibit transparency toward high-frequency EM waves.¹⁵

According to Sushmita and co-workers, the introduction of structural defects in CNTs and the utilization of suitable functional groups that can enhance interactions with the host polymer play significant roles in improving shielding performance.¹⁶ CNTs are preferred because of their low cost, small diameter, ease of percolation, excellent electrical conductivity, and high mechanical strength.¹⁷ Another study found that covalent bonding between a polyaniline matrix and graphene aerogel fillers enhances electron transfer, effectively converting EM energy into heat energy, thereby improving microwave absorption capacity.¹⁸ Due to their exceptional electron mobility ($15,000 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$), large aspect ratios, high thermal conductivity ($5000 \text{ W m}^{-1} \text{ K}^{-1}$), tunable electrical properties, abundance of functional groups, and excellent mechanical characteristics, 2D graphene materials have emerged as popular alternatives for EMI shielding.¹⁹ However, the irreversible aggregation of graphene can significantly diminish its intrinsically high electrical conductivity. Fortunately, the problem of graphene agglomeration can be successfully addressed by using three-dimensional (3D) networks.²⁰

Recently, a variety of micro-structural designs of filler including segregate structures,^{21,22} porous structures,^{23,24} and multi-layer structures^{25,26} have been reported in the literature achieving valuable EMI shielding performance with low filler loading. Researchers have increasingly focused on fabricating porous structures of carbonaceous materials, such as foams²⁷ or aerogel,¹⁸ due to their lightweight, high specific surface area, and high dielectric loss.^{28,29} Generally speaking, various methods have been proposed to introduce porous structures, including chemical foaming,³⁰ freeze-drying,¹⁸ and supercritical carbon dioxide (scCO₂) foaming³¹ among others. For instance, Thomassin *et al.* used scCO₂ to foam polycaprolactone (PCL)/MWNTs composites, resulting in a shielding efficiency of (60–80) dB with low reflectivity at low MWNTs loading of 0.25 vol.%.³² In a different technique, Wang *et al.*³³ employed a new type of carbon tube with a whisker structure and excellent electrical conductivity. They used GO as a bridging and foaming agent to achieve a porous film via a vacuum-assisted filtration and annealing process. This yielded a remarkable near-field shielding performance of 25–50 dB in the frequency range of 1–9 GHz. This confirmed that the porous structures induced by GO during the thermal reduction process changed the polarization direction and distribution of the electric field of attenuated EM waves. However, graphene's excess dielectric constants limit its ability to achieve appropriate impedance matching. One approach to overcome this issue is to dope heteroatoms into graphene materials, thereby introducing significant structural defects and inducing extra polarization relaxation.³⁴ Dopant atoms, in fact, introduce charge into the electron system of sp²-bonded carbon materials.³⁵ Because of the presence of these dopant atoms, free charges become trapped by defects caused by doping under EM fields, leading to an asymmetric distribution

of charges around these defects. When exposed to an alternating EM field, these defects frequently act as polarization centers, resulting in polarization relaxation. Notably, the defective sites serve as permanent dipolar polarization centers, resulting in additional dielectric polarization.³⁶

It can be argued that heteroatom doping is a commonly used method to adjust electronic structure and other material characteristics. It serves three main functions: (1) it increases the number of polarizing centers, (2) it tunes the bandgap, and (3) it enhances dielectric properties.³⁷ Therefore, may assist in improving the performance of graphene for particular applications. Nitrogen (N) is often used as a doping element because it can strengthen EM attenuation capacity due to dielectric loss, lower electrical conductivity, enhance the magnetization performance,³⁸ and enhance the multiple scatterings as well as the balanced impedance match.³⁹ In fact, the size of the N atom is very similar to that of a carbon (C) atom, and it has five valence electrons, which can easily bond with C atoms.³⁷ Furthermore, the difference in electronegativity between the C atom (2.55) and the N atom (3.04) disrupts the electro-neutrality of graphene material and causes the change in atomic charge distribution and spin density. These changes may serve as the source of the magnetic property of graphene.⁴⁰ On the other hand, introducing sulfur (S) into a graphene sheet can improve its magnetic characteristics. As S has two additional electrons, incorporating these two unpaired electrons by each S atom into the conducting band — where they become delocalized among the S and C atoms, — can enhance the magnetic properties.⁴¹

As reported in the literature, the development of nitrogen-doped graphene composites for the purpose of microwave absorption enhancement was achieved very successfully.^{42,43} Besides, there are numerous studies related to the dual doping of graphitic materials with the S and the N which potentially possess promising properties for various applications.^{44–47} However, there remains a dearth of research on the effect of the N and the S dual-doping of graphene or GO and their composites on EMI shielding performance. Therefore, this work focuses on studying how heteroatom doping affects dielectric properties and EMI shielding effectiveness (SE) of N-co-S enrich reduced graphene oxide (rGO) filled polymer nanocomposites. Herein, we employed a simple functionalization technique for inserting heteroatoms into the surface of GO followed by a hot-pressing method to fabricate the shielding nanocomposite film. These techniques were chosen in an attempt to overcome challenges in manufacturing shielding materials, such as complicated preparation processes, high costs, and environmental harm.

2. Experimental Results

2.1. Materials

The graphite powder used in this study was supplied by Zen Chemicals, while the polyethersulfone (PES) was purchased

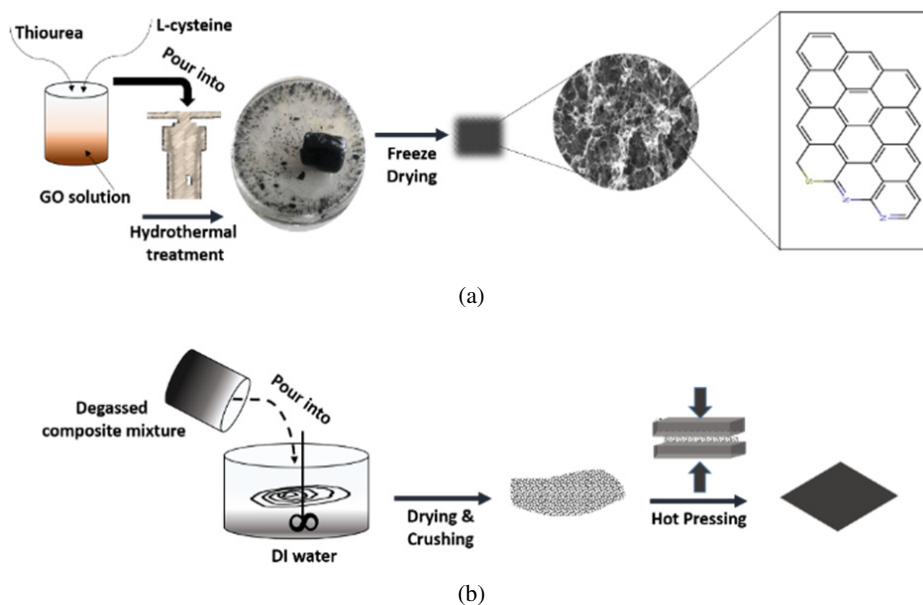


Fig. 1. Schematic illustration of (a) the synthesis procedure of the dual doping of rGO foams and (b) the fabrication of nanocomposite film.

from BASF, Ludwigshafen, Germany. Multiwall carbon nanotubes (MWCNTs) were provided by Sigma Aldrich. The l-cysteine (L-Cys) was purchased from Beijing Solarbio Science & Technology. Thiourea (with a purity greater than 98%), N, N-Dimethyleformamide (DMF), and sulfuric acid (H_2SO_4) with a purity of about 98% were supplied by R&M Chemicals. Sodium nitrate (NaNO_3) with a purity greater than 98%, was also obtained from the same supplier. Hydrogen peroxide (H_2O_2) was provided by Bendosen Laboratory Chemicals, while potassium permanganate (KMnO_4) with a purity greater than 99% was supplied by Chemiz (M) Sdn. Bhd.

2.2. Preparation of graphene oxide

GO was prepared by chemical exfoliation of graphite powder according to a modified Hummer's method. In a typical process, 2 g graphite powder was dispersed in 46 ml H_2SO_4 and 1 g NaNO_3 . The mixture was then placed into an ice bath ($0\text{--}5^\circ\text{C}$) under continuous stirring. 6 g KMnO_4 was added into the mixture portion-wise, and the reaction was kept for at least 2 h and then maintained below 15°C , (the color became green). The ice bath was removed, and the reaction was kept under stirring for 2–3 h at 35°C (the color altered to dark green). After that, 100 ml deionized (di)-water was added and the reaction was maintained at $90\text{--}98^\circ\text{C}$ (the color changed to yellowish brown), and the reaction was kept under this temperature for 30 min, then another 200 ml di-water was added portion wise under stirring. The temperature was decreased to 70°C and then 20 ml H_2O_2 (30 wt.%) was added into the solution. GO was obtained after centrifugation at 7000 rpm.

2.3. Synthesis of N, S dual-doped rGO foams

In a typical process as described in Fig. 1(a), 0.2 g GO was dispersed by ultra-sonication in 100 ml of deionized water for 2 h. In contrast, 0.30 g of a 1:1 mixture of L-Cys and thiourea was dispersed by ultra-sonication in 30 ml of the GO dispersion. The mixture was then transferred into an autoclave with a Teflon liner for hydrothermal treatment lasting 16 h at 170°C . The temperature was maintained at approximately 170°C for the duration of the treatment. Following this, the resulting product was thoroughly rinsed with di-water and then freeze-dried. The resulting nitrogen and sulfur-enriched rGO foam was denoted as NS-rGO. Oxidized CNT was prepared from neat CNTs in our lab as mentioned in our previous work.⁴⁶ Nitrogen and sulfur dual-doped oxidized CNT (NS-OCNT) was prepared under the same hydrothermal conditions.

2.4. Nanocomposite fabrication

Figure 1(b) illustrates the preparation procedure of the nanocomposite samples. Typically, PES and NS-rGO foams were dissolved separately in DMF by ultra-sonication before being mixed together to form the composite mixture. Then, the mixture was degassed for 30 min and slowly poured into di-water under mild stirring. It was then dried at 80°C for 48 h. The dried NS-rGO/PES nanocomposite was crushed into small pieces using a crushing machine. The fabrication of nanocomposite film was performed by a tablet pressing machine (XH-406B) under a pressure of 15 MPa for 10 min at 180°C . The resulting nanocomposite films, which had a thickness of 3 mm, were denoted as follows: GO@PES1 (Control sample, with 0.2 wt.% GO loading), NS-rGO@PES2

(with 0.2 wt.% NS-rGO loading), NS-rGO@PES3 (with 0.5 wt.% NS-rGO loading). For comparison, a nanocomposite with 0.5 wt.% NS-OCNT loading was also prepared and named NS-OCNT@PES4.

2.5. Measurements

The phase patterns of the doped samples were examined using an X-ray diffractometer (XRD) (D2 PHASER – Bruker, Germany) with Cu $K\alpha$ radiation, Cu anode, and a scanning rate of $10^\circ/\text{min}$. To ascertain the final chemical structure of the doped samples, Fourier-transform infrared spectroscopy (FT-IR, Nicolet iS50, USA) and Raman spectroscopy (Renishaw inVia, Germany) were utilized. The morphology and elemental analysis of the doped samples were studied using a scanning electron microscope (SEM, JSM-5600LV, Japan) coupled with an energy-dispersive X-ray spectroscopy (EDS). For quantitative elemental analysis, approximately 3 mg of the dried samples were subjected to a Thermo Scientific elemental analyzer (Flash smart CHNS, Italy), and the atomic percentages of nitrogen, carbon, hydrogen, and sulfur were determined. The microwave absorption measurements were performed using a vector network analyzer (VNA) (Agilent-N5227A, USA).

3. Results and Discussions

3.1. Physicochemical characterization of doped samples

The XRD Phase patterns for NS-rGO are depicted in Fig. 2. The patterns display a broad and wide peak, centered at 24.4° , spanning from 20° to 33° . This can be attributed to the slightly larger interlayer spacing of the (002) diffraction planes of amorphous carbons, which is likely the result of the nitrogen and sulfur doping. Moreover, a clear peak can be observed at $2\theta \approx 25.3^\circ$ in the XRD pattern of neat CNTs

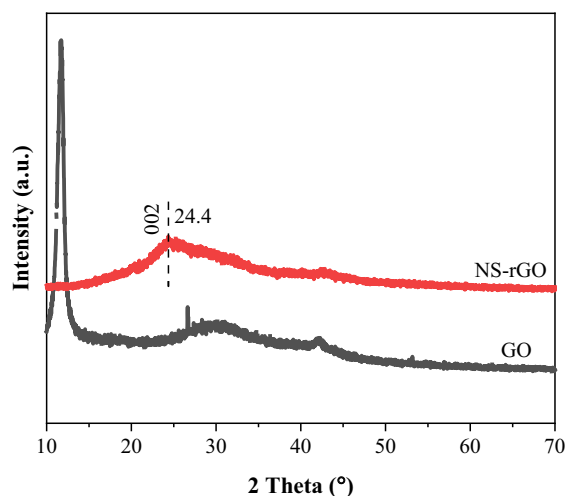
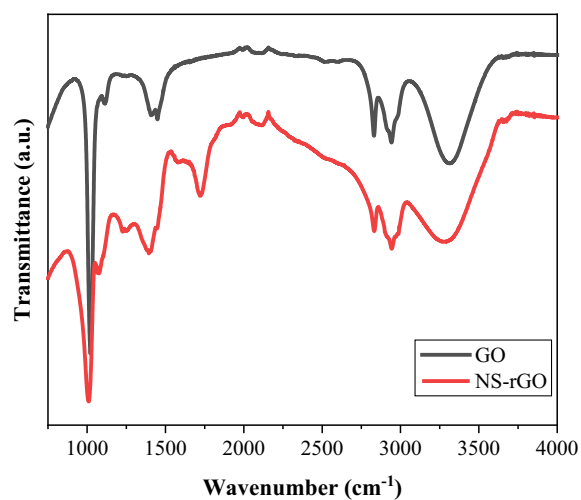


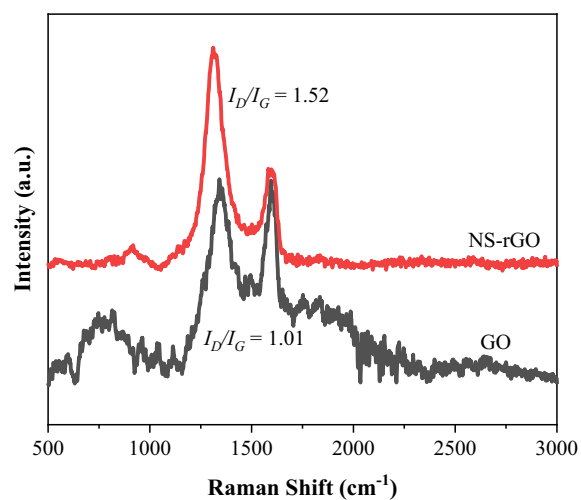
Fig. 2. XRD pattern for the sample of as-prepared GO and dual-doped rGO.

(refer to Fig. S1). However, the pattern of NS-OCNT shows a diminished intensity of the (002) peak compared to the neat CNTs. This could be attributed to the disruption of the graphite structure caused by the insertion of the doping elements, N, and S.⁴⁹

FT-IR spectra of the graphene oxide in Fig. 3(a), GO sample shows wide and strong peaks at about 3314 cm^{-1} corresponding to the stretching vibration of the O–H bond, due to the adsorbed H_2O molecules on GO.⁵⁰ Other absorption bands are noted between 2800 cm^{-1} and 3200 cm^{-1} representing the hydroxyl group in GO network.⁵¹ Another strong peak at around 1021 cm^{-1} is also observed corresponding to C–O–C (epoxy) groups.⁵¹ The intensities of these peaks reduced after the hydrothermal treatment demonstrated the doing of heteroatoms. Besides, a characteristic peak at 1448 cm^{-1} corresponding to the carboxyl group is



(a)



(b)

Fig. 3. (a) FTIR spectra for the sample of as-prepared GO and dual-doped rGO (b) Raman spectra for the sample of GO and dual-doped rGO.

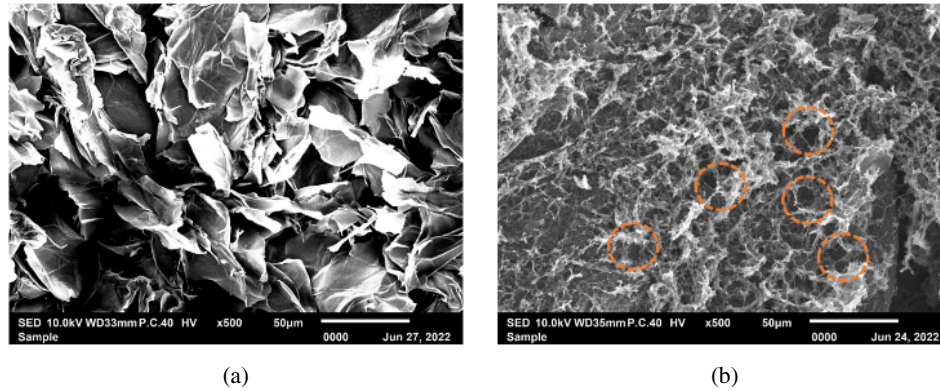


Fig. 4. SEM images of as-prepared (a) GO and (b) NS-rGO.

also observed. After reacting with L-cysteine and thiourea for 16 h, this peak disappeared or shifted to a lower wavenumber (~ 1394), indicating the insertion of N element.⁵² Two absorption peaks at 1576 cm^{-1} and 1720 cm^{-1} were noted on the NS-rGO sample representing the vibrations of $\text{C}=\text{C}$, $\text{C}=\text{N}$, and $\text{C}=\text{O}$.⁵³ More importantly, the insertion of the S element is confirmed based on the feature of an absorption peak around 1200 cm^{-1} which is ascribed to the stretching vibration of a $\text{C}-\text{S}$,⁵³ and peak at 1075 cm^{-1} that corresponded to stretching vibrations of $\text{S}=\text{O}$ bond in graphene sheet.⁵²

Raman spectroscopy provides additional insights into the sulfur-co-nitrogen functionalization of rGO. As shown in Fig. 3(b), the Raman spectra of both GO and NS-rGO exhibit two prominent peaks near 1340 cm^{-1} and 1580 cm^{-1} . These peaks correspond to the corresponding to the D band, which arises from structural defects, and the G band, originating from the sp^2 -bonded graphitic carbon atoms, respectively. Generally, a higher intensity of the D band compared to the G band signifies a higher degree of disorder within the carbon material. The sample GO shows a peak at approximately 2640 cm^{-1} that is related to the second-order two-phonon mode 2D-band and is caused by the disordered property of carbon material.⁵⁴ The NS-rGO sample eliminated this peak. The G band of NS-rGO is slightly displaced to higher wavenumbers (approximately 10 cm^{-1}) than that of GO, which can be attributed to the development of p-type doped graphene. This phenomenon has also been observed in the literature and may be caused by the displacement of dopants in graphene sheets.⁵⁵

Moreover, the ratios of the D to G peak intensities (I_D/I_G) are often used to estimate the level of defects in synthetic materials. For GO and NS-rGO, these ratios are observed to be 1.52 and 1.01, respectively, while the NS-OCNT exhibits an (I_D/I_G) ratio of 1.23 (refer to Fig. S2).

These findings suggest a higher number of structural defects in NS-rGO compared to both GO and NS-OCNT, which can be attributed to a high level of heteroatom doping.^{56,57} Overall, the results from both XRD and Raman spectroscopy collectively indicate that the proposed material

has been successfully doped with nitrogen and sulfur through a one-step hydrothermal process.

Figure 4 displays the morphological observation of GO samples before and after hydrothermal treatment. Figure 4(a) presents the morphology of pure GO, which is characteristically composed of stacked and interlaced ultrathin nanosheets.⁵⁸ In contrast, Fig. 4(b) depicts that NS-rGO foams composed of closely cross-linked graphene sheets that form a looser, porous 3D network.⁵⁴ This network is characterized by spherical holes, remnants of air bubbles, which pile on top of each other and are evenly distributed throughout the foam framework. It is suggested that this 3D porous network structure could enhance microwave absorption. The microporous framework inside the material can cause electromagnetic waves to repeatedly reflect off the walls in each individual closed void. Concurrently, the conductive network greatly enables the movement of electrons, leading to energy loss of the electromagnetic waves.²⁷ Actually, the conductivity of rGO increases from $12.65\ \mu\text{S}$ to $174.70\ \mu\text{S}$ in methanol medium and $25.40\ \mu\text{S}$ to $72.35\ \mu\text{S}$ in Distilled Water (DI) in our case (measured by standard conductivity meter probe) by incorporating dopant elements (see Table 1).

EDS mapping images, as displayed in Fig. S3, unambiguously demonstrate the presence of C, oxygen (O), and S elements. These elements are uniformly distributed across the field of view, which confirms the structural integrity of the 3D graphene network after treatment. These observations suggest successful sulfur functionalization of rGO.

Nitrogen, on the other hand, is not detected by SEM-EDX. This is likely due to the fact that nitrogen tends to have a very

Table 1. Electrical conductivity for the dual-doped sample.

Samples	Solvent	Conductivity (μS)
rGO	Methanol	12.65
	DI	25.40
NS-rGO	Methanol	174.70
	DI	72.35

Table 2. Elemental content for the dual-doped sample.

Sample name	Element %			
	Carbon	Hydrogen	Nitrogen	Sulfur
GO	44.69	3.09	0.09	1.06
NS-rGO	39.78	3.09	8.93	13.19

faint response, making its detection unreliable in most materials.⁵⁹ The results from CHNS analysis confirm the dual incorporation of S and N elements in rGO foams. As seen in Table 2, the NS-rGO sample exhibits a lower C content than the untreated GO sample, but it has higher N and S contents, which are 8.93% and 13.19%, respectively. This verifies successful heteroatom doping.

3.2. Electromagnetic characterization of polymer-based nanocomposites filled with NS-rGO foams

To study the effects of heteroatom doping, variations in the amount of the modified fillers, and the influence of pore structure on the microwave absorption properties and EMI shielding performance of the synthesized nanocomposites, two different loading amounts of NS-rGO in the PES matrix were used, namely, 0.2 wt.% and 0.5 wt.%.

The scattering parameters, S_{11} (reflection) and S_{21} (transmission) were measured across a frequency range of 8 to 12 GHz. The reflection coefficient (R) and transmission coefficient (T), as well as the EMI shielding effectiveness (SE), the absorption shielding effectiveness (SE_A), and the reflection shielding effectiveness (SE_R) of the nanocomposites can be computed from the scattering parameters (S_{11} and S_{21} , or S_{12} and S_{22}) using the following equations^{60,61}:

$$R = |S_{11}|^2 = |S_{22}|^2, \quad (1)$$

$$T = |S_{21}|^2 = |S_{12}|^2, \quad (2)$$

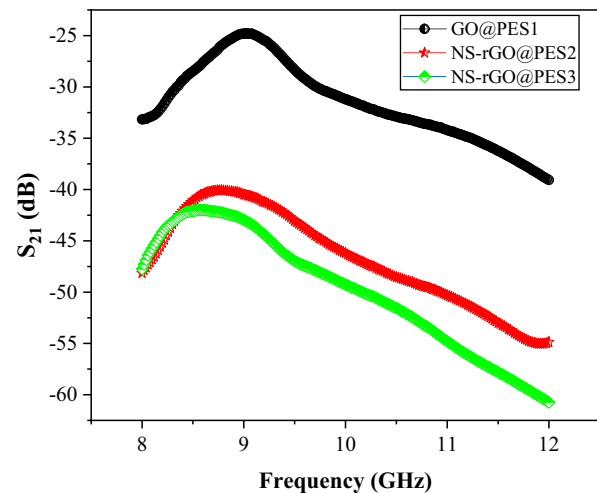
$$\begin{aligned} SE_R &= -10 \log(1 - R) \\ &= 10 \log\left(\frac{1}{1 - R}\right) \\ &= 10 \log\left(\frac{1}{1 - |S_{11}|^2}\right). \end{aligned} \quad (3)$$

$$\begin{aligned} SE_A &= -10 \log\left(\frac{T}{1 - R}\right) \\ &= 10 \log\left(\frac{1 - R}{T}\right) \\ &= 10 \log\left(\frac{1 - |S_{11}|^2}{|S_{21}|^2}\right). \end{aligned} \quad (4)$$

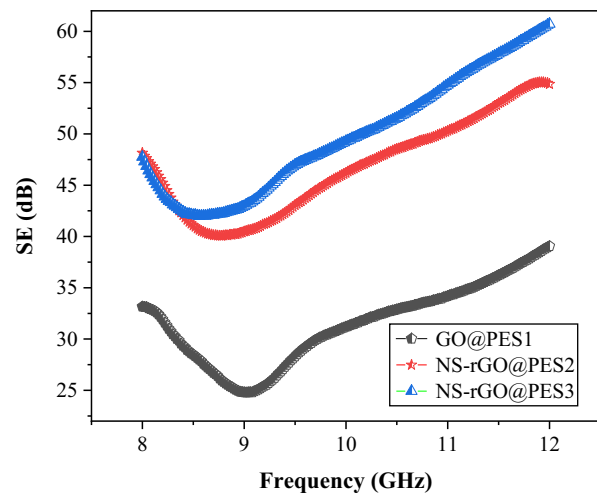
$$SE = SE_R + SE_A = 10 \log \frac{1}{T} = 10 \log \left(\frac{1}{|S_{21}|^2} \right). \quad (5)$$

Figure 5(a) illustrates the transmission spectra of nano-composite film. It is observed that compared to the control sample (GO@PES1), the doped samples exhibit a wider transmission window or bandpass feature at low frequencies. This finding suggests that the hydrothermal doping of GO significantly affects the electromagnetic response of the nanocomposites. Notably, lower transmission magnitude is exhibited by the NS-rGO@PES3 sample, suggesting its optimal doping capacity. Figure 5(b) shows that the total SE decreases at low frequencies and then increases after 9 GHz. This increase is typically associated with a polarization process occurring in the interfacial region due to the presence of a heterogeneous interface, characteristic of a one-phase heterogeneous structure of rGO fillers.⁶²

From the results, the SE values for GO@PES1, NS-rGO@PES2, and NS-rGO@PES3 were found to be 38 dB, 54 dB, and 60 dB, respectively, at a thickness of 3 mm.



(a)



(b)

Fig. 5. (a) Transmission spectra S_{21} and (b) shielding effectiveness (SE) for the nanocomposite films.

In contrast, a nanocomposite fabricated from NS-OCNT with 0.5% loading exhibited an SE of about 45 dB (refer to Fig. S3). Remarkably, the NS-rGO@PES3 sample, which has an rGO loading amount of 0.5 wt.% demonstrates the optimal SE that is 25% higher than that of the NS-OCNT@PES4 nanocomposite with a similar loading amount. This suggests that the enhancement of the SE with optimal loading is attributable to the synergistic effects of S and N dual-doping and the reduction of GO in the hydrothermal process. Moreover, the pores present in the porous foams could enable multi-level reflections of the EM wave within the material, thus resulting in improved EMI shielding performance.⁶³ The electromagnetic shielding performance of a nanocomposite film (is generally accomplished via microwave absorption and/or reflection. For reflection-based shielding to occur, the material must contain movable charge carriers, such as electrons or holes, which are capable of interacting with the incident electromagnetic waves. Materials with high permeability can enhance absorption-based shielding.⁶²

Interfacial polarization, stemming from disparities in permittivity, conductivity, and relaxation time of charge carriers across the interface, can result in significant SE from both absorption and reflections.⁶⁰ Indeed, the shielding performance achieved through absorption is intensified due to the nitrogen/sulfur doping of rGO. This doping process modulates the electrical and magnetic characteristics of the material, thus enhancing its shielding capabilities.⁶² Moreover, the reduction that occurs during the treatment also plays a significant role in the high absorption performance of the nanocomposite. Reduced GO, produced during the hydrothermal process, is a superior option for microwave absorption compared to graphene and GO.³⁸

The frequency-dependent electromagnetic parameters of the nanocomposite film samples are displayed in Fig. 6.

There was only a slight change noted in the frequency range of 8.5–11 GHz. The optimum value of real permittivity (ϵ') was achieved by NS-rGO@PES3, which exhibits a notable peak value of 2.3 at around 9.12 GHz. The increase in the dielectric constant can be attributed to influences such as the Maxwell–Wagner–Sillars (MWS) polarization or interfacial polarization.⁶⁴ According to Debye's theory, the permittivity of free space (ϵ_0) and imaginary permittivity (ϵ'') follows the equation below⁶⁵:

$$\epsilon'' = \epsilon_p'' + \epsilon_c'' = (\epsilon_s - \epsilon_\infty) \frac{2\pi f \tau}{1 + (2\pi f)^2} + \frac{\sigma}{2\pi f \epsilon_0}. \quad (6)$$

Herein, ϵ_p'' and ϵ_c'' are the polarization relaxation and conductive loss. ϵ_s , ϵ_∞ , f , τ , and σ represent the static permittivity, relative permittivity at the high-frequency limit, frequency, relaxation time, and electrical conductivity, respectively. As per the above equation, the ϵ'' is closely related to the ϵ_p'' and ϵ_c'' of nanocomposite samples.

From Eq. (6), the ϵ_c'' should decrease with increasing frequency. However, as shown in Fig. 6(a), an increase in the value of ϵ'' can be noted in the frequency range of 9–12 GHz. This increase could be attributed to the strong polarization relaxation in the resulting nanocomposites.⁶⁶ On the basis of the free electron theory, it can be deduced that the higher the electrical conductivity, the higher the ϵ'' specimen will have.⁶⁷

Therefore, the NS-rGO@PES3 displays slightly larger ϵ' and ϵ'' , indicating the improved storage capability of electric energy and dielectric loss.⁶⁷ Besides, negative values of ϵ'' were observed at low frequencies, while altered to positive for both NS-rGO@PES2 and NS-rGO@PES3 samples when the frequency reached 11.70 GHz. The phenomenon of positive ϵ' and negative ϵ'' was also reported in the literature.⁶⁸ Generally, the term "metacomposites" refers to nanocomposites with the

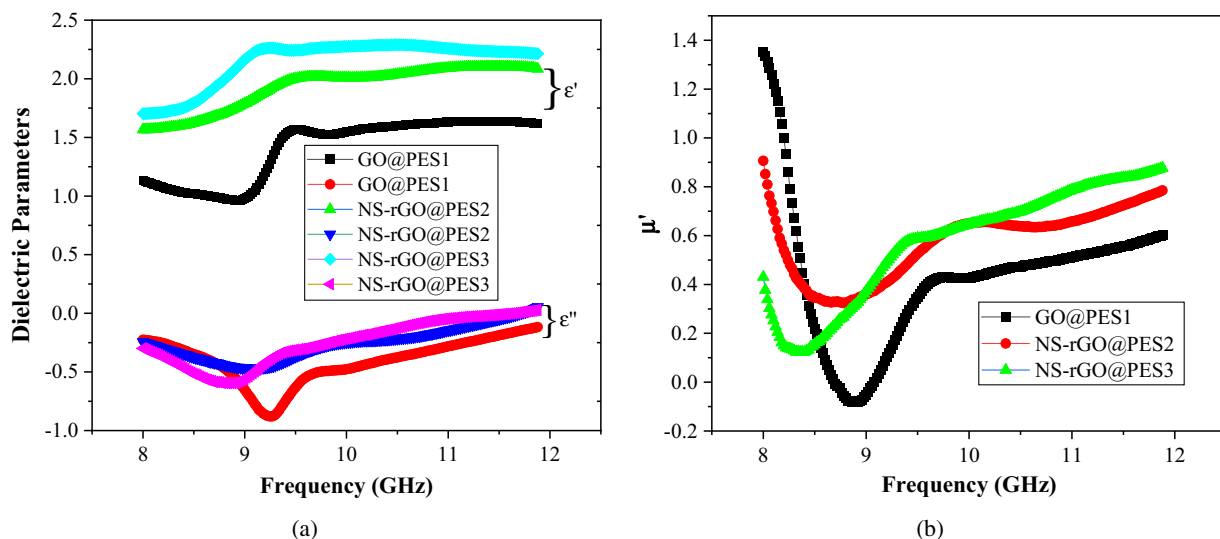


Fig. 6. Frequency dispersion characteristics of (a) real permittivity and imaginary permittivity and (b) real part of permeability of the nanocomposite films.

unique characteristic of negative permittivity. This property depends on the loading, the frequency, and the nanofillers' morphology, and can be adjusted by altering these factors. Metacomposites have a wide range of potential applications, such as super lenses, wave filters, remote aerospace applications, and superconductors. In addition, the negative permittivity in the graphene nanocomposites could be assigned to the unique electronic energy dispersions (also known as surface plasmons). The positive permittivity may be due to the low plasmon resonance conductivity at a relatively high frequency.⁶⁹ This finding suggests that NS-rGO@PES2 and NS-rGO@PES3 samples have better conductivity than GO@PES1, because of the removal of most oxygen-containing groups, demonstrating the major influence of doping treatment on the EM shielding efficiency.³⁹ Thus, the conductivity of doped GO has a strong influence on its dielectric properties, which is consistent with the results presented in Table 1.

As shown in Fig. 6(b), all samples exhibit a slight decreasing trend in the real part of permeability (μ') at low frequency, with notable peaks observed between 8 and 9.50 GHz.

Attenuation loss, which includes dielectric loss and magnetic loss, is critical for microwave absorption. The dielectric loss tangent ($\tan\delta_\epsilon = \epsilon''/\epsilon'$) and magnetic loss tangent ($\tan\delta_\mu = \mu''/\mu'$) can be utilized to estimate the dielectric loss and magnetic loss capacity of microwave absorbers. As shown in Fig. 7, the NS-rGO@PES2 and NS-rGO@PES3 samples demonstrate better $\tan\delta_\epsilon$ compared to NS-GO@PES1, suggesting their enhanced dielectric loss against the incident microwaves. This enhancement can be linked with the occurrence of interfacial polarization, as a small amount of electron tunneling can occur.⁷⁰ In fact, heteroatoms and other sort of defects in graphene sheets like the remaining oxygen-containing groups as well as restacking of graphene layers that are likely to occur during the reduction process can serve as a polarization center to elevate the material loss, hence dissipate microwave energy.³⁹

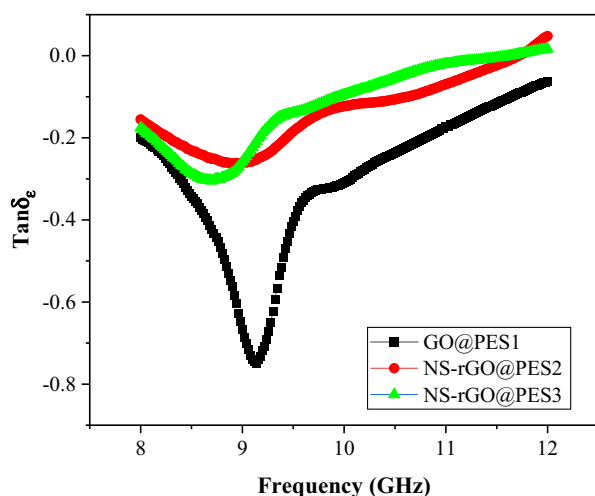


Fig. 7. Frequency dispersion characteristics of dielectric loss tangent of the nanocomposite films.

4. Conclusion

The nitrogen-co-sulfur-doped reduced graphene-oxide (NS-rGO) films with 3D porous structures were synthesized through a hydrothermal method, employing l-cysteine and thiourea as doping precursors. Subsequently, nanocomposite films were fabricated by incorporating the NS-rGO foams into polyethersulfone (PES) through the hot-pressing technique. The examination of elemental content confirmed the successful insertion of sulfur and nitrogen functionalities into rGO. The microwave characterization of the resulting shielding film demonstrated excellent electromagnetic interference shielding performance, which can be attributed to the existence of the porous structure and the heteroatom insertion. The highest shielding effectiveness was achieved at the loading of 0.5 wt.% of NS-rGO. Overall, this work provides guidance for the functional composites and the engineering of pore structures in carbon nanomaterials. The proposed material exhibits great potential for applications in microwave absorption. This type of microwave absorption material represents a favorable solution for environmental protection in the field of EMI shielding.







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Supplementary Material

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