Preparation of a Chemically Reduced Graphene Oxide Reinforced Epoxy Resin Polymer as a Composite for Electromagnetic Interference Shielding and Microwave-Absorbing Applications

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Abstract: The preparation of chemically reduced graphene oxide (rGO) and the optimization of epoxy resins’ properties using micro or nanofillers are now common practices. rGO nanoparticles (60 nm) based on an epoxy resin polymer were prepared at the concentrations of 0, 1, 2, 3, 4, and 5% weight percentage with fixed 6-mm thicknesses. The dielectric properties of the composites were measured by the reflection/transmission technique in connection with a vector network analyser (VNA) at a frequency range of 8–12 GHz. The microwave absorption and shielding effectiveness properties were calculated by using the reflection $S_{11}$ and transmission $S_{21}$ results. The microstructure and morphology of the polymer and the rGO/cured epoxy composites were studied by field emission scanning electron microscopy (FE-SEM), Fourier-transform infrared (FT-IR) spectroscopy, and the X-ray Diffraction (X-RD) technique for characterizing crystalline materials. The dielectric and other properties of the rGO/cured epoxy composites were investigated based on the filler load and frequency. It was found that the applied frequency and the filler concentrations affected the dielectric properties of the rGO/cured epoxy composites. The results showed that the introduction of rGO particles to the composites increased their dielectric properties smoothly. The study of the dependence on frequency of both the dielectric constant $\varepsilon'$ and the dielectric loss $\varepsilon''$ showed a decrease in both quantities with increasing frequency, indicating a normal behaviour of the dielectrics. Cole–Cole plots were drawn with $\varepsilon'$ and $\varepsilon''$. A theoretical simulation in terms of the Cole–Cole dispersion law indicates that the Debye relaxation processes in the rGO/cured epoxy composites are improved due to the presence of the rGO filler. Moreover, with the addition of rGO as a filler into the Epoxy matrix, it now exhibits promise as a lightweight material for microwave absorption as well as an effective electromagnetic interference (EMI) shielding material.

Keywords: reduced graphene oxide; epoxy resins; permittivity; permeability; Cole–Cole

1. Introduction

Electromagnetic interference (EMI) is one of the most undesirable by-products of telecommunication devices and high-frequency electronic systems. Any device that utilizes, processes,
transmits, or distributes electrical energy of any form may emit signals of an electromagnetic nature and interfere with nearby systems and equipment in their normal operation [1]. Such signals may also have negative effects on human health. Efforts to reduce pollution of an electromagnetic nature have been made with the use of EMI shielding materials [2]. Signals are attenuated by such materials through absorption and/or reflection of the radiation power [3]. Traditionally, magnetic materials and metals have been utilized for the shielding of EMI due to their good mechanical properties and high effectiveness in shielding. However, among the drawbacks that these materials have, they are heavy, susceptible to corrosion, and difficult to process. The metals’ high conductivity also limits their application as absorbers of electromagnetic waves due to their shallow skin depth; thus, almost all of the power is reflected on the surface [4]. There is an increasing need to develop practical and effective EMI shielding materials that overcome the conventional metal-based shields’ shortcomings, which has increased the interest of researchers in these novel materials [5].

Polymer composites for applications in EMI shielding have attracted the attention of many researchers. This is mainly because of their characteristics, which usually include low density, good mechanical properties, high dimensional and thermal stability, and also special and less common features, such as electromagnetic absorption [6]. Also, the disadvantages that are associated with the use of metals, including corrosion resistance, light weight, flexibility, as well as processing difficulty, are addressed by these materials [7]. Carbon black [8], Carbon-based particles, such as carbon fibre [9], and carbon nanotubes [10], have been investigated as effective fillers for preparing absorbing materials and conducting polymer composites for EMI shielding applications, military affairs, commerce, and electronic instruments in industry. Most of the materials that are used for microwave absorption are composed of powders of magnetic loss, which include nickel, ferrite, and cobalt, and materials for dielectric loss, such as conducting polymers and carbon nanotubes. Although the carbon nanotubes’ ability for microwave absorption is extremely weak, previous reports have indicated that magnetic material and carbon nanotube composites would exhibit excellent properties for microwave absorption [11].

Recently, graphene’s microwave absorption and electromagnetic shielding abilities have been investigated. Various materials with graphene particles have therefore been highlighted [12] and have shown interesting electromagnetic absorption outcomes. For instance, the improved electromagnetic wave absorption results of chemically reduced graphene oxide (rGO) with residual defects via additional relaxation processes, namely dielectric, polarization, and dipole relaxations, showed that a 6.9 dB microwave reflection loss had been achieved at 7 GHz [12]. Graphene-poly methyl methacrylate nanocomposite microcellular foams were found to have high EMI shielding effectiveness (13–19 dB at 8–12 GHz) by the support of multi-reflections and the scattering of the incident microwaves into the foam samples [13]. Graphene layers with polyaniline nanorods embedded in a paraffin matrix exhibited a microwave reflection loss that was lower than 20 dB from 7.0 to 17.6 GHz when the Debye relaxation process was amended [14]. Graphene nanoplatelets in epoxy resin exhibited a 14.5 dB maximum reflection loss at 18.9 GHz, which was mainly attributed to the charge multipoles at the polarized interfaces into the composite material [15]. Also, graphene was incorporated into an epoxy matrix to study EMI shielding phenomena at the (X-band) frequency, and a 21 dB shielding efficiency was achieved for a 15 wt % (8.8 vol %) loading, indicating that it may be used as an effective and light weight EMI shielding material [16].

Furthermore, reduced graphene oxide has been incorporated into epoxy resin. The results showed shielding effectiveness (SE) values that were higher than the target value (20 dB). As it turned out, at a lower filler loading, and if suitably modified, the surface of the graphene preserves its conductivity, and an EMI SE for practical applications can be achieved [17]. On the other hand, there is a lack of proper research about rGO’s electromagnetic wave absorbing property. rGO shows an improvement in microwave absorption when compared to carbon nanotubes and graphite, is expected to present superior absorption to high-quality graphene, and exhibits promise as microwave absorbing material [18]. Nevertheless, there are some studies that have investigated, and contributed to the
knowledge of, the effect of other factors, such as the particle size of the graphene oxide, on the
electromagnetic properties, where the properties can be tuned by controlling the size of the graphene
particles. The more the particle size increases, the more the dielectric constant increases. The reason
for this might be associated with the discrepancy in the interfacial polarization effect that results from
different particle sizes [19].

Organic polymer materials, such as epoxy resins, have excellent adhesion to different materials,
high resistance to chemical attacks, and excellent mechanical and electrical properties. Besides this,
the hardener HY951 provides the best binding property with an epoxy resin [20]. In addition to
the abovementioned properties of epoxy resins, they are used in this research due to their diverse
characteristics, such as high strength, good stiffness, good thermal stability, antibacterial properties,
low contractibility, and strong adherence chemical resistance. They are used widely in industrial
applications, such as coatings, adhesives, aerospace structures, electronics, potting, composites,
laminates, and the encapsulation of semiconductor devices. Because of their excellent and attractive
mechanical and chemical properties, epoxies are the dominant matrix material for structural composites
of light-weight polymer–matrix [21]. There have been so many reports on the blending of epoxies
with fillers in recent years [22], including the incorporation of nanoferrites, nanoferroelectrics, carbon
nanoparticles, single-walled carbon nanotubes (SWNTs), and multi-walled carbon nanotube (MWNTs)
into epoxy resins as an absorbing material for EM waves [23].

The present study has two main goals: (a) to prepare chemically reduced graphene oxide (rGO)
due to its scalability and high yield, and (b) to prepare rGO/cured epoxy composites and then
investigate their structure, morphology, and dielectric properties. The study of the composites’ physical
and chemical properties were performed through Fourier-transform infrared spectroscopy (FT-IR)
and X-ray diffraction (XRD). Field-emission scanning electron microscopy (FESEM) was conducted to
study the structure of the composite.

2. Experimental Details

2.1. Materials

The materials used in this work are: a polymer matrix (epoxy resin LY1316) and hardener (HY1208,
supplied by Buehler, Lake Bluff, IL, USA), Graphite Oxide (prepared by using a modified Staudenmaier
method), and reduced Graphite Oxide (manufactured by utilizing a weak base, namely Ammonia
(NH$_3$) as the reducing agent). The NH$_3$ was supplied by Sigma Aldrich (Sarasota, FL, USA).

2.2. Preparation of Reduced Graphene Oxide (rGO)

The preparation method for the synthesis and chemical reduction of rGO paper and then powder
involved two major steps. The first step is the synthesis of Graphite Oxide (GO) using the Staudenmaier
Method. To perform the reduction process of GO to rGO, about 400 mg of the obtained GO was placed
in a cellulose extraction thimble (30 by 100 mm) and then placed in the Soxhlet extraction unit.
Approximately 150 mL of 30% Ammonia solution (NH$_3$) was used as a reducing agent. The heating
temperature was set at 90 °C, and the exposure period was investigated (namely 5 h) when the GO
powder had direct contact with the ammonia vapour as well as the condensed liquid. The experimental
setup and measurement is as shown in Figure 1.
2.3. Preparation of rGO/Cured Epoxy Composites

The composites were manufactured in this work using three kinds of materials: epoxy resin (Er), hardener (H), and reduced graphene oxide (rGO), for different filler and polymer percentages which is clearly presented in Table 1. Er, H, and rGO were mixed using a mini-mechanical vortex mixer for 15 min to homogenize the resulting materials. The mixture was poured into rectangular aluminum molds with a thickness of 6 mm, and the coating was allowed to cure in air for 48 h or by utilizing an oven at a temperature between 80 and 140 °C. Figure 2a,b shows the specimens’ preparation process and the measurement of the electromagnetic properties.

Table 1. The compositions of the nanocomposites.

| Sample        | wt % rGO | wt % Cured Er | Mass (gm) rGO | Mass (gm) Cured Epoxy | Mass (gm) Er | Mass (gm) H | Mass (gm) rGO/Cured Epoxy |
|---------------|----------|---------------|---------------|-----------------------|--------------|-------------|--------------------------|
| rGO + cured epoxy 1 | 99 | 0.05 | 4.95 | 4.455 | 0.495 | | |
| 2 | 98 | 0.1 | 4.9 | 4.41 | 0.49 | | |
| 3 | 97 | 0.15 | 4.85 | 4.365 | 0.485 | | |
| 4 | 96 | 0.2 | 4.8 | 4.32 | 0.48 | | |
| 5 | 95 | 0.25 | 4.75 | 4.275 | 0.475 | | 5 gm |

Er, epoxy resin; H, hardener.
2.4. Characterisation

All of the characterizations of the pure material under study (rGO, epoxy resin, hardener, and the cured epoxy composites) at different percentages of filler were carried out as follows.

2.4.1. X-ray Diffraction (XRD)

The analysis of the samples’ phase structure was performed using X-ray diffraction (XRD, XD-3, Cu Ka radiation) under ambient conditions with a Lynx Eye detector using a Bruker diffractometer (Yuseong, Daejeon, Korea) over a 5–90° 2θ range. Cu-Kα radiation with a wavelength of 1.54 Å (nickel filtered) was the X-ray beam and operated at 40 kV generator voltage and 35 mA current value settings. The rGO samples were in the form of a fine powder, while the rGO/cured epoxy composite samples were cut from the specimens that were prepared in advance in a solid form.

2.4.2. Fourier-Transform Infrared (FT-IR) Spectroscopy

In order to obtain appropriate information on the functional groups that were present in the modified reduced graphene oxide, epoxy/hardener, and rGO/cured epoxy composites, an FT-IR
analysis was conducted. The spectra were measured by mixing about 0.05–0.1 wt % of the fine ground modified graphite oxide samples with KBr powder, which was then compressed to form a pellet. Using the micro die method, the FTIR spectra of the pellet were measured using a Perkin-Elmer spectrum (Waltham, MA, USA) 100 at 400–4000 cm\(^{-1}\).

2.4.3. Field Emission Scanning Electron Microscopy (FE-SEM)

The morphology of the prepared rGO/cured epoxy composite samples was studied and observed by field-emission scanning electron microscopy (FE-SEM, JEOL JSM-7001F) with a field emission cathode backscatter detector and a resolution of 1 nm at 30 kV up to a 700,000 \( \times \) magnification and using an Accelerator voltage between 0.1 and 30 kV. All composite samples were coated with gold before analysis.

2.4.4. Electrical Properties

The s-scattering parameters of the reflection (\( S_{11} \)) and transmission (\( S_{21} \)) coefficients were measured, and the dielectric properties of the rGO/cured epoxy composites at room temperature in the frequency range of 8–12 GHz were investigated by utilizing transmission line techniques. A rectangular specimen were inserted into a 22.86 \( \times \) 10.14 \( \times \) 6 mm\(^3\) aluminum sample holder that was connected between the waveguide flanges of an Agilent E8362B network analyser. A full two-port calibration was performed along with the sample holder to default any loss and power redistribution due to the sample holder. All of the composite samples were tested, and, for each sample, 201 data points were taken within the specified frequency range.

2.4.5. EMI Shielding Property

EMI Shielding Mechanisms

The electromagnetic interference (EMI) shielding was calculated by utilizing the S-parameter results [24]. Electromagnetic interference (EMI) shielding is defined as the attenuation of electromagnetic radiation by reflection and/or absorption of the incident power. The incident electromagnetic radiation on a shield can be resolved into three parts; namely, transmittance (T), absorption (A), and reflection (R), with the sum (T + A + R) equal to 1. Hence, the \( SE_{\text{total}} \) is the sum of contributions from the absorption loss (\( SE_A \)), the reflection loss (\( SE_R \)), and multiple reflections (\( SE_M \)), i.e.,

\[
SE_{\text{total}} = SE_A + SE_R + SE_M. \tag{1}
\]

The \( SE_{\text{total}} \) of a shielding material can be written as

\[
SE_{\text{total}} = 10 \log_{10} (1/S_{21}^2) \tag{2}
\]

where \( S_{21}^2 \) is the transmittance value (T), which can be measured from \(-\left(P_i/P_t\right)\) [25]. The \( P_i \) and \( P_t \) are, respectively, the incident and transmitted powers, considering the effective absorbance (\( A_{\text{eff}} \)), which is defined as

\[
A_{\text{eff}} = (1 - S_{11}^2 - S_{21}^2)/(1 - S_{11}^2). \tag{3}
\]

With regard to the power of the incident electromagnetic wave inside the shielding material, the \( SE_{\text{total}} \) can be rewritten and described as the sum of the two terms of effective absorbance and reflectance:

\[
SE_{\text{total}} = 10 \log_{10} (1/(1 - S_{11}^2)) + 10 \log_{10} ((1 - S_{11}^2)/S_{21}^2) = SE_R + SE_A. \tag{4}
\]

Using these equations, the total \( SE \) was resolved into absorption and reflection loss.
3. Results and Discussion

3.1. Powder X-ray Diffraction

X-ray diffraction measurements were utilized to investigate the phase composition and the crystalline structure of the synthesized samples. Figure 3 illustrates the X-ray diffraction pattern of the cured epoxy, the rGO, and the rGO/cured epoxy composites at the filler weight percentages of 1%, 3%, and 5%. The XRD spectra of the neat epoxy showed the appearance of a wide diffraction when compared with the 1% and 3% rGO/cured epoxy composites.

The curve indicates a series of diffraction peaks at 2θ corresponding to the (002), (100), (101), (102), and (004) planes, respectively, of the hexagonal carbon structure and crystal planes (JCPDS no. 19-0629) [27]. A crystalline behaviour of rGO having the highest peak appeared at an angle of 44.87°, which indicates a high degree of crystallinity. The degree of crystallinity is the most important basic parameter for characterizing crystalline polymers. On the other hand, Figure 3 shows the diffraction pattern of the rGO/cured epoxy composite containing 1%, 3%, and 5% filler by weight of resin. X-ray diffraction revealed that the composite was crystalline, as the highest peaks were observed at the angles of 72.63°, 44.71°, and 44.67°. Another interesting observation from Figure 3 was the intensity beside the obvious broad diffraction peak at 2θ = 18.18. Because of the amorphous state of the cured epoxy structure in the composite, the intensity of this diffraction peak is relatively weak. The change of the characteristic peak for rGO particles in the rGO/cured epoxy composites can be correlated to fully exfoliate the rGO particles in the polymer matrix. The composite shows a varying diffraction pattern for each rGO percentage, which could be due to the homogeneous dispersion and complete exfoliation of rGO in the cured epoxy matrix. It is clear that the diffraction peak of the 5% rGO/cured epoxy composite at 2θ = 73.02° became less sharp when compared with the 1% and 3% rGO/cured epoxy composites.

![Figure 3. The XRD patterns of cured epoxy, rGO, and rGO/cured epoxy at different percentages of rGO.](image)

The average crystallite size of the rGO nanoparticles (≈ 60 nm) was calculated by the XRD line broadening using the Scherer equation:

\[
D = \frac{k\lambda}{\beta \cos \theta}
\]
where $D$ is the average crystallite size in nm, $k$ is the shape factor (normally 0.9 for cubic shapes), $\lambda$ is the wavelength of the X-ray, $\theta$ is Bragg’s diffraction angle, and $\beta$ is the broadening of the diffraction line measured at full-width half-maximum intensity in radians (FWHM data converted to radians).

3.2. Field Emission-Scanning Electron Microscopy

The results of Field Emission Scanning electron microscopy (FE-SEM) (structural and morphological characterization) for the rGO as well as the rGO/cured epoxy composites at different rGO loadings (1%, 3%, and 5% rGO) are shown in Figure 4a–d with 25,000–50,000× magnification. It can be seen from Figure 4a that the pure rGO has a wrinkled, irregular, folding, and many-folded layered structure with a lateral size of several micrometres [28]. The layered structure was formed by rGO particles and it is interesting to see that the layers are continually cross-linked in a flaky textured form without any amorphous or other kinds of crystallized phase particles, as described by [29]. To understand the dispersion of rGO in the epoxy matrix, the FE-SEM image of a rGO/cured epoxy composite is shown in Figure 4b–d. Figure 4b shows the homogeneity of the rGO particles’ dispersion in the cured epoxy resin for the 1% rGO to 99% cured epoxy composite. The rGO particles are observed as small particles that are scattered throughout the polymer matrix of the composite, as well as fully incorporated within the epoxy resin matrix embedded in the polymer matrix. On the other hand, after increasing the percentage of rGO (3% and 5%) in the epoxy composite, the dispersion of rGO powder in the matrix in Figure 4c,d showed obvious differences. Figure 4c shows that the rGO powder was dispersed in the epoxy resin as merged particles of a large size. Figure 4d shows that the rGO powder was dispersed in the epoxy resin in the form of agglomerates. These phenomena are quite different from the rGO dispersion at a low percentage of rGO shown in Figure 4b, suggesting the re-agglomeration of rGO during its addition to the epoxy resin. A similar phenomenon was also found in carbon nanotube (CNT)/epoxy composites [30]. The morphology figures indicate that rGO particles have indeed reacted with the cured epoxy to produce rGO/cured epoxy composites.

3.3. The Morphology of the rGO and rGO/Cured Epoxy Composites

Fourier-transform infrared (FT-IR) spectroscopy was used to determine the nature of the functional groups present in the surface of the prepared rGO powder, the Epoxy, the Hardener, and the rGO/cured epoxy composites at different percentages of rGO filler. FT-IR spectra were recorded on a Bruker Vertex 70 spectrometer (Waltham, MA, USA) at room temperature (27 °C) over a frequency range of 400–4000 cm$^{-1}$. Figure 5 presents the typical FT-IR spectra of the rGO powder, the Epoxy, the Hardener, and the rGO/cured epoxy composites. Figure 5 shows absorption bands corresponding to C–O stretching at 1039.46 cm$^{-1}$, C–OH stretching at 1388.03 cm$^{-1}$, phenolic O–H deformation vibrations at 1494.24 cm$^{-1}$, C=C stretching at 1502.38 cm$^{-1}$, C–O–C stretching at 1228.66 cm$^{-1}$, and C–O stretching of ethers at 1040.64 cm$^{-1}$, C–O–C stretching of the oxirane group at 820.33 cm$^{-1}$, and rocking CH$_2$ at 555.46 cm$^{-1}$ [34].

On the other hand, the prepared rGO/Epoxy hardener composites at different percentages of rGO filler compound were confirmed by the identification of characteristic absorption peaks. The IR spectrum of the rGO/cured epoxy composites at 5 wt% displays strong absorption bands corresponding to O–H stretching vibrations at 3451.57 cm$^{-1}$, C–H of CH$_2$ stretching at 2948.03 cm$^{-1}$, aromatic C–C stretching at 1502.38 cm$^{-1}$, C–O–C stretching at 1235.15 cm$^{-1}$, C–O–C stretching at 1228.66 cm$^{-1}$, C–O–C stretching of ethers at 1040.64 cm$^{-1}$, C–O–C stretching of the oxirane group at 820.33 cm$^{-1}$, and rocking CH$_2$ at 555.46 cm$^{-1}$. A shifting of peaks is observed in the IR spectra when their blends have a strong interaction, such as hydrogen bonding or any another bonding.
The morphology of the rGO and rGO/cured epoxy composites was investigated. The FE-SEM micrographs in Figure 4 show the dispersion of rGO particles in the epoxy matrix. As the percentage of rGO increases from 1% to 5%, the rGO particles become more dispersed and embedded within the polymer matrix. Figure 4a-d presents the micrographs at different rGO loadings, from 1% to 5%, respectively.

FT-IR spectroscopy was used to characterize the functional groups present in the rGO and the composites. Figure 5 displays the FT-IR spectra of the neat Epoxy, the Hardener, and the rGO/cured epoxy composites at different rGO loadings (1%, 3%, and 5%). The spectra exhibit characteristic bands corresponding to various functional groups, such as C=O stretching at 1717 cm⁻¹, C–O–C stretching at 3451.57 cm⁻¹, and O–H stretching at 3389.86 cm⁻¹.

The electromagnetic properties of the composites were also investigated. Figure 5 illustrates the absorption bands at different wavelengths, which provide insights into the material's interaction with electromagnetic fields. These properties are crucial for applications requiring electromagnetic shielding or absorption.

In summary, the study highlights the dispersion and morphological changes in rGO/cured epoxy composites at varying rGO loadings. The results are supported by microscopic and spectroscopic analyses, offering a comprehensive understanding of the material's properties.
3.4. Electromagnetic Properties

The electromagnetic properties of a material, namely the magnetic permeability ($\mu$) and the electrical permittivity ($\varepsilon$), define the material’s response to electromagnetic waves. Permeability ($\mu = \mu' - j\mu''$) and permittivity ($\varepsilon = \varepsilon' - j\varepsilon''$) are complex numbers and a determinant on the nature of the material’s interaction with the magnetic and electrical fields of the wave, respectively. These interactions have two parts, namely: the storage and dissipation of energy parts. The storage of energy is due to the lossless energy exchange between the material and the real parts of the field. The dissipation of energy happens when material’s absorbed electromagnetic energy is converted to imaginary heat parts [35,36]. The loss tangent ($\tan \theta$) that is commonly used to describe dielectric losses is calculated by using $\tan \theta = \frac{\varepsilon''}{\varepsilon'}$.

3.4.1. The Dielectric Properties of the rGO/Cured Epoxy Composites

In order to investigate the intrinsic reasons for the EMI shielding effectiveness of the composites, the dielectric properties of the epoxy resin, the hardener, the cured epoxy, and the rGO/cured epoxy composites at different percentages of rGO loading at the frequency range of 8–12 GHz (X-band frequency) were measured. Figure 6 illustrates the results of the dielectric properties for the epoxy, the hardener, and the cured epoxy (90% epoxy: 10% hardener) in the frequency range of 8–12 GHz. It can be observed that the $\varepsilon'$ and $\varepsilon''$ decreased for all samples as the frequency increased. Figure 6a,b shows that the hardener has the highest values of $\varepsilon'$ and $\varepsilon''$ of 3.94 and 1.48 at 8 GHz, then it progressively decreased to 3.63 and 1.39 at 12 GHz, respectively. Then, the $\varepsilon'$ and $\varepsilon''$ values of the epoxy decreased from 3.67 and 1.36 at 8 GHz to 3.42 and 0.98 at 12 GHz, respectively. The values of $\varepsilon'$ and $\varepsilon''$ for all samples at different percentages of rGO loading at the frequency range of 8–12 GHz are depicted in Figure 6.

![Figure 6](image_url). Relative permittivity for the epoxy, the hardener, and the cured epoxy at the X-band frequency.

At very high frequencies, the molecules do not have enough time to orient themselves in the direction of the alternating field [37]. Furthermore, the reduction of the value of complex permittivity can be explained by the conductive and capacitive properties of the liquid form of the epoxy resin, which are induced by the ions and dipoles. When the liquid form of the epoxy resin is exposed to an...
external electric field, ions and dipoles easily align themselves because of the mobility acquired by the low viscosity of the liquid form. Also, the curing causes a decline in the number of dipolar groups, which decreases the loss factor and the dielectric constant. The relation between the viscosity and the dielectric constant is inversely proportional [38]. It was found that both the real and imaginary part of the dielectric constant, which regularly decreased as the curing reaction proceeded, were mainly affected by the disappearance of specific dipolar species, whose relaxation time did not change significantly.

The effects of filler functionality and volume loading on the dielectric properties of the rGO/cured epoxy composites were studied. Figure 7a–c shows the dependence of the \( \varepsilon' \), \( \varepsilon'' \), and \( \tan \delta \) for the rGO/cured epoxy composites at different percentages of filler on the frequency. These figures show that, as more rGO particles were added to the epoxy matrix, the dielectric constant and loss factor gradually increase, with a high dielectric constant but a low loss factor comparable to that of a neat epoxy. The dielectric permittivity increment can be described as interfacial polarization, also known as the Maxwell–Wagner–Sillars (MWS) effect. In the composites, at 1% rGO, there was little distribution of rGO 88–92 particles; thus, there is a weak interaction with the matrix. When the rGO concentration is raised, clusters of filler particles are formed. A cluster may be considered as a region in the epoxy matrix where the particles are in contact or very close to each other as illustrated in the results of the scanning electron microscopy (SEM).

![Figure 7](image-url)

**Figure 7.** The frequency dependence of (a) The real part of permittivity (\( \varepsilon' \)), (b) The imaginary part of permittivity (\( \varepsilon'' \)), and (c) The tangent loss (\( \tan \delta \)) at various rGO loadings.

The average polarization that is associated with a cluster is larger than that of an individual particle because of an increase in the dimensions of the composite inclusion and, hence, greater interfacial area [39], which leads to a greater average polarization and, thus, a greater contribution to the dielectric permittivity. On the other hand, Figure 7a–c shows the dependence of the dielectric constant, loss factor, and loss tangent for the rGO/epoxy composites on the frequency. The dielectric constant of the composites decreases with the increase in the frequency. Due to the relatively high \( \varepsilon' \) of
rGO, the $\varepsilon'$ of the composites increased slightly from 3.15 for 1 vol % to 3.5 for 5 vol %. Furthermore, due to the relatively high $\varepsilon''$ of rGO, the $\varepsilon''$ of the composites increased gradually from 0.198 for 1 vol % to 0.246 for 5 vol %. In Figure 7c, it can be seen that the values of the loss tangent ($\tan\delta$) increase sequentially with increasing amounts of rGO in the composites, and were found to be 0.059 for 1 wt %, and 0.067 for 5 wt %. It could be obviously observed that the $E'$, $E''$, and $\tan\delta$ of the rGO composites with different volume fractions decreased as the frequency increased from 8 to 12 GHz. The result is due to the space-charge polarization, which originates from the conductor–insulator interfaces.

3.4.2. EMI Shielding Effectiveness

Figure 8a–d shows the $SE_R$, $SE_A$, and $SE_{total}$ values of the composites with different percentage fractions of rGO loading. Figure 8a shows that the $SE_R$ values of the rGO/cured epoxy composites decreased in frequency from 3.59 dB at 8 GHz to 0.45 dB at 12 GHz. The $SE_A$ values of the rGO/cured epoxy composites increase significantly as shown in Figure 8b. Clearly, the $SE_A$ values of the composites increased from 1.32 dB at 8 GHz to 24.78 dB at the 12 GHz frequency. Moreover, the $SE_A$ value was enhanced with an increase in frequency, while the $SE_R$ values decreased obviously. Based on the results, it can be concluded that absorption was the primary EMI shielding mechanism. The electron motion hysteresis in these dipoles under an alternating electromagnetic field induced additional polarization relaxation process which were favourable in enhancing microwave absorption attenuation [40]. Therefore, the rGO filler mixture with cured epoxy made a contribution to improving the EMI SE absorbing ability of the composites.
The SE\textsubscript{total} values of the rGO/cured epoxy composites are shown in Figure 8c. The results confirm that the addition of rGO to the matrix improved the EMI shielding property of the composites, which increased with increasing the rGO content. This result can be attributed to the increase in the polarity of all blends due to the increase of rGO concentration, which led to the improvement of the shielding ability by vast numbers of mobile charge carriers (electrons or holes) that made the major mobility interaction with an external EM field possible and easier. Therefore, adding rGO to the matrix lead to the convergence of the composite particles with each other, which facilitates the process for moving mobile charge carriers that could move freely along this convergence [41]. The SE\textsubscript{total} values of the composites with 1%, 2%, 3%, 4%, and 5% mass fractions of rGO at 8, 9, 10, 11 and 12 GHz are presented in Table 2.

Table 2. The SE\textsubscript{total} (dB) values for the rGO composites dependent on the frequencies and percentages of filler.

| Freq (GHz) | 1 wt % | 2 wt % | 3 wt % | 4 wt % | 5 wt % |
|-----------|--------|--------|--------|--------|--------|
| 8         | 5.233  | 5.061  | 4.912  | 4.451  | 4.927  |
| 9         | 7.309  | 7.322  | 7.038  | 7.239  | 7.476  |
| 10        | 9.516  | 9.556  | 9.872  | 10.109 | 10.692 |
| 11        | 12.038 | 13.226 | 13.779 | 14.444 | 15.991 |
| 12        | 16.046 | 19.434 | 20.226 | 21.933 | 25.748 |

We now discuss the surface chemistry of rGO and the reinforcing mechanisms of rGO in the polymer matrices. The rGO nanoparticles were distributed uniformly throughout the whole cured epoxy matrix. This distribution can greatly enhance the utilization ratio of rGO, leading to a reduced rGO loading and an increase in the electrical conductivity of the composite. More importantly, densely packed graphene networks at the interfaces can effectively interact with incident radiation, leading to a very high EMI SE. To better understand the shielding mechanism, the rGO/cured epoxy composite can be considered as a “skin” that is composed of closely packed cells, with dense rGO layers as highly conductive “membranes”. As shown schematically in Figure 9, incident electromagnetic microwaves entering the “skin” are attenuated by reflecting, scattering, and adsorption many times by the multiple layers of membranes. The “cells” of the rGO/cured epoxy composite lead to a great number of membranes such that it is very difficult for waves to penetrate this functional skin.

Figure 9. A schematic representation of microwave transfer across the rGO/cured epoxy composite with a thickness of 6 mm.
3.4.3. Cole–Cole Plot Analysis

The dielectric relaxation as a whole is the result of the movement of dipoles (dielectric relaxation) and electric charges (ionic relaxation) due to an applied alternating electric field. The Debye relaxation model has been widely employed to describe the response of molecules to an applied field. The rGO with a lower loss factor ($\varepsilon''$) has a stronger microwave absorption for three reasons: First, based on previous reports [42], apart from magnetic loss and dielectric loss, another concept of importance relating to the absorption of microwaves is the impedance match characteristic. Having an absorber permittivity that is too high harms the impedance match, resulting in weak absorption and strong reflection [43]. That explains the reasons for the exhibition of stronger microwave absorption by low-permittivity rGO. The second reason is that electronic spin is involved in the microwave band energy transition, which means that for microwave absorption, there is a requirement for greater spin states.

Near the Fermi level as documented, localized states could be created via the introduction of lattice defects. Additionally, electromagnetic energy absorption by a transition to the Fermi level from contiguous states can take place on the absorber surface when irradiation is incident on it [44]. Thus, the existence of defects in rGO favours the absorption of electromagnetic energy, which is an additional reason for the rGO’s exhibition of a better microwave absorption ability. The third reason is the dielectric loss material’s electromagnetic wave absorption mechanism, which arises from the process of relaxation. According to the expression of the equation for Debye relaxation in its simplest form, a single relaxation time ($\tau$) was assumed for the complex permittivity, which can be written using the equation as expressed in [45,46]

$$\varepsilon_r = \varepsilon_\infty + \frac{\varepsilon_s - \varepsilon_\infty}{1 + \left(\frac{\omega \tau}{\pi}ight)^2} = \varepsilon' - j\varepsilon''$$  \hspace{1cm} (6)

where $\varepsilon_s$, $\varepsilon_\infty$, $\omega$, and $\tau$ are the static dielectric constant and the dielectric constant at infinite frequency, the frequency, and polarization relaxation time, respectively. Thus, $\varepsilon'$ and $\varepsilon''$ can be described by

$$\varepsilon' = \varepsilon_\infty + \frac{\varepsilon_s - \varepsilon_\infty}{1 + \omega^2 \tau^2}$$  \hspace{1cm} (7)

$$\varepsilon'' = \frac{\omega \tau (\varepsilon_s - \varepsilon_\infty)}{1 + \omega^2 \tau^2}.$$  \hspace{1cm} (8)

Based on Equations (4) and (5), the relationship between $\varepsilon'$ and $\varepsilon''$ can be deduced as

$$\left(\varepsilon' - \frac{\varepsilon_s - \varepsilon_\infty}{2}\right)^2 + (\varepsilon'')^2 = \left(\frac{\varepsilon_s - \varepsilon_\infty}{2}\right)^2.$$  \hspace{1cm} (9)

Figure 10 presents the $\varepsilon'$ versus $\varepsilon''$ curve characteristic, which is also called the Cole–Cole semicircle [47]. The figures with a clear segment present the rGO’s three overlapped Cole–Cole semicircles, but only a single semicircle at the different filler percentages for the rGO/cured epoxy. This suggests that for rGO/cured epoxy and tripartite relaxation processes, there is a sole relaxation process for rGO, with one Debye relaxation process assigned to each semicircle. For rGO/cured epoxy, the sole process of relaxation may appear as follows: under the lag of induced charges that counters the externally applied field that results in the relaxation, the alternating electromagnetic field converts the electromagnetic energy to heat energy, so the attenuation of microwaves occurs [48,49]. In the rGO/cured epoxy, due to many delocalized electrons, this process of dielectric relaxation is obvious, and a big Cole–Cole semicircle protrudes, as shown in Figure 10, for different percentages of filler. Hence, the major reason for the microwave absorption of the rGO/cured epoxy is the dielectric relaxation.
4. Conclusions

The preparation of rGO was carried out using an ammonia solution method. Cured epoxy composites homogenized with the produced rGO at different loadings have been successfully fabricated using a mini-mechanical vortex mixer. The effect of the rGO powder and their loading on the dielectric properties and mechanical properties was investigated and compared. Various characterizations, including FT-IR, XRD, and FE-SEM, were performed on the samples. The FE-SEM images showed the layered and porous structure of rGO. The dielectric properties of the rGO/cured epoxy composites were measured in the frequency range from 8 to 12 GHz. The rGO in the cured epoxy matrix, even at the lowest concentration of 1%, has been found to show low values of dielectric properties. The calculated shielding effectiveness value of the composite with 5% rGO by weight in the cured epoxy matrix is quite high, i.e., 25.748 dB at 12 GHz at a thickness of 6 mm. The Cole–Cole plots showed the presence of only one process of dielectric relaxation for the rGO/cured epoxy composites with a poor impedance match characteristic, which resulted from the weak ability for...
under the influence of an alternating electromagnetic field, which supports the enhancement of the microwave absorbing ability.

4. Conclusions

The preparation of rGO was carried out using an ammonia solution method. Cured epoxy composites homogenized with the produced rGO at different loadings have been successfully fabricated using a mini-mechanical vortex mixer. The effect of the rGO powder and their loading on the dielectric properties and mechanical properties was investigated and compared. Various characterizations, including FT-IR, XRD, and FE-SEM, were performed on the samples. The FE-SEM images showed the layered and porous structure of rGO. The dielectric properties of the rGO/cured epoxy composites were measured in the frequency range from 8 to 12 GHz. The rGO in the cured epoxy matrix, even at the lowest concentration of 1%, has been found to show low values of dielectric properties. The calculated shielding effectiveness value of the composite with 5% rGO by weight in the cured epoxy matrix is quite high, i.e., 25.748 dB at 12 GHz at a thickness of 6 mm. The Cole–Cole plots showed the presence of only one process of dielectric relaxation for the rGO/cured epoxy composites with a poor impedance match characteristic, which resulted from the weak ability for microwave absorption. However, the groups and residual defects in rGO can not only enhance the impedance match characteristic, but also introduce a transition from contiguous states to the Fermi level, polarization relaxation of electronic dipole groups, and defect polarization relaxation, all of which favours electromagnetic wave absorption and penetration.

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References

1. Wang, L.L.; Tay, B.K.; See, K.Y.; Sun, Z.; Tan, L.K.; Lua, D. Electromagnetic interference shielding effectiveness of carbon-based materials prepared by screen printing. *Carbon* **2009**, *47*, 1905–1910. [CrossRef]

2. Choudary, V.; Dhawan, S.K.; Saini, P. Polymer based nanocomposites for electromagnetic interference (EMI) shielding. *In EMI Shielding Theory and Development of New Materials; Research Signpost: Kerala, India, 2012; pp. 67–100.

3. Thomassin, J.M.; Jerome, C.; Pardoen, T.; Bailly, C.; Huynen, I.; Detrembleur, C. Polymer/carbon based composites as electromagnetic interference (EMI) shielding materials. *Mater. Sci. Eng. R. Rep.* **2013**, *74*, 211–232. [CrossRef]

4. Phang, S.W.; Tadokoro, M.; Watanabe, J.; Kuramoto, N. Synthesis, characterization and microwave absorption property of doped polyaniline nanocomposites containing TiO<sub>2</sub> nanopar-ticles and carbon nanotubes. *Synth. Met.* **2008**, *158*, 251–258. [CrossRef]

5. Liu, C.; Wang, X.; Huang, X.; Liao, X.; Shi, B. Absorption and Reflection Contributions to the High Performance of Electromagnetic Waves Shielding Materials Fabricated by Compositing Leather Matrix with Metal Nanoparticles. *ACS Appl. Mater. Interfaces* **2018**, *16*, 14036–14044. [CrossRef] [PubMed]

6. Kashi, S.; Gupta, R.K.; Baum, T.; Kao, N.; Bhattacharya, S.N. Morphology, electromagnetic properties and electromagnetic interference shielding performance of poly lactide/graphene nanoplatelet nanocomposites. *Mater. Sci.* **2016**, *95*, 119–126. [CrossRef]

7. Karteri, I.; Altun, M.; Gunes, M. Electromagnetic interference shielding performance and electromagnetic properties of wood-plastic nanocomposites with graphene nano platelets. *J. Mater. Sci. Mater. Electron.* **2017**, *28*, 6704–6711. [CrossRef]
8. Deng, B.; Hsu, P.C.; Chen, G.; Chandrashekar, B.N.; Liao, L.; Ayitimuda, Z.; Wu, J.; Guo, Y.; Lin, L.; Zhou, Y.; et al. Roll-to-roll encapsulation of metal nanowires between graphene and plastic substrate for high-performance flexible transparent electrodes. *Nano Lett.* **2015**, *15*, 4206–4213. [CrossRef] [PubMed]

9. Mondal, S.; Nayak, L.; Rahaman, M.; Aldalbahai, A.; Chaki, T.K.; Khastgir, D.; Das, N.C. An effective strategy to enhance mechanical, electrical, and electromagnetic shielding effectiveness of chlorinated polyethylene-carbon nanofiber nanocomposites. *Compos. Part B Eng.* **2017**, *109*, 155–169. [CrossRef]

10. Al-Saleh, M.H.; Saadeh, W.H.; Sundararaj, U. EMI shielding effectiveness of carbon based nanostructured polymeric materials: A comparative study. *Carbon* **2013**, *60*, 146–156. [CrossRef]

11. Gao, C.; Zhang, S.; Wang, F.; Wen, B.; Han, C.; Ding, Y.; Yang, M. Graphene networks with low percolation threshold in ABS nanocomposites: Selective localization and electrical and rheological properties. *ACS Appl. Mater. Interfaces* **2014**, *15*, 12252–12260. [CrossRef] [PubMed]

12. Wang, C.; Han, X.; Xu, P.; Zhang, X.; Du, Y.; Hu, S.; Wang, J.; Wang, X. The electromagnetic property of chemically reduced graphene oxide and its application as microwave absorbing material. *Appl. Phys. Lett.* **2011**, *98*, 072906. [CrossRef]

13. Zang, H.B.; Yan, Q.; Zheng, W.G.; He, Z.; Yu, Z.Z. Tough graphene-polymer microcellular foams for electromagnetic interference shielding. *ACS Appl. Mater. Interfaces* **2011**, *3*, 918–924. [CrossRef] [PubMed]

14. Yu, H.; Wang, T.; Wen, B.; Lu, M.; Xu, Z.; Zhu, C.; Chen, Y.; Xue, X.; Sun, C.; Cao, M. Graphene/polyaniline nanorod arrays: Synthesis and excellent electromagnetic absorption properties. *J. Mater. Chem.* **2012**, *22*, 21679–21685. [CrossRef]

15. Wang, Z.; Luo, J.; Zhao, G.L. Dielectric and microwave absorption properties of graphene nanoplatelet-epoxy composites. *AIP Adv.* **2014**, *4*, 017139. [CrossRef]

16. Liang, J.; Wang, Y.; Huang, Y.; Ma, Y.; Liu, Z.; Cai, J.; Zhang, C.; Gao, H.; Chen, Y. Electromagnetic interference shielding of graphene/epoxy composites. *Carbon* **2009**, *47*, 922–925. [CrossRef]

17. Chhetri, S.; Samanta, P.; Murmu, N.C.; Srivastava, S.K.; Kuila, T. Electromagnetic interference shielding and thermal properties of non-covalently functionalized reduced graphene oxide/epoxy composites. *AIMS Mater.* **2016**, *4*, 61–74. [CrossRef]

18. Ameer, S.; Gul, I.H. Influence of Reduced Graphene Oxide on Effective Absorption Bandwidth Shift of Hybrid Absorbers. *PLoS ONE* **2016**, *11*, 0153544. [CrossRef] [PubMed]

19. Zhang, X.J.; Wang, G.S.; Cao, W.Q.; Wei, Y.Z.; Liang, J.F.; Guo, L.; Cao, M.S. Enhanced microwave absorption property of reduced graphene oxide (RGO)-MnFe$_2$O$_4$ nanocomposites and polyvinylidene fluoride. *ACS Appl. Mater. Interfaces* **2014**, *6*, 7471–7478. [CrossRef] [PubMed]

20. Khurana, G.; Kumar, N.; Kooriyattil, S.; Katiyar, R.S. Structural, magnetic, and dielectric properties of graphene oxide/Zn$_{1-x}$Fe$_x$O$_2$ compositions. *J. Appl. Phys.* **2015**, *117*, 17E106. [CrossRef]

21. Li, Y.; Chen, C.; Li, J.T.; Zhang, S.; Li, C.; Cao, J.; Huang, J. Enhanced dielectric constant for efficient electromagnetic shielding based on carbon-nanotube-added styrene acrylic emulsion based composite. *Nanoscale Res. Lett.* **2010**, *5*, 1170–1176. [CrossRef] [PubMed]

22. Poh, C.L.; Mariatti, M.; Fauzi, M.A.; Ng, C.H.; Chee, C.K.; Chuaah, T.P. Tensile, dielectric, and thermal properties of epoxy composites filled with silica, mica, and calcium carbonate. *J. Mater. Sci. Mater. Electron.* **2014**, *25*, 2111–2119. [CrossRef]

23. Guo, J.; Song, H.; Liu, H.; Luo, C.; Ren, Y.; Ding, T.; Khan, M.A.; Young, D.P.; Liu, X.; Zhang, X.; et al. Polyppyrole-interface-functionalized nano-magnetite epoxy nanocomposites as electromagnetic wave absorbers with enhanced flame retardancy. *J. Mater. Chem. C* **2017**, *5*, 5334–5344. [CrossRef]

24. Idris, F.M.; Hashim, M.; Abbas, Z.; Ismail, I.; Nazlan, R.; Ibrahim, I.R. Recent developments of smart electromagnetic absorbers-based polymer-composites at gigahertz frequencies. *J. Magn. Magn. Mater.* **2016**, *405*, 197–208. [CrossRef]

25. Chen, J.; Wu, J.; Ge, H.; Zhao, D.; Liu, C.; Hong, X. Reduced graphene oxide deposited carbon fiber reinforced polymer composites for electromagnetic interference shielding. *Compos. Part A* **2016**, *82*, 141–150. [CrossRef]

26. Ouyang, X.; Huang, W.; Cabrera, E.; Castro, J.; Lee, I.J. Graphene-graphene oxide-graphene hybrid nanopapers with superior mechanical, gas barrier and electrical properties. *AIP Adv.* **2015**, *5*, 017135. [CrossRef]

27. Wan, Y.J.; Gong, L.X.; Tang, L.C.; Wu, L.B.; Jiang, J.X. Mechanical properties of epoxy composites filled with silane-functionalized graphene oxide. *Compos. Part A* **2014**, *64*, 79–89. [CrossRef]
28. Singh, V.K.; Shukla, A.; Patra, M.K.; Saini, L.; Jani, R.K.; Vadera, S.R.; Kumar, N. Microwave absorbing properties of a thermally reduced graphene oxide/nitrile butadiene rubber composite. *Carbon* 2012, 50, 2202–2208. [CrossRef]

29. Liu, P.; Yao, Z.; Zhou, J. Preparation of reduced graphene oxide/\(\text{Ni}_0.4\text{Zn}_{0.6}\text{Co}_{0.2}\text{Fe}_2\text{O}_4\) nanocomposites and their excellent microwave absorption properties. *Ceram. Int.* 2015, 41, 13409–13416. [CrossRef]

30. Dikin, D.A.; Stankovich, S.; Zimney, E.J.; Piner, R.D.; Dommett, G.H.; Evmenenko, G.; Nguyen, S.T.; Ruoff, R.S. Preparation and characterization of graphene oxide paper. *Nature* 2007, 448, 457. [CrossRef] [PubMed]

31. Tang, L.C.; Wan, Y.J.; Yan, D.; Pei, Y.B.; Zhao, L.; Li, Y.B.; Wu, L.B.; Jiang, J.X.; Lai, G.Q. The effect of graphene dispersion on the mechanical properties of graphene/epoxy composites. *Carbon* 2013, 60, 16–27. [CrossRef]

32. Dato, A.; Lee, Z.; Jeon, K.J.; Erni, R.; Radmilovic, V.; Richardson, T.J.; Frenklach, M. Clean and highly ordered graphene synthesized in the gas phase. *Chem. Commun.* 2009, 40, 6095–6097. [CrossRef] [PubMed]

33. Matchi, R.; Venkatatalapathy, S.; Srinivas, B.A. Synthesis, stability, transport properties, and surface wettability of reduced graphene oxide/water nanofluids. *INT J. Therm. Sci.* 2015, 97, 17–25. [CrossRef]

34. Moghaddam, M.B.; Goharshadi, E.K.; Entezari, M.H.; Nancarrow, P. Preparation, characterization, and rheological properties of graphene–glycerol nanofluids. *Chem. Eng. J.* 2013, 231, 365–372. [CrossRef]

35. González, M.G.; Cabanelas, J.C.; Basela, J. Applications of FTIR on epoxy resins-identification, monitoring the curing process, phase separation and water uptake. In *Infrared Spectroscopy-Materials Science, Engineering and Technology*; InTech: New York, NY, USA, 2012; ISBN 978-953-51-0537-4.

36. Singh, A.P.; Garg, P.; Alam, F.; Singh, K.; Mathur, R.B.; Tandon, R.P.; Chandra, A.; Dhawan, S.K. Phenolic resin-based composite sheets filled with mixtures of reduced graphene oxide, \(\gamma\)-\(\text{Fe}_2\text{O}_3\) and carbon fibers for excellent electromagnetic interference shielding in the X-band. *Carbon* 2012, 50, 3868–3875. [CrossRef]

37. Wei, J.; Liu, J.; Li, S. Electromagnetic and microwave absorption properties of \(\text{Fe}_2\text{O}_4\) magnetic films plated on hollow glass spheres. *J. Magn. Magn. Mater.* 2007, 321, 414–417. [CrossRef]

38. Li, Z.; Sheng, G.; Jiang, X.; Tanaka, T. Effects of inorganic fillers on withstanding short-time breakdown and long-time electrical aging of epoxy composites. *IEEE Trans. Electr. Electron. Eng.* 2017, 12. [CrossRef]

39. Park, H. Dielectric cure determination of a thermosetting epoxy composite prepreg. *J. Appl. Polym. Sci.* 2017, 134. [CrossRef]

40. Zhou, W.; Yu, D. Thermal and dielectric properties of the aluminum particle/epoxy resin composites. *J. Appl. Polym. Sci.* 2010, 118, 3156–3166. [CrossRef]

41. Kong, L.; Yin, X.; Zhang, Y.; Yuan, X.; Li, Q.; Ye, F.; Cheng, L.; Zhang, L. Electromagnetic wave absorption properties of reduced graphene oxide modified by maghemite colloidal nanoparticle clusters. *J. Phys. Chem. C.* 2013, 117, 19701–19711. [CrossRef]

42. Wu, J.; Ye, Z.; Ge, H.; Chen, J.; Liu, W.; Liu, Z. Modified carbon fiber/magnetic graphene/epoxy composites with synergistic effect for electromagnetic interference shielding over broad frequency band. *J. Colloid Interface Sci.* 2017, 506, 217–226. [CrossRef] [PubMed]

43. Xu, P.; Han, X.; Wang, C.; Zhou, D.; Lv, Z.; Wen, A.; Wang, X.; Zhang, B. Synthesis of electromagnetic functionalized nickel/polypyrrole core/shell composites. *J. Phys. Chem. B* 2008, 112, 10443–10448. [CrossRef] [PubMed]

44. Wu, N.; Liu, X.; Zhao, C.; Cui, C.; Xia, A. Effects of particle size on the magnetic and microwave absorption properties of carbon-coated nickel nanocapsules. *J. Alloys Compd.* 2016, 656, 628–634. [CrossRef]

45. Hsiao, S.T.; Ma, C.C.M.; Tien, H.W.; Liao, W.H.; Wang, Y.S.; Li, S.M.; Yang, C.Y.; Lin, S.C.; Yang, R.B. Effect of coherent modification of graphene nanosheets on the electrical property and electromagnetic interference shielding performance of a water-borne polyurethane composite. *ACS Appl. Mater. Interfaces* 2015, 7, 2817–2826. [CrossRef] [PubMed]

46. Zhao, B.; Shao, G.; Fan, B.; Sun, B.; Guan, K.; Zhang, R. Facile synthesis and novel microwave electromagnetic properties of flower-like Ni structures by a solvothermal method. *J. Mater. Sci. Mater. Electron.* 2014, 25, 3614–3621. [CrossRef]

47. Zhang, X.J.; Wang, G.S.; Wei, Y.Z.; Guo, L.; Cao, M.S. Polymer-composite with high dielectric constant and enhanced absorption properties based on graphene–CuS nanocomposites and polyvinylidene fluoride. *J. Mater. Chem. A* 2013, 1, 12115–12122. [CrossRef]

48. Bayat, M.; Yang, H.; Ko, F.K.; Michelson, D.; Mei, A. Electromagnetic interference shielding effectiveness of hybrid multifunctional \(\text{Fe}_2\text{O}_4\)/carbon nanofiber composite. *Polymer* 2014, 55, 936–943. [CrossRef]
49. Imholt, T.J.; Dyke, C.A.; Hasslacher, B.; Pérez, J.M.; Price, D.W.; Roberts, J.A.; Scott, J.B.; Dhawan, A.; Ye, Z.; Tour, J.M. Nanotubes in microwave fields: Light emission, intense heat, outgassing, and reconstruction. Chem. Mater. 2003, 15, 3969–3970. [CrossRef]

50. Vázquez, E.; Prato, M. Carbon nanotubes and microwaves: Interactions, responses, and applications. ACS Nano 2009, 3, 3819–3824. [CrossRef] [PubMed]

51. Wen, B.; Cao, M.; Lu, M.; Cao, W.; Shi, H.; Liu, J.; Wang, X.; Jin, H.; Fang, X.; Wang, W.; et al. Reduced graphene oxides: Light-weight and high-efficiency electromagnetic interference shielding at elevated temperatures. Adv. Mater. 2014, 26, 3484–3489. [CrossRef] [PubMed]

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