


Cite this: *RSC Adv.*, 2025, 15, 40252

# Structural properties and electromagnetic shielding performance of bidirectional carbon fabric and Fe<sub>2</sub>O<sub>3</sub> nano filler reinforced epoxy composites

R. Suresha,<sup>a</sup> H. K. Sachidananda,<sup>b</sup> B. Shivamurthy,<sup>\*c</sup> Gibin George<sup>d</sup> and Sampath Parasuram<sup>e</sup>

This study investigates the structural and electromagnetic interference (EMI) shielding performance of bidirectional carbon fabric/epoxy composites reinforced with Fe<sub>2</sub>O<sub>3</sub> nanoparticles for aerospace applications. Composites with varying Fe<sub>2</sub>O<sub>3</sub> loadings (1–3 wt%) were fabricated using a hand lay-up method, followed by mechanical and electromagnetic shielding characterisation in the X-band frequency range (8.2–12.4 GHz). Tensile testing revealed that 1–2 wt% Fe<sub>2</sub>O<sub>3</sub> loading enhanced the stiffness and tensile strength of the composite due to improved fibre–matrix interfacial bonding, while 3 wt% caused agglomeration and reduced strength. EMI shielding measurements showed absorption-dominated performance across all configurations, with multiple layers significantly improving total shielding effectiveness (SE<sub>T</sub>). The highest SE<sub>T</sub> (25.3 dB) was achieved for a two-layer laminate with 3 wt% Fe<sub>2</sub>O<sub>3</sub>, attributed to synergistic dielectric and magnetic losses and enhanced internal reflections. The results demonstrate that optimised Fe<sub>2</sub>O<sub>3</sub> content and laminate layering can deliver lightweight, structurally robust composites with effective EMI shielding, making them suitable for advanced aerospace structures requiring mechanical integrity and electromagnetic compatibility. These findings highlight that optimized nanoparticle loading, and laminate architecture can yield lightweight composites that unite mechanical robustness with effective EMI shielding, offering strong potential for aerospace structures demanding both structural performance and electromagnetic compatibility.

Received 22nd August 2025  
Accepted 15th October 2025

DOI: 10.1039/d5ra06245d

rsc.li/rsc-advances

## 1. Introduction

Modern avionics and onboard systems integrate dense digital avionics, networked sensors, and wireless links, heightening susceptibility to electromagnetic interference from both onboard systems and passenger devices. Uncontrolled EMI degrades signal integrity and can trigger malfunctions or failures in safety-critical functions, undermining operational reliability and flight safety.<sup>1</sup>

In addition to onboard sources, aircraft encounter external EMI from thunderstorms, electrostatic discharges, and solar activity, making shielding essential for flight safety.<sup>2</sup> At the same time, the drive for lightweight, high-performance structures has accelerated the use of carbon fibre-reinforced epoxy (CF/E) composites, which combine high specific strength, fatigue resistance, thermal stability, and corrosion resistance, positioning them as attractive alternatives to traditional metals in aerospace applications.<sup>3</sup>

Although carbon fiber/epoxy composites provide excellent structural performance, their semiconducting nature limits intrinsic EMI shielding. Consequently, research increasingly targets enhancing their effectiveness in the 8–12 GHz X-band frequency, a range vital for radar, satellite communication, navigation, and wireless networks, where reliable EMI protection is essential for uninterrupted operation.<sup>4</sup>

Bidirectional carbon fabric/epoxy composites offer superior EMI shielding compared to unidirectional ones, but excessive nanofiller addition can cause agglomeration and degrade mechanical strength, underscoring the need for optimized loading.<sup>5</sup> Recent studies emphasize strategies such as incorporating magnetic nanofillers, forming conductive networks, and

<sup>a</sup>School of Engineering & Information Technology, Department of Electrical & Electronics Engineering, Manipal Academy of Higher Education, Dubai, United Arab Emirates. E-mail: suresha@manipaldubai.com

<sup>b</sup>School of Engineering & Information Technology, Department of Mechanical Engineering, Manipal Academy of Higher Education, Dubai, United Arab Emirates. E-mail: sachidananda@manipaldubai.com

<sup>c</sup>Department of Mechanical & Industrial Engineering, Manipal Institute of Technology, Manipal Academy of Higher Education, Manipal-576104, India. E-mail: shiva.b@manipal.edu

<sup>d</sup>Department of Mechanical Engineering, SCMS School of Engineering and Technology, Karukutty, Kerala, India. E-mail: gibingeorge@scmsgroup.org

<sup>e</sup>Department of Materials Engineering, Indian Institute of Science, Bangalore, India. E-mail: parasurams@iisc.ac.in



designing multilayer structures to enhance shielding effectiveness in the X-band (8.2–12.4 GHz).

Ahmad *et al.*<sup>6</sup> showed that alternating carbon fiber plies reinforced with MWCNTs and Fe<sub>2</sub>O<sub>3</sub> enhanced EMI shielding *via* combined dielectric and magnetic losses. Waseem *et al.*<sup>7</sup> developed a hybrid layered composite using ZrB<sub>2</sub>-SiC, Fe<sub>3</sub>O<sub>4</sub>-loaded carbon fabric, and carbonized cotton fibre, achieving lightweight, thermally stable, absorption-dominated shielding. Shao *et al.*<sup>8</sup> reported a Janus-structured CNT/Fe<sub>3</sub>O<sub>4</sub>/MXene membrane (~84.9 μm) with 44.56 dB shielding in the X-band, along with excellent thermal conductivity, flexibility, and low weight. Likewise, Yu *et al.*<sup>9</sup> created a nacre-inspired Fe<sub>3</sub>O<sub>4</sub>/CNT composite with alternating conductive and magnetic layers, delivering durable, absorption-dominant shielding that retained 96% efficiency after physical damage.

Zhang *et al.*<sup>10</sup> engineered ultralight rGO-CNT-epoxy aerogels with Fe<sub>3</sub>O<sub>4</sub> nanoparticles, achieving a reflection loss of −58.13 dB at 12.08 GHz through a 3D porous structure that enhanced scattering and impedance matching. Wilson *et al.*<sup>11</sup> reported TPU laminates with graphite and CoFe<sub>2</sub>O<sub>4</sub>, where the G/F/G configuration reached approximately 54 dB shielding from synergistic dielectric and magnetic losses. Durmaz *et al.*<sup>12</sup> developed bio-based PA11/PLA composites with 30 wt% carbon fibres, yielding 28 dB SE at 10 GHz through reflection-dominated shielding. Ahmad *et al.*<sup>13</sup> demonstrated graphene-Fe<sub>2</sub>O<sub>3</sub> polymer composites with broadband EMI attenuation from combined dielectric reflection and magnetic absorption. Duan *et al.*<sup>14</sup> fabricated CF/GO/Fe<sub>3</sub>O<sub>4</sub>/epoxy multilayers that improved impedance matching and promoted absorption-based shielding. Fallah *et al.*<sup>15</sup> achieved 36.6 dB SE at 8.2 GHz in epoxy nanocomposites with Fe<sub>2</sub>O<sub>3</sub> and carbon black, showing tunable frequency response. Bhaskaran *et al.*<sup>16</sup> developed Fe<sub>3</sub>-O<sub>4</sub>@GNP hybrids with ~9.6 dB SE at 1 mm, outperforming individual fillers through synergistic effects. Xu *et al.*<sup>17</sup> produced epoxy-cotton fibre scaffolds with GNPs and Fe<sub>3</sub>O<sub>4</sub>, delivering 33.1 dB SE<sub>T</sub> and absorption-dominant shielding with structural stability for defence and electronics.

Veeramani *et al.*<sup>18</sup> studied carbon fiber/epoxy composites fabricated *via* vacuum-assisted resin infusion, incorporating polyphenylene ether (PPE) as a toughening agent. PPE dispersed in the interlaminar region enhanced toughness, with DMA results showing a notable increase in glass transition temperature. In another work, Veeramani *et al.*<sup>19</sup> developed self-healing composites by embedding urea-formaldehyde-encapsulated dicyclopentadiene (DCPD) microcapsules with epoxy, chopped CF, and CNTs. These systems enabled self-detection and prevention of microcrack propagation, with potential applications in aerospace, wind energy, and marine structures. Eken *et al.*<sup>20</sup> investigated two-layer CFRP composites reinforced with hematite (Fe<sub>2</sub>O<sub>3</sub>) and goethite (FeO(OH)) produced by hand lay-up, evaluating EMI shielding across 700–6000 MHz and impact resistance. They reported overall improvements in both shielding effectiveness and impact strength compared to unreinforced samples.

Fe<sub>2</sub>O<sub>3</sub> was selected in this study due to its lower cost, environmental benignity, chemical stability, and strong magnetic loss response in the X-band compared to fillers like Fe<sub>3</sub>O<sub>4</sub> and

CoFe<sub>2</sub>O<sub>4</sub>. Unlike earlier works that broadly combined carbon-based conductive and magnetic fillers, the novelty of this research lies in optimizing Fe<sub>2</sub>O<sub>3</sub> loading within carbon fabric/epoxy laminates to simultaneously balance structural performance and absorption-dominated EMI shielding. The exclusive focus on the X-band is justified by its critical importance in aerospace applications, particularly radar, satellite communication, and navigation systems, where reliable EMI protection is essential for operational safety.

Previous studies on conductive and magnetic nanofillers have advanced EMI shielding but often overlooked structural robustness. Next-generation aerospace, defence, and wearable systems require materials that are both lightweight and multifunctional. This study addresses that gap by investigating Fe<sub>2</sub>O<sub>3</sub>-reinforced carbon fabric/epoxy composites, linking shielding effectiveness with mechanical performance. By integrating structural strength with absorption-dominated shielding, the work proposes a multifunctional composite system that meets aerospace demands for both mechanical integrity and electromagnetic compatibility, thereby contributing to the design of advanced materials capable of balancing performance requirements often neglected in earlier research.

## 2. Materials and methods

In this research work, the epoxy matrix is reinforced with bidirectional carbon fabric and Fe<sub>2</sub>O<sub>3</sub> nanoparticles to prepare composites.

### 2.1. Materials

The bi-directional oriented plain weave type carbon fabric supplied by M/s Bhor Chemical & Plastics Pvt. Ltd, Nasik, India, is used as primary reinforcement. The material batch number: W21A31/3298, the area density of the carbon fabric used, and the type of fibre were 160 g m<sup>−2</sup> ± 5%, and 3k, respectively. Fig. 1(a) is the digital photograph of the carbon fabric used for this study. Bisphenol-A epoxy, combined with a suitable hardener supplied by M/s Addnano, was used as matrix material. Fe<sub>2</sub>O<sub>3</sub> Nanofillers (Fig. 1(b)) are procured from M/s Addnano, used as secondary reinforcement. The density of the oxide is about 5.3 g cm<sup>−3</sup>, molecular weight 159.6 g mol<sup>−1</sup>, and melting point about 1565 °C. All these components are used as received for processing the composite. In this study, we used diglycidyl ether of bisphenol-A (DGEBA, LY 556 grade, Huntsman) as the epoxy resin, cured with aromatic amine hardener (HY 951) at a 100:20 resin-to-hardener weight ratio, following an amine addition curing mechanism. Ethanol was chosen as the dispersant for Fe<sub>2</sub>O<sub>3</sub> nanoparticles due to its low toxicity, high volatility, and good compatibility with both the epoxy system and nanoparticle surfaces, which facilitated uniform dispersion while minimizing health and environmental hazards compared to DMF or acetone. To ensure complete removal of residual ethanol prior to curing, the nanoparticle-resin mixtures were subjected to continuous magnetic stirring and mild heating at 60 °C, followed by vacuum drying, thereby eliminating solvent traces that could otherwise affect curing or composite integrity.



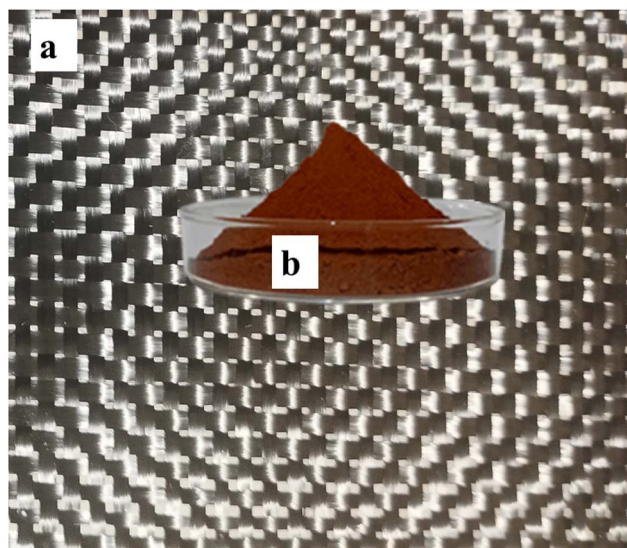


Fig. 1 Digital photographs of (a) carbon fabric, (b) Fe<sub>2</sub>O<sub>3</sub>.

## 2.2. Preparation of the Fe<sub>2</sub>O<sub>3</sub> dispersed epoxy matrix

When preparing Fe<sub>2</sub>O<sub>3</sub> nanoparticle-loaded carbon fibre-reinforced epoxy composites, it is paramount that the nanofillers are evenly dispersed within the epoxy matrix. Nanofiller dispersion in the matrix allows for an efficient loading of the reinforced area, which can positively affect load transfer from the matrix to the reinforcements. This eventual loading establishes and extends the interfaces between the nanofillers and polymer matrix, leading to improved strength of the matrix, in addition to other functional attributes. However, high viscosity and low solubility of the epoxy polymer are significant hurdles to effective nanofiller dispersion in the polymer matrix. Solvents such as tetrahydrofuran (THF), dimethylformamide (DMF), acetone, ethanol, water, dichloromethane (DCM), and methyl ethyl ketone

(MEK) have been utilised as dispersants to allow the nanofiller to be dispersed into the epoxy matrix. The most common methods used for dispersion have used techniques including tip sonication, bath sonication, mechanical mixing, shear mixing, and three-roll calendaring. Most of these methods of dispersion have been successful. In this study, ethanol was used as the solvent to disperse the Fe<sub>2</sub>O<sub>3</sub> nanoparticles in the epoxy resin. Incorporating Fe<sub>2</sub>O<sub>3</sub> nanoparticles into multifunctional epoxies like TGDDM or TGPAP would likely enhance EMI shielding stability at high temperatures, strengthen interfacial polarization effects, and improve mechanical/thermal endurance compared to DGEBA-based systems. The only challenges are related to brittleness and nanoparticle dispersion. These multifunctional epoxies provide a more robust and reliable platform for advanced EMI shielding composites, particularly for aerospace and defence applications where performance under extreme conditions is required.

The preparation of the Fe<sub>2</sub>O<sub>3</sub>-filled epoxy resin is illustrated in Fig. 2. First, equal amounts of ethanol and epoxy were mixed with a magnetic stirrer for 30 minutes and termed as solution A. In a separate container, Fe<sub>2</sub>O<sub>3</sub> nanoparticle was dispersed in ethanol *via* a probe-type ultrasonicator for 2 hours, and this mixture was termed solution B. After a uniform dispersion of Fe<sub>2</sub>O<sub>3</sub> in ethanol was achieved, solution B was mixed with solution A and ultrasonic dispersion was performed for an additional 2 hours. This mixture was termed solution C. Solution C was mixed with hardener and mechanically stirred for 5 minutes to prepare the hybrid matrix for composite laminate fabrication. This procedure for matrix preparation was repeated for different Fe<sub>2</sub>O<sub>3</sub> nanoparticle concentrations of 1 wt%, 2 wt%, and 3 wt%. The material compositions of the hybrid matrix prepared using the previously described method are summarised in Table 1.

## 2.3. Fabrication of composite laminates

The bi-directional plain weave carbon fabric that the manufacturer provided was cut into 100 mm × 100 mm squares and

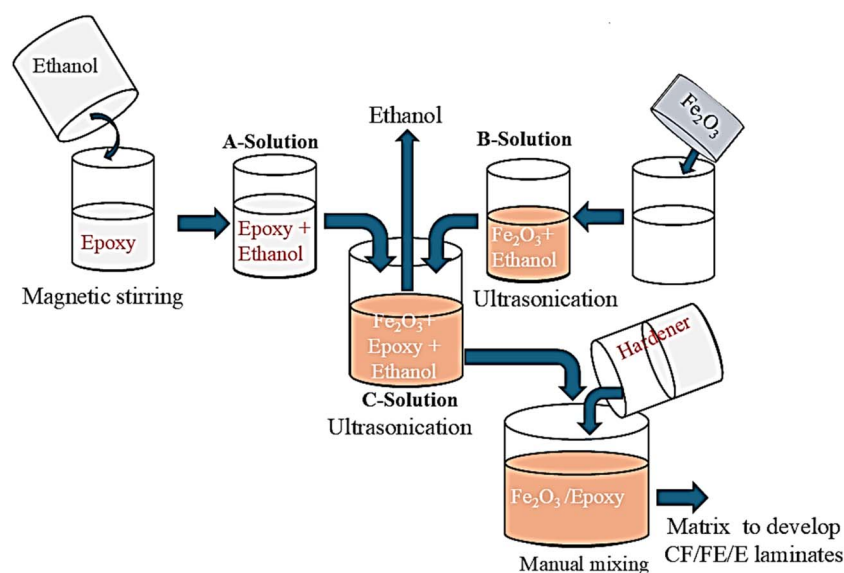


Fig. 2 Fe<sub>2</sub>O<sub>3</sub>-filled epoxy matrix preparation.



Table 1 Material constituents of the Fe<sub>2</sub>O<sub>3</sub>-filled epoxy resin

Sl no.	Sample designation	Fe <sub>2</sub> O <sub>3</sub> (wt%)	Epoxy (vol%)	Hardener (vol%)
1	E	0	100	20
2	1FE/E	1	100	20
3	2FE/E	2	100	20
4	3FE/E	3	100	20

immersed in acetone to remove sizing agents, dirt, contaminations, amorphous carbon, and grease. After soaking in the acetone, the mass was rinsed with deionized water and then placed in the hot air oven for 10 minutes to remove residual moisture.

The cleaned carbon fabric was then placed on a glass plate, and the prepared hybrid epoxy matrix was spread over the surface of the carbon fabric using a roller and brush. Similarly, fabrics were stacked one above the other, with the matrix applied between the fabric layers and then consolidated by pressing with a roller. Before placing the carbon fabric on the glass plate, a releasing film was used to facilitate easy removal of the laminate. Likewise, after completing the layup, a releasing film was placed on the top along with the glass plate. The entire stack, impregnated with resin, was kept at room temperature for curing, which took about 24 hours, resulting in the production of the laminates. Table 2 presents the material composition and designation of the composite sample laminates.

## 2.4. Characterization

The morphology of the Fe<sub>2</sub>O<sub>3</sub> nanofiller was studied by scanning electron microscopy (SEM; CARL ZEISS, Germany). The crystalline structure of the Fe<sub>2</sub>O<sub>3</sub> nanoparticles was identified using powder X-ray diffraction (XRD; Rigaku MiniFlex 600, Japan) which ran at scanning speed in a  $2\theta$  range of 20° to 80°. Fourier Transform Infrared spectroscopy (IR Spirit, Shimadzu, Japan) was performed on the composite and each of its components to capture any interactions. For mechanical characterization, the test samples were cut using an abrasive water jet machining as per ASTM standards. The samples for the tensile test were prepared according to ASTM D-638.<sup>21</sup>

**2.4.1. Electromagnetic interference shielding test.** The shielding effectiveness of the composite laminates was evaluated using the coaxial transmission line method, wherein the scattering parameters (*S*-parameters: *S*<sub>11</sub>, *S*<sub>12</sub>, *S*<sub>21</sub>, and *S*<sub>22</sub>) were measured with a two-port Vector Network Analyzer (VNA), model Keysight Technologies N1991A MY58312077, integrated with a WR-90 waveguide specimen holder.

Among the measured parameters, *S*<sub>11</sub> (reflection coefficient at port 1) quantifies the portion of incident power reflected back from the material, while *S*<sub>21</sub> (transmission coefficient from port 1 to port 2) indicates the amount of electromagnetic power transmitted through the specimen. Parameters *S*<sub>12</sub> and *S*<sub>22</sub>, typically associated with reverse transmission and reflection at port 2, were considered less critical for this application and thus excluded from the analysis. The *S*<sub>21</sub> parameter is of primary importance in determining shielding effectiveness, as it directly correlates with the transmitted power and, therefore, the attenuation capability of the shielding material.

The specimens for EMI shielding (22.86 mm × 10.16 mm) were cut with an abrasive water jet machine from the prepared CF/E and CF/FE/E laminates with strict dimensional tolerances and high edge quality. The samples were mounted in a WR-90 rectangular waveguide sample holder, where the scattering parameters (*S*-parameters) were measured at frequencies between 8 and 12 GHz. Each specimen was inserted into the measurement system after performing a total two-port calibration and recording the *S*-parameters from a baseline for each specimen to minimise systematic error.

As shown in Fig. 3, the block diagram depicts the experimental setup for evaluating EMI shielding effectiveness (SE) in the X-band frequency range (8.2–12.4 GHz) using a rectangular waveguide. The vector network analyzer (VNA) generates and receives microwave signals using two coaxial ports. The signals

Table 2 Constituents and designations of the prepared composite laminates

Sl. no.	Fe <sub>2</sub> O <sub>3</sub> (wt%)	Number of layers	Epoxy : hardener (vol, %)	Sample designation
1	0	1	100 : 20	CF/E-1Layer
2		2		CF/E-2Layer
3		4		CF/E-4Layer
4		1		CF/1FE/E-1Layer
5	1	2		CF/1FE/E-2Layer
6		4		CF/1FE/E-4Layer
7		1		CF/2FE/E-1Layer
8		2		CF/2FE/E-2Layer
9	2	4		CF/2FE/E-4Layer
10		1		CF/3FE/E-1Layer
11		2		CF/3FE/E-2Layer
12		4		CF/3FE/E-4Layer



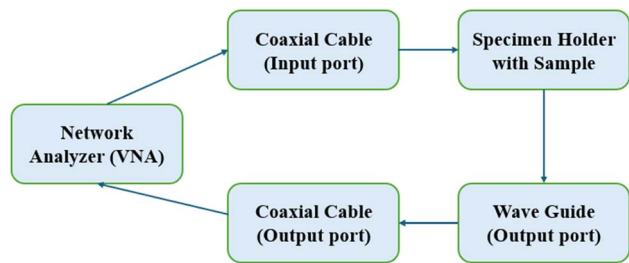


Fig. 3 Experimental setup for measuring electromagnetic interference (EMI) shielding effectiveness.

are generated on port 1 and transmitted *via* a coaxial cable into the waveguide input, where the test specimen is mounted in a custom holder. The attenuated or transmitted signals exit through the waveguide output and are then directed to port 2 of the VNA. In this setup, it is now possible to measure the scattering parameters ( $S_{11}$  and  $S_{21}$ ) to evaluate a specimen's shielding performance. Total shielding effectiveness was computed from the measured forward transmission as given in eqn (1) and (2) of the manuscript consistent with waveguide SE formulations reported for planar samples. Because the WR-90 fixture with a flat, isotropic specimen is a passive reciprocal two-port,  $S_{21} = S_{12}$  by reciprocity and the diagonal terms are dominated by the same port reflections; thus, reporting  $S_{11}$  and  $S_{21}$  suffices and is standard practice in SE studies using waveguides/VNAs. We now cite representative sources and general standards to support these choices.

All EMI shielding measurements were conducted in a controlled laboratory environment at  $23 \pm 2$  °C and  $50 \pm 5\%$  relative humidity, with specimens conditioned under the same conditions for 48 h prior to testing. The vector network analyzer (VNA) was calibrated using a thru-reflect-line (TRL) standard provided with the WR-90 waveguide kit to eliminate systematic errors from cables, connectors, and fixtures. Calibration was repeated before each measurement series to ensure stability. By reporting these environmental controls and calibration

procedures, we provide greater transparency and confidence in the accuracy of the shielding effectiveness data.

The shielding effectiveness due to absorption ( $SE_A$ ) and reflection ( $SE_R$ ) was determined from the measured  $S$ -parameters using eqn (1) and (2), respectively:

$$SE_A(\text{dB}) = 10 \log_{10} \left[ \frac{1 - S_{11}^2}{S_{21}^2} \right] \quad (1)$$

$$SE_R(\text{dB}) = 10 \log_{10} \left[ \frac{1}{1 - S_{11}^2} \right] \quad (2)$$

The total shielding effectiveness ( $SE_T$ ) was then computed by summing the contributions from absorption and reflection, as shown in eqn (3).

$$SE_T = SE_A + SE_R \quad (3)$$

### 3. Results and discussion

#### 3.1. Morphology of $\text{Fe}_2\text{O}_3$

The morphology of the  $\text{Fe}_2\text{O}_3$  nanofillers was investigated using SEM at various magnifications, and the obtained images are presented in Fig. 4. Fig. 4(a) shows the SEM image of  $\text{Fe}_2\text{O}_3$  at  $25\,000\times$  and Fig. 4(b) at  $100\,000\times$  magnification. It is found that the  $\text{Fe}_2\text{O}_3$  nanofillers used in this study are a mixture of spherical and irregular morphologies.

At  $250\,00\times$  magnification (refer Fig. 4(a)), the  $\text{Fe}_2\text{O}_3$  nanoparticles exhibit a highly agglomerated and irregular cluster-like structure. The particles appear to form dense aggregates, likely due to strong van der Waals forces and magnetic interactions characteristic of iron oxide nanoparticles. The surface is rough, with visible granularity indicating the polycrystalline nature of  $\text{Fe}_2\text{O}_3$ . At  $100\,000\times$  magnification (refer Fig. 4(b)), the fine nanostructures within the agglomerates become clearer. The particles

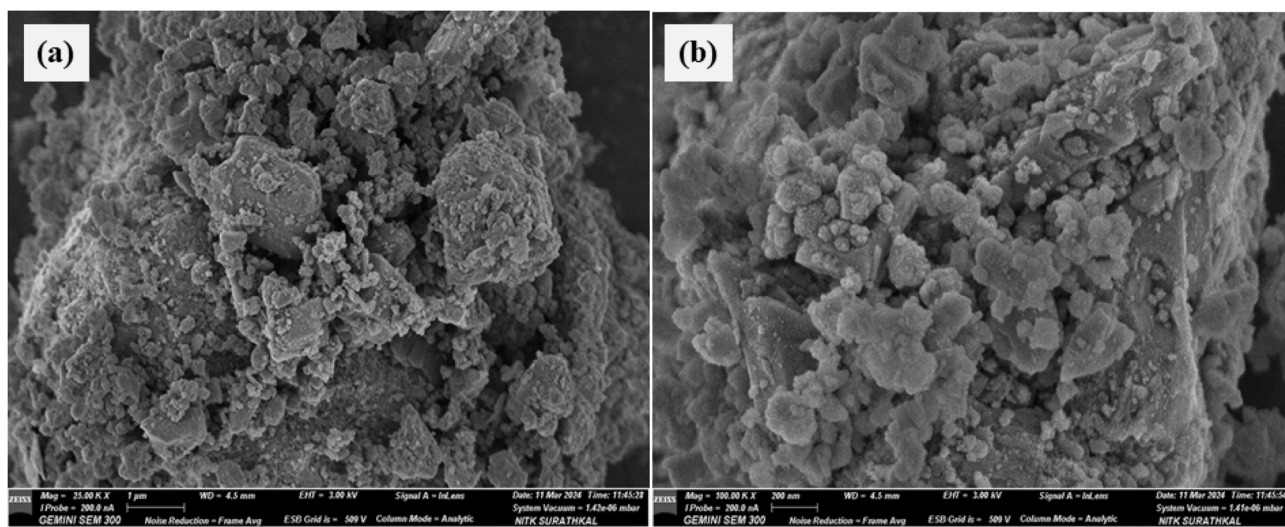


Fig. 4 FESEM images of  $\text{Fe}_2\text{O}_3$  nano filler at (a)  $25\,000\times$  and (b)  $100\,000\times$  magnification.



appear to have a roughly spherical to polyhedral shape, with an average size likely in the 20–80 nm range (based on visual approximation). The higher magnification reveals that these clusters are formed from much smaller primary nanoparticles.

The primary particle size (in Fig. 4(b)) seems to be in the nanometer range, but the secondary clusters (visible in Fig. 4(a)) are significantly larger, spanning micrometre scales. This hierarchical structure such as small primary nanoparticles forming larger agglomerates is typical for  $\text{Fe}_2\text{O}_3$  due to high surface energy and magnetic dipole–dipole interactions. The particle distribution appears non-uniform, with some densely packed regions and others showing voids or gaps, which may affect dispersion if used as a filler in composites.

### 3.2. XRD analysis of $\text{Fe}_2\text{O}_3$

Powder X-ray diffraction (XRD) was used to verify the phase and crystal structure of the  $\text{Fe}_2\text{O}_3$  nanoparticles employed in this study. The XRD pattern of the  $\text{Fe}_2\text{O}_3$  nanoparticles as received is shown in Fig. 5. The observed diffraction peaks at  $21.2^\circ$ ,  $33.4^\circ$ ,  $36.7^\circ$ ,  $43.3^\circ$ ,  $49.04^\circ$ ,  $53.3^\circ$ ,  $59.14^\circ$ ,  $62.7^\circ$ , and  $64.2^\circ$ , correspond to the characteristic reflections of hematite with a hexagonal crystal system (space group  $R\bar{3}c$ ), with lattice parameters of  $a = b = 5.034 \text{ \AA}$  and  $c = 13.748 \text{ \AA}$  (JCPDS no. 033-0664). As no additional diffraction peaks were identified, this also shows that the  $\text{Fe}_2\text{O}_3$  nanoparticles are of high purity, and this agrees with the stated 99% purity.

### 3.3. FTIR spectra

The FTIR spectrum of  $\text{Fe}_2\text{O}_3$ /carbon fabric/epoxy (CF/3FE/F) composite is compared with the FTIR spectra of the individual components  $\text{Fe}_2\text{O}_3$ , carbon fabric, and epoxy, respectively, as shown in Fig. 6. Table 3 presents the peak

assignments, and the respective functional groups present in the samples. One can see that the FTIR spectrum of the CF/3FE/F composite reveals the successful incorporation of the peaks from all three constituents without any substantial chemical degradation or change in the functional groups. The IR spectrum of the composites incorporates peaks originating from CF and  $\text{Fe}_2\text{O}_3$ , such as O–H, C–H, and Fe–O vibrations, indicating successful integration of the reinforcements in the composite. The absence of any significant peak shifts or new, strong bands indicates that the interaction of the constituents in the composite is mainly due to the physical interaction. However, the minor broadening and intensity variations at certain regions (*e.g.*,  $3380\text{--}3400 \text{ cm}^{-1}$  and  $1200\text{--}1500 \text{ cm}^{-1}$ ) imply the presence of interfacial hydrogen bonding and potential interactions between  $\text{Fe}_2\text{O}_3$  and the polymer matrix. These interactions between the epoxy- $\text{Fe}_2\text{O}_3$  can strengthen the epoxy-CF adhesion and can expect an enhancement in the mechanical properties as compared to composites without  $\text{Fe}_2\text{O}_3$  nanoparticles.

The observed bands in the  $470\text{--}580 \text{ cm}^{-1}$  region corroborate Fe–O vibrations of  $\alpha\text{-Fe}_2\text{O}_3$ , distinguishing physical adsorption from chemical bonding at the polymer–filler interface requires thermal or surface-energetic evidence. We therefore now frame the FTIR discussion as indicative rather than conclusive and note that DSC/TGA and contact-angle/XPS measurements would be appropriate to substantiate interfacial mechanisms in future work.

### 3.4. Tensile behaviour of composites

In this previous study, we presented tensile behaviour of pristine epoxy, which was reinforced with bi-directional carbon fabric composite (CF/E) and value as the control material. These

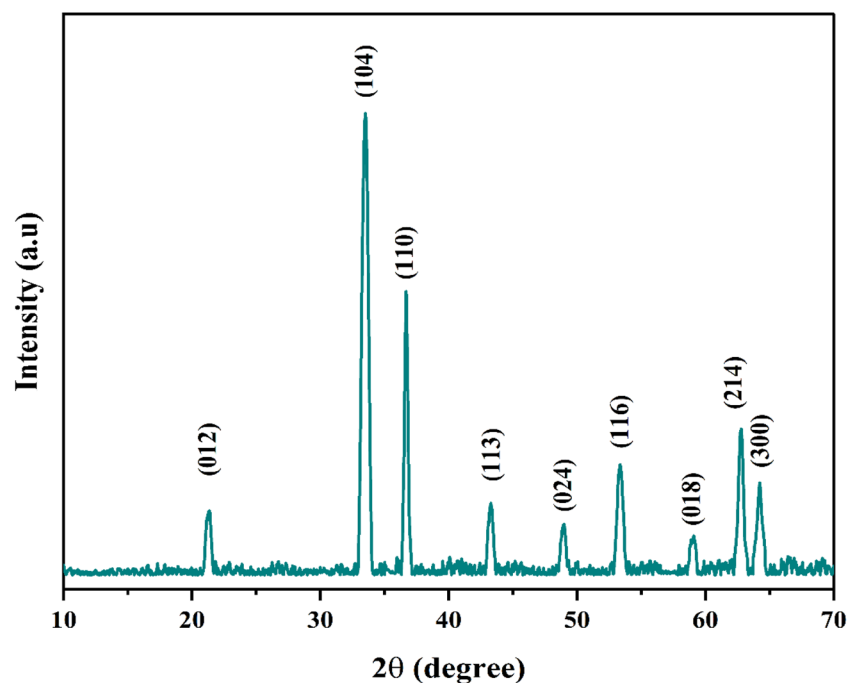


Fig. 5 X-RD spectra of  $\text{Fe}_2\text{O}_3$  nanoparticles.

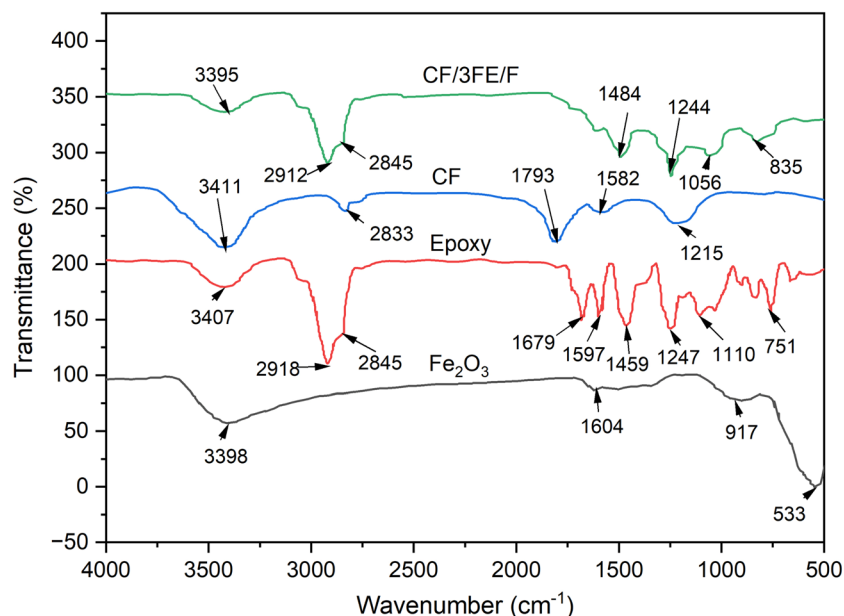


Fig. 6 FTIR spectra of  $\text{Fe}_2\text{O}_3$  nanoparticles, epoxy, carbon fabric and carbon fabric/ $\text{Fe}_2\text{O}_3$ /epoxy (CF/3FE/F) composite.

Table 3 FTIR peak assignments and functional group analysis

Wavenumber ( $\text{cm}^{-1}$ )	Functional group/vibration mode	Origin
3400	O-H stretching	$\text{Fe}_2\text{O}_3$ , CF, epoxy, composite
2912, 2845	C-H stretching	CF, epoxy, composite
1793	C=O stretching	CF
1679–1604	O-H bending/aromatic	$\text{Fe}_2\text{O}_3$ , epoxy
1582	C=C stretching	CF, composite
1484–1459	Aromatic C-H, ring	Epoxy, composite
1247–1244, 1215	C-O-C stretching (epoxy)	Epoxy, composite
1110–1056	C-O stretching	Epoxy, composite
917, 835	Fe-O vibration/aromatic ring	$\text{Fe}_2\text{O}_3$ , composite
751–533	Fe-O vibration/aromatic	$\text{Fe}_2\text{O}_3$ , epoxy, composite

results are presented here in relation to those of epoxy carbon fabric composites with 1, 2, or 3 wt%  $\text{Fe}_2\text{O}_3$  (CF/1FE/E, CF/2FE/E and CF/3FE/E), which are displayed in Fig. 7. The concentration of the loaded  $\text{Fe}_2\text{O}_3$  nanoparticles substantially influences the mechanical properties of the composites. At low concentrations of  $\text{Fe}_2\text{O}_3$  (1 wt%), tensile properties of the composite improved due to better interfacial bonding between the carbon fibres and epoxy matrix. When interfacial bonding strength improves, load transfer is improved and early failure of composites is delayed through the mechanism predominantly through interfacial hydrogen bonding as evidenced by FTIR. Nanoparticles also assist in uniform stress distribution within the polymer matrix that can potentially restrict crack initiation and propagation. At higher concentrations (3 wt%), filler agglomeration leads to a decline the tensile strength. Sun *et al.* reported that 1 wt% of  $\text{Fe}_2\text{O}_3$  nanoparticles yielded the best tensile properties in carbon fabric/epoxy composites, whereas higher concentrations resulted in diminished mechanical performance due to poor dispersion.<sup>22</sup> The present study shows a similar trend. Overall, adding  $\text{Fe}_2\text{O}_3$  nanoparticles to carbon fabric epoxy composites

generally increases tensile strength and modulus, with optimal results around 1–2 wt% nanoparticle loading. However, strain-to-failure and tensile strength slightly decrease at loadings above 2 wt% because of reduced ductility and agglomeration. At 3 wt%  $\text{Fe}_2\text{O}_3$ , agglomeration may occur, reducing reinforcement effectiveness and decreasing the composite's tensile properties.

Neat epoxy is highly flaw-sensitive under tensile loading, small variations in cure, micro voids, or surface defects act as critical flaws that trigger failure, producing wide specimen-to-specimen scatter. Also, processing related microstructure often differs between neat and filled systems that directly affect tensile strengths. Small amounts of nanoparticles can reduce effective void content by occupying free volume will increase if dispersion is poor. In these cases, both voids and partial cure reduce tensile strength and increase test-to-test variability. Similarly, well dispersed nanoparticles can stabilize the microstructure and narrow the scatter.  $\text{Fe}_2\text{O}_3$  additions can create competing mechanisms that make the filled curves look “trendless” unless dispersion is tightly controlled. At very low loadings, well-dispersed  $\text{Fe}_2\text{O}_3$  can increase tensile strength *via*



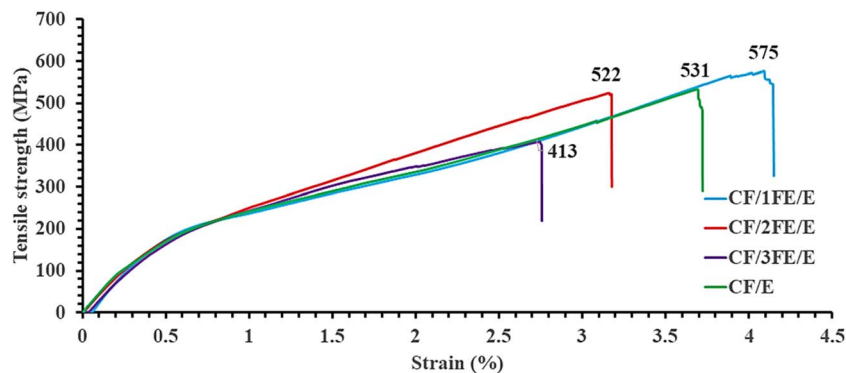


Fig. 7 Tensile behaviour of carbon fabric and  $\text{Fe}_2\text{O}_3$  reinforced epoxy (CF/FE/E) composites.

crack deflection/pinning and improved interfacial polarization-induced energy dissipation, at higher loadings, agglomerates behave as stress concentrators that depress strength and reintroduce scatter. Therefore, the trend depends upon the balance between dispersion-assisted toughening and agglomeration-induced embrittlement.

Referring to the tensile failure sample of CF/E composites previously reported by our group, which is shown in Fig. 8(a),<sup>23</sup> the tensile failure of CF/FE/E composite samples in this study is displayed in Fig. 8(b), (c), and (d). It is observed that the CF/E and CF/FE/E samples exhibited similar behaviour, with less delamination and good integrity of filler, fibre, and matrix, as

shown in Fig. 8(a) and (b). The sample with 2 wt%  $\text{Fe}_2\text{O}_3$  filler showed matrix cracking with delamination, as shown in Fig. 8(c), and this failure was more severe in the 3 wt% filler-loaded sample, as shown in Fig. 8(d). This indicates that at higher  $\text{Fe}_2\text{O}_3$  concentrations, the material failed due to brittle fracture of the matrix and increased delamination. In the current study, the failure analysis presented in Fig. 8 is qualitative, showing features such as river markings, fibre pull-out, and matrix cracking that are indicative of interfacial adhesion and filler-matrix interactions. While this provides useful insight, we acknowledge the limitation that no quantitative

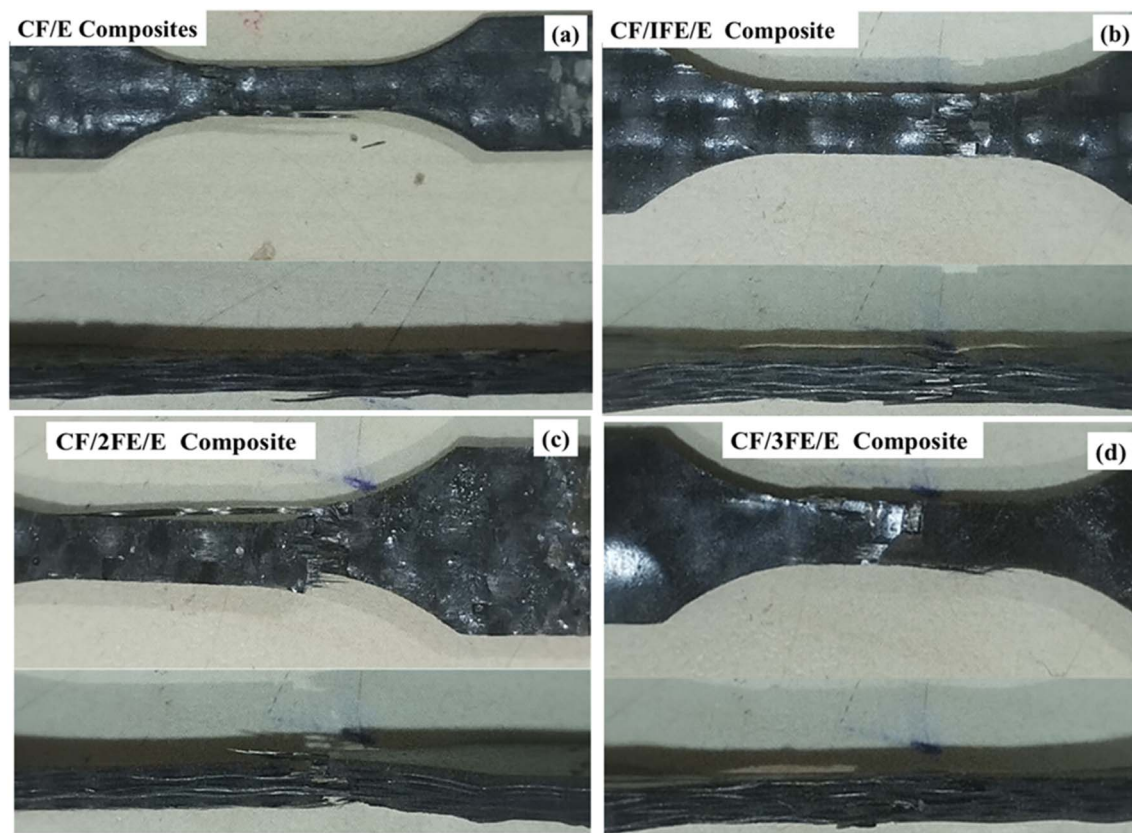


Fig. 8 Fractured tensile specimen of (a) CF/E composite, (b) CF/1FE/E composite, (c) CF/2FE/E composite, (d) CF/3FE/E composite.



fractographic metrics (e.g., crack propagation length, void density, or fibre-matrix debonding area) were extracted.

From the tensile stress *versus* strain plot, the Young's modulus of the neat CF/E composite is computed and found to be 26.83 GPa and a maximum tensile strength of 531 MPa. Incorporation of Fe<sub>2</sub>O<sub>3</sub> significantly influenced stiffness, with a maximum modulus of 35.20 GPa observed for CF/2FE/E, representing a ~31% increase over the neat composite shown in Fig. 9. This enhancement is attributed to improved stress transfer due to strong interfacial adhesion between the Fe<sub>2</sub>O<sub>3</sub>-modified epoxy matrix and the CF reinforcement, consistent with findings by Shukla (2019) and Ahmad *et al.* (2021), where optimal nanofiller content improved modulus *via* synergistic reinforcement mechanisms.<sup>4,6</sup> In terms of tensile strength, the highest value (575 MPa) was recorded for CF/1FE/E, marginally surpassing the neat composite. However, strength declined at higher loadings, with CF/3FE/E showing the lowest value (413 MPa). This reduction is attributed to nanoparticle agglomeration, void formation, and reduced fibre-matrix interfacial integrity at excessive filler content, as similarly reported in studies on nanoparticle-reinforced polymer composites.<sup>8,24</sup> Overall, the results indicate that moderate filler addition (1–2 wt%) optimizes stiffness, while excessive loading compromises tensile strength despite maintaining high modulus. This trend aligns with literature highlighting the balance between dispersion quality, interfacial bonding, and filler content in determining mechanical performance.<sup>24,25</sup>

## 4. Electromagnetic shielding effectiveness

The electromagnetic interference shielding effectiveness (EMI SE) of a material represents its capacity to attenuate or block incident electromagnetic waves. The calculated values of SE<sub>A</sub>, SE<sub>R</sub> and SE<sub>T</sub> for CF/1FE/E, CF/2FE/E, and CF/3FE/E composite laminates in single-layer, double-layer, and four-layer configurations were plotted over the 8–12 GHz frequency range. The

corresponding results are presented in Fig. 10(a–c), 11(a–c), and 12(a–c), respectively.

From previous studies on CF/E composites in the X-band, we determined that the SE<sub>A</sub> of the single- and double-layer samples remained relatively stable throughout the frequency range, while the four-layer samples exhibited serious frequency-dependent variability in SE<sub>A</sub> due to the skin effect. The skin depth, defined as the extent to which an electromagnetic wave will penetrate a material, is inversely proportional to the square root of frequency ( $f$ ) which is displayed in (4). Note that  $\delta$  represents the skin depth,  $\mu$  is magnetic permeability (H m<sup>−1</sup>),  $\sigma$  is electrical conductivity (S m<sup>−1</sup>), and  $f$  is the frequency of the impinging electromagnetic field.

$$\delta = \sqrt{\frac{1}{\pi\mu\sigma f}} \quad (4)$$

According to eqn (4), as the skin depth  $\delta$  decreases as frequency increases, means that, at higher frequencies, EM waves tend to travel along the surface rather than penetrate the material. At lower frequencies, larger  $\delta$  allows for deeper penetration and greater absorption, which explains the superior low-frequency performance of four-layer samples. Increasing the number of layers enhances absorption-dominated shielding, as multiple carbon fabric layers better attenuate EM waves through successive absorption and reduced reflection.

Fig. 10(a–c) presents the SE<sub>A</sub>, SE<sub>R</sub>, and SE<sub>T</sub> of CF/1FE/E composites with single-, double-, and four-layer structures in the X-band region. As depicted in Fig. 10(d), the SE<sub>A</sub> contribution increases from 61.2% to 67.6% with the transition from a single to a two-layer configuration. A comparable trend in SE<sub>A</sub> and SE<sub>R</sub> is observed for the two- and four-layer samples. Notably, the SE<sub>R</sub> of the CF/1FE/E composite is lower than that of the CF/E system, which can be attributed to the enhanced absorption of EM radiation by the Fe<sub>2</sub>O<sub>3</sub> fillers. Therefore, incorporation of Fe<sub>2</sub>O<sub>3</sub> promotes absorption-dominated shielding effectiveness in the composite.

SE<sub>A</sub>, SE<sub>R</sub>, and SE<sub>T</sub> of CF/2GNP/E composites with one-, two-, and four-layer structures in the X-band are shown in Fig. 11(a–c), and the percentage SE is illustrated in Fig. 11(d).

CF/2FE/E composites demonstrate absorption-dominated EMI shielding in all configurations, with SE<sub>T</sub> and SE<sub>A</sub> values increasing as the number of layers increases. The four-layer laminate achieves the highest shielding effectiveness (~22 dB), confirming the benefit of layer stacking in enhancing performance. SE<sub>A</sub> contributes more than 66% to the overall shielding across all samples, indicating that absorption is the primary shielding mechanism in the X-band region.

In the X-band region, the SE<sub>A</sub>, SE<sub>R</sub>, and SE<sub>T</sub> values of the CF/3FE/E composites with single, double-, and four-layer configurations are represented in Fig. 12(a–c) and the corresponding percentage SE is given in Fig. 12(d). The presence of 3 wt% Fe<sub>2</sub>O<sub>3</sub> as a filler increased the absorption-related shielding effectiveness due to a more efficient conductive network formed in the epoxy; in addition, multilayered structures also augmented absorption-dominated shielding. The two-layer CF/

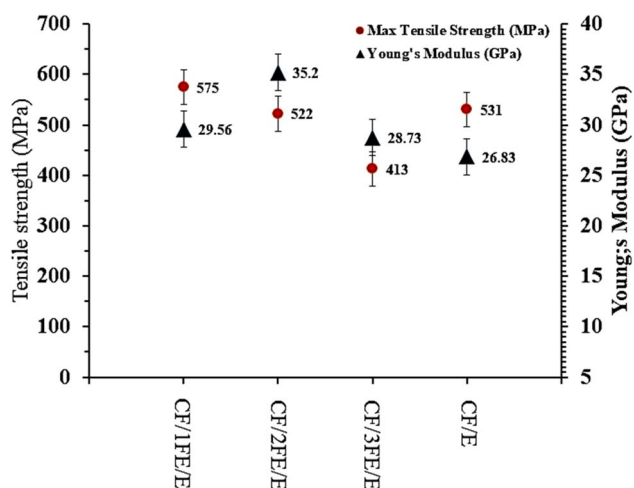


Fig. 9 Comparison of tensile strength and Young's modulus of CF/E with CF/FE/E composites.



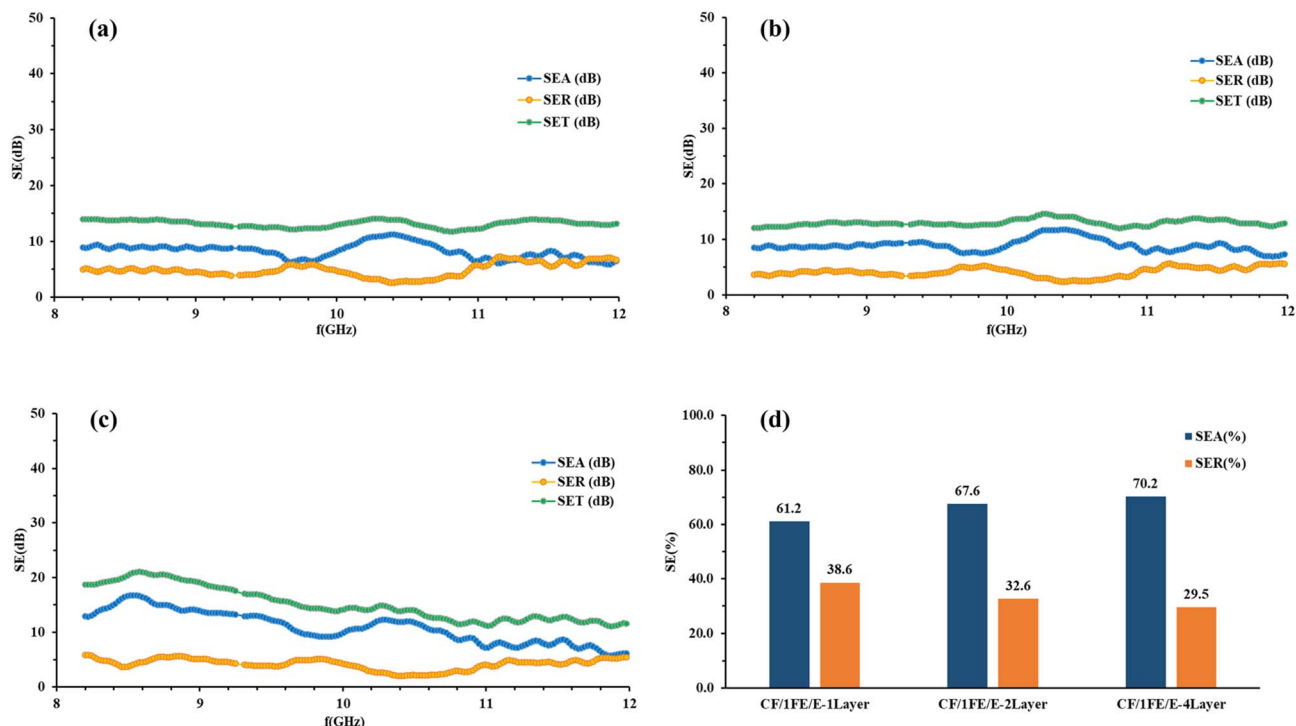


Fig. 10  $SE_A$ ,  $SE_R$ , and  $SE_T$  of (a) CF/1FE/E single-layer, (b) CF/1FE/E double-layer, (c) CF/1FE/E four-layer composites, and (d) the corresponding  $SE_A$  and  $SE_R$  percentage contributions for all CF/1FE/E samples.

3FE/E demonstrably the best overall shielding effectiveness at 25.3 dB among the tested samples, which indicates significant potential for practical use in the X-band frequency range.

The shielding effectiveness and mechanisms in case of GNP-based epoxy composites is that graphene nanoplatelets contribute primarily through high electrical conductivity and the formation of percolated conductive networks. In this case

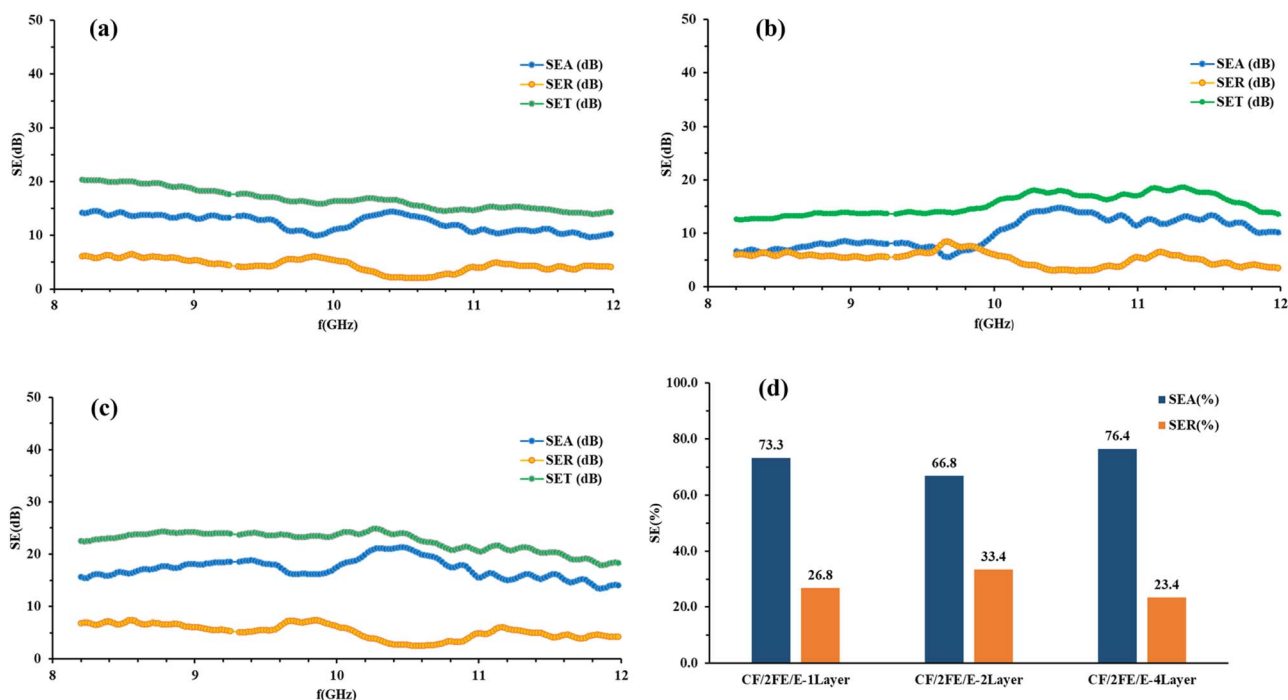


Fig. 11  $SE_A$ ,  $SE_R$ , and  $SE_T$  of (a) CF/2FE/E single-layer, (b) CF/2FE/E double-layer, (c) CF/2FE/E four-layer composites, and (d) the corresponding  $SE_A$  and  $SE_R$  percentage contributions for all CF/2FE/E samples.

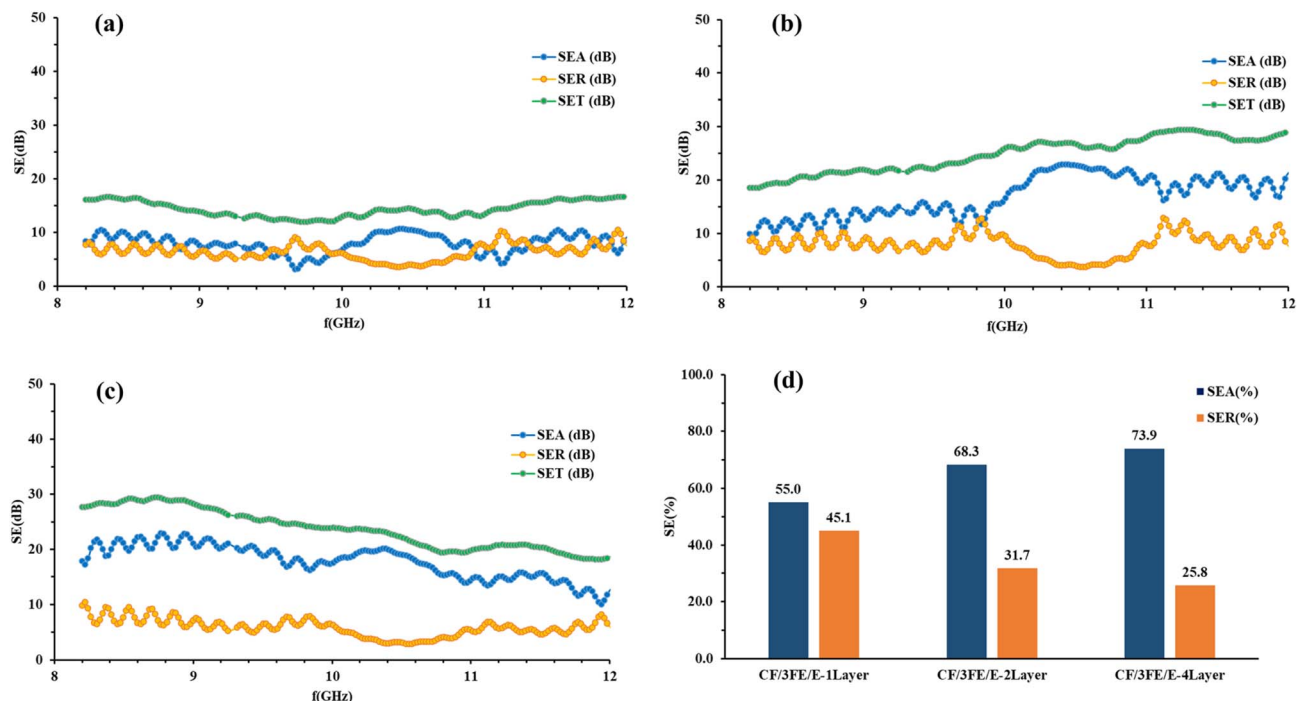


Fig. 12 SE<sub>A</sub>, SE<sub>R</sub>, and SE<sub>T</sub> of (a) CF/3FE/E single-layer, (b) CF/3FE/E double-layer, (c) CF/3FE/E four-layer composites, and (d) the corresponding SE<sub>A</sub> and SE<sub>R</sub> percentage contributions for all CF/3FE/E samples.

the dominant shielding mechanism is reflection due to impedance mismatch and is supplemented by some absorption *via* multiple scattering inside the conductive network. So, these systems achieve relatively high SE values at moderate loadings. In case of carbon fabric with Fe<sub>2</sub>O<sub>3</sub> nanoparticle-epoxy composites, carbon fibers provide a long range continuous conductive pathways and mechanical reinforcement. Also, Fe<sub>2</sub>O<sub>3</sub> nanoparticles introduce magnetic loss (natural resonance, eddy current loss) and enhance interfacial polarization. Thus, shielding arises from a synergistic combination of

reflection from conductive carbon fabric, absorption from Fe<sub>2</sub>O<sub>3</sub>-induced magnetic/dielectric losses, and multiple scattering due to heterogeneous interfaces. So, compared with GNP alone, this dual-action usually yields higher absorption-dominated shielding, which is desirable in applications where weight reduction, structural integrity and EMI performance must be simultaneously achieved.

Fig. 13 illustrates the comparative analysis of CF/1FE/E, CF/2FE/E, and CF/3FE/E composites across single-, double-, and four-layer configurations, revealing a clear trend of increasing

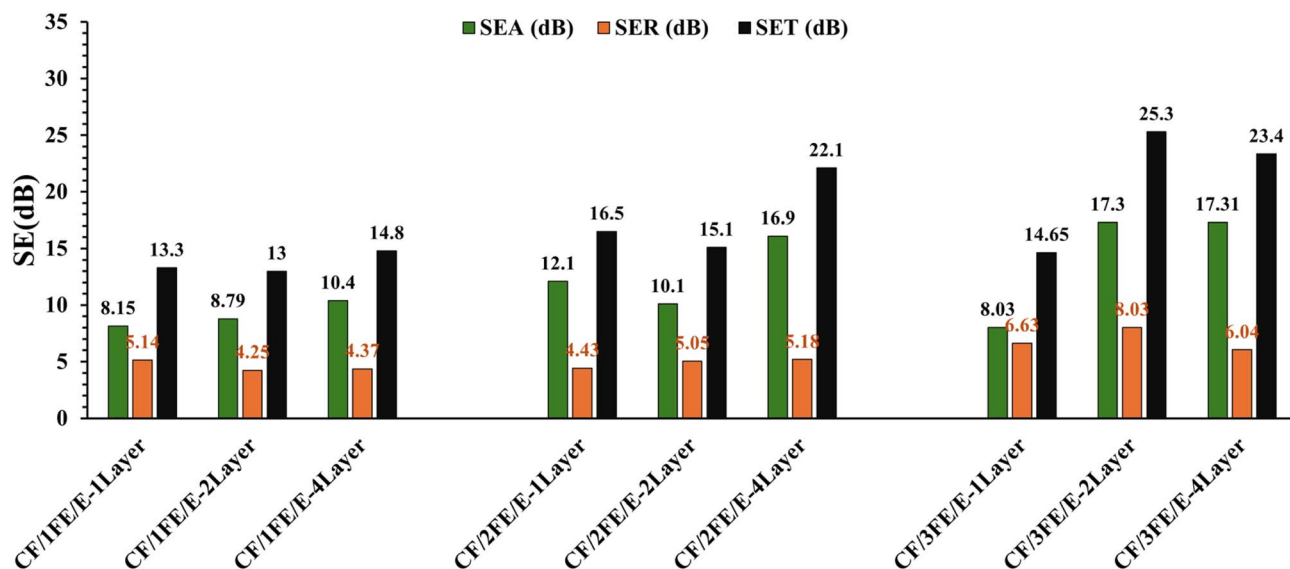


Fig. 13 SE<sub>A</sub>, SE<sub>R</sub> and SE<sub>T</sub> of CF/FE/E composites.

Table 4 Comparison of the EMI SE of various carbon fabric composites from previous studies and the present work

Matrix, primary reinforcement & secondary reinforcement			SE							
Fibre/fabric/matrix	Filler	Concentration (wt%)	Process	N	t (mm)	$\sigma$ (S cm <sup>-1</sup> )	SE <sub>A</sub> (dB)	SE <sub>R</sub> (dB)	SE <sub>T</sub> (dB)	Ref.
T800-CF C fabric-plain weave PTFE + CNT dual NF	MWCNT + Fe <sub>2</sub> O <sub>3</sub>	10	Prepreg + compression molding	20	3.0	35.0	~50	~10	59.0	6
	Fe <sub>3</sub> O <sub>4</sub>	60	Solution casting + hot pressing	1	1.54	1.17	~47.1	~3.8	~58.93	7
	Fe <sub>3</sub> O <sub>4</sub> + MXene	50 + 70	Shear-induced <i>in situ</i> fibrillation + vacuum filtration	2	0.0849	4.0278	~24.8	~19.8	44.56	8
	Fe <sub>2</sub> O <sub>3</sub>	<10	Matrix-directed mineralisation + lamination	~100	<0.001	—	~21–25	~4–7	~27–29	9
rGCE-aerogel TPU PA11/PA + SCF	Fe <sub>3</sub> O <sub>4</sub>	10	<i>In situ</i> chemical precipitation	—	2.5	—	—	—	~58.13	10
	Graphite + CoFe <sub>2</sub> O <sub>4</sub>	40 + 40	Melt mixing + compression molding	3 (G/F/G)	2.4	—	—	—	54	11
	—	30	Extrusion + compression molding	1	1	~2.28 × 10 <sup>-5</sup>	~22.7	~5.3	28	12
	Gp + Fe <sub>2</sub> O <sub>3</sub>	4 + 6	Hand lay-up followed by compression molding)	16 [0/90] <sub>8</sub>	2.5	G-4 F-6	~90.5	~5	95.57	13
UDCF/E prepreg (T800, 12 K) C fabric/E	GO + Fe <sub>3</sub> O <sub>4</sub>	8	Layer-by-layer deposition + hot press curing	8	~1.35	~3.6	22.8	10.7	32.9	14
Epoxy resin Epoxy resin Cotton fibre/epoxy C fabric-plain weave	Fe <sub>3</sub> O <sub>4</sub> + CB	15 + 50	Mixing + casting	—	0.7	—	11.85	~17.6	36.6	15
	GNP + Fe <sub>3</sub> O <sub>4</sub>	12 + 4	Melt blending	1	1	—	—	—	~7	16
	Fe <sub>3</sub> O <sub>4</sub> decorated GNP	12 + 4	Epoxy impregnation + hot press	2	2.68	—	~28.1	~5.0	33.1	17
	Fe <sub>3</sub> O <sub>4</sub> + GNP	5 + 30	Hand lay-up method	1	—	—	7.8	6.0	13.9	23
Present study	Fe <sub>2</sub> O <sub>3</sub> /CF	1	Hand lay-up method	2	—	—	10.2	8.8	19	
		4		12.6	8.0	20.6				
		1		8.15	5.14	13.3				
		2		8.79	4.25	13				
	2	4	10.4	4.37	14.8					
		1	12.1	4.43	16.5					
		2	10.1	5.05	15.1					
		4	16.9	5.18	22.1					
	3	1	8.03	6.63	14.6					
		2	17.3	8.03	25.3					
		4	17.3	6.04	23.4					



total shielding effectiveness ( $SE_T$ ) with higher filler content and additional layers. Among all, CF/3FE/E-2Layer exhibits the maximum  $SE_T$  of 25.3 dB, closely followed by CF/3FE/E-4Layer (23.4 dB).  $SE_A$  consistently dominates over  $SE_R$  in all samples, indicating absorption as the primary EMI shielding mechanism, enhanced by  $Fe_2O_3$ -induced magnetic losses and carbon fabric conductivity. Stacking layers boosts  $SE_A$  via repeated internal reflections; however, the higher  $Fe_2O_3$  content strengthens dielectric and magnetic losses, resulting in excellent X-band shielding performance. In aerospace and defence sectors, EMI shielding materials are generally expected to achieve  $\geq 30$  dB attenuation, which corresponds to  $\sim 99.9\%$  reduction of incident electromagnetic radiation across the relevant frequency ranges (typically 8–12 GHz or even broader). For critical avionics enclosures or military-grade applications, the requirement is even stricter, often  $\geq 60$ –80 dB to protect sensitive electronic systems and ensure electromagnetic compatibility (EMC). Civil aircraft structural composites are usually benchmarked against the 30 dB threshold, balancing performance with weight reduction and cost. The 25.3 dB  $SE_T$  value reported here corresponds to  $\sim 99.7\%$  attenuation. While this is a strong result for a lab-developed epoxy composite with  $Fe_2O_3$  nanoparticles and carbon fabric, it falls slightly below the minimum 30 dB level generally recommended for aerospace-grade shielding panels. This study achieves near-threshold performance with a structural composite (carbon fabric reinforced epoxy +  $Fe_2O_3$ ) is significant because it demonstrates dual functionality of mechanical integrity and EMI shielding which is a crucial step toward lightweight multifunctional aerospace materials. Table 4 compares the EMI-SE of various carbon fabric composites reported previously and in the present study. The present study reports a maximum SE of 25.3 dB, this value should be interpreted relative to filler loading, laminate thickness, and electrical conductivity. Unlike several literature examples where high SE values ( $>50$  dB) are achieved at substantially higher filler concentrations (e.g., 40–60 wt%  $Fe_3O_4$ , CNT/graphite systems, or dual nanofillers) and thicker laminates ( $>20$ –100 layers), our design intentionally maintains low filler content (1–4 wt%  $Fe_2O_3$ ) and relatively thin laminates (1–3 layers,  $\sim 2$ –3 mm thickness) to balance mechanical integrity with processability. Consequently, the lower conductivity of our composites compared to highly conductive matrices (e.g., MXene- or CNT-based systems) inherently limits SE, as predicted by classical shielding theory where  $SE \propto \sigma^{0.5} \times t$ . However, the present study demonstrates that even with minimal filler loading and simpler hand lay-up processing, SE values in the 13–25 dB range are achievable, which is significant for lightweight EMI shielding applications. Furthermore, our results are consistent with other low-filler, low-thickness  $Fe_2O_3$ -based systems, which typically yield SE values in the 20–30 dB range, underscoring the material's comparative efficiency.

The improvement in tensile strength observed at moderate  $Fe_2O_3$  loadings reflects effective stress transfer at the fiber-matrix interface, which simultaneously promotes the formation of continuous conductive or semi-conductive networks essential for EMI attenuation. Conversely, the reduction in mechanical properties at higher filler contents can be attributed to

agglomeration and interfacial defects, which not only act as stress concentrators but also disrupt the uniformity of the shielding pathways, thereby limiting shielding effectiveness despite increased filler volume. Quantitatively, this trade-off can be rationalized by considering that both load transfer efficiency ( $\sigma \propto V_f \times \tau_i$ , where  $\tau_i$  is interfacial shear strength) and shielding effectiveness ( $SE \propto \sigma^{0.5} \times t$ , where  $\sigma$  is conductivity and  $t$  is thickness) are maximized under conditions of uniform dispersion and strong interfacial bonding. By correlating tensile strength retention with SE trends across different filler loadings, we demonstrate that optimal performance arises not simply from increasing filler concentration, but from achieving synergistic dispersion and interfacial compatibility. This mechanistic perspective situates our findings within broader composite design principles and highlights pathways for future optimization (e.g., surface functionalization of  $Fe_2O_3$ , hybrid filler strategies, or controlled laminate stacking) to simultaneously advance mechanical robustness and EMI shielding efficiency.

## 5. Conclusions

This work established that bidirectional carbon fabric/epoxy composites reinforced with  $Fe_2O_3$  nanoparticles can simultaneously deliver mechanical robustness and effective electromagnetic shielding in the X-band frequency range. Optimal filler loading of 1–2 wt% improved tensile strength and stiffness through enhanced fiber-matrix interfacial bonding, while higher loading led to agglomeration and reduced strength. All composites exhibited absorption-dominated EMI shielding, with a maximum  $SE_T$  of 25.3 dB achieved in a two-layer laminate containing 3 wt%  $Fe_2O_3$ . The findings demonstrate that careful control of nanofiller concentration and laminate configuration enables the design of lightweight, multifunctional composites suitable for aerospace applications requiring structural integrity and electromagnetic compatibility.

The carbon fabric/ $Fe_2O_3$  nanoparticle/epoxy (CF/3FE/E-Layer) composite developed in this study shows multifunctional characteristics as good mechanical integrity from carbon fabric, magnetic and dielectric loss from  $Fe_2O_3$ , and structural reliability from epoxy. While the aerospace sector is an obvious target, several other industries could benefit from such composites. Which included automotive and electric vehicles (EV) with increasing use of sensors, radar, and advanced driver-assistance systems (ADAS), EMI shielding is crucial for safety and reliability. Also, the composites can replace heavier metallic shielding, lower vehicle weight and improve fuel efficiency. These composites can also be used in EMI-safe enclosures for EV batteries and control units, reducing electromagnetic crosstalk. The carbon fabric/ $Fe_2O_3$  nanoparticle/epoxy (CF/3FE/E-Layer) composite developed in this study also can be used in structural panels for antenna housings, base stations, and smart towers where both strength and EMI shielding are needed. Also, it can be used in laptops, tablets, and high-frequency devices that benefit from lightweight shielding to prevent interference.



To further improve the shielding effectiveness ( $SE_T$ ) beyond the current 25.3 dB, future work should focus on multilayer or gradient architectures that enhance multiple reflections, hybridization of  $Fe_2O_3$  with conductive 2D fillers such as graphene or MXenes to combine magnetic and conductive losses, and surface functionalization of  $Fe_2O_3$  to improve dispersion and interfacial polarization. In addition, thickness optimization guided by skin-depth calculations and the introduction of porous or foam-like sublayers can significantly increase absorption without adding excessive weight. Together, these strategies are expected to raise  $SE_T$  above the aerospace benchmark of 30 dB and broaden the applicability of the composites to advanced structural and electronic shielding applications.

## Conflicts of interest

There are no conflicts of interest to declare.

## Data availability

Data supporting the results reported in this manuscript are included in this article.

## References

- 1 R. B. J. Chandra, B. Shivamurthy, S. D. Kulkarni and M. S. Kumar, Hybrid polymer composites for EMI shielding application – a review, *Mater. Res. Express*, 2004, **6**(8), 1–29, DOI: [10.1088/2053-1591/aaff00](#).
- 2 S. L. Stupar, M. M. Vuksanovic, D. Ž. Mijin, B. C. Milanovic, V. J. Joksimovic, T. S. Barudzija and M. R. Knezevic, Multispectral electromagnetic shielding and mechanical properties of carbon fabrics reinforced by silver deposition, *Mater. Chem. Phys.*, 2022, **289**, 126495, DOI: [10.1016/j.matchemphys.2022.126495](#).
- 3 D. D. L. Chung, Electromagnetic interference shielding effectiveness of carbon materials, *Carbon*, 2001, **39**(2), 279–285, DOI: [10.1016/S0008-6223\(00\)00184-6](#).
- 4 V. Shukla, Review of electromagnetic interference shielding materials fabricated by iron ingredients, *Nanoscale Adv.*, 2019, **1**(7), 1640–1671, DOI: [10.1039/c9na00108e](#).
- 5 M. S. Hareesh, P. Joseph and S. George, Electromagnetic interference shielding: a comprehensive review of materials, mechanisms, and applications, *Nanoscale Adv.*, 2025, **5**, 4510, DOI: [10.1039/d5na00240k](#).
- 6 J. Ahmad, A. Maryam, H. S. Manzoor, N. Nasir, Y. Nawab and H. S. Ahmad, Enhancement of EMI shielding effectiveness in carbon fiber-reinforced composite structures impregnated with MWCNT and  $Fe_2O_3$  nanofillers through optimized laminating sequences, *Mater. Sci. Eng., B*, 2025, **313**, 117980, DOI: [10.1016/j.mseb.2025.117980](#).
- 7 M. Waseem, X. Zhang, W. Wang, S. Ameen, K. Li, X. Guo and J. Chen, Lightweight and high EMI shielding performed  $ZrB_2$ -SiC/CF@Epoxy- $Fe_3O_4$  composites with carbonated cotton fibers, *J. Alloys Compd.*, 2025, **1030**, 180915, DOI: [10.1016/j.jallcom.2024.180915](#).
- 8 R. Shao, G. Wang, J. Chai, J. Lin, G. Zhao, Z. Zeng and G. Wang, Multifunctional Janus-structured polytetrafluoroethylene-carbon nanotube- $Fe_3O_4$ /MXene membranes for enhanced EMI shielding and thermal management, *Nano-Micro Lett.*, 2025, **17**, 136, DOI: [10.1007/s40820-025-01647-x](#).
- 9 C. X. Yu, Y. F. Meng, B. Yang, J. Pang, X. S. Meng, Z. Y. Zhao, Q. Y. Wang, L. B. Mao, Z. K. Wu and S. H. Yu, Iron oxide/CNT-based artificial nacre for electromagnetic interference shielding, *Nano Res.*, 2024, **17**(7), 6560–6566, DOI: [10.1007/s12274-024-6567-7](#).
- 10 L. Zhang, G. Song, Y. Song, J. Li, Z. Li, X. Yang, Z. Li, X. Li and Z. Jia, Ultralight 3D cross-linked reinforced graphene@ $Fe_3O_4$  composite aerogels for electromagnetic wave absorption, *Mater. Res. Bull.*, 2024, **173**, 112696, DOI: [10.1016/j.materresbull.2024.112696](#).
- 11 N. H. Wilson, A. Masař, M. Machovsky, D. Skoda, P. Urbanek, M. Urbanek, M. Poschl, J. Vilcakova, I. Kuritka and R. S. Yadav, Excellent electromagnetic interference shielding of multi-layered thermoplastic polyurethane nanocomposites with  $CoFe_2O_4$  nanoparticles and graphite, *Adv. Compos. Hybrid Mater.*, 2025, **8**, 64, DOI: [10.1007/s42114-024-01155-3](#).
- 12 B. U. Durmaz, A. O. Salman and A. Aytac, Electromagnetic interference shielding performances of carbon-fiber-reinforced PA11/PLA composites in the X-band frequency range, *ACS Omega*, 2023, **8**(25), 22762–22773, DOI: [10.1021/acsomega.3c01656](#).
- 13 H. S. Ahmad, T. Hussain, Y. Nawab and H. Awais, Graphene and  $Fe_3O_4$  filled composites for mitigation of electromagnetic pollution and protection of electronic appliances, *Compos. Sci. Technol.*, 2023, **240**, 110097, DOI: [10.1016/j.compscitech.2023.110097](#).
- 14 N. Duan, Z. Shi, J. Wang, X. Zhang, C. Zhang, C. Zhang and G. Wang, Multi-layer-structured carbon fiber fabric/graphene oxide/ $Fe_3O_4$ /epoxy composite for highly efficient mechanical and electromagnetic interference shielding, *Appl. Surf. Sci.*, 2023, **613**, 156038, DOI: [10.1016/j.apsusc.2022.156038](#).
- 15 R. Fallah, S. Hosseinabadi and G. Pourtaghi, Influence of  $Fe_3O_4$  and carbon black on the enhanced electromagnetic interference (EMI) shielding effectiveness in the epoxy resin matrix, *J. Environ. Health Sci. Eng.*, 2022, **20**, 113–122, DOI: [10.1007/s40201-021-00759-x](#).
- 16 K. Bhaskaran, R. K. Bheema and K. C. Etika, The influence of  $Fe_3O_4$ @GNP hybrids on enhancing the EMI shielding effectiveness of epoxy composites in the X-band, *Synth. Met.*, 2020, **265**, 116374, DOI: [10.1016/j.synthmet.2020.116374](#).
- 17 J. Xu, R. Chen, Z. Yun, Z. Bai, K. Li, S. Shi, J. Hou, X. Guo, X. Zhang and J. Chen, Lightweight epoxy/cotton fiber-based nanocomposites with carbon and  $Fe_3O_4$  for electromagnetic interference shielding, *ACS Omega*, 2022, **7**(15), 15215–15222, DOI: [10.1021/acsomega.2c01293](#).
- 18 N. Veeramani, A. M. D. James, A. N. Alapati, A. Bolluk and D. Ray, Enhancing fracture toughness of carbon fiber/



- epoxy composites using polyphenylene ether as a modifier, *J. Appl. Polym. Sci.*, 2024, **141**, 1–11, DOI: [10.1002/app.55388](https://doi.org/10.1002/app.55388).
- 19 N. Veeramani, D. R. Kumar, N. T. Manikandanath, A. S. Ganesh, S. Siju and G. Siju, Investigation of microcapsules based self-healing composites embedded with carbon nanotubes for improved healing efficiency, *J. Polym. Res.*, 2024, **31**(312), 1–11, DOI: [10.1007/s10965-024-04155-5](https://doi.org/10.1007/s10965-024-04155-5).
  - 20 T. Y. Eken, C. Kaykilarli, B. Kucukelyas and M. B. Tabakcioglu, Electromagnetic Shielding Effectiveness and Impact Test Performance of Carbon Fiber Reinforced Polymer Composites with Hematite and Goethite, *Macromol. Mater. Eng.*, 2023, **309**, 2300271, DOI: [10.1002/mame.202300271](https://doi.org/10.1002/mame.202300271).
  - 21 S. A. Kumar and S. N. Yeole, Tensile testing and evaluation of 3D-printed PLA specimens as per ASTM D638 type-IV standard, in *Innovative Design and Development Practices in Aerospace and Automotive Engineering (IDAD 2018), Lecture Notes in Mechanical Engineering*, ed. V. Sridhar, M. Vasudevan and V. Mohanavel, Singapore, Springer, 2018, pp. 79–95, DOI: [10.1007/978-981-13-2718-6\\_9](https://doi.org/10.1007/978-981-13-2718-6_9).
  - 22 T. Sun, Y. Wang, Y. Yang, H. Fan, M. Liu and Z. Wu, A novel Fe<sub>2</sub>O<sub>3</sub>@APFS/epoxy composite with enhanced mechanical and thermal properties, *Compos. Sci. Technol.*, 2020, **193**, 108146, DOI: [10.1016/j.compscitech.2020.108146](https://doi.org/10.1016/j.compscitech.2020.108146).
  - 23 R. Suresha, H. K. Sachidananda, B. Shivamurthy, S. Kumar, N. Kumar and S. Parasuram, Mechanical properties and electromagnetic shielding effectiveness of carbon fabric/GNP reinforced epoxy composites, *Sci. Rep.*, 2025, **15**, 15735, DOI: [10.1038/s41598-025-00634-x](https://doi.org/10.1038/s41598-025-00634-x).
  - 24 B. Le, J. Khaliq, D. Huo, X. Teng and I. Shyha, A review on nanocomposites. Part 1: Mechanical properties, *J. Manuf. Sci. Eng.*, 2020, **142**(10), 100801, DOI: [10.1115/1.4047047](https://doi.org/10.1115/1.4047047).
  - 25 A. Manohar, T. Suvarna, S. Sangaraju, A. A. Almunyif and K. H. Kim, Enhancing electromagnetic interference shielding: The role and advances of ferromagnetic materials, *J. Alloys Compd.*, 2025, **1032**, 180988, DOI: [10.1016/j.jallcom.2025.180988](https://doi.org/10.1016/j.jallcom.2025.180988).

