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# One-step *in situ* synthesis of strontium ferrites and strontium ferrites/graphene composites as microwave absorbing materials†

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Strontium ferrites/graphene (SrFe<sub>12</sub>O<sub>19</sub>/G) composites were synthesized by a facile one-step surfactant assisted hydrothermal method. The structure, magnetic properties and electromagnetic parameters of the as-prepared composites were evaluated. Uniform SrFe<sub>12</sub>O<sub>19</sub> nanoparticles with a diameter of 60–70 nm were well distributed on the graphene. The formation of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles and the reduction of graphene oxide were accomplished simultaneously. The re-stacking of graphene sheets and the aggregation of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles were inhibited. The as-obtained composites exhibited better absorbing properties than the pristine SrFe<sub>12</sub>O<sub>19</sub> particles due to the synergistic effect of SrFe<sub>12</sub>O<sub>19</sub> and graphene. The minimum reflection loss (RL) of SrFe<sub>12</sub>O<sub>19</sub>/G composites with the thickness of 3 mm reached –34.8 dB at 13.6 GHz, and the effective absorption bandwidth (RL < –10 dB) was 5.7 GHz. However, the minimum RL of the SrFe<sub>12</sub>O<sub>19</sub> particles only reached –13.5 dB, and the effective absorption bandwidth was 4.6 GHz. A possible mechanism for the enhanced absorbing performance was proposed.

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## Introduction

The wide application of electromagnetic technology facilitates people's work and life. However, the electromagnetic pollution which results from the overuse of electromagnetic technology has become a new form of environmental pollution in addition to those such as noise, waste water, exhaust gas and solid waste. Over the last few decades, researchers have been trying to find effective ways to prevent electromagnetic pollution and protect information security and human health.<sup>1</sup> Consequently, absorbing materials have emerged, which can absorb the incident electromagnetic waves and attenuate them in the form of heat.<sup>2</sup> The traditional absorbing materials cannot meet all of the requirements, including tiny thickness, light weight, broad bandwidth and strong absorption. Therefore, combining two or more kinds of functional materials is widely regarded as a feasible way to improve the absorbing performances.<sup>3,4</sup>

As a kind of useful electromagnetic functional material, ferrites have aroused enormous research interests in many fields, with applications such as magneto-optical devices, magnetic memories, sensitive elements, catalysts, biology, absorbing materials and so on.<sup>5–11</sup> As a typical hard magnetic material discovered in 1950s, strontium ferrite (SrFe<sub>12</sub>O<sub>19</sub>) is of

great importance due to its low cost production, high saturation magnetization and good chemical stability.<sup>12</sup>

A number of different methods, including co-precipitation method,<sup>13</sup> high temperature calcinations,<sup>14</sup> sol-gel method<sup>15</sup> and hydrothermal reaction followed by post-calcination,<sup>16</sup> have been used to develop SrFe<sub>12</sub>O<sub>19</sub> based materials. However, the preparation of SrFe<sub>12</sub>O<sub>19</sub> often suffers from the agglomeration, which is caused by the magnetic dipole interaction between ferrite particles.<sup>17</sup> Moreover, single component functional material cannot meet the ever-increasing demands for absorbing materials. Hence, combining SrFe<sub>12</sub>O<sub>19</sub> with another suitable component would be a feasible way to solve the above mentioned problems.

Graphene, a single layer of carbon atoms patterned in a hexagonal lattice, has attracted worldwide attentions owing to its two dimensional structure, high specific surface area, super electronic, mechanical and thermal properties.<sup>18,19</sup> Considering the excellent properties of graphene and SrFe<sub>12</sub>O<sub>19</sub>, SrFe<sub>12</sub>O<sub>19</sub>/graphene (SrFe<sub>12</sub>O<sub>19</sub>/G) hybrids can be very attractive for microwave absorbing applications. On one hand, graphene with planar structure provides an ideal platform for the growth of SrFe<sub>12</sub>O<sub>19</sub>, suppressing the aggregation of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles. On the other hand, SrFe<sub>12</sub>O<sub>19</sub> nanoparticles on the surface of graphene inhibit the re-stacking of graphene. The structural stability guarantees the good electromagnetic properties.

Luo *et al.*<sup>20</sup> prepared reduced graphene oxide/strontium ferrite/polyaniline (RGO/SF/PANI) composites by a three-step method. Firstly, they obtained SrFe<sub>12</sub>O<sub>19</sub> nanoparticles by co-precipitation and post-calcination method. Secondly, they

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mixed SrFe<sub>12</sub>O<sub>19</sub> nanoparticles with graphene oxide (GO) solution and obtained RGO/SrFe<sub>12</sub>O<sub>19</sub> hybrids. Finally, *in situ* polymerization method was used to obtain RGO/SF/PANI composites. Zhao *et al.*<sup>24</sup> synthesized RGO/SrFe<sub>12</sub>O<sub>19</sub> nanoflakes composites by a two-step method. They separately prepared SrFe<sub>12</sub>O<sub>19</sub> nanoflakes and RGO, and subsequently mixed them to obtain the composites. The absorption properties were also investigated. However, there are few reports focused on the construction of SrFe<sub>12</sub>O<sub>19</sub>/graphene composites by a one-step *in situ* method.

Herein we synthesize the SrFe<sub>12</sub>O<sub>19</sub>/G composites by a one-step hydrothermal reaction with the assistance of the surfactant. The formation of SrFe<sub>12</sub>O<sub>19</sub> and the reduction of GO are accomplished simultaneously, which is much different from two-step method.<sup>20,21</sup> Also, post-calcinations are not needed in our experiment, avoiding the introduction of impurities, and saving energy and time.<sup>16</sup> Moreover, the planar structure of graphene and the use of surfactant restrain the aggregation of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles.<sup>3</sup> In addition, the applications of pristine ferrites are limited due to large densities and narrow absorbed frequency. Graphene is light-weight and highly conductive, which offers the possibility of satisfying the ever-increasing demands for absorbing materials.<sup>22</sup> The structure, morphology and absorbing properties of as-prepared hybrids were investigated. The minimum reflection loss (RL) of SrFe<sub>12</sub>O<sub>19</sub>/G composites at the thickness of 3 mm reached -34.8 dB at 13.6 GHz, which was much better than that of the pristine SrFe<sub>12</sub>O<sub>19</sub> particles (-13.5 dB). It is believed that the as-synthesized composites can find applications in the microwave absorbing field.

## Experimental

### Materials

Natural flake graphite (325 mesh) was purchased from Beijing Creative Biological Engineering Materials Co. Ltd. Strontium chloride (SrCl<sub>2</sub>·6H<sub>2</sub>O, 99.9%) and Iron(III) chloride hexahydrate (FeCl<sub>3</sub>·6H<sub>2</sub>O, 99.9%) were purchased from Aladdin. All other reagents and solvents were purchased from Beijing Chemicals. All chemicals used in our experiments were reagent grade and used without further purification.

### Preparation of the SrFe<sub>12</sub>O<sub>19</sub>/G composites

Graphene oxide (GO) was synthesized from natural flake graphite by a modified Hummers method as reported elsewhere.<sup>17</sup> The detailed procedure is described as follows: FeCl<sub>3</sub>·6H<sub>2</sub>O and SrCl<sub>2</sub>·6H<sub>2</sub>O (the atomic ratio of Fe/Sr = 12) were mixed in 40 mL of deionized water under magnetic stirring. After complete dissolution, 0.5 g of polyvinylpyrrolidone (PVP) was added into the above solution with ultrasonication for 0.5 h. Then, 0.05 g of GO was slowly added into the mixture solution and dispersed by ultrasonication for 0.5 h. Subsequently, 4 g of NaOH was slowly added into 20 mL of ammonia solution (28% mass concentration, purchased and used without further processing) with mild agitation in an ice bath. Then the aqueous solution containing 0.1 mol of NaOH and 0.294 mol of ammonia was obtained. The

freshly prepared mixed solution containing NaOH and ammonia was used to adjust the pH value of above precursor solution to 13. Finally, the mixture was transferred to a 80 mL Teflon-lined autoclave, sealed and maintained at 220 °C for 6 h. The autoclave was then naturally cooled to room temperature and the black precipitate was collected, washed, and dried under vacuum at 80 °C for 12 h.

### Characterization

The field emission scanning electron microscopy (FESEM) was performed with a Hitachi S-4800 microscope operated at 15 kV. Transmission electron microscopy (TEM) was carried out using a Hitachi HT7700 microscope. Fourier transform infrared spectroscopy (FTIR) was recorded on a Bruker VECTOR 22 spectrometer in the frequency range of 4000–500 cm<sup>-1</sup>. X-ray diffraction (XRD) patterns were recorded on a X-ray diffractometer (Ultima IV) at 40 kV and 150 mA with Cu K<sub>α</sub> radiation. The composition and chemical states of samples were investigated by X-ray photoelectron spectroscopy (XPS, PHI 5300X). The magnetic properties were measured using a vibrating sample magnetometer (VSM, Lakeshore 7300) in the field of 20 kOe. Vector network analyzer (HP 8722ES) was used to measure the complex permittivity and permeability of the composites in 1–18 GHz.

## Results and discussion

Fig. 1 displayed representative TEM and HRTEM images of the GO, SrFe<sub>12</sub>O<sub>19</sub> particles and SrFe<sub>12</sub>O<sub>19</sub>/G composites. It can be seen from Fig. 1a that the GO had a two dimensional planar structure with wrinkles and folds on the edges. The pure SrFe<sub>12</sub>O<sub>19</sub> particles showed the typical irregular agglomerated particle morphology in Fig. 1b. The sizes of SrFe<sub>12</sub>O<sub>19</sub> particles reached up to several microns. As revealed by the TEM images of as-prepared composites (Fig. 1c and d), the whole graphene sheets were decorated by plenty of homogeneous nanoparticles with a diameter of 60–70 nm. And no obvious aggregation was observed, which was different from the pure SrFe<sub>12</sub>O<sub>19</sub> particles.

Fig. 1e illustrated the HRTEM image of GO and the lattice spacing of GO was 0.83 nm. Both the lattice fringes of graphene and SrFe<sub>12</sub>O<sub>19</sub> nanoparticles were clearly observable in Fig. 1f. The lattice spacing of graphene was 0.36 nm, smaller than that of GO, indicating the removal of the oxygen-containing functional groups.<sup>1</sup> The lattice fringe distances of about 0.26 nm and about 0.27 nm were indexed to the (114) and (107) planes of SrFe<sub>12</sub>O<sub>19</sub>, respectively. The clear lattice indicated that SrFe<sub>12</sub>O<sub>19</sub> nanoparticles were of high crystallinity.

Graphene with planar structure provided an ideal platform for the growth of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles, and the existence of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles inhibited the re-stacking of graphene. Therefore, it was safe to deduce that the introduction of graphene overcame the poor dispersion of SrFe<sub>12</sub>O<sub>19</sub> particles in the composites. The structural uniformity indicated the good electromagnetic properties and potential microwave absorbing applications of the SrFe<sub>12</sub>O<sub>19</sub>/G composites.

Fig. 2 showed FTIR spectra of the GO and SrFe<sub>12</sub>O<sub>19</sub>/G composites. The peaks of GO located at 3409, 1738, 1625, 1399



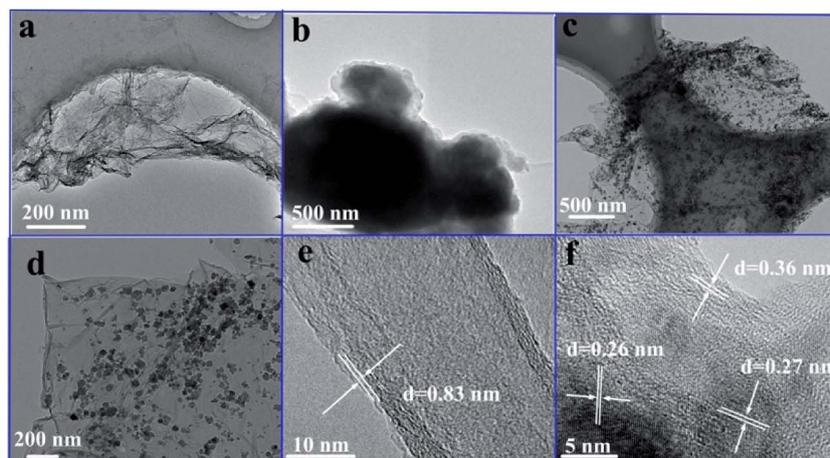


Fig. 1 TEM images of the GO (a), SrFe<sub>12</sub>O<sub>19</sub> particles (b), SrFe<sub>12</sub>O<sub>19</sub>/G composites (c and d), HRTEM images of the GO (e) and SrFe<sub>12</sub>O<sub>19</sub>/G composites (f).

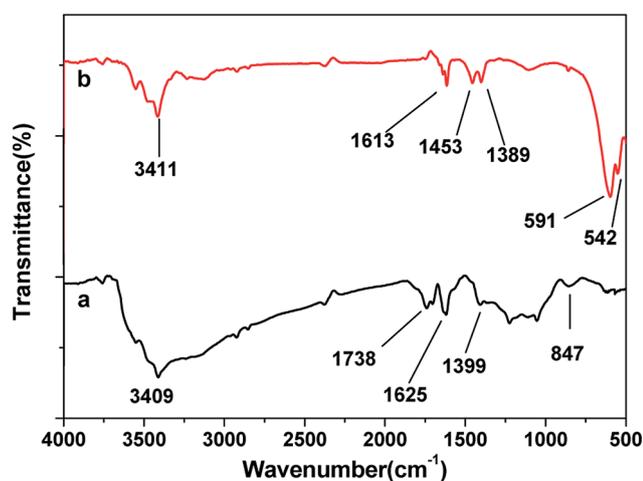


Fig. 2 FTIR spectra of the (a) GO and (b) SrFe<sub>12</sub>O<sub>19</sub>/G composites.

and 847 cm<sup>-1</sup> were due to the vibration and deformation bands of O–H, C=O stretching vibrations from carbonyl groups, C=C configurable vibrations from the aromatic zooms, C–OH stretching vibrations, C–O vibrations from epoxy groups and alkoxy groups, respectively.<sup>18</sup> However, most of the peaks related to the oxygen-containing functional groups disappeared in the FTIR spectrum of SrFe<sub>12</sub>O<sub>19</sub>/G composites (Fig. 2b), revealing that these oxygen-containing functional groups were almost removed after the hydrothermal reaction. The hydrothermal condition with ammonia provided thermal and chemical factors to reduce the GO. The peaks at 1613 cm<sup>-1</sup> were attributed to the C=C configurable vibrations from the graphene, peaks at 1453 and 1389 cm<sup>-1</sup> exhibited the stretching vibration and deformation vibration of hydroxy group. Moreover, an additional peak at 591 and 542 cm<sup>-1</sup> in Fig. 2b could be ascribed to lattice absorption of Fe–O, further confirming the existence of SrFe<sub>12</sub>O<sub>19</sub>.<sup>19</sup>

As shown in Fig. 3a, a sharp peak at  $2\theta = 10.6^\circ$  can be indexed to the (001) crystallographic plane of the GO. However,

it vanished and a new characteristic (002) diffraction peak appeared at  $2\theta = 25.1^\circ$  after the hydrothermal reduction (Fig. 3b). It was noteworthy that the broadening and weakening of the (002) diffraction peak of graphene was obvious, indicating the reduction process cannot restore graphene to the highly ordered crystal structure of graphite. The SrFe<sub>12</sub>O<sub>19</sub> and SrFe<sub>12</sub>O<sub>19</sub>/G composites exhibited the very similar diffraction peaks, as shown in Fig. 3c and d. And the Rietveld refinement of XRD patterns of SrFe<sub>12</sub>O<sub>19</sub>/G composites was shown in Fig. S1.† The (001) crystallographic plane of the GO disappeared in the XRD pattern of SrFe<sub>12</sub>O<sub>19</sub>/G composites, suggesting the reduction of GO after the hydrothermal reaction.<sup>3</sup> Nevertheless, no (002) diffraction peak of graphene appeared, which may be caused by two reasons. First of all, the *in situ* growth of SrFe<sub>12</sub>O<sub>19</sub> nanoparticles on the graphene inhibited the restacking of graphene layers during the reduction process, weakening its diffraction intensity. Moreover, the diffraction peak of graphene was easy to be suppressed by SrFe<sub>12</sub>O<sub>19</sub> due to the loading content of graphene in the composites.<sup>23,24</sup> All

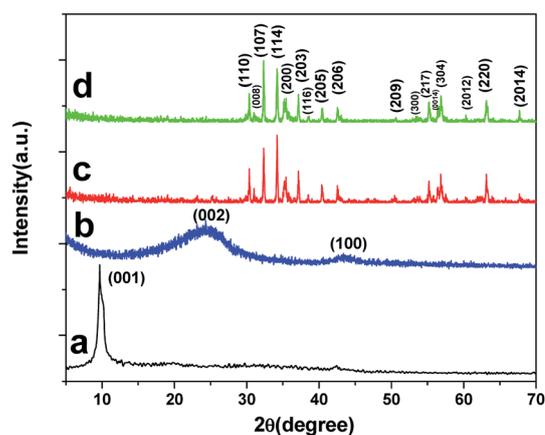


Fig. 3 XRD patterns of the (a) GO, (b) graphene, (c) SrFe<sub>12</sub>O<sub>19</sub> and (d) SrFe<sub>12</sub>O<sub>19</sub>/G composites.



labeled diffraction peaks in the range of 5–70° can be indexed to the hexagonal SrFe<sub>12</sub>O<sub>19</sub> (PDF card no. 33-1340) crystallites.<sup>25,26</sup> The sharp diffraction peaks and high intensity indicated the good crystallinity of as-obtained composites.

XPS was also employed to analyze GO and the SrFe<sub>12</sub>O<sub>19</sub>/G composites. The wide scan XPS spectrum (Fig. 4a) of the SrFe<sub>12</sub>O<sub>19</sub>/G composites showed photoelectron lines at a binding energy of 133.1, 284.6, 530.2 and 710.9 eV attributed to Sr 3d, C 1s, O 1s and Fe 2p, respectively. The inset in Fig. 4b showed the C 1s of GO, which consisted of two main components arising from C–C/C=C (284.6 eV) and C–O (286.6 eV) groups and two minor components from C=O (288.2 eV) and O–C=O (289.4 eV) groups.<sup>3,17</sup> Compared with GO, the C 1s of SrFe<sub>12</sub>O<sub>19</sub>/G composites varied dramatically. The content of C–O group decreased rapidly, and peaks of C=O and O–C=O groups even disappeared, suggesting a remarkable reduction after the hydrothermal reaction. In Fig. 4c, the Sr 3d<sub>5/2</sub> signal appeared at 133.3 eV, and the peak at 134.9 eV was ascribed to the Sr 3d<sub>3/2</sub> level. Fig. 4d showed the spectrum of Fe 2p region. The peaks of Fe 2p<sub>3/2</sub> and Fe 2p<sub>1/2</sub> were located at 711.6 and 724.8 eV, respectively. All the above analysis indicated the successful synthesis of the SrFe<sub>12</sub>O<sub>19</sub>/G composites.

Magnetic hysteresis loops of the SrFe<sub>12</sub>O<sub>19</sub>/G composites and SrFe<sub>12</sub>O<sub>19</sub> particles were shown in Fig. 5. The saturation magnetization and coercivity of the SrFe<sub>12</sub>O<sub>19</sub> particles were 61.2 emu g<sup>-1</sup> and 2312.2 Oe, respectively. It can be observed that the saturation magnetization dropped dramatically with the introduction of nonmagnetic graphene. For the SrFe<sub>12</sub>O<sub>19</sub>/G composites, the saturation magnetization and coercivity were 41.6 emu g<sup>-1</sup> and 1695.6 Oe, respectively.

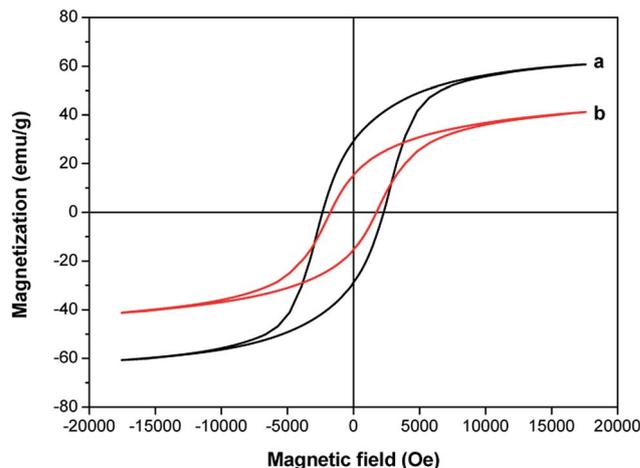


Fig. 5 Magnetic hysteresis loops for (a) SrFe<sub>12</sub>O<sub>19</sub> particles and (b) SrFe<sub>12</sub>O<sub>19</sub>/G composites.

A vector network analyzer was used to measure the electromagnetic parameters of the as-prepared SrFe<sub>12</sub>O<sub>19</sub>/G composites in the frequency range of 1–18 GHz. The complex permittivity and complex permeability of the SrFe<sub>12</sub>O<sub>19</sub>/G composites and the SrFe<sub>12</sub>O<sub>19</sub> particles are shown in Fig. 6 and 7, respectively. As we know, the real part of complex permittivity ( $\epsilon'$ ) and complex permeability ( $\mu'$ ) stand for the storage ability of electromagnetic energy, and the imaginary part of complex permittivity ( $\epsilon''$ ) and complex permeability ( $\mu''$ ) represent the electromagnetic energy loss ability.<sup>27</sup> For the SrFe<sub>12</sub>O<sub>19</sub>/G composites, the  $\epsilon'$  showed an approximately constant around

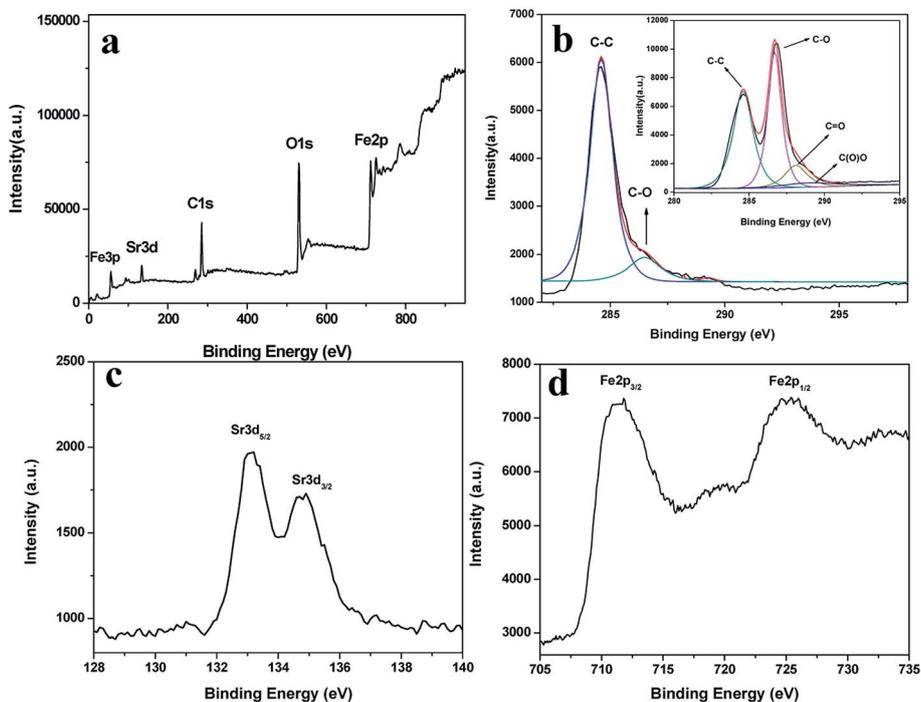


Fig. 4 XPS spectra of the (a) survey scan, (b) C 1s region (the inset shows C 1s of GO), (c) Sr 3d region, (d) Fe 2p region of the SrFe<sub>12</sub>O<sub>19</sub>/G composites.



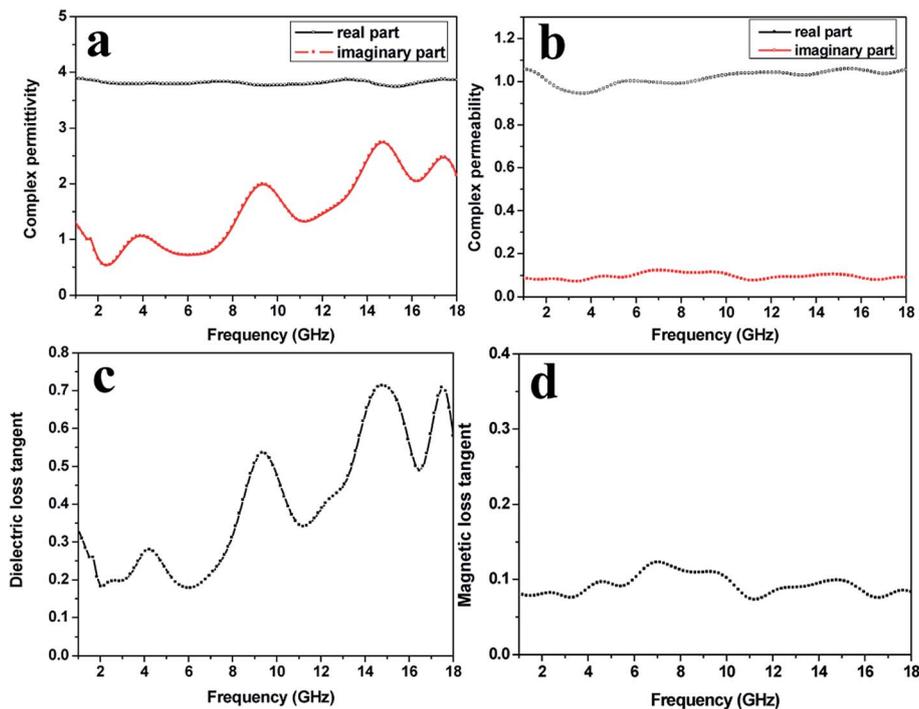


Fig. 6 The complex permittivity (a), complex permeability (b), dielectric loss tangent (c), magnetic loss tangent (d) of the SrFe<sub>12</sub>O<sub>19</sub>/G composites.

3.9 in the range of 1–18 GHz, while the  $\epsilon''$  exhibited four high fluctuations from 0.5 to 2.7 in the whole frequency range. According to free electron theory,<sup>28</sup>  $\epsilon'' \approx 1/2\pi\epsilon_0\rho f$ , where  $\rho$  is the resistivity. It can be speculated that the lower  $\epsilon''$  values represent higher electric resistivity. In general, the high electric resistivity is in favor of building a balance between complex permeability

and complex permittivity, thus decreasing the reflection coefficient of the absorbing materials. Meanwhile, the  $\mu'$  and  $\mu''$  changed gently in the range of 1–18 GHz.  $\mu'$  and  $\mu''$  maintained at around 1.0 and 0.1, respectively. The dielectric loss tangent indicates the inherent dissipation of electromagnetic energy for dielectric materials. It can be seen from Fig. 6c that the

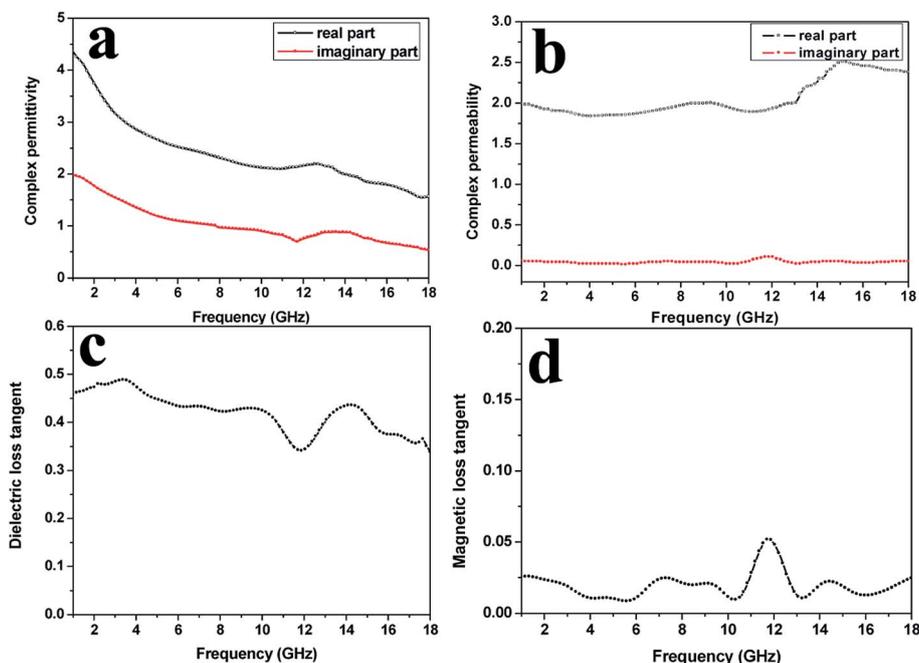


Fig. 7 The complex permittivity (a), complex permeability (b), dielectric loss tangent (c), magnetic loss tangent (d) of the SrFe<sub>12</sub>O<sub>19</sub> particles.



dielectric loss tangent showed a similar variation tendency with  $\epsilon''$ . Four sharp peaks appeared at 4.4, 9.9, 15.1 and 17.5 GHz, and the peak values were 0.29, 0.54, 0.73 and 0.71, respectively. Meanwhile, the magnetic loss tangent increased from 0.08 to 0.13 at the 1.0–7.1 GHz range, and decreased from 0.13 to 0.08 over 7.1–18.0 GHz. Apart from dielectric loss and magnetic loss, another significant factor affecting electromagnetic wave absorption is impedance match characteristic, too high permittivity of absorber is detrimental to the impedance match and results in strong reflection and weak absorption.

The complex permittivity, complex permeability, dielectric loss tangent and magnetic loss tangent of the SrFe<sub>12</sub>O<sub>19</sub> particles are shown in Fig. 7a–d, respectively. Both the  $\epsilon'$  and  $\epsilon''$  showed a downward trend in the range of 1–18 GHz, declining from 4.4 and 2.0 to 1.6 and 0.6, respectively. The  $\mu'$  kept unchanged around 2.0 in the range of 1.0–13.2 GHz, and then went up to 2.3 over 13.2–18.0 GHz. The  $\mu''$  remained constant at about 0.08 in the whole frequency range. Compared with the SrFe<sub>12</sub>O<sub>19</sub>/G composites, the SrFe<sub>12</sub>O<sub>19</sub> particles exhibited much lower dielectric loss tangent (less than 0.5) and magnetic loss tangent (around 0.025). Thus, it would be safe to conclude that the SrFe<sub>12</sub>O<sub>19</sub>/G composites exhibited better electromagnetic properties than the SrFe<sub>12</sub>O<sub>19</sub> particles.

The RL can be calculated from the complex permeability and complex permittivity at the given absorber thickness and frequency using the following equations:

$$RL(\text{dB}) = 20 \log \left| \frac{Z_{\text{in}} - 1}{Z_{\text{in}} + 1} \right| \quad (1)$$

While the normalized input impedance ( $Z_{\text{in}}$ ) is calculated by:

$$Z_{\text{in}} = \sqrt{\frac{\mu_r}{\epsilon_r}} \tanh \left( j \frac{2\pi f d}{c} \sqrt{\mu_r \epsilon_r} \right) \quad (2)$$

where  $f$  is the electromagnetic wave frequency,  $d$  is the thickness of the absorb layer,  $c$  is the velocity of electromagnetic wave in vacuum.  $\epsilon_r$  and  $\mu_r$  are the relative complex permittivity and permeability, respectively, which can be calculated by:

$$\epsilon_r = \epsilon' - j\epsilon'', \quad \mu_r = \mu' - j\mu'' \quad (3)$$

The RL curves of the SrFe<sub>12</sub>O<sub>19</sub> particles and the SrFe<sub>12</sub>O<sub>19</sub>/G composites with different thicknesses are shown in Fig. 8. The

RL is expected to be as low as possible at a given sample thickness. The minimum RL of the SrFe<sub>12</sub>O<sub>19</sub> particles at the thickness of 4 mm reached  $-13.5$  dB, and the effective absorption frequency (RL <  $-10$  dB) ranged from 12.9 to 17.5 GHz. However, for the SrFe<sub>12</sub>O<sub>19</sub>/G composites, the minimum RL of  $-34.8$  dB was observed at 13.6 GHz with a thickness of 3 mm, and the effective absorption frequency ranged from 11.7 to 17.4 GHz. All these analysis suggested that the SrFe<sub>12</sub>O<sub>19</sub>/G composites exhibited much better microwave absorbing properties than the SrFe<sub>12</sub>O<sub>19</sub> particles.

The related literatures based on SrFe<sub>12</sub>O<sub>19</sub> were listed in Table 1. Wang *et al.*<sup>29</sup> synthesized SrFe<sub>12</sub>O<sub>19</sub>-MWCNTs composites with different MWCNTs content by sol-gel method. The results indicated that 6 wt% SrFe<sub>12</sub>O<sub>19</sub>-MWCNTs composites possessed the best magnetic and wave absorbing properties. Poorbafrani *et al.*<sup>30</sup> firstly prepared SrFe<sub>12</sub>O<sub>19</sub> by sol-gel method, then combined with different concentrations of ZnO and obtained SrFe<sub>12</sub>O<sub>19</sub>/ZnO composites. They found the reflection loss (RL) in the Ku frequency band (12–18 GHz) was enhanced by up to 5 dB for 15% of ZnO in the nanocomposites. Chen *et al.*<sup>31</sup> synthesized SrFe<sub>12</sub>O<sub>19</sub>/ZnFe<sub>2</sub>O<sub>4</sub> composites with different weight ratios of SrFe<sub>12</sub>O<sub>19</sub> to ZnFe<sub>2</sub>O<sub>4</sub> using sol-gel technique. The composite powder with 15 wt% ZnFe<sub>2</sub>O<sub>4</sub> possessed good microwave absorption properties. The minimum RL of the composite powder reached  $-37$  dB at the thickness of 2.2 mm. He *et al.*<sup>32</sup> synthesized a new type of conductive and magnetic PANI/Ag/SrFe<sub>12</sub>O<sub>19</sub> composites *via* three-step method. The RL of the composites was below  $-10$  dB between 8.7 GHz and 12.1 GHz, with a minimum loss value of

Table 1 Overview of absorbing properties for different SrFe<sub>12</sub>O<sub>19</sub> based composites

Samples	Parameters		
	Minimum RL (dB)	Frequency (GHz)	Effective bandwidth (GHz)
SrFe <sub>12</sub> O <sub>19</sub> -MWCNTs	-19.7	3.2	Not mentioned
SrFe <sub>12</sub> O <sub>19</sub> /ZnO	-15.47	8.37	Not mentioned
SrFe <sub>12</sub> O <sub>19</sub> /ZnFe <sub>2</sub> O <sub>4</sub>	-37	8.7	3.9
PANI/Ag/SrFe <sub>12</sub> O <sub>19</sub>	-14.86	9.98	3.4
Sr <sub>1-x</sub> La <sub>x</sub> Fe <sub>12</sub> O <sub>19</sub>	-41.7	9.0	4
This work	-34.8	13.6	5.7

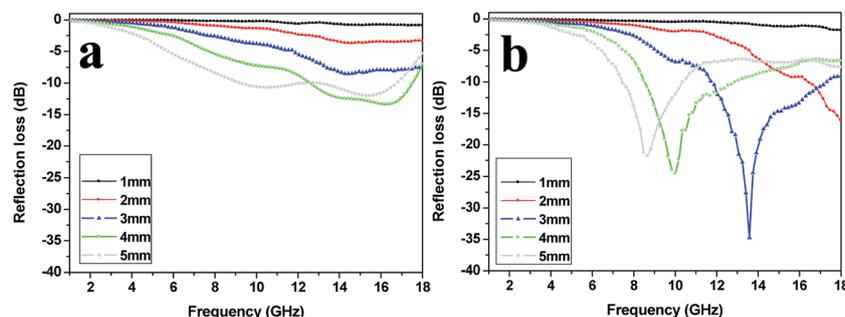


Fig. 8 The calculated reflection loss of (a) the SrFe<sub>12</sub>O<sub>19</sub> particles, (b) the SrFe<sub>12</sub>O<sub>19</sub>/G composites with different thicknesses.



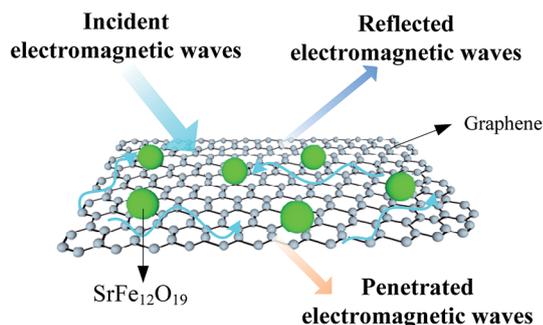


Fig. 9 Schematic diagram of absorbing mechanism for SrFe<sub>12</sub>O<sub>19</sub>/G composites.

−14.86 at 9.98 GHz. Chen *et al.*<sup>33</sup> prepared lanthanum substituted strontium ferrite Sr<sub>1-x</sub>La<sub>x</sub>Fe<sub>12</sub>O<sub>19</sub> ( $x = 0.05, 0.1, 0.15, 0.2$ ) by a sol-gel process. The results showed that the minimum RL of the Sr<sub>1-x</sub>La<sub>x</sub>Fe<sub>12</sub>O<sub>19</sub> ( $x = 0.15$ ) reached −41.7 dB with the −10 dB bandwidth over 4 GHz at the thickness of 2.3 mm.

The enhanced absorbing properties of as-obtained composites can be attributed to three reasons. First of all, the conductive graphene with two dimensional planar structures constructed an effective conductive network in the composites, providing high dielectric loss tangent.<sup>17</sup> Moreover, the well dispersion of homogenous SrFe<sub>12</sub>O<sub>19</sub> nanoparticles on the graphene support ensured the pronounced absorbing performances. In addition, interface scattering could be generated owing to the difference in complex permittivity between graphene and SrFe<sub>12</sub>O<sub>19</sub> nanoparticles, leading to more efficient absorption of electromagnetic waves. The absorbing mechanism diagram of SrFe<sub>12</sub>O<sub>19</sub>/G composites was illustrated in Fig. 9.

## Conclusions

In summary, the SrFe<sub>12</sub>O<sub>19</sub>/G composites were synthesized by a one-step *in situ* surfactant-assisted hydrothermal method. The homogeneous SrFe<sub>12</sub>O<sub>19</sub> nanoparticles with a typical diameter of 60–70 nm were well anchored on graphene without aggregation. The minimum RL of −34.8 dB was observed at 13.6 GHz with a thickness of 3 mm, and the effective absorption bandwidth was 5.7 GHz. The synergistic effect of SrFe<sub>12</sub>O<sub>19</sub> and graphene made a great contribution to the improved absorbing properties. The SrFe<sub>12</sub>O<sub>19</sub>/G composites exhibited better absorbing performance than the SrFe<sub>12</sub>O<sub>19</sub> particles, and showed a promising prospect as absorbing materials.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 M. Fu, Q. Z. Jiao and Y. Zhao, *Mater. Charact.*, 2013, **86**, 303.
- 2 V. M. Petrov and V. V. Gagulin, *Inorg. Mater.*, 2001, **37**, 93.
- 3 M. Fu, Q. Z. Jiao and Y. Zhao, *J. Mater. Chem. A*, 2013, **1**, 5577.
- 4 W. Chen, Q. Y. Liu, X. X. Zhu and M. Fu, *Appl. Organomet. Chem.*, 2017, DOI: 10.1002/aoc.4017.
- 5 Y. W. Wang, L. D. Zhang and G. W. Meng, *J. Phys. Chem. B*, 2002, **106**, 2502.
- 6 Q. Y. Liu, Y. T. Yang, H. Li, R. R. Zhu, Q. Shao, S. G. Yang and J. J. Xu, *Biosens. Bioelectron.*, 2015, **64**, 147.
- 7 L. F. Sun, Y. Y. Ding and Y. L. Jiang, *Sens. Actuators, B*, 2017, **239**, 848.
- 8 S. Ruhle, A. Y. Anderson, H. N. Barad, B. Kupfer, Y. Bouhadana, E. Rosh-Hodesh and A. Zaban, *J. Phys. Chem. Lett.*, 2012, **3**, 3755.
- 9 D. Sando, Y. R. Yang, E. Bousquet, C. Carrétéro, V. Garcia, S. Fusil, D. Dolfi, A. Barthélémy, P. Ghosez, L. Bellaiche and M. Bibes, *Nat. Commun.*, 2016, **7**, 1.
- 10 M. M. Shirolkar, C. Hao, X. Dong, T. Guo, L. Zhang, M. Li and H. Wang, *Nanoscale*, 2014, **6**, 4735.
- 11 S. H. Hsieh, R. S. Solanki, Y. F. Wang, Y. C. Shao, S. H. Lee, C. H. Yao, C. H. Du, H. T. Wang, J. W. Chiou, Y. Y. Chin, H. M. Tsai, J. L. Chen, C. W. Pao, C. M. Cheng, C. W. Chen, H. J. Lin, J. F. Lee, F. C. Chou and H. M. Tsai, *Sci. Rep.*, 2017, **7**, 161.
- 12 A. L. Xia, C. H. Zuo, L. Chen, C. G. Jin and Y. H. Lv, *J. Magn. Magn. Mater.*, 2013, **332**, 186.
- 13 G. Asghar and M. Anis, *J. Alloys Compd.*, 2012, **526**, 85.
- 14 X. S. Liu, L. Fernandez-Garcia, F. Hu, D. R. Zhu, M. Suárez and J. L. Menéndez, *Mater. Chem. Phys.*, 2012, **133**, 961.
- 15 A. Ghasemi, *J. Magn. Magn. Mater.*, 2012, **324**, 1375.
- 16 J. F. Wang, C. B. Ponton and I. R. Harris, *J. Magn. Magn. Mater.*, 2001, **234**, 233.
- 17 X. Tang and Y. G. Yang, *Appl. Surf. Sci.*, 2009, **255**, 9381.
- 18 K. S. Novoselov, A. K. Geim, S. V. Morozov, E. W. Hill, P. Blake, M. I. Katsnelson and K. S. Novoselov, *Science*, 2004, **306**, 666.
- 19 A. K. Geim, *Science*, 2009, **324**, 1530.
- 20 J. Luo, P. Shen, W. Yao and C. Jiang, *Nanoscale Res. Lett.*, 2016, **11**, 141.
- 21 C. Y. Zhao, M. Y. Shen, Z. X. Li, R. Sun, A. L. Xia and X. G. Liu, *J. Alloys Compd.*, 2016, **689**, 1037.
- 22 M. Fu, Q. Z. Jiao, Y. Zhao and H. S. Li, *J. Mater. Chem. A*, 2014, **2**, 735.
- 23 A. V. Murugan, T. Muraliganth and A. Manthiram, *Chem. Mater.*, 2009, **21**, 5004.
- 24 X. A. Fan, J. G. Guan, X. F. Cao, W. Wang and F. Z. Mou, *Eur. J. Inorg. Chem.*, 2010, **2010**, 419.
- 25 C. S. Lin, C. C. Hwang, T. H. Huang, G. P. Wang and C. H. Peng, *Mater. Sci. Eng., B*, 2007, **139**, 24.
- 26 M. M. Hessien, M. M. Rashad and K. El-Barawy, *J. Magn. Magn. Mater.*, 2008, **320**, 336.
- 27 X. F. Zhang, P. F. Guan and X. L. Dong, *Appl. Phys. Lett.*, 2010, **96**, 223111.



- 28 X. F. Zhang, X. L. Dong, H. Huang, B. Lv and C. J. Choi, *J. Phys. D: Appl. Phys.*, 2007, **40**, 5383.
- 29 W. T. Wang, Q. L. Li and C. B. Chang, *Synth. Met.*, 2011, **161**, 44.
- 30 A. Poorbafrani, P. Kameli and H. Salamati, *J. Mater. Sci.*, 2013, **48**, 186.
- 31 N. Chen, G. H. Mu, X. F. Pan, K. K. Gan and M. Y. Gu, *Mater. Sci. Eng., B*, 2007, **139**, 256.
- 32 Z. He, S. H. Qi, X. L. Zhong, H. Ma, P. Wang and H. Qiu, *J. Alloys Compd.*, 2015, **621**, 194.
- 33 N. Chen, K. Yang and M. Y. Gu, *J. Alloys Compd.*, 2010, **490**, 609.

