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Controlling the electric permittivity of honeycomb-like core-shell Ni/CuSiO₃ composite nanospheres to enhance microwave absorption properties†

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Controlling impedance matching and enhancing absorption properties are crucial for developing next-generation microwave absorbing materials. Herein, we report the facile preparation of honeycomb-like core-shell nickel/copper silicate (Ni/CuSiO₃) composite nanospheres with controlled shell (CuSiO₃) thickness to control impedance matching and enhance absorption properties. The nanospheres with a shell thickness of about 30 nm had greatly improved microwave absorption performance. They exhibited a strong reflection loss of −39.5 dB (9.6 GHz) at the absorber thickness of 2.5 mm, with a wide effective absorbing bandwidth of 4.8 GHz (7.8–12.6 GHz). This remarkable microwave absorption performance is attributed to a novel structure that enhanced interfacial polarization and controlled shell thickness, resulting in good impedance matching of electromagnetic properties (ϵ , μ) and magnetic loss. Our permittivity-controlled core-shell Ni/CuSiO₃ composite nanospheres could be used to fabricate novel high-performance microwave absorbers.

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1 Introduction

Extensive research has been conducted on the design and fabrication of highly effective electromagnetic (EM) wave absorbers to suppress unwanted EM wave reflections. These absorbers are of great interest for electric and electronic devices, and in military and industrial applications.^{1–5} Ferromagnetic materials are the most promising candidates for EM absorbers because of their high EM loss efficiency.^{1–3} Notably, nickel (Ni)-based absorbers are easy to prepare and display high permeability in the GHz range.^{6–9} However, severe eddy current loss in the GHz range caused by high conductivity results in impedance mismatch between the material and air space, which decreases microwave absorption performance. Eddy current loss can also decrease the permeability, in which case there is a deterioration of the matching between the permittivity (ϵ) and permeability (μ).^{9,10} Additionally, oxidation and corrosion occur when Ni is used at high temperatures and in corrosive environments. These shortcomings can be overcome using

inorganic/nonmagnetic coating materials. Core-shell composite microstructures can also improve microwave absorption performance by controlling the impedance mismatch.^{4,11,12}

Various types of core-shell composite microstructures have been reported based on Ni as the core material and different shell materials including polymers, C, SnO₂, TiO₂, SiO₂, ZnO, ZnS, CuO, and graphene composites.^{9,13–20} Zhao *et al.* reported ZnS-wrapped Ni spheres that exhibited a reflection loss (RL) of −42.4 dB with high effective bandwidth.¹⁷ Deng *et al.* prepared ZnO nanorod-coated Ni spheres and reported an RL of −30.2 dB.²⁰ Zhao *et al.* reported Ni@TiO₂ and Ni@SiO₂ core-shell composites and found that the SiO₂ shell had better microwave absorption properties and anti-oxidation capability than pure Ni or Ni@TiO₂.⁹ These reports concluded that the coating material plays a decisive role in obtaining good EM wave absorption properties through impedance matching, cooperation between the EM parameters (ϵ , μ), core and shell synergistic effects, and interfacial polarizations.

Copper silicate (CuSiO₃) has been used as a shell material for magnetic core materials because of its good dielectric and anti-oxidation properties, ease of preparation, and low cost.^{21,22} Core-shell composites having CuSiO₃ as the shell have been used as microwave absorbers due to the enhanced polarization obtained with the highly dielectric CuSiO₃ shell and a magnetic core.^{21–25} The combination of CuSiO₃ as the shell with a Ni magnetic core would be advantageous because more interfaces could be created, along with enhanced polarization. Furthermore, a design with a variable shell thickness would facilitate

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impedance matching between the EM properties (ϵ and μ) of the core-shell nanospheres because the shell thickness has a strong influence on ϵ and μ . Notably, the ϵ value can be easily tuned by controlling the shell (CuSiO_3) thickness of Ni/ CuSiO_3 composite nanospheres. No integrated CuSiO_3 and Ni core-shell microstructures have been reported to date.

Herein, we report the facile synthesis of honeycomb-like core-shell Ni/ CuSiO_3 composite nanospheres. The shell thickness was readily controlled by varying the concentration of Ni spheres in the intermediate step of the sol-gel (Stöber) method used in their preparation. The Ni/ CuSiO_3 composite nanospheres with a 30 nm-thick shell exhibited enhanced EM absorption properties, including strong RL and wide effective absorption bandwidth. This behavior is attributed to the novel honeycomb-like structure, well-controlled ϵ value that resulted in higher microwave attenuation, and good impedance matching. These core-shell Ni/ CuSiO_3 nanospheres with an adjustable shell thickness have great potential as high-performance EM wave absorbers.

2 Experimental

Materials

Nickel(II) chloride hexahydrate ($\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$), copper nitrate trihydrate ($\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$), sodium hypophosphate (NaH_2PO_2), tetraethyl orthosilicate (TEOS), sodium hydroxide (NaOH), sodium acetate (NaAc), sodium citrate, and glycerol were purchased from Sigma-Aldrich and used as received.

Preparation of Ni nanospheres, Ni/ SiO_2 nanospheres, and honeycomb-like core-shell Ni/ CuSiO_3 composite nanospheres

Nickel spheres were synthesized using a hydrothermal method. Briefly, 0.6 g of $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ was dissolved in a mixture of glycerol (15 mL) and deionized (DI) water (5 mL). To this mixture was added 1.5 g of NaAc and 0.1 g of sodium citrate; the solution was stirred until homogeneous. Meanwhile, another solution was prepared by dispersing NaOH (0.8 g) and NaH_2PO_2 (0.6 g) in DI water (10 mL). Then, the two solutions were combined and transferred into an autoclave for heat treatment at 140 °C for 15 h. The resultant gray-colored product was separated from the reaction solution using a permanent magnet and washed with ethanol and DI water. The powdered product (Ni) was obtained after vacuum drying at 60 °C. These Ni spheres were then used to prepare Ni/ SiO_2 spheres according to the sol-gel procedure reported elsewhere.²² The Ni spheres (0.5 g) were dispersed in an ethanol/DI water mixture (8 : 2 volume ratio), followed by addition of ammonia solution (2 mL). Then, 0.6 mL of TEOS was added dropwise to the solution, which was then stirred for about 10 h. The obtained products were washed with DI water and ethanol three times by magnetic decantation and dried in a vacuum oven. These Ni/ SiO_2 spheres were treated hydrothermally to obtain the final honeycomb-like Ni/ CuSiO_3 core-shell composites, as previously described.²² Typically, 0.03 g of Ni/ SiO_2 spheres was dissolved in 30 mL of DI water by ultrasonication. Under ultrasonication, 1.5 mL of ammonia solution and 1.5 mL of 0.1 M aqueous solution of $\text{Cu}(\text{NO}_3)_2 \cdot 3\text{H}_2\text{O}$ were

added to the solution. The mixed solution was transferred to an autoclave for hydrothermal treatment at 120 °C for about 12 h. The final products were obtained after washing three times with ethanol and DI water using magnetic decantation and dried in a vacuum oven.

Characterization

The size and morphology of the nanospheres were observed by field-emission scanning electron microscopy (FE-SEM; S-4800; Hitachi, Japan), transmission electron microscopy (TEM; Tecnai G2 F30; FEI, USA), high-resolution TEM (HRTEM), and bright-field high-angle annular dark-field scanning transmission electron microscopy (BF/HAADF-STEM). The elemental composition was determined using energy-dispersive X-ray spectroscopy (EDS). The crystal structure and phases were both studied by X-ray diffraction (XRD; D8 Advance; Bruker, Germany) and Micro diffraction using two-dimensional detectors (XRD; Discover; Bruker, Germany). Magnetization (M - H) curves were obtained using a vibrating sample magnetometer (VSM; Lakeshore 7304; Lakeshore Cryotronics, Inc., USA). The EM properties were studied using a network analyzer (N5245A; Agilent, USA) in the microwave range (2–18 GHz) for evaluation of RL. A toroidal sample (diameter (ϕ)_{outer}: 6.95 mm; (ϕ)_{inner}: 3.05 mm) was prepared by blending powders (30 wt%) with paraffin wax. The RL of a composite absorber can be calculated from using eqn (1) and (2):²

$$\text{RL} = 20 \log |(Z_{\text{in}} - Z_0)/(Z_{\text{in}} + Z_0)| \quad (1)$$

$$Z_{\text{in}} = Z_0 \sqrt{\mu_r/\epsilon_r} \tanh \left\{ j(2\pi f d/c) \sqrt{\mu_r/\epsilon_r} \right\} \quad (2)$$

where Z_{in} and Z_0 are the input impedances of the absorber and the free space impedance, respectively, c is the velocity of light, and d is the absorber thickness. ϵ_r and μ_r are the relative complex permittivity and permeability, respectively, and are expressed as $\epsilon_r = \epsilon' - j\epsilon''$ and $\mu_r = \mu' - j\mu''$.^{2,9,17,20}

3 Results and discussion

The synthesis of the core-shell Ni/ CuSiO_3 composite nanospheres is illustrated in Fig. 1. The Ni nanospheres prepared by the hydrothermal treatment appeared as walnut-like spheres with rough surfaces. These are obtained by the reduction of Ni^{2+} ions in alkaline medium with sodium hypophosphite (NaH_2PO_2) at 140 °C for 15 h. By considering the ecological, economical and handling factors, NaH_2PO_2 is better reducing agent than the hydrazine (N_2H_4) which was commonly used in preparation of Ni particles.²⁶ Thus, in this work we select NaH_2PO_2 for the reduction purpose of Ni^{2+} ions. At first, Ni^{2+} ions formed from the dissociation of NiCl_2 in water and glycerol mixed solvent medium then it will reduced to metallic Ni with reducing agent (NaH_2PO_2) at strong alkali medium. This process has been represented as follows by eqn (3).

Fig. 1 Schematic illustration of the synthesis of the core-shell nickel/copper silicate (Ni/CuSiO₃) composite nanospheres.

Alkali medium (pH $\sim$ 10) was created from the NaOH and Na-citrate and will be regulated and maintained by the presence of NaAc, which acts as stabilizing buffer. The heating temperature (140 °C) and alkali medium favor to accelerating the kinetics for the homogeneous nucleation of Ni. Then the Ni spheres are obtained through the self-assembly of Ni particles into a walnut-like fashion at reaction temperature of 140 °C, keeping for 15 h. They were coated with SiO₂ using the Stöber sol-gel process. The SiO₂ coating protected the Ni core from air

oxidation and modified the surface by minimizing the curvature thereof.^{18,24} The Ni/SiO₂ spheres were treated hydrothermally in basic medium and with a copper salt to form CuSiO₃ shells by the reaction of silicate anions from SiO₂ in the alkaline solution with the copper ions from dissolved Cu(NO₃)₂·3H₂O.

The morphology and size of the products at each stage are shown in Fig. 2. Fig. 2(a–c) shows that the walnut-like-shaped Ni nanospheres were about 350 nm in diameter and had curved surfaces. The Ni/SiO₂ nanospheres displayed a uniform SiO₂

Fig. 2 Scanning electron microscopy (SEM) and transmission electron microscopy (TEM) images. (a–c) Ni nanospheres, (d–f) Ni/SiO₂ nanospheres, and (g–i) Ni/CuSiO₃ composite nanospheres.

Fig. 3 Scanning transmission electron microscopy (STEM) images of the Ni/CuSiO₃ composite nanospheres. (a) Image showing the core-shell structure, (b) elemental overlay image, (c) energy-dispersive X-ray spectroscopy line spectra, and (d–g) elemental distributions.

coating (about 30 nm-thick) with smoother surface morphology (Fig. 2(d–f)) than the Ni nanospheres. The BF/HAADF-STEM images and elemental maps revealed that the Ni spheres were

completely covered with an SiO₂ layer (Fig. S1†). The thickness of this layer could be regulated by simply varying the Ni concentration. For example, 0.3, 0.5, and 0.6 g of Ni gave 50-, 30-, and

Fig. 4 (a) X-ray diffraction (XRD) patterns of the Ni nanospheres, Ni/SiO₂ nanospheres, and Ni/CuSiO₃ composite nanospheres. (b) Vibrating sample magnetometer hysteresis loops of the same materials. (c) Two-dimensional XRD pattern of the Ni/CuSiO₃ composite nanospheres.

Fig. 5 Transmission electron microscopy (TEM) images of the Ni/CuSiO₃ composite nanospheres having different shell thicknesses. (a and b) NCS-1, (c and d) NCS-2, and (e and f) NCS-3. (g) High-resolution TEM image. (h) Selected area electron diffraction pattern.

20 nm-thick SiO₂ layers, respectively (Fig. S2†). The final composite nanospheres had a core-shell structure (Fig. 2(g–i)). The inset SEM image (Fig. 2(g)) and TEM image (Fig. 2(h and i)) shows that the honeycomb-like CuSiO₃ layer was composed of many CuSiO₃ flakes, which aggregate in a honeycomb-like fashion. The resulting honeycomb-like core-shell Ni/CuSiO₃ composite nanospheres had diameters equal to those of the Ni/SiO₂ nanospheres (about 380 nm).

The BF/HAADF-STEM images of the Ni/CuSiO₃ composite nanospheres further confirmed the microstructure and elemental distribution. The composite nanospheres had a core–

shell structure (Fig. 3(a)); the elemental composition is shown in Fig. 3(b). The line-EDS spectra revealed the presence of all four elements (Fig. 3(c)). Individual elemental maps (Fig. 3(d–h)) revealed that the Ni was in the core (Fig. 3(d)), while Cu, Si, and O were mainly in the shell. Moreover, the composition of the shell was in sharp contrast to that of the CuSiO₃ formed by a combination of Cu, Si, and O.

The XRD diffraction peaks of the three products were in good agreement with the face-centered cubic (FCC) phase of Ni (marked by circle symbol in Fig. 4); they were centered at $2\theta = 44.7^\circ$, 52.0° , and 77.6° (JCPDS#04-0850).¹⁵ The final product exhibited additional peaks; the peaks marked by circles indexed to CuSiO₃ (PDF#32-0346).^{22,25} However, Ni/SiO₂ core-shell nanospheres shows XRD peaks for only Ni, and absence for SiO₂. Whereas, STEM image and elemental mapping of Ni/SiO₂ core-shell nanospheres (Fig. S1†) indicated that the silicon, and oxygen atoms are present in shell. It is noted that there is no evidence for crystalline SiO₂ in XRD but SiO₂ existing in the Ni/SiO₂ core-shell nanospheres as in amorphous state. The same behavior was also observed in our previous works.²² Fig. 4(b) shows the two-dimensional XRD pattern of the final composite nanospheres. The intense lines of high contrast correspond to the Ni phase, while the less intense lines correspond to crystalline CuSiO₃. Therefore, the XRD results confirmed the presence of both Ni and CuSiO₃ in the composite structure. Hysteresis curves demonstrated that all three samples had ferromagnetic characteristics (Fig. 4(c)). The saturated magnetization (Ms) values of Ni, Ni/SiO₂, and Ni/CuSiO₃ were 48, 37, and 30 emu g^{−1}, respectively. Lower Ms values were measured for Ni/SiO₂ and Ni/CuSiO₃ because of the non-magnetic shell layers in their core-shell structures.^{27,28} This feature enhanced EM absorption.

Fig. 5(a–f) displays TEM images of the prepared Ni/CuSiO₃ composite nanospheres with shell thicknesses of 50, 30, and 20 nm; these materials are denoted as NCS-1, NCS-2, and NCS-3, respectively. The SiO₂ layer thicknesses of the precursor Ni/SiO₂ nanospheres are shown in Fig. S2.† The HRTEM images of the composite nanospheres (Fig. 5(g)) revealed that the core and shell were highly crystalline. The selected area electron diffraction image (Fig. 5(h)) taken from the square area in Fig. 5(g) indicated that the Ni was highly crystalline.

Electromagnetic parameters (ϵ_r , μ_r) are crucial to assess EM absorption properties. This is because the real parts (ϵ' , μ') and imaginary parts (ϵ'' , μ'') of the permittivity and permeability represent the storage and loss ability of EM energy, respectively.^{21,29} Fig. 6 illustrates how the EM properties were influenced by the shell thickness of the composite nanospheres. Fig. 6(g) shows the permittivity (ϵ_r) of Ni nanospheres is lower than its composites. It has been increased and tuned *via* the shell thickness. As the shell thickness increase the ϵ' value increased. On the other hand, the permeability (μ_r) values are not in big difference for these samples. Hence, controlling of ϵ_r is favor to reach the good matching between the electromagnetic parameters. Fig. 6(a, c and e) clearly shows that ϵ_r decreased with decreasing shell thickness; the thickness was highest for NCS-1 and lowest for NCS-3. While the real part of the permittivity (ϵ') decreased with increasing frequency and

Fig. 6 Electromagnetic properties of (a and b) NCS-1, (c and d) NCS-2, (e and f) NCS-3, and (g and h) Ni nanospheres.

suddenly declined at high frequency, ϵ'' increased slowly, and then abruptly at about 12–18 GHz. Despite the increasing trend in ϵ'' , there were strong fluctuations related to strong interfacial polarizations in the composite nanospheres. These polarizations are due to high current lag at the interface, which increases ϵ'' .^{15,25} The dipole and enhanced interfacial polarizations account for the dielectric loss.^{2,15} The dipole polarizations derive from defects in the core and shell materials, whereas interfacial polarizations result from charge accumulation at the interfaces due to the presence of very different materials (Ni and CuSiO₃) in the composite nanospheres and are verified from cole-cole plot (Fig. S3†).^{21,30,31} Fig. 6(b, d and f) reveals that the trends in μ_r and ϵ_r were opposite. The μ' value steadily decreased with increasing frequency, except for a slight fluctuation in the 16–18 GHz region. The μ'' displayed peaks at low frequency correspond to natural resonance of the Ni

nanospheres, and the peaks at high frequency correspond to exchange resonance or eddy current loss.^{15,32} The coating of the Ni with a dielectric material (CuSiO₃) could have been responsible for the eddy current loss.^{18,33} The dielectric loss ($\tan \delta_e = \epsilon''/\epsilon'$) was slightly higher than the magnetic loss ($\tan \delta_m = \mu''/\mu'$) due to enhanced polarization; this indicated that dielectric loss was the dominant loss mechanism in the Ni/CuSiO₃ composite nanospheres (Fig. S3†).

The microwave absorption performance of the absorbers can be obtained from analysis of the RL curves.^{29,31–34} Ni show very poor performance than Ni@SiO₂ core-shell nanospheres which has the RL of −18 dB at 8.3 GHz ($d = 2.5$ mm) (Fig. S4†). This is mainly arise from the dielectric contribution of SiO₂ layer in core-shell nanospheres. Since the CuSiO₃ shell is more efficient than simple SiO₂ layer on Ni and has been introduced to further enhance the microwave absorption properties of Ni composites.

Fig. 7 Reflection loss (RL) as a function of frequency for different thicknesses of (a and b) NCS-1, (c and d) NCS-2, and (e and f) NCS-3.

The NCS-1, NCS-2, and NCS-3 absorbers exhibited significantly enhanced microwave absorption from lower to higher frequencies (Fig. 7(a–f)). The greatest RLs were -23.5 dB at 11.3 GHz ($d = 3$ mm), -39.5 dB at 9.6 GHz ($d = 2.5$ mm), and -22.3 dB at 8.1 GHz ($d = 3$ mm) for NCS-1, NCS-2, and NCS-3, respectively. These results indicate that the microwave

absorption properties can be easily tailored by controlling the shell thickness of the prepared composite nanospheres. The NCS-2 absorber had the highest RL and the widest effective absorbing bandwidth (RL below -10 dB), of about 4.8 GHz (7.8 to 12.6 GHz), at the thickness of 2.5 mm. Table 1 compares the enhanced microwave absorption performance (stronger RL and

Table 1 Comparison of microwave absorption performance among Ni-based core-shell composites

Sample	Max. RL (dB)	Thickness (mm)	Filler weight (%)	Effective bandwidth (< -10 dB) (GHz)	Ref.
Ni/SnO ₂	-18.6	7	70	1.5	33
Ni/polyaniline	-35	5	50	0.3	34
PS/PPy/Ni	-20	2	50	4.6	35
Ni/carbon	-32	2	40	4.3	36
Ni/graphene	-13	2	20	2.6	37
Ni/ZnS	-42.4	2.2	40	4.2	17
Ni/ZnO	-30.2	2.2	40	2.5	20
Ni/SiO ₂	-40	1.5	70	3.5	9
Ni/TiO ₂	-35.4	4	70	1.0	9
Ni/CuSiO ₃	-39.5	2.5	30	4.8	This work

Fig. 8 Microwave attenuation constant for NCS-1, NCS-2, and NCS-3; inset shows the impedance matching coefficient for NCS-2.

Fig. 9 Proposed mechanism of microwave absorption by the Ni/CuSiO₃ composite nanospheres.

wider effective bandwidth) of the present composite nanospheres with that of other reported Ni-based composites.^{9,17,20,34–39} The superior performance of the prepared absorbers suggests that the honeycomb-like core-shell structure contributed significantly to the microwave absorption. Dielectric loss, magnetic loss, shell thickness, and geometry affected performance. The present composite nanospheres displayed higher, and tunable, performance at lower filler weight and with thinner thickness. Such tunable microwave absorption properties are desirable in many applications.

The superior microwave absorption performance of our new absorbers resulted from the good microwave impedance matching by the honeycomb-like core-shell Ni/CuSiO₃ composite nanospheres, and strong microwave attenuation.

The impedance matching coefficient (η) for the composites is given by eqn (4):⁴⁰

$$\eta = \frac{\mu''\epsilon'}{\mu'\epsilon''} \quad (4)$$

The η of the Ni/CuSiO₃ composite nanospheres was constant over the entire frequency range, which indicated good impedance matching (Fig. 8(b)). This was due to the balance of EM parameters in the composite nanospheres. The microwave attenuation constant (α) was calculated using eqn (5):^{31,33}

$$\alpha = \left(\frac{\sqrt{2}\pi f}{c} \right) \sqrt{(\mu''\epsilon'' - \mu'\epsilon')^2 + (\mu''\epsilon' - \mu'\epsilon'')^2 + (\epsilon'\mu'' - \epsilon''\mu')^2} \quad (5)$$

NCS-2 have reached the higher microwave attenuation constant (α) value than other two composites and NCS-3 attains the lowest value. The higher value of the attenuation constant observed for NCS-2 was due to good impedance matching (Fig. 8(b)); its excellent attenuation indicated good microwave absorption. Therefore, the Ni/CuSiO₃ composite nanospheres provided strong RL (−39.5 dB) and wide effective bandwidth (4.8 GHz) with reduced thickness. These advantages offer new opportunities for Ni/CuSiO₃ composite nanospheres as tunable, high-performance microwave absorbers.

The high performance of the composite nanospheres was clearly associated with the unique honeycomb-like core-shell structure. This type of structure can provide multiple scatterings/reflections from the interspaces between the core and shell, resulting in a longer pathway for the EM wave in the absorber, and a higher rate of heat dissipation.^{2,15} The many interfaces of the honeycomb structure, and the thin flakes constituting the shell, increase the path length. Additionally, interfacial polarizations associated with the relaxation process are enhanced in the core-shell structures, which have highly curved surfaces at the interfaces.^{21,41} The Ni/CuSiO₃ composite nanospheres having enhanced interfacial polarizations strongly attenuated microwave energy. Moreover, they also exhibited magnetic loss from the ferromagnetic Ni. Fig. 9 shows the probable microwave absorption mechanism for the Ni/CuSiO₃ composite nanospheres. These can be considered as high-performance microwave absorption nanomaterials having the advantages of tunable and strong RL, wide absorbing bandwidth, and novel structural characteristics.

4 Conclusions

Novel honeycomb-like core-shell Ni/CuSiO₃ composite nanospheres displayed strong microwave absorption performance because of the novel structure of a Ni core and a honeycomb shell composed of small CuSiO₃ flakes. The CuSiO₃ shell thickness was tuned using different thicknesses of the precursor SiO₂ layer (Ni/SiO₂) formed by adjusting the Ni concentration in the sol-gel preparative process. However, the composite nanospheres with a 30 nm-thick shell displayed particularly strong microwave absorption properties, *i.e.*, a strong RL of −39.5 dB (9.6 GHz) and a wide effective absorption bandwidth of about 4.8 GHz. Notably,

the novel structure effectively dissipated microwave energy because of the combination of high dielectric loss, high magnetic loss, and enhanced interfacial polarizations. The abundant interfaces between the Ni core and honeycomb-like CuSiO₃ shell ensured efficient multiple scattering/reflection. Thus, the novel design of the structure and tunability of the shell thickness are expected to assist development of other high-performance microwave absorbers.

Conflicts of interest

The authors declare no conflict of interest.

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