

Effect of zinc-nickel ferrite nanoparticles content on microwave absorbing ability for the X-band frequency range

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Abstract

In this paper, the absorbing paint consists of carbon black/Zn_{0.8}Ni_{0.2}Fe₂O₄/SiO₂/epoxy (20 wt.% of carbon black constantly) had been fabricated. The effect of Zn_{0.8}Ni_{0.2}Fe₂O₄ content on the microwave-absorbing ability was studied for X-band frequency range. Using XRD, TEM, VSM to study the structure, morphology and magnetic properties of zinc nickel ferrite nanoparticles and zinc nickel ferrite nanoparticles covered by SiO₂ matrix. Network Analyzer (Agilent E5071C ENA series) was used to measured the reflection loss (RL) of samples with different Zn_{0.8}Ni_{0.2}Fe₂O₄ content (0 – 1.25 wt.%). The absorption percentage of samples increase by increasing the Zn_{0.8}Ni_{0.2}Fe₂O₄ content and reach to 98.4 % (1.25 wt.% Zn_{0.8}Ni_{0.2}Fe₂O₄).

Keywords: Zinc-nickel ferrite, carbon black, microwave-absorbing materials, absorbing paint.

1. Introduction

Microwave-absorbing materials are currently in high demand for civil engineering and military engineering applications. Dielectric and magnetic materials have microwave-absorbing properties. Forms of microwave-absorbing materials commonly known as sheets, bricks, paints....[1] These paints include an absorber such as carbon black, graphene, ferrite, iron carbonyl or a polymer matrix. The spinel ferrites have been chosen as absorbing materials in various forms for many years. Epoxy can play an important role in realizing these microwave absorbers with additional functionalities like elasticity for the mixture. Some works are mixture of glass fabric/epoxy and carbon black [2,3], carbon nanotubes in a matrix of strontium and nickel ferrite, composites of carbon nanotubes and polyurethane [4], iron carbonyl and carbon black in an epoxy resin/rubber matrix [5,6] The content of absorbers

are 20–25 wt.% carbon black [1, 5], 60–80 wt.% ferrite [7,8] or 25–50 wt.% iron carbonyl [5, 6]. The microwave-absorbing ability depended on the absorber content and coating thickness. The thinner and lighter coating with high microwave absorption was the aim of the application.

In this study, microwave-absorbing samples were fabricated using 20 wt.% carbon black, super-paramagnetic Zn_{0.8}Ni_{0.2}Fe₂O₄ dispersed in SiO₂ matrix with different contents (0, 5, 15 and 25%) corresponding to 0, 0.25, 0.75, 1.25 wt%, of Zn_{0.8}Ni_{0.2}Fe₂O₄ and epoxy resin to test their micro wave absorption at 8 – 12 GHz (X band range).

2. Materials and Method

The structure of zinc-nickel ferrite cover by SiO₂ layers samples were analysed by X-ray diffraction (XRD) using Bruker AXS D8 diffractometer with Cu-Kα radiation. The morphological study was carried out by transmission electron microscope (TEM, JEOL- 1400). Magnetic measurements of samples were taken out at room temperature using vibrating sample magnetometer, (VSM, MICRO SENSE 3474-140). To study the reflection loss of samples, carbon black (20 wt.%) was mixed with different Zn_{0.8}Ni_{0.2}Fe₂O₄ contents and epoxy resin, hardener to form microwave-absorbing paints. Then, the paint was coated onto the steel plates (200 × 200 × 3 mm) and dried for three days. The mixture proportions was shown in Table 1. The reflection loss were measured thorough Network Analyzer (Agilent E5071C ENA series) in the frequency range of 8 –12 GHz at room temperature.

The prepared samples with Zn_{0.8}Ni_{0.2}Fe₂O₄ contents of 0, 5, 15 and 25 wt.% were signed as samples 1, 2, 3 and 4.

Table 1. The four prepared samples with different $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ content

Samples	Carbon black (wt.%)	Zinc-nickel ferrite covered by SiO_2 layers (wt.%)	Epoxy resin (wt.%)
1	20	0 (0)	80
2	20	5 (0.25)	75
3	20	15 (0.75)	65
4	20	25 (1.25)	55

The super-paramagnetic $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content of the $CB/Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ /epoxy resin mixture is indicated in parentheses.

3. Results and Discussion

3.1 Structure analysis

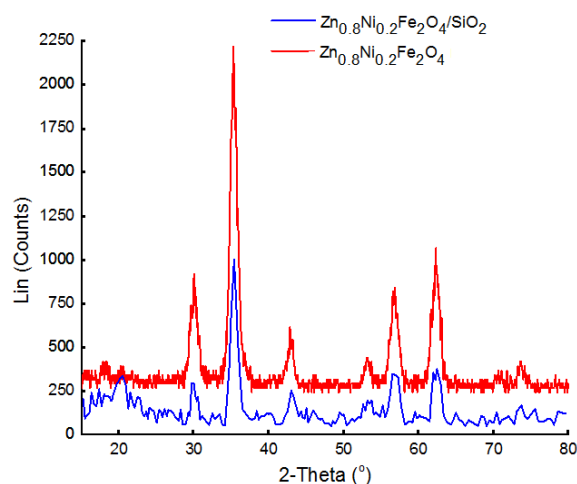


Fig. 1 XRD spectra of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ and $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$

The zinc nickel ferrite nanoparticles were obtained after annealing at $140^\circ C$ for 6 hours, all peaks (220), (311), (400), (422), (511) and (440) corresponding to zinc nickel ferrite spectrum (JCPDS 52-0277) shown in figure 1. Besides that, the XRD pattern of the $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ composite displays only five peaks corresponding to (220), (311), (400), (511) and (440) lattice planes that are less intense than those of $Zn_{0.8}Ni_{0.2}Fe_2O_4$. This could be explained by an amorphous SiO_2 layer cover on the surface of particles.

3.2 Morphology of $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ and carbon black nanoparticles

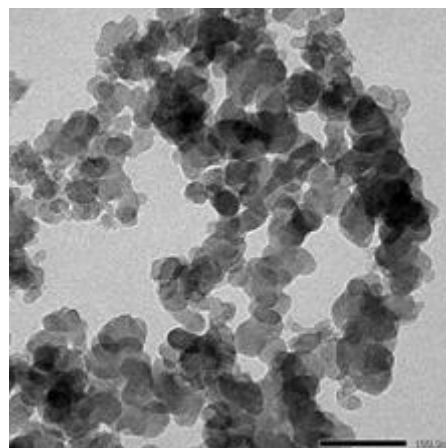


Fig.2 TEM image of carbon black particles

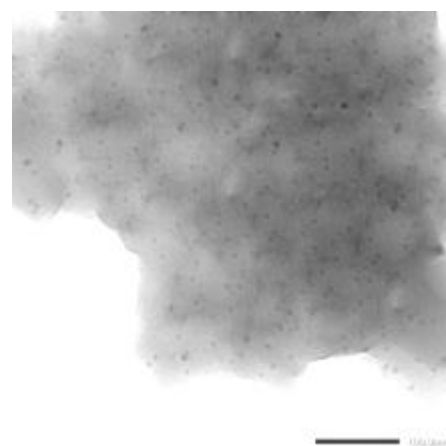


Fig. 3 TEM micrograph of zinc nickel nanoparticles dispersed into SiO_2 matrix

Figure 2 shows the average size of carbon black is around of 29 nm. The TEM image of figure 3 indicates that zinc nickel nanoparticles are dispersed into SiO_2 matrix, the average particles size is about 5 nm and the morphology is nearly spherical.

3.3 Magnetic properties of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ and $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$

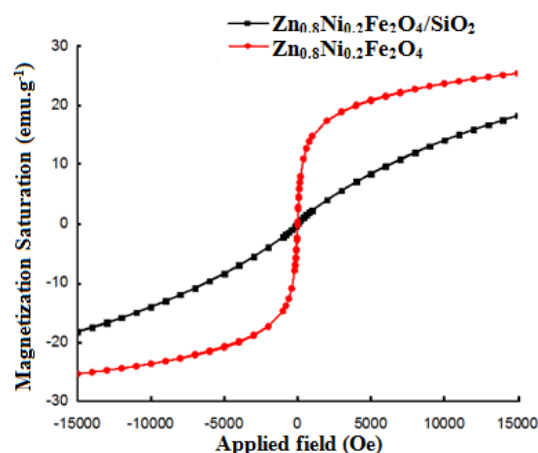


Fig. 4 Magnetisation curves of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ and $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$

Table 2. Magnetic properties of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ and $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ nanoparticles

Sample name	Remanent magnetisation M_r ($emu \cdot g^{-1}$)	Saturation magnetisation M_s ($emu \cdot g^{-1}$)	Coercivity H_c (Oe)
$Zn_{0.8}Ni_{0.2}Fe_2O_4$	≈ 0	25.39	≈ 0
$Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$	≈ 0	18.95	≈ 0

Figure 4 and table 2 show the magnetic properties of the $Zn_{0.8}Ni_{0.2}Fe_2O_4$ nanoparticles and $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$. All of samples demonstrate the coercivity (H_c) and remnant magnetisation (M_r) are nearly zero, clearly show the super-paramagnetic state. The saturation magnetisation (M_s) of the $Zn_{0.8}Ni_{0.2}Fe_2O_4$ nanoparticles is $25.39 \text{ emu} \cdot g^{-1}$, higher than sample of $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ which expressed only $18.95 \text{ emu} \cdot g^{-1}$. The nonmagnetic SiO_2 layers is the main cause of the saturation magnetisation (M_s) decrease.

3.4 Microwave-absorbing ability of four samples for X band range (8 – 12 GHz)

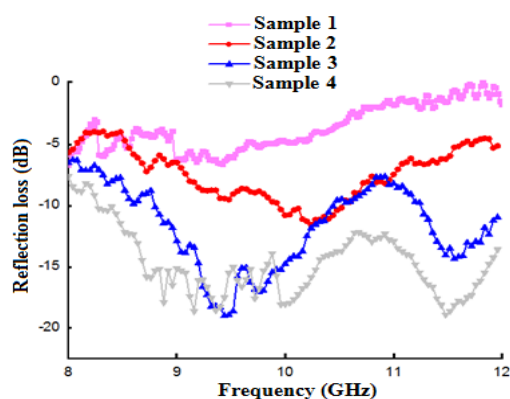


Fig. 5 Reflection loss (RL) of sample 1, 2, 3 and 4 for X-band range

 Table 3. RL and absorption percentage of samples based on $Zn_{0.8}Ni_{0.2}Fe_2O_4$ in a super-paramagnetic or ferri-magnetic state

Samples	RL (dB) at 10-GHz centred frequency	Absorption percentage (%) at 10-GHz centred frequency
1	-4.83	67.2
2	-10.85	91.8
3	-14.82	96.7
4	-18.07	98.4

Figure 5 and table 3 indicate that measured samples exhibit the RLs and absorption percentages of sample 1, 2, 3 and 4 at the X-band centre frequency of 10 GHz are -4.83, -10.85, -14.82, -18.07 dB, corresponding to 0, 0.25, 0.75 and 1.25 wt.% $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content of absorbing paint. The increase in $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content led to an increase in the RL of all samples. The small content of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ (0.25 wt.%) reaches to high RL (91.8%). So, the RLs of the samples at 10 GHz are almost dependent on the $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content.

4. Conclusions

In summary, the absorbing paint based on using carbon black/ $Zn_{0.8}Ni_{0.2}Fe_2O_4/SiO_2$ /epoxy are successfully fabricated. The effect of super-paramagnetic $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content on the microwave-absorbing ability was carried out in the X-band frequency range. It found that the increase of $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content (from 0 to 1.25 wt.%) led to an increase of RL. Sample 1 with no $Zn_{0.8}Ni_{0.2}Fe_2O_4$ content reach only 67.2 % of absorption percentage. However, sample 4 contained 1.25 wt.% $Zn_{0.8}Ni_{0.2}Fe_2O_4$ exhibited the high microwave absorption of 98.4% power attenuation at 10 GHz.

Acknowledgments

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