

Supporting Information

Morphology-controlled synthesis and excellent microwave absorption performance of ZnCo₂O₄ nanostructures via a self-assembly process of flake units

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1. Synthesis of ball-like ZnCo₂O₄

Briefly, ZnCl₂ (1 mmol) and CoCl₂·6H₂O (2 mmol) were dissolved in a mixed solvent (50 ml EG and 5 ml deionized water) under magnetic stirring at room temperature. Then, 30 mmol of NH₄HCO₃ were added into the mixture. After being stirred for 30 min, the resultant solution was transferred into a Teflon lined stainless-steel autoclave with a capacity of 100 mL, and heated at 200°C for 24 h in an oven. The ball-like ZnCo₂O₄ precursors were collected by centrifuging, washed with water three times and ethanol four times, and finally dried in a vacuum oven at 60°C for 8 h. Finally, the black ball-like ZnCo₂O₄ materials were obtained after calcination at 600°C for 4 h in air. The sample was ready for further characterization.

2. Synthesis of hydrangea-like ZnCo₂O₄

Briefly, Zn(CH₃COO)₂·2H₂O (1 mmol) and Co(CH₃COO)₂·4H₂O (2 mmol) were dissolved in 40 ml EG under magnetic stirring at room temperature. After being stirred for 30 min, the resultant solution was transferred into a Teflon lined stainless-steel autoclave with a capacity of 50 mL, and heated at 180°C for 12 h in an oven. The hydrangea-like ZnCo₂O₄ precursors were collected by centrifuging, washed with water three times and ethanol four times, and finally dried in a vacuum oven at 60°C for 12 h. Finally, the black cabbage-like ZnCo₂O₄ materials were obtained after calcination at 400°C for 2 h in air. The sample was ready for further characterization.

3. Synthesis of pineapple-like ZnCo₂O₄

Briefly, ZnCl_2 (1 mmol) and $\text{CoCl}_2 \cdot 6\text{H}_2\text{O}$ (2 mmol) were dissolved in a mixed solvent (50 ml EG and 5 ml distilled water) under magnetic stirring. Then, 30 mmol of NH_4HCO_3 and 10 mmol of urea were added to the above solution. After being stirred for 30 min, the resultant solution was transferred into a Teflon lined stainless-steel autoclave with a capacity of 100 mL, and heated at 200°C for 24 h in an oven. The pineapple-like ZnCo_2O_4 precursors were collected by centrifuging, washed with water three times and ethanol four times, and finally dried in a vacuum oven at 60°C for 8 h. Finally, the black pineapple-like ZnCo_2O_4 materials were obtained after calcination at 600°C for 4 h in air. The sample was ready for further characterization.

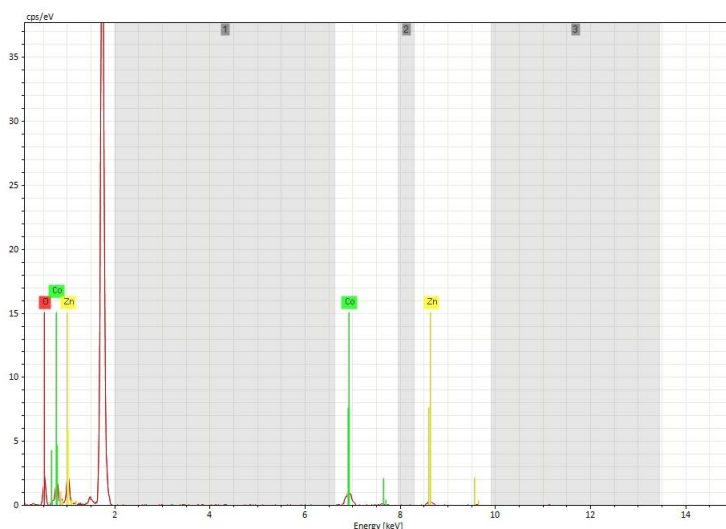


Figure S1. The EDS of cabbage-like ZnCo_2O_4 samples.

EDS quantitative analysis shows that the atomic ration of Zn, Co and O is 1:1.96:3.68, implying a certain degree of oxygen vacancy in cabbage-like ZnCo_2O_4 spinel crystal.

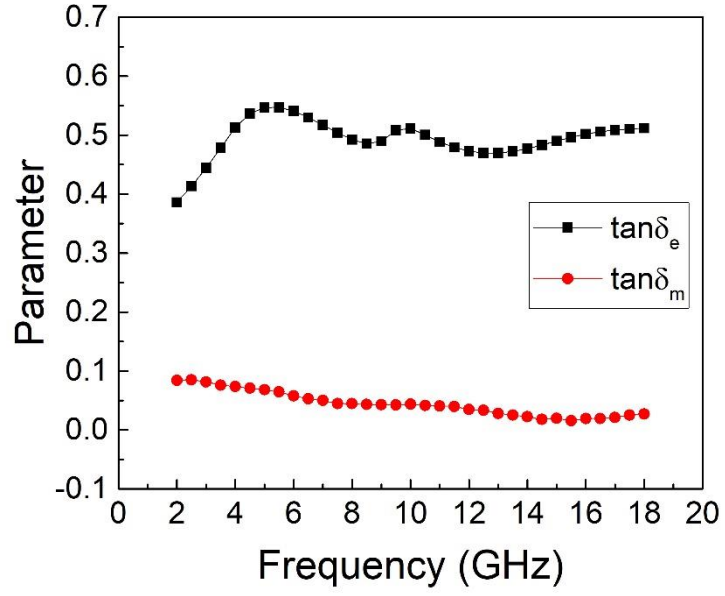


Figure S2. The $\tan \delta_e$ and $\tan \delta_m$ of as-prepared ZnCo_2O_4 samples.

Because of the calculated $\tan \delta_e$ values are much higher than that of $\tan \delta_m$, as-prepared ZnCo_2O_4 samples can be considered as a dielectric loss-type microwave absorber.