

Synthesis of $\gamma\text{-Fe}_2\text{O}_3\text{@SiO}_2\text{@Polypyrrole}$ Core/Shell/Shell Nanospheres with Flexible
Controllability of Electromagnetic Properties

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Supporting Information

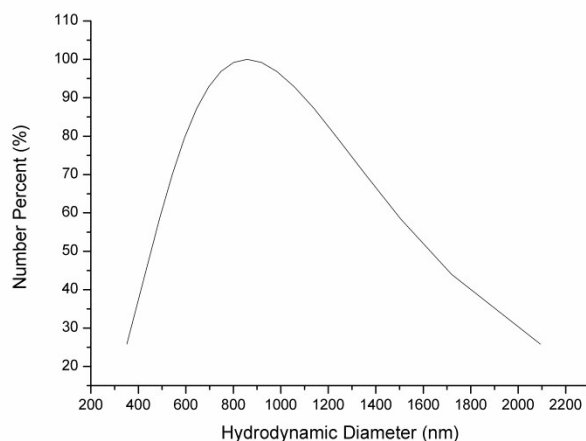


Figure S1. Dynamic light scattering profile of FSHP_{1.0-2} in alcohol. The average hydrodynamic diameter is 858 nm, PDI: 0.34.

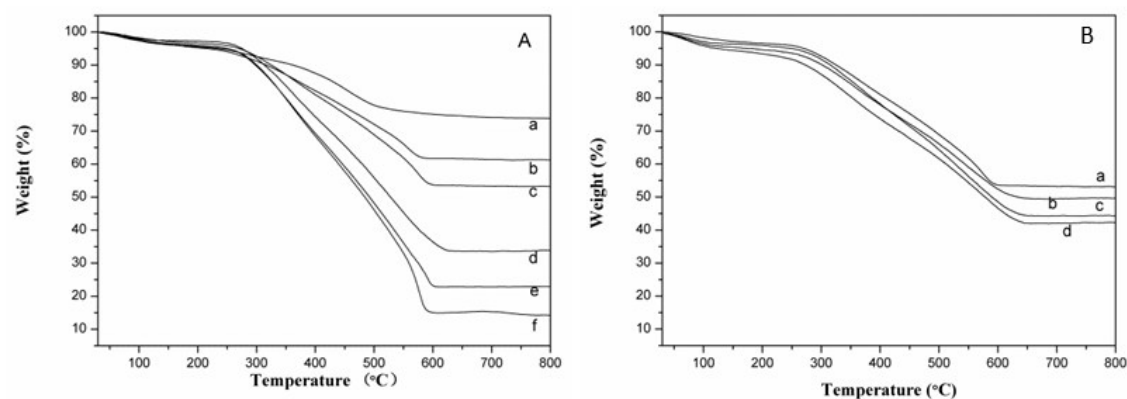


Figure S2. TG curves registered under air atmosphere: A: FSHP (a), and FSHP_{x-2} samples with different FSHP dosage: FSHP_{1.0-2} (b), FSHP_{0.6-2} (c), FSHP_{0.2-2} (d), FSHP_{0.1-2} (e), FSHP_{0.05-2} (f); B: FSHP_{0.6-y} samples with different polymerization conditions: FSHP_{0.6-2} (a), FSHP_{0.6-3} (b), FSHP_{0.6-4} (c), FSHP_{0.6-1} (d).

Table S1. FSHP mass fraction of the FSHP_{x-2} and FSHP_{0.6-y} samples calculated by the residual mass fractions at 800 °C.

Sample	Residual mass fraction (wt%)	FSHP mass fraction (wt%)*
FSHP	73.8	\
FSHP _{1.0-2}	61.3	83.1
FSHP _{0.6-2}	53.2	72.1
FSHP _{0.2-2}	33.9	45.9
FSHP _{0.1-2}	23.0	31.2
FSHP _{0.05-2}	14.2	19.2
FSHP _{0.6-1}	42.3	57.3
FSHP _{0.6-3}	49.6	67.2

FSHP _{0.6-4}	44.3	60.0
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* It is reasonable to consider that the residual mass at 800 °C was attributed to the residue of $\gamma\text{-Fe}_2\text{O}_3\text{@SiO}_2$ for each sample. In other words, the residues at 800 °C are the same substance for each sample. So we propose a method to calculate the FSH mass fraction of the FSHP_{x-2} and FSHP_{0.6-y} samples. Here we designate the FSH mass fraction of FSHP samples as **x**, mass of FSHP as **a** g and residual mass fraction of FSH and FSHP samples at 800 °C as **b** and **c**, respectively. So there is:

$$a \cdot x \cdot b = a \cdot c$$

$$x = \frac{c}{b}$$

(b and c are given by TGA data)