

Supplementary data

for

Fenton-like degradation of bisphenol A by Fe₃O₄ rhombic dodecahedrons

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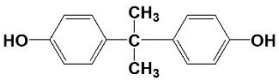
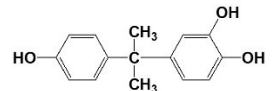
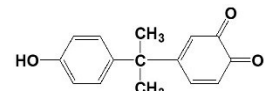
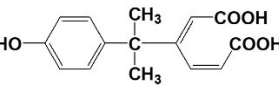
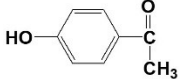
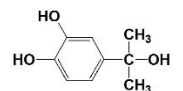
Table S1 Basic properties of Fe₃O₄-R before and after reaction in the cycle experiments.

Samples	BET surface	Pore Volume	Pore size	Surface Fe(II)/Fe(III)
	area (m² g⁻¹)	(cm³ g⁻¹)	(nm)	ratio
Fresh Fe ₃ O ₄ -R	19.99	0.041	2.12	63.51%
Used Fe ₃ O ₄ -R	18.56	0.035	1.65	50.53%

Table S2 Comparison of Fe₃O₄-R with other Fe₃O₄ catalysts in references.

Catalysts	Catalyst dosage (g L ⁻¹)	H ₂ O ₂ dosage (mM)	Pollutant concentration (mM)	Initial pH	Removal efficiency	Ref.
Superparamagnetic Fe ₃ O ₄	5	1.2	phenol (1)	6.0-7.0	60% (6 h)	1
Fe ₃ O ₄ nanoparticles	1.0	12	2,4-dichlorophenol (0.61)	3.0	51% (180 min)	2
Nano-Fe ₃ O ₄	1.0	0.05	4-chlorocatechol (10 ⁻³)	6.5	100% (3 h)	3
Nano-sized Fe ₃ O ₄	0.25	40	phenol (6.38)	4.0	98% (90 min)	4
Nanosized Fe ₃ O ₄	0.5	UV-Fenton (11.8)	catechol (0.9)	3.0	84% (240 min)	5
magnetite	0.2	UVA-Fenton (1)	phenol (0.1)	3.0	100% (4 h)	6
Fe ₃ O ₄ magnetic nanoparticles	0.585	Sono-Fenton (160)	BPA (0.09)	3.0	100% (500 min)	7
Fe ₃ O ₄ -R	0.1	0.2	BPA (0.1)	5.0	100% (30 min)	This work

Table S3 LC/MS intermediates obtained during Fenton-like degradation of BPA using Fe₃O₄-R.

Product	Experimental mass [M-H] ⁺ m/z	RT	Molecular Formula	Tentative structure
BPA	227.1045	2.64	C ₁₅ H ₁₆ O ₂	
TP243	243.0993	2.01	C ₁₅ H ₁₆ O ₃	
TP241	241.0838	1.70	C ₁₅ H ₁₄ O ₃	
TP275	275.0477	1.36	C ₁₅ H ₁₆ O ₅	
TP135	135.0424	1.42	C ₈ H ₈ O ₂	
TP167	167.0683	1.35	C ₉ H ₁₂ O ₃	

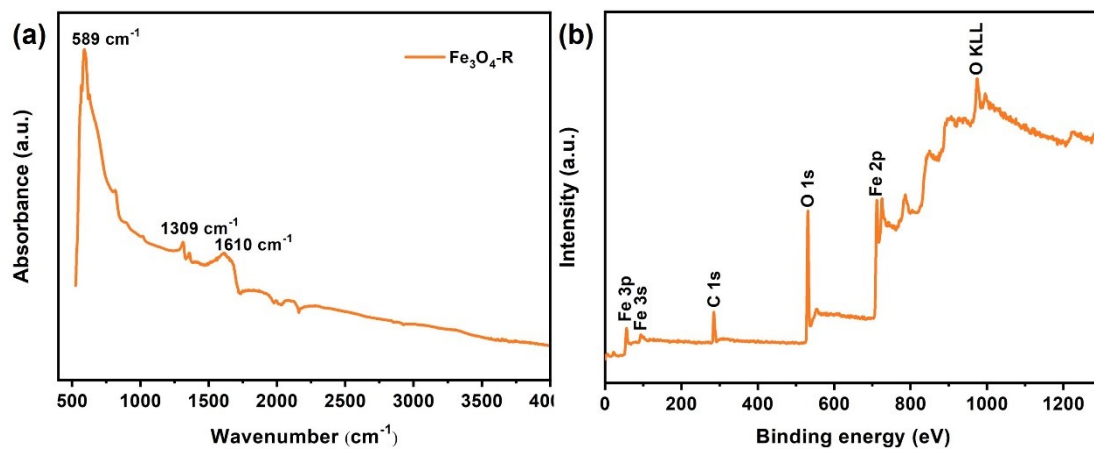


Fig. S1 (a) FTIR spectrum and (b) XPS spectrum of Fe₃O₄-R.

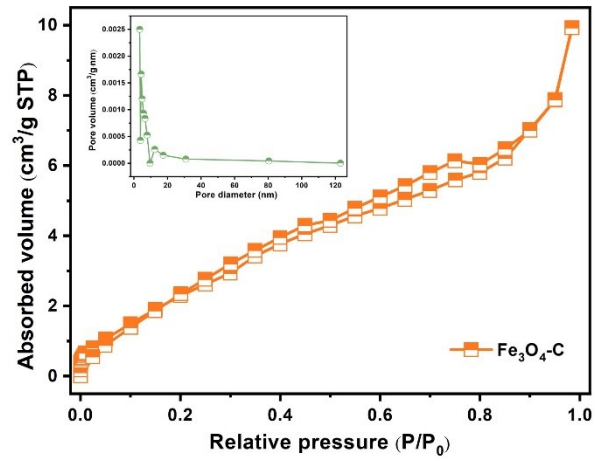


Fig. S2 Nitrogen adsorption-desorption isotherm and its pore size distribution (inset) of Fe₃O₄-C.

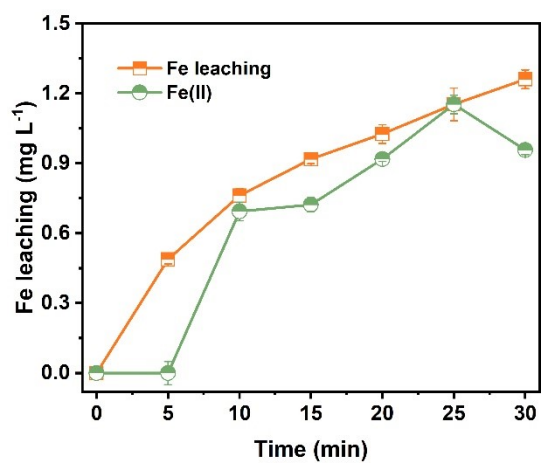


Fig. S3 The concentration of Fe leaching and Fe(II) in the $\text{Fe}_3\text{O}_4\text{-R}/\text{H}_2\text{O}_2$ system.

Conditions: $[\text{catalyst}]_0 = 0.1 \text{ g L}^{-1}$, $[\text{H}_2\text{O}_2]_0 = 5 \text{ mM}$, $[\text{BPA}]_0 = 0.1 \text{ mM}$, and initial pH

= 5.0.

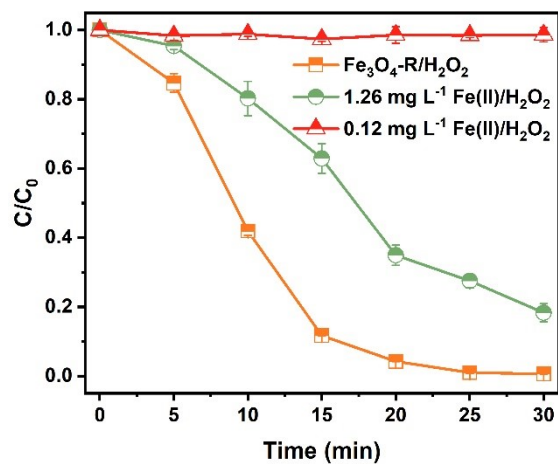


Fig. S4 Estimated contribution of homogeneous and heterogeneous systems to the BPA degradation in the $\text{Fe}_3\text{O}_4\text{-R}/\text{H}_2\text{O}_2$ system. Conditions: $[\text{catalyst}]_0 = 0.1 \text{ g L}^{-1}$, $[\text{H}_2\text{O}_2]_0 = 5 \text{ mM}$, $[\text{BPA}]_0 = 0.1 \text{ mM}$, and initial $\text{pH} = 5.0$.

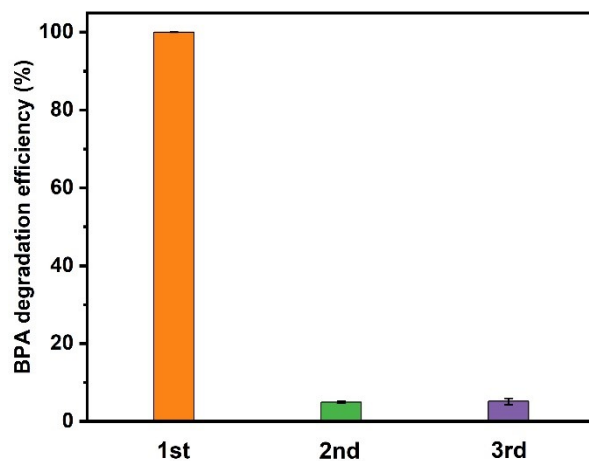


Fig. S5 Reuse performance of Fe₃O₄-R for Fenton-like degradation of BPA.

Conditions: [catalyst]₀ = 0.1 g L⁻¹, [H₂O₂]₀ = 5 mM, [BPA]₀ = 0.1 mM, and initial pH

= 5.0.

Reference

- [1] S. Zhang, X. Zhao, H. Niu, Y. Shi, Y. Cai and G. Jiang, Superparamagnetic Fe_3O_4 nanoparticles as catalysts for the catalytic oxidation of phenolic and aniline compounds, *J. Hazard. Mater.*, 2009, **167**, 560-566.
- [2] L. Xu and J. Wang, Fenton-like degradation of 2,4-dichlorophenol using Fe_3O_4 magnetic nanoparticles, *Appl. Catal. B: Environ.*, 2012, **123-124**, 117-126.
- [3] J. He, X. Yang, B. Men, Z. Bi, Y. Pu and D. Wang, Heterogeneous Fenton oxidation of catechol and 4-chlorocatechol catalyzed by nano- Fe_3O_4 : Role of the interface, *Chem. Eng. J.*, 2014, **258**, 433-441.
- [4] L. Hou, Q. Zhang, F. Jérôme, D. Duprez, H. Zhang and S. Royer, Shape-controlled nanostructured magnetite-type materials as highly efficient Fenton catalysts, *Appl. Catal. B: Environ.*, 2014, **144**, 739-749.
- [5] W. Li, Y. Wang and A. Irini, Effect of pH and H_2O_2 dosage on catechol oxidation in nano- Fe_3O_4 catalyzing UV-Fenton and identification of reactive oxygen species, *Chem. Eng. J.*, 2014, **244**, 1-8.
- [6] M. Minella, G. Marchetti, E. De Laurentiis, M. Malandrino, V. Maurino, C. Minero, D. Vione and K. Hanna, Photo-Fenton oxidation of phenol with magnetite as iron source, *Appl. Catal. B: Environ.*, 2014, **154-155**, 102-109.
- [7] R. Huang, Z. Fang, X. Yan and W. Cheng, Heterogeneous sono-Fenton catalytic degradation of bisphenol A by Fe_3O_4 magnetic nanoparticles under neutral condition, *Chem. Eng. J.*, 2012, **197**, 242-249.