

Electronic Supplementary Information
Direct hydrothermal synthesis of single-crystalline triangular nanoprisms structure of Fe_3O_4

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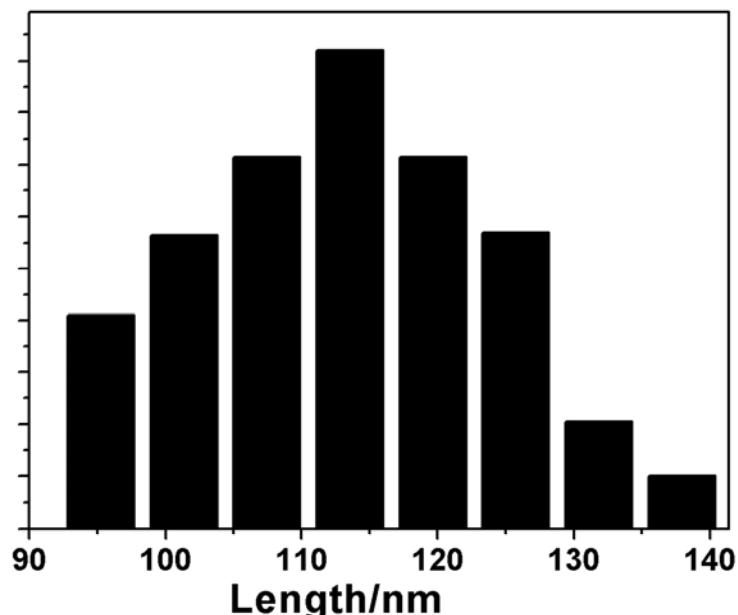


Fig. S1 The edge length distribution of the TNPs.

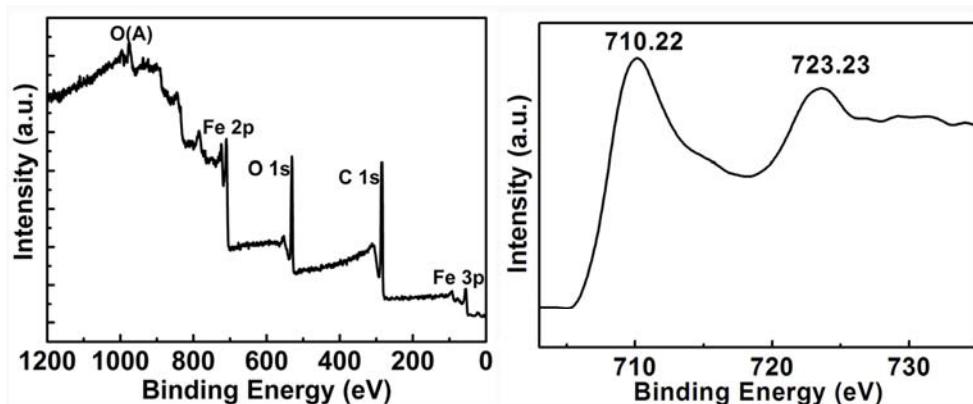


Fig. S2 (a) XPS spectrum of the as-prepared Fe_3O_4 TNPs. (b) The expanded spectrum of Fe 2p.

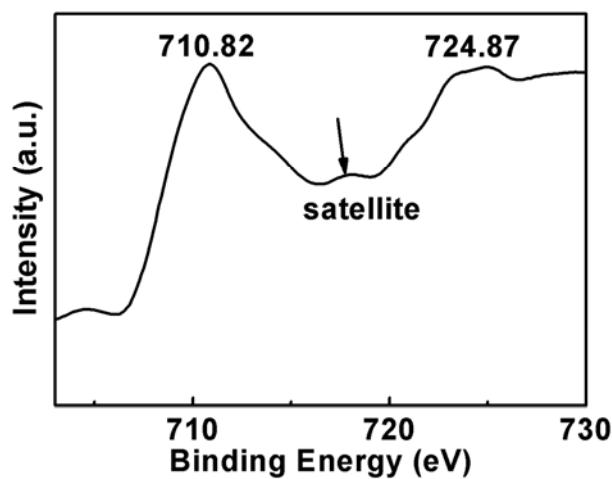


Fig. S3 XPS spectrum of Fe 2p for $\gamma\text{-Fe}_2\text{O}_3$.

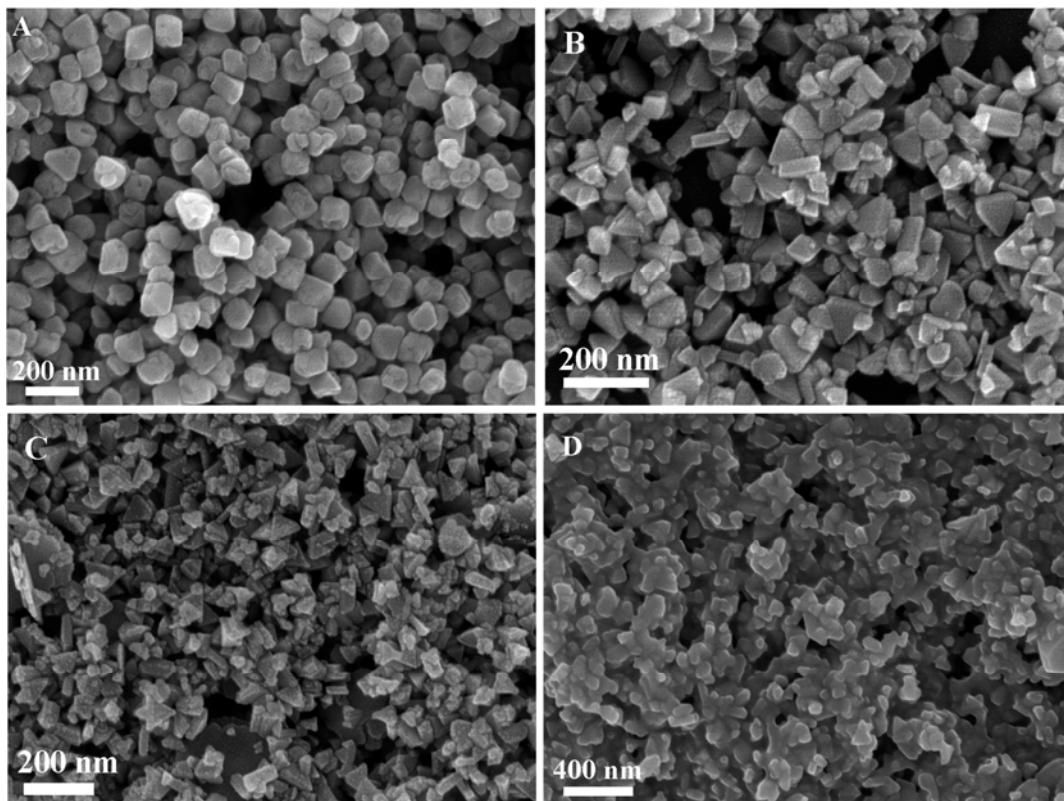


Fig. S4 SEM images of Fe_3O_4 prepared using the quantity of NaAc: (a) 1 g, (b) 2 g, (c) 4 g, (d) 5 g with other experiment parameters kept constant ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ 1 g, EG 20 mL, PDA 10 mL, at 200°C for 12 h).

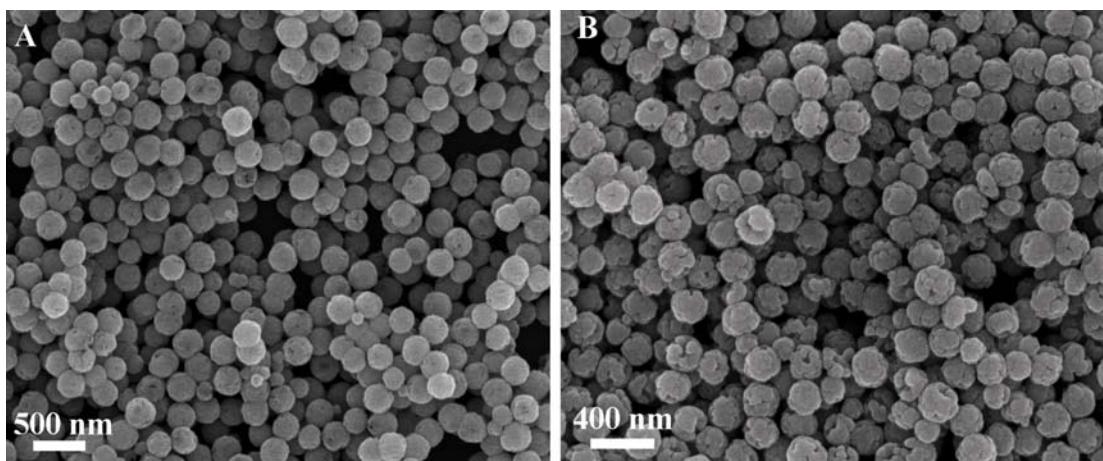


Fig. S5 SEM images of as-prepared Fe_3O_4 NCs using (a) 1,6-hexamethylene (b) ethylenediamine with other experiment parameters kept constant ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ 1 g, NaAc 3 g, EG 20 mL, at 200°C for 12 h).

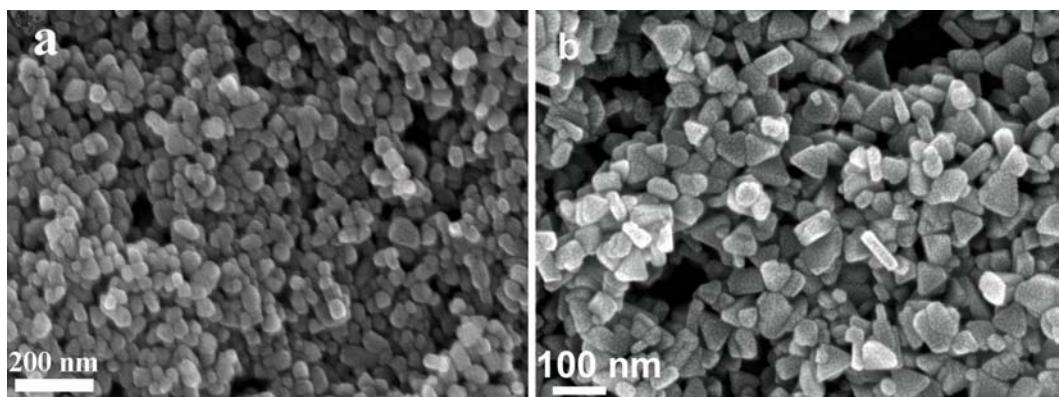


Fig. S6 (a) SEM image of Fe_3O_4 prepared using DEG as solvent with other experiment parameters kept constant ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ 1 g, NaAc 3 g, PDA 10 mL, at 200°C for 12 h). (b) SEM image of Fe_3O_4 prepared with the volume ratio of EG to PDA (mL): 20:2, with other experiment parameters kept constant ($\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ 1 g, NaAc 3 g, at 200°C for 12 h).