

Supporting Information

Ligand-Assisted Solid-State Transformation of Nanoparticles

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Synthesis of β -FeOOH ellipsoids.¹ The synthesis of β -FeOOH ellipsoids is based on a previously reported method with minor revisions. 0.2 M $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ and 10 g CTAC were dissolved in 100 ml water. Then, the solution was transferred into a 250 mL narrow-necked bottle and was heated at 87 °C for 24 hours. The final product was washed with water 3 times and collected by centrifugation. At last, the precipitate was dispersed in 25 mL of water.

Synthesis of β -FeOOH nanorods.² 0.1M $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ was dissolved in 400 mL distilled water. Then the solution was heated at 60 °C for 4 days. In the initial stage of the heating process, the temperature rises at a rate of 0.2 °C·min⁻¹ until it rises to 60 °C. The product was removed by centrifugation and washed with water twice and then dispersed in 25 mL water for future use.

Synthesis of Prussian blue nanocubes.³ The synthesis followed a previously reported method. 113.14 mg potassium ferricyanide and 3 g of PVP (Mw.40,000) were added to a 40 mL 0.1 M HCl solution. After stirring at room temperature for 30 min, the system was heated at 80°C for 20 hours. The blue product was then collected by centrifugation and dispersed in 25 mL water for future use.

Synthesis of $\text{Co}(\text{OH})_2$ nanoplates.⁴ $\text{Co}(\text{OH})_2$ nanoplates were synthesized based on a previous report with some modifications. Typically, 0.48 g of CoCl_2 was dissolved in 0.2 g of PVP (Mw=55,000) and 72 mL of H_2O , followed by adding 0.56 g of hexamethylenetetramine and 8 mL of ethanol. The mixture was heated at 95 °C for 2 h. The products were collected by centrifugation, washed with H_2O and redispersed in 40 mL of H_2O .

Synthesis of $\text{Ni}(\text{OH})_2$ nanoplates.⁵ To synthesize the 50 nm $\text{Ni}(\text{OH})_2$ nanoplates, an aqueous solution of 10 mol·mL⁻¹ $\text{Ni}(\text{NO}_3)_2$ and 20 mol·mL⁻¹ $\text{NH}_3 \cdot \text{H}_2\text{O}$ was heated in an oven at 90°C for 2 hours. And then the solution was washed with water several times and dispersed in 25 mL water for the next step.

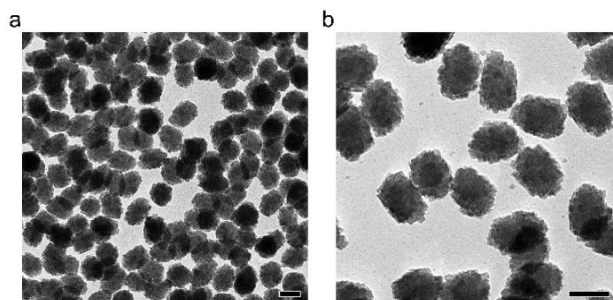


Figure S1. TEM images of the original β -FeOOH ellipsoids. Scale bars: 100 nm.

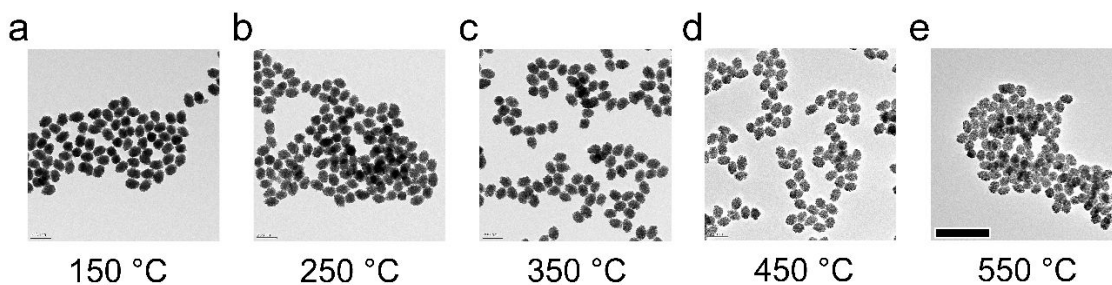


Figure S2. TEM images of the PAA-modified β -FeOOH ellipsoids calcined at various temperatures for 2h. Scale bar: 500 nm.

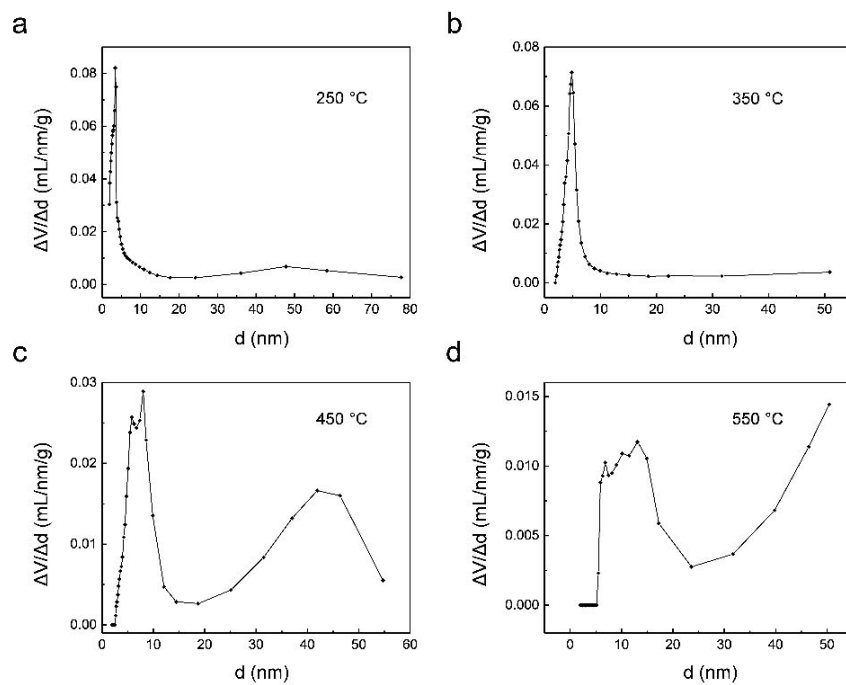


Figure S3. Pore size distribution curves of PAA-modified β -FeOOH ellipsoids calcined at various temperatures for 2h.

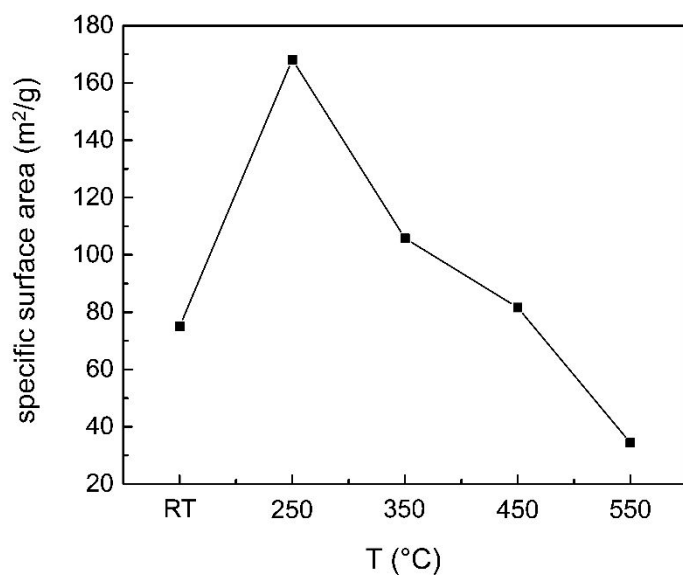


Figure S4. Specific surface area of the original β -FeOOH ellipsoids and the samples calcined in air at various temperatures for 2h.

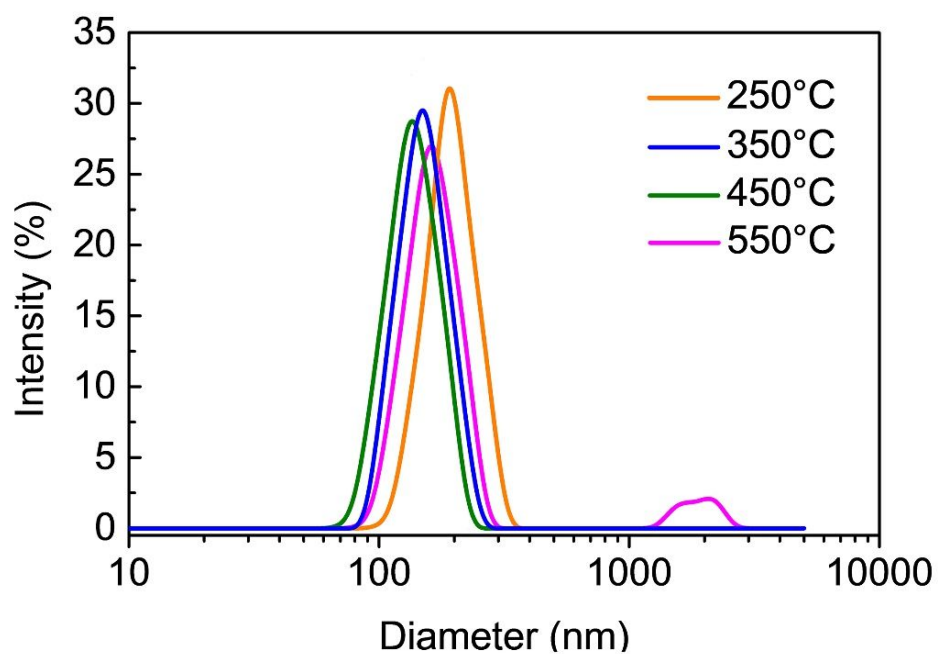


Figure S5. The size distributions of PAA-modified ellipsoids calcined for 2h at 250 °C, 350 °C, 450 °C and 550 °C.

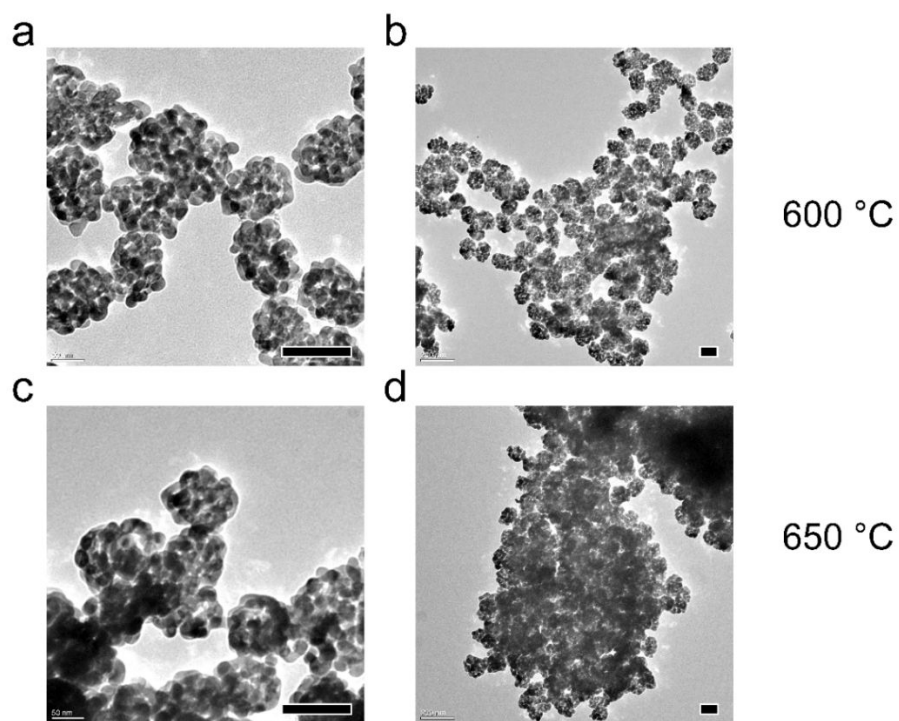


Figure S6. Morphologies of the β -FeOOH ellipsoids calcined at 600 °C (**a** and **b**) and 650 °C (**c** and **d**) for 2h. Scale bars: 100 nm.

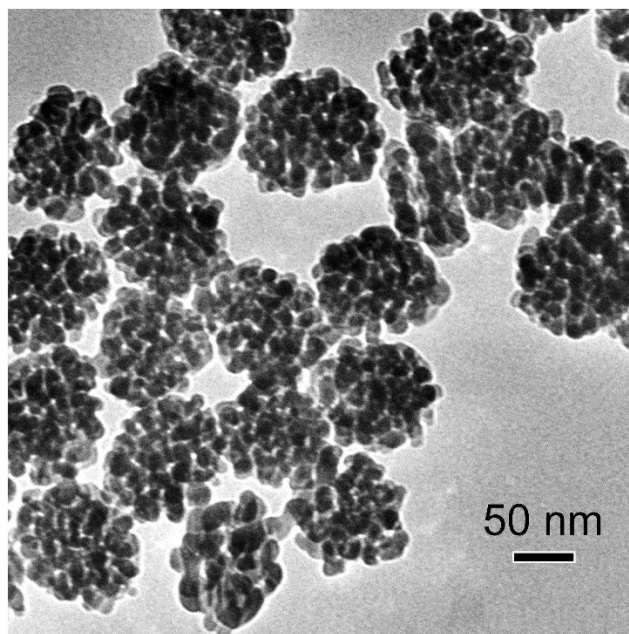


Figure S7. TEM image of the Fe_2O_3 ellipsoids with a secondary PAA modification calcined in air at 550 °C for 2h.

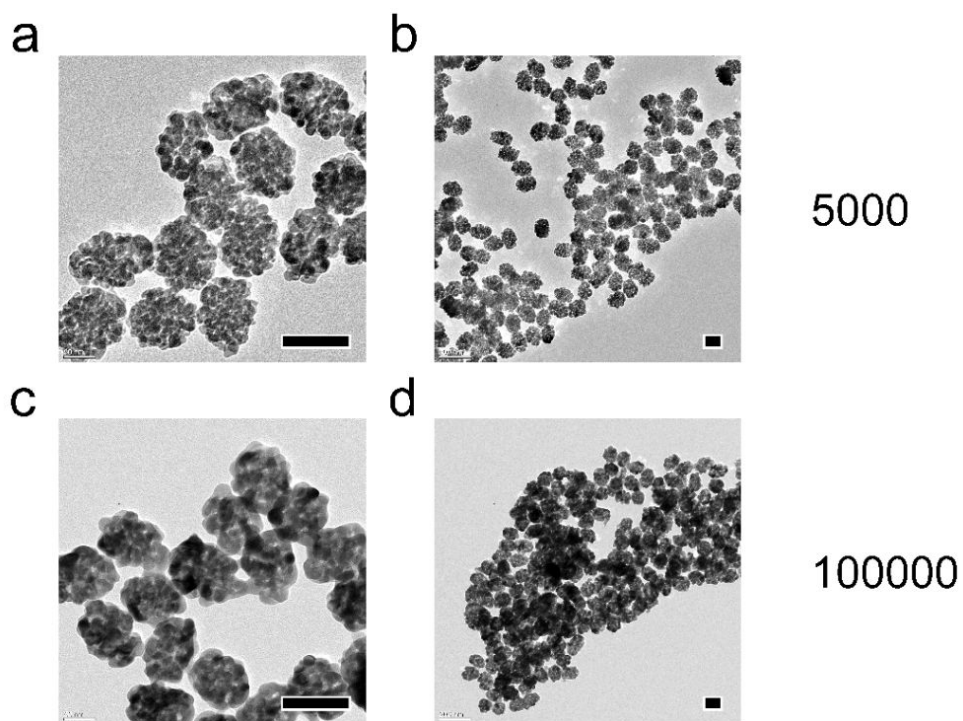


Figure S8. PAA with a molecular weight of 5000 (**a** and **b**) and 100,000 (**c** and **d**) were used to modify the FeOOH ellipsoids. Then these nanoparticles were calcined at 600 °C for 2h. Scale bars: 100 nm.

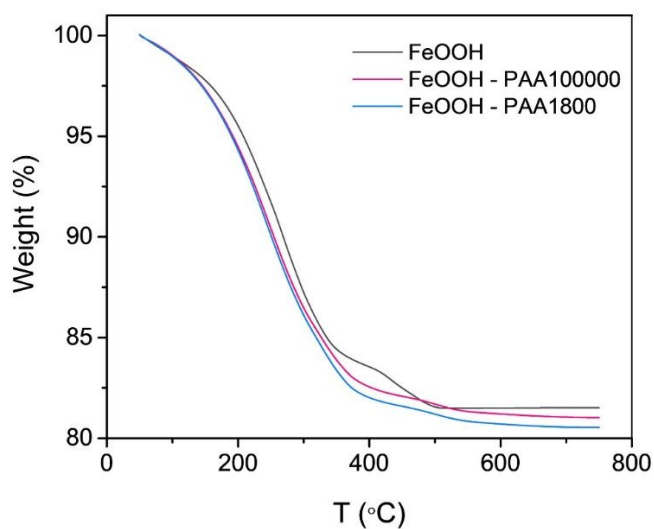


Figure S9. TGA measurements of FeOOH ellipsoids and FeOOH ellipsoids modified by PAA with molecular weight of 1800 and 100000.

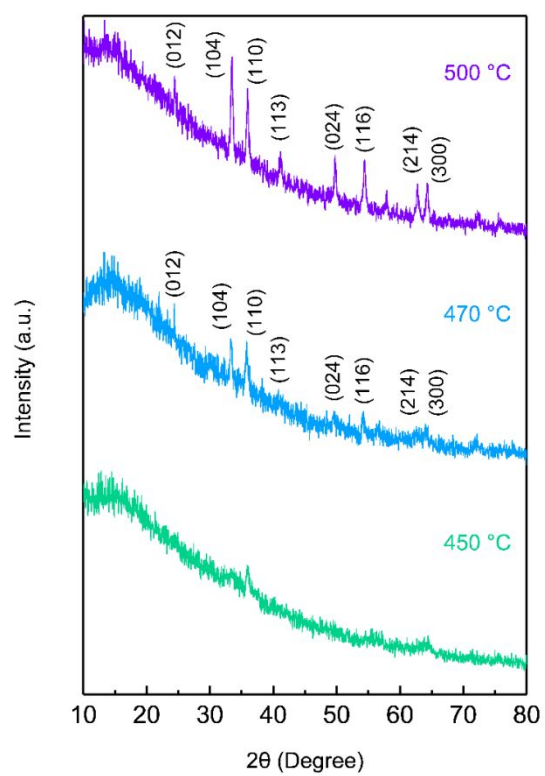


Figure S10. XRD patterns of PAA-modified β -FeOOH ellipsoids calcined at 450 °C, 470 °C, and 500 °C.

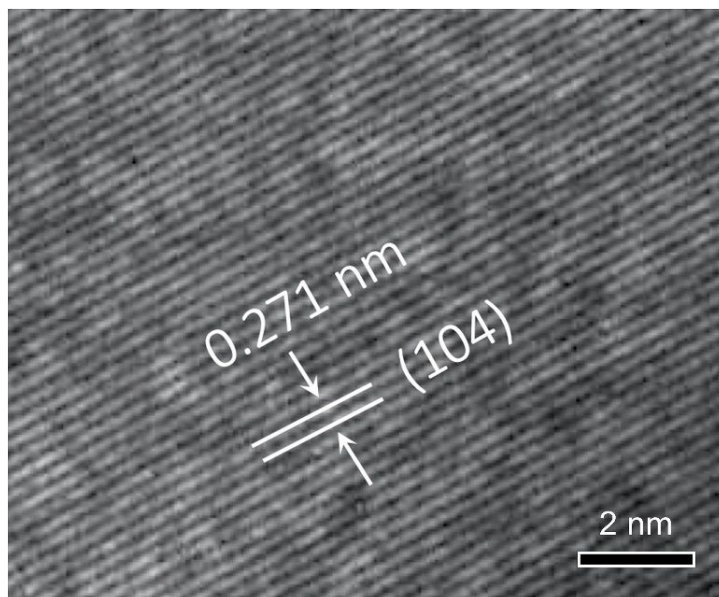


Figure S11. HRTEM image of Fe_2O_3 ellipsoids.

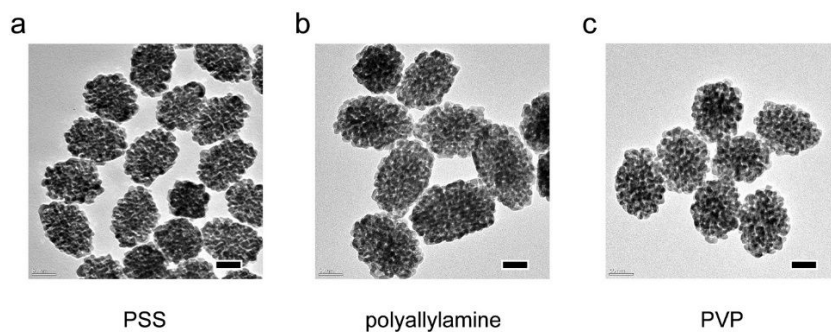


Figure S12. β -FeOOH ellipsoids were modified by PSS (a), polyallylamine (b) and PVP (c) and then calcined at 350 °C for 2h. Scale bars: 50 nm.

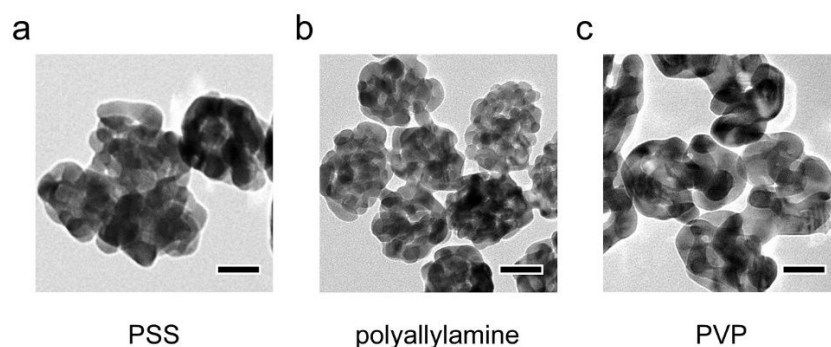


Figure S13. β -FeOOH ellipsoids were modified by PSS (a), polyallylamine (b) and PVP (c) and then calcined at 550 °C for 2h. Scale bars: 50 nm.

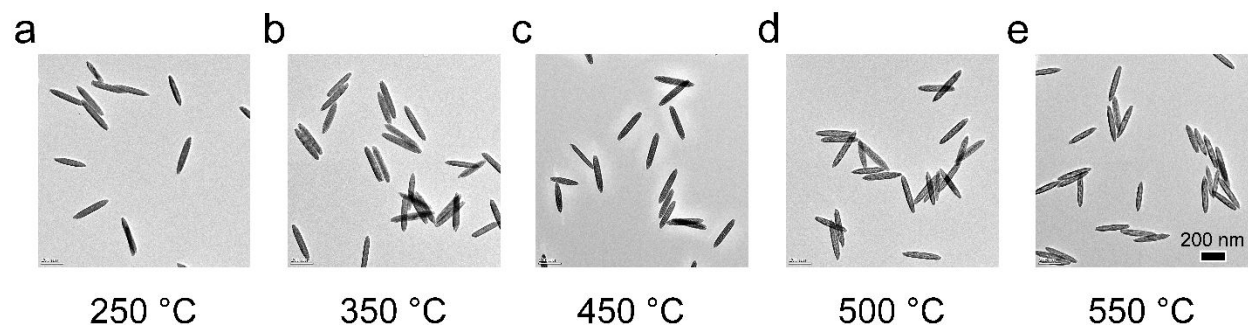


Figure S14. The morphology evolution of the β -FeOOH rods after calcination for two hours at a gradient temperature from 250 °C to 550 °C with an interval of 100 °C.

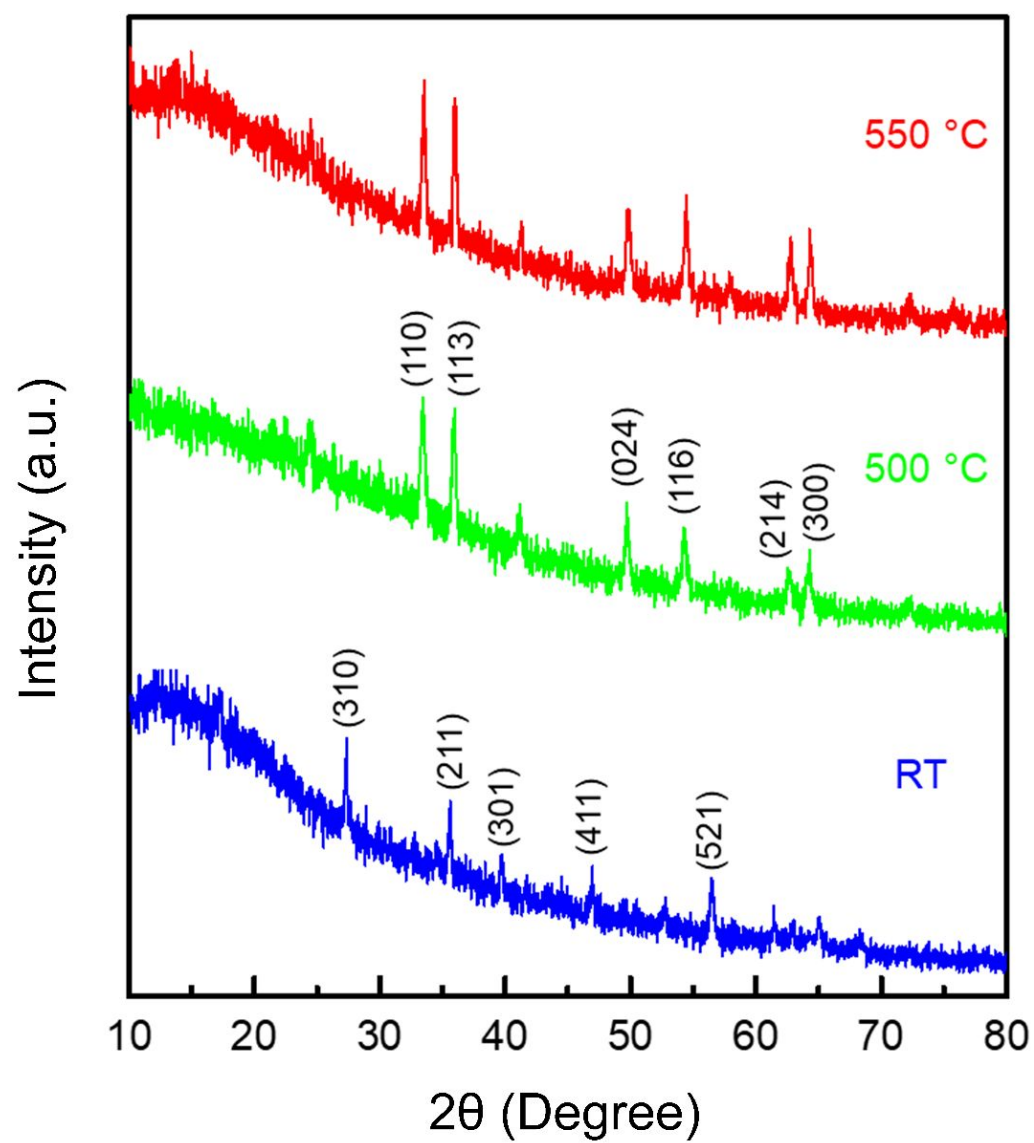


Figure S15. XRD patterns of PAA-modified β -FeOOH rods and the samples calcined at 500 °C and 550 °C.

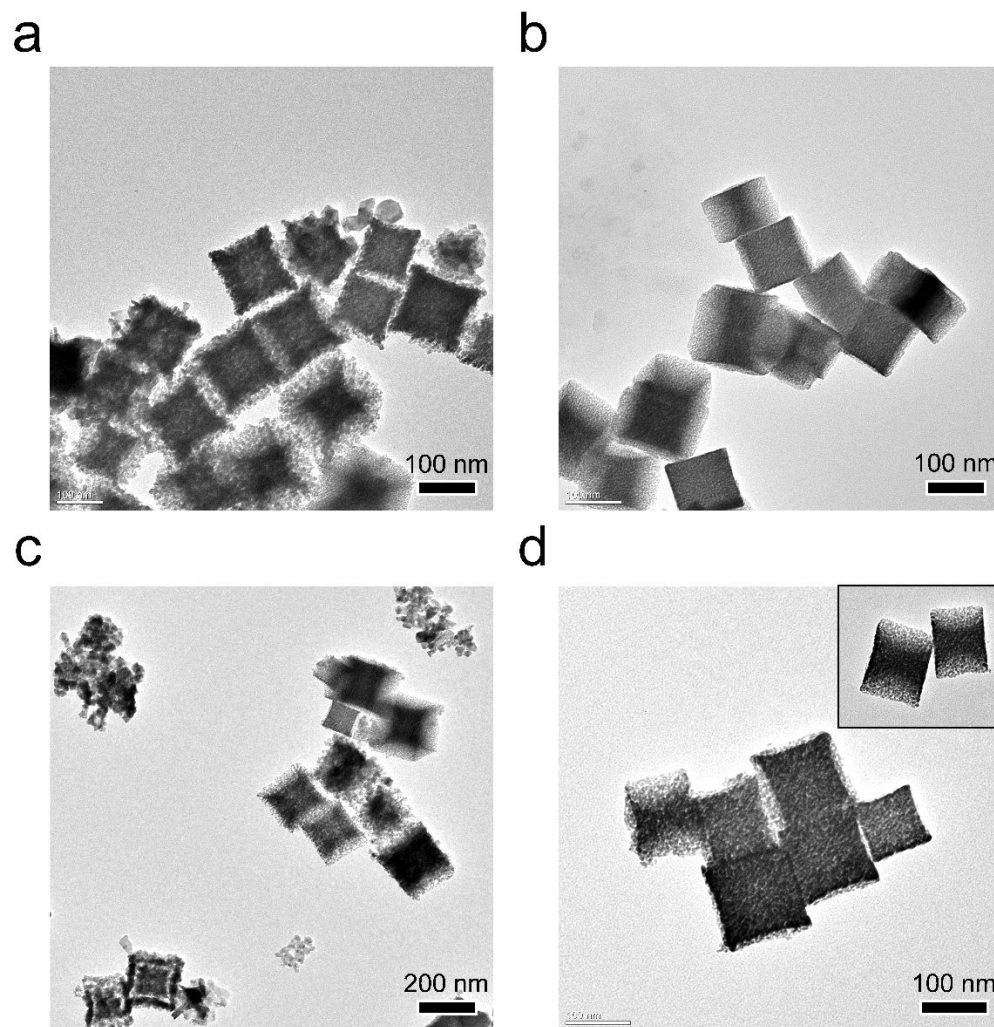


Figure S16. **a** and **c** are TEM images of original PB nanocubes calcined at 250 °C and 350 °C, respectively. PB nanocubes capped with PAA were also heated at 250 °C (**b**) and 350 °C (**d**).

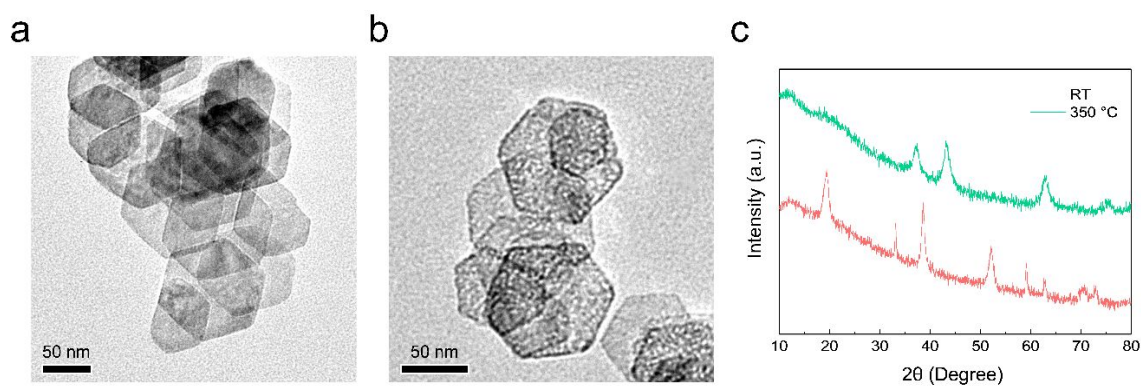


Figure S17. Morphology of PAA-modified Ni(OH)₂ nanoplates before (**a**) and after (**b**) calcination at 350 °C for 2h. **c**, XRD patterns of PAA-modified Ni(OH)₂ nanoplates annealed at 350 °C.

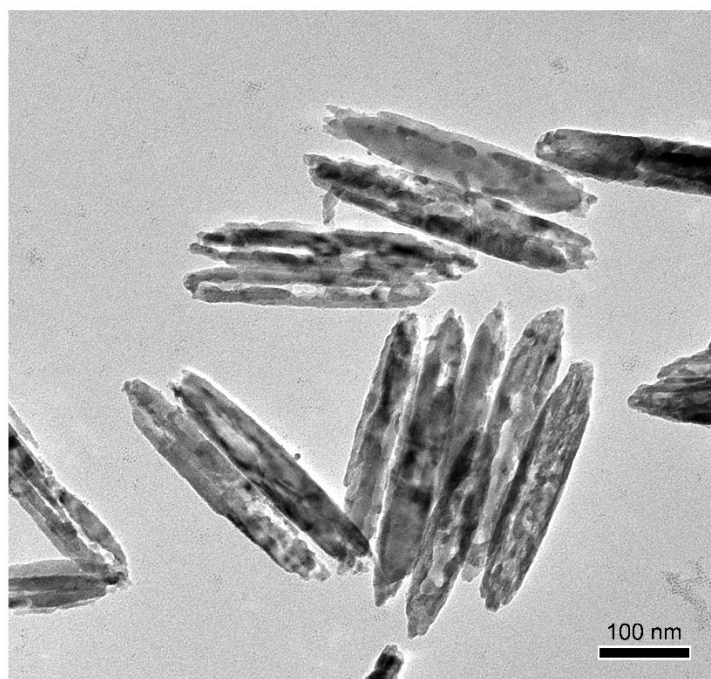
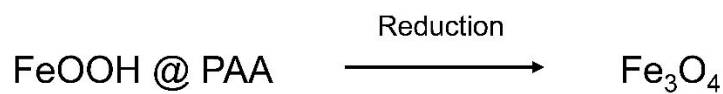


Figure S18. Direct reduction of PAA-modified FeOOH rods in forming gas at 330 °C for 2h.

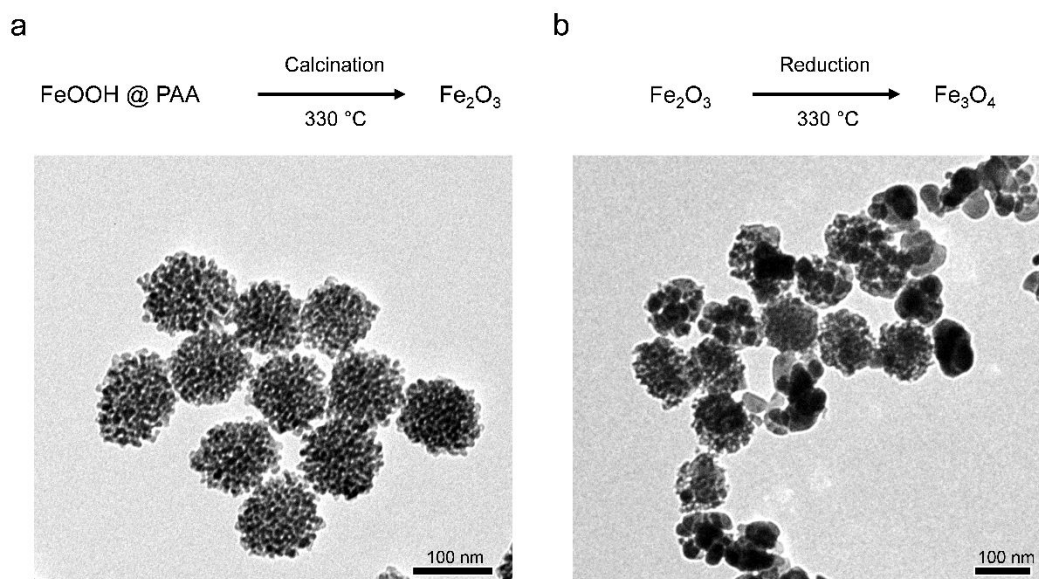


Figure S19. PAA-modified FeOOH ellipsoids were heated in air at 330 °C for 2h. Then the obtained nanoparticles (**a**) were reduced in forming gas directly without a secondary PAA modification (**b**).

References:

1. Wang, M.; He, L.; Xu, W.; Wang, X.; Yin, Y. *Angew Chem Int Ed Engl* **2015**, 54, (24), 7077-81.
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