## **Supporting Information**

## Fe<sub>3</sub>O<sub>4</sub> Nanoparticles entrapped in the inner Surfaces of N-doped Carbon

## microtubes with enhanced biomimetic activity

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## Synthesis of MoO<sub>3</sub>@PPy@FeOOH and NCMTs@Fe<sub>3</sub>O<sub>4</sub>

The preparation of MoO<sub>3</sub>@PPy@FeOOH was based on a similar method for synthesizing MoO<sub>3</sub>@FeOOH. Then, the as-synthesized MoO<sub>3</sub>@PPy@FeOOH (50 mg) and ammonia aqueous solution (1 mL) were dispersed in 20 mL deionized water. After stirring for 10 min, the products were collected and washed with water and ethanol for several times and then dried at 60 °C overnight. Subsequently, the obtained PPy@FeOOH microtubes were annealed at 700 °C temperatures under N<sub>2</sub> gas for 5 h to obtain the NCMTs@Fe<sub>3</sub>O<sub>4</sub> composites.



Fig. S1. SEM images of MoO<sub>3</sub> (A, B), MoO<sub>3</sub>@FeOOH (C, D), MoO<sub>3</sub>@FeOOH@PPy (E, F), FeOOH@PPy (G, H), and Fe<sub>3</sub>O<sub>4</sub>@NCMTs-700 (I, J).



**Fig. S2.** XRD patterns of different products: (A) MoO<sub>3</sub> (B) MoO<sub>3</sub>@FeOOH(a), MoO<sub>3</sub>@FeOOH@PPy(b), FeOOH@PPy(c), Fe<sub>3</sub>O<sub>4</sub>@NCMTs-700(d).



Fig. S3. SEM images of Fe<sub>3</sub>O<sub>4</sub>@NCMTs-500 (A, B) and Fe<sub>3</sub>O<sub>4</sub>@NCMTs-900 (D, E); TEM images of Fe<sub>3</sub>O<sub>4</sub>@NCMTs-500 (C) and Fe<sub>3</sub>O<sub>4</sub>@NCMTs-900 (F).