

Supporting information

Construction of NCMTs@MoO₂/FeNi₃ Hierarchical Tubular Heterostructures for Enhanced Performance in Catalysis and Protein adsorption

Hongxin Wang^a, Lixian Guo^{b*}, Jianmin Pan^a, Jingli Xu^a, Xue-Bo Yin^a, Min Zhang^{*a}

^a College of Chemistry and Chemical Engineering, Shanghai University of Engineering Science, Shanghai 201620, China.

^b Jinan Children's Hospital, Jinan 250022, China.

Preparation of NCMTs@MoO₂/FeNi₃-1

In a typical reaction, 50mg of the as-prepared FeOOH@NiMoO₄ were dispersed in 25 mL methanol under ultrasound for 15 minutes. Then another solution containing 23 mg of hexachloro cyclophosphazene and 52 mg of 4, 4'-sulfonyldiphenol in 6 mL of methanol was added drop by drop. After stirring for 5 minutes, 80 μL of triethylamine was added drop wise and the solution was continued to stir for 8 h. The product was collected by centrifugation and washed with water and ethanol for several times before drying at 60 °C overnight. The obtained product was denoted as FeOOH@NiMoO₄@PZS. Then, the as-prepared FeOOH@NiMoO₄@PZS powder was placed in a ceramic boat at the middle of a horizontal tube furnace. After heating at 150°C for 1 h and continuously increasing to 500°C and maintaining for 5 h with a ramp rate of 2°C min⁻¹ in N₂ gas, the obtained black powder was NCMTs@MoO₂/FeNi₃-1.

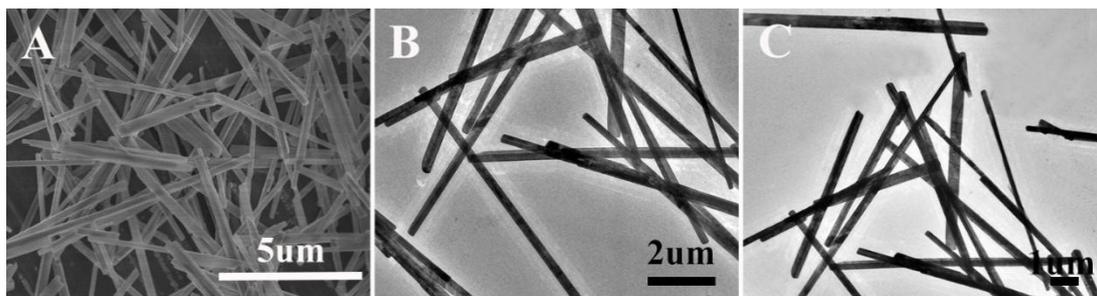


Fig. S1. SEM and TEM images of MoO_3

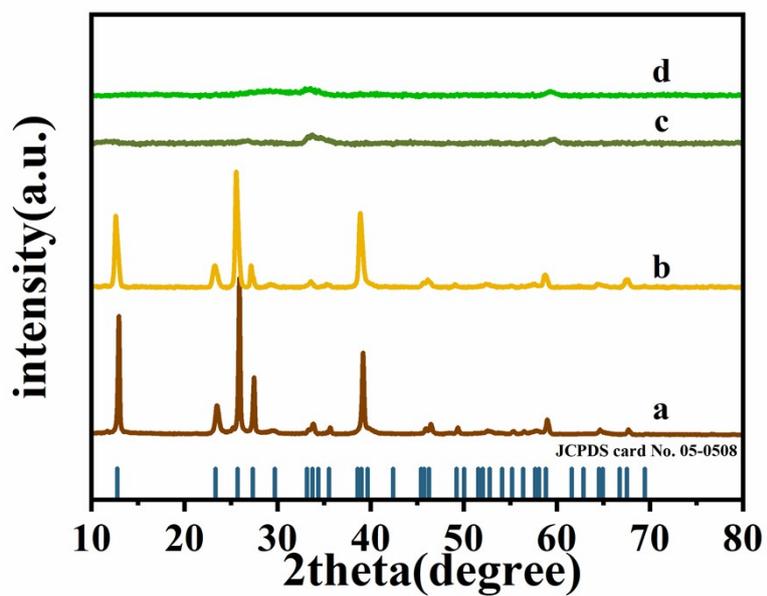


Fig. S2. XRD patterns of MoO_3 (a) and $\text{MoO}_3/\text{FeOOH}$ (b); $\text{FeOOH}/\text{NiMoO}_4$ (c); $\text{FeOOH}/\text{NiMoO}_4@\text{PDA}$ (d)

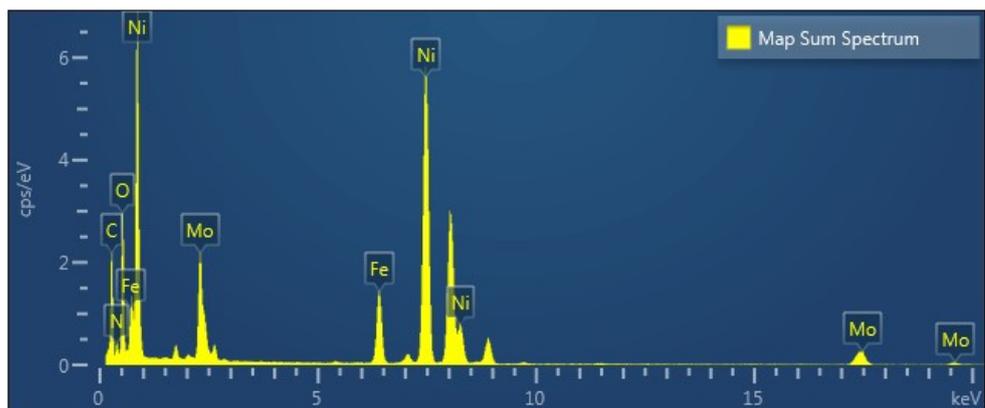


Fig. S3. Energy-dispersive X-rays spectrum of NCMTs@MoO₂/FeNi₃.

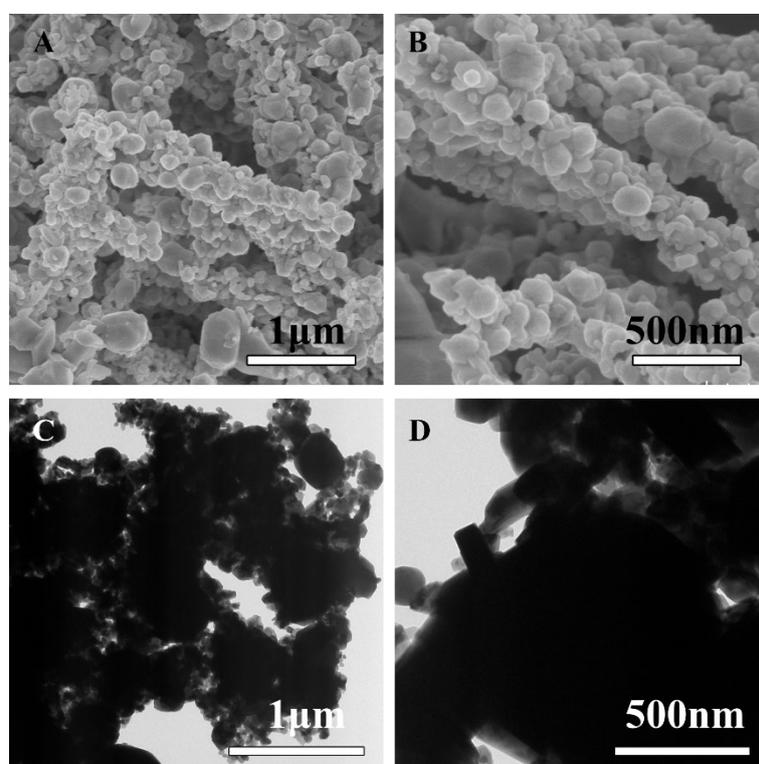


Fig. S4 (A, B) SEM images and (C, D) TEM images of Mo₂C@Fe_{0.64}Ni_{0.36}/Ni-900.

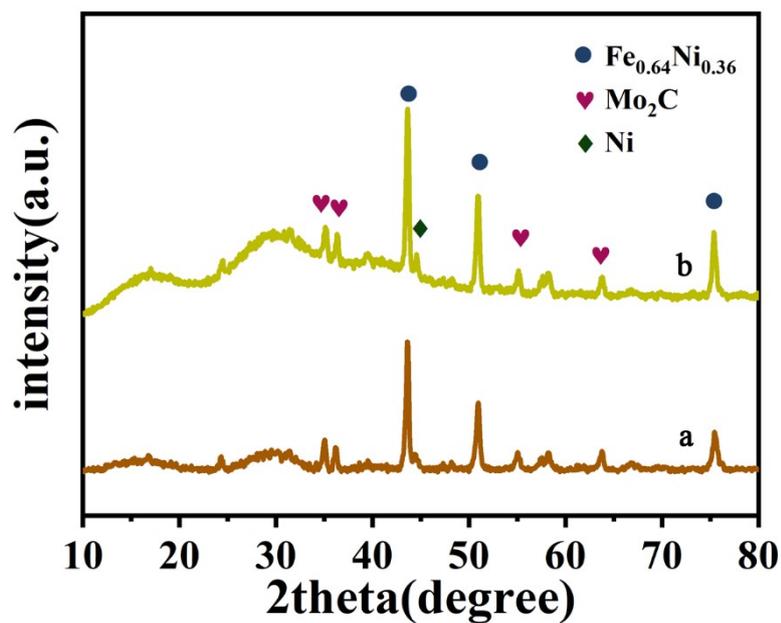


Fig. S5. XRD patterns of (a) $\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-700$ and (b) $\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-900$

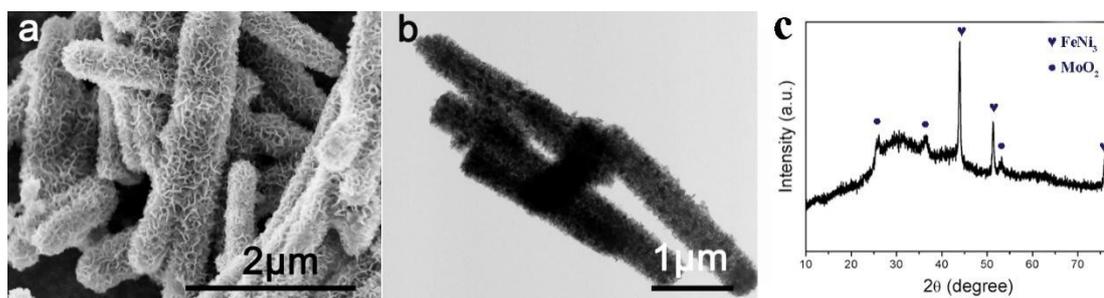


Fig. S6 SEM(a), TEM(b) images and XRD patterns(c) of $\text{NCMTs}@MoO_2/FeNi_3-1$

Table S1 ICP data and catalysis activity of $\text{NCMTs}@MoO_2/FeNi_3$, $\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-700$ and $\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-900$

Samples	Nickel content ($\mu\text{g}/\text{mg}$)	K ($\times 10^{-3}\text{s}^{-1}$)	κ ($\times 10^{-3}\text{mg}^{-1}\text{s}^{-1}$)
$\text{NCMTs}@MoO_2/FeNi_3$	429.11	17.12	39.90
$\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-700$	215.67	6.94	32.18
$\text{Mo}_2\text{C}@Fe_{0.64}Ni_{0.36}/Ni-900$	373.97	3.04	8.13

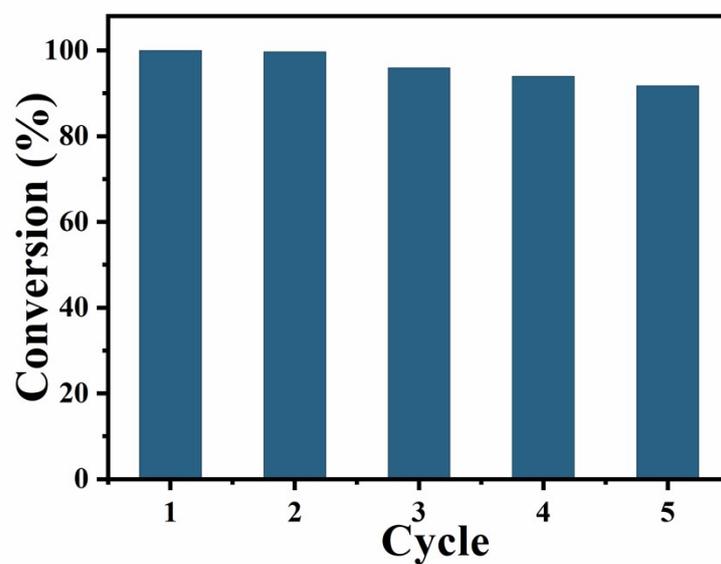


Fig. S7 The recyclability of the NCMTs@MoO₂/FeNi₃ as the catalyst for 4-nitrophenol

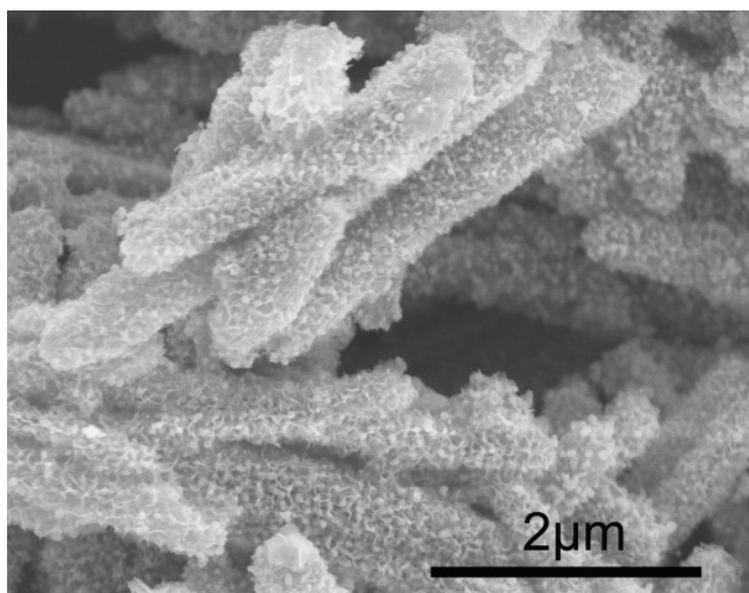


Fig. S8 SEM images of NCMTs@MoO₂/FeNi₃ after five catalytic reactions

Table S2. the estimate of Langmuir model and Freundlich model

Langmuir			Freundlich		
Q _m	b	R ²	Q _m	n	R ²
943.40	0.0319	0.9979	58.74	1.87	0.9540

