

Surface-modified Pt₁Ni₁-Ni(OH)₂ Nanoparticles with Abundant Pt-Ni(OH)₂ Interfaces Enhance Electrocatalytic Properties

Yuchen Qin ^{a,b*}, Hongying Zhuo ^b, Xiaoyu Liang ^a, Kuomiao Yu ^b, Yao Wang ^b, Daowei Gao ^{c*}, Xin Zhang ^{b*}

^a College of Science, Henan Agricultural University, Zhengzhou 450000, China.

^b State Key Laboratory of Heavy Oil Processing, College of Chemical Engineering, China University of Petroleum, Beijing 102249, China.

^c School of Chemistry and Chemical Engineering, University of Jinan, Jinan 250022, China.

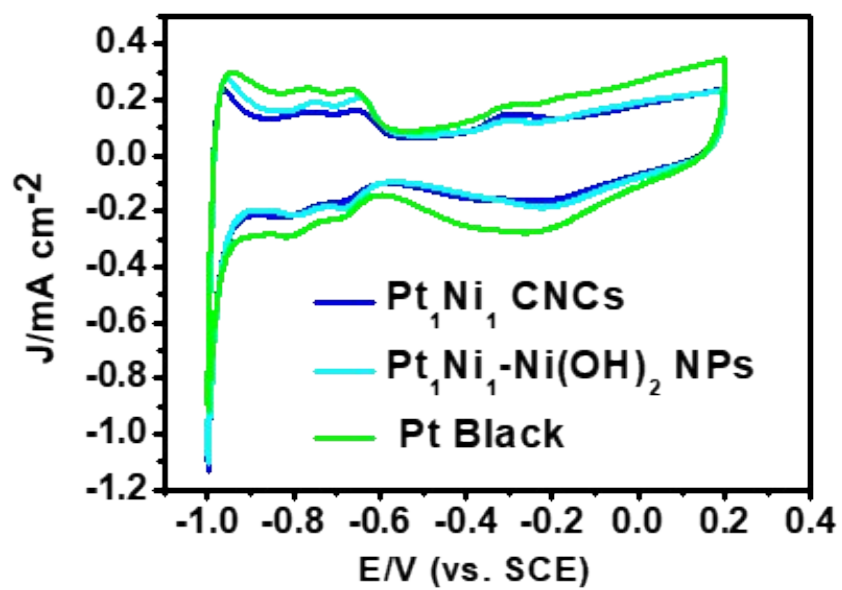


Fig. S1. CV curves of the Pt_1Ni_1 CNCs and $\text{Pt}_1\text{Ni}_1\text{-Ni(OH)}_2$ NPs and Pt Black in a N_2 -Purged 0.1M NaOH solution at a scan rate of 50 mV s^{-1} .

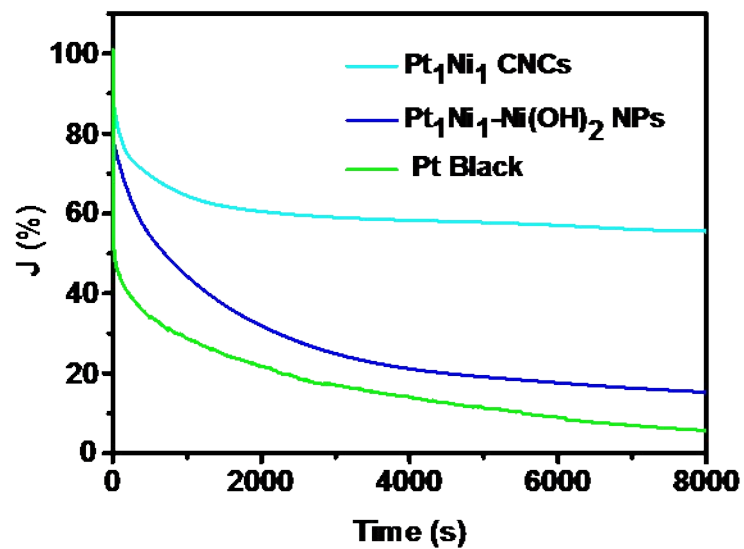


Fig. S2. I-t curves at 0.6 V vs. RHE for methanol electrooxidation in a mixture of 0.1 M NaOH and 1 M CH₃OH of as-prepared Pt₁Ni₁ CNCs, Pt₁Ni₁-Ni(OH)₂ NPs and commercial Pt Black.

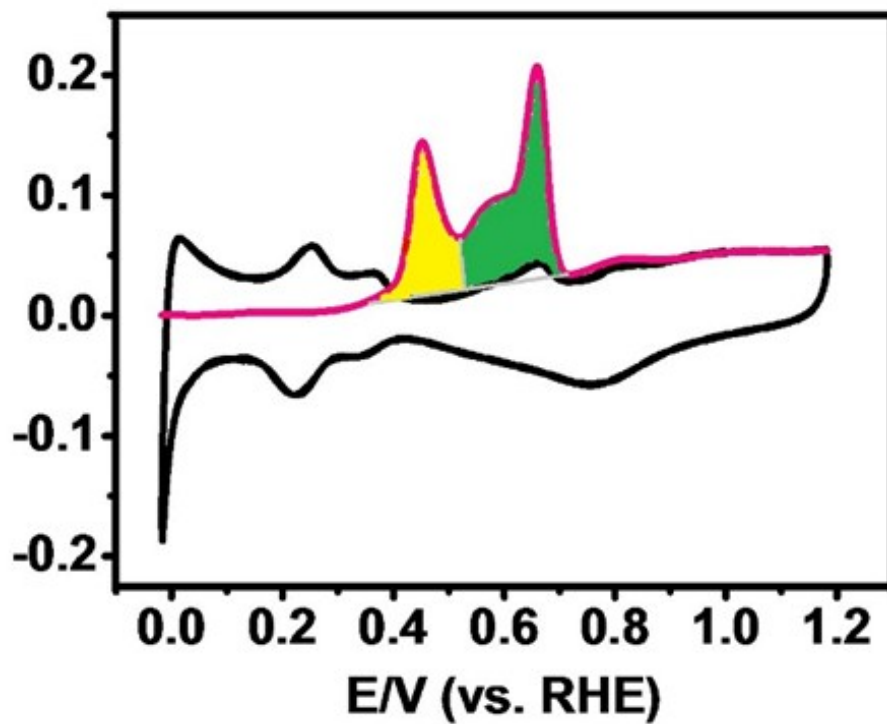


Fig. S3. The integration of CO oxidation peak areas of Pt₁Ni₁-Ni(OH)₂ NPs.

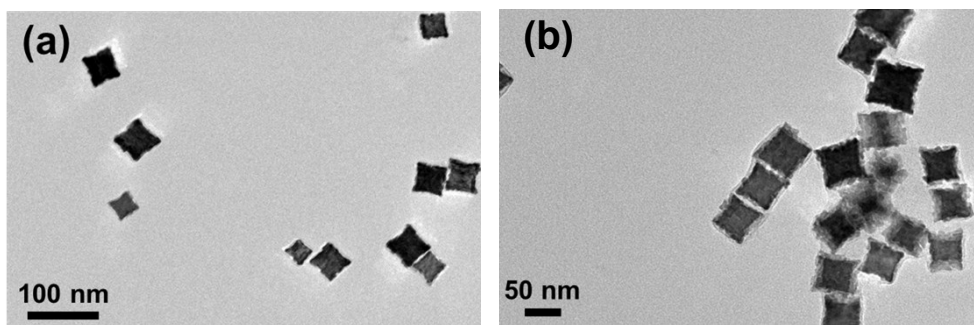


Fig. S4. Representative TEM images of (a) Pt₃Ni CNCs and (b) Pt₃Ni-Ni(OH)₂ NPs.

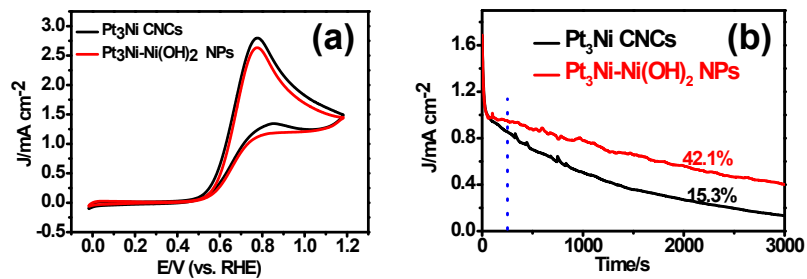


Fig. S5. (a) Cyclic voltammograms and (b) i-t curves (at 0.6 V vs. RHE) of as-prepared Pt₃Ni CNCs and Pt₃Ni-Ni(OH)₂ CNCs of methanol electrooxidation in a mixture of 0.1 M NaOH and 1 M CH₃OH at a scan rate of 50 mV s⁻¹.

Table S1. The components of as-prepared catalysis measured by ICP-OES

Samples	Pt/Ni molar ratio
Pt ₁ Ni ₁ -Ni(OH) ₂ NPs	62.7/37.3
Pt ₃ Ni-Ni(OH) ₂ NPs	81.4/18.6

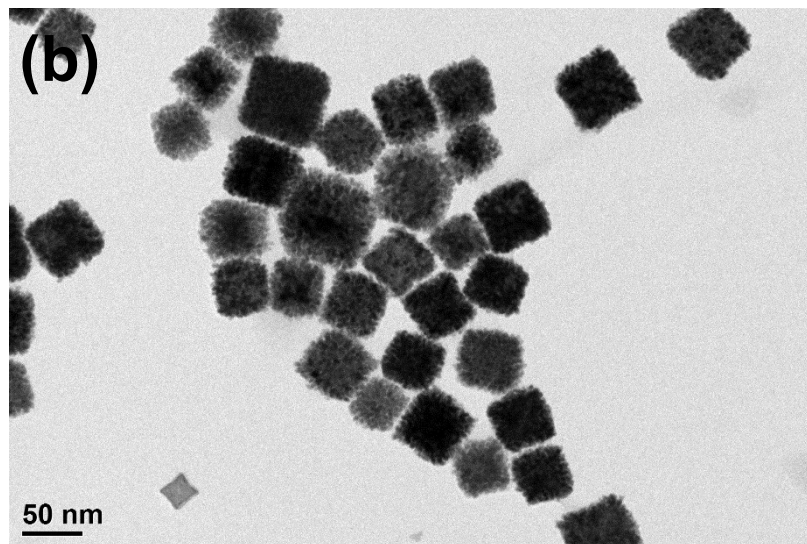
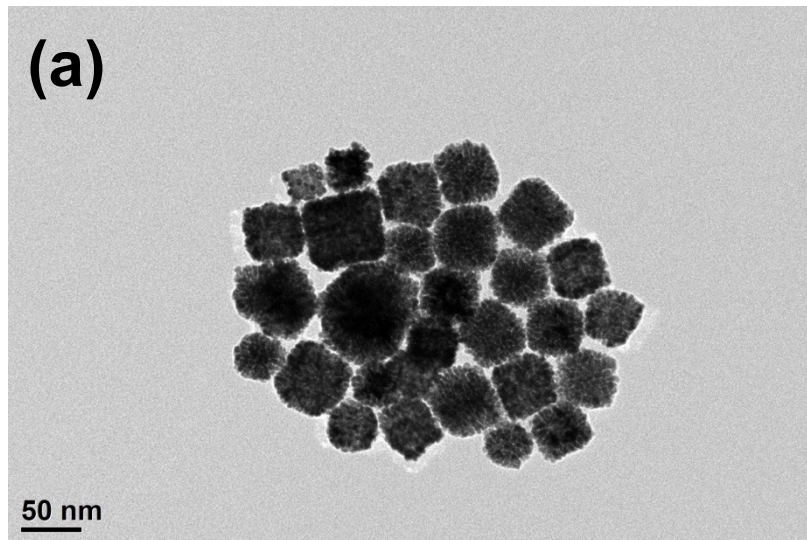


Fig. S6. (a) and (b) TEM images of PtNi₃ NPs.