

Bimetallic Pt-M (M = Fe, Co, Ni) nanobunches assembled by ultrathin nanowires with strong synergy and rich surface defects for enhanced methanol oxidation electrocatalysis

Suwen Li^{a,b,1}, Yu Zhang^{a,b,1}, Yulan Liu^{a,b}, Fangfei Lv^{a,b}, Yiming Yan^{a,b}, Baocang Liu^c,
Lili Huo^{a,b*}

^aChemical Engineering College, Inner Mongolia University of Technology, 49 Aimin Street, Hohhot 010051, P. R. China

^bKey Laboratory of CO₂ Resource Utilization at Universities of Inner Mongolia Autonomous Region, 49 Aimin Street, Hohhot 010051, P. R. China

^cInner Mongolia Engineering and Technology Research Center for Catalytic Conversion and Utilization of Carbon Resource Molecules, School of Chemistry and Chemical Engineering, Inner Mongolia University, Hohhot 010021, P. R. China

*Corresponding Author: L. Huo, Email: HuoLL101@imut.edu.cn

Table S1. Actual contents of elements in different catalysts as determined by EDS, XPS and ICP-MS results

Catalysts	Pt/M			
	Designed ratio	EDS	ICP	XPS
Pt ₃ Co	75.00/25.00	79.80/20.20	75.91/24.09	71.15/28.55
PtCo	50.00/50.00	52.61/47.39	51.87/48.13	46.06/53.94
PtCo ₃	25.00/75.00	21.67/78.33	23.62/76.38	20.15/79.85
Pt ₃ Ni	75.00/25.00	77.16/22.84	72.49/27.51	70.33/29.67
PtNi	50.00/50.00	37.86/62.84	37.37/62.63	34.37/65.63
PtNi ₃	25.00/75.00	21.32/78.68	23.35/76.65	19.71/80.29
Pt ₃ Fe	75.00/25.00	69.62/30.38	68.39/31.61	64.23/35.77
PtFe	50.00/50.00	45.92/54.08	46.28/53.72	42.34/57.66
PtFe ₃	25.00/75.00	29.47/70.53	28.66/71.34	31.98/68.02

Table S2. The MOR performance of the catalysts in acidic and alkaline media

Catalyst	Media	Onset potential/ V	$j_p/$ $\text{mA}\cdot\text{mg}^{-1}_{\text{Pt}}$	$j_p/$ $\text{mA}\cdot\text{cm}^{-2}$	ECSA $(\text{m}^2\cdot\text{g}^{-1}_{\text{Pt}})$
Pt₃Co	0.5 M H ₂ SO ₄	0.489	173.83	33.33	22.42
	1.0 M KOH	0.465	208.80	40.03	-
PtCo	0.5 M H ₂ SO ₄	0.449	632.96	78.90	74.75
	1.0 M KOH	0.375	589.85	67.50	-
PtCo₃	0.5 M H ₂ SO ₄	0.439	419.56	22.78	40.08
	1.0 M KOH	0.485	120.09	6.52	-
Pt₃Ni	0.5 M H ₂ SO ₄	0.469	223.57	42.37	24.15
	1.0 M KOH	0.475	274.15	51.96	-
PtNi	0.5 M H ₂ SO ₄	0.419	908.08	84.34	103.61
	1.0 M KOH	0.395	1162.21	107.65	-
PtNi₃	0.5 M H ₂ SO ₄	0.519	13.15	0.7	1.59
	1.0 M KOH	0.565	6.31	0.34	-
Pt₃Fe	0.5 M H ₂ SO ₄	0.479	225.82	39.09	23.17
	1.0 M KOH	0.455	348.92	60.39	-
PtFe	0.5 M H ₂ SO ₄	0.469	502.60	57.35	46.87
	1.0 M KOH	0.395	543.85	67.30	-
PtFe₃	0.5 M H ₂ SO ₄	0.539	140.74	12.15	18.78
	1.0 M KOH	0.415	360.48	31.06	-
Pt/C	0.5 M H ₂ SO ₄	0.499	255.90	34.48	44.45
	1.0 M KOH	0.465	222.30	29.95	-

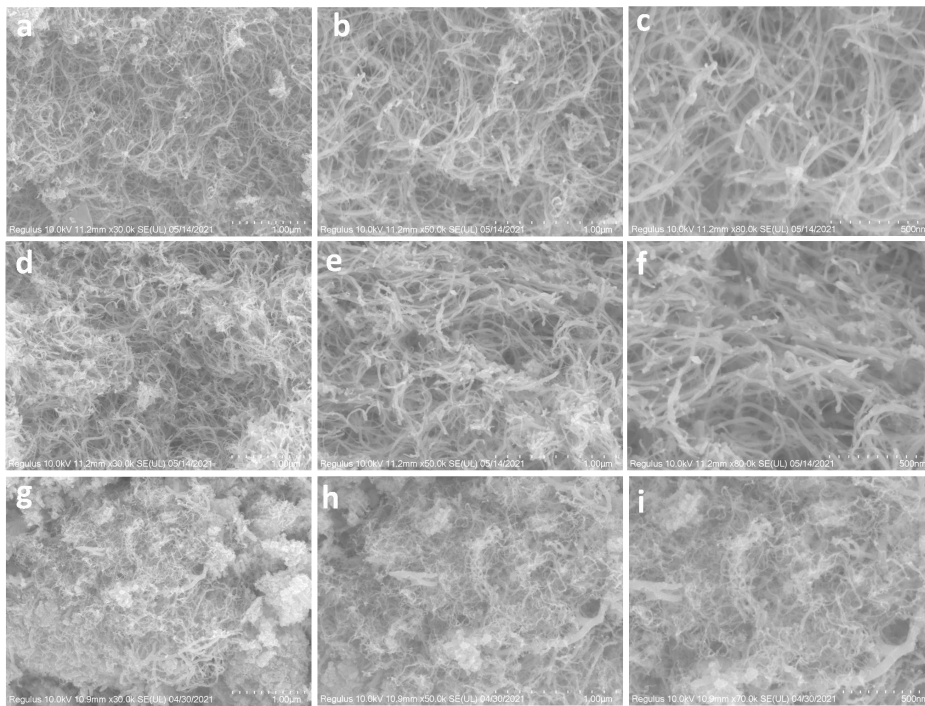


Figure S1. SEM images of Pt₃Co (a-c), Pt₃Fe (d-f) and Pt₃Ni NBs.

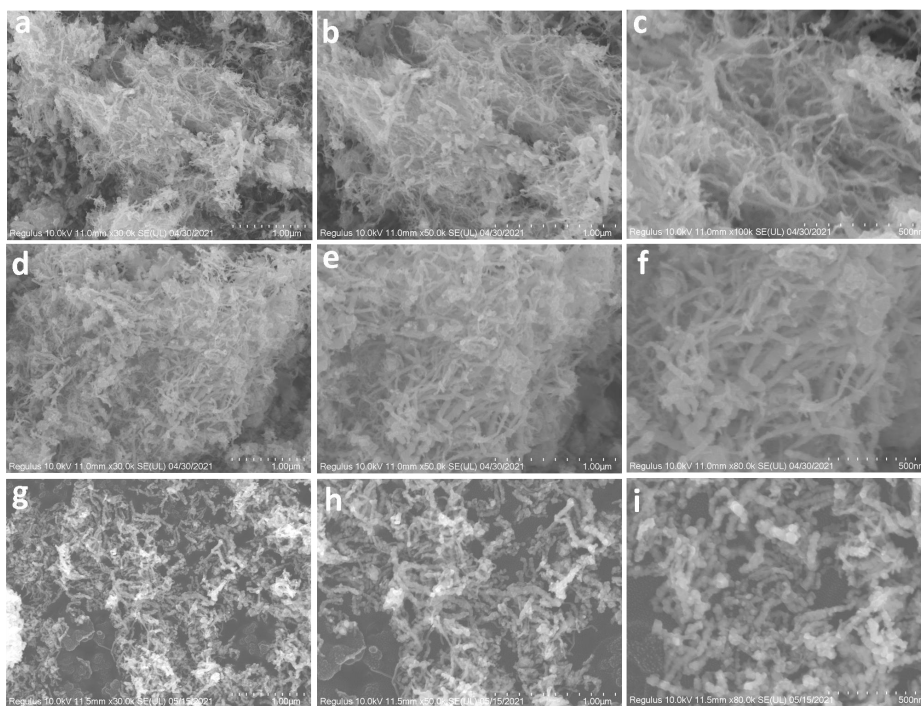


Figure S2. SEM images of PtCo₃ (a-c), PtFe₃ (d-f) and PtNi₃ NBs.

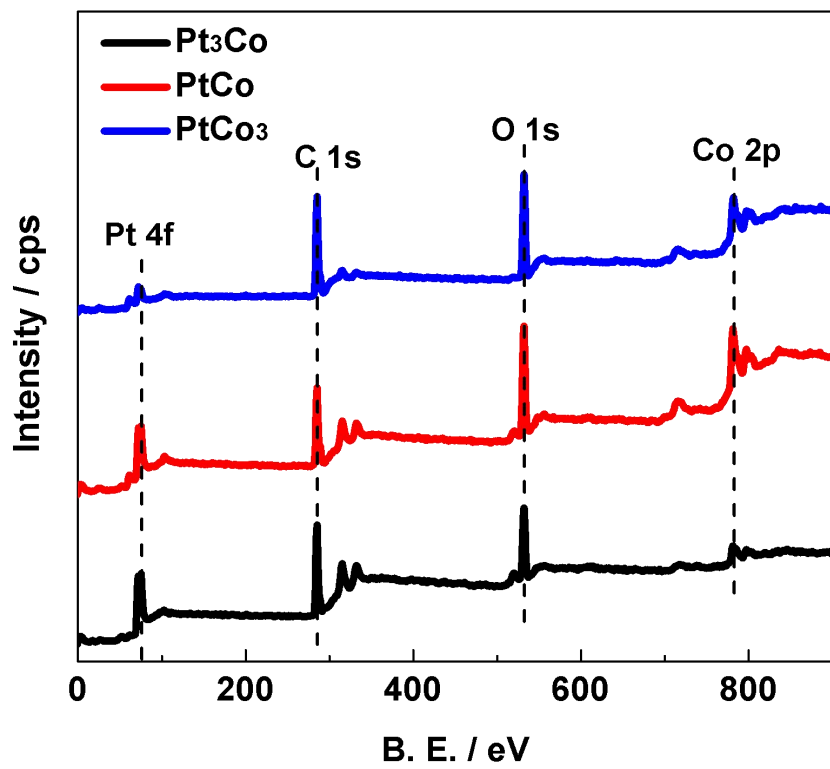


Figure S3. XPS survey spectra of PtCo NBs.

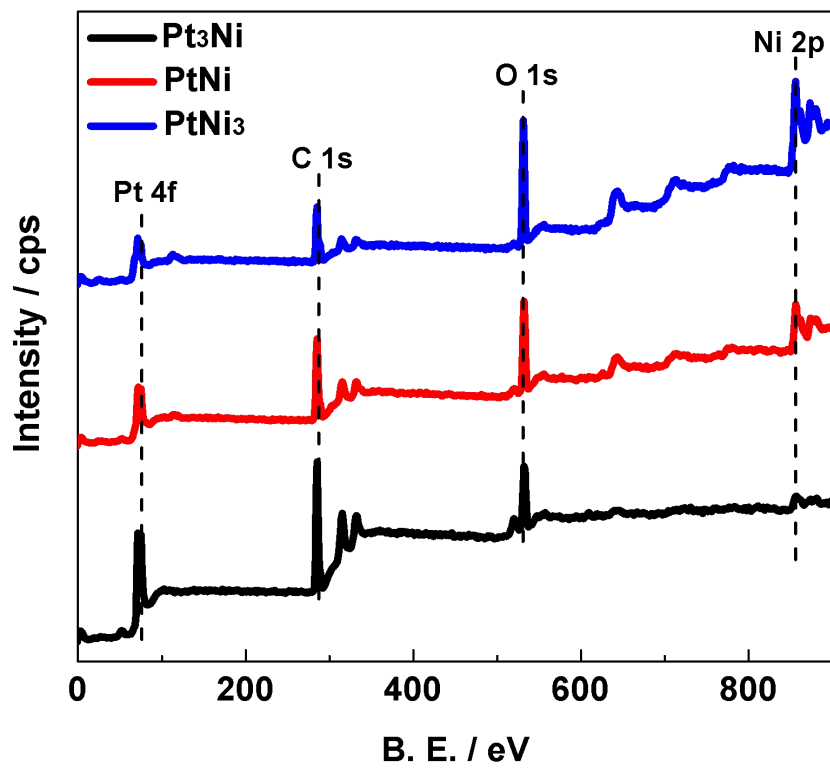


Figure S4. XPS survey spectra of PtNi NBs.

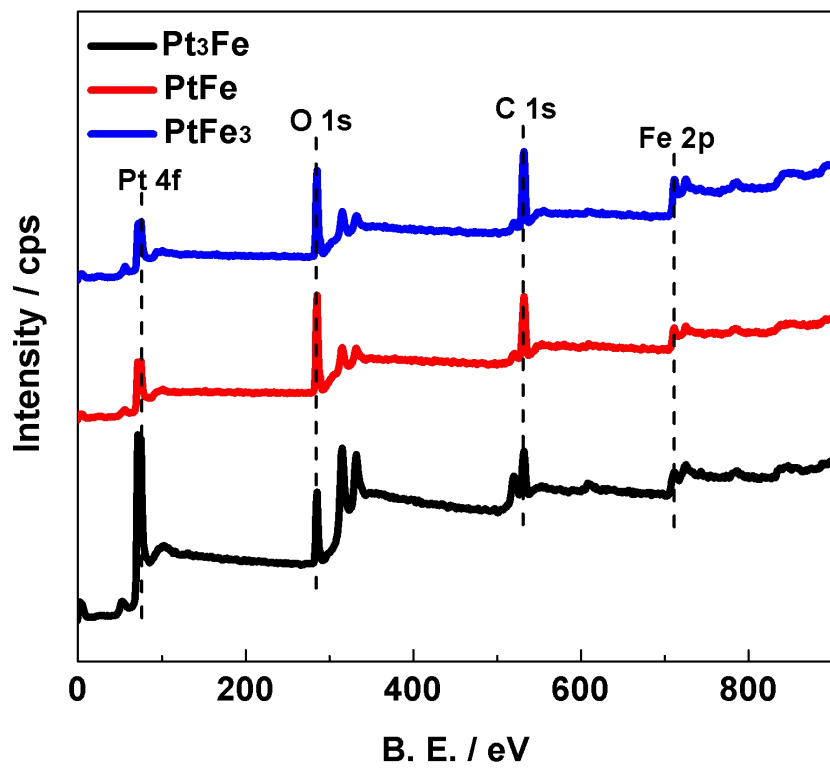


Figure S5. XPS survey spectra of PtFe NBs.

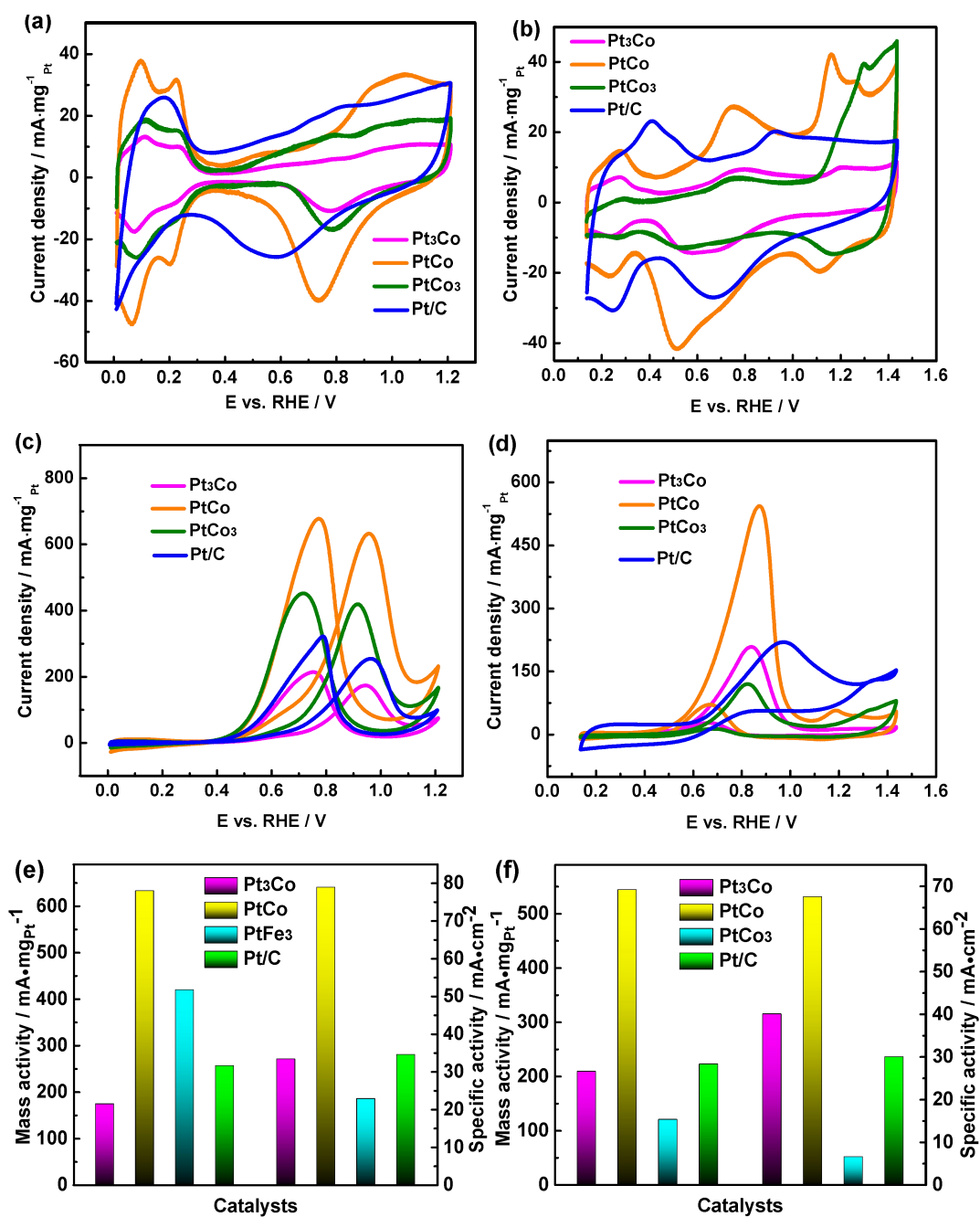


Figure S6. CVs of Pt-Co NBs in (a) 0.5 M H₂SO₄, (b) 0.5 M H₂SO₄ + 1.0 M CH₃OH, (c) 1.0 M KOH, and (d) 1.0 M KOH + 1.0 M CH₃OH. Mass- and area-specific activities of Pt-Co NBs at 0.929 and 0.835 V (vs. RHE) in (e) 0.5 M H₂SO₄ + 1.0 M CH₃OH and (f) 1.0 M KOH + 1.0 M CH₃OH, respectively.

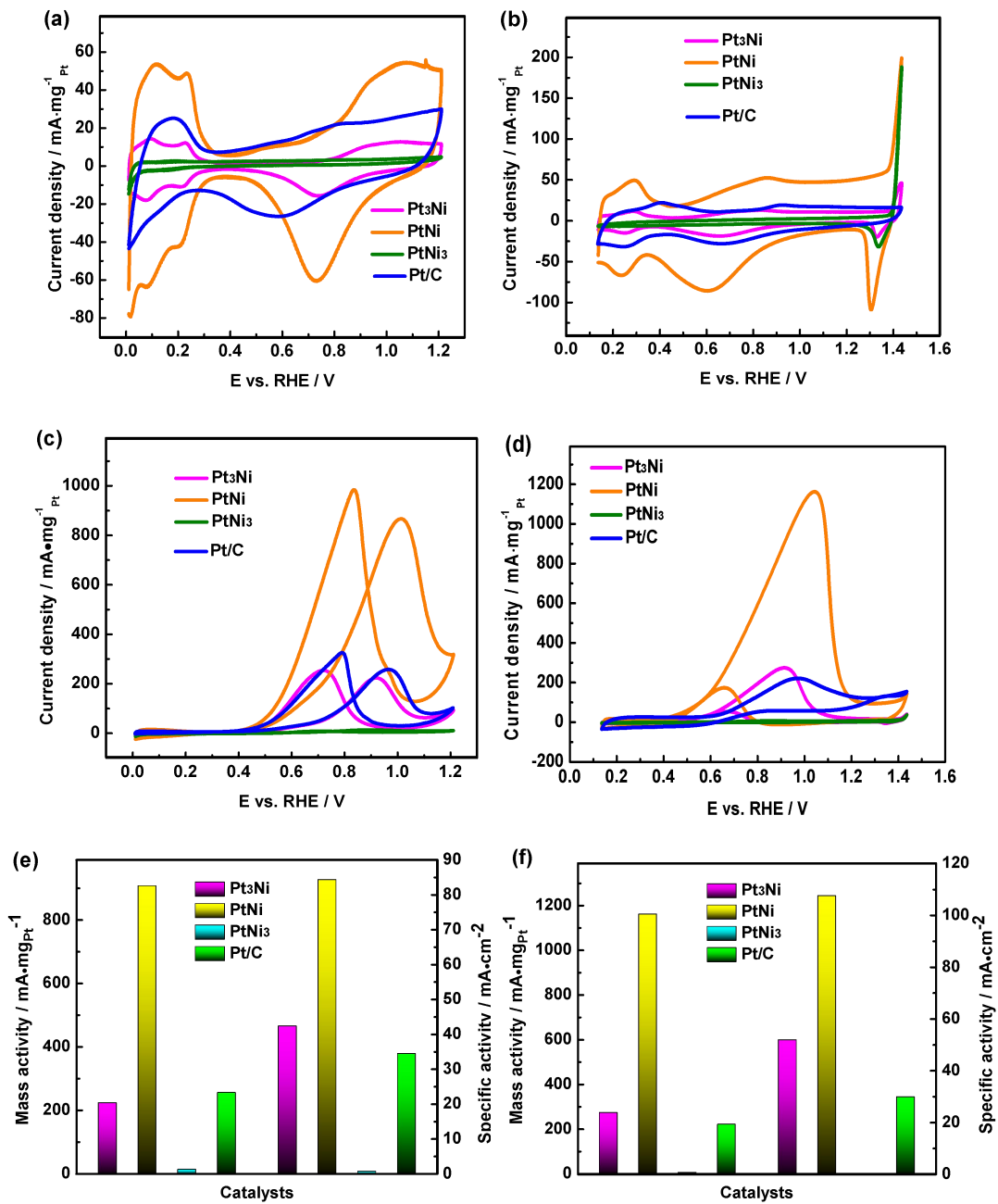


Figure S7. CVs of Pt-Ni NBs in (a) 0.5 M H₂SO₄, (b) 0.5 M H₂SO₄ + 1.0 M CH₃OH, (c) 1.0 M KOH, and (d) 1.0 M KOH + 1.0 M CH₃OH. Mass- and area-specific activities of Pt-Ni NBs at 0.929 and 0.835 V (vs. RHE) in (e) 0.5 M H₂SO₄ + 1.0 M CH₃OH and (f) 1.0 M KOH + 1.0 M CH₃OH, respectively.

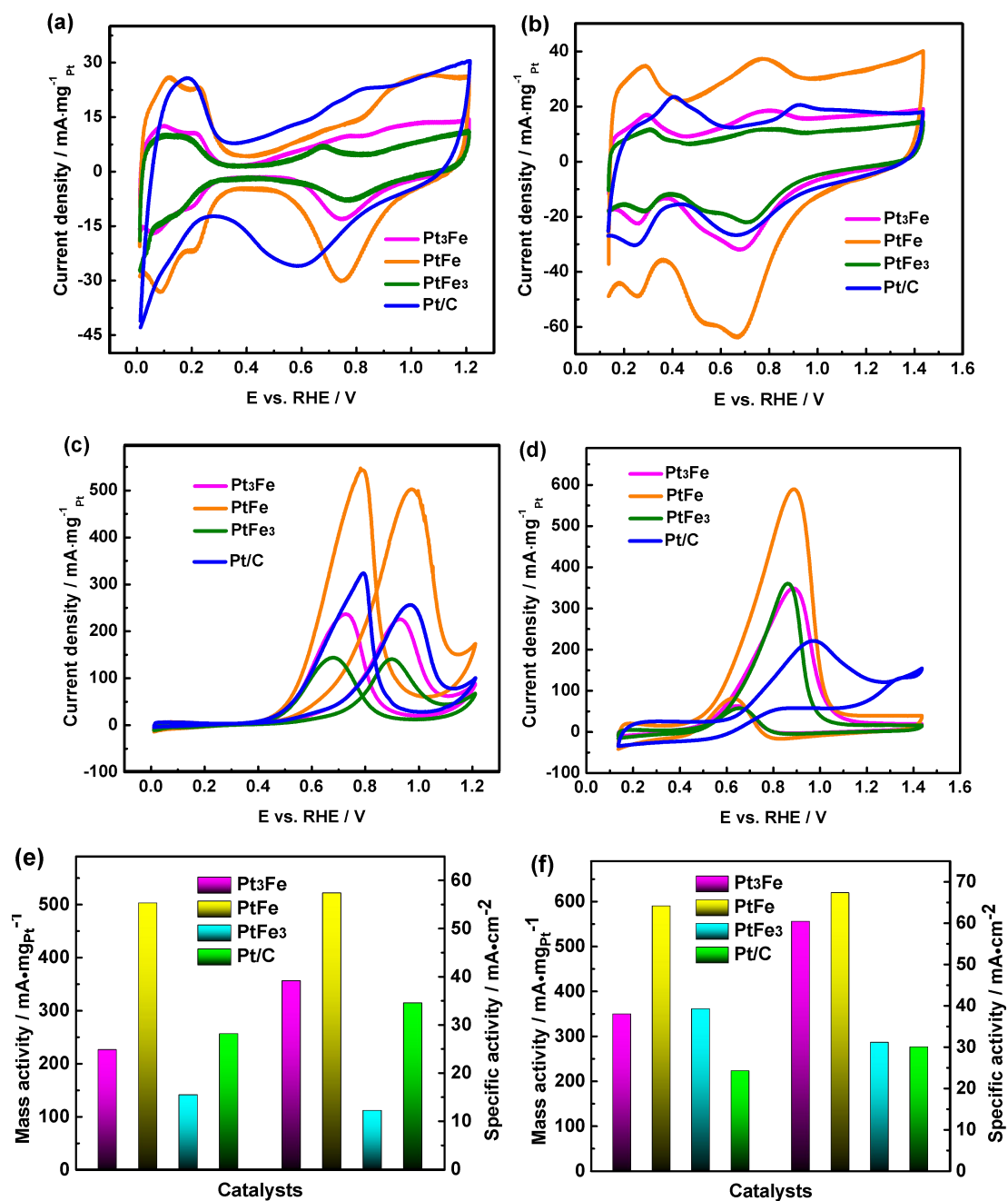


Figure S8. CVs of Pt-Fe NBs in (a) 0.5 M H₂SO₄, (b) 0.5 M H₂SO₄ + 1.0 M CH₃OH, (c) 1.0 M KOH, and (d) 1.0 M KOH + 1.0 M CH₃OH. Mass- and area-specific activities of Pt-Fe NBs at 0.929 and 0.835 V (vs. RHE) in (e) 0.5 M H₂SO₄ + 1.0 M CH₃OH and (f) 1.0 M KOH + 1.0 M CH₃OH, respectively.

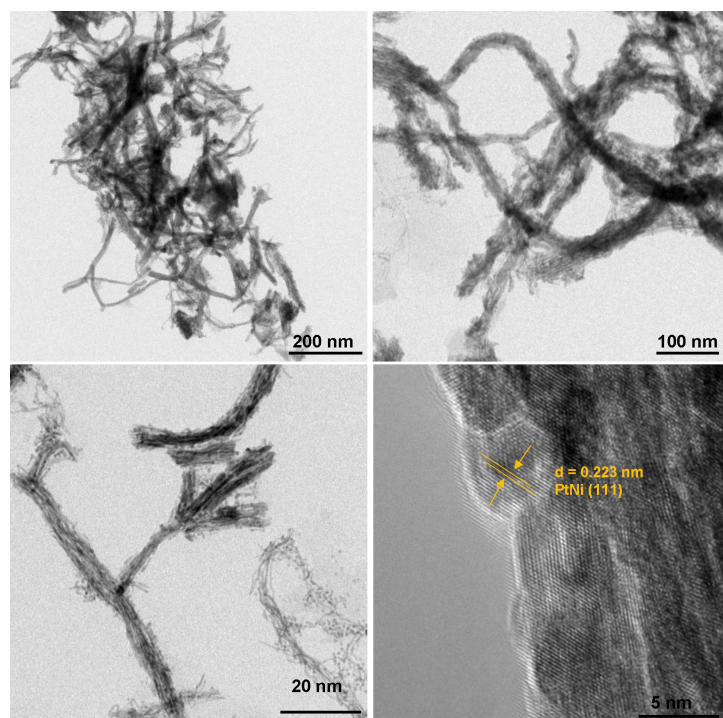


Figure S9. TEM and HRTEM images of PtNi NBs after 2000 cycles in 0.5 M H₂SO₄ + 1.0 M CH₃OH .