

Selective hydrogenolysis of furfural to 1,2-pentanediol over Pt-Fe/MT catalyst under mild conditions

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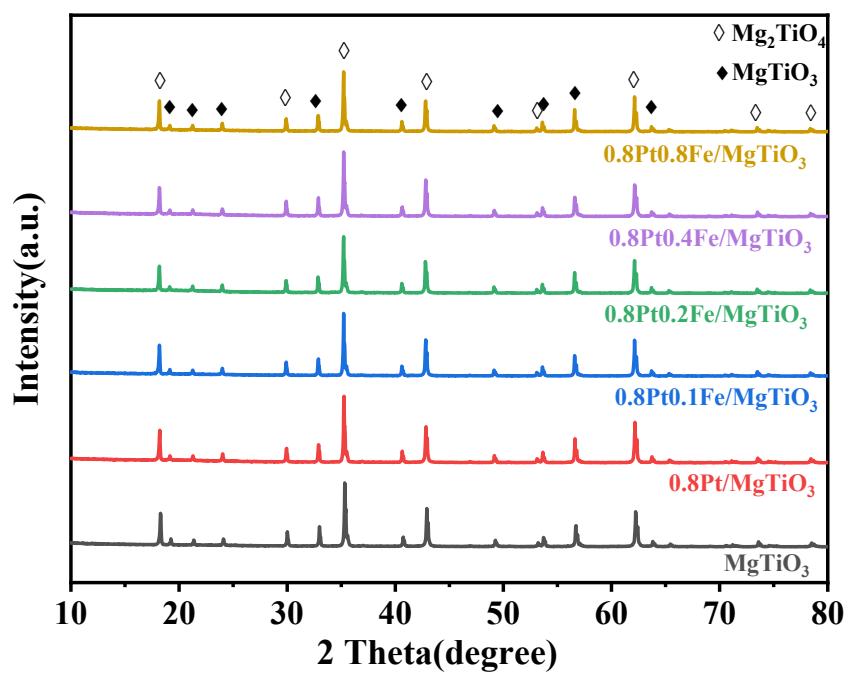


Figure S1 XRD patterns of the series of x Pt- y Fe/MT catalysts.

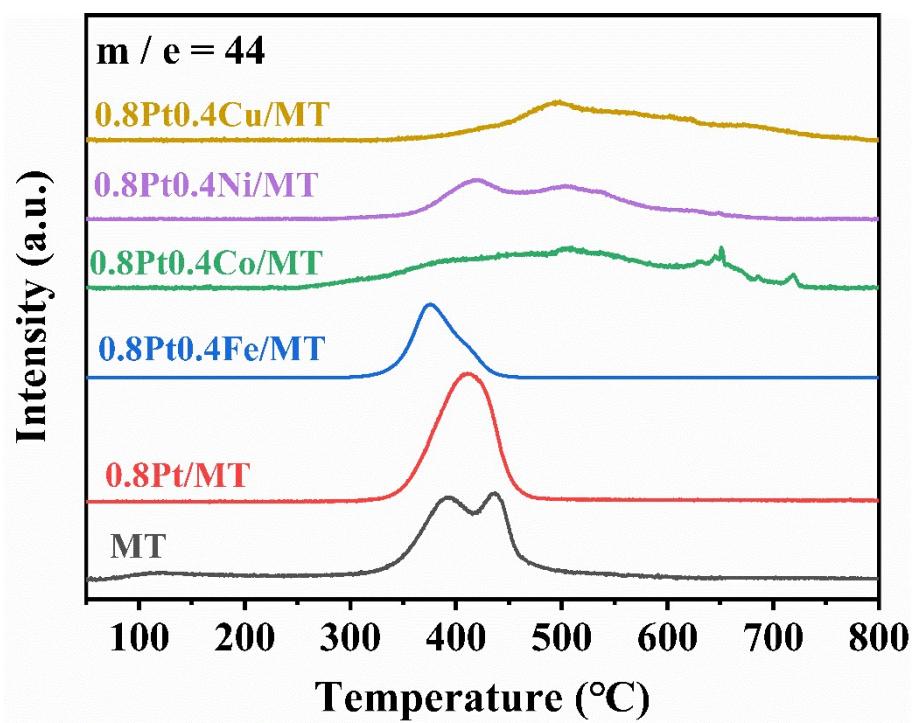


Figure S2 CO₂-TPD profiles for the MT, 0.8Pt/MT and 0.8Pt-0.4M/MT catalysts.

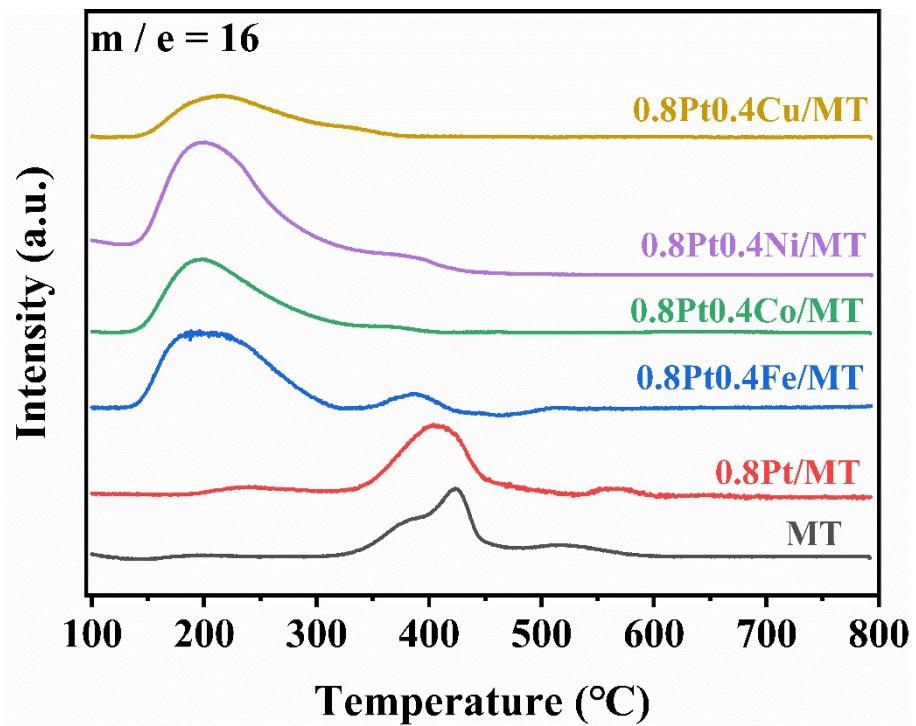


Figure S3 NH₃-TPD profiles for the MT, 0.8Pt/MT and 0.8Pt-0.4M/MT catalysts.

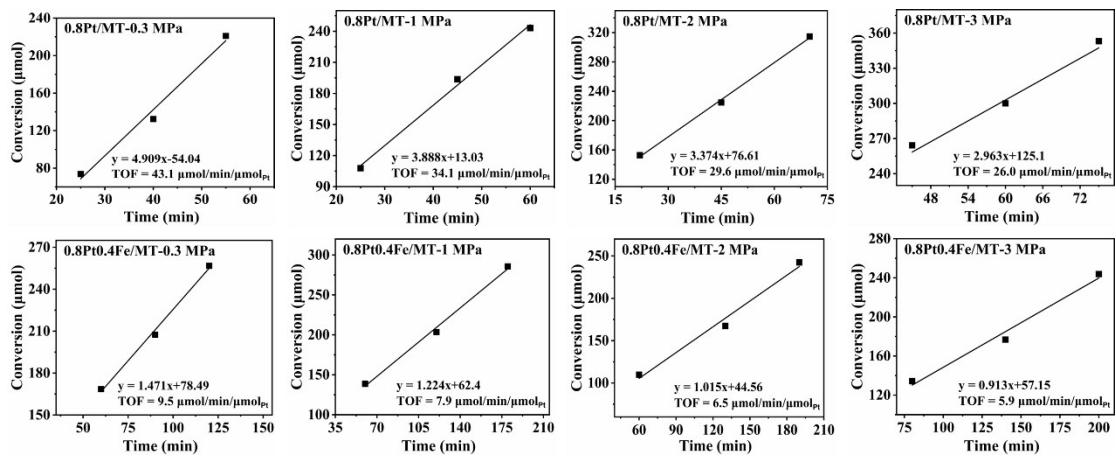


Figure S4 TOF values under different reaction pressure over 0.8Pt/MT and 0.8Pt0.4Fe/MT catalysts.

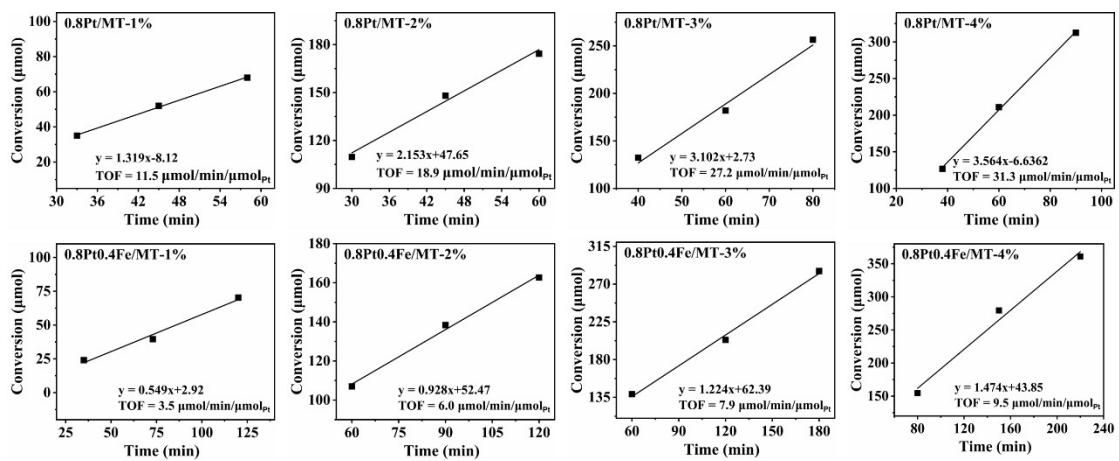


Figure S5 TOF values under different FFA concentration over 0.8Pt/MT and 0.8Pt0.4Fe/MT catalysts.

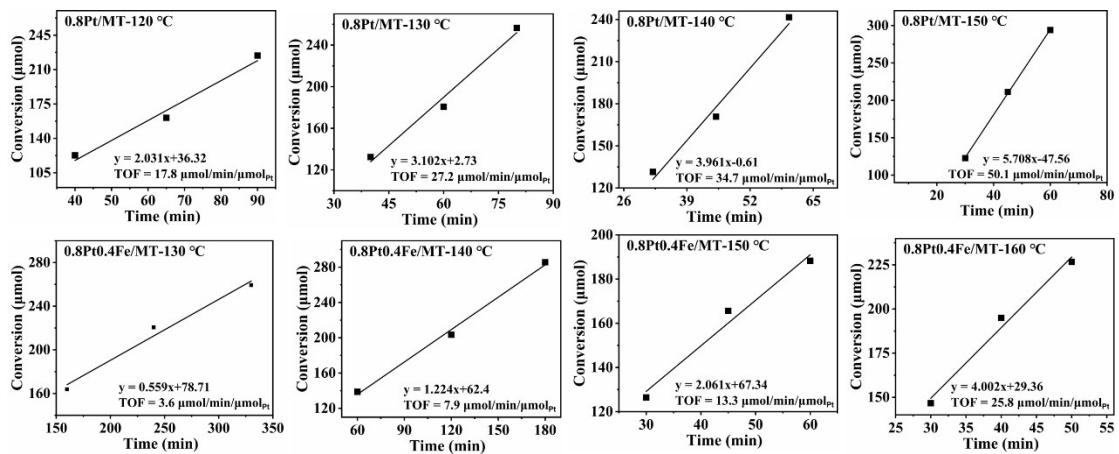


Figure S6 TOF values under different temperature over 0.8Pt/MT and 0.8Pt0.4Fe/MT catalysts.

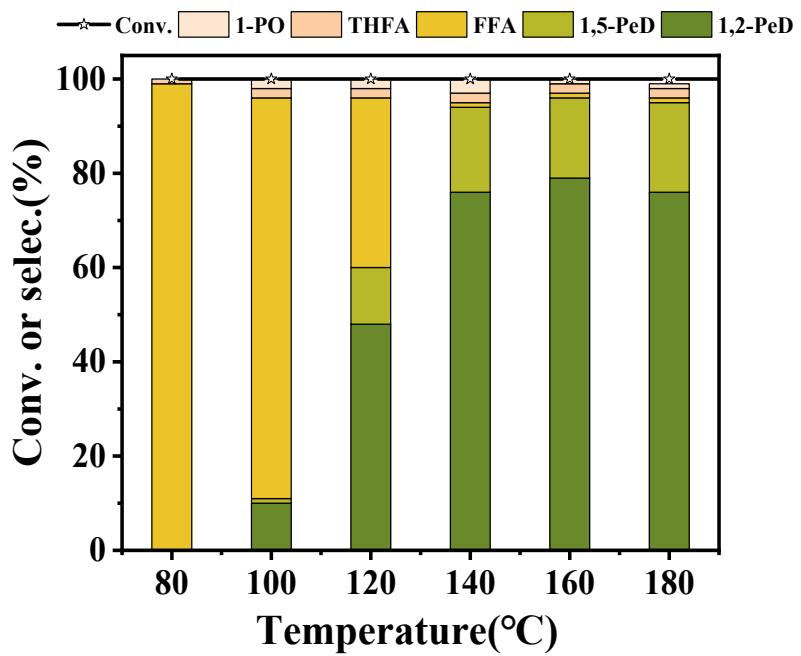


Figure S7 Catalytic performance under different temperature over 0.8Pt0.4Fe/MT catalyst.
Reaction conditions: 0.1 g FA in 5 g water, 100 mg catalyst, 1 MPa H₂, 10 h.

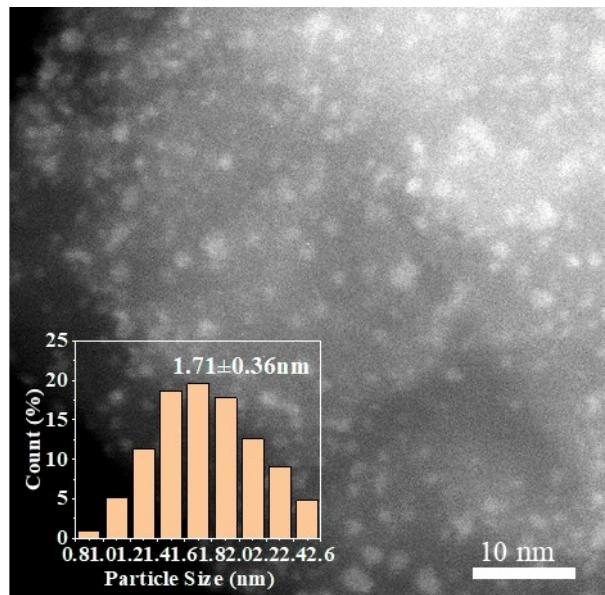


Figure S8 HAADF-STEM image of the 0.8Pt0.4Fe/MT-used catalyst.

Table S1 The amount of Mg²⁺ in the impregnation solution.

Catalyst	Concentration of Mg ²⁺
0.8Pt/MT	247.5 mg/L

Add 0.343 g of an aqueous solution of H₂PtCl₆·6H₂O (Pt: 1.166 wt %) into a beaker, followed by the addition of water to achieve a total mass of 7 g, then introduce 0.5 g of MT. After stirring the mixture of metal precursor and support for 24 hours, it was filtered, and the filtrate was subjected to ICP testing. The leaching content of Mg is 0.3 wt %.

Table S2 The alkalinity and acidity of 0.8Pt0.4M/MT catalysts.

Catalysts	CO ₂ uptake ^a (μmol _{CO2} /g _{cat})	NH ₃ uptake ^b (μmol _{NH3} /g _{cat})
0.8Pt0.4Co/MT	22.3	13.2
0.8Pt0.4Ni/MT	25.8	19.9
0.8Pt0.4Cu/MT	18.7	7.7

^a Determined by CO₂-TPD using a MS signal m / e = 44.

^b Determined by NH₃-TPD using a MS signal m / e = 16

Table S3 Catalytic performance of the 0.8Pt0.4Fe/MT catalyst for the THFA hydrogenolysis.

Entry	Conversion (%)	Selectivity (%)	
		1,2-PeD	1,5-PeD
1	0	0	0

Reaction conditions: 0.1 g THFA in 5 mL H₂O, 100 mg catalyst, 1 MPa H₂ and 140 °C for 10 h.

Table S4 Comparison with the literature on the production rate of 1,2-PeD by hydrogenolysis of FA or FFA over noble metal catalyst.

Sub.	Cat.	T (°C)	P (MPa)	Time (h)	Sel. /%			Production rate (mol _{1,2-PeD} / (h * mol _{noble metal}))	Ref
					THFA	1,2-PeD	1,5-PeD		
FF-A	4Ru/MnOx	150	1.5	6	30	42.1	-	3.6	1
FF-A	10Ru/Al ₂ O ₃	200	10	1	57	32	-	20.4	2
FA	1Ru-5Sn/ZnO	140	3.5	6	0	84.5	12.4	168	3
FF-A	5Ru/MgO	190	3	1	51	42	2.9	144	4
FA	20Ru/PVP	125	2	48	64	36	-	1.5	5
FA	3Pd/MMT-K	220	3.5	5	13	66	-	47	6
FA	1Rh/OMS-2	160	3	8	0	87	-	33	7
FA	1.9Pt/HT	150	3	4	14	73	8	19	8
FA	5Pt/CeO ₂	165	1.5	4	22.8	59.9	3.1	6	9
FF-A	4.5%Pt/CeO ₂	165	2	24	11.7	77.1	7.3	3	10
FF-A	1.67Pt/Mg(Al)O@Al ₂ O ₃ -IR	200	3	0.12 h ⁻¹	6	86	5	12.5	11
FA	0.8Pt0.4Fe/MT	140	0.1	10	2	81	15	53 ^a	
FA	0.2Pt0.1Fe/MT	140	0.1	10	6	67	20	128 ^a	This work
FA	0.1Pt0.05Fe/MT	140	0.1	10	2	43	17	178 ^a	

^a: These data were obtained after a reaction time of 2 h.

Table S5 The effect of solvents.^a

Entry	Solvent	Conversion. (%)	Selectivity (%)				
			1,2-PeD	1,5-PeD	FFA	THFA	1-PO
1	Methanol	100	0	0	100	0	0
2	Ethanol	100	0	0	100	0	0
3	N,N-Dimethylformamide	100	0	0	100	0	0
4	Water	100	76	18	1	2	3

^a Reaction conditions: 0.1 g of FA in 5 mL H₂O, 100 mg of 0.8Pt0.4Fe/MT, 1 MPa H₂, and 140 °C for 10 h.

Table S6 Catalytic performance of the 0.8Pt/MT and the 0.8Pt0.4Fe/MT catalyst for the furan and its derivatives hydrogenolysis.

Entry	Substrate	Catalyst	Selectivity (%)	
			Tetrahydrofuran	1-Butanol
1	Furan	0.8Pt/MT	47	53
2	Furan	0.8Pt0.4Fe/MT	15	85
3	2,5-Dihydrofuran	0.8Pt0.4Fe/MT	93	7
4	2,3-Dihydrofuran	0.8Pt0.4Fe/MT	90	10

Reaction conditions: 0.1 g of substrate in 5 g solvent (3 g H₂O and 2 g ethanol), 100 mg of catalyst, 1 MPa H₂, and 140 °C for 10 h.

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