

Supplementary information

Modification of Composite Catalytic Material $\text{Cu}_m\text{V}_n\text{O}_x@\text{CeO}_2$ Core-shell Nanorod by Tungsten for NH_3 -SCR

Xiaosheng Huang^{a,b}, Fang Dong^a, Guodong Zhang^a and Zhicheng Tang^{a,c*}

(*a. State Key Laboratory for Oxo Synthesis and Selective Oxidation, National Engineering Research Center for Fine Petrochemical Intermediates, Lanzhou Institute of Chemical Physics, Chinese Academy of Sciences, Lanzhou 730000, PR China*

b. University of Chinese Academy of Sciences, Beijing 100039, PR China)

c. Dalian National Laboratory for Clean Energy, Dalian Institute of Chemical Physics, Chinese Academy of Sciences, Dalian, 116023, China

*Corresponding author. Tel.: +86-931-4968083, Fax: +86-931-8277088, E-mail address:
tangzhicheng@licp.ac.cn (Z.Tang).

Fig. S1 TG/DTA curves of PVP (K90).

Fig. S2 TEM images of $\text{Cu}_m\text{V}_n\text{O}_x$ samples calcined at different temperatures.

Fig. S3 FTIR spectrum (a) and XRD patterns (b) of Ce-MOF, $\text{Cu}_m\text{V}_n\text{O}_x$ -NF/Ce and $\text{Cu}_m\text{V}_n\text{O}_x$ -NF@Ce-MOF.

Fig. S4 N_2 sorption curves and pore size distribution of all as-prepared catalysts.

Fig. S5 XPS spectra of $\text{Cu}_m\text{V}_n\text{O}_y$ @ CeO_2 - WO_x calcined at different temperatures.

Table S1 Quantity results of surface acid sites.

Table S2 Peak integration area of pyridine FTIR and NH_3 -TPD.

Table S3 XPS results of surface composition

Table S4. Reaction rate constant k of $\text{Cu}_3(\text{VO}_4)_2$, CeO_2 , $\text{Cu}_3(\text{VO}_4)_2$ @ CeO_2 and $\text{Cu}_3(\text{VO}_4)_2$ @ CeO_2 - WO_x .

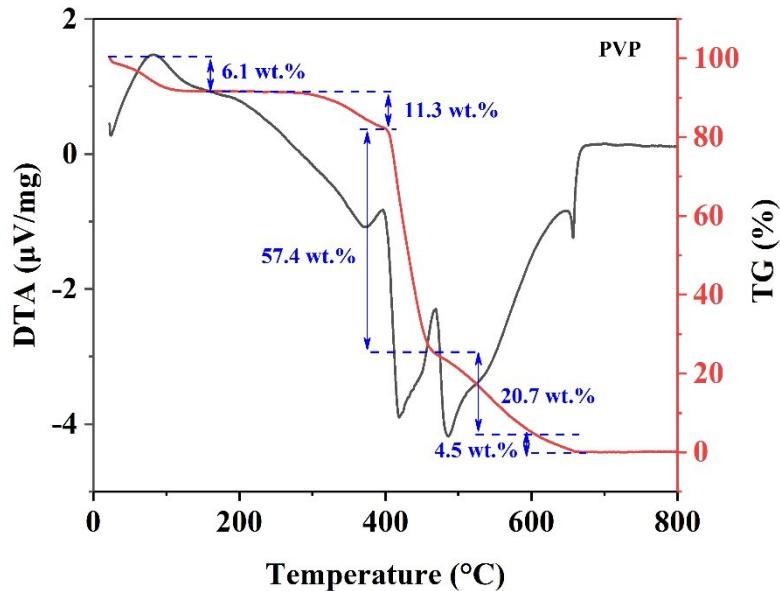


Fig. S1 TG/DTA curves of PVP (K90).

As shown in above TG/DTA curves, the decomposition of PVP (k90) mainly occurred in $290 \sim 410$ °C with several exothermic peaks, which implied that the side and main chains were decomposed step by step. All residuum was completely consumed above 663 °C. Above information could provide references for other samples' TG/DTA analyses in text part.

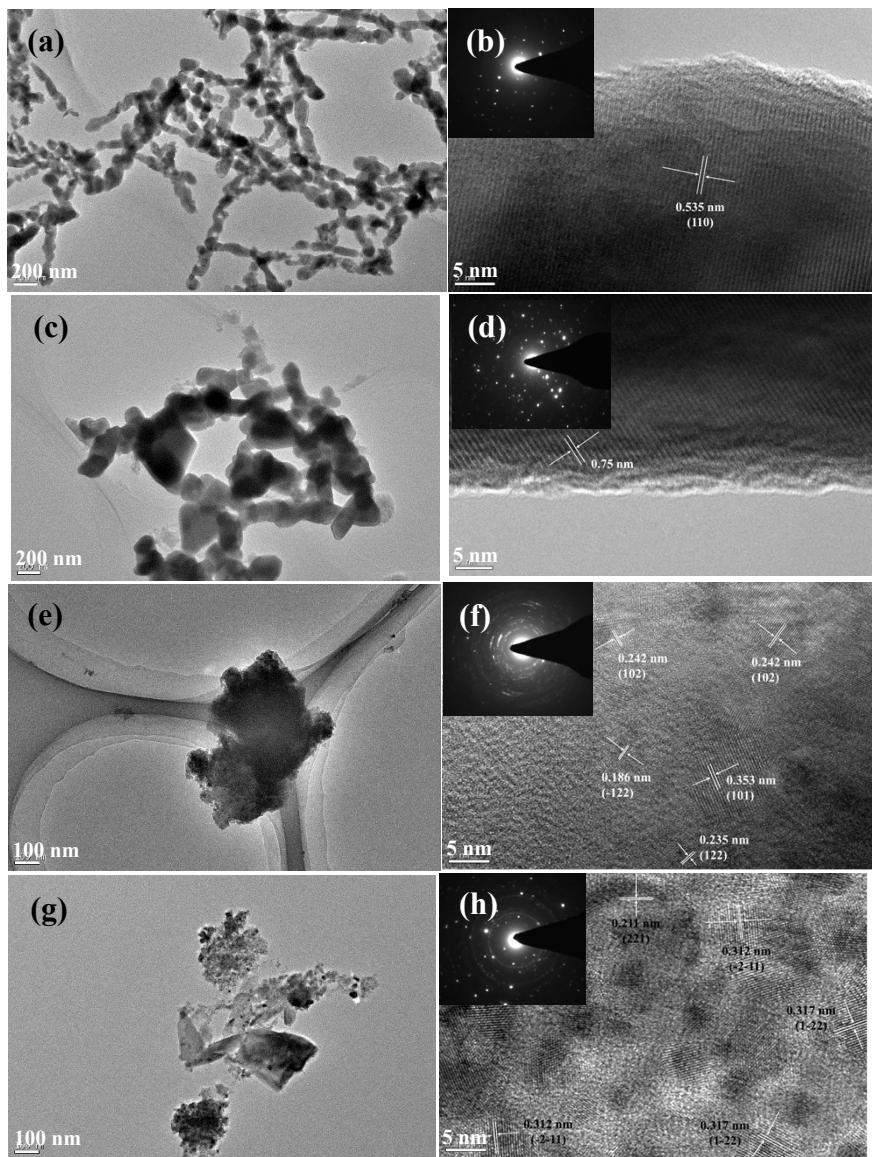


Fig. S2 TEM images of $\text{Cu}_m\text{V}_n\text{O}_x$ samples calcined at 400 °C (a, b), 500 °C (c, d), 600 °C (e, f) and 700 °C (g, h).

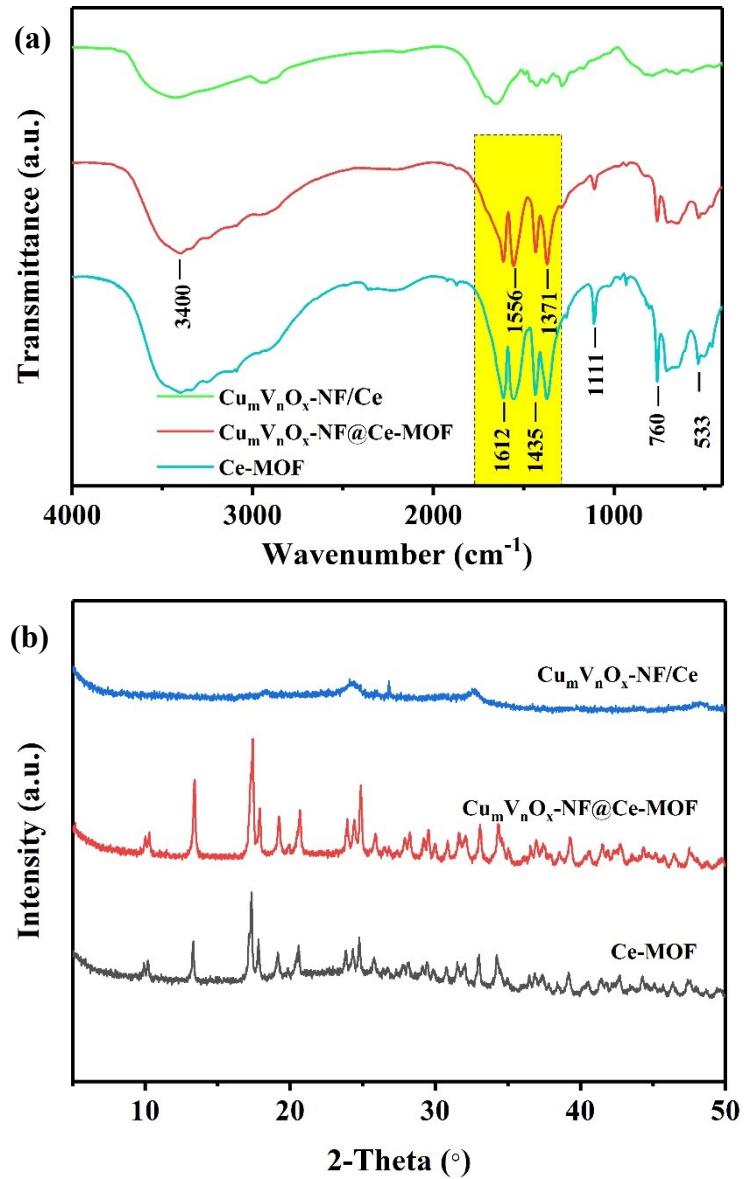


Fig. S3 FTIR spectrum (a) and XRD patterns (b) of Ce-MOF, $\text{Cu}_m\text{V}_n\text{O}_x\text{-NF@Ce-MOF}$ and $\text{Cu}_m\text{V}_n\text{O}_x\text{-NF/Ce}$.

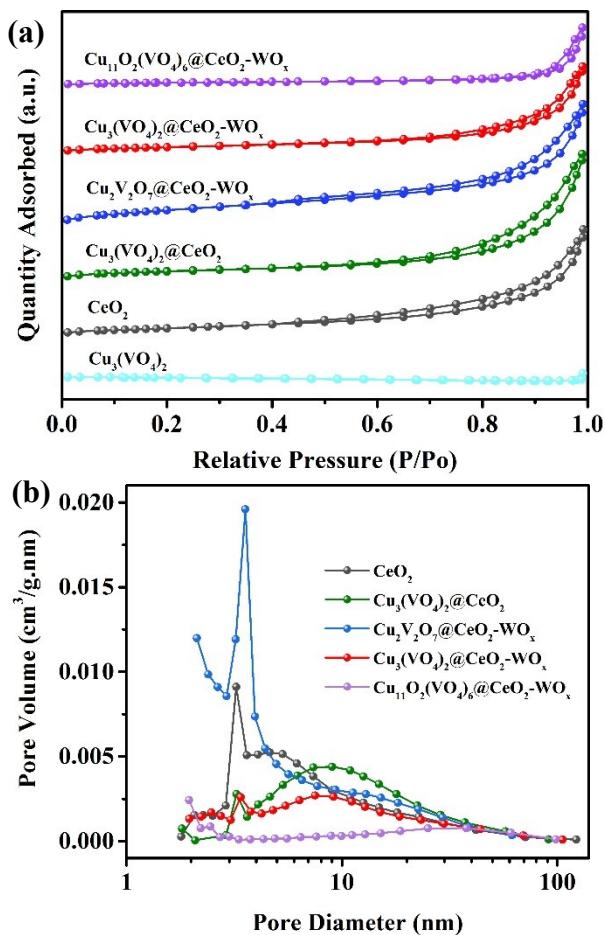


Fig. S4 N_2 sorption curves (a) and pore size distribution (b) of all as-prepared catalysts.

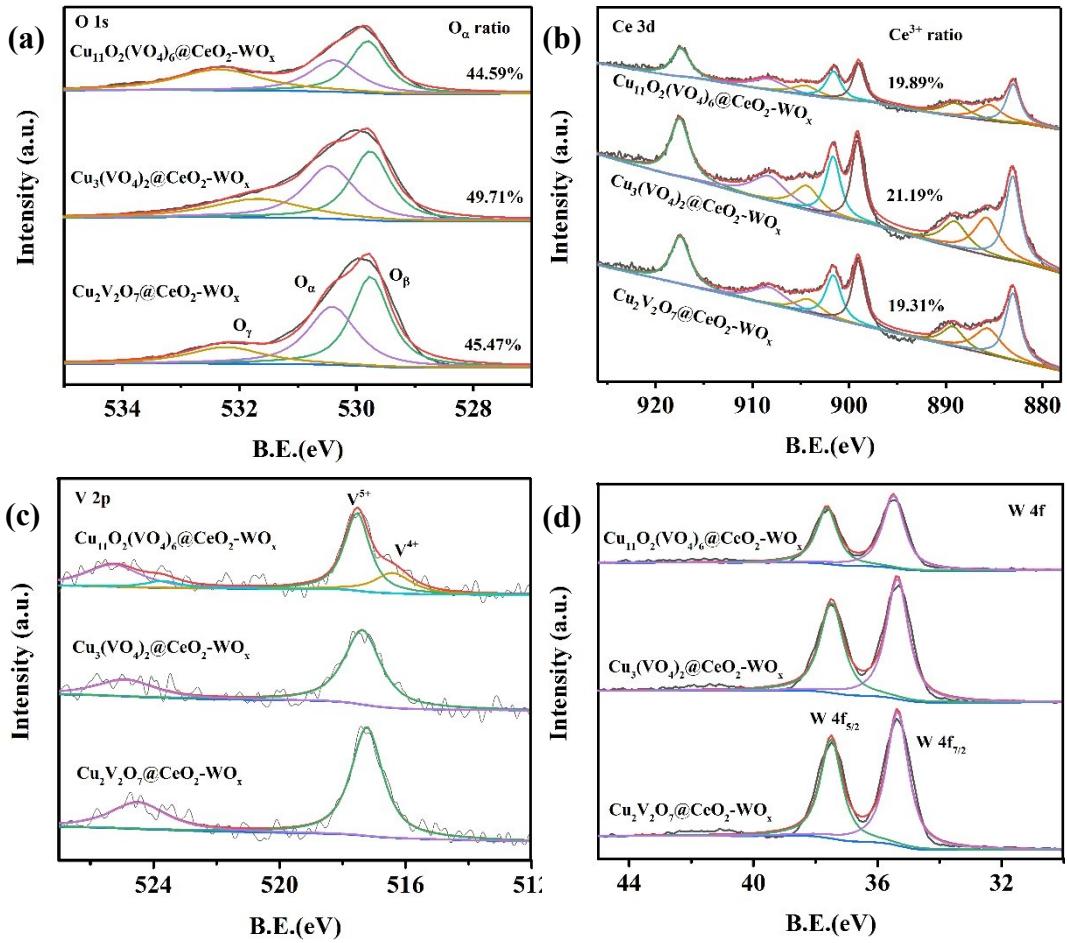


Fig. S5 XPS spectra of $\text{Cu}_m\text{V}_n\text{O}_y@\text{CeO}_2\text{-WO}_x$ calcined at different temperatures.

Table S1 Quantity results of surface acid sites.

samples	NH ₃ uptake (mmol/g _{cat})		
	Weak acid (<240 °C)	Medium strong acid (>240 °C)	total
Cu ₃ (VO ₄) ₂	-	1.15	1.15
CeO ₂	0.48	-	0.48
Cu ₃ (VO ₄) ₂ @CeO ₂	-	1.04	1.04
Cu ₂ V ₂ O ₇ @CeO ₂ -WO _x	0.67	0.88	1.55
Cu ₃ (VO ₄) ₂ @CeO ₂ -WO _x	1.30	1.29	2.59
Cu ₁₁ O ₂ (VO ₄) ₆ @CeO ₂ -WO _x	0.65	0.24	0.89

 Table S2 Peak integration area of pyridine FTIR and NH₃-TPD.

Cu ₃ (VO ₄) ₂ @CeO ₂ -WO _x	Peak integration area (a.u.)		Area ratio
	Lewis acid	Brønsted acid	
Pyridine FTIR			0.96 ^a
	12.3	11.8	
NH ₃ -TPD	Weak acid	Medium strong acid	0.99 ^b
	145.2	144.8	

^a the area ratio of Brønsted acid to Lewis acid

^b the area ratio of Medium strong acid to Weak acid

Table S3 Catalysts surface composition (%)

	Cu	V	O	Ce	W	O _a ratio	Ce ³⁺ ratio
Cu ₃ (VO ₄) ₂	12.42	28.99	58.59	-	-	49.12	-
Cu ₃ (VO ₄) ₂ @ CeO ₂	3.11	20.12	61.97	14.80	-	38.73	19.07
Cu ₂ V ₂ O ₇ @ CeO ₂ -WO _x	2.76	19.95	60.93	10.61	5.75	45.47	19.31
Cu ₃ (VO ₄) ₂ @ CeO ₂ -WO _x	2.59	19.37	60.04	12.86	5.14	49.71	21.19
Cu ₁₁ O ₂ (VO ₄) ₆ @ CeO ₂ -WO _x	2.61	20.66	63.15	8.76	4.81	44.59	19.89

Table S4. Reaction rate constant k of Cu₃(VO₄)₂, CeO₂, Cu₃(VO₄)₂@CeO₂ and

Cu₃(VO₄)₂@CeO₂-WO_x.

Temperatur e (°C)	Reaction rate constant k (L·g ⁻¹ ·min ⁻¹)			
	Cu ₃ (VO ₄) ₂	CeO ₂	Cu ₃ (VO ₄) ₂ @CeO ₂	Cu ₃ (VO ₄) ₂ @CeO ₂ -WO _x
100	0.02	0.02	0.03	0.09
140	0.04	0.03	0.10	0.39
180	0.07	0.06	0.22	1.02
220	0.10	0.14	0.42	1.61