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Templating synthesis of metal oxides by an incipient wetness

impregnation route and their activities for CO oxidation

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Catalyst	Amount of H2 desorbed (µmol/g)	Dispersion (%)
Co/SBA-16_80	310.2	27
Co/SBA-16_100	259.4	20
Co/SBA-16_120	226.9	15

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Table 2. Comparison in the CO oxidation activity of the herein materials with that in literature

Catalyst ^a	Reaction conditions	$T_{ig}(^{\circ}C)$	T ₁₀₀ (°C)	Ref.
Co ₃ O ₄ -16_80	50 mg catalyst; 50 mL min ⁻¹ ;	90	140	This work
$Ce_{0.7}Cu_{0.3}O_2-15_{100}$	0.5% CO-7.5% O ₂ /Ar	100	210	
Co ₃ O ₄ -40	200 mg catalyst; 60 mL min ⁻¹ ;	-30	30	1
Co ₃ O ₄ -100	1% CO in air	-20	80	
Co ₃ O ₄ -130		5	150	
Meso-Co-250	100 mg catalyst; 20 mL min ⁻¹ ;	-67	-35	2
Meso-Co-350	1% CO-1% O ₂ /N ₂	-75	-60	
Meso-Co-450		-45	0	
m-Co ₃ O ₄	50 mg catalyst; 37 mL min ⁻¹ ;	-80	-55	3
	1% CO in air			
Co ₃ O ₄ -450	100 mg catalyst; 33.3 mL min ⁻¹ ;	< -100	-37	4
Co ₃ O ₄ -500	1% CO-21% O ₂ /He	< -100	-49	
Co ₃ O ₄ -600		< -100	-29	
(CuO)5%/CeO2,	1500 mg catalyst; 200 mL min ⁻¹ ;	85	131	5
(CuO)10%/CeO2,	1% CO-10% O ₂ /N ₂	65	111	
(CuO) 15%/CeO2,		85	151	
CuCe _{0.33}	50 mg catalyst; 25 mL min ⁻¹ ;	60	71	6
CuCe _{0.46}	1% CO in air	62	81	
CuCe _{0.22}		80	84	

^a for the meaning of the catalyst's name, see the respective literature

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Figure S1. The small-angle XRD patterns (A) and the wide-angle XRD patterns (B) of NiO, CeO₂ and Ce_{0.7}Cu_{0.3}O₂ replicated from SBA-15



Figure S2. N₂ adsorption–desorption isotherms of (A) the silica templates SBA-16_T and (B) the Co₃O₄-silica composites, and the corresponding pore size distribution for each.



Figure S3. TEM images of the SBA-15_100 and SBA-16_T templates



Figure S4. TEM images of (A) NiO-15_100, (B) CeO₂-15_100 and (C) Ce_{0.7}Cu_{0.3}O₂-15_100.



Figure S5. SEM images of the Co₃O₄/SBA-16_100 and Co₃O₄/SBA-15_100



Figure S6. SEM images of the replicated Co₃O₄-16_100, Co₃O₄-16_120 and Co₃O₄-15_100



Figure S7. UV-Raman spectra of the replicated Co₃O₄-16_T and Co₃O₄-15_100



Figure S8. CO conversion obtained from NiO-15_100, CeO₂-15_100 and Ce_{0.7}Cu_{0.3}O₂-15_100. Reaction conditions: 50 mg Catalyst, 0.5% CO-7.5% O₂/Ar, flow rate of 50 mL min⁻¹.



Figure S9. Comparison in the CO oxidation activity of Co_3O_4 -16_80 pre-treated with and without 8% O₂/Ar at 120 °C for 30 min.