



## Supporting Information

for

### **Synthesis of MnO<sub>2</sub>–CuO–Fe<sub>2</sub>O<sub>3</sub>/CNTs catalysts: low-temperature SCR activity and formation mechanism**

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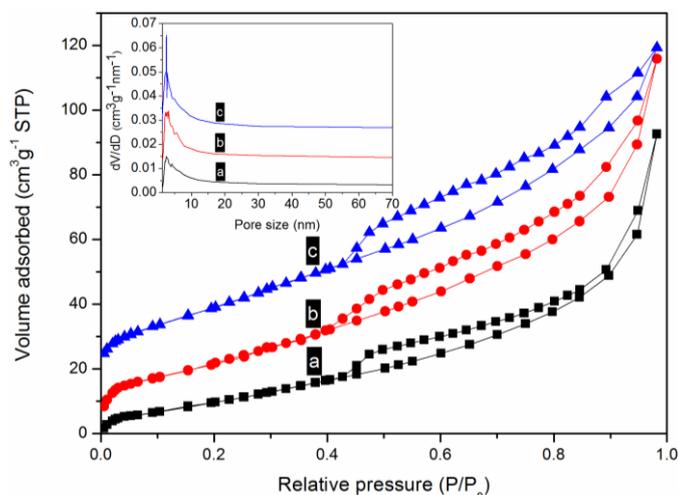
## Additional experimental data

**Table S1:** BET surface area and pore volume for the pristine CNTs, acid-treated CNTs and as-prepared catalysts

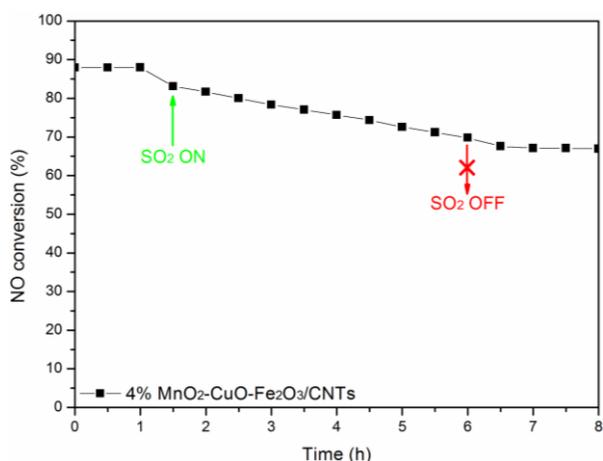
sample	$S_{\text{BET}}$ ( $\text{m}^2 \cdot \text{g}^{-1}$ )	pore volume ( $\text{cm}^3 \cdot \text{g}^{-1}$ )
pristine CNTs	63.11	0.1457
acid-treated CNTs	71.11	0.1585
1% Mn–CuO–Fe <sub>2</sub> O <sub>3</sub> /CNTs	61.29	0.1319
2% Mn–CuO–Fe <sub>2</sub> O <sub>3</sub> /CNTs	86.40	0.1660
4% Mn–CuO–Fe <sub>2</sub> O <sub>3</sub> /CNTs	95.66	0.1841
6% Mn–CuO–Fe <sub>2</sub> O <sub>3</sub> /CNTs	130.54	0.2447
Mn–Cu–FeO <sub>x</sub> /CNTs-IWIM	111.05	0.1673

**Table S2:** Relative oxygen content of the as-prepared catalysts.

sample	O(%)	
	O <sub>L</sub>	O <sub>S</sub>
4% MnO <sub>2</sub> –CuO–Fe <sub>2</sub> O <sub>3</sub> /CNTs	33.3	66.7
Mn–Cu–FeO <sub>x</sub> /CNTs-IWIM	63.2	36.8



**Figure S1:** N<sub>2</sub> adsorption–desorption isotherm and pore size distribution (inset) of (a) acid-treated CNTs, (b) 4% MnO<sub>2</sub>–CuO–Fe<sub>2</sub>O<sub>3</sub>/CNTs, and (c) Mn–Cu–FeO<sub>x</sub>/CNTs-IWIM.



**Figure S2:** SO<sub>2</sub> tolerance of 4% MnO<sub>2</sub>-CuO-Fe<sub>2</sub>O<sub>3</sub>/CNTs catalyst. Reaction conditions: [NO] = [NH<sub>3</sub>] = 400 ppm, [SO<sub>2</sub>] = 50 ppm (when used), [O<sub>2</sub>] = 5%, N<sub>2</sub> as balance gas, WHSV=280 L·g<sub>cat</sub><sup>-1</sup>·h<sup>-1</sup>, 0.15 g catalyst.

### BET surface area data

The results of BET surface area measurements (Table S1) show that the surface area of the pristine CNTs was 63.11 m<sup>2</sup>·g<sup>-1</sup>, whereas it became larger after being loaded with catalyst, indicating the even distribution of metal-oxide catalysts on the CNTs. It should be noted that although the 4% MnO<sub>2</sub>-CuO-Fe<sub>2</sub>O<sub>3</sub>/CNTs catalyst showed the best catalytic activity among all samples, it possessed a smaller surface area (95.66 m<sup>2</sup>·g<sup>-1</sup>) than Mn-Cu-FeO<sub>x</sub>/CNTs-IWIM (111.05 m<sup>2</sup>·g<sup>-1</sup>), revealing that the surface area is not the only important factor in SCR activity [1].

## **N<sub>2</sub> adsorption-desorption curves**

Figure S1 shows the N<sub>2</sub> adsorption–desorption isotherm and the pore size distribution of the acid-treated CNTs and the as-synthesized catalysts. All samples present a typical type-IV isotherm along with a type-H4 loop, verifying the mesoporous structure [2,3]. The pore sizes of the samples were between 2.5 and 5.0 nm.

## **SO<sub>2</sub> tolerance**

SO<sub>2</sub>, a common component of flue gas, can inhibit the catalytic activity. Therefore, the catalysts need to be resistant against SO<sub>2</sub>. In Figure S2, it is shown that the denitration efficiency of 4% MnO<sub>2</sub>–CuO–Fe<sub>2</sub>O<sub>3</sub>/CNTs catalyst is ca. 87.9% in the absence of SO<sub>2</sub>. When 50 ppm SO<sub>2</sub> is added to the gas flow for 5 h, there is a decline in catalytic efficiency to ca. 67.6%. Afterwards, the catalytic activity remains stable at around 67.1%. Hence, the 4% MnO<sub>2</sub>–CuO–Fe<sub>2</sub>O<sub>3</sub>/CNTs exhibits SO<sub>2</sub> tolerance, which is favorable for its practical application.

## **References**

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