## CO preferential oxidation over ordered mesoporous Co<sub>3</sub>O<sub>4</sub>-based composite catalysts

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Preferential oxidation of carbon monoxide (CO-PROX) has got a great attention during decades, especially in proton exchange membrane fuel cell (PEMFC) field due to its simplicity and cost-effectiveness for the removal of CO in the  $H_2$  fuel. This is because CO gas in the  $H_2$  fuel contaminates Pt electrodes in PEMFC system, thus the performance of PEMFC diminished. [1,2]

Among various kinds of materials, noble metals (Pt, Rh, Ru, and Au) were recognized as the promising candidates for effective CO-PROX reaction, however noble metal catalysts were not attractive due to their high costs.

Transition metal oxides (Co<sub>3</sub>O<sub>4</sub>, MnO<sub>2</sub>, etc.) have been investigated for escaping from noble metal catalysis system. Co<sub>3</sub>O<sub>4</sub> is a wellknown CO-PROX catalyst which exhibits high catalytic performance (CO oxidation by O<sub>2</sub>) at low temperature. In the preliminary research, ordered mesoporous Co<sub>3</sub>O<sub>4</sub> showed high catalytic performances with T<sub>50</sub> of 68 °C due to its high surface area (that is, large active sites on the catalyst). [3]

In this work, we enhanced the CO-PROX performance of ordered mesoporous  $Co_3O_4$  using 2 strategies: Co metal incorporation with ordered mesoporous  $Co_3O_4$  framework and other transition metal doping. There were the studies of Co metal promotion in noble metal catalysts for improving CO-PROX activity. [4–6] Also, other transition metal additives (such as Cu, Fe, etc.) in Co<sub>3</sub>O<sub>4</sub> framework enhanced on the catalytic activity of ordered mesoporous Co<sub>3</sub>O<sub>4</sub> due to the change of surface electronic structures. [7]

Thus, we prepared ordered mesoporous CoO, Co<sub>3</sub>O<sub>4</sub>, Co/Co<sub>3</sub>O<sub>4</sub>, and transition metal (Cu, Fe, etc.) and Co<sub>3</sub>O<sub>4</sub> nanocomposites for CO-PROX by nano-casting method using KIT-6 silica hard template. The physicochemical properties of prepared catalysts were evaluated with X-ray diffraction (XRD), N<sub>2</sub> adsorption-desorption analysis, electron microscopy (SEM and TEM) with energy-dispersive X-ray spectroscopy (EDX), and temperature programmed experiments (TPR and TPD). The CO-PROX activity was improved when Co metal was incorporated with Co<sub>3</sub>O<sub>4</sub> framework.



Fig.1 XRD patterns of prepared materials.



Fig.2 SEM with HR-SEM (in-set images) of prepared materials.

## REFERENCES

[1] S.Y. Hwang, E. Yurchekfrodl, C. Zhang and Z. Peng, ChemCatChem, 8 (2016) 97.

[2] M. Jin, J.-N. Park, J.K. Shon, J.H. Kim, Z. Li, Y.-K. Park and J.M. Kim, Catal. Today, 185 (2012) 183.

[3] S. Sun, Q. Gao, H. Wang, J. Zhu and H. Guo, Appl. Catal. B: Environmental, 97 (2010) 284.

[4] A. Tömcrona, M. Skoglundh, P. Thormählen, E. Fridell and E. Jobson, Appl. Catal. B: Environmental, 14 (1997) 131.

[5] W.S. Epling, P.K. Cheekatamarla and A.M. Lane, Chem. Eng. J., 93 (2003) 61.

[6] J. Choi, C.B. Shin and D.J. Suh, Catal. Commun., 9 (2008) 880.

[7] S. Varghese, M.G. Cutrufello, E. Rombi, C. Cnnas, R. Monaci and I. Ferino, Appl. Catal. A: General, 443–444 (2012) 161.