

Synthesis of Fe₃O₄-CuO@meso-SiO₂ Nanostructure as a Magnetically Recyclable and Efficient Catalyst for Styrene Epoxidation

Xiaowei Zhang,^a Ge Wang,^{*a} Mu Yang,^a Yi Luan,^a Wenjun Dong,^{*b} Rui Dang,^{a,b}
Hongyi Gao^a and Jie Yu^a

^a School of Materials Science and Engineering, University of Science and Technology
Beijing, Beijing 100083, China. Fax: +86-10-62327878; Tel: +86-10-62333765; E-mail:
gewang@mater.ustb.edu.cn

^b Center for Nanoscience and Nanotechnology, Department of Physics, Zhejiang Sci-tech
University, Hangzhou 310018, China. Fax: +86-571-86843587; Tel: +86-571-86843587;
E-mail: wenjundong@zstu.edu.cn

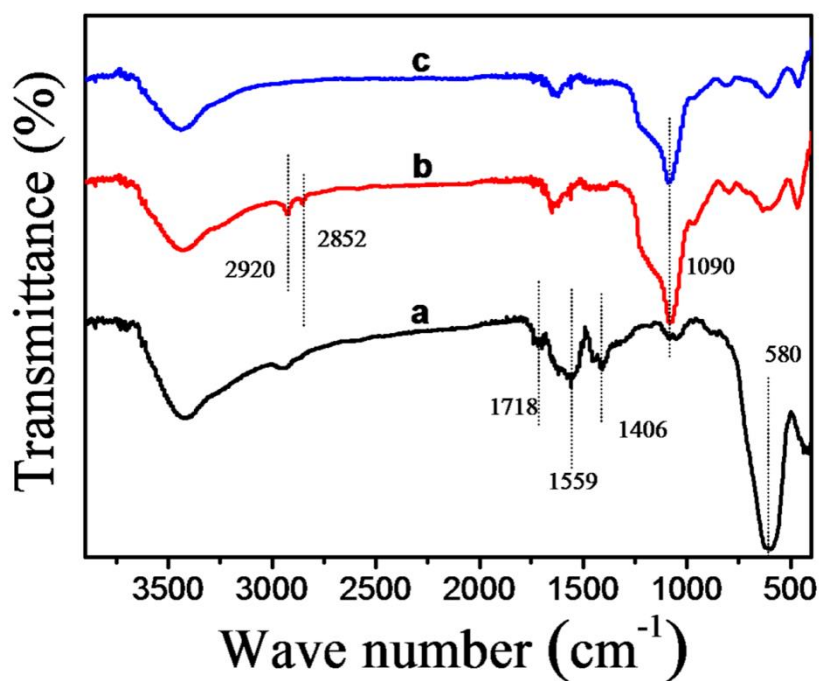


Fig. S1 FT-IR spectra of (a) $\text{Fe}_3\text{O}_4(\text{PAA})$, (b) $\text{Fe}_3\text{O}_4\text{-CuO@meso-SiO}_2(\text{CTAB})$ without the treatment of acetone and (c) $\text{Fe}_3\text{O}_4\text{-CuO@meso-SiO}_2$ after the treatment of acetone.

The existence of PAA (curve a) was confirmed by the C-O bond at 1406 cm^{-1} , C=C bond at 1559 cm^{-1} and C=O bond at 1718 cm^{-1} from carboxyl groups.¹ For $\text{Fe}_3\text{O}_4\text{-CuO@meso-SiO}_2$ with CTAB (curve b), the bonds at around 2852 cm^{-1} and 2920 cm^{-1} correspond to the vibrations of $-\text{CH}_2$ from CTAB. The disappearance of the two peaks at 2852 cm^{-1} and 2920 cm^{-1} (curve c) after the extraction treatment with acetone, indicate the complete removal of the CTAB template.²

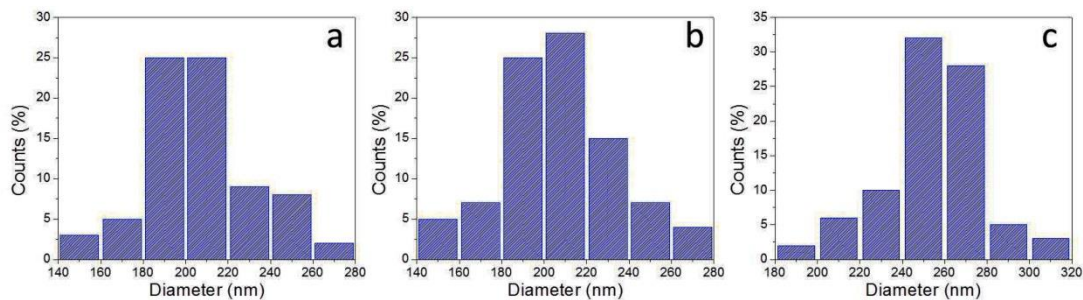


Fig. S2 The particle size distributions of (a) Fe_3O_4 , (b) $\text{Fe}_3\text{O}_4\text{-CuO}$ and (c) $\text{Fe}_3\text{O}_4\text{-CuO@meso-SiO}_2$.

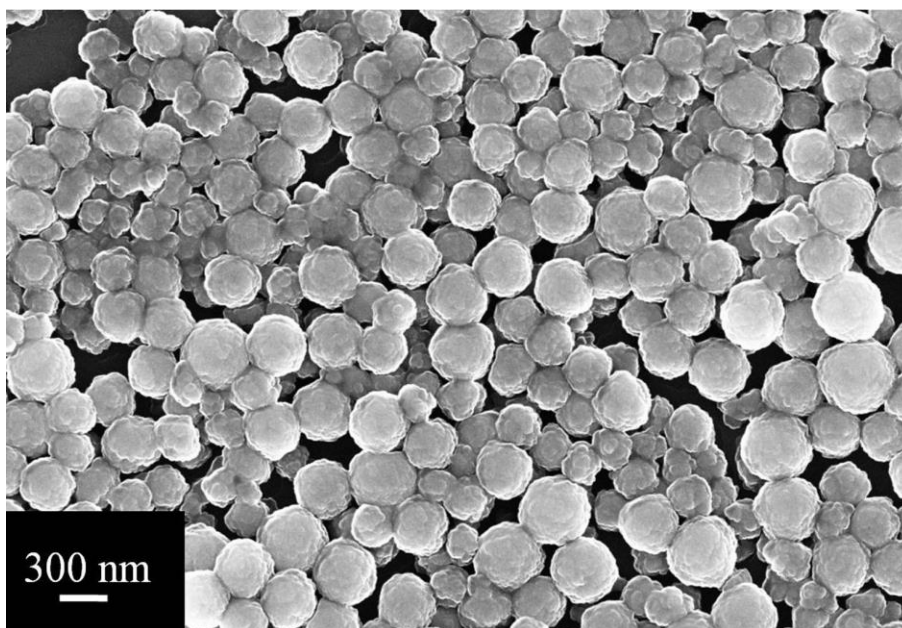


Fig. S3 SEM photograph of the as-synthesized CuO nanoparticles.

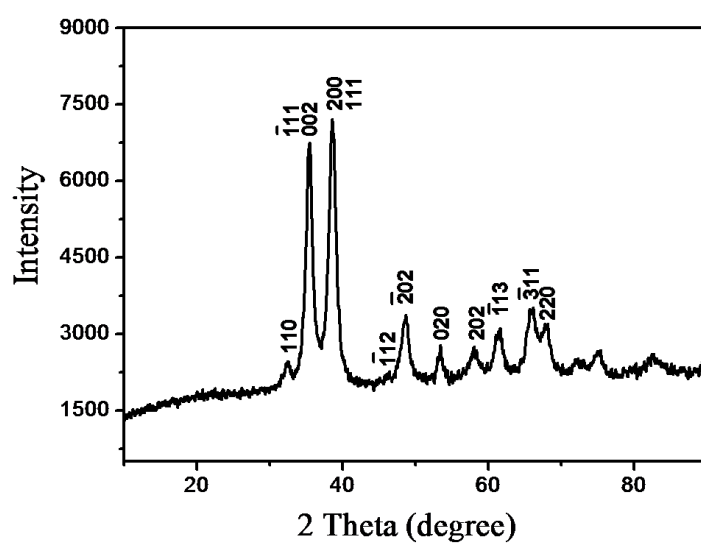


Fig. S4 X-ray diffraction patterns of the as-synthesized CuO nanoparticles.

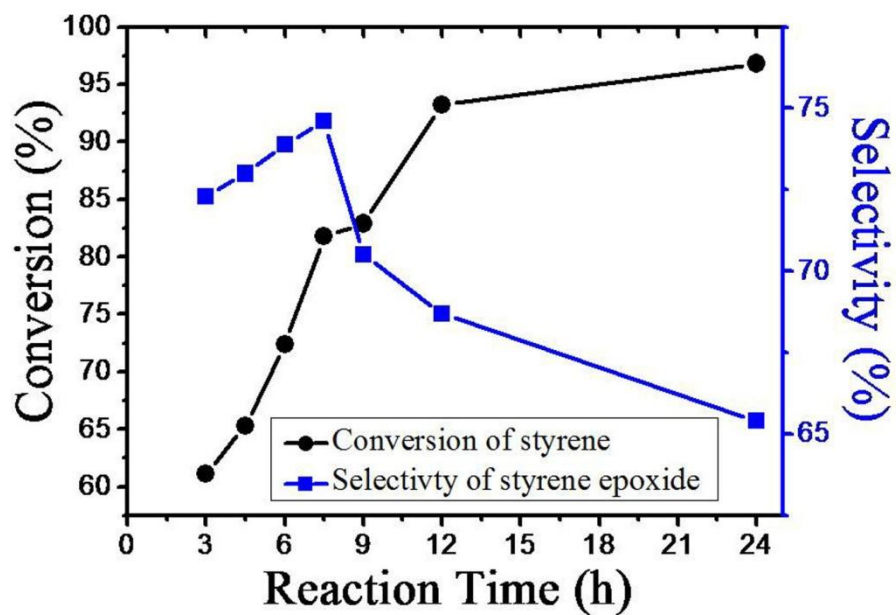


Fig. S5. Conversion of styrene and selectivity of styrene epoxide versus reaction time catalyzed by as-prepared CuO nanoparticles.

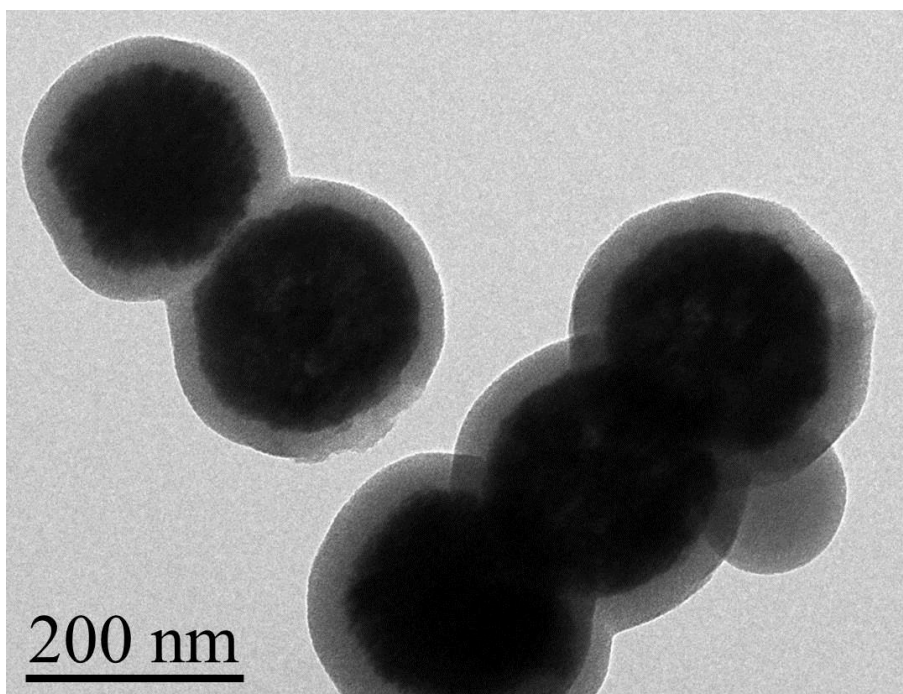


Fig. S6 HRTEM photograph of the $\text{Fe}_3\text{O}_4\text{-CuO@meso-SiO}_2$ catalyst after eight runs of styrene epoxidation.

References:

1. X. L. Huang, J. Zhuang, D. Chen, H. Y. Liu, F. Q. Tang, X. Y. Yan, X. W. Meng, L. Zhang and J. Ren, *Langmuir*, 2009, **25**, 11657.
2. Y. H. Deng, D. W. Qi, C. H. Deng, X. M. Zhang and D. Y. Zhao, *J. Am. Chem. Soc.*, 2008, **130**, 28.