

Supplementary Information for

CO₂-in-PEG emulsion-templating synthesis of poly(acrylamide)s with controllable porosity and their use as efficient catalyst supports

Zhimin Xue,^{a*} Weihong Chang,^a Yan Cheng,^b Jing Liu,^b Jian Li,^b Wancheng Zhao^c and Tiancheng Mu^{c*}

^a Beijing Key Laboratory of Lignocellulosic Chemistry, College of Materials Science and Technology, Beijing Forestry University, Beijing 100083, China. Email: zmxue@bjfu.edu.cn

^b Key Laboratory of TCM Quality Control Technology, Shandong Analysis and Test Center, 250014 Jinan, China.

^c. Department of Chemistry, Renmin University of China, Beijing 100872, China. Tel: 86-10-62514925, Email: tcmu@ruc.edu.cn

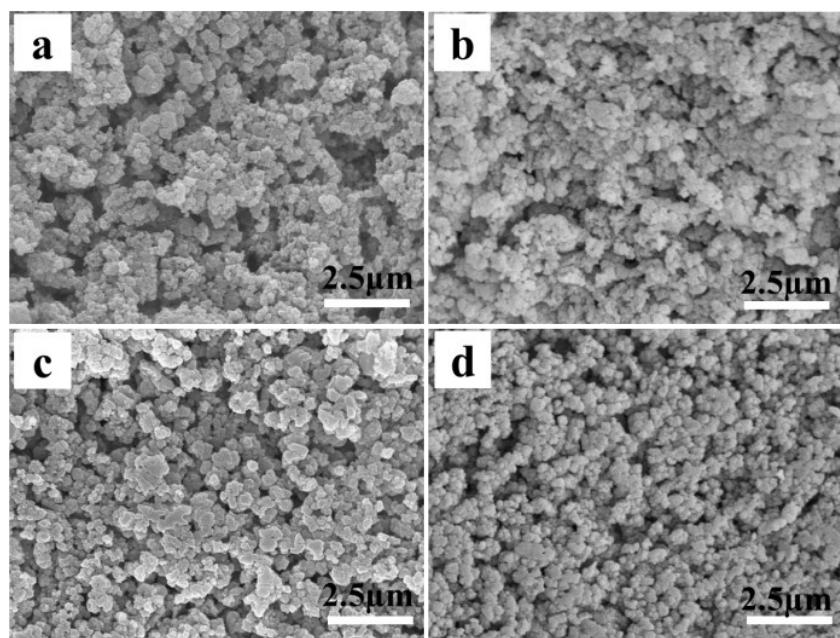


Fig. S1. SEM images of polymer synthesized in CO₂-in-PEG emulsion at 8.0 MPa (a), 10.0 MPa (b), 12.0 MPa (c) and 14.0 MPa (d).

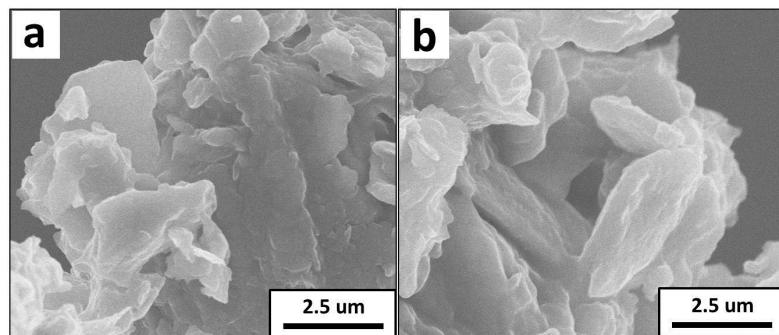


Fig. S2. SEM images of polymer synthesized in PEG alone (a) or the system of PEG and P104 (b) in the absence of CO₂.

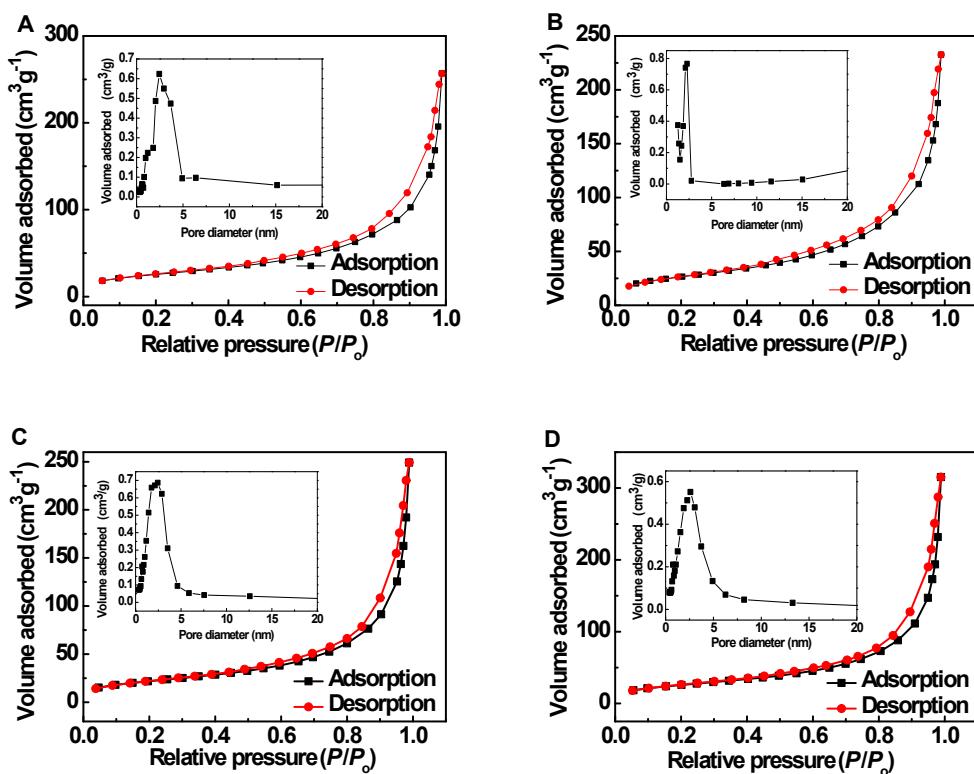


Fig. S3. Mesoporous size distribution of the PAM synthesized at 8 MPa (A), 10 MPa (B), 12 MPa (C) and 14 MPa (D). The inset graphics were the pore size distribution calculated using BJH model.

Table S1 Effect of pressure on the porosity, the total macropore volume (V_{pore}), median macropore diameter (D_{macro}), BET surface area (S_{BET}), total pore volume (V_t) and mesopore diameter (D_{meso}) of PAM synthesized in CO₂-in-PEG emulsion.

Pressure /MPa	Porosity /%	V _{pore} /cm ³ g ⁻¹	S _{BET} /m ² g ⁻¹	V _t /cm ³ g ⁻¹	D _{meso} /nm
8.0	73.3	2.5	74.3	0.29	2.16
10.0	79.9	2.5	83.6	0.33	2.28
12.0	78.1	2.8	90.4	0.41	2.41
14.0	79.6	2.9	94.3	0.48	2.49

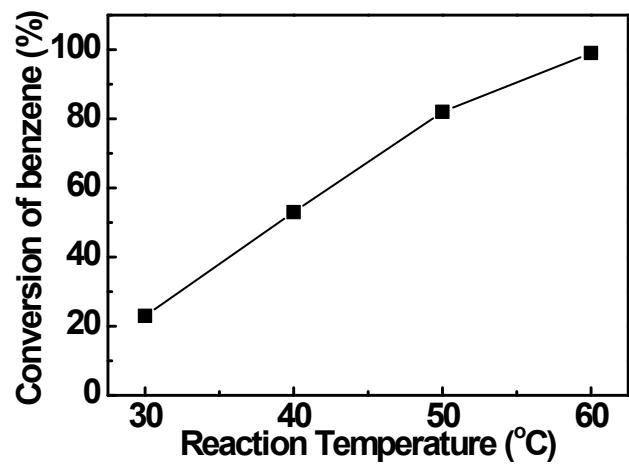


Fig. S4. Effect of reaction temperature on the conversion of benzene over Ru/PAM. Reaction conditions: benzene 1.1 mL; benzene/Ru (mol/mol) = 5000; reaction time 2.5 h.