

Supporting information

In Situ fabrication of core-shell-structured Beta@Silicalite-1 catalysts by a novel steam-assisted crystallization strategy

Lanjian Xu,^{ab} Yangyang Yuan,^{*a} Jie Zhang,^c Yanfei Zhang,^{ab} Xiaomin Zhang,^a Lei Chen,^{ab} Lei Xu^{*a}

^a National Laboratory for Clean Energy, Dalian Institute of Chemical Physics, Chinese Academy of Sciences, Dalian 116023, People's Republic of China.

^b University of Chinese Academy of Sciences, Beijing 100049, People's Republic of China.

^c College of Chemistry and Molecular Engineering, Peking University, Beijing 100871, People's Republic of China.

*E-mail addresses: leixu@dicp.ac.cn, yuanyangyang@dicp.ac.cn

Table S1 The SiO₂/Al₂O₃ molar ratio of the pure Beta(800) zeolite (A), the solid obtained (B) treated by only TPAOH via SAC method, core-shell sample of 4 h (C) and 24 h (D) measured by ICP

Fig. S1 The mass change curve of obtained solid for different crystallization time.

Fig. S2 The XRD pattern of the core Beta zeolites (A) and the solid obtained (B) treated by only TPAOH via SAC method.

Fig. S3 The S_{EBT} (A) and Coverage (B) of core-shell zeolite for different crystallization time

Fig. S4 SEM images of core-shell zeolite Beta(600)@Silicalite-1

Fig. S5 The XRD patterns of Beta(800)@Silicalite-1 synthesized by TEOS (A) and CS (B)

Fig. S6 The N₂ adsorption isothermals Beta(800)@Silicalite-1 synthesized by TEOS (A) and CS (B)

Fig. S7 NH₃-TPD profiles of core Beta(800) (A) and core-shell zeolite Beta(800)@Silicalite-1 (B)

Table S2 Acidity characteristic of core Beta(800) (A) and core-shell zeolite Beta(800)@Silicalite-1 (B)

Table S3 Catalytic performance of Beta(800) (A) and Beta(800)@Silicalite-1 (B) in conversion 1, 3, 5-trimethylbenzene (TMB) and benzyl alcohol (BA)

Table S1 The SiO₂/Al₂O₃ molar ratio of the pure Beta(800) zeolite (A), the solid obtained (B) treated by only TPAOH via SAC method, core-shell sample of 4 h (C) and 24 h (D) measured by ICP

Sample	SiO ₂ /Al ₂ O ₃
A	727
B	717
C	772
D	701

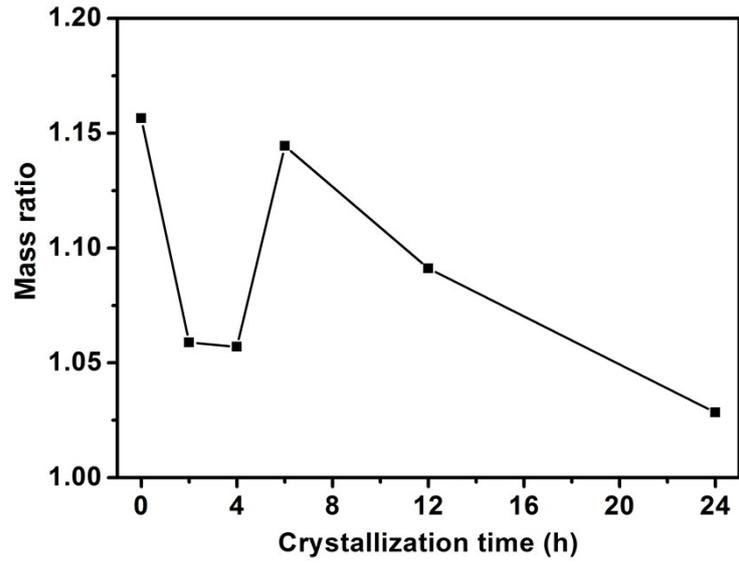


Fig. S1 The mass change curve of obtained solid for different crystallization time.

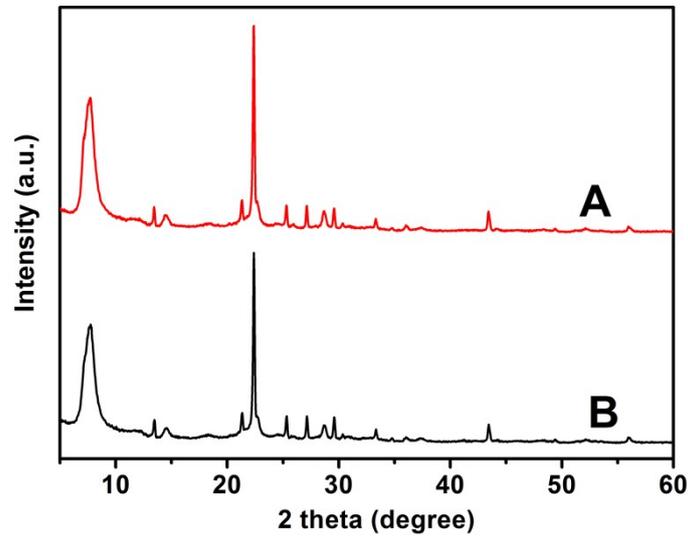


Fig. S2 The XRD pattern of the core Beta zeolites (A) and the solid obtained (B) treated by only TPAOH via SAC method.

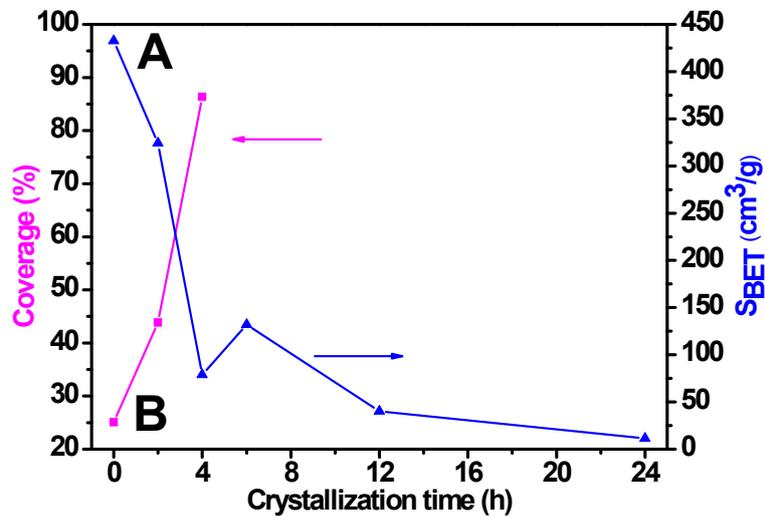


Fig. S3 The S_{EBT} (A) and Coverage (B) of core-shell zeolite for different crystallization time.

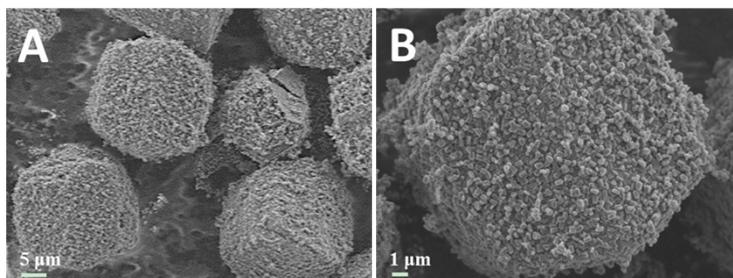


Fig. S4 SEM images of core-shell zeolite Beta(600)@Silicalite-1.

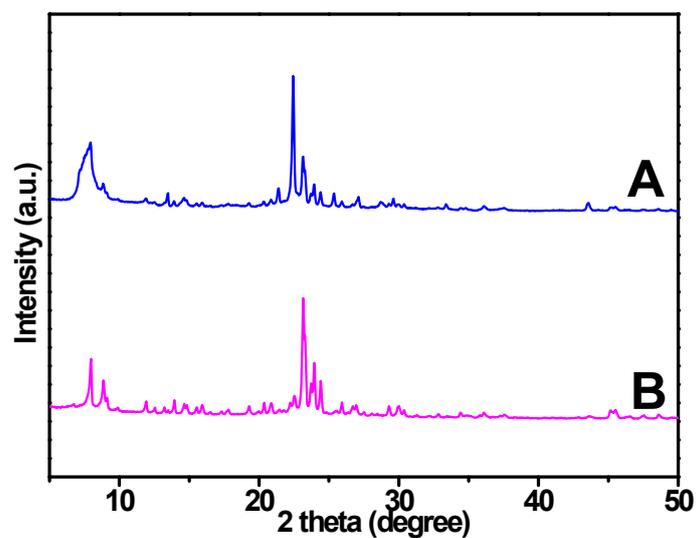


Fig. S5 The XRD patterns of Beta(800)@Silicalite-1 synthesized by TEOS (A) and CS (B).

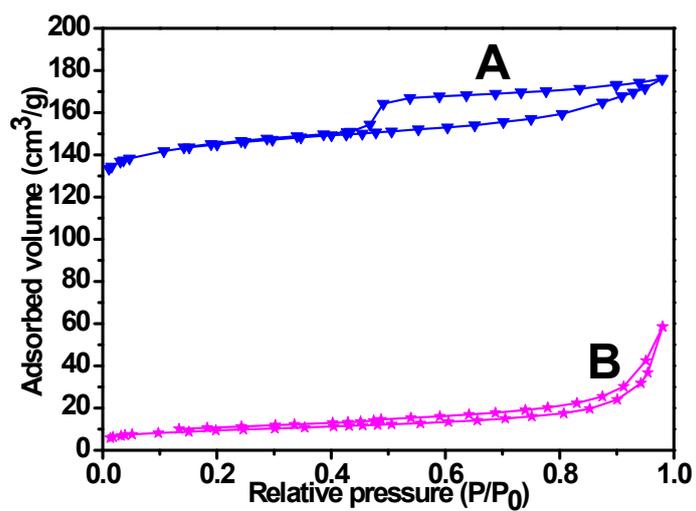


Fig. S6 The N_2 adsorption isothermals Beta(800)@Silicalite-1 synthesized by TEOS (A) and CS (B).

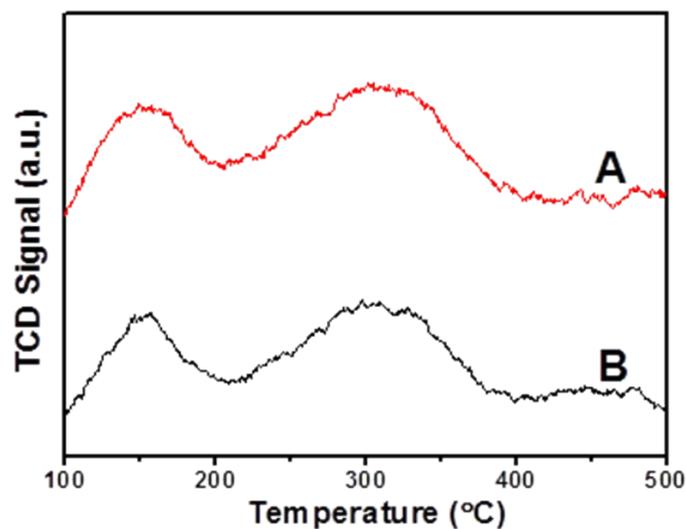


Fig. S7 NH₃-TPD profiles of core Beta(800) (A) and core-shell zeolite Beta(800)@Silicalite-1 (B).

Table S2. Acidity characteristic of core Beta(800) (A) and core-shell zeolite Beta(800)@Silicalite-1 (B)

Sample	T _{peak} (°C)		Acid amount ^b (μmol·g ⁻¹)		
	LT peak ^a	HT peak ^a	Total acidity	Weak acidity	Strong acidity
A	148.5	308.9	47.8	14.9	32.9
B	151.0	308.4	35.7	10.8	24.9

^a The LT peak represents a low temperature desorption peak. The HT peak represents a high temperature desorption peak.

^b Calculated with Gaussian function fit.

Table S3. Catalytic performance of Beta(800)@Silicalite-1 (A) and Beta(800) (B) in conversion 1, 3, 5-trimethylbenzene (TMB) and benzyl alcohol (BA)

Sample	Time (h)	α (2C _{DE} /C _{TMBB})	Conversion (%)
A	2	90.25	4.0
	4	89.71	5.3
	6	87.80	6.5
	8	85.17	7.1
B	2	23.40	3.5
	4	22.89	7.3
	6	21.12	8.3
	8	20.47	9.8