

Electronic Supplementary Information

Rapid detemplation of nanozeolite β : microwave-assisted Fenton-like oxidation

Yuanyuan Hu, Yahong Zhang* and Yi Tang

The detailed calculation of $\text{H}_2\text{O}_2/\text{TEA}^+$ and $\text{Fe}^{3+}/\text{H}_2\text{O}_2$ are showed as follow:

For the as-prepared nanozeolite β :

$$\frac{\text{H}_2\text{O}_2(\text{mol})}{\text{TEA}^+(\text{mol})} = \frac{\frac{V_{\text{H}_2\text{O}_2} \text{ ml} \times 1.1 \text{ g/ml} \times 30\%}{34 \text{ g/mol}}}{\frac{150 \text{ mg} \times 0.8 \times 10^{-3}}{147 \text{ g/mol}}} = 11.89 V_{\text{H}_2\text{O}_2} \quad (1)$$

Herein, the amount of TEA^+ in the solution is determined by the volume of as-prepared β nanozeolite solution and the mass of TEAOH in the synthesis receipts, which means that $\text{TEAOH(g)}/\text{Nanozeolite}_{\text{as-prepared}}(\text{g})$ is 0.8 and the mass of β nanozeolite added in the system is 150 mg. $V_{\text{H}_2\text{O}_2}$ presents the addition volume of H_2O_2 in the as-prepared β nanozeolite system.

For the washed β nanozeolite:

$$\frac{\text{H}_2\text{O}_2(\text{mol})}{\text{TEA}^+(\text{mol})} = \frac{\frac{V'_{\text{H}_2\text{O}_2} \text{ ml} \times 1.1 \text{ g/ml} \times 30\%}{34 \text{ g/mol}}}{\frac{150 \text{ mg} \times 0.2 \times 10^{-3}}{130 \text{ g/mol}}} = 42.06 V'_{\text{H}_2\text{O}_2}$$

The mass of TEA^+ in the washed nanozeolite solution was determined by the TG analysis (Fig. 2b-I). It was indicated (Fig. 2b-I) that TEA^+ in the micropores of the as-prepared β could reach about 20 wt%, i.e., $\text{TEA}^+(\text{g})/\text{Nanozeolite}_{\text{washed}}(\text{g}) = 0.2$, and the mass of β nanozeolite added in the system is 150 mg. $V'_{\text{H}_2\text{O}_2}$ presents the addition volume of H_2O_2 in the washed β nanozeolite system.

$$\frac{Fe^{3+}(\mu mol)}{H_2O_2(mol)} = \frac{\frac{V_{Fe^{3+}} \mu L \times 0.2 mol / L}{V_{H_2O_2}(V'_{H_2O_2}) ml \times 1.1 g / ml \times 30\%}}{34 g / mol} = 20.606 \frac{V_{Fe^{3+}}}{V_{H_2O_2}(V'_{H_2O_2})}$$

$V_{Fe^{3+}}$ presents the addition volume of Fe (NO₃)₃ of 0.2 M in the as-prepared (or washed) β nanozeolite system.

Table S1 Fe/Si ratios in Fenton-treated β with different compositions before and after acid wash step.

Sample ^a	8	5	9	10
Fe ³⁺ / H ₂ O ₂ (10 ⁻⁶ mol/mol)	15.45	9.27	4.64	2.32
Fe/Si before acid wash ^b	2.6×10 ⁻³	2.1×10 ⁻³	1.9×10 ⁻³	1.5×10 ⁻³
Fe/Si after acid wash ^b	n.d. ^c	n.d.	n.d.	n.d.

a: the No. of sample and reaction condition are corresponding to those in Table 1. b: The Fe/Si ratios of nanozeolites were detected by EDX (Philips XL 30). c: n.d. = not detected.

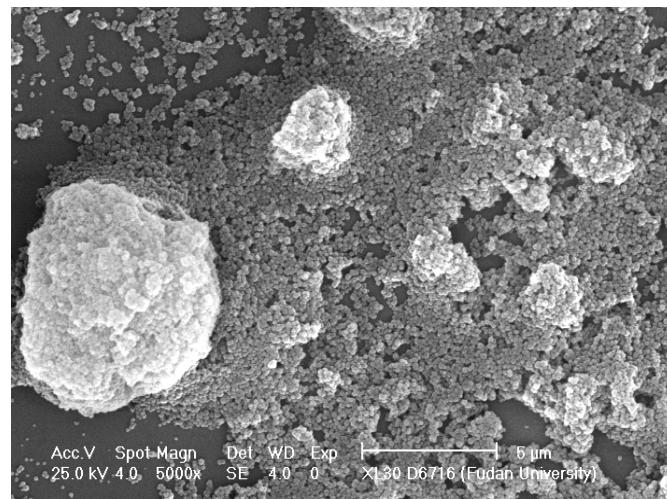


Fig. S1 SEM image of calcined β nanozeolite.

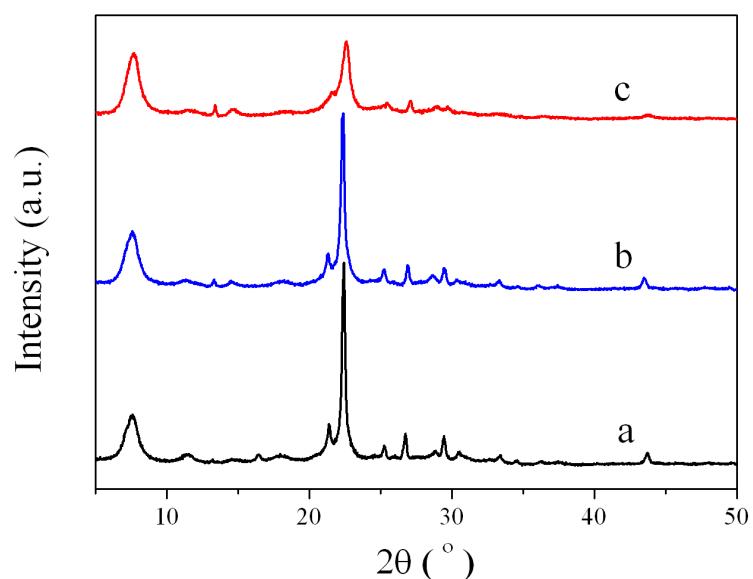


Fig. S2 XRD patterns of as-prepared β (a), Fenton-treated β (b) and calcined β (c).

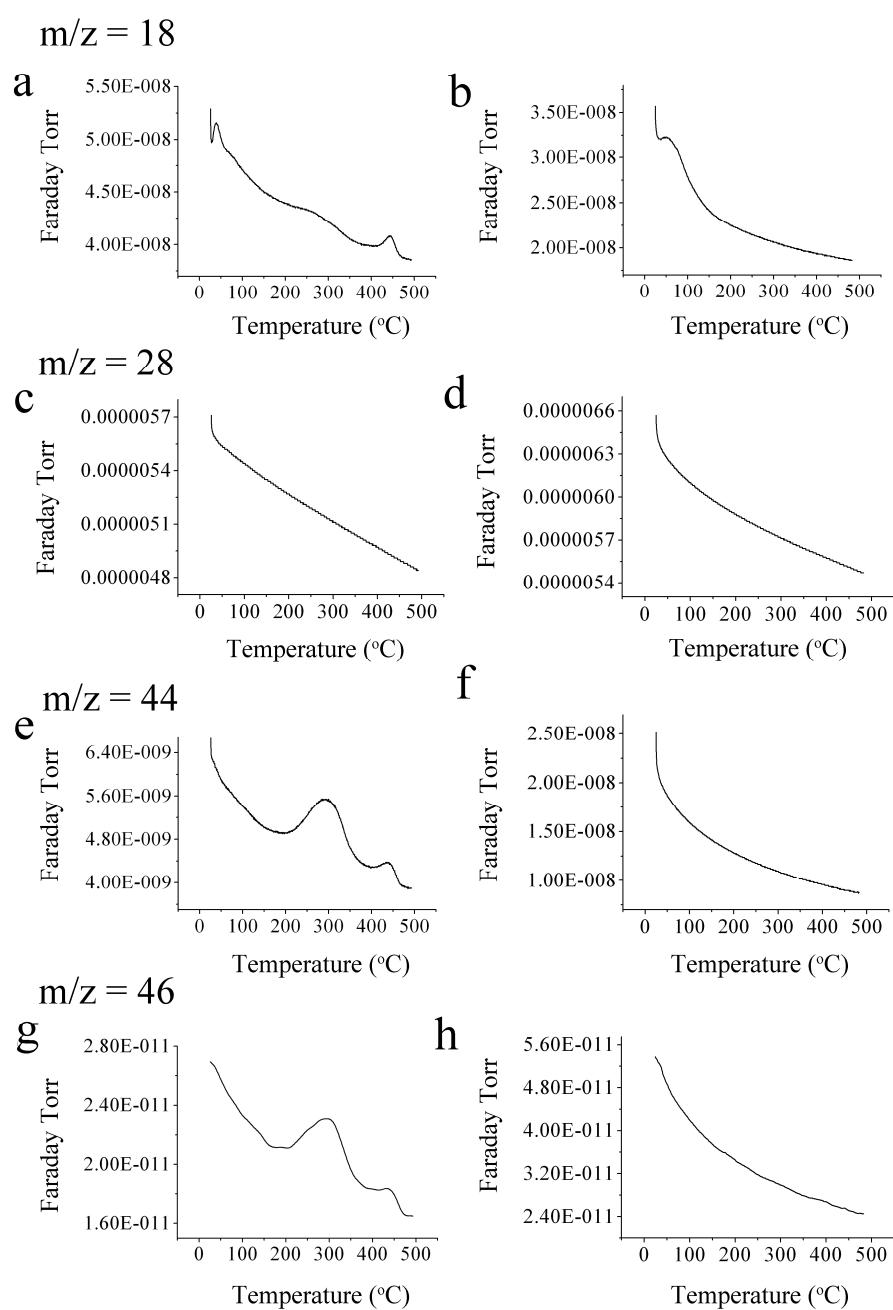


Fig. S3 MS signals of H₂O ($m/z = 18$), CO ($m/z = 28$), CO₂ ($m/z = 44$) and NO₂ ($m/z = 46$) evolved when the as-prepared β (a, c, e, g) and Fenton-treated β (b, d, f, h) heated from 25 to 480 °C in air

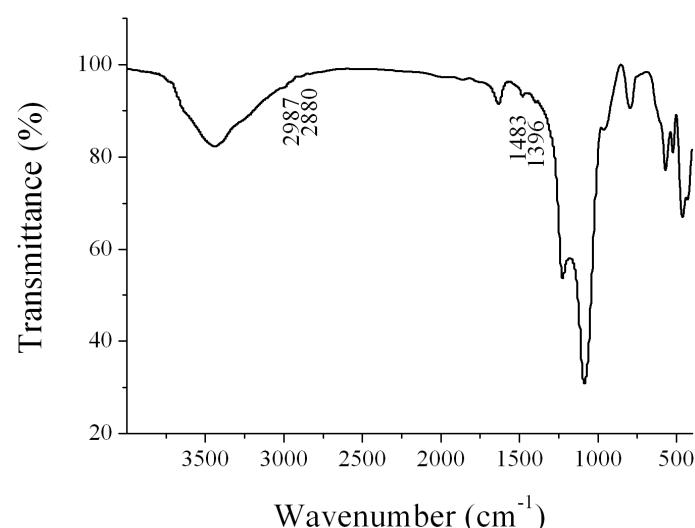


Fig. S4 FTIR spectrum of the Fenton-treated sample by an electric oven at 170 °C for 120 min. The C-H stretching (2987 and 2880 cm^{-1}) and bending (1483 and 1396 cm^{-1}) vibrations corresponding to SDAs are still observed.