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Supplementary Information

Catalytic properties and deactivation behavior of H-MCM-22 in the conversion of methanol to hydrocarbons

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Fig. S1 NH₃-TPD profiles of the parent and oxalic acid-treated H-MCM-22 samples; (a) H-MCM-22-15, (b) H-MCM-22-15-o, (c) H-MCM-22-25, (d) H-MCM-22-25-o, (e) H-MCM-22-37, (f) H-MCM-22-37-o and (g) H-MCM-22-50.



Fig. S2 Deconvolved ²⁷Al MAS NMR spectra of H-MCM-22-15 and H-MCM-22-50 (the dash line is the simulated spectra).



Fig. S3 Product selectivity obtained over the parent and the oxalic acid-treated H-MCM-22 samples (Reaction conditions: 450 °C, WHSV of 2 h⁻¹); (A) H-MCM-22-15; (B) H-MCM-22-15-o; (C) H-MCM-22-25; (D) H-MCM-22-25-o; (E) H-MCM-22-37; (F) H-MCM-22-37-o; (G) H-MCM-22-50; (\blacksquare) ethane and propane, (\bullet) butane, (\blacktriangle) C6+ (aliphatic and aromatics hydrocarbons).



Fig. S4 DTA profiles of the coked (a) H-MCM-22-15, (b) H-MCM-25-22, (c) H-MCM-22-37 and (d) H-MCM-22-50.



Fig. S5 *p*-Xylene adsorption isotherms at 25 °C on the fresh and coked H-MCM-22-x and H-MCM-22-x-o.



Fig. S6 GC-MS chromatogram of the coke species extracted with CH_2Cl_2 from (A) H-MCM-22-15, (B) H-MCM-22-25, (C) H-MCM-22-25-0 and (D) H-MCM-22-37 at different reaction time (Reaction conditions: 450 °C, WHSV = 2 h⁻¹. The asterisk (*) represents the internal standard (C_2Cl_4)).



Fig. S7 Raman spectra of the coke species deposited on the (a) H-MCM-22-25 and (b) H-MCM-22-37.



Scheme S1 MCM-22 framework with 8 crystallographically inequivalent T sites.