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## **Supporting Information**

for

Small-pore molecular sieves SAPO-57 and SAPO-59: synthesis, characterization, and

## catalytic properties in methanol-to-olefins conversion

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Material	Relative amount (%) of total acid sites <sup>a</sup>	Relative distribution (%)		
		Weak acid site $(250 - 280 \text{ °C})^b$	Intermediate acid sit $(320 - 370 \text{ °C})^b$	te Strong acid site $(400 - 450 \text{ °C})^b$
H-SAPO-57( <i>l</i> )	4.78	45	37	18
H-SAPO-57( <i>h</i> )	4.59	48	32	20
H-SAPO-59( <i>l</i> )	1.00	57	24	19
H-SAPO-59( <i>h</i> )	2.80	43	33	24
H-SAPO-34	3.09	12	41	47
H-SAPO-35	3.84	15	42	43

**Table S1** Distributions of acid sites with different degrees of strength in a series of SAPO catalysts determined from deconvolution of the NH<sub>3</sub> TPD data

<sup>*a*</sup> Reported relative to the total area of  $NH_3$  desorption peaks from H-SAPO-59(*l*). <sup>*b*</sup> Temperature range of the desorption peak maxima corresponding to the respective acid sites determined from deconvolution of the  $NH_3$  TPD profiles.

## **Figure Captions**

- Fig. S1 Powder XRD patterns and SEM images of the as-made form of (a) SAPO-34 and(b) SAPO-35 molecular sieves synthesized in this work.
- **Fig. S2** <sup>27</sup>Al MAS NMR spectra of the as-made (left) and proton (right) forms of (a) SAPO-57(*l*), (b) SAPO-57(*h*), (c) SAPO-59(*l*), and (d) SAPO-59(*h*).
- Fig. S3 <sup>31</sup>P MAS NMR spectra of the as-made (left) and proton (right) forms of (a) SAPO-57(*l*), (b) SAPO-57(*h*), (c) SAPO-59(*l*), and (d) SAPO-59(*h*).
- Fig. S4 Powder XRD patterns of (a) H-SAPO-57(*l*), (b) H-SAPO-57(*h*), (c) H-SAPO-59(*l*), and (d) H-SAPO-59(*h*) after MTO at 350 °C and 0.67 h<sup>-1</sup> WHSV for 600 min on stream.



Fig. S1



Fig. S2



Fig. S3



