Electronic Supplementary Information

Sol Preparation

One pure silica sol was prepared using a single-step acid-catalysed hydrolysis of TEOS (Si(OC₂H₅)₄) in ethanol (C₂H₅OH). Nitric acid (HNO₃) was chosen as catalyst. These reactants were mixed together with distilled water (H₂O), in the molar ratio TEOS : C₂H₅OH : HNO₃ : H₂O = 1 : 3.8 : 0.085 : 6.5. This mixture was refluxed for 3 hours at 60°C to yield the pure SiO₂ sol.

One methylated silica sol was prepared analogously to the silica sol. MTES $(Si(OC_2H_5)_3CH_3)$ was used as a source for the methyl groups. A mixture with the same composition as above was refluxed for 2 hours and 45 minutes. At this stage MTES was added in the molar ratio MTES : TEOS = 1 : 1. The mixture thus obtained was further refluxed for another 15 minutes.

Several methylated silica sols with different methyl concentrations were prepared using a two-step process. A mixture with molar ratio TEOS : C_2H_5OH : HNO_3 : $H_2O = 1$: 3.8 : 0.0036 : 2.0 was refluxed at 60°C for 90 minutes. At this stage MTES was added with a molar ratio TEOS : MTES = 10 : 1, 10 : 3, 10 : 4, or 10 : 6, and the reflux was continued for another 30 minutes. This was followed by the addition of more nitric acid and water so that the final molar ratio was TEOS : C_2H_5OH : HNO_3 : $H_2O = 1 : 6.2 : 0.0108 : 6.0$ with variable MTES content. Finally, this was refluxed for 3 more hours, before it was diluted with ethanol. We were unable to prepare membranes for the sols with TEOS : MTES = 10 : 4 and 10 : 6.



Gel Permeation Chromatography.

Figure 1S: Particle size distribution as a function of the elusion time of the sols prepared with the 'single-step' procedure and the sol prepared with the 'two-step' procedure with TEOS:MTES = 10:6. We were unable to prepare a membrane from this last sol.

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Figure 2S: Particle size distribution as a function of the elusion time of the sols prepared with the 'two-step' procedure with TEOS:MTES = 10:1; 10:3, and 10:6. We were unable to prepare a membrane from this last sol.

The molar mass (M_w in kg/mol) can be calculated from the elusion time (t, min) using $M_w = 10^{(-0.66t + 6.26)}$.

Nuclear Magnetic Resonance

Solid state ²⁹Si NMR spectra were collected on free-standing films prepared analogously to the membranes. The spectra were recorded on a Bruker MSL300 and Bruker MSL400, using a spinning rate of 4 kHz and one-pulse experiments with 90° pulses and 100 s recycle delay time. The number of acquisitions varies between 200 and 700, which corresponds to a measurement time of 5 to 19 hours. The summed concentration of T groups is equivalent to the methyl concentration.

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TEOS : MTES	10:1	10:3	10:6	10:10
preparation route	two-step	two-step	two-step	single-step
CH ₃ /Si	0.09	0.23	0.37	0.50
T _{tot}	0.13	0.22	0.36	0.45
T_2	0.02	0.05	0.13	0.22
T ₃	0.11	0.17	0.23	0.23
Q _{tot}	0.88	0.79	0.65	0.55
Q ₂	0.04	0.05	0.05	0.02
Q ₃	0.31	0.32	0.32	0.15
Q ₄	0.52	0.42	0.28	0.38

Table 1S: The concentration of tetrafuntional (Q_i, from TEOS) and trifunctional (T_i, from MTES) units in free-standing films prepared from various sols.



Figure 3S: Sample NMR spectrum including fit and numerical results for a methylated silica film prepared from a 'two-step' sol with initial TEOS : MTES ratio of 10 : 3.

Scanning Electron Microscopy





Figure 4S: Two SEM micrographs showing typical surface appearance of a membrane before testing in pervaporation. The great majority of the membrane surface looks like the micrograph in a). Imperfections on the µm scale, as depicted in b), are present.