Supporting Information for:

Synthesis and Characterization of Functionalized Mesoporous Silica by Aerosol-Assisted Self-Assembly

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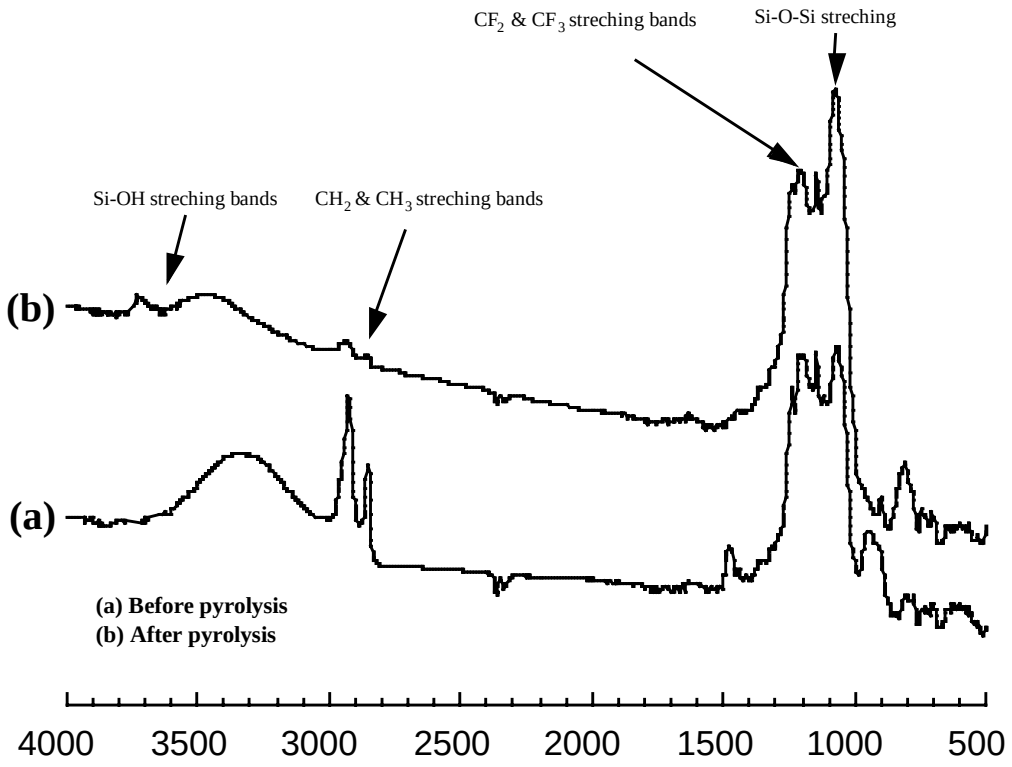

Figure S1. FT-IR spectra of the TFTS-modified silica particles before and after heat treatment at 350 °C for 5 hr in a nitrogen atmosphere.

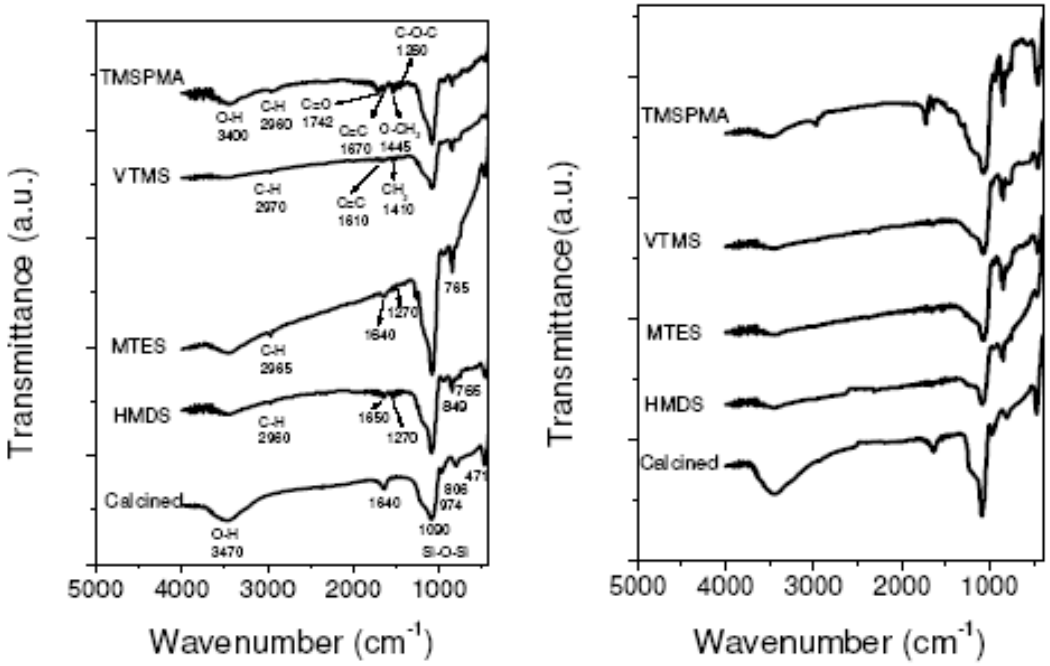

Figure S2. FT-IR spectra for organic functionalized silica particles, (a) Brij-56, (b) CTAB, (c) P123 as a template.

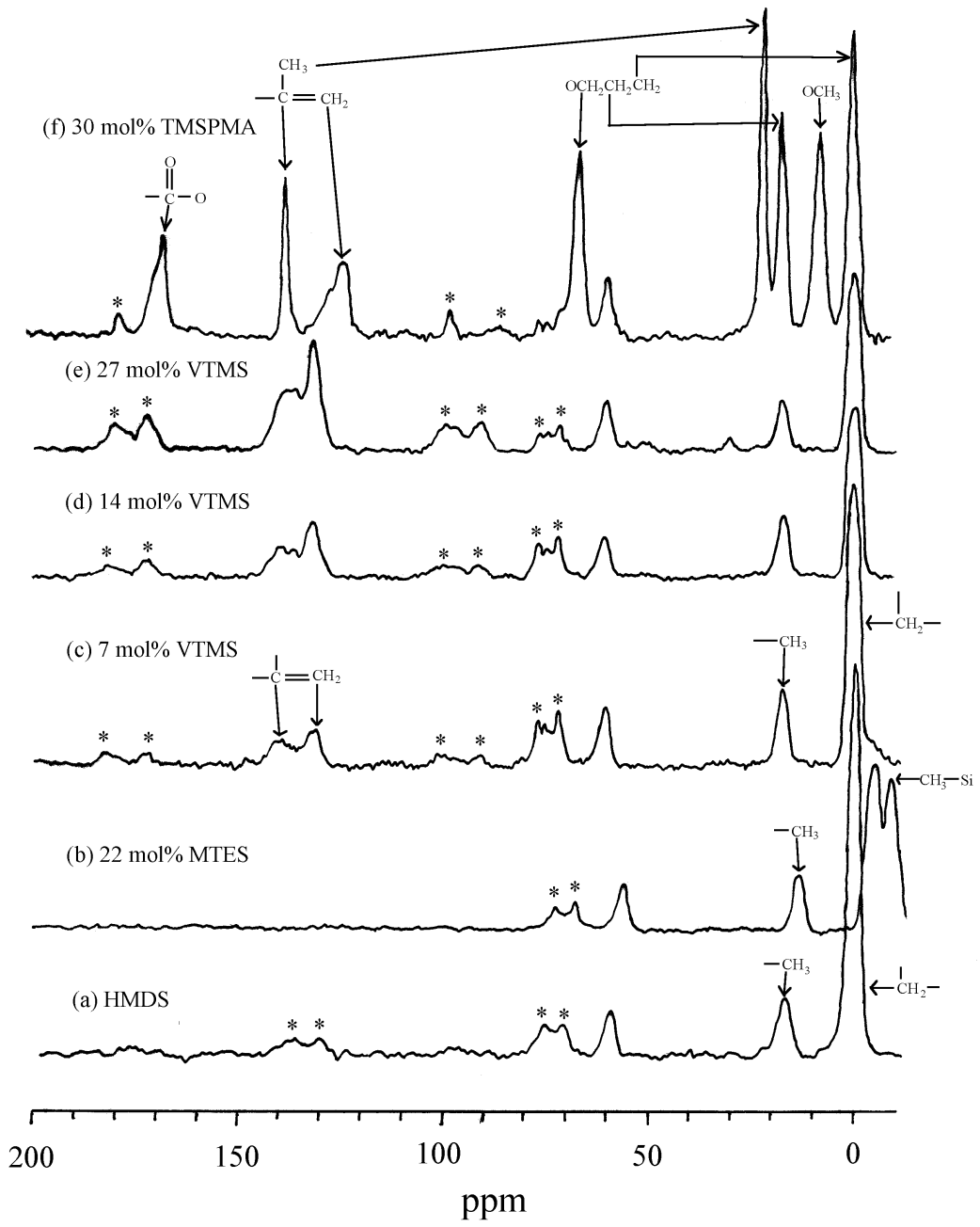

Figure S3. Solid-state ^{13}C CP/MAS NMR spectra, (a) HMDS, (b) 22 mol% MTES, (c) 7 mol% VTMS, (d) 14 mol% VTMS, (e) 27 mol% VTMS, (f) 30 mol% TMSPMA, * spinning side bands.

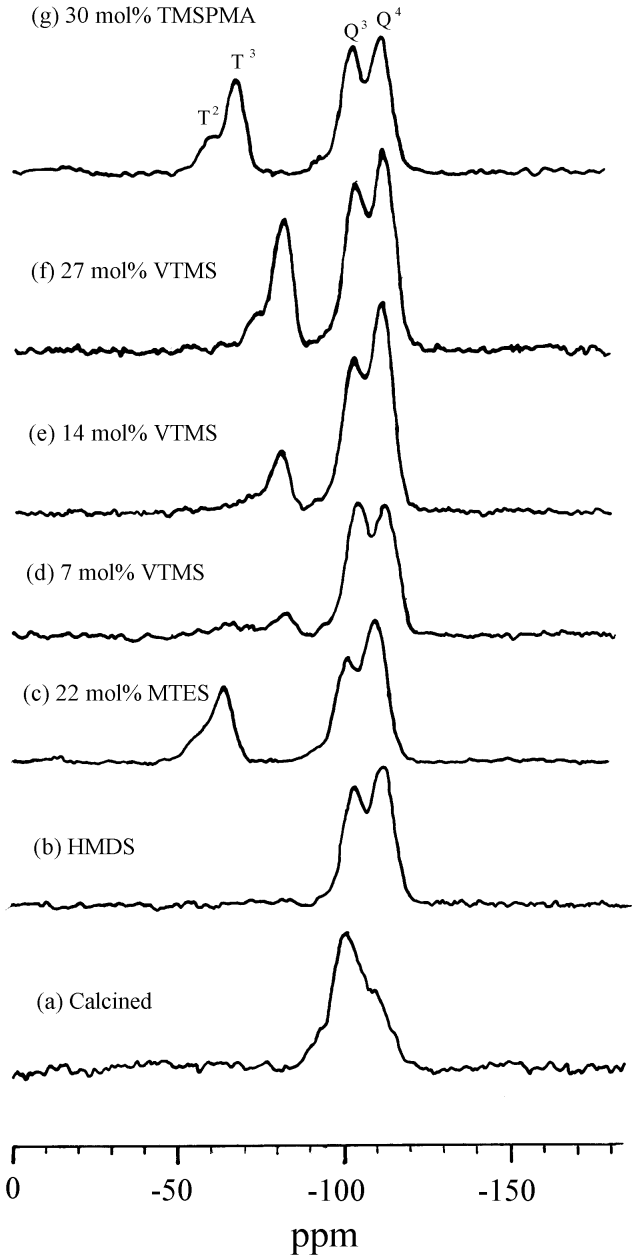

Figure S4. Solid-state ^{29}Si MAS NMR spectra, (a) calcined at 450^{o}C for 4 h, (b) HMDS, (c) 22 mol% MTES, (d) 7 mol% VTMS, (e) 14 mol% VTMS, (f) 27 mol% VTMS, (g) 30 mol% TMSPMA. (The cross-polarization (CP) contact time used in this study was 8 ms)

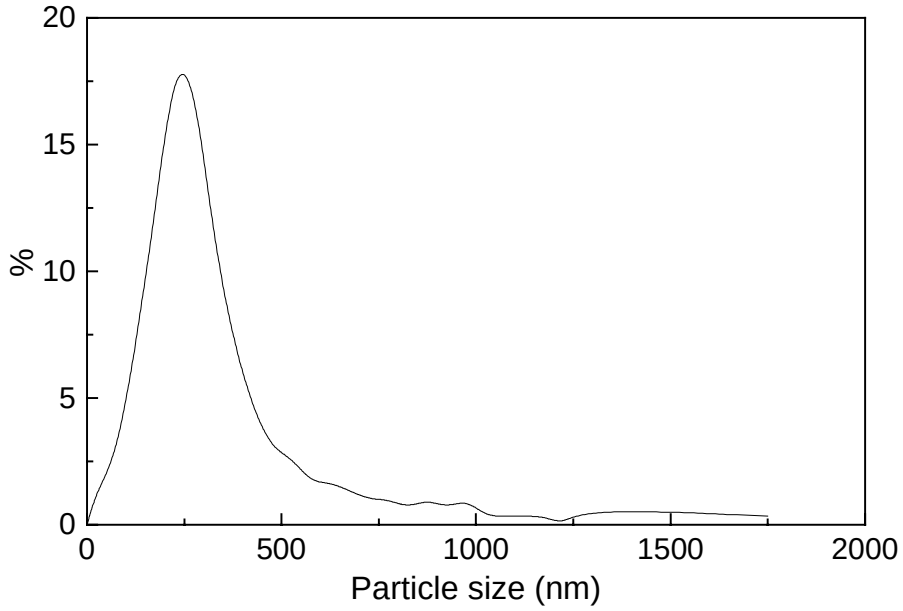

Figure S5 Particle size distribution of samples using P123 as a template