SUPPORTING INFORMATION

Characterization of ZnE (E = S, Se, Te) Materials Synthesized Using Silylated–Chalcogen Reagents in Mesoporous MCM-41

Elizabeth A. Turner[†], Harald Rösner[‡], Dieter Fenske[‡], Yining Huang^{†*} and John F.

Corrigan^{†*}

†Department of Chemistry, The University of Western Ontario, London, Ontario N6A

5B7, Canada and ‡Institüt für Nanotechnologie, Forschungszentrum Karlsruhe GmbH,

76344 Eggenstein-Leopoldshafen, Germany

Supplementary Material.

Figure S1. TGA curves for functionalized MCM-41 monitored over a 96 hour time period. Percentage indicates percent loss during heating.

Figure S2. TGA curves for Zn-MCM-41 monitored over a 96 hour time period. Percentage indicates percent loss during heating.

Figure S3. Raman spectra of TPED, F-MCM-41 and Zn-MCM-41. C–H deformation of ethylene diamine (×).

Table S1. Peak area ratios between $S(SiMe_3)_2$, Me_3SiOAc and $O(SiMe_3)_2$ and the internal standard (toluene) taken at specified time intervals during the synthesis of ZnS-MCM-41. Shown are the ratios observed when an excess of $S(SiMe_3)_2$ and an approximately stoichiometric (1:0.5) amount of $S(SiMe_3)_2$ are used in the synthesis.



Figure S1.



Figure S2.

Raman Spectra

(×) C–H deformation of ethylene moiety



Figure S3.

Table S1.

| | Peak Area Ratios Between Designated Product and Toluene Standard | | | | | |
|---------|--|-----------|-----------------------|-----------|------------------------------------|-----------|
| Time | S(SiMe ₃) ₂ | | Me ₃ SiOAc | | O(SiMe ₃) ₂ | |
| [hours] | Excess | No-excess | Excess | No-excess | Excess | No-excess |
| 2 | 1.5 | 0.3 | 0.4 | 0.3 | 0.0 | 0.0 |
| 4 | 1.3 | 0.2 | 0.4 | 0.4 | 0.0 | 0.0 |
| 22 | 1.1 | 0.0 | 0.4 | 0.4 | 0.2 | 0.2 |
| 70 | 1.0 | 0.0 | 0.4 | 0.4 | 0.5 | 0.3 |
| 90 | 0.9 | 0.0 | 0.4 | 0.27 | 0.9 | 0.3 |
| 120 | 0.8 | 0.0 | 0.4 | 0.11 | 1.1 | 0.3 |
| 160 | 0.4 | 0.0 | 0.4 | 0.0 | 1.8 | 0.4 |