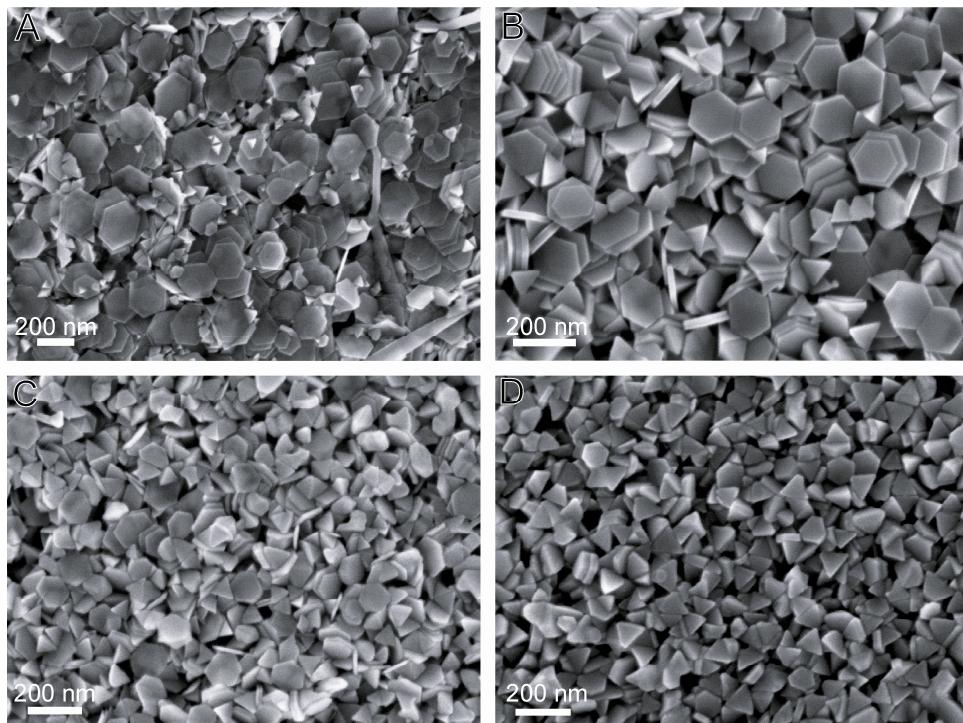


## Supporting Information for Synthesis of Metal Sulfide Nanomaterials via Thermal Decomposition of Single-Source Molecular Precursors

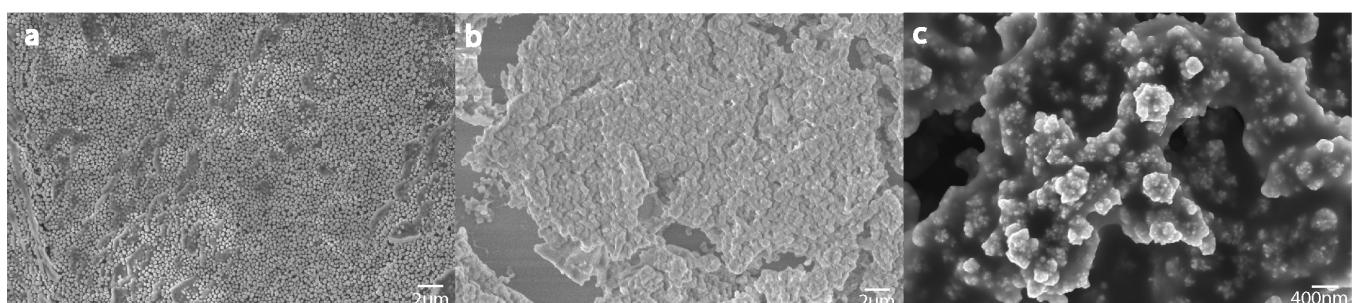
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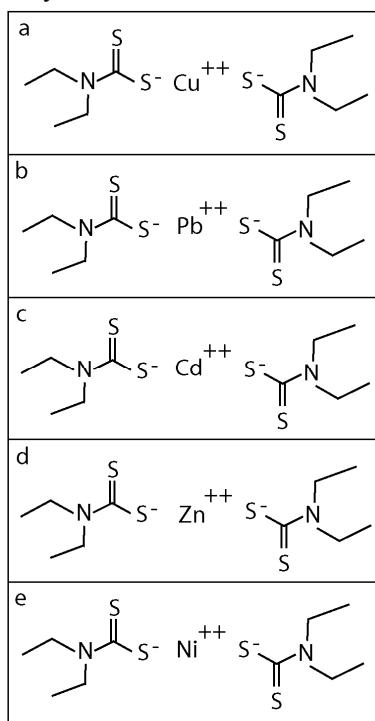
<sup>‡</sup>Department of Chemistry; <sup>§</sup>Department of Chemical Engineering, University of California, Berkeley; CA 94720



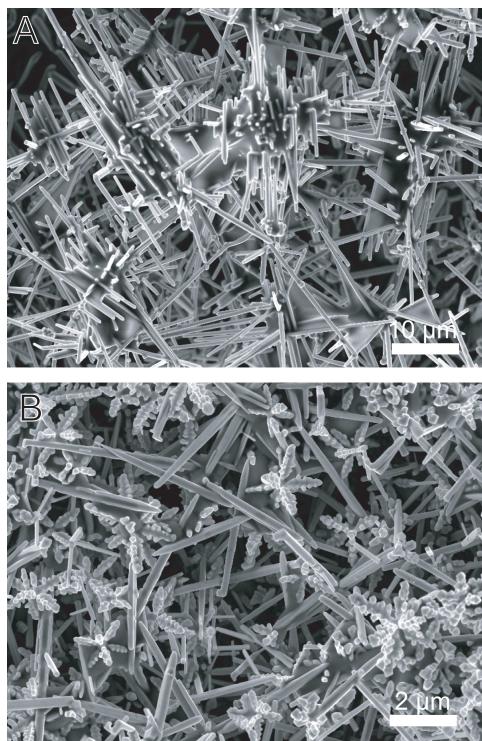
**Figure S1.** Cuprous sulfide ( $\text{Cu}_2\text{S}$ ) hexagonal plates formed from  $\text{Cu}(\text{II})[\text{S}_2\text{CNC}_4\text{H}_{10}]_2$  in trioctylphosphine. Thermal decomposition reactions were performed in solution under standard air-free Schlenk technique. a)  $\text{Cu}(\text{II})[\text{S}_2\text{CNC}_4\text{H}_{10}]_2$  is 0.055M in trioctylphosphine. b) 0.11M c) 0.22M d) 0.33M As the initial concentration increases, the yield of triangular plates to hexagonal plates increases. All structures are the low-chalcocite monoclinic crystal phase.



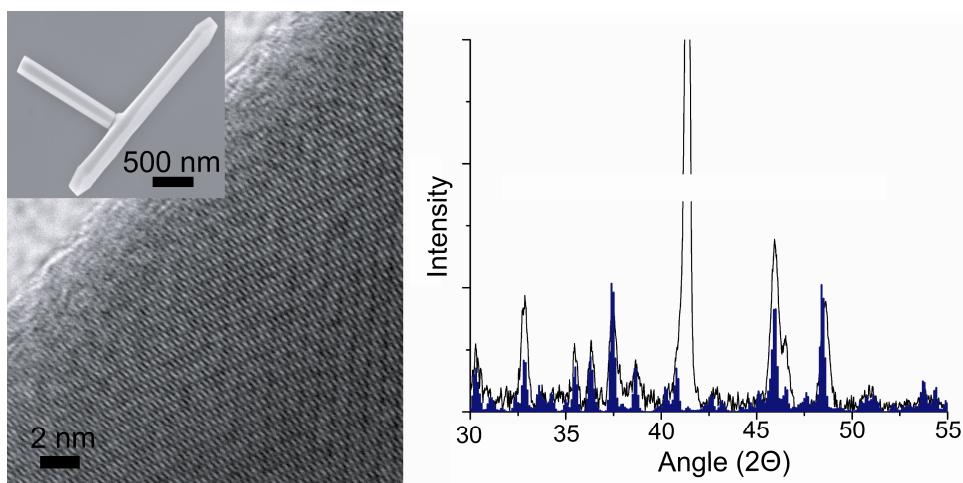
**Figure S2.** Thermal decomposition products of other metal sulfide single-source molecular precursors from trioctylphosphine on substrates. a)  $\text{CdS}$  nanoparticles. b)  $\text{ZnS}$  c)  $\text{Ni}_3\text{S}_2$



**Figure S3.** Single-source molecular precursors used. a-e) Copper, lead, cadmium, zinc, and nickel bis(diethyldithiocarbamate). All synthesized from the metal chloride salt dissolved in deionized water mixed with sodium diethyldithiocarbamate. The product is vacuum filtered, dried, and recrystallized from hot chloroform.



**Figure S4.** Lower magnification SEM images the Cu<sub>2</sub>S and PbS nanowire substrates. Cuprous sulfide nanowires show highly branched structures.



**Figure S5.** A) High-resolution TEM with inset SEM of Cu<sub>2</sub>S an individual nanowire in combination with B) X-ray diffraction confirms the crystal structure of the Cu<sub>2</sub>S nanowires to be the same as those of the nanocrystals, monoclinic.