A General Single-Pot Heating Method for Morphology, Size and Luminescence-Controllable Synthesis of Colloidal ZnO Nanocrystals

Xin Liu and Mark T. Swihart*

Department of Chemical and Biological Engineering, University at Buffalo (SUNY), Buffalo, NY 14260, USA

*Address Correspondence to swihart@buffalo.edu

Supporting Information:



Figure S1. TEM image (left panel) showing quasi-spherical ZnO NPs synthesized in the presence of TOPO, benzyl ether and hexadecanediol. Selected-area electron diffraction (SAED, right panel) showed that these ZnO NPs had poor crystallinity.



Figure S2. HRTEM image showing that the ZnO NPs with pinecone morphology were composed of small ZnO nanocrystals.



Figure S3. HRTEM image showing dendritic ZnO NCs synthesized by thermolysis of 300 mg $Zn(acac)_2$ in the presence of 2.2 g TOPO, 7 mL benzyl ether and 736 mg octadecanol at 290 °C

Synthesis of ZnO NPs using only oleylamine as ligand

150 mg Zn(acac)₂, 2.625 mL oleylamine, 7 mL benzyl ether and 367 mg hexadecanediol were mixed together. The mixture was heated to 250 $^{\circ}$ C and kept at this temperature for 1hr.



Figure S4. TEM and HRTEM (inset) of ZnO NPs synthesized using only OAm as ligand.

Synthesis of ZnO NPs using only dodecylamine as ligand

150 mg $Zn(acac)_2$ precursor was mixed with 738 mg dodeclyamine, 367 mg hexadecanediol and 7 ml benzyl ether. The solution was heated to 295 °C and kept at this temperature for 60 min.



Figure S5. TEM of ZnO NPs synthesized using only dodecylamine as ligand.

Synthesis of ZnO NPs using TOPO and OAm as the ligands

150 mg Zn(acac)₂ precursor was mixed with 1.55 g TOPO, 2.5 ml OAm and 735 mg hexadecanediol. The solution was heated to 205 $^{\circ}$ C and kept at this temperature for 30 min.



Figure S6. TEM of ZnO NPs synthesized using TOPO and OAm as ligands.