Supplementary Material

Sonochemical synthesis of cadmium(II) coordination polymer nanospheres as precursor for cadmium oxide nanoparticles

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<i>D</i> —НА	<i>D</i> —Н	НА	D A	<i>D</i> —НА
N2-H2BO1W	0.79	1.93	2.869(7)	163
O1W-H1WAN6 ⁱⁱⁱ	0.91	1.97	2.687(7)	172
O12-H12CO3 ^{iv}	0.97	1.75	2.712(7)	171
O1W-H1WBO10 ⁱ	0.92	2.32	3.162(7)	152
C2-H2AO10 ⁱ	0.95	2.44	3.322(8)	155
C14-H14AO7	0.95	2.52	3.392(8)	153
C7-H7AO1W ⁱⁱ	0.95	2.67	3.239(8)	119

Table S1. Hydrogen bond parameters in 1.

Symmetry codes: (i) -*x*-2, *y*-1/2, -*z*; (ii) *x*, *y*, *z*; (iii) -*x*-1, *y*+1/2, -*z*; (iv) -*x*-2, *y*-1/2, -*z*-1.



Branched Tube Synthesis/Crystallization Technique

Fig. S1. Schematic representation of the branched tube synthetic method.



Fig. S2. Fragment of the crystal packing pattern of **1** (view along the *a* axis). CH hydrogen atoms are omitted for clarity. Cd (yellow balls), N (blue), O (red), C (gray), H (dark gray).



Fig. S3. Topological representation of a 3D H-bonded network in 1 showing a binodal 3,5-connected net with the hms (3,5-conn) topology and point symbol of $(6^3)(6^9.8)$. (A) View along the *b* axis. (B) View along the *c* axis. Cd2-based [Cd(L)(NO₃)(H₂O)] nodes (gray balls), centroids of 5-connected Cd1-based [Cd(HL)(NO₃)₂] nodes (yellow balls).



Fig. S4. Crystal packing diagram of 1 along the *b* axis (H bonds are shown as red and blue lines).



Fig. S5. TGA of compound 1.



Fig. S6. Size distribution histograms for nanoparticles of 1 (A) and CdO (B).



Fig. S7. Comparison between PXRD patterns of (A) nanoparticles 1 and (B) solid formed in methanol (synthesized using a procedure similar to 1 but in MeOH).



Fig. S8. FT-IR spectra of (A) compound **1** and (B) solid formed in methanol (synthesized using a procedure similar to **1** but in MeOH).