Supporting Information:

Sonochemistry-assisted synthesis and optical properties of mesoporous ZnS nanomaterials

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Ultrasonic time /min	Crystallite size /nm	$S_{\rm BET}/{\rm cm}^2~{\rm g}^{-1}$	$V_{\rm pore}/{\rm cm}^3 {\rm g}^{-1}$	$D_{\rm DFT}/{\rm nm}$	k / \min^{-1}
0 min	7.2	160	0.42	4.9	0.0138
1 min	5.9	197	0.41	3.6	0.0154
3 min	4.5	226	0.36	5.9	0.0188
5 min	3.8	263	0.31	5.1	0.0245
20 min	3.8	258	0.32	5.1	0.0242

Table S1 Summary of the physicochemical properties and photodegradation rate constant of the synthesized s-ZnS materials in ethanol system after various duration of ultrasonic irradiation.

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Fig. S1 XRD patterns of the ZnS nanomaterials synthesized in the ethanol solutions with the different ultrasonic irradiation time.



Fig. S2 N_2 sorption isotherms (left) and the corresponding pore size distribution curves (right) of the s-ZnS nanomaterials synthesized in the ethanol solutions with the different ultrasonic irradiation time. The volume adsorbed was shifted by 500, 380, 240 and 120, and dV/dD value was shifted by 0.25, 0.20, 0.14, and 0.08 for the curves of the s-ZnS samples after 0, 1, 3, 5 and 20 min ultrasonic treatment, respectively.

Electronic Supplementary Material (ESI) for Journal of Materials Chemistry A This journal is $\ensuremath{\mathbb{O}}$ The Royal Society of Chemistry 2013



Fig. S3 Time-dependent PL emission spectra of the ZnS materials synthesized in ethanol system with the different ultrasonic irradiation time (0, 1, 3, 5, 20 min), showing ultrasonic-time-dependent increase of emission intensity.



Fig. S4 PL emission spectra of the ZnS materials synthesized in the ethanol after 5 min sonication. Initial solution volume and concentration: $Zn(NO_3)_2$ (30 ml): 30, 35 and 48 mmol L⁻¹, Na₂S (30ml): 30 mmol L⁻¹.



Fig. S5 (a) Photoactivities of the ZnS materials synthesized in ethanol system with the different ultrasonic irradiation time for RhB degradation under UV-light irradiation. (b) Plots of $\ln(C_0/C)$ versus the irradiation time, showing the fitting results using the pseudo-first-order reaction.