Supporting Information

Hollow PbWO₄ Nanospindles via a Facile Sonochemical Route

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1. Comparative experimental results under vigorous electric stirring

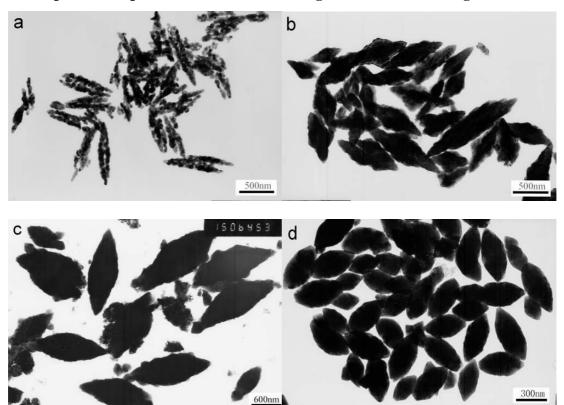


Figure S1. TEM images of PbWO₄ sample prepared under vigorous electric stirring in different P123 concentration systems: (a) $2 \text{ g} \cdot \text{L}^{-1}$, (b) $3 \text{ g} \cdot \text{L}^{-1}$, (c) $4 \text{ g} \cdot \text{L}^{-1}$, and (d) $5 \text{ g} \cdot \text{L}^{-1}$.

2. Effect of sonication time on the final products

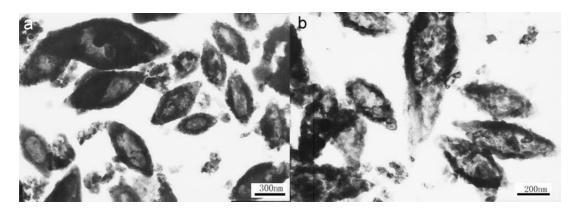


Figure S2. TEM images of samples obtained after sonication for (a) 60 min, and (b) 120 min.

3. Effect of concentrations of Pb(CH₃COO)₂ and Na₂WO₄

The experimental results show that the uniform hollow spindles could only be obtained at the concentration of 0.5-1 M. When the concentration of Pb(CH₃COO)₂ and Na₂WO₄ were changed to 0.1 M (1/5 of optimal concentration), the product was a mixture of hollow spindles and nanorods, and the yield was much lower (Figure S3a). When the concentration of Pb(CH₃COO)₂ and Na₂WO₄ were changed to 2 M (4 times of optimal concentration), the morphology changed to be small nanoparticles together with irregular aggregation (Figure S3b). We think when the concentrations of Pb(CH₃COO)₂ and Na₂WO₄ were relatively low (the P123 concentration was relatively higher), the copolymer might also work as capping reagent to restrict the growth of PbWO₄ crystals resulting the formation of nanorods. While under higher reactant concentration, nucleation occurred at an outburst speed right after the reaction solutions were mixed, leading to a large quantity of PbWO₄ precipitation, so small nanoparticles with their aggregation were obtained under this situation.

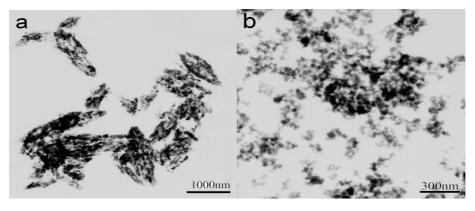


Figure S3. TEM images of samples obtained with the concentration of $Pb(CH_3COO)_2$ and Na_2WO_4 of (a) 0.1 M, and (b) 2 M.