

Supporting Information

Ultralong Hydroxyapatite Microtubes: Solvothermal Synthesis and Application in Drug Loading and Sustained Drug Release

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Supplementary Figures

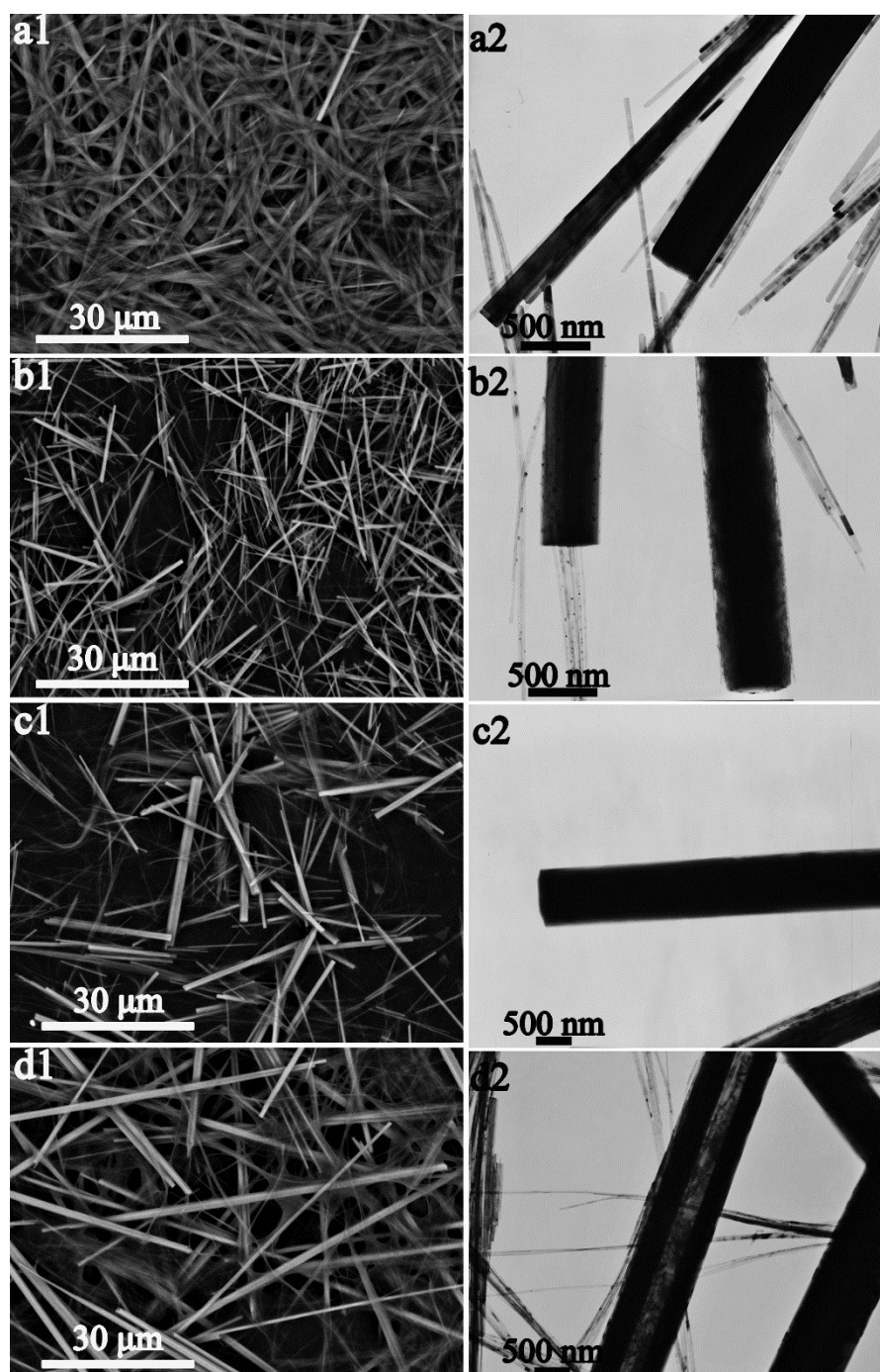


Figure S1. SEM and TEM micrographs of the control samples prepared under the same conditions as sample a2 except for using other phosphorus source. (a1, a2) $\text{Na}_3\text{PO}_4 \cdot 12\text{H}_2\text{O}$; (b1, b2) $\text{Na}_2\text{HPO}_4 \cdot 12\text{H}_2\text{O}$; (c1, c2) $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$; (d1, d2) $(\text{NaPO}_3)_3$.

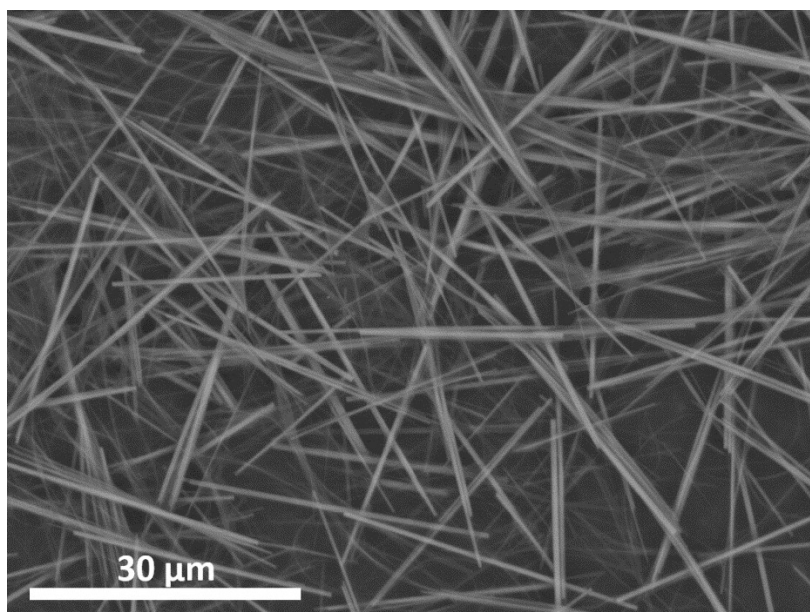


Figure S2. SEM micrograph of the control sample used for drug loading and drug release, which was prepared under the same conditions as sample b3 except for using $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$ (10 mL H_2O , 0.3638 g $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$) instead of $(\text{NaPO}_3)_6$ as the phosphorus source.

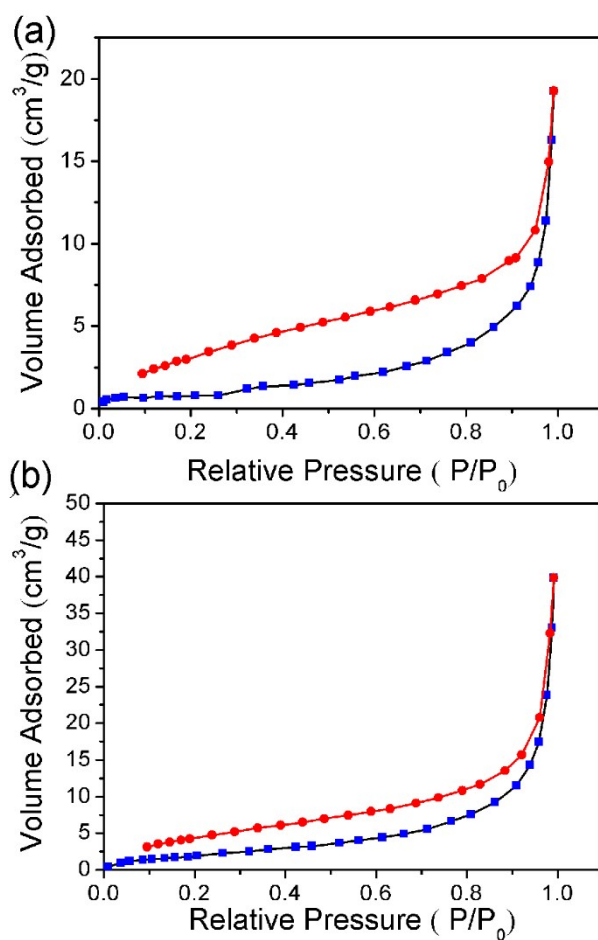


Figure S3. (a) Nitrogen adsorption–desorption isotherms of HAP microtubes (sample b3). (b) Nitrogen adsorption–desorption isotherms of the control sample synthesized under the same conditions as HAP microtubes (sample b3) except for using $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$ (10 mL H_2O , 0.3638 g $\text{NaH}_2\text{PO}_4 \cdot 2\text{H}_2\text{O}$) instead of $(\text{NaPO}_3)_6$ as the phosphorus source.