## **Electronic supplementary information**

## Hydrothermal Synthesis of Titanate Nanotubes from TiO<sub>2</sub> Nanorods Prepared via a Molten Salt Flux Method as an Effective Adsorbent for Strontium Ion Recovery

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## Table S1

Particle size and powder yield of nanomaterials.

Samples	Nanorod s	TNTs- 1D	TNTs- 2D	TNTs- 3D
<sup>a</sup> Nano powder yield (in gm)	0.920	0.506	0.560	0.610
<sup>b</sup> External diameter (or width) of single particle (in nm)	160–300	20–35	16–25	30–60

<sup>a</sup> Nanopowder yield after washing and drying in an oven overnight (g).

<sup>b</sup> Nanomaterial external diameter or width of a particle, based on FESEM images (nm).

## **Figure captions**

**Fig. S1** Effect of the HT reaction time on the formation of nanotubes from nanorods (in 10 M NaOH at 160°C), FESEM images of (a) molten salt NRs, (b) 6 h and (c) 12 h samples; nanotubes formed on the surfaces of the NRs, (d) 24 h (1D); most NRs had transformed into nanotubes, (e) 48 h (2D); NRs completely disappeared, (f) 72 h (3D); nanotubes folded together due to a prolonged reaction, revealing solid, rod-like structures.

**Fig. S2** FE-SEM images of admixture nanoparticles (both untransformed NRs and TNTs) following the reaction over (a) 6 h, (b) 12 h, (c) 1D, (d) 2D, and (e) 3D at 160°C.

Fig. S3 (A) BJH pore size distribution, and (B) nitrogen adsorption and desorption isotherms obtained from the  $TiO_2$  nanorods.

**Fig. S4** Effect of mixed cations  $(Sr^{2+}/Ca^{2+})$  solution over strontium uptake by TNTs-2D. The varying Ca<sup>2+</sup> concentrations are (0, 100, 200, 300, 400 mg L<sup>-1</sup>) and fixed Sr<sup>2+</sup> (10 mg L<sup>-1</sup>) with 30 min contact time. Inset image: Equimolar Sr (mmol/g) and Ca (mmol/g) adsorption curve for TNT-2D nanomaterial adsorbents in seawater.

**Fig. S5** Effect of calcination on the TNT-2D adsorbents over time, and  $Sr^{2+}$  adsorption as a function of the  $Sr^{2+}$  concentration.

Fig. S6 Comparison of the  $k^2$ -weighted Fourier transform of the theoretical scattering patterns from the Ti-Na and Ti-Sr<sup>2+</sup> interactions in the titanate structure.



Fig. S1





Fig. S3



Fig. S4



Fig. S5



Fig. S6