## SUPPLEMENTARY INFORMATION

## A safe, convenient liquid phase pre-sodiation method for titanium-based SIB materials

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Materials synthesis:

 $Na_2Ti_6O_{13}$  were synthesized via a hydrothermal method. 20 ml 2.5 M sodium hydroxide was slowly dropped in a mixture solution of 0.68 g tetrabutyl titanate and 20 ml ethylene glycol and stirred for 1.5 h. Then the solution was transferred in a Teflon vessel and placed in autoclave at 200 °C for 10 h. The sample was washed and calcinated at 400 °C for 5 h.

 $Na_2Ti_3O_7$  was synthesized through a simple solid-state reaction. The precursors of 6 mmol TiO<sub>2</sub> (P25, Acros) and 0.222 g anhydrous  $Na_2CO_3$  (>99.995%, Aldrich) was mixtured with 10 g NaCl. These mixtures were calcinated at 800 °C for 24 h. Then the as-prepared sample was washed with deionized water and dried at 80 °C.

TiO2: P25, purchased from Acros.

Figure S1. The chemical process of the dissolution of sodium metal and its sodiation mechanism.



**Figure S2.** (a) Cycle performance at a current of 100 mAg<sup>-1</sup>. (b) Rate discharge capabilities of the pristine and pre-sodiated electrode at the current density of 100, 200, 400, 600, 800, 1000 mAg<sup>-1</sup> respectively. (c) Galvanostatic discharge and charge profiles of full cell before and after pre-sodiation. (d) Cycle performance of full cell before and after pre-sodiation with 4M Na-Naph-DME solution.



Figure S3. XRD pattern and crystal structure of as prepared  $TiO_2$  (a, c) and  $Na_2Ti_3O_7$  (b, d) powder. The first galvanostatic discharge and charge profiles of  $TiO_2$  (e) and  $Na_2Ti_6O_{13}$  (f) before and after pre-sodiation.

