

Supporting Information for

Controlled Growth of SnO₂@Fe₂O₃ Double-sided Nanocombs as Anodes for Lithium-Ion Batteries

Weiwei Zhou,^a Yee Yan Tay,^b Xingtao Jia,^a Denis Yu Yau Wai,^c Jian Jiang,^a Huey Hoon Hng,^b and Ting Yu^{a,c,d,*}

^a *Division of Physics and Applied Physics, School of Physical and Mathematical Sciences, Nanyang Technological University, 21 Nanyang Link, 637371 Singapore.*

^b *School of Materials Science and Engineering, Nanyang Technological University, 639798 Singapore.*

^c *Energy Research Institute @ NTU (ERIAN), 50 Nanyang Drive, 637553 Singapore.*

^d *Department of Physics, Faculty of Science, National University of Singapore, 117576 Singapore .*

* Address correspondence to yuting@ntu.edu.sg

Experimental

Synthesis: The SnO₂@Fe₂O₃ double-sided nanocombs were prepared by a simple two-step process, which can be easily scaled up. The Fe₂O₃ nanoflakes were firstly synthesized by heating the polished (by sand paper) and cleaned (with distilled water) Fe foil (Aldrich, 0.1 mm thick, 99.9+ %) on a hotplate at 400 °C for 24 h in air. Then, a piece of as-obtained Fe foil covered by Fe₂O₃ nanoflakes was placed in a 50 mL sealed Teflon autoclave filling with 30 mL aqueous solution containing 0.174 g of SnCl₄ · 2H₂O and 0.534 g of NaOH. After growing at 220 °C for 3 h, the substrate was removed from the solution, rinsed with distilled water, and dried with N₂ blow.

Material Characterization: Samples were characterized by powder X-ray diffraction (XRD) (Bruker D-8 Avance), scanning electron microscopy (SEM) (JSM-6700F, 10.0 kV), transmission electron microscopy (TEM) (JEM-2010, 200 kV).

Electrochemical Characterization: Electrochemical measurements were performed using two-electrode CR2016 coin-type cells with lithium foil serving as both counter and reference electrodes at room temperature. The SnO₂@Fe₂O₃ nanocombs on Fe foil were directly used as working electrode. Besides, the Fe₂O₃ nanoflakes on Fe foil were also tested for comparison. The electrolyte used was 1.0 M LiPF₆ in a mixture of ethylene carbonate and diethyl carbonate (1:1 by volume). Cell assembly was carried out in an argon-filled glovebox with both moisture and oxygen contents below 0.1 ppm. Galvanostatic charge/discharge tests were performed using a NEWARE battery tester at a voltage window of 0.005–3 V. Cyclic voltammetry (CV, 0.005–3 V, 0.5 mV s⁻¹) and Impedance measurement (0.1-30000 Hz) were performed on the as-made cells using an electrochemical workstation (CHI 760D).

Supplementary Figures

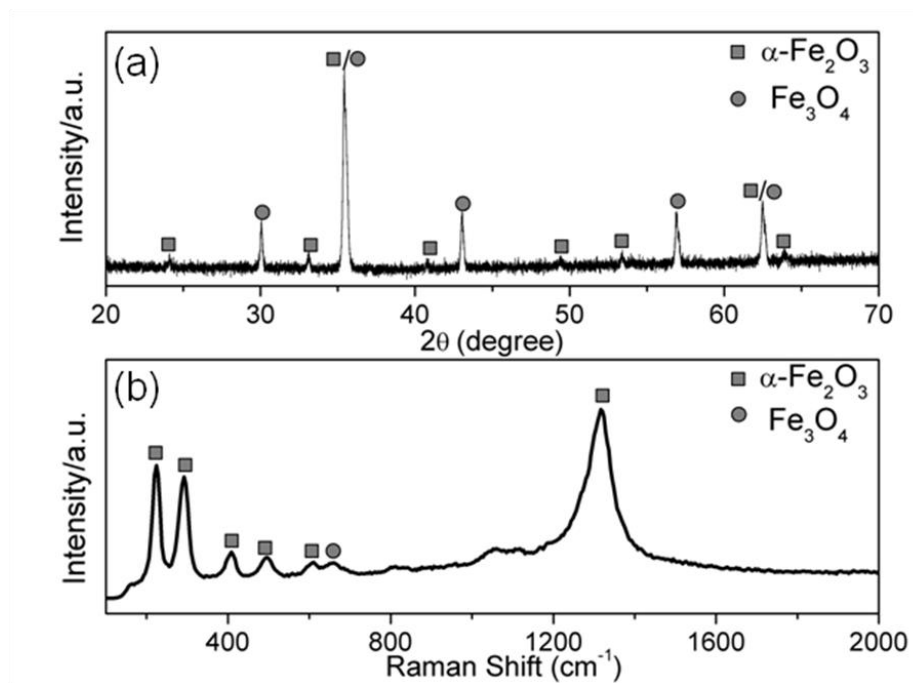


Fig. S1. (a) XRD pattern and (b) Raman spectrum of the as-obtained Fe_2O_3 nanoflakes.

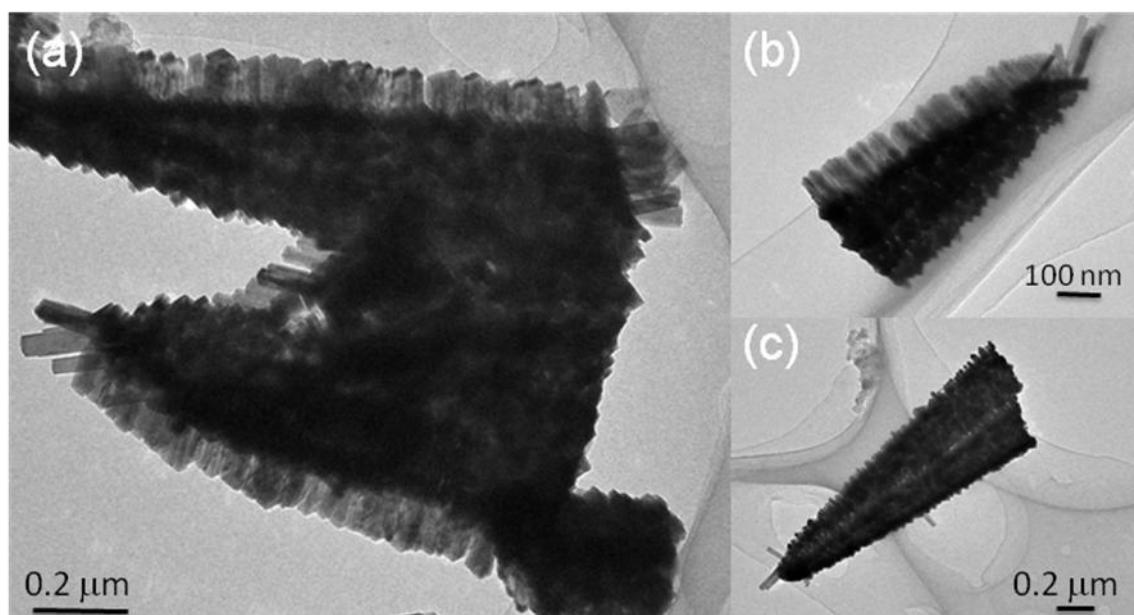


Fig. S2. TEM images of $\text{SnO}_2@\text{Fe}_2\text{O}_3$ nanocombs after being sonicated for 0.5 h, suggesting the strong adherent forces between the SnO_2 branches and Fe_2O_3 cores.

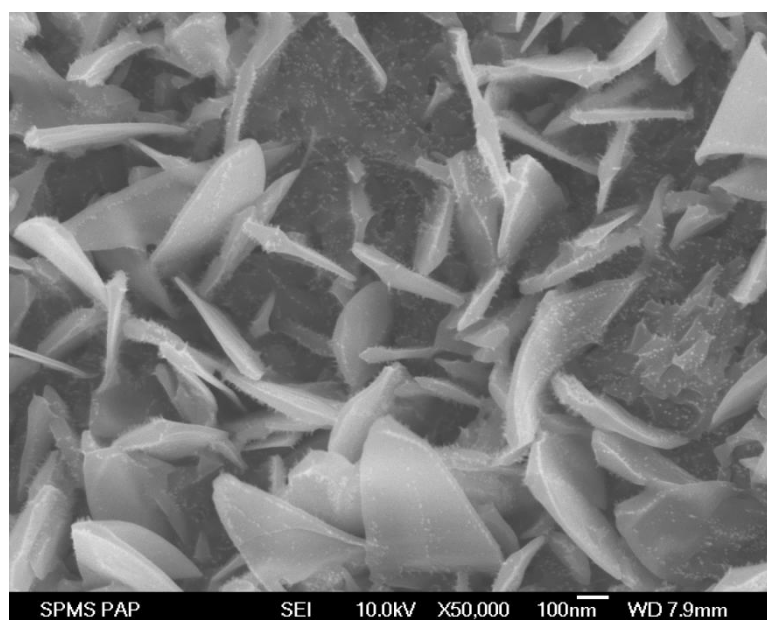


Fig. S3. SEM image of the product gained after 2 h of hydrothermal growth of SnO_2 branches, showing the initial nuclei of SnO_2 .

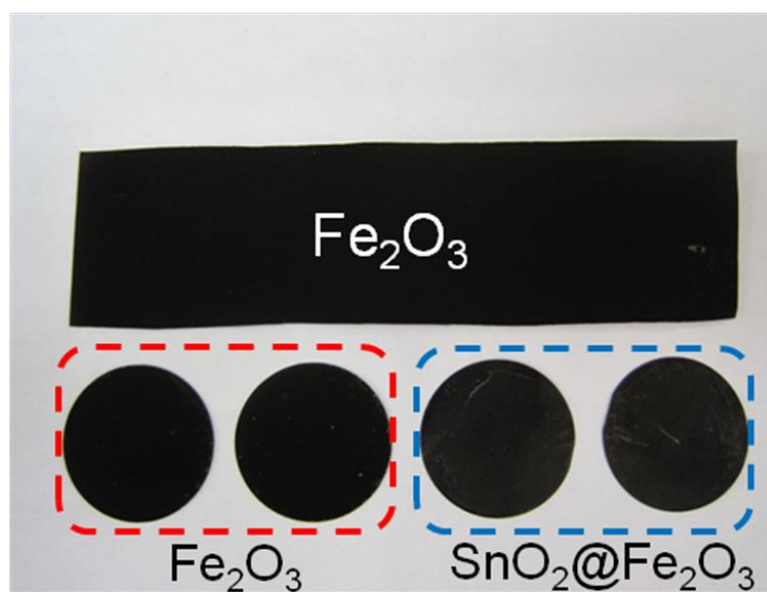


Fig. S4. Optical image of our samples, showing that the working electrodes, namely Fe_2O_3 and $\text{SnO}_2@\text{Fe}_2\text{O}_3$, can be simply obtained by cutting the as-gained products into platelets with certain sizes.