## Supplementary Information

Controllable formation of multilayered SnO<sub>2</sub>@Fe<sub>2</sub>O<sub>3</sub> sandwich cubes as high-performance anode for Li-ion batteries

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**Figure S1.** XRD patterns of the as-obtained products after (a) fast precipitation, (b) anneal, and (c) etching steps, respectively.



Figure S2. Digital photographs of (a) the pristine  $SnO_2$  HNCs after the etching process and (b) final  $SnO_2@Fe_2O_3$  SNCs.



**Figure S3.** (a) Low-magnification FESEM image of the as-obtained ZHS singleshelled hollow precursor. (b-e) EDX mapping images of an individual ZHS singleshelled hollow cube. The inset of (a) is a typical SAED pattern taken from a ZHS cube.



**Figure S4.** (a) Low-magnification FESEM and (c) TEM images of  $Zn_2SnO_4/SnO_2$ double-shelled hollow cubes, (b) enlarged FESEM image of a broken  $Zn_2SnO_4/SnO_2$ hollow cube, and (d) typical SAED pattern from a  $Zn_2SnO_4/SnO_2$  hollow cube.



Figure S5. (a) FESEM image of single  $SnO_2@Fe_2O_3$  particle, and (b-d) the corresponding elemental mapping showing the dispersion of Sn, O, and Fe elements in  $SnO_2@Fe_2O_3$  particle.



Figure S6. XPS survey spectra of the as-prepared  $SnO_2$  HNCs and  $SnO_2@Fe_2O_3$ SNCs. The upper inset is the typical high-resolution C 1s XPS spectra of  $SnO_2$  HNCs.



**Figure S7.** XRD patterns of the as-prepared samples without hydrothermal process (a) and within the time-dependent hydrothermal process of (b) 0.5 h, (c) 1.0 h, and (d) 1.5 h, respectively.



**Figure S8.** FESEM images of (a,b) the as-prepared  $SnO_2$  HNCs without hydrothermal process and (c,d)  $SnO_2@Fe_2O_3$  SNCs after the  $Fe_2O_3$ -loaded hydrothermal process.



Figure S9. The following figure shows the CV curves of  $SnO_2$  HNCs electrode. There exists a large cathodic peak at 0.72 V at the first scan, which can be attributed to the typical voltage trend of  $SnO_2$  during the lithium intercalation, as shown in the following irreversible reaction (1). The discriminable anodic peaks centered at 0.57 V and 1.22 V could be identified for the lithium extraction in Sn materials, as shown in the reaction (2). The obvious difference between the first and subsequent cycles could be attributed to the irreversible reaction and the formation of SEI layer. In addition, it is clearly discerned after the first cycle that the CV curves with peak intensity and integrated area during the reductive and oxidative polarization processes are clearly reduced, indicating an unsatisfied reversibility of lithium insertion/extraction reaction. The main electrochemical processes of SnO<sub>2</sub> electrode can be described as follows [1-3].

$$SnO_2 + 4Li^+ + 4e^- \rightarrow Sn + 2Li_2O \tag{1}$$

$$Sn + xLi^{+} + xe^{-} \leftrightarrow Li_{x}Sn \ (0 \leq x \leq 4.4)$$
(2)



Figure S10. Galvanostatic discharge-charge voltage profiles of  $SnO_2$  HNCs for the 1st, 2nd and 10th cycles at a current density of 100 mA g<sup>-1</sup>.



Figure S11. Nyquist plots of  $SnO_2$  HNC and  $SnO_2@Fe_2O_3$  SNC electrodes after 5 cycles.



Figure S12. TEM and FESEM images of (a,c)  $SnO_2$  HNC and (b,d) $SnO_2@Fe_2O_3$ 

SNC electrodes after 20 cycles at a current density of 100 mA  $g^{-1}$ .

## Notes and references

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