

Supporting Information

Partially Phosphorized SnO₂/Graphene Nanocomposite for Highly Reversible Pseudocapacitive Lithium-Ion Storage

Ying Yang,^{a, b} Xu Zhao,^b Hong-En Wang,^b Malin Li,^b Ce Hao,^a Min Ji,^a Suzhen Ren,^{* a} Guozhong Cao^{*}

^{a, b}

^a College of Chemistry, Dalian University of Technology, Dalian 116024, China.

^b Department of Materials Science and Engineering, University of Washington, Seattle, WA 98195, USA

*Corresponding author: rensz@dlut.edu.cn; gzcao@u.washington.edu

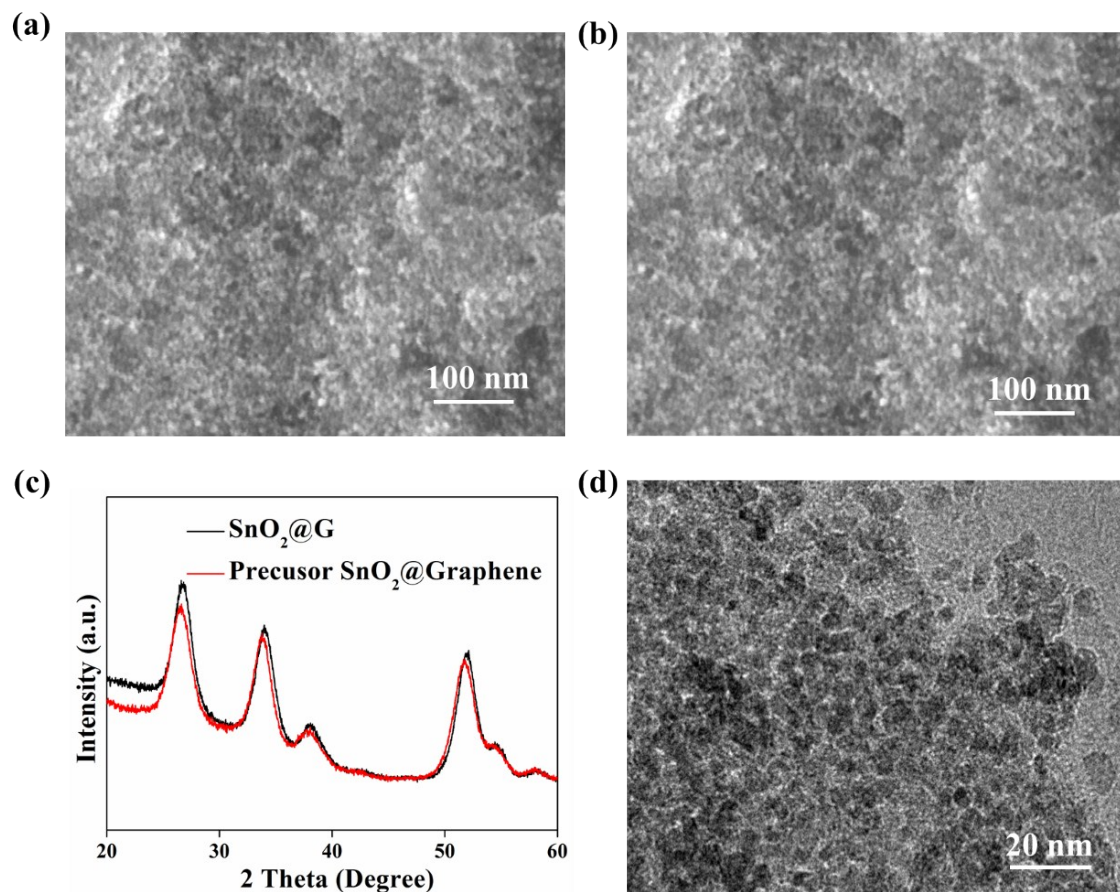


Figure S1. Scanning electron microscope (SEM) images of (a) SnO₂@G-Pre composite and (b) SnO₂@G composite after annealing; (c) XRD curves of SnO₂@G composite before and after annealing; (d) transmission electron microscope image (TEM) of SnO₂@G after annealing.

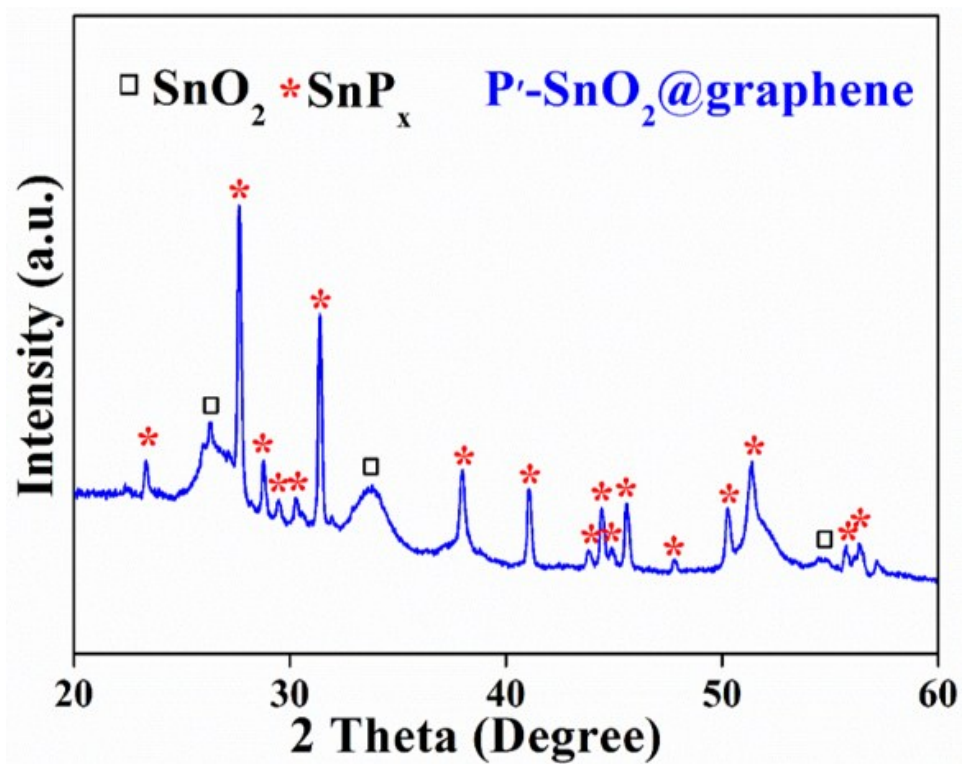


Figure S2. XRD pattern of P-SnO₂@G-10 phosphorized at low molar ratio of Sn and P (1:10).

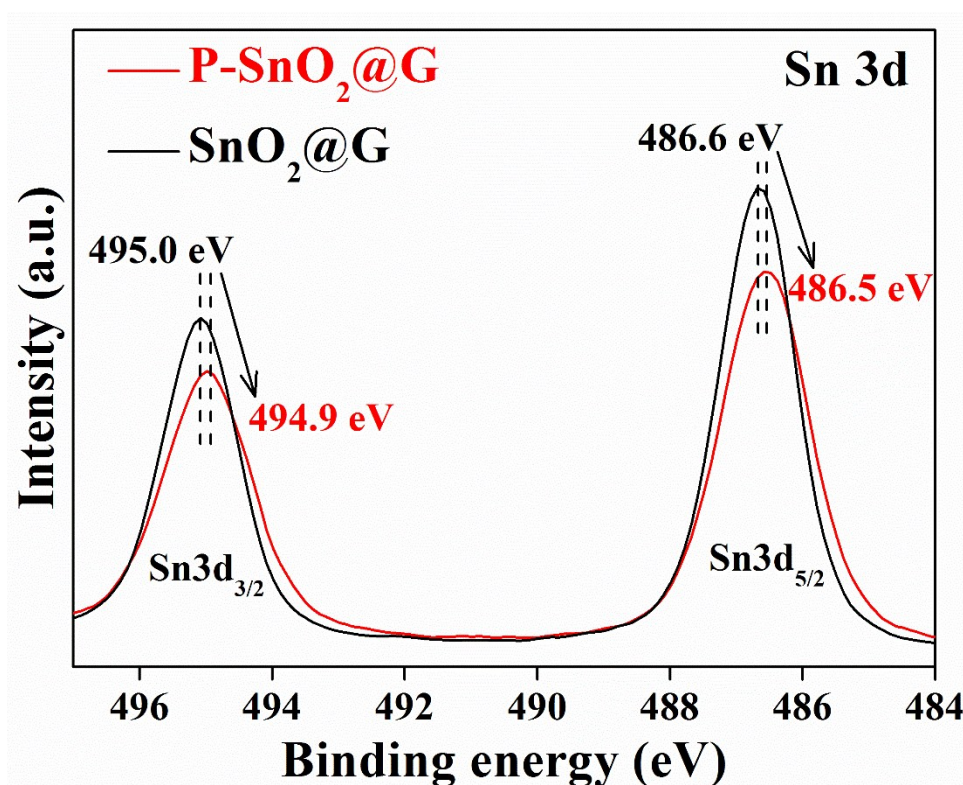


Figure S3. High-resolution XPS spectra of Sn 3d in SnO₂@G and P-SnO₂@G.

Table S1. The element composition of P-SnO₂@G estimated based on Energy dispersive spectrometer (EDS).

Element	Weight%	Atomic%
C	11.52	35.88
N	2.15	1.14
O	13.31	29.98
P	10.43	12.60
Sn	62.59	20.41

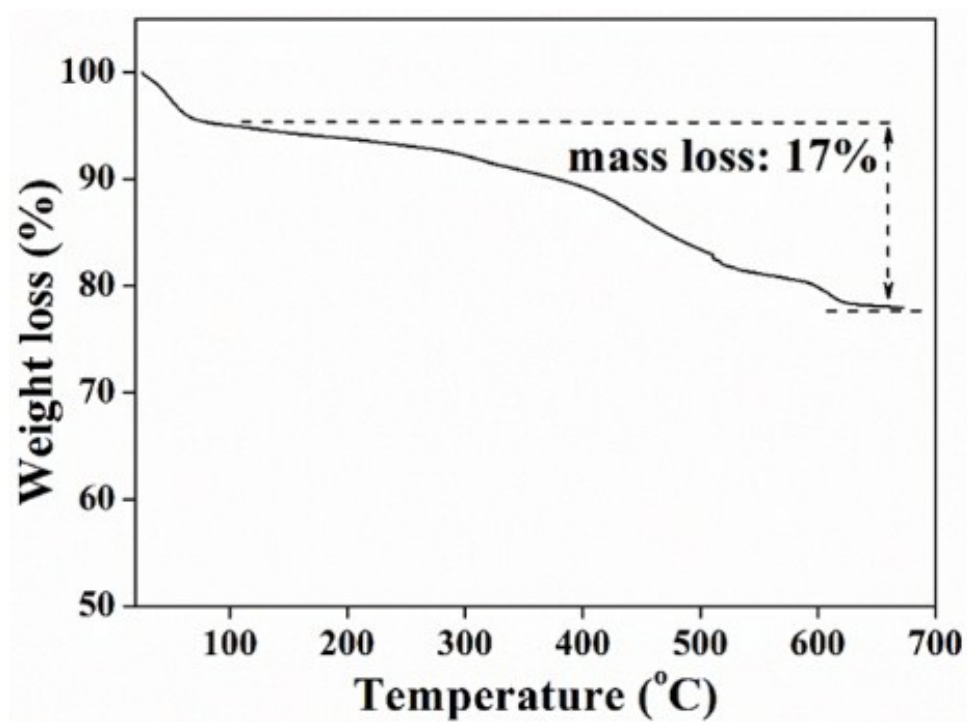


Figure S4. Thermogravimetric analysis curve of as-synthesized $\text{SnO}_2@\text{G-Pre}$ composite

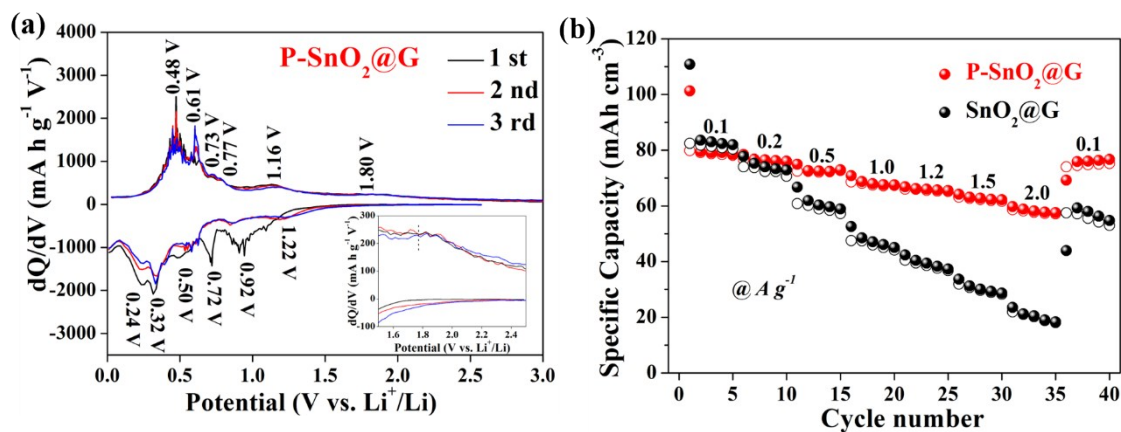


Figure S5. (a) Differential charge - discharge capacity plots of as-synthesized P-SnO₂@G composite electrode at a current density of 0.1 mA g⁻¹ for the first three cycles; (b) Volumetric capability of SnO₂@G and P-SnO₂@G obtained at various current densities from 0.1 to 2 A g⁻¹, then back to 0.1 A g⁻¹ in the potential window of 0.01 ~ 3.0 V.

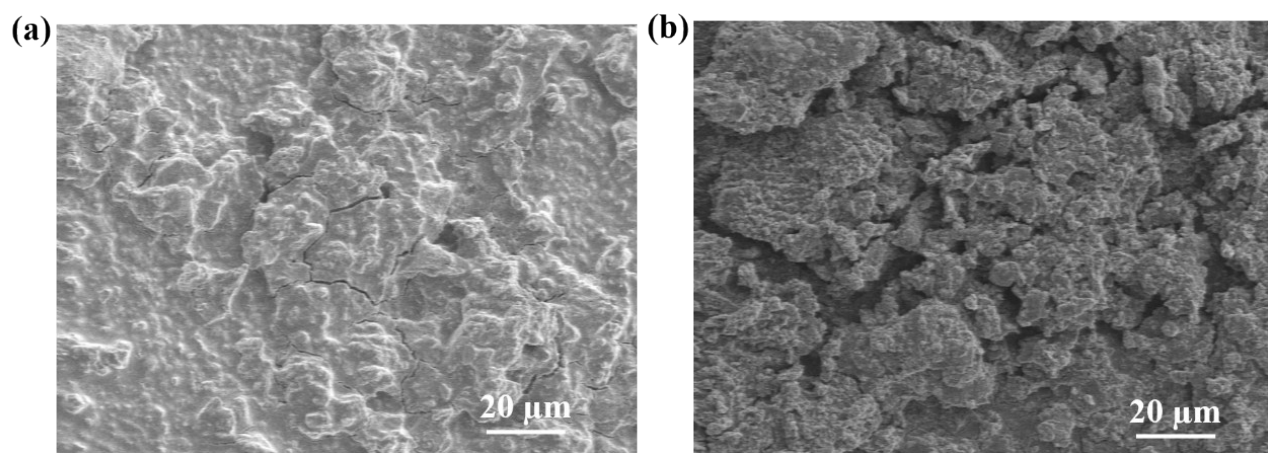


Figure S6. Ex situ SEM images of (a) P-SnO₂@G; (b) SnO₂@G after cycling

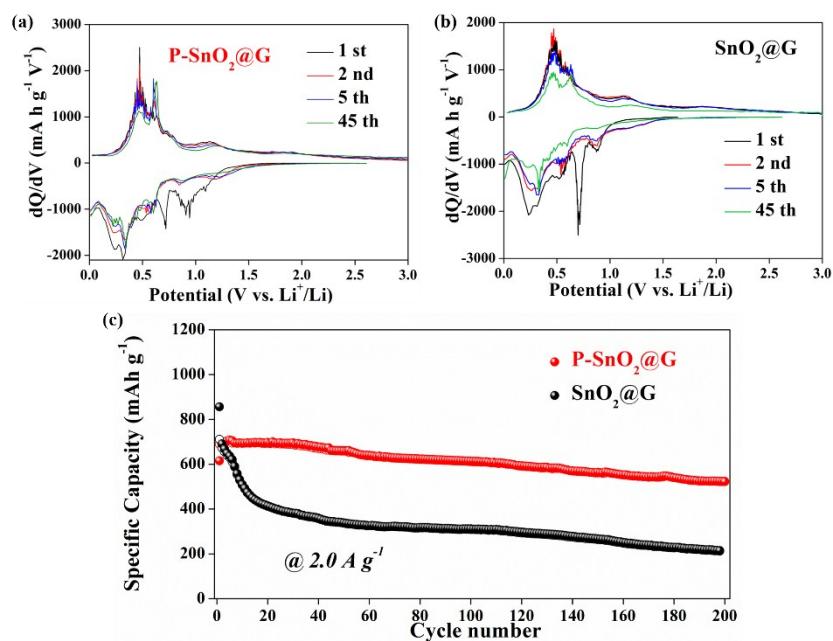


Figure S7. Differential discharge and charge capacity plots (DCPs) for the 1st, 2th, 5th and 45th cycles of (a) SnO₂@G; (b) P-SnO₂@G; (c) Cycling stability (200 cycles) of SnO₂@G and P-SnO₂@G at 2.0 A g⁻¹ in the potential of 0.01 ~ 3.0 V.

Table S2. The R_s , R_f , and R_{ct} and Li^+ diffusion coefficients of $\text{SnO}_2@\text{G}$ and $\text{P-SnO}_2@\text{G}$ electrode after 50 cycles at current density of 1.0 A g^{-1} .

Sample	$\text{SnO}_2@\text{G}$	$\text{P-SnO}_2@\text{G}$
$R_s (\Omega)$	1.43	4.17
$R_f (\Omega)$	47.71	115.80
$R_{ct} (\Omega)$	92.73	157.30
$D_{\text{Li}^+} (\text{cm}^2 \text{ s}^{-1})$	6.50×10^{-14}	1.36×10^{-13}

(Note: R_s represents the combined ohmic resistance of the battery; R_f is resistance of surface film; R_{ct} is the value of the charge transfer resistance, fitted from Figure 5a)

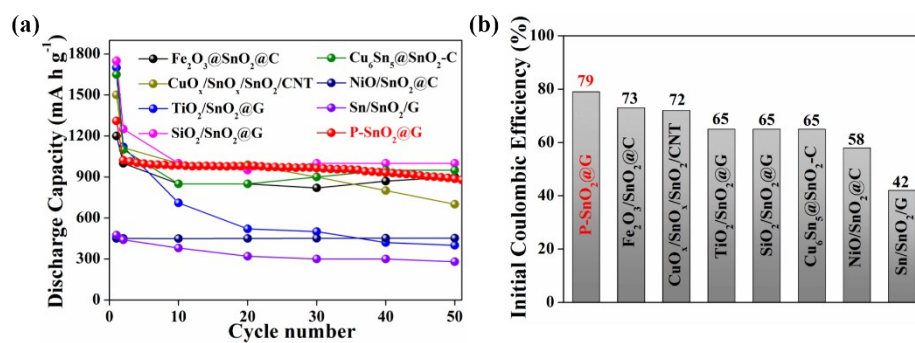


Figure S8. (a) Comparison of capacity retention and (b) initial coulombic efficiency of P-SnO₂@G (0.01-3 V, 0.1 A g⁻¹) with multiple components composite anodes including Fe₂O₃@SnO₂@C (0.01-3 V, 0.2 A g⁻¹), CuO_x/SnO_x/SnO₂/CNT (0.005-2.7 V, 0.1 A g⁻¹), TiO₂/SnO₂/C (0.01-3 V, 0.1 A g⁻¹), SiO₂/SnO₂@G (0.01-2.5 V, 0.5 A g⁻¹), Cu₆Sn₅@SnO₂-C (0.05-3V, 0.2 A g⁻¹), NiO/SnO₂/C (0.01-3 V, 0.8 A g⁻¹), Sn/SnO₂@C (0.05-1.5 V, 0.05 A g⁻¹). References is same with the main article.