

Supporting Information

3D Graphene Network Encapsulating SnO₂ Hollow Spheres for High-performance Anode Material of Lithium-ion Batteries

3D Graphene Network Encapsulating SnO₂ Hollow Spheres for High-performance Anode Material of Lithium-ion Batteries

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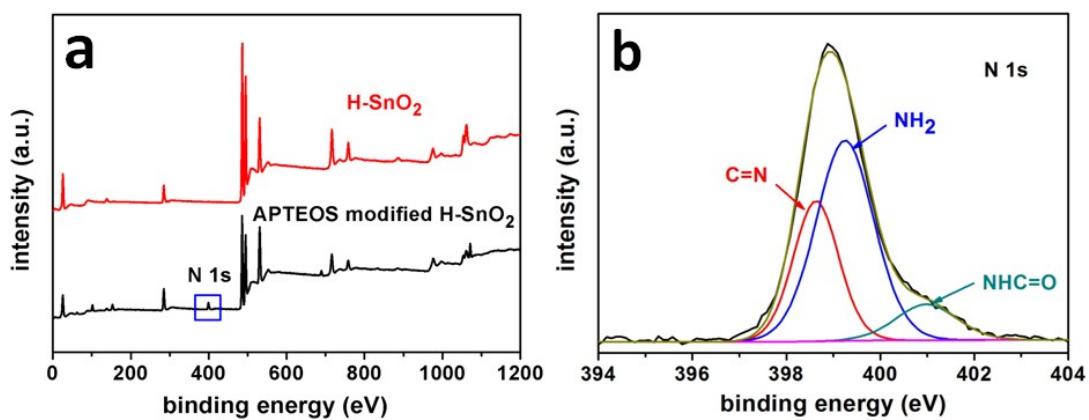


Figure S1. XPS survey spectra (a) of APTEOS modified H-SnO₂ and pure H-SnO₂ samples, and high resolution XPS spectrum of N 1s (b) for the APTEOS modified H-SnO₂ samples. The N 1s peak shows that the primary amines (NH₂, 399.3 eV) and secondary amines (N=C, 398.6 eV) along with oxidized species as amides (NHC=O, 400.9 eV) in the high resolution XPS spectrum. The abundant amino groups can serve as deposition places to coat graphene nanosheets, resulting in graphene fully enwrapped H-SnO₂ nanospheres by electrostatic adsorption.

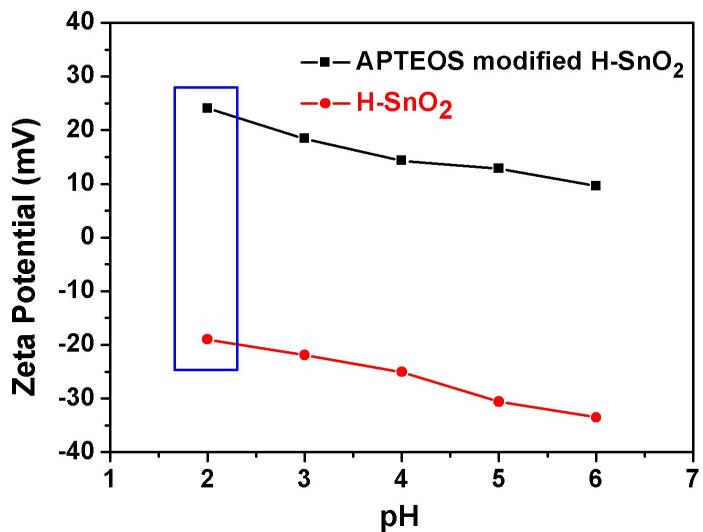


Figure S2. Zeta potentials of APTEOS modified H-SnO₂ (black square) and graphene oxide (red circle) in aqueous solution at different pH values. The blue rectangle indicates that the assembly process can be spontaneous at pH 2 where the maximum electrostatic interactions are achieved between the APTEOS modified H-SnO₂ and graphene oxide.



Figure S3. Photographs of the assembly process of APTEOS modified H-SnO₂ and graphene oxide in aqueous solutions at pH 2.

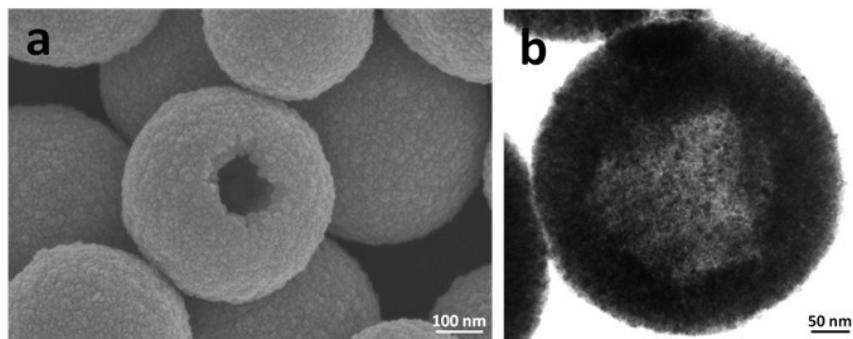


Figure S4. The magnified a) FESEM and b) TEM images of the as-prepared H-SnO₂.

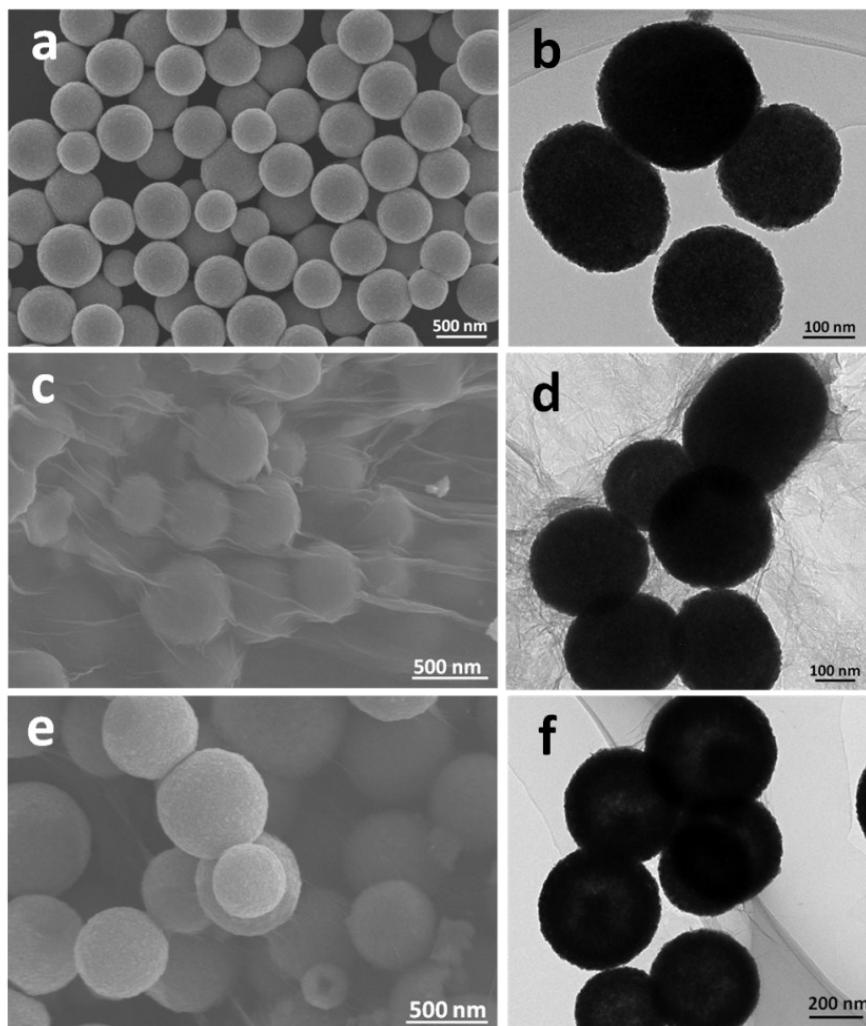


Figure S5. a) FESEM and b)TEM images of the solid SnO₂ nanospheres (S-SnO₂) prepared at 150 °C for 6 h. c) FESEM and d) TEM images of after self-assembled wrapping of interconnected graphene networks (S-SnO₂@rGO). e) FESEM and f)TEM images of graphene loading hollow SnO₂ nanospheres (H-SnO₂/rGO).

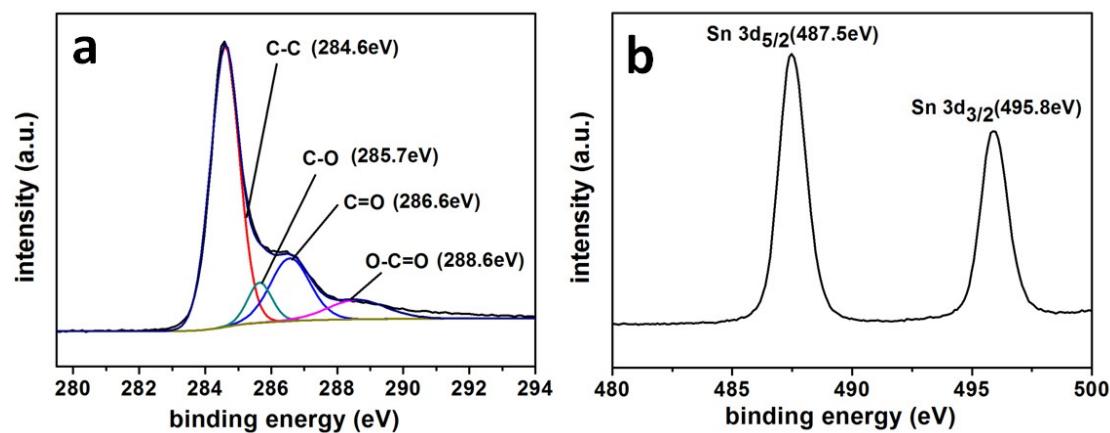


Figure S6. a) XPS of C 1s and b) XPS Sn 3d fine scan spectrum of H-SnO₂@rGO.

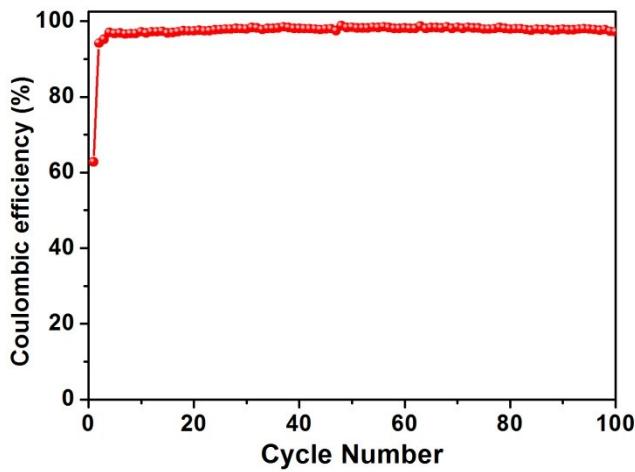


Figure S7. The coulombic efficiency of the H-SnO₂@rGO electrode materials at a current density of 0.1 A g⁻¹.

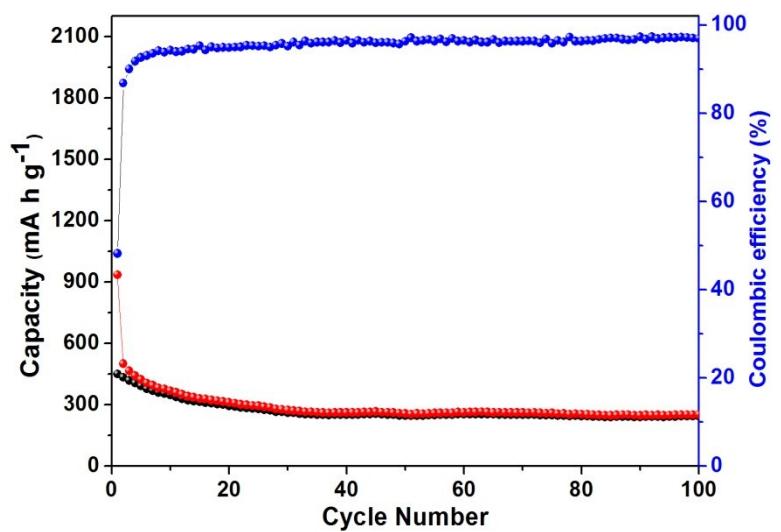


Figure S8. The electrochemical performance of rGO at a current density of 0.1 A g^{-1} .

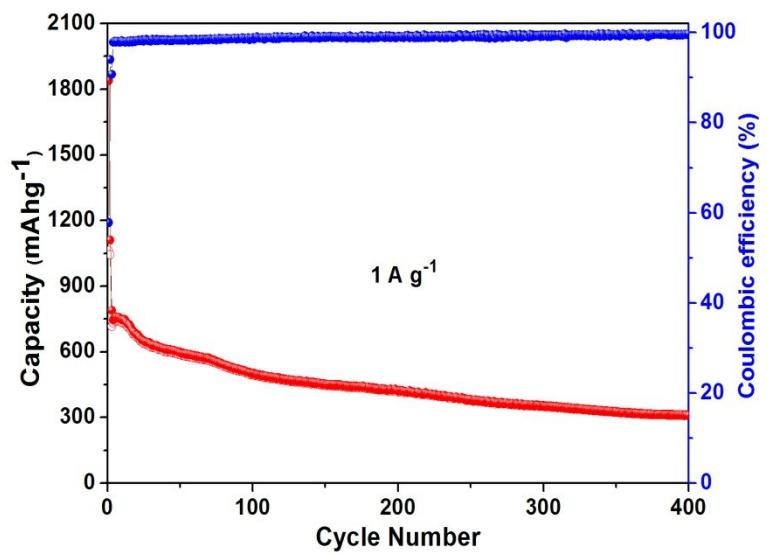


Figure S9. Cycling performance and Coulombic efficiency of S-SnO₂@rGO for 400 cycles at the current density of 1 A g⁻¹.

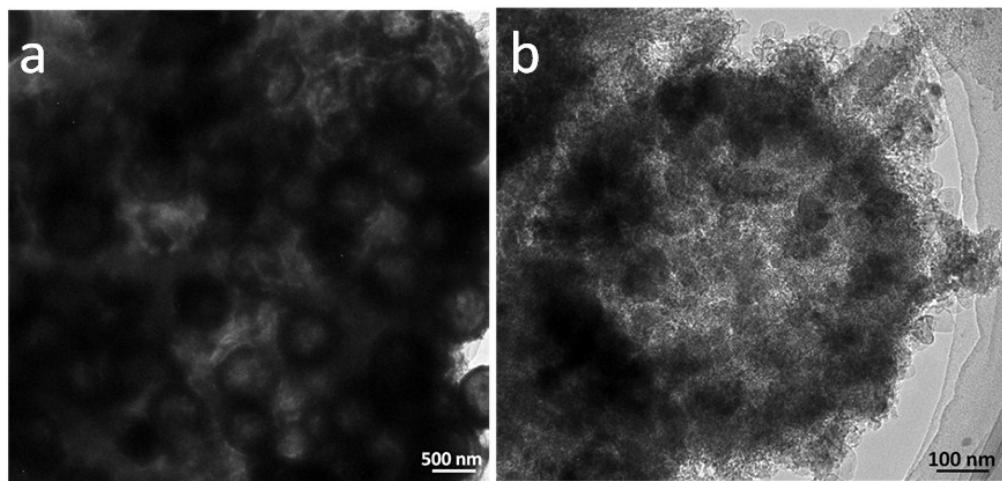


Figure S10. a,b) Typical TEM image of a fully charged H-SnO₂@rGO electrode after 100 cycles at a current density of 100 mA g⁻¹.

Table S1. Electrochemical performance comparison of H-SnO₂@rGO with previously reported graphene-based SnO₂ composites with different morphologies or compositions.

Materials	Voltage range(V)	Current density (mA g ⁻¹)	Cycle number	Specific capacity (mAh g ⁻¹)	Reference
H-SnO ₂ @rGO	0.01-3.0	100	100	1107	Our Work
		1000	500	552	
S-SnO ₂ @rGO	0.01-3.0	100	100	744	Our Work
Graphene-based mesoporous SnO ₂	0.01-3.0	78	50	848	1
SnO ₂ /GNS	0.005-2.0	50	30	570	2
3D SnO ₂ /graphene	0.01-3.0	200	50	845	3
Dually fixed SnO ₂ /G@Pani	0.01-3.0	100	100	770	4
SnO ₂ -GO hybrid	0.005-2.5	100	200	800	5
Graphene nanoribbons/SnO ₂	0.01-2.5	100	50	825	6
3D-G/SnO ₂ @C	0.005-3	100	100	820	7
N-doped G-SnO ₂ Sandwich Papers	0.005-3	50	50	910	8
SnO ₂ Quantum Dots@GO	0.01-3	100	100	1121	9
SnO ₂ /graphene	0.01-2.5	100	200	830	10

SnO ₂ @C@GS	0.01-2	200	100	830	11
SnO ₂ -HNS/G	0.005-3	500	300	696	12
Polyaniline @SnO ₂ @Graphene	0.01-3	1000	100	560	13
SnO ₂ nanosheets @graphene sheets	0.01-1.2	160	50	518	14
SnO ₂ /RGO/C foam	0.01-1.5	130	100	717	15
SnO ₂ quantum dots/RGO	0.01-3	100	200	924	16
rGO/SnO ₂	0.01-3	100	100	536	17
polydopamine- coated RGO/SnO ₂	0.01-2	100	200	718	18
SnO ₂ @G@G	0.01-2	80	120	591	19
graphene/C-SnO ₂	0.005-3	100	50	502	20

References

1. S. Yang, W. Yue, J. Zhu, Y. Ren and X. Yang, *Advanced Functional Materials*, 2013, 23, 3570-3576.
2. S. M. Paek, E. Yoo and I. Honma, *Nano letters*, 2009, 9, 72-75.
3. R. Tian, Y. Zhang, Z. Chen, H. Duan, B. Xu, Y. Guo, H. Kang, H. Li and H. Liu, *Scientific reports*, 2016, 6, 19195.
4. Y. Dong, Z. Zhao, Z. Wang, Y. Liu, X. Wang and J. Qiu, *ACS applied materials & interfaces*, 2015, 7, 2444-2451.
5. H. Song, N. Li, H. Cui and C. Wang, *Journal of Materials Chemistry A*, 2013, 1, 7558.
6. J. Lin, Z. Peng, C. Xiang, G. Ruan, Z. Yan, D. Natelson and J. M. Tour, *ACS nano*, 2013, 7, 6001-6006.
7. B. Luo, T. Qiu, L. Hao, B. Wang, M. Jin, X. Li and L. Zhi, *J. Mater. Chem. A*, 2016, 4, 362-367.

8. X. Wang, X. Cao, L. Bourgeois, H. Guan, S. Chen, Y. Zhong, D.-M. Tang, H. Li, T. Zhai, L. Li, Y. Bando and D. Golberg, *Advanced Functional Materials*, 2012, 22, 2682-2690.
9. K. Zhao, L. Zhang, R. Xia, Y. Dong, W. Xu, C. Niu, L. He, M. Yan, L. Qu and L. Mai, *Small*, 2016, 12, 588-594.
10. S. J. Prabakar, Y. H. Hwang, E. G. Bae, S. Shim, D. Kim, M. S. Lah, K. S. Sohn and M. Pyo, *Adv Mater*, 2013, 25, 3307-3312.
11. X. Zhou, W. Liu, X. Yu, Y. Liu, Y. Fang, S. Klankowski, Y. Yang, J. E. Brown and J. Li, *ACS applied materials & interfaces*, 2014, 6, 7434-7443.
12. X. S. Zhou, Y. X. Yin, L. J. Wan and Y. G. Guo, *Journal of Materials Chemistry*, 2012, 22, 17456-17459.
13. F. Ye, B. Zhao, R. Ran and Z. Shao, *Journal of Power Sources*, 2015, 290, 61-70.
14. S. Ding, D. Luan, F. Y. C. Boey, J. S. Chen and X. W. Lou, *Chemical communications*, 2011, 47, 7155.
15. H. Tao, S. Zhu, L. Xiong, X. Yang and L. Zhang, *ChemElectroChem*, 2016, 3, 1063-1071.
16. C. Zhu, S. Zhu, K. Zhang, Z. Hui, H. Pan, Z. Chen, Y. Li, D. Zhang and D. W. Wang, *Scientific reports*, 2016, 6, 25829.
17. D. Zhou, W.-L. Song, X. Li and L.-Z. Fan, *Electrochimica Acta*, 2016, 207, 9-15.
18. L. Wang, D. Wang, Z. Dong, F. Zhang and J. Jin, *Nano letters*, 2013, 13, 1711-1716.
19. J. Zhu, G. Zhang, X. Yu, Q. Li, B. Lu and Z. Xu, *Nano Energy*, 2014, 3, 80-87.
20. M. Shahid, N. Yesibolati, M. C. Reuter, F. M. Ross and H. N. Alshareef, *Journal of Power Sources*, 2014, 263, 239-245.