## Electronic Supplementary Information

## In-situ preparation of uniform SnO<sub>2</sub> nanocrystals anchored

## within mesoporous carbon network as advanced anode materials

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Fig. S1. a) Photograph of seaweeds. b) Chemical Structure of sodium alginate.



**Fig. S2.** SEM images of the SnO<sub>2</sub>@SAMC composite with different magnifications: a) 2 um. b) 400 nm, c) 200 nm, d) 100 nm.



Fig. S3. Higher magnification TEM images of  $SnO_2@SAMC$  composite before calcination.



**Fig. S4.** a) Thermogravimetric analysis of sodium alginate and glucose under nitrogen atmosphere. Heating rate:  $10^{\circ}$ C min<sup>-1</sup>. b) N<sub>2</sub> adsorption and desorption isotherms of SnO<sub>2</sub>@GC composite, the inset shows the pore-size distribution.



Fig. S5. Raman spectra for SnO<sub>2</sub>@SAMC and SnO<sub>2</sub>@GC composites.

Materials	Methods	Reactants	Morphology	Particle Sizes	References
Sn/SnO <sub>2</sub> particles within mesoporous carbon	hard-template and liquid impregnation	mesoporous SBA-15; Pluronic P123; sucrose; HF; H <sub>2</sub> SO <sub>4</sub> ; SnCl <sub>2</sub> ·2H <sub>2</sub> O; etc.	702) 0.226m (101)	not reported	1
SnO <sub>2</sub> particles within micro/mesoporous carbon	soft-template and liquid impregnation	phloroglucinol; Pluronic F127; organic solvent; HCl; H <sub>2</sub> O <sub>2</sub> ; SnCl <sub>4</sub> .5H <sub>2</sub> O; etc.		0.8-4 nm	2
SnO <sub>2</sub> particles within N-doped graphene sheets	freeze-drying and vapor reduction	SnCl <sub>4</sub> · 5H <sub>2</sub> O; graphite oxide and hydrazine monohydrate.	SnO <sub>2</sub> (110) N-RGO	4-5 nm	3
Polydopamine-coated SnO <sub>2</sub> particles	ATRP, hydrolysis, hydrothermal reaction and polydopamine coating;	Hydroxypropyl cellulose, Na <sub>2</sub> SnO <sub>3</sub> ·3H <sub>2</sub> O, dopamine hydrochloride, etc.	PDA coating ↑ 0 0 10 nm	average size of approximately 5 nm	4
SnO <sub>2</sub> @SAMC	ion exchange, hydrothermal and calcination	sodium alginate and SnCl <sub>2</sub> ·2H <sub>2</sub> O	Carton metwork	2.2-3.8 nm	this work

**Table S1.** Comparison of synthesizing methods for confinement of 0D crystalline  $SnO_2$  particles within carbonaceous materials



**Fig. S6.** TEM images of  $SnO_2@GC$  composite: a, b) Freshly prepared electrode before cycling with different magnification, and c, d) different magnification after cycling at 200 mA g<sup>-1</sup> for 200 cycles.



Fig. S7. a and b) TEM images of  $SnO_2$  sheets, inset of b) shows the SAED. c) XRD pattern for  $SnO_2$  sheets, d) Cycle performance at specific current of 200 mA g<sup>-1</sup>.



**Fig. S8.** SEM images of  $SnO_2@SAMC$  composite: a, b) Freshly prepared electrode before cycling with different magnification, and c, d) different magnification after cycling at 200 mA g<sup>-1</sup> for 300 cycles. TEM images of  $SnO_2@SAMC$  composite: e) Freshly prepared electrode before cycling, and f) after cycling at 200 mA g<sup>-1</sup> for 300 cycles.

## References

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