

## Supporting information

### An oxygen doped Porous hard carbon derived from durian shell for high-performance sodium ion storage

Jianliang Guo,<sup>†,‡</sup> Zhihua Sun,<sup>‡</sup> Wenzheng Zhu,<sup>‡</sup> Lei Li,<sup>‡</sup> Song Han,<sup>†,\*</sup> Hongxun Yang,<sup>‡,\*</sup>

<sup>†</sup>*National Energy Group Science and Technology Research Institute Co., Ltd., Nanjing 210008, Jiangsu, China;* <sup>‡</sup>*School of Environmental & Chemical Engineering, Jiangsu University of Science and Technology, Zhenjiang 212003, Jiangsu, China*

\*Corresponding author:

E-mail: yhongxun@126.com;

F-mail: 12068942@ceic.com

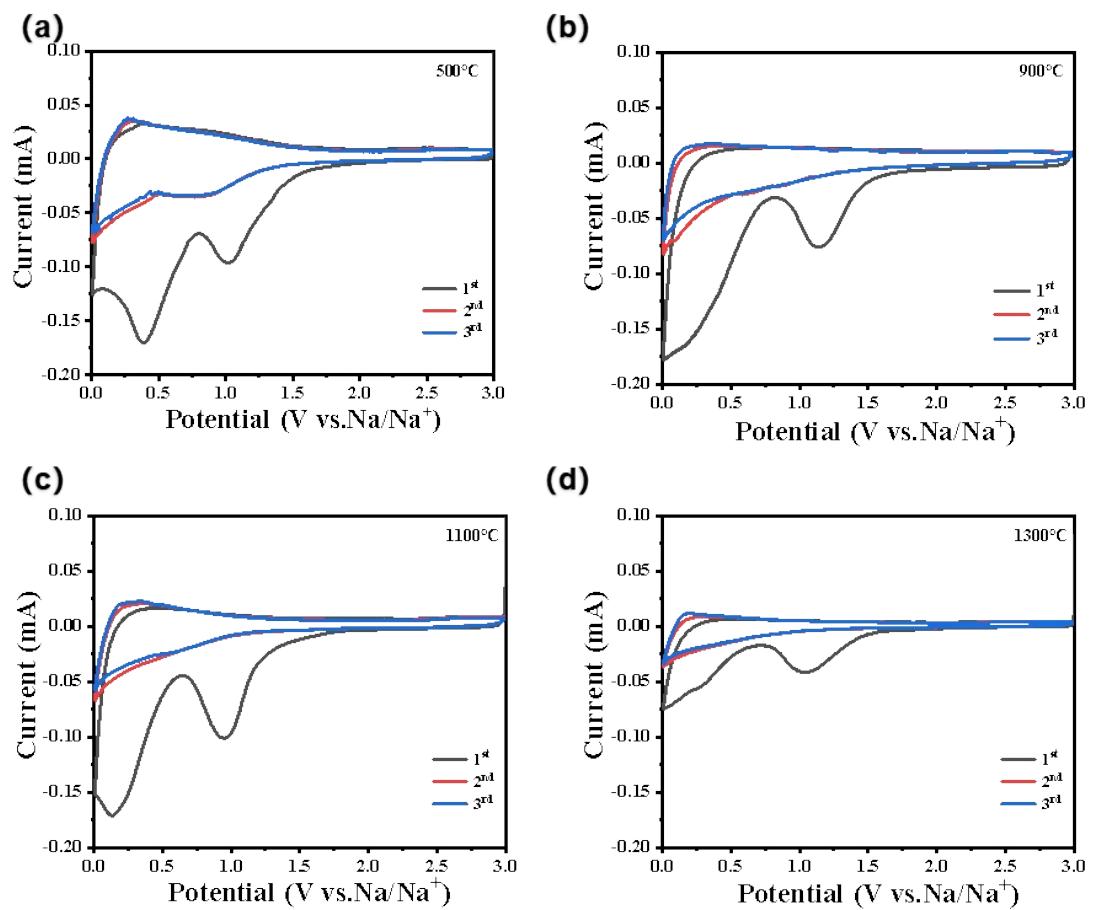


Fig. S1 (a-d) The CV curves of the first four electrodes of DSHC500, DSHC900, DSHC1100 and DSHC1300 at a scanning rate of  $0.1 \text{ mV s}^{-1}$ .

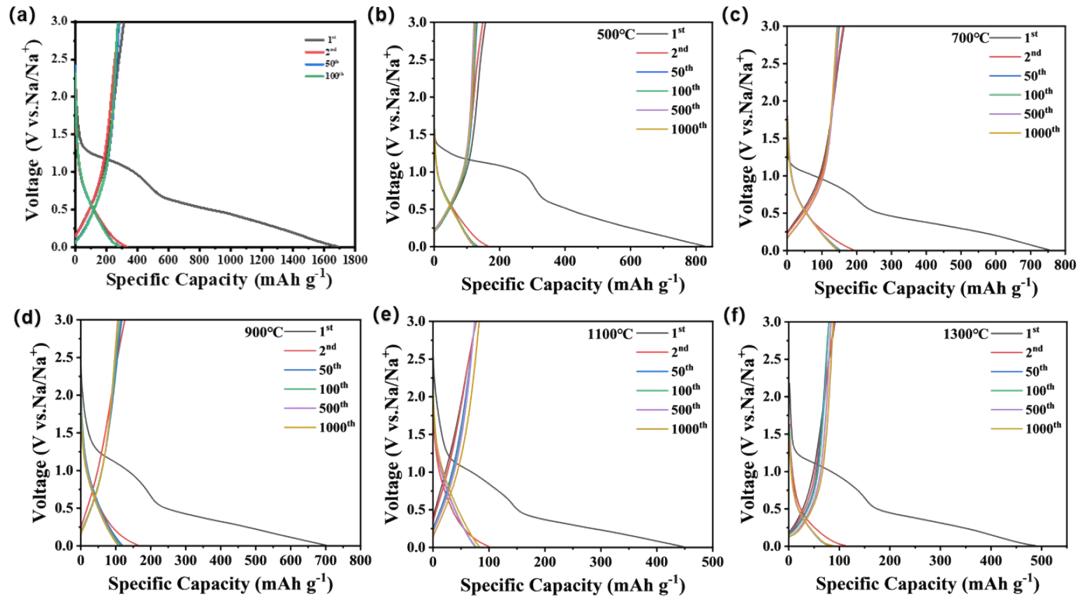


Fig. S2 (a) The 1<sup>st</sup>, 2<sup>nd</sup>, 50<sup>th</sup> and 100<sup>th</sup> charge-discharge curves of DSHC700 at 25  $\text{mA g}^{-1}$ ; (b-f) The 1<sup>st</sup>, 2<sup>nd</sup>, 50<sup>th</sup>, 100<sup>th</sup>, 500<sup>th</sup> and 1000<sup>th</sup> charge-discharge curves of DSHC500, DSHC700, DSHC900, DSHC1100 and DSHC1300 of different temperature at 500  $\text{mA g}^{-1}$ .

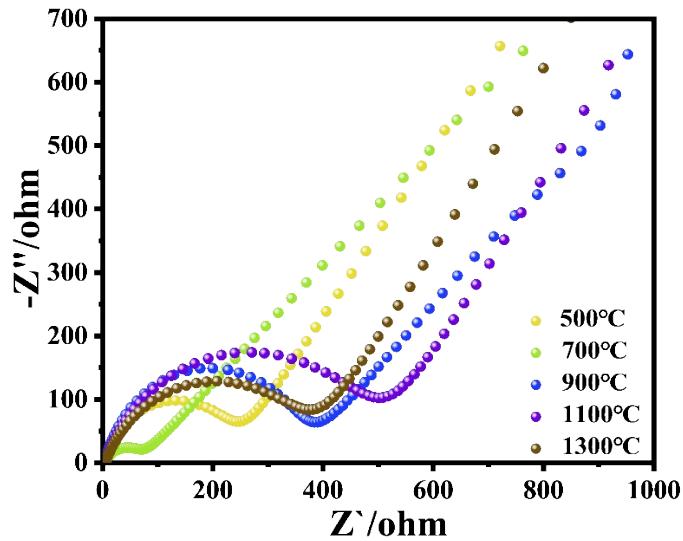


Fig. S3. The EIS spectra of DSHC-x after cycling.

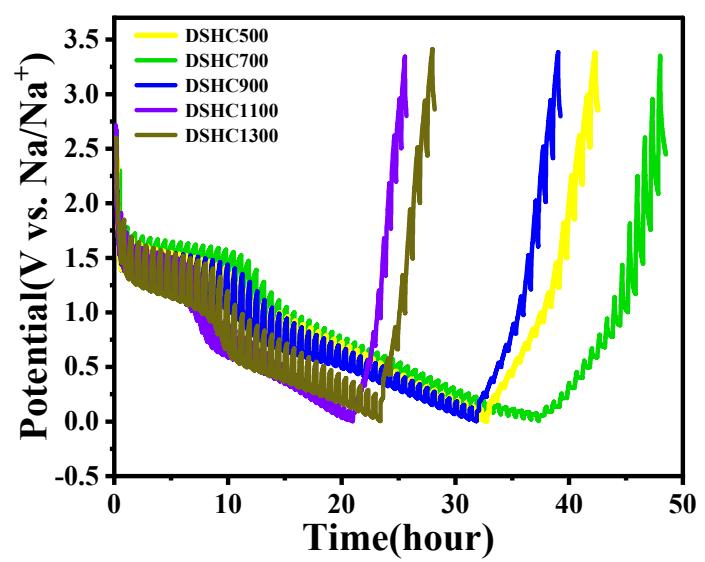


Fig. S4. GITT curves of DSHC-x.

Table S1.  $d_{002}$ ,  $I_D/I_G$ ,  $L_a$  of DSHC-x series.

	DSHC500	DSHC700	DSHC900	DSHC1100	DSHC1300
$d_{002}$ (nm)	0.378	0.382	0.391	0.367	0.362
$I_D/I_G$	1.03	1.013	1.003	1.11	1.08
$L_a$ (nm)	19.697	18.978	19.167	17.319	17.801

Table S2. Comparsion of the electrochemical performance of DSHC700 in this work and the previously reported hard carbon materials based anode materials

Samples	Initial coulumbic efficiency	Reversible capacity (mAh <sup>-1</sup> )	Current Density (mA g <sup>-1</sup> )	Cycle number	Ref.
DSHC700	18.5%	280.2	25	100	<b>This work</b>
CDHC-1300	80%	250.3	25	100	43
LPC-800	\	277.5	25	100	44
PSDHC-600	\	256.5	25	100	45
RPHC-1200	64%	224.3	25	100	46
CPP	59.8%	221.5	100	200	47
KHC-1300	64%	205	200	300	48
SC-800	78.2%	230	50	50	49
RRP-800-N	99.91%	223.7	31	100	50
HC-SC	92.3%	261	50	100	51
SAL	58.9%	245.3	50	100	52
SCL	88.1%	207.2	50	100	52
ECL	69.7%	227.1	50	100	52
CSHP2	44.1%	215	100	100	53

## Reference:

[1] Y. Tang, J. He, J. Peng, J. Yang, Z. Wu, P. Liu, K. Zhou, S. Hu, L. Hu, X. Wang, Electrochemical Behavior of the Biomass Hard Carbon Derived from Waste Corncob as a Sodium-Ion Battery Anode, *Energy Fuels*, 38(2024) 7389-7398.

[2] M. Yan, Y. Qin, L. Wang, M. Song, D. Han, Q. Jin, S. Zhao, M. Zhao, Z. Li, X. Wang, L. Meng, X. Wang, Recent Advances in Biomass-Derived Carbon Materials for Sodium-Ion Energy Storage Devices, *Nanomaterials*, 12(2022) 936.

[3] W. Lv, F. Wen, J. Xiang, J. Zhao, L. Li, L. Wang, Z. Liu, Y. Tian, Peanut shell derived hard carbon as ultralong cycling anodes for lithium and sodium batteries, *Electrochim. Acta*, 176(2015) 533-541.

[4] Z. Zhang, A. Zhang, S. Wang, J. Sun, L. Hou, C. Yuan, Biomass-derived hard carbon with tunable microstructures for sustainable and high-rate sodium-ion batteries, *New. J. Chem.* 49 (2025) 6277-6287.

[5] Z. Zhang, X. Li, P. Dong, G. Wu, J. Xiao, X. Zeng, Y. Zhang, X. Sun, Honeycomb-like hard carbon derived from pine pollen as high-performance anode material for sodium-ion batteries, *ACS Appl. Mater. Interfaces*, 10 (2018) 42796-42803.

[6] P. Wang, X. Zhu, Q. Wang, X. Xu, X. Zhou, J. Bao, Kelp-derived hard carbons as advanced anode materials for sodium-ion batteries, *J. Mater. Chem. A*, 5 (2017) 5761-5769.

[7] M. Kim, J. F. S. Fernando, Z. Li, A. Alowasheir, A. Ashok, R. Xin, D. Martin, A. K. Nanjundan, D. V. Golberg, Y. Yamauchi, N. Amiralian, J. Li, Ultra-stable sodium ion storage of biomass porous carbon derived from sugarcane, *Chem. Eng. J.*, 445 (2022) 136344.

[8] F. Lou, J. Wang, X. Wang, M. Zhang, J. Yuan, Unraveling multi-level porous carbon negative electrode materials based on Rosa roxburghii pomace for high-performance sodium-ion batteries, *RSC Adv.*, 15 (2025) 20872-20880.

[9] C. Wei, W. Dang, M. Li, X. Ma, M. Li, Y. Zhang, Hard-soft carbon nanocomposite prepared by pyrolyzing biomass and coal waste as sodium-ion batteries anode material, *Mater. Lett.*, 330 (2023) 133368.

[10] H. Wei, H. Cheng, N. Yao, G. Li, Z. Du, R. Luo, Z. Zheng, Invasive alien plant

biomass-derived hard carbon anode for sodium-ion batteries, *Chemosphere*, 343 (2023) 140220.

[11] S. Chen, K. Tang, F. Song, Z. Liu, N. Zhang, S. Lan, X. Xie, Z. Wu, Porous hard carbon spheres derived from biomass for high-performance sodium/potassium-ion batteries, *Nanotechnology*, 33 (2022) 055401.

Table S3. Impedance fitting values of DSHC-x electrodes before and after cycling.

	Before cycling		After cycling	
	$R_s$ ( $\Omega$ )	$R_{ct}$ ( $\Omega$ )	$R_s$ ( $\Omega$ )	$R_{ct}$ ( $\Omega$ )
DSHC500	2.19	647.2	1.33	266.3
DSHC700	2.55	436.8	0.92	93.2
DSHC900	2.16	230.6	1.01	200.2
DSHC1100	1.55	335.4	1.34	318.8
DSHC1300	4.54	316.3	1.38	277.2

## Text S1

(1) Calculation formula for  $d_{002}$

$$d_{002} = 0.5\lambda / \sin\theta \quad (1)$$

Where  $\lambda$  is the  $K_{\alpha}$  wavelength of Cu ( $\lambda = 0.15416$  nm),  $\theta$  is the diffraction angle.

## Text S2

(2) Calculation formula for  $L_a$

$$L_a(\text{nm}) = \left( 2.4 \times 10^{-10} \right) \times \lambda^4 \left( \frac{I_D}{I_G} \right)^{-1} \quad (2)$$

Where  $\lambda = 532$  nm is the laser wavelength in nanometers.

### Text S3

(3) Relationship between peak current  $i$  and scan rate  $v$

$$I = a * v^b \quad (3)$$

$$\log i = \log a + b * \log v \quad (4)$$

Where  $a$  and  $b$  are adjustable empirical constants, the value of  $b$  can be obtained from the slope of  $\log(i)$  and  $\log(v)$ . Generally speaking,  $a$  and  $b$  value close to 0.5 indicates that the diffusion process mainly controls the electrochemical reaction, while  $a$  and  $b$  value close to 1 represents an ideal capacitive process.

### Text S4

(4) Contribution equation of pseudo-capacitance behavior to sodium cell

$$i(V) = k_1 v + k_2 v^{0.5} \quad (5)$$

$$i(V)/v^{0.5} = k_1 v^{0.5} + k_2 \quad (6)$$

Where  $i$  is the current,  $v$  is the sweep speed,  $k_1$  and  $k_2$  are constants under a certain voltage,  $k_1 v$  and  $k_2 v^{1/2}$  represent the current contributed by diffusion behavior and pseudocapacitance behavior respectively. At a specific sweep speed, the corresponding values of  $k_1$  and  $k_2$  at a certain voltage can be obtained by fitting the linear relationship in formula (4), so as to calculate the current value contributed by the diffusion behavior and pseudocapacitance behavior.

### Text S5

(5) According to Fick's second law,  $D_{Na^+}$  can be calculated using the following simplified equation.

$$D_{Na^+} = \frac{4}{\pi \tau} \left( \frac{m_B V_m}{M_B S} \right)^2 \left( \frac{\Delta E_s}{\Delta E_\tau} \right)^2 \quad (7)$$

where  $m_B$  represents the weight of the active material on the electrode,  $V_m$  represents the molar volume of carbon,  $M_B$  represents the molecular mass of carbon,  $S$  represents the area of the HCRH-1200 electrode,  $\tau$  represents the pulse time, and  $\Delta E_s$  represents the voltage difference when the voltage reaches a steady state during a single GITT process.

