## Supporting Information

## A sustainable approach to hierarchically porous carbons from tannic acid and their utilization in supercapacitive energy storage

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Table S1. Che	emical composition	of the hierarchically	porous carbons.
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Sample code	Elemental composition (wt.%)					(O/C) <sub>at</sub> a
	С	Н	Ν	S	0	()
CK-750	90.9	0.2	0.1	0.0	8.9	0.07
CK-800	93.2	0.2	0.1	0.0	6.5	0.05
CK-850	93.8	0.1	0.2	0.0	6.1	0.05
CK-900	93.5	0.1	0.3	0.0	6.2	0.05

<sup>a</sup> O/C atomic ratio.

**Table S2**. Textural properties and yield of tannic acid-derived carbons produced by a variety of conditions.

Sample code	Activating agent/template	Textural properties			
	(wt. ratio)ª	S <sub>BET</sub> (m <sup>2</sup> g <sup>-1</sup> )	V <sub>Total</sub> (cm <sup>3</sup> g <sup>-1</sup> )	V <sub>&lt;2nm</sub> (cm <sup>3</sup> g <sup>-1</sup> )	
ТК	-/KCI (0/6.7)	510	0.48	0.16	26.1
CK	K <sub>2</sub> CO <sub>3</sub> /- (1/0)	1770	0.86	0.64	35.2
BK	KHCO <sub>3</sub> /KCI (1/6.7)	2180	0.91	0.82	36.3
OK	K <sub>2</sub> C <sub>2</sub> O <sub>4</sub> /KCl (1/6.7)	1990	0.84	0.72	38.7
CK-N	K <sub>2</sub> CO <sub>3</sub> /NaCl (1/6.7)	1890	0.91	0.69	35.0
CK-C	K <sub>2</sub> CO <sub>3</sub> /Na <sub>2</sub> CO <sub>3</sub> (1/6.7)	1830	0.95	0.68	32.4
CK-F	K <sub>2</sub> CO <sub>3</sub> /KCI (1/6.7)	2340	0.98	0.86	36.1
CK-M	K <sub>2</sub> CO <sub>3</sub> /KCI (1/6.7)	2130	0.87	0.78	35.2

<sup>a</sup> Weight ratio with respect to 1 part of tannic acid.

<sup>b</sup> Yield calculated by dividing the weight of porous carbon by the weight of tannic acid in the mixture prior to pyrolysis.

Carbon precursor	Activating agent	Textural properties		Carbon violda (9/)	Deference
		S <sub>BET</sub> (m <sup>2</sup> g <sup>-1</sup> )	V <sub>Total</sub> (cm <sup>3</sup> g <sup>-1</sup> )		Relefence
Tannic acid	K <sub>2</sub> CO <sub>3</sub>	2740	1.39	32.1	[This work]
Lignin	K <sub>2</sub> CO <sub>3</sub>	1950	0.93	39.0	[37]
Waste tea	K <sub>2</sub> CO <sub>3</sub>	1722	0.95	15.9	[40]
Coconut shell	K <sub>2</sub> CO <sub>3</sub>	1430	0.65	48.0	[41]
Chickpea husk	K <sub>2</sub> CO <sub>3</sub>	1780	0.65	13.0	[42]
Palm shell	K <sub>2</sub> CO <sub>3</sub>	1170	-	19.0	[43]
Rice husks	K <sub>2</sub> CO <sub>3</sub>	1165	0.78	14.2	[44]
Tobacco stems	K <sub>2</sub> CO <sub>3</sub>	2557	1.65	16.7	[45]
Tannin-F hydrogel	KOH	1800	0.65	21.0	[31]
Sugar cane pulp	KOH	2910	2.05	11.0	[7]
Lignite	KOH	2810	1.35	9.7	[6]
Gulfweed	KOH	2862	1.62	31.4	[5]
Glucose	KNO₃	1912	0.93	9.1	[46]
Glucosamine	$K_2C_2O_4$	2680	1.49	11	[47]
soya flour	$K_2C_2O_4$	2924	2.15	5.0	[47]
Sodium Glutamate	none	1010	0.56	33.0	[22]

**Table S3**. Textural properties and carbon yield of porous carbons obtained by using different biomass-based carbon precursors and activating agents.

<sup>a</sup> Yield calculated by dividing the weight of porous carbon by the weight of biomass-based precursor.



**Figure S1**. EDX analysis of a carbonized product (a) before and (b) after washing.



Figure S2. SEM images of CK-750 (a), CK-800 (b), CK-850 (c) and CK-900 (d).



**Figure S3**. SEM images of sample TK prepared in the absence of  $K_2CO_3$  (a and b) and CK carbon obtained in the absence of KCI (c and d).



**Figure S4**. XRD patterns of carbons prepared from tannic acid and KCI (TK), tannic acid and  $K_2CO_3$  (CK) and the ternary mixture (CK-800) using the same carbonization temperature (800 °C).



Figure S5. SEM images of BK (a) and OK (b) carbons.



**Figure S6**. N<sub>2</sub> adsorption isotherms (a) and pore size distributions (b) of carbons prepared at 800 °C using  $K_2CO_3$ , KHCO<sub>3</sub>,  $K_2C_2O_4$  and no activating agent. N<sub>2</sub> adsorption isotherms (c) and pore size distributions (d) of carbons prepared at 800 °C using  $K_2CO_3$ , Na<sub>2</sub>CO<sub>3</sub>, Na<sub>3</sub>CO<sub>3</sub>, Na<sub>2</sub>CO<sub>3</sub>, Na<sub>2</sub>



Figure S7. SEM images of CK-800 (a), CK-C (b) and CK-N (c) carbons.



Figure S8. SEM images of CK-F (a) and CK-M (b) carbons.



**Figure S9**. Chemical structure of tannic acid. Figure reproduced with permission.<sup>1</sup> Copyright, Elsevier, 2015.



**Figure S10.** Cyclic voltammograms at different scan rates of the porous carbons in (a)  $1M H_2SO_4$ , (b)  $1 M TEABF_4$  and (c) EMImTFSI/AN.



**Figure S11.** Nyquist plots (above) and Bode plots (below) for the porous carbons in 1 M  $H_2SO_4$  (a and d), 1 M TEABF<sub>4</sub> (b and e) and EMIMTFSI/AN (c and f).



**Figure S12**. Cycling stability of the electrodes in (a) 1 M  $H_2SO_4$ , (b) 1 M TEABF<sub>4</sub> and (c) EMIMTFSI/AN.

1. Z. Xia, A. Singh, W. Kiratitanavit, R. Mosurkal, J. Kumar and R. Nagarajan, *Thermochim. Acta*, 2015, **605**, 77-85.