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# Promising activated carbon derived from sugarcane tip as electrode material for high-performance supercapacitors†

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We present a simple, low-cost method for producing activated-carbon materials from sugarcane tips (ST) via two-step pre-carbonization and KOH activation treatment. After optimizing the amount of KOH, the resulting ST-derived activated carbon prepared with a KOH to PC-ST mass ratio of 2 (ACST-2) contained 17.04 wt% oxygen and had a large surface area of 1206.85 m<sup>2</sup> g<sup>-1</sup>, which could be attributed to the large number of micropores in ACST-2. In a three-electrode system, the ACST-2 electrode exhibited a high specific capacitance of 259 F g<sup>-1</sup> at 0.5 A g<sup>-1</sup> and good rate capability with 82.66% retention from 0.5 to 10 A g<sup>-1</sup>. In addition, it displayed a high capacitance retention of 89.6% after 5000 cycles at a current density of 3 A g<sup>-1</sup>, demonstrating excellent cycling stability. Furthermore, the ACST-2//ACST-2 symmetric supercapacitor could realize a high specific energy density of 7.93 W h kg<sup>-1</sup> at a specific power density of 100 W kg<sup>-1</sup> in 6 M KOH electrolyte. These results demonstrate that sugarcane tips, which are inexpensive and easily accessible agricultural waste, can be used to create a novel biomass precursor for the production of low-cost activated carbon materials for high-performance supercapacitors.

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## Introduction

The demand for green and efficient energy storage technologies has become increasingly urgent because of the depletion of traditional energy sources and the serious environmental issues associated with their use.<sup>1,2</sup> Supercapacitors (SCs), as advanced energy storage devices, have attracted increasing attention for many applications, such as portable electronics, hybrid electric vehicles, and biomedical devices owing to their high-power density, fast charge and discharge processes, long cycle life, convenient fabrication, and wide operating temperature.<sup>3–6</sup> Activated carbon materials (ACs) are widely utilized in SCs because they have excellent conductivity, low cost, and chemical stability. ACs with biological waste as the precursors are widely used in double-layer supercapacitors (EDLCs) because of their advantages that include a large specific surface area and being renewable and abundant.<sup>7</sup> To date, ACs derived from biological waste show good electrochemical performance, including materials derived from waste tea-leaves,<sup>8</sup> pomelo peel,<sup>9</sup> lotus seed-pods,<sup>10</sup> mangosteen peel,<sup>11</sup> cotton fabric,<sup>12</sup> and cotton straw.<sup>13</sup>

In principle, the charge storage of EDLCs can be attributed to electrostatic charge accumulated at the electrode/electrolyte interfaces; therefore, it is highly dependent on the effective surface area of the electrode materials that is accessible to the electrolyte ions.<sup>14</sup> In general, the electrode materials of supercapacitors are usually processed via an activation process to increase the specific surface area, which can lead to enhanced electrochemical performance. However, previous studies have found that the electrochemical performance and specific surface area are not linearly related. To obtain high electrochemical performance, it is necessary not only to increase the specific surface area but also have a suitable pore structure.<sup>15</sup> Normally, chemical activation is a clear pathway to prepare activated carbon. To achieve the above conditions, chemical activating agents such as KOH, NaOH, K<sub>2</sub>CO<sub>3</sub>, and H<sub>3</sub>PO<sub>4</sub> have been used to prepare ACs by changing the activation conditions. Among these reagents, KOH is the most effective and commonly used activator for chemical activation.<sup>7</sup>

The sugarcane tip is formed by the top 2–3 shoots of the sugarcane, accounting for ~10% of the sugarcane biomass, and is the main lignocellulosic waste after agricultural sugarcane harvest. Although bagasse, a waste product of sugarcane, has been widely used for the preparation of carbon materials, the application of sugarcane tips in this field has not been widely investigated. Because sugarcane tips are cheap, plentiful, and contain 43% cellulose, 27% hemicellulose, and 17% lignin,<sup>16</sup> they also serve as an effective activated-carbon precursor. Every year, 7 million tons of sugarcane tip waste is generated in Guangxi, China, of which, only a small portion is used for seed

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retention and feed, and the majority is left in the field to be burned, causing not only pollution and safety risks but is also a waste of resources. Therefore, the use of sugarcane tips to prepare activated-carbon materials can be one effective way to address these issues.

In this work, we demonstrate a facile strategy to synthesize activated carbon materials with a high specific surface area *via* a two-step method using ST as the biomass precursor. The effect of the activator (KOH) content on the morphology, crystallinity, and porosity of the produced activated carbon along with their electrochemical performance was investigated. ACST-2 had a high surface area of 1206.85 m<sup>2</sup> g<sup>−1</sup>, a synergism of porosity and graphitization degree. Benefiting from its unique properties, ACST-2 presented a high specific capacitance of 259 F g<sup>−1</sup> at 0.5 A g<sup>−1</sup> in a three-electrode system with 6 M KOH electrolyte. Furthermore, the ACST-2//ACST-2 symmetric supercapacitor device showed excellent energy storage performance with a high specific energy density of 7.93 W h kg<sup>−1</sup> at a specific power density of 100 W kg<sup>−1</sup>. Hence, this simple synthesis method has significant application prospects to prepare sugarcane-tip-derived activated carbon for use as supercapacitor electrode materials.

## Experimental

### Materials

The sugarcane tip used in this work was from Guangxi University. All chemicals in this study were of analytical reagent grade and used without further purification.

### Preparation of sugarcane-tip-derived activated carbon

The sugarcane tip was washed with distilled water and dried at 105 °C overnight and subsequently crushed into small particles to obtain a fine powder. Sugarcane tip powder was first carbonized at 500 °C for 2 h under vacuum with a heating rate of 5 °C min<sup>−1</sup>. The pre-carbonized powders are denoted as PC-ST (pre-carbonized sugarcane tip). Then, the PC-ST sample was mixed with KOH with a weight ratio of 1 : 1, 1 : 2, and 1 : 3. Thereafter, the resultant mixtures were calcined at 600 °C for 2 h under vacuum at a heating rate of 5 °C min<sup>−1</sup> in a vacuum muffle furnace. After cooling down to room temperature, the as-obtained products were washed with HCl solution (2 M) and deionized water several times until they obtained a pH of 7. They were then dried in an oven at 100 °C overnight. The final prepared activated carbon materials were named ACST-*x*, where *x* represents the mass ratio of KOH to PC-ST (1, 2, and 3). For comparison, carbonized sugarcane tip (CST) without any activation was obtained by direct carbonization at 600 °C for 2 h under vacuum conditions.

### Materials characterization

The morphology characteristics and microstructure of the materials were determined by field emission scanning electron microscopy (FESEM, SU8020) and transmission electron microscopy (TEM, JEOL JEM 2100). The crystalline attributes of the as-obtained products were determined by X-ray diffraction

(XRD, Rigaku D/Max 2500 diffractometer) using Cu K $\alpha$  radiation (40 kV, 100 mA,  $\lambda = 1.5406$  Å) in a scanning angle range of 5°–80° at a scanning rate of 5° min<sup>−1</sup>. Raman spectroscopy of the as-obtained products were recorded with a laser Raman spectroscope (inVia Reflex, 523 nm laser wavelength). Nitrogen adsorption and desorption isotherms were obtained at −196 °C with an adsorption setup (Micromeritics ASAP 2020 2.02). The specific surface area was determined *via* the Brunauer–Emmett–Teller (BET) method and the pore size distribution was calculated using a nonlocalized density functional theory (NLDFT) model. The surface elemental composition and chemical states were obtained by an X-ray photoelectron spectrometer (XPS, Thermo Fisher Scientific) with a monochromatic Al K $\alpha$  X-ray source (1486.6 eV).

### Electrochemical measurements

All the electrochemical characterization measurements including cyclic voltammetry (CV), galvanostatic charge-discharge (GCD), and electrochemical impedance spectroscopy (EIS) were carried out using a CHI 660E electrochemical workstation (Shanghai Chenhua, China) in aqueous 6 M KOH electrolyte. To prepare the working electrode, the ACST-*x*, acetylene black, and polytetrafluoroethylene (PTFE) were mixed at a mass ratio of 80 : 10 : 10 using *N*-methyl-2-pyrrolidone (NMP) as the solvent. The obtained slurry was coated on a piece of nickel foam (1.0 × 1.0 cm<sup>2</sup>) and then dried at 80 °C under vacuum for 12 h. An Hg/HgO reference electrode and Pt wire counter electrode were used in the three-electrode systems. CV and GCD measurements were carried out in the potential range of −1 to 0 V. EIS measurements were performed within the frequency range of 100 000 Hz to 0.01 Hz with an amplitude of 5 mV. Symmetrical supercapacitors were assembled from the same activated carbon material as the positive and negative electrodes. For the two-electrode system, the specific capacitance from the GCD curve was calculated from the following equation:

$$C_m = \frac{I \times \Delta t}{m \times \Delta v}, \quad (1)$$

where  $C_m$  is the specific capacitance (F g<sup>−1</sup>),  $m$  is the weight of the active substance,  $\Delta v$  is the discharge potential window (V),  $I$  is the discharging current (A), and  $\Delta t$  is the discharge time (s).

The specific capacitance from GCD was calculated with the symmetrical-electrode method calculated using equation:

$$C = \frac{4I \times \Delta t}{m \times \Delta v}, \quad (2)$$

where  $C$  is the specific capacitance,  $I$ ,  $\Delta t$  and  $\Delta v$  are the discharging current, the discharge time, and the discharge potential window, respectively, and  $m$  is the total mass of electrode materials.

The specific energy density ( $E$ , W h kg<sup>−1</sup>) and specific power density ( $P$ , W kg<sup>−1</sup>) of the symmetrical supercapacitor were calculated from the following equations:

$$E = \frac{1}{2 \times 4 \times 3.6} \times C \times \Delta V^2, \quad (3)$$



$$P = \frac{E \times 3600}{\Delta t} \quad (4)$$

## Results and discussion

A schematic for the preparation process to obtain the ACST is given in Scheme 1. The pre-carbonization temperature was determined by TGA, as shown in Fig. S1.† The sugarcane tip powder was subjected to a pre-carbonization process at 500 °C for 2 h in vacuum condition to yield PC-ST. To determine the optimal capacitor electrode material, the effect of the activation temperature on the performance of the supercapacitor was investigated, and the results are shown in Fig. S2 and S3.† To further refine the pore properties and specific surface area, PC-ST was activated with KOH (1 : 1, 1 : 2, and 1 : 3 mass ratio) at the optimized temperature of 600 °C for 2 h. Then, the obtained products were washed and dried. Significantly, the specific area and porous nature of the products were remarkably affected by the activator ratio.

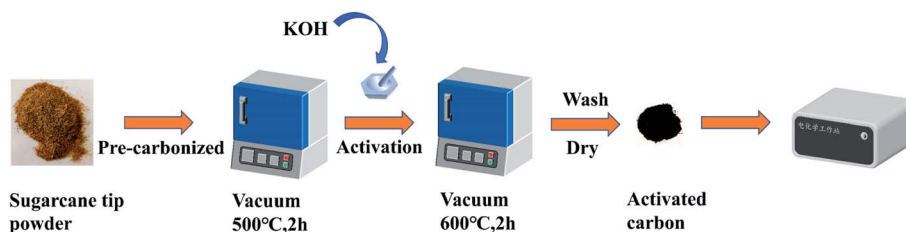
### Structural, morphological, and elemental analysis

The surface microscopic properties of the obtained products were characterized by FESEM and TEM. SEM images of CST and ACST-2 with low magnification are shown in Fig. S4.† The microstructure of CST is shown in Fig. 1a. The sample exhibits a relatively smooth surface with some small debris, however, no obvious pores appear on the surface. The microscopic structure of ACST-2 is shown in Fig. 1b, which clearly shows that etching the substance with KOH results in an extremely rough surface. Moreover, the surface is wrinkled and larger fragments are produced because of the added activator. The crumpled structure and rough surface of ACST-2 can effectively improve the open surface area, provide more active sites for storage of a large amount of charge, and increase electron transportability to facilitate the formation of the double electric layer.<sup>17</sup> Simultaneously, high-magnification images demonstrated that a significant number of pores could be observed on the surface of the ACST-2, as shown in the inset of Fig. 1b, confirming that carbon reacting with KOH effectively creates plentiful pores on the surface of the carbon materials. These pores embedded in the material are important for electron diffusion, and therefore, improving the electrochemical properties of materials. Furthermore, the EDS-mapping image of ACST-2, which is shown in Fig. 1c–e, further confirms the uniform distribution of C, O, and N atoms throughout the carbon framework.

The microstructure of ACST-2 was further characterized *via* TEM. Fig. 1f shows the presence of plate-like structure with relatively rough edges and abundant pores in ACST-2, which is consistent with the SEM results. Fig. 1g exhibits a high-resolution TEM image, from which a large number of micropores can be clearly observed, demonstrating that the material mainly consists of amorphous carbon. Furthermore, an unclear diffraction ring in the SAED diagram (inset in Fig. 1g) indicates that the sample has a low degree of crystallinity.<sup>18</sup>

Fig. 2a shows the XRD spectrums of the as-obtained ACST-1, ACST-2, and ACST-3. All products display two broad diffraction peaks at approximately 23° and 43° that correspond to the (002) and (100) planes of graphite,<sup>11,19</sup> respectively, suggesting the presence of amorphous carbon;<sup>20,21</sup> these results are in good agreement with the TEM and SAED results. Furthermore, it is noteworthy that clear intensity associated with ACST-2 and ACST-3 could be observed at a low-angle scattering peak. The shift of the intensity from high to low at a low angle can be explained by the presence of a certain amount of micropores in the sample.<sup>22,23</sup> The results for the degree of graphitization of the as-prepared carbon materials were further confirmed by Raman spectroscopy, as shown in Fig. 2b. The two dominant peaks at 1345 and 1590 cm<sup>−1</sup> correspond to the D-band and G-band of carbon materials, respectively. The D-band is associated with defects in the carbon material, while the G-band can be attributed to the vibration of sp<sup>2</sup> carbon in the graphite crystallites.<sup>24</sup> Generally, the intensity ratio of the D-band and G-band (*I*<sub>D</sub>/*I*<sub>G</sub>) is used to reflect the graphitic degree of the samples, and a higher value of *I*<sub>D</sub>/*I*<sub>G</sub> indicates the presence of more defects in the carbon materials.<sup>12,25</sup> Here, the measured *I*<sub>D</sub>/*I*<sub>G</sub> ratios of ACST-1, ACST-2, and ACST-3 were 0.84, 0.87, and 0.88, respectively. It can be observed that as the amount of KOH increased, the *I*<sub>D</sub>/*I*<sub>G</sub> ratio gradually increased, demonstrating increased disordered portions and defects in ACST-*x*. The presence of defects is beneficial for promoting the specific surface area and charge transport.<sup>26</sup> However, the electrical conductivity of the materials may decline with the existence of the defects.<sup>27</sup> Therefore, determining a suitable *I*<sub>D</sub>/*I*<sub>G</sub> ratio is beneficial for enhancing electrochemical performance.

To further study the porous texture, the as-obtained carbon materials were investigated *via* N<sub>2</sub> adsorption–desorption measurements. As shown in Fig. 2c, all the samples revealed typical type I isotherms, and a sharp increasing adsorption isotherm could be observed at low relative pressures <0.1. This increase indicated that the ACST-*x* contained a large number of micropores because the adsorption and desorption curves



Scheme 1 Schematic illustration of synthesis procedure to obtain activated carbon derived from sugarcane tip (ACST).





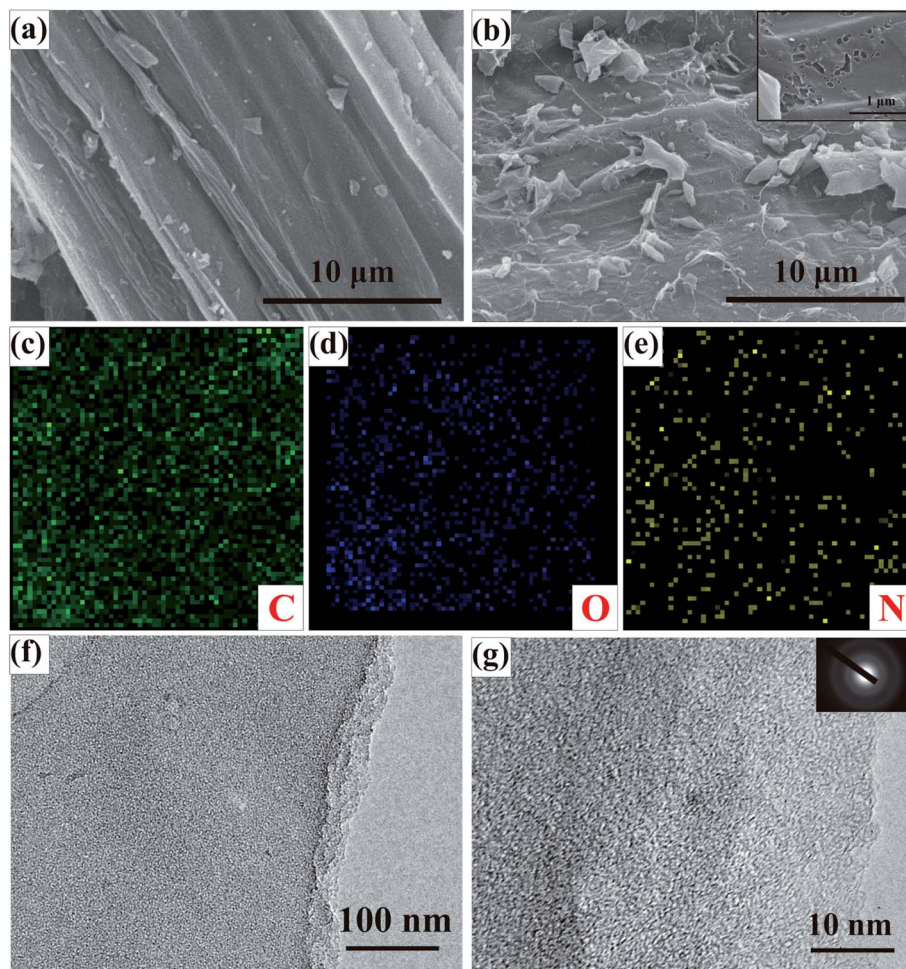


Fig. 1 FESEM images of (a) CST; (b) ACST-2 (the inset is the high magnification image); (c–e) the elemental mapping of ACST-2 of C; O; N; (f and g) TEM images of ACST-2. The inset in (g) shows the corresponding SAED pattern.

almost coincided with each other.<sup>28,29</sup> Fig. 2d shows the corresponding pore size distribution curves of ACST-*x* based on the NLDFT method with the desorption curve. It can be seen that all the samples had micropores primarily less than 2 nm, which is suitable for charge accumulation, and thus, increases the electric double-layer capacitance. Details of the porous texture of the ACST-*x* samples are summarized in Table 1. It can be noted that the specific surface area and total pore volume increased rapidly as the activator content increased. The specific surface area of ACST-1, ACST-2, and ACST-3 were determined to be 496.48, 1206.85, and 1438.01 m<sup>2</sup> g<sup>−1</sup> and the total pore volumes were 0.28, 0.52, and 0.61 cm<sup>3</sup> g<sup>−1</sup>, respectively. Intriguingly, this trend was similar to that of the  $I_D/I_G$  ratio with increasing amounts of KOH. These results indicate increased defect generation due to the reaction between KOH and the carbon skeleton, which facilitates utilization of the porosity and increases the surface area by generating a large number of micropores and enlarging the pore size.

Fig. 3 displays the chemical components and functional groups of CST and ACST-2 analyzed *via* XPS. Three peaks with binding energies of 285.2, 401.1, and 533.0 eV can be observed

from the survey of CST and ACST-2, as shown in Fig. 3a, which can be assigned to the C 1s, N 1s, and O 1s peaks, respectively. The C : O : N content was determined to be 84.35 wt% : 13.73 wt% : 1.91 wt% and 81.28 wt% : 17.04 wt% : 1.68 wt% for the CST and ACST-2 samples, respectively. Fig. 3b exhibits the high-resolution spectrum of C 1s, which could be deconvoluted into four individual peaks at 284.8, 286.3, 287.2, and 288.4 eV that could be attributed to C–C, C–O, C=O, and O–C=O, respectively.<sup>30</sup> The high-resolution spectrum of O 1s is given in Fig. 3c, and three peaks could be observed and attributed to C=O, C–O, and O=C–O bonds at the binding energies of 531.8, 532.7, and 533.6 eV, respectively.<sup>31,32</sup> Previous literature reported that double and single bonded oxygen could enhance the hydrophilicity of carbon materials and markedly improve the total specific capacity by enhancing the faradaic pseudo-capacitance.<sup>33,34</sup> In addition, the N 1s spectra (Fig. 3d) revealed three major peaks that could be attributed as pyridinic N (398.9 eV, N-6), pyrrolic N (400.4 eV, N-5), and graphitic N (402.1 eV, N-Q).<sup>35</sup> It has been widely demonstrated that pyridinic N and pyrrolic N display excellent electrochemical capability in an alkaline solution and generate pseudo-

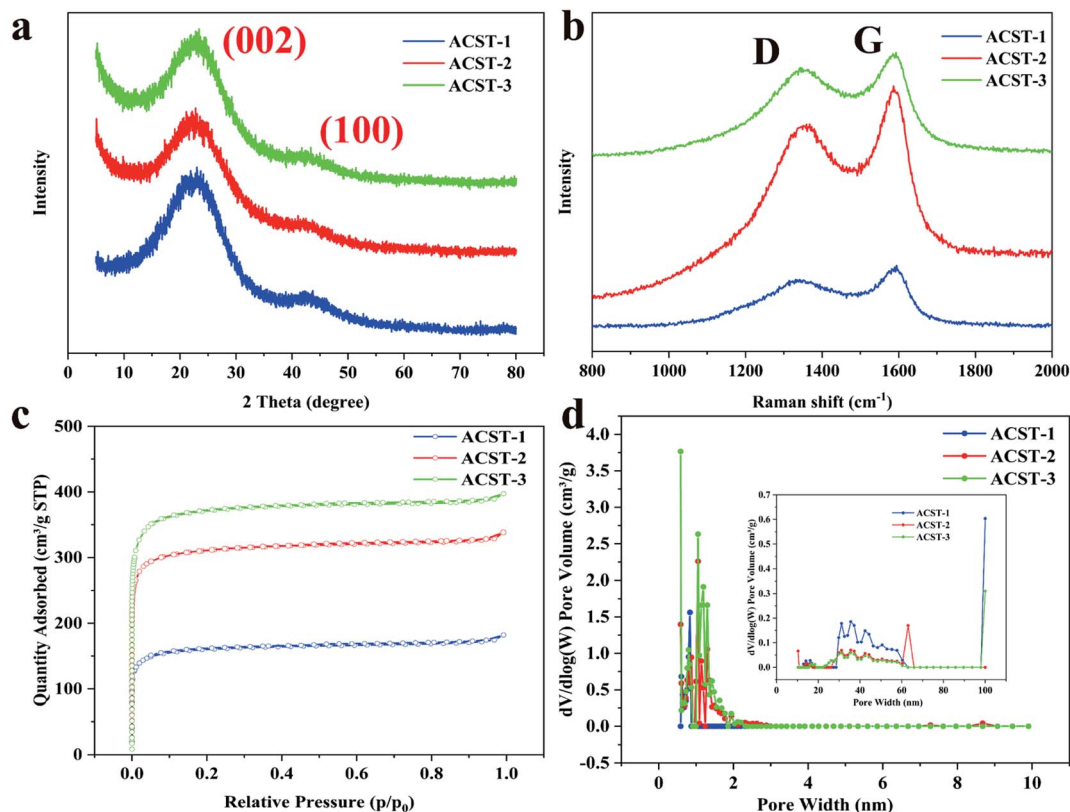


Fig. 2 (a) XRD patterns; (b) Raman spectra; (c)  $N_2$  adsorption–desorption isotherms, and (d) the pore size distribution curves of the ACST-1, ACST-2, and ACST-3.

capacitance.<sup>36,37</sup> Graphitic N can be embedded in the carbon matrix and combined with carbon atoms, contributing to electron diffusion and improving the conductivity of the electrode.<sup>38</sup>

### Electrochemical performance

To check the electrochemical properties of CST, ACST-1, ACST-2, and ACST-3, CV was employed based on a three-electrode configuration in an aqueous electrolyte of 6 M KOH solution within a potential window ranging between  $-1$  and  $0$  V vs. Hg/HgO at a scan rate of  $10 \text{ mV s}^{-1}$ . The results show that  $600^\circ\text{C}$  is the optimum activation temperature. Fig. 4a displays the CV curve of CST, which resembles a triangle; the region under the curve is small, which indicates weak charge storage performance and a relatively low capacitance value. The CV curve after activation of the sample clearly exhibits an approximate quasi-rectangular shape, demonstrating excellent capacitive behavior

and a relatively small internal resistance. Compared with the other samples, the CV curve of ACST-2 shows the largest enclosed area, indicating the highest specific capacitance of ACST-2. Moreover, a wide peak could be observed in the voltage range of  $-0.8$  to  $-0.3$  V, which is due to pseudo-capacitance from the faradaic processes of the  $\text{C}=\text{O}$ ,  $\text{N}-5$ , and  $\text{N}-6$  functional groups on the electrode surface.<sup>39</sup> Fig. 4b shows the CV curves for the ACST-2 electrode at various scan rates. The CV curves continue to have a nearly rectangular shape and display slight variations in the scan rate range of  $5$ – $100 \text{ mV s}^{-1}$ , demonstrating rapid charge transport capability and easy ion diffusion within its pore channel.<sup>40</sup> Nevertheless, the curve was distorted as the scan rate increased to  $200 \text{ mV s}^{-1}$ . This could be attributed to the ohmic resistance for electrolyte motion in the carbon pore channels, in which distributed charge storage has been reported for the double layer formation mechanism.<sup>8,41</sup>

Table 1 BET textural characteristics of the samples

Sample	$S_{\text{BET}}^a$ ( $\text{m}^2 \text{ g}^{-1}$ )	$S_{\text{mic}}^b$ ( $\text{m}^2 \text{ g}^{-1}$ )	$V_{\text{total}}^c$ ( $\text{cm}^3 \text{ g}^{-1}$ )	$V_{\text{mic}}^b$ ( $\text{cm}^3 \text{ g}^{-1}$ )	$D_{\text{aver}}^d$ (nm)
ACST-1	496.48	403.74	0.28	0.21	4.42
ACST-2	1206.85	1062.26	0.52	0.42	3.83
ACST-3	1438.01	1266.56	0.61	0.50	3.50

<sup>a</sup> Total surface area calculated by the BET method. <sup>b</sup> The specific surface area and volume of micropore measured by the  $t$ -plot method. <sup>c</sup> Total pore volume calculated at  $P/P_0 = 0.99$ . <sup>d</sup> Average pore diameter determined by BJH desorption.



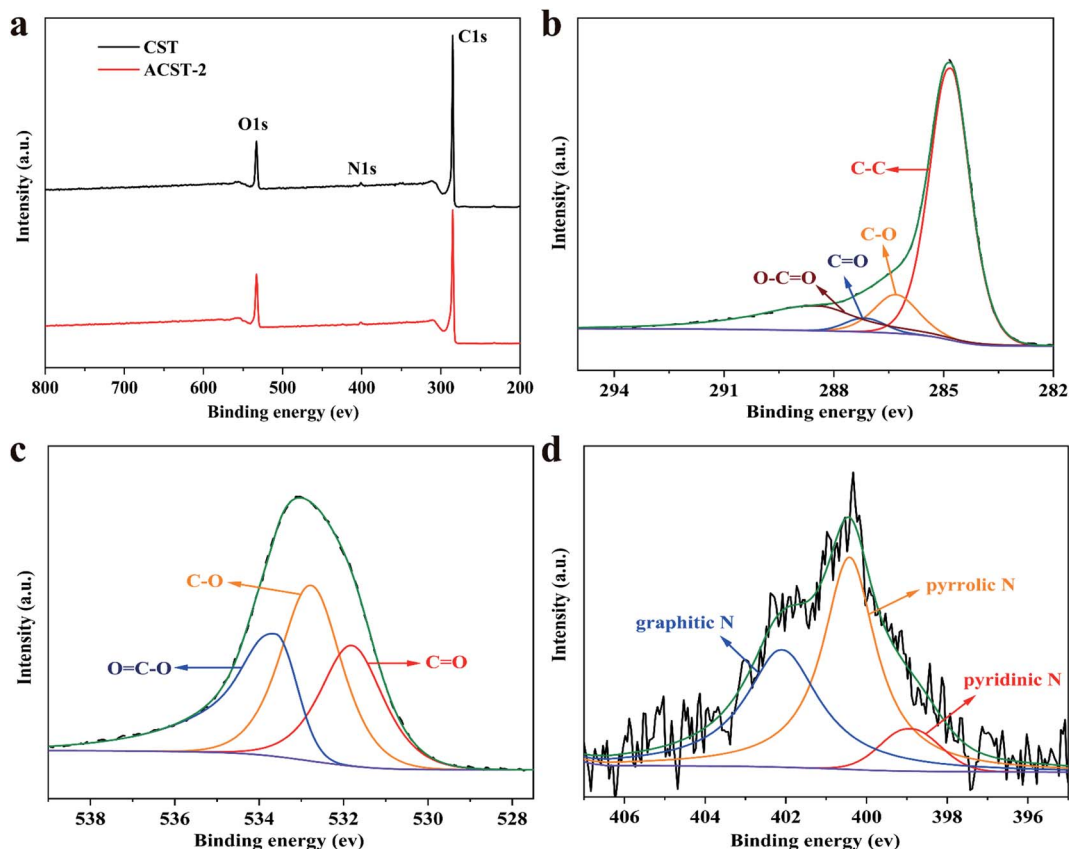


Fig. 3 XPS (a) survey spectrum of CST and ACST-2; high-resolution spectrum of ACST-2 of (b) C 1s; (c) O 1s; and (d) N 1s.

Fig. 4c exhibits the GCD profiles of the ST-derived electrodes under a current density of  $0.5 \text{ A g}^{-1}$ . The profiles display a quasi-triangle shape instead of a symmetric triangle, which indicates an associative feature of electrical double-layer capacitance and pseudocapacitive behaviors for the N and O active species.<sup>42</sup> As illustrated in literature, N-5 and N-6 play a vital role in affording pseudo-capacitance and thus leading to curve distortion.<sup>43</sup> Notably, the ACST-2 electrode displays the longest discharge time, while ACST-1 and ACST-3 show slightly reduced discharge times, and the CST electrode presents the shortest discharge time. This reveals that the ACST-2 electrode exhibits the largest specific capacitance ( $259 \text{ F g}^{-1}$ ), followed by ACST-1 ( $243.8 \text{ F g}^{-1}$ ), ACST-3 ( $211.9 \text{ F g}^{-1}$ ), and CST ( $19.5 \text{ F g}^{-1}$ ), which agreed well with the CV results at  $10 \text{ mV s}^{-1}$ . Although the specific capacitance of ACST-2 is the largest, the surface area of the carbon materials did not follow the same sequence (ACST-3 > ACST-2 > ACST-1). This demonstrates that the capacitance is not only related to the specific surface area but also associated with other characteristics, such as pore size distribution or functional groups.<sup>44</sup> Importantly, the surface area-normalized capacitance of ACST-2 was  $0.21 \text{ F m}^{-2}$ , which is higher than ACST-1 ( $0.15 \text{ F m}^{-2}$ ), exhibiting a higher surface area utilization rate for ACST-2 than ACST-3. The superior specific capacitance for the ACST-2 can be attributed to the relatively high effective surface area with abundant exposed ion absorption sites for charge accumulation in electric double-layer.<sup>45</sup>

The GCD profiles of the ACST-2 electrode displayed an approximately triangular shape at various current densities, as shown in Fig. 4d. Additionally, the GCD profile could still be maintained as a triangle under a high current density of  $10 \text{ A g}^{-1}$ , demonstrating desirable capacitive behavior. In addition, only a slight IR drop ( $200 \text{ mV}$ ) was detected at this high current density, indicating a high charge-discharge efficiency and low equivalent series resistance,<sup>10,46</sup> which can explain the distortion of the CV curves.

The specific capacitances of all the samples gradually decreased with increasing current densities from  $0.5$  to  $10 \text{ A g}^{-1}$ , as shown in Fig. 4e. This decrease could be attributed to the slow migration of ions between the electrolyte and surface of electrode at low current density. All hydrated ions could entirely diffuse within the pore, enhancing the ion storage capacity and thus increasing the specific capacitance. However, at high current densities, the electrolyte ions do not have sufficient time to access the microporous surface, and thus, the specific capacitance at high current densities is lower than that at low current densities.<sup>40</sup> In addition, at a current density of  $10 \text{ A g}^{-1}$ , the capacitance of ACST-1, ACST-2, and ACST-3 remains at  $79.70\%$  ( $194.3 \text{ F g}^{-1}$ ),  $82.66\%$  ( $214.1 \text{ F g}^{-1}$ ), and  $91.65\%$  ( $194.2 \text{ F g}^{-1}$ ) of the initial capacitance at  $0.5 \text{ A g}^{-1}$ . These results demonstrate that ACST-2 is a promising carbon electrode material for supercapacitors. The cycling stability imply that the number of times the electrolyte ions could migrate into the





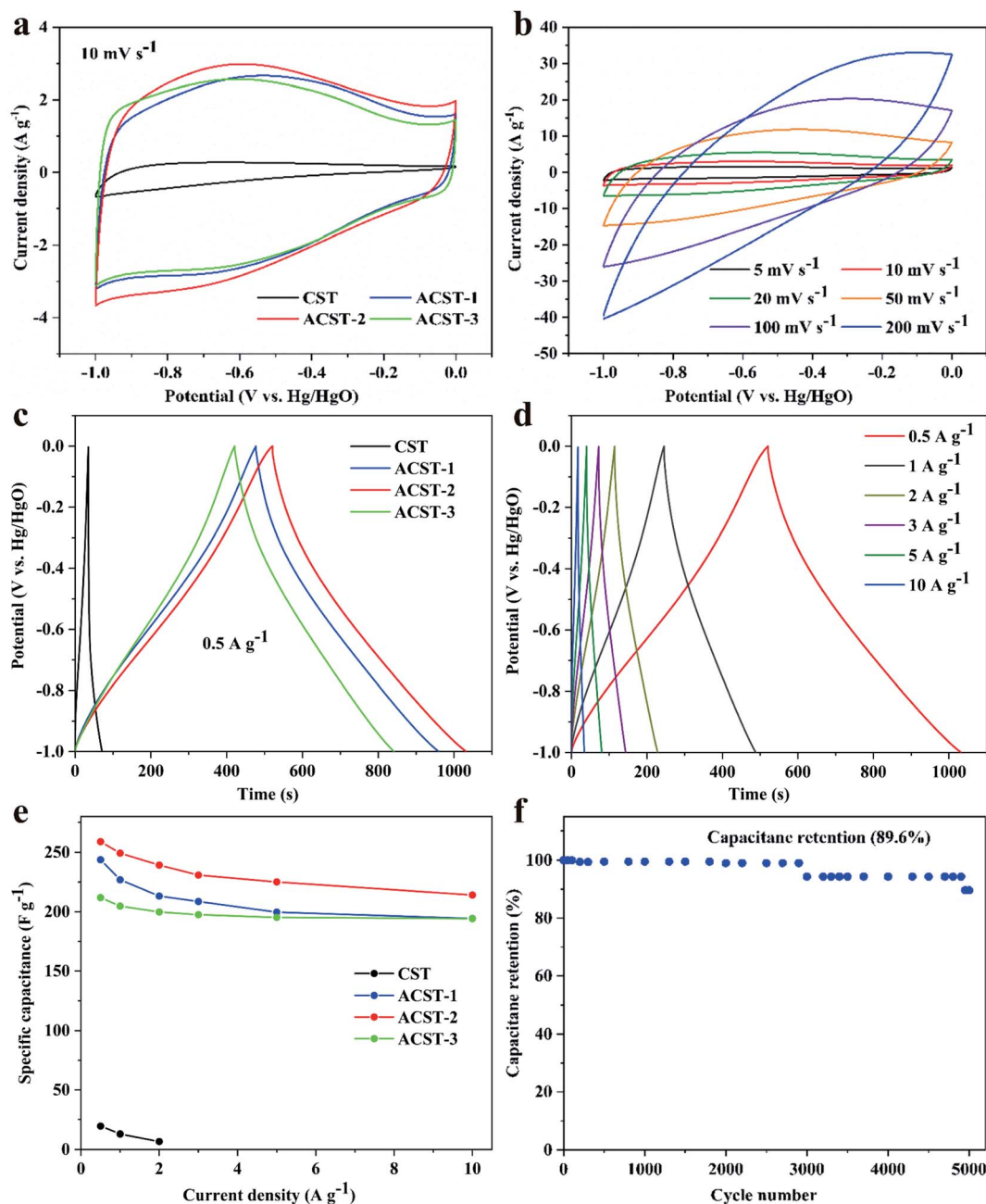


Fig. 4 (a) CV curves of CST and ACST-*x* at a scan rate of  $10 \text{ mV s}^{-1}$ ; (b) CV curves at various scan rates of ACST-2; (c) GCD profiles of CST and ACST-*x* at a current density of  $0.5 \text{ A g}^{-1}$ ; (d) GCD profiles of ACST-2 under various current densities; (e) specific capacitance at different current densities of CST, ACST-1, ACST-2, and ACST-3-derived electrodes; (f) cycling stability of ACST-2 at  $3 \text{ A g}^{-1}$ .

available pores for the repeated number of cycles. Fig. 4f shows the long cycling stability of the ACST-2 electrode as demonstrated *via* GCD measurement at  $3 \text{ A g}^{-1}$  for 5000 cycles. After cycling, the capacitance value decreased, which may be due to the increase of internal resistance. The ACST-2 electrode still retained 89.6% of the initial capacitance after 5000 cycles, demonstrating excellent cycling stability and high reversibility.

The electrochemical impedance spectroscopy (EIS) of ACST-1, ACST-2, and ACST-3 were further studied from 0.01 to 100 000 Hz in a 6 M KOH solution. Fig. 5a shows the Nyquist plots of the ACST-1, ACST-2, and ACST-3 electrodes and an

equivalent electric circuit to fit the Nyquist plot. All samples were divided into three different sections: a semicircle, a straight line of  $45^\circ$ , and a vertical line in the high, medium, and low frequencies. As shown in the inset of Fig. 5a, the intercept of the semicircle at the real axis ( $Z'$ ) indicates the equivalent series resistance ( $R_s$ ), which contains the intrinsic electroactive material resistance, ionic transport resistance of the electrolyte, and the contact resistance at the interface of the electrode materials and current collectors.<sup>47</sup> The  $R_s$  value of ACST-1, ACST-2, and ACST-3 were 0.32, 0.53, and 0.33  $\Omega$ , respectively, indicating negligible contact resistance and good



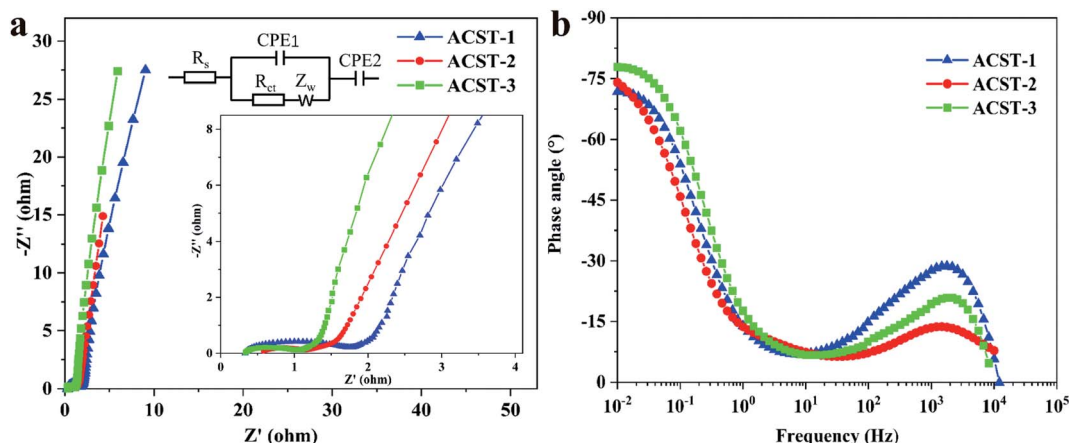


Fig. 5 (a) Nyquist plots and (b) Bode plots of the ACST-1, ACST-2, and ACST-3 electrodes.

conductivity.<sup>48</sup> The diameter of the semicircle is related to the interfacial charge transfer resistance ( $R_{ct}$ ). The  $R_{ct}$  values for ACST-1, ACST-2, and ACST-3 were 1.24, 0.54, and 0.62  $\Omega$ , respectively, which demonstrated minimal charge transfer resistance for the ACST-2 electrode. In addition, the short line with the 45° slope is called the Warburg resistance, which is associated with ion diffusion/transport into the active

materials.<sup>49</sup> At low frequencies, the straight line is nearly vertical—close to 90°—suggesting better capacitive behavior and rapid ion diffusion in the carbon electrode structure.<sup>50</sup> Fig. 5b displays the Bode phase plots of ACST-1, ACST-2, and ACST-3, where the phase angle shifts to negative values of  $-71.77^\circ$ ,  $-74.04^\circ$  and  $-77.80^\circ$ , respectively, at a low frequency of 0.01 Hz. This indicates that the results are relatively close to

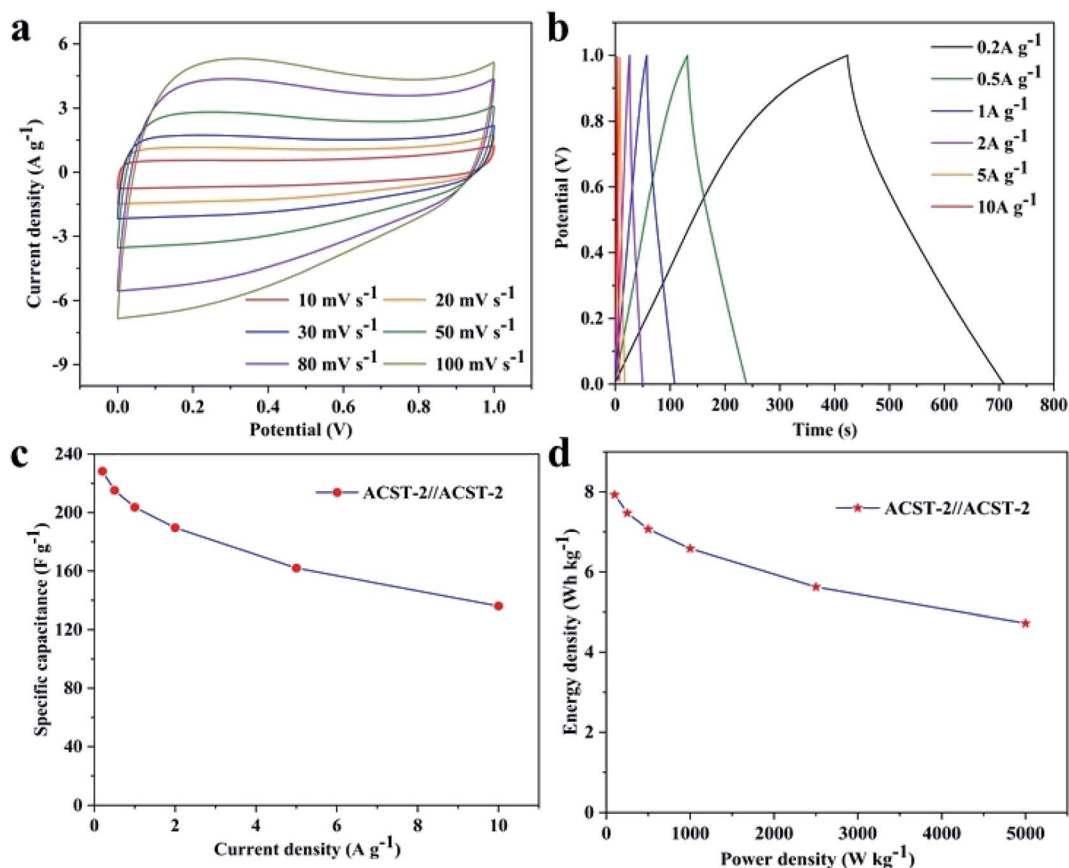


Fig. 6 Electrochemical performance of ACST-2 in a two-electrode system using 6.0 M KOH electrolyte: (a) CV curves at scanning rates from 10 to 100  $\text{mV s}^{-1}$ ; (b) GCD curves at 0.2 to 10  $\text{A g}^{-1}$ ; (c) specific capacitance at different current densities for ACST-2; (d) Ragone plot of ACST-2 in a symmetrical supercapacitor.





that of an ideal supercapacitor ( $-90^\circ$ ), which suggests a primarily double-layer charge storage contribution with some pseudo-capacitive contribution for the material.<sup>51</sup>

To study the practical application of the ACST-2 sample, the ACST-2//ACST-2 symmetric supercapacitor was assembled in 6.0 M KOH electrolyte with an operating voltage range of 0–1 V. Fig. 6a exhibits the CV curves of the symmetric capacitor, the electrode exhibits a rectangular shape at a scanning rate of  $10 \text{ mV s}^{-1}$ , indicating a typical double-layer capacitor behavior at the electrode and electrolyte interface. In addition, the CV curve remains approximately rectangular at a high scan rate of  $100 \text{ mV s}^{-1}$ , indicating an ideal electrochemical capacitance behavior and excellent rate performance. The GCD curves were tested at different current densities ( $0.2$  to  $10 \text{ A g}^{-1}$ ) and are shown in Fig. 6b. All the curves exhibited an approximately symmetric linear behavior, demonstrating the existence of bilayer behavior with favorable reversibility. From the GCD curves, specific capacitances of 228.3, 215.2, 203.6, 189.6, 162, and  $136 \text{ F g}^{-1}$  were obtained at 0.2, 0.5, 1, 2, 5, and  $10 \text{ A g}^{-1}$ , respectively. Fig. 6c shows the relationship between the specific capacitance and the current density, a specific capacitance of 59.6% was still retained even though the current density increased 50 times, indicating good rate performance for the ACST-2 supercapacitor. Fig. 6d shows the Ragone plot of the ACST-2//ACST-2 symmetric supercapacitor. The specific energy density of the symmetry supercapacitor was measured at  $7.93 \text{ W h kg}^{-1}$  for a specific power density of  $100 \text{ W kg}^{-1}$ . For practical applications, it is important to measure the leakage current and self-discharge of the device, the results of our studies in this direction are shown in Fig. S5.† These results demonstrated that ACST-2 is a promising electrode material for supercapacitor applications.

## Conclusions

A novel sugarcane-tip-derived activated-carbon material was successfully prepared *via* a two-step method of pre-carbonization and KOH activation. The optimized activated carbon (ACST-2) displayed a large number of micropores, a high specific surface area of  $1206.85 \text{ m}^2 \text{ g}^{-1}$ , and a high oxygen content of 17.04 wt%. The ACST-2, as an electrode material, exhibited a high specific capacitance of  $259 \text{ F g}^{-1}$  at a current density of  $0.5 \text{ A g}^{-1}$  in a three-electrode system. Moreover, the material displayed good rate capability with 82.66% retention from 0.5 to  $10 \text{ A g}^{-1}$  and excellent cycling stability with 89.6% capacitance retention after 5000 cycles at current density of  $3 \text{ A g}^{-1}$ . Furthermore, the symmetric ACST-2//ACST-2 device exhibited excellent electrochemical properties in KOH electrolyte with a wide operating voltage range of 0–1 V. These outstanding electrochemical results confirm that sugarcane tip-derived activated carbon is a promising electrode material for energy-storage device application.

## Author contributions

Bo Wei: date curation, methodology, and writing the original draft. Tiantian Wei: investigation. Caifeng Xie:

conceptualization, review and editing. Kai Li: grammar check & data curation. Fangxue Hang: review, editing, and funding acquisition.

## Conflicts of interest

There are no conflicts to declare.

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