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Capacitive performance of porous carbon nanosheets derived from biomass cornstalk†

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Porous carbon nanosheets (aCS) formed by carbonization and chemical activation of biomass cornstalk are presented for capacitive energy storage. The obtained carbon possesses a two-dimensional (2D) sheet-like structure which is mainly composed of abundant micropores. Depending on the amount of KOH used in the activation step, the specific surface area of the carbon sheets varies from 388 to 1736 m² g^{−1}, whereas the lateral size is in the range of 50–100 μm, and sheet thickness is around 100 nm. The capacitive performance of aCS-5 in both KOH and Na₂SO₄ aqueous electrolytes is investigated. The aCS-5 electrode in KOH electrolyte exhibits a remarkable high-rate capability due to the dramatically shortened ion pathway and more exposed external surface. While, in Na₂SO₄ it shows relatively lower capacitance retention but a stable and extended voltage window (0–1.6 V), thus delivering a specific energy of 20.2 W h kg^{−1} at a power density of 398 W kg^{−1}. The aCS-5 with a large aspect ratio (200–500) can also serve as an electrode for assembling a solid-state capacitor (SSC) using PVA/KOH gel as a polymer electrolyte. The SSC retains the normal capacitive performance under bent and twisted states and exhibit an unusual rate capability with an areal capacitance of 136 mF cm^{−2} at 0.5 mA cm^{−2}.

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1. Introduction

Supercapacitors have attracted considerable attention in recent years due to their high power density, fast charge–discharge rates and excellent cycling stability.^{1–4} Different from the batteries that store energy through ion insertion in the bulk of electrodes, carbon-based supercapacitors rely on fast ion adsorption at the carbon surface to store energy.^{5,6} Such an energy storage mechanism makes supercapacitors store and release energy very quickly. However, energy stored in supercapacitors is about one order of magnitude lower than that of batteries. Over the past few decades, tremendous efforts have been devoted to boosting the energy density of a supercapacitor without lowering its power density. As the energy density of a supercapacitor is proportional to the specific capacitance of electrode and square of the cell voltage, improving the energy density of a supercapacitor can be readily achieved by enhancing the specific capacitance of carbon electrode through increasing its ion-accessible surface area and optimizing its pore structure for better ion transport.^{7,8} On the other hand, using non-aqueous electrolytes,^{9–11} or configuring an aqueous-based asymmetric supercapacitor^{4,12} have demonstrated to be effective solution to get a high energy density.

Carbonization of various cheap biomasses represents one of most cost-effective methods for scalable production of activated carbon towards energy storage applications.^{13–20} The capacitive performances of carbon electrodes are strongly affected by their ion-accessible surface area and pore textures. Hierarchical porous carbons with interconnected and well-balanced micro-meso-macropores ratio are considered to be highly desirable because the macropores facilitates electrolyte transport while the micropores improve the charge storage capacity.²¹ Besides the pore structure, the macroscopic morphology of the carbon electrodes also plays a vital role in boosting its capacitive performance. For example, unlike the conventional activated carbons which are dominated by long and tortuous micropores, hollow carbon fibers derived from carbonization and chemical activation of natural cotton can potentially shorten the ion-transport length and facilitate the utilization of carbon electrode for double-layer capacitance.²² Most recently, two-dimensional (2D) carbon materials have emerged as promising candidates for high-performance supercapacitor in view of their high aspect ratio, short ion-transport length and more exposed external surface area.^{23–26} The current methods for fabrication of 2D carbon materials include chemical vapor deposition of carbon films on flat inorganic template,^{26,27} pyrolysis of gelatin on boron oxides,²⁵ chemical activation of ion-exchanged resin,²⁸ or using 2D graphene precursor as structure-direct agents to make graphene-sandwiched carbon.^{29–33} In spite of the improved capacitive performance of the 2D carbon materials, their synthetic methods usually involve the complicated process or require some expensive carbonaceous precursors. However,

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scalable preparation of sheet-like nanocarbons through carbonization of renewable and low-cost biomasses still remains a challenge.^{17,34}

Cornstalk is a kind of agricultural byproducts which are abundant in China countryside. Although carbonization of cornstalk for supercapacitor electrodes have been reported as electrode materials for supercapacitor or Li-ion batteries,^{35–37} the overall synthetic procedure involves the use of Fe or Ni to catalyze the graphitization of carbon. Moreover, the obtained carbon materials suffer from either low energy density or poor cycling stability. Herein, inspiring by the fact that cornstalk pith is inherently composed of flake-like cellulose, we report the sheet-like porous carbon by carbonization of cornstalk pith, followed by chemical activation with KOH. Depending on the amount of KOH used in the activation step, the as-made activated carbon samples (aCS) remain sheet-like morphologies, while exhibiting specific surface area (SSA) in the range of 388–1736 m² g^{−1}. When used as supercapacitor electrode, the aCS-5 exhibits a specific capacitance as high as 301 F g^{−1} in 6.0 M KOH with a capacitance retention of 82% at 60 A g^{−1} (247 F g^{−1}) in spite of its pore size in the range of 1.0–4.0 nm and pore volume dominated by micropores. More interestingly, a symmetric capacitor built with aCS as electrode can reversibly operate in the cell voltage of 0–1.6 V in neutral Na₂SO₄ aqueous electrolyte, delivering an energy density of 20.2 W h kg^{−1} at 398 W kg^{−1}. Moreover, the 2D structure with large aspect ratio (200–500) enables aCS to serve as solid-state capacitor electrode, which shows flexible property without noticeable change of capacitive performance under normal, bent and twisted states. The aCS electrode can deliver an areal capacitance of 136 mF cm^{−2} at 0.5 mA cm^{−2} and retain 92% of its initial capacitance after 1000 cycles of charge discharge.

2. Experimental

2.1 Sample preparation

The dried cornstalk pith were cut into small pieces with typical size of 1.5 cm × 0.5 cm × 0.5 cm (Fig. 1a). These pith were

firstly subject to pyrolysis at 600 °C for 90 min under nitrogen atmosphere to get carbonized products (Fig. 1a). In the following chemical activation step, 0.12 g of carbonized products were immersed into 20 mL water containing 0.6 g of KOH. The mixture was magnetically stirred for 24 h, followed by evaporation of water at 80 °C. Subsequently, the obtained carbon/KOH mixture was placed in a nickel boat and then subjected to chemical activation at 800 °C for 90 min. After washing with 1.0 M HCl and copious water to remove residual impurities, the black products were collected and dried at 80 °C for 10 h. They were donated as aCS-*x* with *x* representing the mass ratio of KOH to preliminarily carbonized cornstalk.

2.2 Characterization methods

The structure of the samples were examined by field-emission scanning electron microscopy (FESEM) on SU8020 and transmission electron microscopy (TEM) on a JEM-2100 TEM (JOEL) at acceleration voltage of 200 kV. Nitrogen adsorption/desorption isotherms were measured at 77 K on a micro-meritics ASAP 2460 analyzer. Samples were degassed at 180 °C for 8 h. The SSA of the samples was calculated using the Brunauer–Emmett–Teller (BET) method with the adsorption data in the relative pressure (*P/P*₀) range of 0.05–0.2. Total pore volumes were estimated at *P/P*₀ of 0.99. Pore size distribution (PSD) curves were derived from adsorption branch using the nonlocal density functional theory (NLDFT) model assuming the slit pore geometry. X-ray diffraction (XRD) measurements were recorded on a DX-2700 X-ray diffractometer with Cu Kα (*k* = 0.154 nm). X-ray photoelectron spectroscopy (XPS) spectra were measured on an AXIS ULTRA spectrometer (Kratos Analytical) using a monochromatized Al Kα X-ray source (1486.71 eV). Raman spectra were collected on a Renishaw inVia Raman microscope with an excitation wavelength of 532 nm.

2.3 Electrochemical measurements

The electrochemical performances of the electrode materials were characterized by cyclic voltammetry (CV), galvanostatic

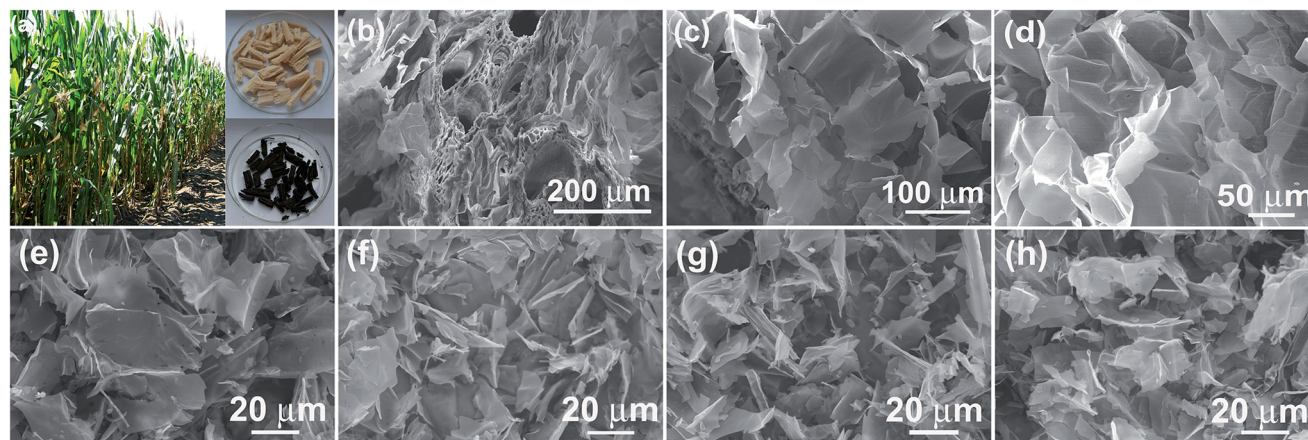


Fig. 1 (a) Digital photo of pith separated from the dried cornstalk and the carbonized pith. (b–d) SEM image showing the interior structure of pith. (e–h) SEM images of aCS (e), aCS-4 (f), aCS-5 (g) and aCS-6 (h).



charge–discharge and electrochemical impedance spectroscopy (EIS) on Gamary Reference 3000 electrochemical workstation. Both 6.0 M KOH and 1.0 M Na₂SO₄ aqueous electrolyte were used as electrolyte. The working electrode was prepared by mixing an active material (90 wt%) with carbon black (5 wt%) and polytetrafluoroethylene (5 wt%) in water. The slurry of the mixture was painted between two pieces of nickel foam (1.2 cm²) and then pressed under a pressure of 100 kg cm⁻². The mass loading for each electrode is typically of 1.0 mg cm⁻². In a two electrode cell, a Swagelock-type capacitor was configured with non-woven fabric as the separator. The specific capacitance was calculated according to the equation:³⁸ $C_s = 4 \times I \times \Delta t / (\Delta V \times m)$, where I , Δt and ΔV are discharge current (A), discharge time (s) and the voltage change (V) excluding the IR drop during the discharge process, respectively. Whereas, m is the total mass of the active material (g) in two electrodes. The ion kinetics within an electrode material is investigated by electrochemical impedance spectroscopy (EIS) with an amplitude of 10 mV at the frequency range of 0.005 Hz to 100 kHz. Energy and power density of a supercapacitor were determined by using $E = 0.5 \times CV^2/3.6$ and $P = 3600 \times E/t$, where E and P are energy (W h kg⁻¹) and power density (W kg⁻¹), C is the specific capacitance measured using a two-electrode cell (F g⁻¹), and t is the discharging time (s), respectively.^{39,40}

2.4 Fabrication of solid-state supercapacitor

In fabricating a solid-state supercapacitor, the commercially available carbon cloth (CC) was used as the current collector. Typically, the aCS-5 slurry of 1.5 mg was coated onto CC of 1.0 cm², and then dried at 60 °C for 3 h. Afterward, two pieces of aCS-5-coated CC was immersed into the KOH/PVA sol for 3 min, followed by solidification at room temperature for 1 h. A solid-state capacitor was then assembled by sandwiching two electrodes with Celgard@3501 and sealed with parafilm film. The areal capacitance was calculated according to the equation: $C_s = 4 \times I \times \Delta t / (\Delta V \times S)$, where S is the electrode area coated on the CC.

3. Results and discussion

Fig. 1a shows the digital photo of the lightweight pith separated from the dried cornstalk. The sectional SEM image shows that the pith is naturally formed by many porous texture with macropore size over tens of micrometers. These macropores provide numerous channels for the efficient access of activation agent into the inner of aCS for subsequent chemical activation. A SEM image taken on the pith surface reveal that it is actually composed of many cellulose nanosheets which display disordered and curved structure. The lateral size of the individual cellulose nanosheets varies in the range of 50 to 100 μm (Fig. 1c and d). These sponge-like pith is used as the starting materials for preparation porous carbon sheets. It is noted that our experiment results show that a higher temperature of 850 °C with longer time (90 min) would yield little products, while lower temperature of 700 °C leads to product with lower specific surface area due to the insufficient activation process. Fig. 1e–h

shows the SEM images of the resultant carbon products prepared by chemical activation of aCS at 800 °C for 90 min with various amount of KOH. It is found that the resulting aCS- x breaks into smaller pieces with the increase of the KOH activation agents. This observation suggests the successive reaction between carbon atoms and KOH proceeding as:⁴¹ $6\text{KOH} + \text{C} \rightarrow 2\text{K} + 3\text{H}_2 + 2\text{K}_2\text{CO}_3$. The progressive reaction of carbon atoms with KOH results in the breakage of large sheets into small ones. Fig. 2 shows the SEM and TEM images of aCS-5 sample. Like the initial aCS without activation treatment, the aCS-5 is composed of carbon sheets with lateral size of ~50 μm and thickness around 100 nm (Fig. 2a and b and inset). TEM image shown in Fig. 2c reveals that the aCS-5 is mainly made up of abundant micropores. A HRTEM image (Fig. 2d) presents an interlayer spacing of ~0.39 nm, which is larger than 0.34 nm of pure graphite, thus suggesting a disordered arrangement of carbon layers in aCS-5.

The structure of aCS was characterized by powder XRD and Raman spectroscopy. As shown in Fig. 3a, all aCS samples present broad and weak diffraction peaks at $2\theta = 23.7^\circ$, which can be attributed to the (002) reflections of graphitic carbon.⁴² The less ordered graphitic structure in aCS- x sample as compared with previous cornstalk-derived carbon is clearly due to the absent of Fe or Ni catalysts during the activation step.^{35,36} Raman spectroscopy shown in Fig. 3b presents two bands at 1340 cm⁻¹ (D band) and 1594 cm⁻¹ (G band) for each sample. The former band is related to the defective or disordered sp³ carbon atoms, while the latter is associated with the sp²-hybridized carbon atoms in graphitic layers.²² The defect density of carbon materials, as manifested by the intensity ratio of D band to G band (I_D/I_G),⁴⁰ is determined to be 0.94 for pristine aCS and ~1.0 for aCS- x samples after KOH activation (Table S1†). The surface element composition of aCS and aCS-5 probed by XPS gives only C 1s and O 1s signals (Fig. S1a†), suggesting the high purity of the products. The deconvoluted C 1s spectra of aCS and aCS-5 show four peaks. They can be ascribed to the C–C (284.6 eV), C=O (285.6 eV) and –COOH (288.6 eV) bonds, respectively (Fig. S1b†). XPS quantitative

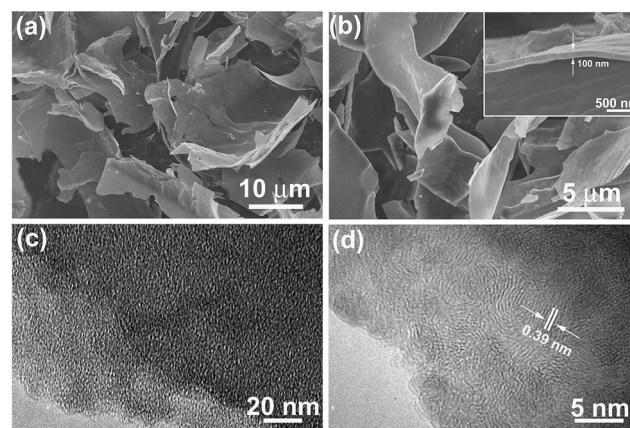


Fig. 2 SEM (a and b) and TEM images (c and d) of aCS-5 with different magnifications. Inset in panel (b) showing the thickness of individual aCS-5.



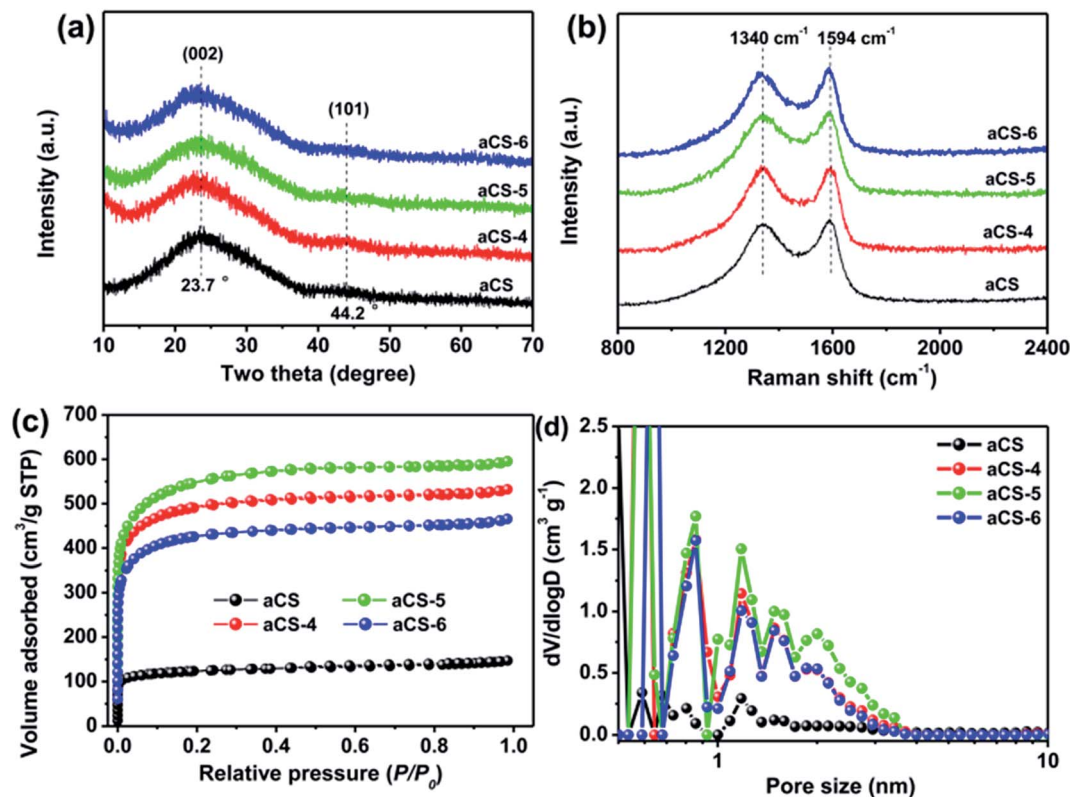


Fig. 3 Powder XRD pattern (a), Raman spectra (b), N₂ adsorption isotherms (c), and corresponding NLDFT pore size distribution (d) of aCS and aCS-*x* samples.

analysis reveals that the O atomic concentration increases from 3.1% of aCS to 9.1% of aCS-5, in consistence with more edge defects caused by KOH activation. The progressive oxidation process could leave behind many vacancies that extend into abundant micropores in aCS, as confirmed by N₂ adsorption-desorption measurement in Fig. 3c. All the isotherms display the characteristic type I adsorption curves, meaning existence of abundant micropores in the carbon products.⁴³ However, as compared with pristine aCS, aCS-*x* shows an dramatic increase of N₂ uptake, indicating a significantly enhanced surface area and pore volume. Fig. 3d shows the NLDFT pore size distribution. Compared with the nearly nonporous aCS, pore size in aCS-*x* products distributes mainly in the range of 1.0–4.0 nm, in good accordance with the TEM in Fig. 2c and d. Depending on the amount of KOH utilized in the activation step, the SSA of aCS-*x* varies from 1331 to 1736 m² g⁻¹ (Table S1†), which is much higher than 388 m² g⁻¹ of pristine aCS without activation treatment, confirming an important role of KOH activation agents in etching carbon atoms to create numerous nanopores on aCS.

The electrocapacitive performance of the aCS-*x* samples in 6.0 M KOH electrolyte was investigated by CV and galvanostatic charge-discharge in a three-electrode cell. As can be seen in Fig. S2a,† all the aCS-*x* electrodes present quasi-rectangular CV profiles with a broad redox peak occurring in the potential range from -0.9 to -0.5 V (vs. Ag/AgCl), suggesting predominated double-layer capacitance due to ion adsorption and

additional pseudocapacitance arising from the interfacial reactions between KOH and surface oxygen-containing functional groups.⁴⁴ Based on the symmetric charge-discharge curves at constant current density of 2.0 A g⁻¹ (Fig. S2b†), the aCS-5 electrode has the largest discharge time, an indicative of best charge storage capability among all the electrodes investigated. The specific capacitance of all aCS-*x* electrodes at varying current densities are plotted in Fig. S2c.† The aCS-5 electrode not only achieves the highest capacitance of 301 F g⁻¹ at 2.0 A g⁻¹, but also exhibits excellent rate capacity with capacitance retention of 82% at 60 A g⁻¹ (247 F g⁻¹). These electrocapacitive performances are superior to previous cornstalk-derived carbon,^{35,36} and other biomass-derived carbon electrodes in term of the specific capacitance or the rate capability.^{18,45–49}

To demonstrate the superior performance of aCS-5, we fabricated a Swagelok-type capacitor using aCS-5 as symmetric electrodes. Fig. S3† shows the CV profiles of capacitor over a wide scan rates in different aqueous electrolytes. Interestingly, the capacitor in KOH still remains an ideally rectangular CV profile even at a fast scan rate of 1.0 V s⁻¹ (Fig. S3b†). This result is in sharp contrast to the conventional activated carbon electrodes which are characteristic of tortuous micropores with micrometer-sized ion diffusion length, thus limiting the effective ion transport and lowering the capacitance performance. Our results clearly shows that in spite of dominant micropores, the aCS-5 electrode still exhibits remarkably high rate capability



due to its unique sheet-like structure, enabling aCS-5 to have more exposed external surface and substantially reduced ion pathway.²³ Fig. 4a compares the CV curves of aCS-5-based capacitor at scan rate of 100 mV s^{-1} in the two electrolytes. In contrast to the alkaline KOH electrolyte, the capacitor in neutral Na_2SO_4 electrolyte has relatively narrowed CV loop, suggesting a lower capacitance in Na_2SO_4 electrolyte. Whereas, the operating cell voltage in Na_2SO_4 electrolyte can extend up to 1.6 V without a significant increase of the anodic current at both low and high scan rate (Fig. 4a and S3c†), suggesting that the electrolyte is quite stable without obvious decomposition in the range of applied voltage due to the high over-potential for dihydrogen evolution.^{12,50} The galvanostatic charge–discharge curves of aCS-5-based capacitors in two electrolytes are shown in Fig. 4b. Both curves display symmetric charge and discharge profiles which are characteristic of double-layer capacitance. The specific capacitance of aCS-5 at various current densities is plotted in Fig. 4c. At the low current density of 0.5 A g^{-1} , aCS-5 exhibits approximately the same capacitance value, 234 F g^{-1} in KOH and 231 F g^{-1} in Na_2SO_4 electrolyte. However, at higher current density, capacitance in Na_2SO_4 electrolyte drops markedly. This behavior agrees well with the CV profiles recorded at high scan rate (100 mV s^{-1}) and indicates that Na_2SO_4 electrolyte with larger hydrated ions (3.59 \AA for Na^+ and 7.33 \AA for SO_4^{2-}) and high desolvation energy (108 kJ mol^{-1} when desolvating one H_2O molecule from SO_4^{2-})⁵¹ is difficult to access into the small micropores of aCS-5 electrode, leading to less contribution of double-layer capacitance at high scan rates.

The diffusion kinetics of electrolyte ions within the nanopores of aCS-5 were also investigated by EIS. Fig. 4d shows the Nyquist plots of aCS-5-capacitor in KOH and Na_2SO_4 electrolytes. The cell displays a vertical line in low-frequency range, indicating ideal double-layer capacitive behavior in both electrolytes. Whereas, in the middle- and high-frequency range (inset in Fig. 4d), capacitor in Na_2SO_4 shows a larger solution resistance (R_s) and longer Warburg region. This observation is attributed to the lower ion conductivity of Na_2SO_4 ($50 \text{ vs. } 540 \text{ mS cm}^{-1}$ of KOH)^{52,53} and larger hydrated ions of Na^+ and SO_4^{2-} , which may suffer from relatively larger resistance when accessing into the nanopores of aCS electrode. To quantitatively describe the capacitive behavior of aCS-5 in the two electrolytes, the dependence of phase angle on the frequency are shown in Fig. S4.† The relaxation time constant τ_0 ($\tau_0 = 1/f_0$ at a phase angle of -45°), marking the point where the capacitive impedance is equal to resistive impedance,⁵⁴ can be extracted from Fig. S4.† It is 0.82 s for KOH and 2.08 s for Na_2SO_4 electrolyte. Note the time constant of aCS-5 in KOH electrolyte is even comparable to that of chemically activated graphene aerogel (0.73 s),⁴⁰ and graphene nanomesh (0.47 s),⁵⁵ but much smaller than that of commercial activated carbon (2.45 s for RP20).⁷ These results are indicative of the high frequency response of aCS-5 electrode due to its unique sheet-like structure which really promotes fast ion diffusion into the interior electrode.

The cycling stability is another important character which is highly desirable for practical application. Fig. 4e shows the cycling performance of the aCS-5-capacitor operating in KOH and Na_2SO_4 electrolytes. The aCS-5-based supercapacitor

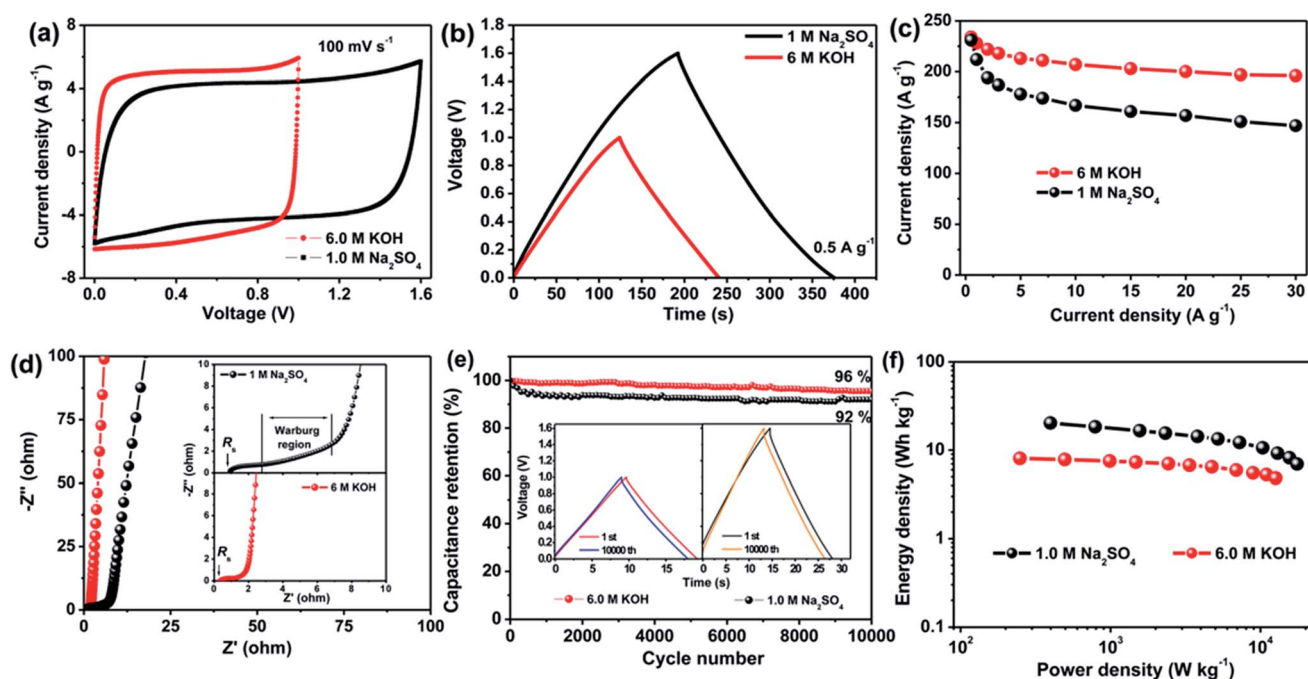


Fig. 4 Performance of aCS-5-based capacitor in 6.0 M KOH and 1.0 M Na_2SO_4 . (a) CV curves at scan rate of 100 mV s^{-1} , (b) galvanostatic charge–discharge profiles at current density of 0.5 A g^{-1} , (c) capacitance retention in current density range of $1\text{--}30 \text{ A g}^{-1}$, (d) Nyquist plots with inset showing the plots in high-frequency range, (e) cycling stability of aCS-5-based capacitor at 5.0 A g^{-1} with inset showing the charge–discharge curves of the 1st and the 10 000th cycles, (f) Ragone plots of aCS-5-based capacitor in the two electrolytes.



retains 96% and 92% of its initial capacitance in KOH and Na_2SO_4 electrolyte, respectively, after 10 000 cycles of continuous charge–discharge at constant current density of 5.0 A g^{-1} . The nearly identical charge–discharge curves of the first and last cycle along with the negligible changes of the voltage drop (IR drop) (inset in Fig. 4e and S5†) provide further evidences that the capacitor does not show noticeable increase in the internal resistance, confirming the electrochemically stable Na_2SO_4 electrolyte in the voltage range of 0–1.6 V. Such an extended cell voltage enable the aCS-5-capacitor to deliver an energy density as high as 20.2 W h kg^{-1} at power density of 398 W kg^{-1} , which is higher than 8.1 W h kg^{-1} (249 W kg^{-1}) of the counterpart in KOH electrolyte (Fig. 4f). Besides delivering a high energy, another merit of using Na_2SO_4 electrolyte is that it has a higher ion conductivity as compared with most of organic electrolyte. This inherent character ensures a high power capability. For example, at power density of 17.6 kW kg^{-1} , an energy density as high as 7.0 W h kg^{-1} can still be delivered.

The 2D structure of aCS-5 with large aspect ratio (200–500) enables us to fabricate a flexible solid-state capacitor (SSC) with aCS-5 as electrolyte and PVA–KOH as gel electrolyte. The SSC was assembled by coating aCS-5 slurry on commercial carbon cloth (CC), followed by immersing the CC-aCS-5 electrode into the PVA/KOH sol and solidification at room temperature. Fig. 5a shows the CV profile of SSC over a wide range of scan rates (100–500 mV s^{-1}). Interestingly, the CV curve presents a quasi-rectangular shape even at a high scan rate 500 mV s^{-1} . Considering the very high scan rate for SSC and the low ion conductivity of PVA/KOH electrolyte, the observed unusual CV behavior clearly reveals that sheet-like

structure of aCS-5 is particularly beneficial for fast ion transport. Like the capacitive behavior in aqueous KOH electrolyte, the aCS-5-based SSC stores charges through double-layer capacitance as evidenced by the symmetric and triangular profiles of the charge–discharge curves (Fig. 5b). The areal capacitance of aCS-5 is measured to be 136 mF cm^{-2} (142 F g^{-1}) at 0.5 mA cm^{-2} , and it decreases to 84 mF cm^{-2} as the current density increase to 20 mA cm^{-2} (Fig. 5c). The areal capacitance reported herein is higher than 43.4 mF cm^{-2} of functionalized carbon fabric,⁵⁶ 130 mF cm^{-2} of graphene/carbon nanotube film,⁵⁷ 87 mF cm^{-2} of rGO/carbon fabric,⁵⁸ and even comparable to 138.7 mF cm^{-2} of ZnO/ MnO_2 /CC hybrid.⁵⁹ Moreover, the aCS-5 based SSC remains 92% of its initial areal capacitance after 1000 cycles of charge–discharge at 5 mA cm^{-2} (inset in Fig. 5c).

In addition to the high rate performance and satisfying areal capacitance, the aCS-5 SSC also displays good flexibility as confirmed by the nearly identical CV curves measured under normal, bend and twisted state (Fig. 5d and inset). To demonstrate the practical application of SSC, two identical capacitors were fabricated and connected in series to improve the overall voltage output. The CV curves for single cell and two cells connected in series are present in Fig. 5e. As compared with the single cell, two cells in series exhibit the similar rectangular CV profile and charge–discharge behavior except the cell voltage extended to 2.0 V and the current density reduced by 50% (Fig. 5e and f). The two cells connected in series could easily light up a red light-emitting-diode (LED), demonstrating the potential of the aCS-5 electrode materials for electrochemical energy storage.

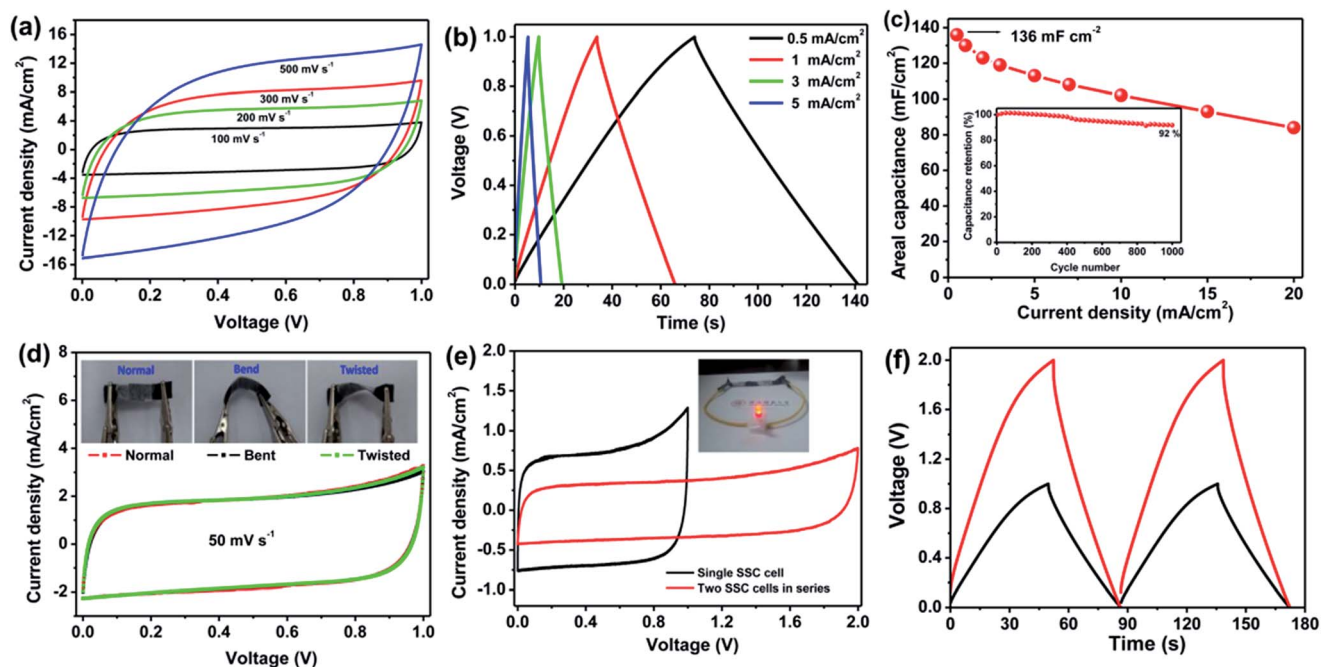


Fig. 5 Performance of aCS-5-based SSC with PVA/KOH as polymer gel electrolyte. (a) CV profiles over a wide range of scan rates, (b) galvanostatic charge–discharge curves at different current densities, (c) dependence of areal capacitance on the current density with the inset showing the cycling stability measured at 5.0 mA cm^{-2} , (d) CV profiles measured under normal, bend and twisted states, (e) CV and (f) galvanostatic charge–discharge profiles of single SSC cell and two cells connected in series.



4. Conclusions

In summary, we have developed a facile yet cost-effective method for scalable production of 2D porous carbon nanosheets through carbonization and chemical activation of the pith naturally abundant in biomass cornstalk. The obtained carbon nanosheets are characterized by dominant micropores, thin thickness (~ 100 nm), controllable SSA and large lateral size (50–100 μm). These structure merits enable the carbon sheets to serve as promising supercapacitor electrodes in both KOH and Na_2SO_4 aqueous electrolyte. Our results also show that the capacitive performance of conventional microporous carbons which are limited by slow ion diffusion kinetics can be largely emended if their morphologies are properly designed to facilitate ion transport.

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