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Winter-jujube-derived carbon with self-doped heteroatoms and a hierarchically porous structure for high-performance supercapacitors

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Novel biomass-derived carbon materials have attracted a lot of attention for application in supercapacitors due to their low cost and environmental friendly properties. Herein, we report a novel winter-jujube-derived carbon material (JC) prepared via a pre-carbonization and KOH activation strategy. The physical and chemical properties of JC were characterized by field-emission scanning electron microscopy (FESEM), X-ray photoelectron spectroscopy (XPS), X-ray diffraction (XRD), Raman spectroscopy, and nitrogen adsorption-desorption isotherms. The electrochemical characteristics are measured using cyclic voltammetry (CV), galvanostatic charge-discharge (GCD), and electrochemical impedance spectroscopy (EIS). The as-obtained carbon skeleton possesses a hierarchical porous structure, abundant heteroatom groups, and good electrical conductivity. It is notable that the use of KOH significantly influenced the properties of JC. When the KOH to carbon weight ratio is 3.5 : 1, the as-prepared sample (JC-3.5) possesses an ultra-huge specific surface area (SSA) of 2286.2 m² g⁻¹. In addition, the electrochemistry results show that JC-3.5 exhibits a high specific capacitance up to 341 F g⁻¹ at 0.5 A g⁻¹ and a considerable rate capability of 76% retention when the current density increases to 20 A g⁻¹. After 5000 cycles at 5 A g⁻¹, the capacitance still has 96.5% retention. In addition, the JC-3.5-based symmetric supercapacitor device possesses an energy density of 30.9 kW h⁻¹ at 0.1 A g⁻¹ and power density of 8.9 kW at 5 A g⁻¹. Therefore, it is quite feasible to make the winter-jujube-derived carbon material into a high performance supercapacitor electrode material.

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1. Introduction

Since the 21st century, the global economy has continued to develop rapidly. At the same time, mankind has made great progress in technology and the quality of life, which has resulted in the increased consumption of resources and serious environmental pollution. There is an urgent need for green, efficient, renewable and advanced energy storage devices and conversion technologies. Currently, lithium ion batteries have gained a lot of attention as a new storage device of secondary batteries.¹⁻⁶ However, the low power density and short charge and discharge life limit the applications of lithium-ion batteries. Supercapacitors, as new energy storage devices among traditional capacitors and batteries, have attracted more and more attention due to their high power density, short charging time, long cycle life, wide operating temperature range, and environmental friendly characteristics.⁷⁻¹⁵ However, unsatisfactory capacitance is the biggest drawback of most

commercial supercapacitors.¹⁶ In order to provide a continuous energy supply, it is necessary to improve the capacitance of a supercapacitor.

There are various ways to improve the capacitance of supercapacitors, and the key issue is the selection and preparation of the electrode materials.¹⁷ Currently, there are two main types of supercapacitors namely faradaic pseudocapacitors, and electric double-layer supercapacitors (EDLCs), which possess different electrode materials. Faradaic pseudocapacitors are composed of transition metal oxide or conductive macromolecule electrodes and usually have a large capacitance. However, as the chemical reactions proceed, there are often irreversible components involved in the electrode reaction, resulting in poor cycling stability. The electrodes of double-layer supercapacitors (EDLCs) are usually constructed from porous activated carbon. The physical energy storage process makes its cycling stability very good. However, their capacitance still needs improvement.¹⁶

It is an important task to combine the advantages of these two supercapacitors, such as the higher energy density and cycling stability. One method is to use an asymmetric hybrid supercapacitor system, that is, one electrode using activated carbon and the other electrode using a pseudo-capacitance

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electrode material or battery electrode material. Another way is to increase the capacitance of the supercapacitor electrode materials, and researchers have carried out a lot of work in this area. Mu *et al.* dispersed Fe_3O_4 nanosheets onto one-dimensional (1D) carbon nanofibers (CNFs) *via* an electrospinning technique and solvent-thermal process.¹⁸ Zhang *et al.* prepared graphene and polyaniline (PANI) nanofiber composites *via* the *in situ* polymerization of an aniline monomer in the presence of graphene oxide under acidic conditions.¹⁹ However, the preparation of carbon materials such as graphene, carbon fibers and carbon nanotubes is always very complex. In addition, the high cost is the biggest drawback for their practical application. In addition, the preparation processes are not environment friendly.

Biomass carbon materials, which are easily accessible, renewable, sustainable and environment friendly, can be simply prepared using carbonization or pyrolysis. Biomass materials usually have an ordered structure, and KOH-activated biomass carbon material can have an ultra-high specific surface area. Moreover, the considerable heteroatom self-doping (such as N, O, and P) makes it a great candidate for supercapacitor electrode materials. Currently, various types of biomass materials have been used as carbon materials.^{20–23} Huang *et al.* produced activated carbon fibers from recycled fir sawdust using steam-KOH reactivation.²⁴ Its specific capacitance is 242 F g^{-1} at 0.5 A g^{-1} due to its mediocre structure and poor heteroatom content. Liang *et al.* synthesized a three-dimensional (3D) honeycomb-like porous carbon (HLPC) using the simple carbonization of pomelo peel.²⁵ The excellent porous structure provides HLPC a high specific capacitance up to 342 F g^{-1} at 0.2 A g^{-1} . However, the low heteroatom content hinders its capacitance from being enhanced. Wei *et al.* made NiCo_2O_4 nanowires grow on a mollusc shell-based macroporous carbon material (MSBPC).²⁶ The obtained NiCo_2O_4 /MSBPC composites showed an ultra-high specific capacitance up to 1696 F g^{-1} at 1 A g^{-1} ; however, the major drawback is external doping, which limits the cycling stability (12% capacitance loss after 2000 cycles) and rate capability (24.9% capacitance remained at 50 A g^{-1}). For the purpose of improving the comprehensive performance of supercapacitors, we need materials that possess considerable self-doped heteroatoms and excellent hierarchical porous structures with large specific surface areas.

Winter-jujube is an abundant and renewable source with plenty of O, N and trace elements. In this study, winter-jujube-derived carbon materials (JC) were prepared *via* a pre-carbonization and KOH activation process. This activated carbon possesses abundant heteroatom (O and N) groups. These groups give JC good electrical conductivity and enable JC to not only provide electrical double-layer capacitance, but also faradaic pseudocapacitance, which combines the contributions from EDLCs and faradaic pseudocapacitors. In addition, KOH activation gave JC an excellent hierarchically porous structure. When the KOH to carbon weight ratio was 3.5 : 1, the as-prepared sample (JC-3.5) possessed an ultra-high specific surface area of $2286.2 \text{ m}^2 \text{ g}^{-1}$ and a high capacitance up to 341 F g^{-1} at 0.5 A g^{-1} . It is quite promising to make JC a candidate carbon source to fabricate biomass-derived carbon

based supercapacitors. In addition, the application of this green resource in supercapacitors is also beneficial for environmental protection and energy demand.

2. Experimental

2.1 Materials

Fresh winter-jujubes were purchased from a local fruit store. The producing place was the Shandong Province of China. The winter-jujubes were washed clean before use without further treatment. KOH and HCl solution were provided by Beijing Chemical Work.

2.2 Preparation of winter-jujube-derived carbon (JC)

JC was prepared using a pre-carbonization and KOH activation strategy. In a typical process, a few winter-jujubes were first freeze-dried using a lyophilizer for about three days. Then, the dry winter-jujubes were placed in a corundum boat to be pre-carbonized in a tube furnace. The pre-carbonization temperature was 400°C at a heating rate of 5°C min^{-1} under a flow of N_2 . The resulting black solid block material was then mixed with a KOH aqueous solution in a beaker with a magnetic stirrer. After that, the black mixture was heated to 120°C until water was removed. Then, the remaining black solid was placed in the tube furnace again to be carbonized. With the protection of N_2 , the procedures were as follows: it was first heated to 450°C , and this temperature was held for 30 minutes; then, the temperature was increased up to 650°C and held for another 30 minutes; finally, the temperature was increased to 800°C and held for 1 hour. After the carbonization procedure, the sample was cooled naturally under a flow of N_2 . Hydrochloric acid solution was used to remove the impurities produced during the carbonization step and the pH of the mixture was adjusted to be weakly acidic. The resulting mixture was enclosed carefully in a dialysis bag. Then, the bag was immersed into deionized water to remove the impurities. A series of samples with different ratios of KOH to carbon weight were prepared to investigate the effect of KOH on the micro-structure and electrochemical performance of the as-prepared JC. We chose varying KOH to carbon weight ratios of 1.5 : 1, 2.5 : 1, 3.5 : 1 and 4.5 : 1, and the corresponding JC samples were named as JC-1.5, JC-2.5, JC-3.5 and JC-4.5, respectively.

2.3 Characterization

The morphology of the pre-carbonized and KOH-activated winter-jujube was analyzed *via* field-emission scanning electron microscopy (FESEM, Hitachi S-4800). The structure of JC was investigated using wide-angle X-ray diffraction (WXR, D/MAX-2500, Rigaku Co.) with 2θ values of $5\text{--}90^\circ$ at a scanning rate of 5° min^{-1} and Raman spectroscopy (inVia Reflex, Renishaw Co.) with an excitation wavelength of 532 nm. X-ray photoelectron spectroscopy was performed on a Thermo-Fisher Scientific Escalab 250 using a monochromatic Al K α source. The porous structure was measured using nitrogen adsorption and desorption isotherms on a JWGB SCI. & TECH. Co. BK200C analyzer. The specific surface area (SSA) was calculated using the Brunauer–Emmet–Teller (BET) equation. The pore-size distribution (PSD) was obtained using the



Horvath–Kawazoe (HK) method (<2 nm) and the Barrett–Joyner–Halenda (BJH) method (>2 nm) using the nitrogen adsorption data. Galvanostatic discharge–charge and cyclic stability tests were performed on a Land ct2001a supercapacitor tester. Cyclic voltammetry (CV) and impedance spectroscopy (EIS) were measured on an electrochemical workstation from Autolab PGSTAT 302N, Metrohm, Netherlands. A standard three-electrode system in $1 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$ aqueous solution was adopted for the electrochemical test. The working electrode was a mixture of JC and poly(vinylidene fluoride) (PVDF) at a mass ratio of 9 : 1 smeared on a platinum foil. The reference electrode was an Ag/AgCl electrode, and the counter electrode was a blank platinum foil. Two-electrode system (supercapacitors device) tests were conducted in $1 \text{ mol L}^{-1} \text{ H}_2\text{SO}_4$ aqueous solution with two nearly identical (by weight and size) JC electrodes and a porous polymeric separator permeable to ion transport. The specific capacitance (C_s) based on the GCD measurement was calculated using the following equation:

$$C_s = \frac{I \times \Delta t}{\Delta V \times m}$$

where I (A) is the charge/discharge current, Δt (s) is the discharge time, ΔV (V) is the voltage change excluding the IR drop during the discharge process, and m (g) is the mass of active material.

The energy density (E , W h kg^{-1}) and power density (P , W kg^{-1}) were calculated using the following equations:

$$E = \frac{1}{2} C_s \Delta V^2$$

$$P = \frac{E}{\Delta t}$$

where C_s (F g^{-1}) is the specific capacitance of the JC supercapacitors, ΔV (V) is the voltage change excluding the IR drop during the discharge process, and Δt (s) is the discharge time.

3. Results and discussion

Fig. 1 shows the surface morphology of the pre-carbonized (Fig. 1a–c) and KOH activated winter-jujube (Fig. 1d–g).

Despite using different KOH-carbon weight ratios, each sample displayed a similar morphology. Fig. 1a is the microstructure of the pre-carbonized winter-jujube. It demonstrates that abundant micrometer pores exist in the JC structure (ranging from 20 to $100 \mu\text{m}$), which may be attributed to the inherent property of winter-jujube. Upon further magnification, it revealed smaller pores sized a few tenths to a $1 \mu\text{m}$ (Fig. 1b) inside of the above-mentioned large pores. In addition, Fig. 1c reveals that the margin and interior of the pores are smooth. After the KOH activation process, plenty of new micropores appeared (Fig. 1d–g), which indicated that the specific surface area of JC had increased. To confirm this assumption, the pore properties of JC were determined using nitrogen adsorption and desorption measurements. Fig. 2a shows the adsorption–desorption isotherms for all four JC samples. The typical hysteresis loop at $P/P_0 > 0.4$ in each sample indicates a combination of Type I and IV isotherms, which indicate the existence of micropores and mesopores. Moreover, the slight rise in high relative pressure ($P/P_0 = 0.95\text{--}1.0$) and the sharp rise in low relative pressure ($P/P_0 = 0.01$) revealed the presence of macropores and

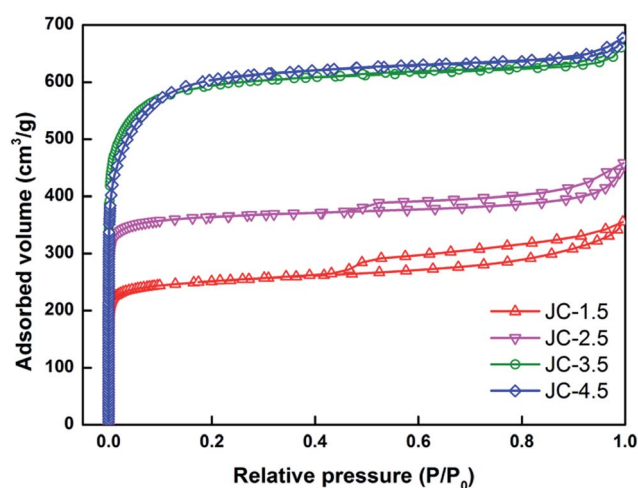


Fig. 2 The nitrogen adsorption–desorption isotherms of the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples.

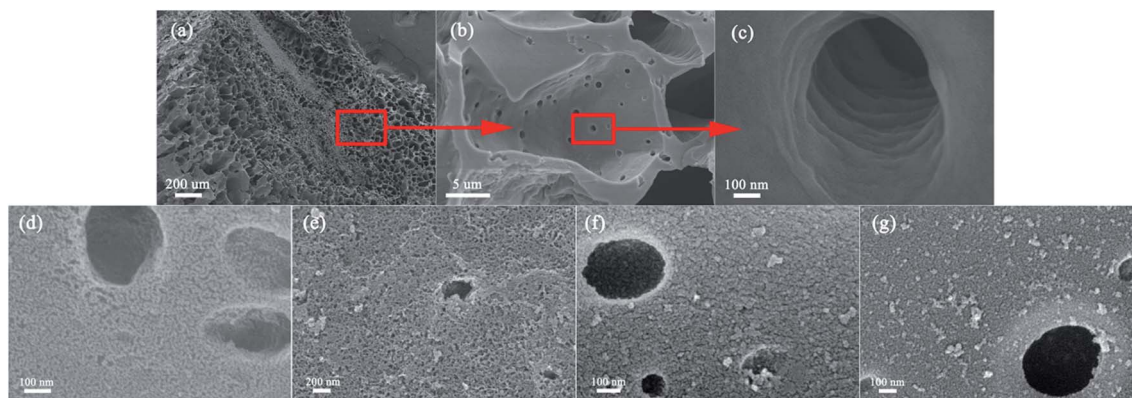


Fig. 1 SEM images of pre-carbonized winter-jujube (a–c) and KOH activated JC-1.5 (d), JC-2.5 (e), JC-3.5 (f) and JC-4.5 (g).



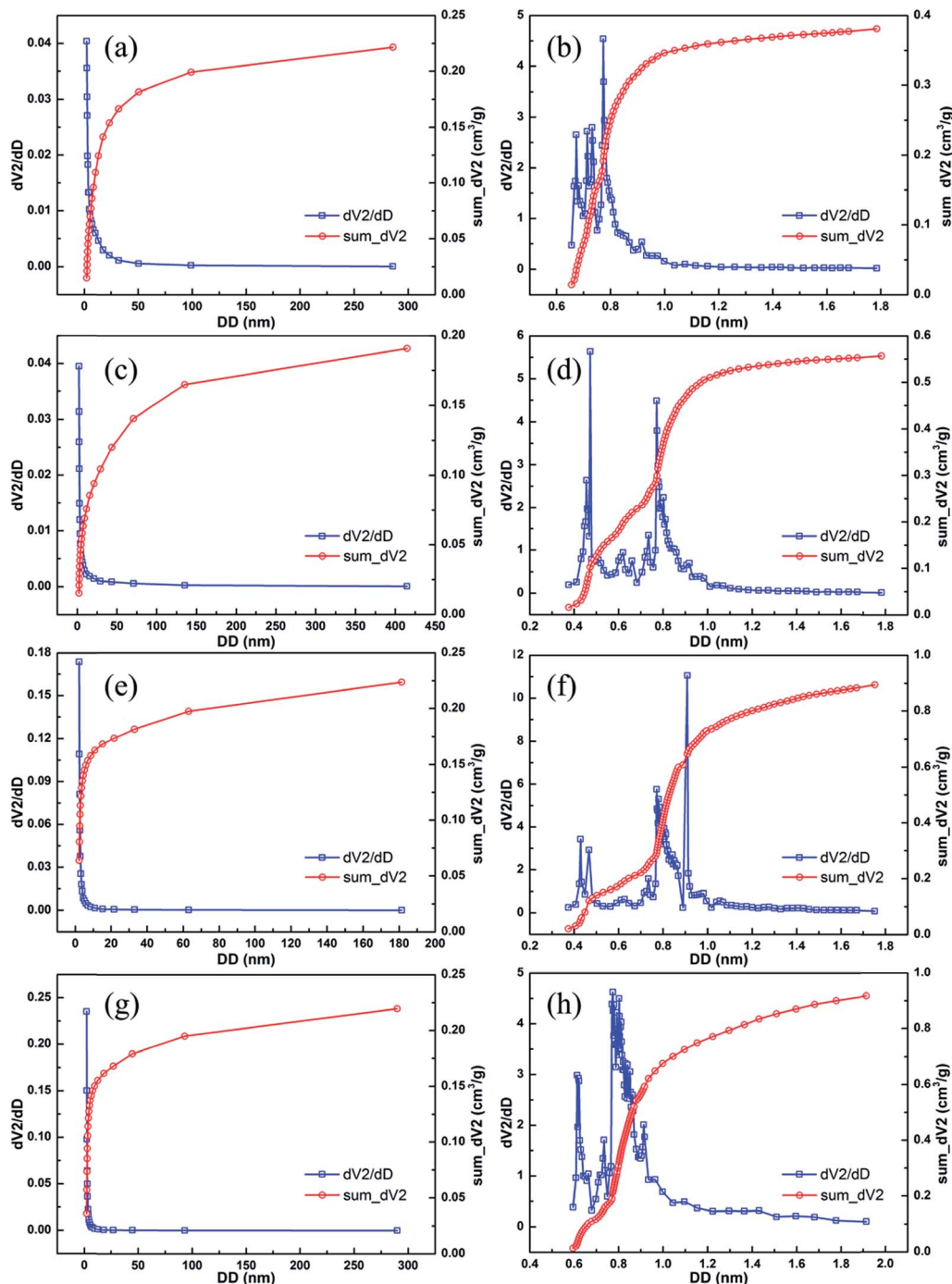


Fig. 3 The pore size distribution and integration in the pore-size ranges of 0–2 nm and 2–400 nm of the JC-1.5 (a and b), JC-2.5 (c and d), JC-3.5 (e and f) and JC-4.5 (g and h) samples.

micropores, respectively.^{27–29} Fig. 3 shows the pore size distribution of JC. From Fig. 3, we can see the existence of micropores (less than 2 nm), mesoporous (2–50 nm), and macropores (more than 50 nm). These various pores constitute a hierarchically porous structure and significantly improve the electrolyte contact and accumulation of electrolyte ions.^{28,30} Table 1 shows the specific surface area and pore structure parameters of the samples, which confirm that KOH makes a vital contribution to

the formation of the pore structure. For these several samples, the SSA increased along with the consumption of KOH. JC-3.5 and JC-4.5 both reveal an ultra-high BET surface area of $2286.2 \text{ m}^2 \text{ g}^{-1}$ and $2303.2 \text{ m}^2 \text{ g}^{-1}$, respectively. This also shows that an excess KOH does not offer further assistance towards pore formation.

Further information on the structure of JC was obtained from X-ray diffraction (XRD) and Raman spectroscopy. The XRD



Table 1 The SSA and pore structure parameters of the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples

Sample	BET SSA (m ² g ⁻¹)	Average pore width (nm)	Pore volume (V _{total} , cm ³ g ⁻¹)	Micropore volume
JC-1.5	987	2.227	0.550	0.381
JC-2.5	1463	1.937	0.709	0.557
JC-3.5	2286	1.787	1.021	0.895
JC-4.5	2303	1.820	1.048	0.917

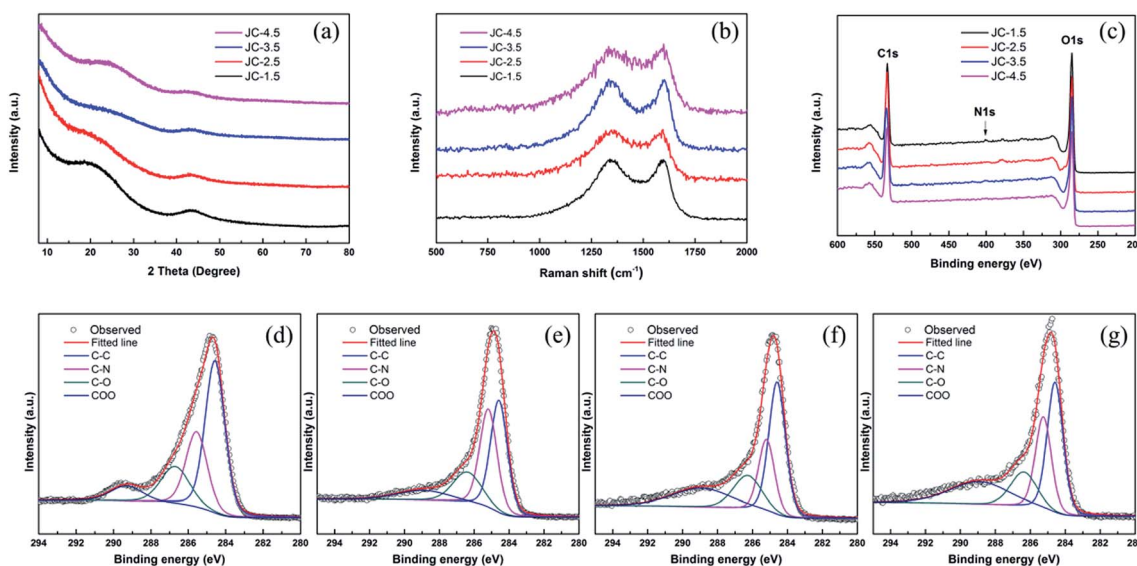
Table 2 The elemental composition of the JC-1.5, JC-2.5, JC-3.5, and JC-4.5 samples

Sample	XPS (at%)		
	C	O	N
JC-1.5	77.8	21.1	1.1
JC-2.5	73.0	25.6	1.3
JC-3.5	74.9	23.9	1.2
JC-4.5	73.5	24.9	1.6

pattern (Fig. 4a) of JC-1.5 shows two weak and broad peaks around $2\theta = 21$ and 43.5° , which correspond to the (002) and (100) planes diffraction of graphitic carbon. The broader peak around 21° was attributed to the amorphous carbon structure, which contributed to the high SSA. In addition, a large increase appears in the low angle area and confirms the microporous structure.³¹ The weak peak located at 43° suggests the existence of a graphitic-type carbon structure in JC. The intensity of the two peaks decreases with an increase in KOH, which reveals that an excess amount of KOH destroys the structure of JC. Simultaneously, the slight shift in the location of the (002) diffraction peaks also indicates the differences in the constituents of the four samples. The Raman spectra, shown in Fig. 4b, show two well-known peaks in carbon materials, which are the D-band located around 1339 cm^{-1} and G-band around 1589 cm^{-1} . In addition, the D-band represents the defects and disordered structure, while the G-band reveals the existence of graphitic structure in the material.³² As an indicator of the degree of graphitization, the D/G intensity ratios approximating to 1 reveal the amorphous characteristics of JC, which are beneficial for the conduction of electrons. The XRD and Raman results are consistent with the SEM results.

Furthermore, XPS was used to identify the elemental composition and surface functional groups of the as-synthesized JC. The results of the survey spectra (Fig. 4c) confirm that the main components of JC are C, O, and N on account of incomplete carbonization. The specific compositions are shown in Table 2. Fig. 4f shows the C1 high resolution spectrum of JC-3.5. The four peaks at 284.6 eV, 285.2 eV, 286.3 eV, and 288.9 eV correspond to C–C, C–N, C–O and COO, respectively.³³ The spectra observed for the three other samples are shown in Fig. 4. The XPS results verify the existence of O and N, and these heteroatoms can process faradaic reactions and increase the pseudo-capacitance.

To evaluate the electrochemical performance of the as-prepared JC applied in a supercapacitor, cyclic voltammetry (CV) and galvanostatic charge/discharge (GCD) measurements were conducted. Fig. 5 shows the CV curves of JC at different scan rates. The shapes of the CV curves are quasi-rectangular, demonstrating that JC behaves as an electrical double-layer capacitor with fast charge/discharge processes. In addition, the rapid growth of current appearing in the switching potential indicates an excellent mass-transfer ability. This is ascribed to the hierarchically porous structure shown in the pore size distribution. As we can see clearly in Fig. 6, the curve

**Fig. 4** The XRD pattern (a), Raman spectra (b) and XPS spectra (c) of the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples. The C1 high-resolution XPS spectra of the JC-1.5 (d), JC-2.5 (e), JC-3.5 (f) and JC-4.5 (g) samples.

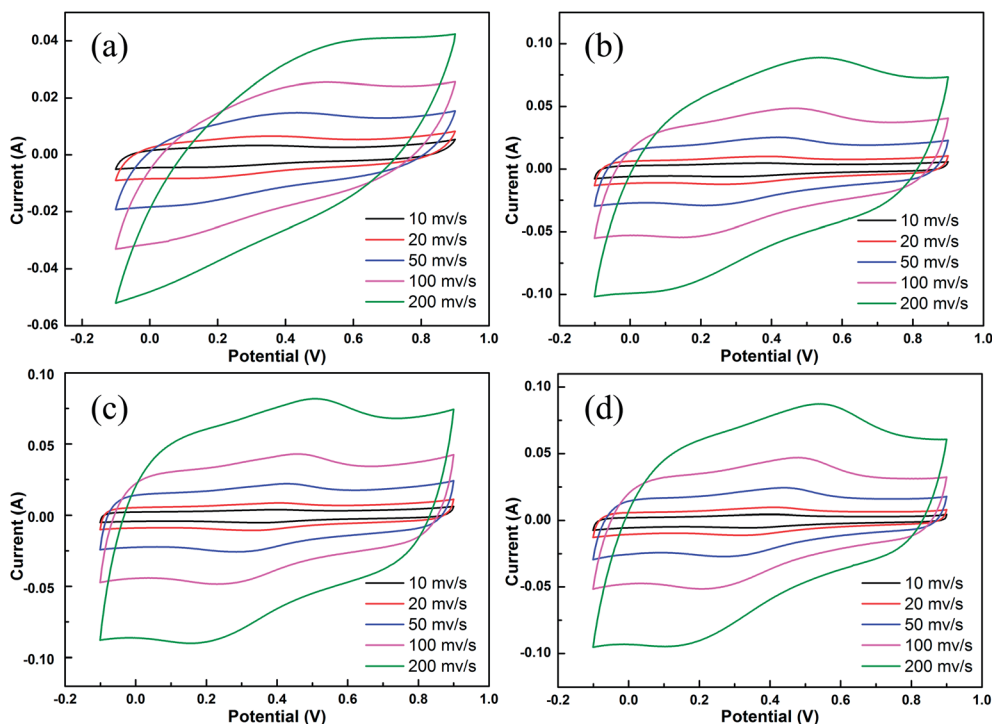


Fig. 5 The CV curves obtained for the JC-1.5 (a), JC-2.5 (b), JC-3.5 (c) and JC-4.5 (d) samples.

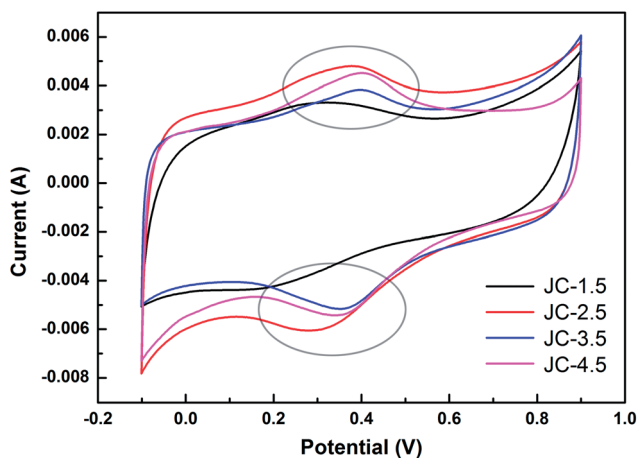


Fig. 6 The CV curves obtained at 10 mV s^{-1} for the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples.

at 10 mV s^{-1} scan rate, a redox reaction caused current increasing appears around 0.3 V in the CV curves. This was ascribed to the N and O functional groups on the surface, which were determined by the XPS spectra. Fig. 7 displays all the galvanostatic charge–discharge curves within a potential range of -0.1 to 0.9 V at different current densities. All of the curves possess a nearly perfect symmetrical triangular shape except a slight deformation. This suggests an electrical double layer capacitance characteristic combined with the existence of pseudocapacitive behavior, which is consistent with the conclusion of CV.

Fig. 8a summarizes the specific capacitances calculated from the CV curves and the influence of the current density to the specific capacitance of JC. The curve shows that the highest specific capacitance was up to 233, 274, 341 and 333 F g^{-1} at a current density of 0.5 A g^{-1} for the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples, respectively. The capacitance increases with an increase in the amount of KOH. When the KOH-carbon ratio was 3.5 : 1, the capacitance reached its highest value. Further increasing the amount of KOH did not increase the capacitance. When the current density was increased to 20 A g^{-1} , the capacitance was still retained at 155, 225, 260, and 250 F g^{-1} with 66.5, 82.1, 76.2, and 75.1% retention, respectively. Moreover, the charge–discharge cycle stability was also investigated at a current density of 5 A g^{-1} for 5000 cycles, and the results are shown in Fig. 8b. After the process, there was still 96.6, 98.8, 96.5, and 96.9% capacitance retention for JC-1.5, JC-2.5, JC-3.5 and JC-4.5, respectively, indicating their excellent cycling stability. Table 3 lists the properties of other biomass-derived carbon materials with JC as a comparison, which indicates the excellent rate capability and specific capacitance of JC.^{34–37}

Electrochemical impedance spectroscopy (EIS) was performed (Fig. 8c and d). A classical Nyquist plot was also obtained in a $1 \text{ M H}_2\text{SO}_4$ aqueous electrolyte using a three-electrode system. The frequency was investigated from 10^{-2} to 10^5 Hz . The Nyquist plots obtained for JC-2.5, JC-3.5, and JC-4.5 show nearly vertical lines in the low-frequency region, which corresponds to ideal capacitive behavior. Moreover, the 45° slope, which represents the ion diffusion ability, is very short to obtain efficient access of the electrolyte ions to the electrode. Corresponding to displaying the highest specific capacitance, sample



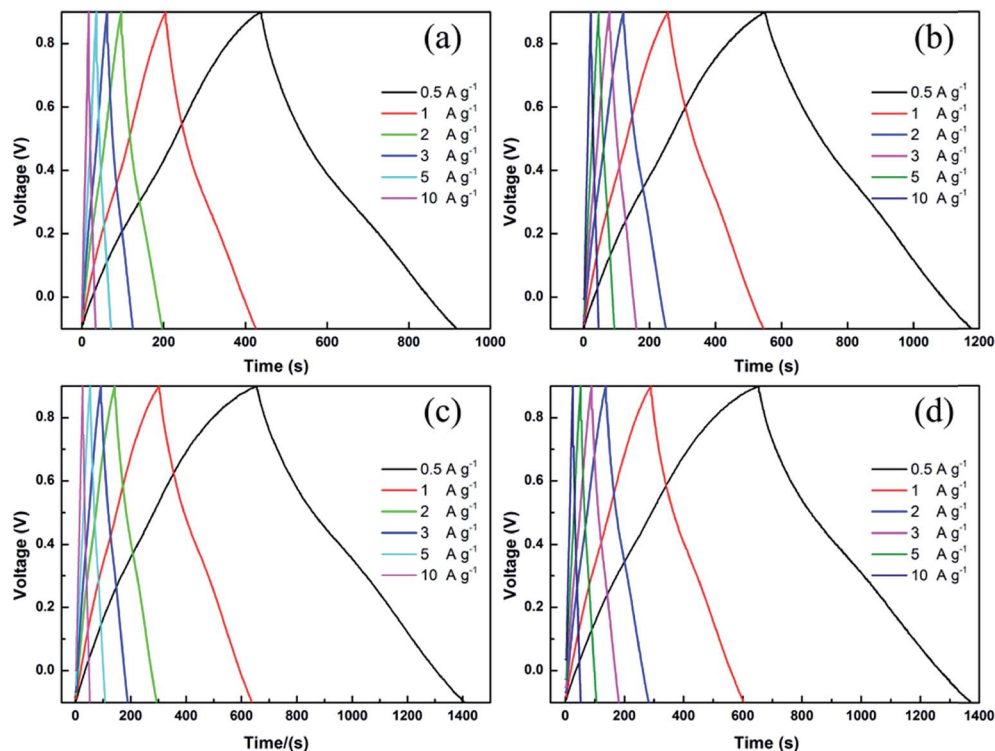


Fig. 7 The galvanostatic charge-discharge curves obtained at different current densities for the JC-1.5 (a), JC-2.5 (b), JC-3.5 (c) and JC-4.5 (d) samples.

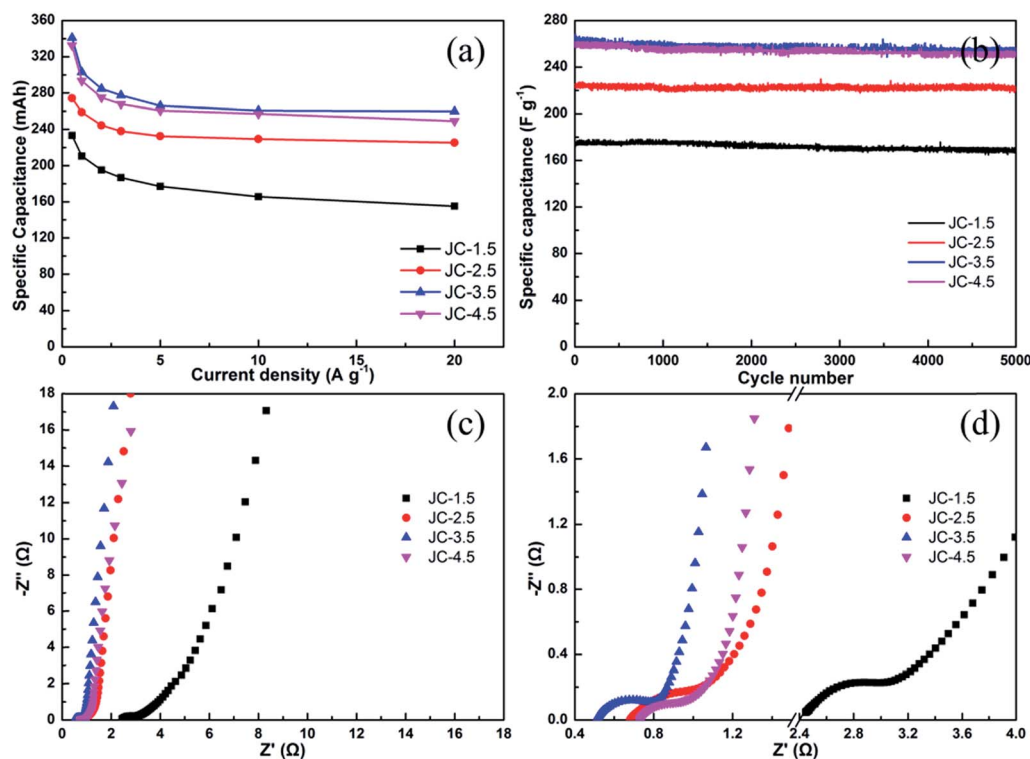


Fig. 8 The specific capacitance at various current densities (a), cycling performance at 5 A g⁻¹ (b) and Nyquist plot (c and d) obtained for the JC-1.5, JC-2.5, JC-3.5 and JC-4.5 samples.



Table 3 A comparison of the properties of other biomass-derived materials with JC

Source	Specific surface area ($\text{m}^2 \text{g}^{-1}$)	Specific capacitance (at current density) (F g^{-1})	Rate capability (from $a \text{ A g}^{-1}$ to $b \text{ A g}^{-1}$)	Ref.
Bamboo	171.5	318 in 1 M H_2SO_4 (0.2)	33.3% (0.2 to 4.0)	20
Peanut shell	1552	245 in 6 M KOH (0.05)	27.8% (0.2 to 20)	21
Sawdust	—	242 in 6 M KOH (0.5)	85.3% (0.5 to 5)	22
Tobacco rods	2115	286.6 in 1 M H_2SO_4 (0.5)	74% (0.5 to 30)	23
Pomelo peel	2725	342 in 6 M KOH (0.2)	62% (0.2 to 20)	25
Rice husk ash	786	260 in 6 M KOH (1)	65.4% (1 to 20)	34
Soybean curd residue	581	215 in 2 M KOH (0.5)	60.5% (0.5 to 10)	35
Tofu	1202	418 in 6 M KOH (1)	>83.7% (1 to 20)	36
Torreya grandis shell	2101	290.5 in 6 M KOH (0.5)	62.6% (0.5 to 20)	37
Winter jujube	2286	341 in 1 M H_2SO_4 (0.5)	76% (0.5 to 20)	This work

JC-3.5 had the smallest R_s values of 0.51Ω (the intercept of real axis), which means the smallest equivalent serial internal resistance. The curves show a semicircle in the high-frequency region, and the small semicircle diameter of JC-2.5, JC-3.5, and JC-4.5 indicate their small charge-transfer resistance. As with JC-1.5, its poor EIS performance when compared with the other samples matches its poor specific capacitance and CV curve.

The practical application of JC-3.5 was evaluated in a two-electrode system. Fig. 9a displays the CV curves obtained for JC-3.5 at different scan rates. The rectangular shape suggests its ideal electrochemical performance, which corresponds to the results obtained from the three-electrode system. EIS

measurement reveals a quite small R_s value of 0.09Ω (Fig. 9b). The high slope of the Nyquist plot in the low-frequency region also demonstrates its outstanding ion diffusion capability. The GCD curves (Fig. 9c) at different current densities were also measured to characterize the specific capacitance. As shown in Fig. 9d, the specific capacitance of JC-3.5 was calculated to be 61.9 F g^{-1} at 0.1 A g^{-1} and it was still maintained at 79.2% (49 F g^{-1}) at 5 A g^{-1} . In addition, as shown in the Ragone plot (Fig. 9d), JC-3.5 possesses energy densities of 30.9 and 24.5 W h kg^{-1} at power densities of 133.5 W kg^{-1} and 8.9 kW kg^{-1} , respectively.

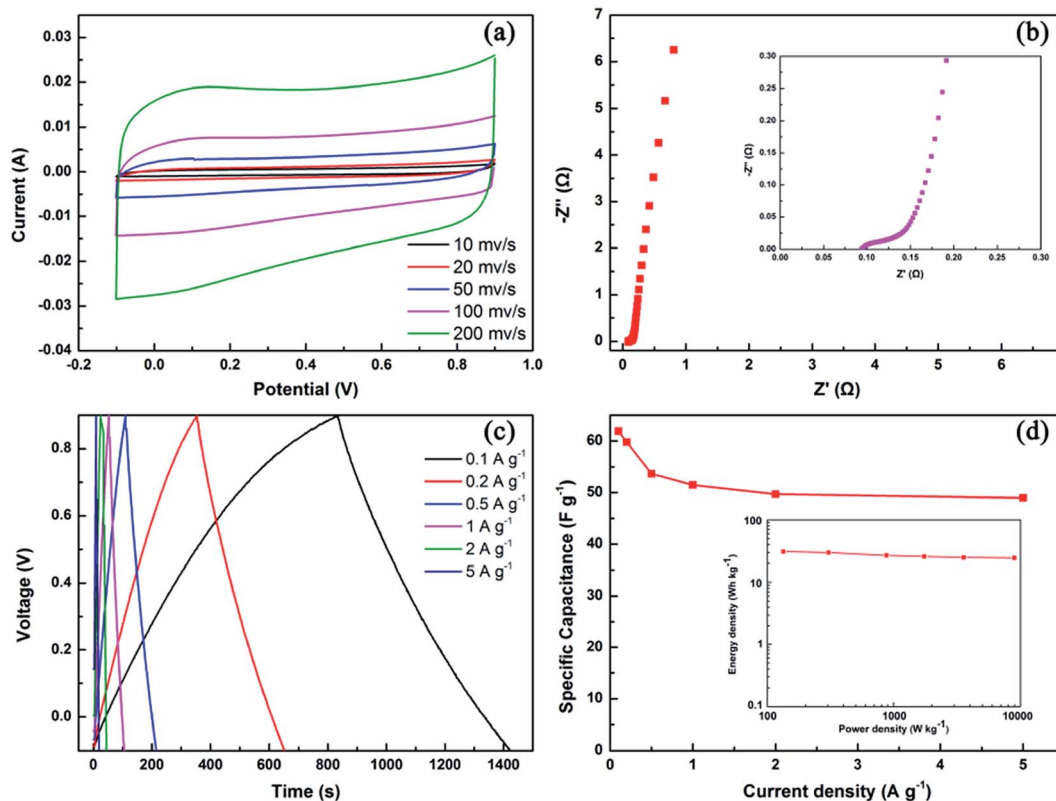


Fig. 9 The CV curves obtained for JC-3.5 at different scan rates (a), the Nyquist plot of JC-3.5 with the inset magnifying the high frequency region (b), galvanostatic charge–discharge curves at different current density of JC-3.5 (c) and specific capacitance at different current density with the inset showing the Ragone plot of JC-3.5 (d).



Based on the above evaluation of the electrochemistry performance, JC-3.5 shows the most superior properties. First of all, suitable KOH consumption makes it possess the largest SSA and a hierarchical porous structure containing abundant micropores. This pore structure ensures excellent electrolyte contact and rapid ionic transportation, and thus enhances the charge storage density. Except for the excellent pore structure, the O and N containing functional groups determined using XPS can also increase the electrical conductivity and help the diffusion of ions and electrons. The self-doped heteroatoms on the surface may have undergone redox reactions to provide pseudo-capacitance behavior. As a supercapacitor electrode material, the JC performance is a combination of EDLC and faradaic pseudocapacitance, and possesses high capacitance, good rate performance, and excellent cycling stability.

4. Conclusions

In this work, we have reported a winter-jujube-derived carbon material (JC) prepared *via* pre-carbonization and KOH-activation. The synthesized JC samples possess abundant self-doped heteroatoms and hierarchically porous structures (the co-existence of micro-, meso-, and macropores). The samples prepared with different KOH consumption possess different properties. In these samples, JC-3.5 with a KOH to carbon weight ratio of 3.5 : 1 had an ultra-high SSA of 2286.2 cm² g⁻¹. JC-3.5 possessed a high capacitance up to 341 F g⁻¹ at 0.5 A g⁻¹. The presence of abundant self-doped heteroatoms O and N made a huge contribution. Besides the high specific capacitance, it also has good rate performance (76% capacitance retained when the current was increased to 20 A g⁻¹ from 0.5 A g⁻¹). As for its cycling stability, there was only a 4.5% capacitance loss after 5000 cycles at 5 A g⁻¹. Besides, the JC-3.5-based symmetric supercapacitor device possessed an energy density of 30.9 kW h⁻¹ at 0.1 A g⁻¹ and power density of 8.9 kW at 5 A g⁻¹. In summary, as a renewable green resource, winter-jujube-derived carbon material is a promising candidate for use as a supercapacitor electrode material.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

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