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# A hierarchical porous P-doped carbon electrode through hydrothermal carbonization of pomelo valves for high-performance supercapacitors†

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Porous carbon materials are synthesized from pomelo valves by the hydrothermal activation of  $\text{H}_3\text{PO}_4$  followed by simple carbonization. The as-synthesized hierarchically porous carbon electrode exhibits a high specific capacitance of  $966.4 \text{ F g}^{-1}$  at  $1 \text{ A g}^{-1}$  and an ultra-high stability of 95.6% even after 10 000 cycles. Moreover, the supercapacitor also demonstrates a maximum energy of  $36.39 \text{ W h kg}^{-1}$  and a maximum power of  $33.33 \text{ kW kg}^{-1}$  with an energy retention of  $25.56 \text{ W h kg}^{-1}$ , which paves the way for the development of high-performance, green supercapacitors for advanced energy storage systems.

## 1. Introduction

The rapidly expanding market for portable electronic devices and hybrid vehicles is stimulating intensive research for developing cost-effective and eco-friendly energy storage devices.<sup>1–3</sup> Energy storage devices play a crucial role in enabling the transition from fossil fuels to renewable energy sources.<sup>4–6</sup> Supercapacitors feature ultrafast charge/discharge rates, high power densities, low maintenance costs, and long cycling lives as promising energy storage devices for electric vehicles, portable electronics, and stationary energy storage devices. They are classified into electrical double layer capacitors (EDLCs) and pseudocapacitors according to the different capacitive mechanisms.<sup>7–10</sup>

Because the capacitance is proportional to the surface area of the electrode, carbon materials usually have good conductivities, abundant morphologies, and tunable surface properties with doping and/or surface functionalization, which are the main options to explore advanced EDLC electrode materials.<sup>11–13</sup> Various carbon materials such as activated carbon (AC), carbon nanotubes (CNTs), and graphene have been intensively studied, and it has been demonstrated that there is still plenty of room to further improve the supercapacitive performance with new

carbon architectures.<sup>14–16</sup> In particular, porous carbon materials derived from biomass have attracted increasing attention due to their availability, chemical composition, environmentally friendly disposal and cost decrease associated with activated carbon production.<sup>17–20</sup> More importantly, most biomass contains abundant hetero-atoms (*e.g.*, oxygen, nitrogen, and sulfur) that can be *in situ* doped into carbon matrices during carbonization processes. These heteroatoms not only increase the wettability of the carbon material, but also contribute pseudo-capacitance.<sup>21–24</sup>

Several studies have dealt with functionalization of activated carbon to achieve the desired properties by grafting different surface groups at appropriate oxidation states.<sup>25–27</sup> Among the methods for modification of AC properties, doping the carbon matrix with heteroatoms, such as oxygen, nitrogen, boron, sulfur, and phosphorus, is the most effective way to tailor the electronic structures and improve the surface properties. Oxygen functionalities are often present on the carbon surface, and their effect on the capacitive performance must be considered as they take part in faradaic interactions and thus significantly increase the specific capacitance of carbons in acidic aqueous supercapacitors. The conjugation of the lone-pair electrons of N and the graphitic  $\pi$ -bonds of carbon materials can further distort the carbon structure to create defects and available active sites, which has been extensively and deeply studied. However, the functional mechanism of the phosphorus configuration in the skeleton of P-doped carbon materials remains unclear.<sup>28–36</sup>

In this context, we report a novel type of porous P-enriched carbon material derived from pomelo valves through hydrothermal carbonization of  $\text{H}_3\text{PO}_4$  (Scheme 1). By the way, the chemical bonding states of phosphorus with carbon and oxygen have been systematically explored and the type of P-doped configuration with the electrochemical performance has also

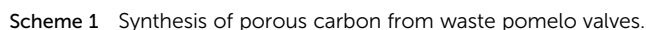
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## 2.1 Materials and methods

**2.1.2 Synthesis of the activated pomelo valve-derived porous carbon (APVPC).** The waste pomelo valves were air-dried and cut into pieces prior to the hydrothermal carbonization process. During the procedure of the hydrothermal treatment, 1 g of pomelo valves were placed in a Teflon-lined stainless steel autoclave wherein 70 mL of a homogeneous  $\text{H}_3\text{PO}_4$  (1 M, 5 M, 10 M, and 16 M) solution was present. Subsequently, the autoclave was sealed and placed in an electric oven at a temperature of 120 °C for 1, 3, and 6 h. After cooling down to room temperature naturally, the samples were centrifuged and freeze-dried. Next, the products were calcined at 700 °C under an Ar flow for 3 h. Finally, the black carbon material was washed with hot water several times and dried in an oven at 80 °C for 12 hours to afford APVPC. The samples were denoted as APVPC-1 (1 h, 16 M  $\text{H}_3\text{PO}_4$ ), APVPC-2 (3 h, 16 M), APVPC-3 (6 h, 16 M), APVPC-4 (3 h, 1 M), APVPC-5 (3 h, 5 M) and APVPC-6 (3 h, 10 M). For comparison, air-dried pomelo valves treated with deionized water were obtained according to a similar procedure and the obtained porous carbon was denoted as PVPC.

## 2.2 Characterization

2100) coupled with energy dispersive spectroscopy (EDS). Elemental analysis was carried out in a LECO CHNS micro-analyzer. Nitrogen sorption isotherms were obtained using an Autosorb-1 (Quantachrome Instruments). The specific surface area was calculated using the modified Brunauer–Emmett–Teller (BET) method. The pore size distribution and the pore volume were analyzed from the adsorption branch isotherms by the density functional theory (DFT) method. Moreover, the total pore volume ( $V_t$ ) was estimated from the amount adsorbed at a relative pressure  $P/P_0$  of 0.990. The micropore volume ( $V_{mic}$ ) and micropore surface area ( $S_{mic}$ ) were determined using  $t$ -plot theory. Raman spectra were acquired with a Jobin-Yvon HR 800 spectrometer. X-ray photoelectron spectroscopy (XPS) measurements were performed on a Thermo Fisher Scientific instrument (Escalab 250xi, USA). Fourier transform infrared (FT-IR) spectra were recorded on a Thermo Scientific Nicolet iS 50 spectrometer.

### 2.3 Electrochemical measurements

For the three-electrode system, the working electrode was fabricated through the dispersion of activated carbon in a mixed solution of Nafion and ethanol (1 : 20) dropped on a glassy carbon electrode; platinum foil and Hg/HgO were used as the counter and reference electrodes, respectively. The electrochemical capacitive performance was analyzed on an electrochemical workstation (Shanghai Chenhua Instrument Co. Ltd, China, CHI660D) at room temperature.

For the three-electrode system, the gravimetric specific capacitance,  $C_{sp}$  (F g<sup>-1</sup>), can be calculated according to the following equations

$$C_{\text{sp}} = I \times \Delta t / m \times \Delta V \quad (1)$$

where  $I$  is the discharge current (A),  $\Delta t$  is the discharge time (s),  $\Delta V$  is the voltage (V), and  $m$  is the mass of the active materials (g).

For the two-electrode system, the working electrodes were prepared by mixing the electroactive material, acetylene black, and polytetrafluoroethylene (PTFE) binder with a weight ratio of 8 : 1 : 1, followed by coating on foamed Ni grids (1 cm × 1 cm) and pressing under a pressure of 20 MPa as well as drying overnight at 80 °C. The mass of the active material loading in each working electrode was measured to be 2–3 mg cm<sup>-2</sup>. The symmetrical capacitor was assembled using two nearly identical working electrodes and a glass fiber membrane (Waterman, GF/B) as a separator as well as a 3 M KOH aqueous solution as the electrolyte.

For the two-electrode system, the gravimetric specific capacitance,  $C_{\text{sp}}$  ( $\text{F g}^{-1}$ ), for a single electrode was calculated from each galvanostatic charge-discharge (GCD) curve using the following equation:

$$C_{\text{sp}} = 2I \times \Delta t/m \times \Delta V \quad (2)$$

where  $C_{\text{sp}}$  ( $\text{F g}^{-1}$ ) is the specific capacitance based on the mass of the activated carbon,  $I$  (A) is the discharge current,  $\Delta t$  (s) is the discharge time,  $\Delta V$  (V) is the voltage change excluding the  $IR$



drop during the discharge process, and  $m$  (g) is the loading of the active material in each working electrode.

The energy density  $E$  ( $\text{Wh kg}^{-1}$ ) and the power density  $P$  ( $\text{W kg}^{-1}$ ) were estimated by using the following equations:

$$E = C_{\text{sp}} V^2 / (2 \times 3.6) \quad (3)$$

$$P = 3600 E / \Delta t \quad (4)$$

where  $C_{\text{sp}}$  ( $\text{F g}^{-1}$ ) is the specific capacitance of a single electrode,  $V$  (V) is the voltage change during the discharge process, and  $\Delta t$  (s) is the discharge time.

### 3. Results and discussion

#### 3.1 Materials characterization

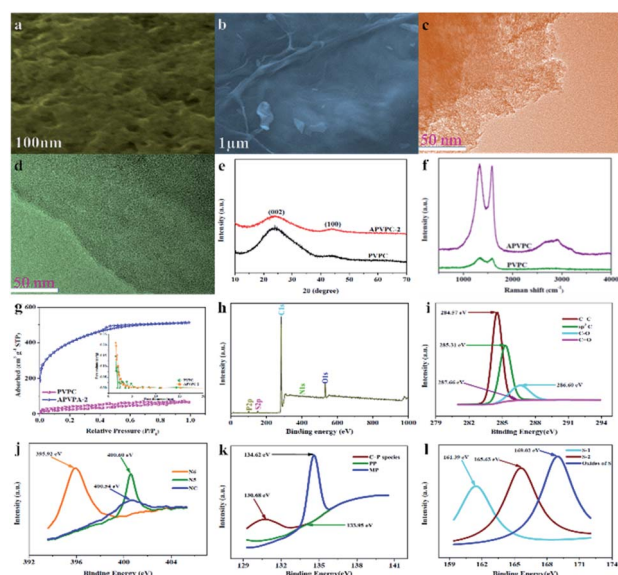
The detailed synthesis procedure of the APVPC is described in Scheme 1. It can be noted that  $\text{H}_3\text{PO}_4$  activation plays an important role in the development of the pore structure in APVPC. In particular, the specific surface area (SSA) and the pore structure of the product are obviously influenced by the concentration of  $\text{H}_3\text{PO}_4$ . The sample produced with 85%  $\text{H}_3\text{PO}_4$  is selected as a typical sample for characterization and used for SCs unless otherwise specified.

The scanning electron microscopy (SEM) image (Fig. 1a) reveals that the APVPC-2 is made up of many carbon flakes with numerous uniformly sized mesopores and micropores through carbonizing the gelatinous tissue of pomelo valves, which are favorable for ion transport. Compared to the SEM image of APVPC-2, the PVPC (Fig. 1b) exhibits a monolithic morphology,

implying that the activation of  $\text{H}_3\text{PO}_4$  could help to form thin sheets of carbon and to create a porous structure. The high resolution TEM image of the APVPC-2 (Fig. 1c) demonstrates that the APVPC-2 consists of many carbon flakes with numerous nanopores of 1–3 nm in size, which contribute to a super high surface area. In addition to the connected thin layers, there are many amorphous carbon particles about 20 nm in diameter interconnected tightly, which are conducive to a high-rate-performance supercapacitor for facilitating the diffusion of electrolyte ions.<sup>37</sup> Compared with that of APVPC-2, the TEM image of PVPC (Fig. 1d) indicates a highly defective micro- and mesoporous morphology surrounded by curved carbon, which confirms a highly disordered structure.<sup>38</sup>

The X-ray diffraction (XRD) (Fig. 1e) patterns of the samples show two characteristic peaks located at  $2\theta$  values of  $23.12^\circ$  and  $43.8^\circ$ , corresponding to the (002) and (100) planes of graphitic carbon, respectively, and barely discernible peaks. The broad diffraction peak of carbon demonstrates the highly amorphous and disordered nature of porous carbon. Further, the high intensity in the low angle region may be ascribed to the existence of abundant micropores.<sup>39</sup> Compared with the spectrum of APVPC-2, we can see that the water residue in the biomass tissue might be effective in increasing the porosity as the spectrum of PVPC shows a broader peak.<sup>40</sup> In the Raman spectrum for APVPC-2 (Fig. 1f), the two peaks located at about  $1331$  and  $1584 \text{ cm}^{-1}$  correspond to the D (disordered carbon) and G (graphitic carbon) bands, respectively. The intensity ratio between D and G bands ( $I_{\text{D}}/I_{\text{G}}$ ) is summarized in Fig. S2.† The  $I_{\text{D}}/I_{\text{G}}$  ratio obviously decreases from 1.051 for PVPC to 1.022 for APVPC-2 (Fig. S2a†), confirming that deeper activation decreased the presence of defects and the disordered section in the APVPC-2 sample,<sup>41</sup> which agrees well with the XRD analysis and TEM images. However, the G band of APVPC-2 shows a higher redshift to  $1584 \text{ cm}^{-1}$  compared to that of pristine PVPC ( $1594 \text{ cm}^{-1}$ ), which could be attributed to n-type substitutional doping of porous carbon and further confirms the heteroatom doping into the carbon framework.<sup>23</sup> As for the effect of reaction time, the  $I_{\text{D}}/I_{\text{G}}$  ratio of APVPC-2 (1.022, 3 h) is obviously higher than that of APVPC-1 (0.948, 1 h) and APVPC-3 (0.96, 6 h) (Fig. S2b and c†), which indicates that the defect intensity gradually becomes stronger, and then turns slightly weaker at longer activation times. In addition, the  $I_{\text{D}}/I_{\text{G}}$  ratio increases along with the concentration of  $\text{H}_3\text{PO}_4$  (1 M vs. 5 M vs. 10 M vs. 16 M  $\text{H}_3\text{PO}_4$ ; APVPC-4 vs. APVPC-5 vs. APVPC-6 vs. APVPC-2; 0.927 vs. 0.941 vs. 0.958 vs. 1.022) (Fig. S2d and e†), which demonstrates that the graphitization intensity gradually becomes weaker. Based on the aforementioned results, the higher values for APVPC-2 indicate the highly disordered nature and heteroatom doping and could further contribute to the increase in electrical conductivity after heteroatom doping, which is due to broadening of van Hove singularities in the density of states making the Fermi level shift towards the conduction bands.<sup>8</sup>

Based on the apparent hysteresis loop at a relative pressure  $P/P_0$  from 0.4 to 0.9 (Fig. 1g), APVPC-2 possesses a well-ordered mesoporous structure, which agrees well with the pore size distribution calculated using the Barrett–Joyner–Halenda (BJH)



**Fig. 1** (a) SEM images of APVPC-2. (b) SEM images of PVPC. (c) TEM images of APVPC-2. (d) TEM images of PVPC. (e) XRD patterns of PVPC and APVPC-2. (f) Raman spectra of PVPC and APVPC-2. (g)  $\text{N}_2$  adsorption/desorption isotherms and the pore size distribution of APVPC-2 and PVPC. (h) XPS survey spectrum of APVPC-2. (i) High resolution C 1s spectrum of APVPC-2. (j) High resolution N 1s spectrum of APVPC-2. (k) High resolution P 2p spectrum of APVPC-2. (l) High resolution S 2p spectrum of APVPC-2.





model. Compared with PVPC, APVPC-2 has the widest pore size distribution, which results in a high nitrogen BET specific surface area of  $1432.125 \text{ m}^2 \text{ g}^{-1}$  (vs.  $96.939 \text{ m}^2 \text{ g}^{-1}$ ) with a pore volume of  $0.718 \text{ cm}^3 \text{ g}^{-1}$  ( $0.104 \text{ cm}^3 \text{ g}^{-1}$ ). The SSA contributed by mesopores is determined to be about  $115 \text{ m}^2 \text{ g}^{-1}$ . It is worth noting that the surface area of APVPC-2 is much higher than that of PVPC, further confirming that the activation of  $\text{H}_3\text{PO}_4$  could effectively create holes and further increase the specific surface area. As shown in Table 1, all samples have both micropores (0–2 nm) and mesopores (2–50 nm), and the pore size is mainly  $\sim 0.718 \text{ nm}$ . Increasing the activation time from 1 h to 3 h generates a huge amount of gas (CO and  $\text{CO}_2$ ) from the activation agent ( $\text{H}_3\text{PO}_4$ ) which favorably increases the textural properties (1 h vs. 3 h;  $245.82 \text{ m}^2 \text{ g}^{-1}$ ,  $0.135 \text{ cm}^3 \text{ g}^{-1}$  vs.  $1432.125 \text{ m}^2 \text{ g}^{-1}$ ,  $0.718 \text{ cm}^3 \text{ g}^{-1}$ ; APVPC-1 vs. APVPC-2). However, further increasing the activation time to 6 h induces the pores to collapse, unfavorably resulting in the decrease of the textural properties (3 h vs. 6 h;  $1432.125 \text{ m}^2 \text{ g}^{-1}$ ,  $0.718 \text{ cm}^3 \text{ g}^{-1}$  vs.  $678.56 \text{ m}^2 \text{ g}^{-1}$ ,  $0.374 \text{ cm}^3 \text{ g}^{-1}$ ; APVPC-2 vs. APVPC-3). Additionally, the textural properties also increase along with the concentration of  $\text{H}_3\text{PO}_4$  (1 M vs. 5 M vs. 10 M vs. 16 M; from  $373.664$  to  $1432.125 \text{ m}^2 \text{ g}^{-1}$ , from  $0.144 \text{ cm}^3 \text{ g}^{-1}$  to  $0.718 \text{ cm}^3 \text{ g}^{-1}$ ; APVPC-4 vs. APVPC-5 vs. APVPC-6 vs. APVPC-2). It is widely accepted that mesopores and macropores can provide electrolyte ion transport and diffusion channels, and the micropores offer space for ion storage energy.<sup>42</sup> The high porosity greatly maximizes the ion adsorption sites, which could lead to high capacitance and provide quick electrolyte diffusion into deeper pores.<sup>43</sup> Therefore, the as-prepared APVPC-2 should have a high specific capacitance and rate capability when applied as the electrode material in supercapacitors.

For more information on the elemental chemical states of the heteroatoms in the APVPC-2, X-ray photoelectron spectroscopy (XPS) (Fig. 1h) is carried out and indicates five peaks at about 283.99, 399.94, 531.49, 128.12 and 163.8 eV corresponding to the C 1s, N 1s, O 1s, P 2p and S 2p peaks, respectively. The relative percentages of the carbon species are given in Table 2. The results of elemental analysis reveal that the APVPC-2 mainly consists of C, N, O, P and S elements, and their contents are 62.86, 3.94, 32.28, 0.74 and 0.17 atom%, respectively, while for PVPC, the XPS spectrum also shows five peaks for C 1s, N 1s, O 1s, P 2p and S 2p at about 283.99, 399.94, 531.49, 128.12 and 163.8 eV, respectively. Meanwhile, the elemental contents for C, N, O, P and S are 91.72, 0.52, 7.63, 0.06 and 0.08%,

Table 2 XPS analysis of the PVPC and APVPC-2

Samples	C (%)	N (%)	O (%)	P (%)	S (%)
PVPC	91.72	0.52	7.63	0.06	0.08
APVPC-2	62.86	3.94	32.28	0.74	0.17

respectively. Other possible metal impurities from the biomass source are below the detection limit; these could have been removed during the post-acid washing process. The high resolution C 1s spectra for APVPC-2 (Fig. 1i) have been resolved into four peaks centered at around 284.57, 285.31, 286.60, and 287.66 eV, which correspond to  $\text{C}=\text{C}$ ,  $\text{sp}^3\text{C}$ ,  $\text{C}-\text{O}$  and  $\text{C}=\text{O}$ ,<sup>44</sup> respectively. Moreover, the deconvoluted N 1s spectrum (Fig. 1j) contains three peaks, pyridinic-N (N-6, 395.92 eV), pyrrolic-N (N-5, 400.60 eV) and amides/amines or nitrile N (NC, 400.54 eV),<sup>45</sup> which indicates the presence of nitrogen and further could improve the wettability and provide a large number of active sites for adsorption. In the  $2\text{p}_{3/2}$  spectrum (Fig. 1k), the three peaks at 130.68 eV, 133.93 eV and 134.62 eV can be assigned to C–P species, pyrophosphate/polyphosphate (PP) and monophosphate/metaphosphate (MP) groups,<sup>46</sup> respectively. After hydrothermal carbonization of pomelo valves through  $\text{H}_3\text{PO}_4$ -activation, the content of P is increased from 0.06 (PVPC) to 0.74 (APVPC-2), which confirms the successful phosphorus-doping of porous carbon. In addition, the high-resolution S 2p spectrum (Fig. 1l) shows three peaks at 161.39, 165.63 and 169.02 eV, which are attributed to  $\text{C}=\text{S}$  and  $\text{C}-\text{S}$  bonds and the oxides of sulfur.<sup>47</sup> Thus, the XPS results also demonstrate the existence of heteroatoms such as N, O, P and S-containing functional groups in the samples. These functional groups may participate in the faradaic reactions that increase the pseudo-capacitance and improve the wettability of the APVPC-based electrodes in the aqueous electrolytes.<sup>48</sup>

Fourier transform infrared (FT-IR) spectroscopy is used to identify the surface functional groups of the as-synthesized samples (Fig. S1†). The absorptions at 3443, 2978, 1654, 1048, 1078 and  $671 \text{ cm}^{-1}$  are attributed to  $-\text{NH}$ ,  $-\text{C}-\text{H}$ ,  $-\text{C}=\text{O}$ ,  $\text{C}-\text{OH}$ ,  $\text{C}=\text{S}$ , and  $\text{C}-\text{P}$ ,<sup>49</sup> respectively. From the above FT-IR analysis, it could be deduced that oxygen-, nitrogen-, phosphorus- and sulphur-containing groups may exist in the samples, which agrees well with the XPS results. Based on the aforementioned characterization, the carbon material APVPC-2

Table 1 Summary of the BET characteristics of activated carbon

Sample	$S_{\text{BET}}^a$ ( $\text{m}^2 \text{ g}^{-1}$ )	$V_{\text{tot}}^b$ ( $\text{cm}^3 \text{ g}^{-1}$ )	$S_{\text{mi}}^c$ ( $\text{m}^2 \text{ g}^{-1}$ )	$S_{\text{me}}^d$ ( $\text{m}^2 \text{ g}^{-1}$ )	$S_{\text{ma}}^e$ ( $\text{m}^2 \text{ g}^{-1}$ )	$V_{\text{mi}}^f$ ( $\text{cm}^3 \text{ g}^{-1}$ )	$V_{\text{me}}^g$ ( $\text{cm}^3 \text{ g}^{-1}$ )	$V_{\text{ma}}^h$ ( $\text{cm}^3 \text{ g}^{-1}$ )
PVPC	96.939	0.104	12.67	63.28	20.989	0.0162	0.0623	0.0255
APVPC-1	245.82	0.135	86.132	105.62	54.068	0.0138	0.0461	0.0751
APVPC-2	1432.125	0.718	349.06	969.377	113.688	0.175	0.486	0.058
APVPC-3	678.56	0.374	213.62	268.53	196.41	0.102	0.153	0.119
APVPC-4	373.664	0.144	121.56	134.89	117.214	0.319	0.157	0.255
APVPC-5	421.694	0.224	163.72	136.26	121.714	0.0631	0.116	0.0449
APVPC-6	631.502	0.624	168.42	251.68	211.402	0.168	0.235	0.221

<sup>a</sup>  $S_{\text{BET}}$ : BET surface area. <sup>b</sup>  $V_{\text{tot}}$ : total volume. <sup>c</sup>  $S_{\text{mi}}$ : micropore surface area. <sup>d</sup>  $S_{\text{me}}$ : mesopore surface area. <sup>e</sup>  $S_{\text{ma}}$ : macropore surface area. <sup>f</sup>  $V_{\text{mi}}$ : micropore volume. <sup>g</sup>  $V_{\text{me}}$ : mesopore volume. <sup>h</sup>  $V_{\text{ma}}$ : macropore volume.



derived from pomelo valves is a more preferential electrode material for supercapacitors by virtue of its high surface area and 3D interconnected porous structure as well as the hetero-atom doping.

Based on the aforementioned investigation, we propose a formation mechanism of porous carbon nanosheets. The pomelo valves possess a net like structure consisting of a large amount of polymeric cellulose. Owing to substantial hydrophilic functional groups such as hydroxy and carbonyl groups, the molecules of the polymeric cellulose in the pomelo valves could adsorb certain amounts of water molecules. In the activation process,  $\text{H}_3\text{PO}_4$  vibrates violently in water to prevent the adjacent cell walls from agglomerating, and then the atoms decomposed from the molecules of polysaccharides connect together *in situ* to form ultrathin nanosheets and abundant pores. During the subsequent high-temperature carbonizing process, the polymeric cellulose is hydrolyzed to oligosaccharides along with gas evolution, which could contribute to forming a porous structure.

### 3.2 Electrochemical behaviors

As shown in Fig. 2a, the cyclic voltammetry (CV) curves of the APVPC-2 electrode have perfect rectangular shapes with obvious humps in the low potential region, which is indicative of the combined effects of electric double-layer capacitance (EDLC) and pseudo-capacitance due to the existence of active hetero-atoms.<sup>45</sup> Even with an increase in scan rates from 5 to 100  $\text{mV s}^{-1}$ , the APVPC-2 electrode still maintains perfect rectangle-shaped profiles, indicating faster kinetics in the porous carbon by virtue of the openness in the structure resulting in fast ion transportation inside the pores.<sup>50</sup> Because the CV curves at 100  $\text{mV s}^{-1}$  are correlated with the capacitance of the electrode, APVPC-2 shows a larger rectangular shape of the CV curves than other materials (Fig. S3a and b†), suggesting the highest capacitance, which is consistent with the capacitance calculated from the GCD curves. This could be attributed to the

broader PSD and higher mesoporosity of APVPC-2 that is conducive to facile, fast electrolyte ion transport and shortens the diffusion distances from the electrolyte to the inner micropore surface.<sup>51</sup>

Electrochemical impedance spectroscopy (EIS) is also a critical technique to further study the electrochemical performance of the samples. According to the Nyquist plots of APVPC-2 (Fig. 2b) in 3 M KOH, the steep linear curve of APVPC-2 in the low-frequency region is closer to a vertical line than that of PVPC, which reveals that the  $\text{H}_3\text{PO}_4$ -activation can improve the capacitive behavior effectively. In contrast to PVPC, APVPC-2 exhibits a larger slope of the straight line in Nyquist plots, which reveals lower diffusive resistance ( $W$ ). At low frequency, electrolyte ions could penetrate inside the porous AC and access more of the electrode surface thereby contributing to the high capacitance value. However, at high frequency, electrolyte ions can only access the surface of carbon materials whereas the inner pores are not accessible, resulting in a sharp decrease in the capacitance.<sup>52</sup> Meanwhile, the internal cell resistance (real  $Z'$  axis in Nyquist plots) is 3.12  $\Omega$  (56.25 kHz) for APVPC-2 and 4.408  $\Omega$  (68.12 kHz) for PVPC. Moreover, pseudocapacitance originating from Faraday reactions owing to the surface functional groups of APVPC-2 has a insignificant effect in comparison to the double-layer capacitance, which agrees well with CV curves.<sup>53</sup> As shown in Fig. S3c,† the internal cell resistance is 3.087  $\Omega$  (82.52 kHz) for APVPC-1 and 3.104  $\Omega$  (68.12 kHz) for APVPC-3. The slope of APVPC-2 parallel to the  $Z''$  axis is higher than those of APVPC-1 and APVPC-3 prepared with different activation times (1 h vs. 3 h vs. 6 h, APVPC-1 vs. APVPC-2 vs. APVPC-3) which reveals less diffusion restriction for electrolyte ions to access the electrode surface.<sup>54</sup> From the slopes of the straight lines in the Nyquist plots, the order of WAPVPC-2 > WAPVPC-1 > WAPVPC-3 could be deduced. Additionally, the internal cell resistance is 6.289  $\Omega$  (68.12 kHz) for APVPC-4, 5.004  $\Omega$  (56.25 kHz) for APVPC-5 and 3.349  $\Omega$  (56.25 kHz) for APVPC-6 (Fig. S3d†). The slope of APVPC-2 is also the higher than those of APVPC-4, APVPC-5 and APVPC-6 synthesized with different concentrations of  $\text{H}_3\text{PO}_4$  (1 M vs. 5 M vs. 10 M vs. 16 M  $\text{H}_3\text{PO}_4$ , APVPC-4 vs. APVPC-5 vs. APVPC-6 vs. APVPC-2), which exhibits the order of WAPVPC-2 > WAPVPC-4 > WAPVPC-6 > WAPVPC-5. From the electrochemical impedance spectroscopy measurements, it could be deduced that the electrode material APVPC-2 has low ionic resistance inside the porous structure which promotes the fast diffusion of electrolyte ions into the porous network. In addition, the electrode material APVPC-2 exhibits good conductivity and low charge transfer resistance that facilitate fast charge transfer between the electrolytes and electrode materials. The results further demonstrate that the APVPC-2 is a promising electrode material for supercapacitor applications.

The capacitive performances of the porous materials are further tested with galvanostatic charge–discharge (GCD) measurements at various current densities in 3 M KOH with the same voltage windows as those in the CV analysis. The GCD curves are almost symmetrical with slight distortion, confirming the good capacitive behavior of electrical double layer capacitance (EDLC) and a little pseudo-capacitance, which

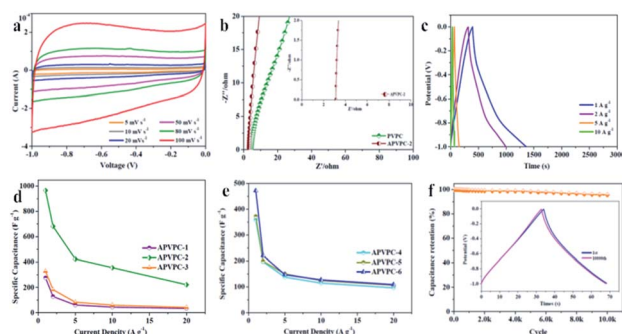


Fig. 2 (a) CV curves of APVPC-2 at different scan rates. (b) Nyquist plots of PVPC and APVPC-2. (c) Galvanostatic charge/discharge curves of APVPC-2 at different current densities. (d) Capacitances of APVPC samples prepared with different reaction times at different current densities. (e) Capacitances of APVPC samples prepared with different concentrations of  $\text{H}_3\text{PO}_4$  at different current densities. (f) Cycling performance of APVPC-2 at 10  $\text{A g}^{-1}$ . The inset shows the GCD curves of the 1st and 10 000th cycle.



coincides with the results of CV. As calculated from the GCD curves, APVPC-2 exhibits a specific capacitance of  $966.4 \text{ F g}^{-1}$  at  $1 \text{ A g}^{-1}$  ( $474.38 \text{ F g}^{-1}$  at  $1 \text{ A g}^{-1}$  for PVPC) and still displays a specific capacitance of  $221.6 \text{ F g}^{-1}$  even at  $20 \text{ A g}^{-1}$ , which reveals the good energy storage-output capacity at high current density and further confirms the effect of the activation agent  $\text{H}_3\text{PO}_4$ . The lower capacitance at higher current densities is also ascribed to the insufficient electrolyte ion diffusion kinetics across the micropores due to the narrow pore size at higher operating current densities, and the reduced accumulation amount of electrolyte ions on the porous electrode interface leads to a decrease in specific capacitance.<sup>49</sup> The APVPC-2 electrode exhibits high specific capacitance due to the synergistic effect of the rich active heteroatom-doping and suitable pore size distribution. This result also illustrates that the conductivity of the electrode is good, but the pore structure is favorable for  $\text{K}^+$  and  $\text{OH}^-$  ion diffusion. In the KOH aqueous electrolyte solution, the micropores with a pore size less than  $0.5 \text{ nm}$  are usually deemed too narrow for double-layer formation; however, pores larger than  $2 \text{ nm}$  are too wide. Consequently, the APVPC-2 that possesses a concentrated pore size distribution of  $0.718 \text{ nm}$  is very applicable for the delivery of the electrolyte ions in the sample porous channels, further achieving high electrosorption efficiency, which also could result in improved specific capacitance.<sup>55</sup> In addition, the surface functionalities and incorporation of N, O, P and S heteroatoms in APVPC-2 which have been investigated by XPS could also contribute to high pseudocapacitance.<sup>56</sup> In this context, the graphitic N in APVPC-2 greatly improves the electron transport by decreasing the intrinsic resistance, and the pyridinic and pyrrolic N in porous carbon strongly favors rapid ion transport kinetics inside the pores.<sup>54</sup> The P-doped configuration of APVPC-2 could inhibit the formation of electrophilic oxygen species but also greatly stabilizes the electrochemical interface of the graphene electrode.<sup>57</sup> The S heteroatoms can greatly enhance space utilization by promoting the electrosorption of electrolyte ions.<sup>58</sup> The presence of heteroatoms increases the polarity of the pores and thereby facilitates easy contact between pores and the electrolyte. In general, the co-doping effect of N, P and S could greatly modify the chemical and surface nature of the porous carbon and improve the ion and electron transfer kinetics of carbon, which could contribute to a superior performance. At the same time, the reaction time and concentration of  $\text{H}_3\text{PO}_4$  could also affect the specific capacitance. Fig. 2d and e compare the capacitance retention of all APVPC samples in a range of current density from  $1$  to  $20 \text{ A g}^{-1}$ . As the reaction time increases (Fig. 2d), the APVPC-2 delivers higher capacitance than APVPC-1 and APVPC-3, which is attributed to the increased pores formed from the activated carbon atoms and the activator with the optimal reaction time ( $1 \text{ h}$  vs.  $3 \text{ h}$  vs.  $6 \text{ h}$ ,  $474.38$  vs.  $966.4$  vs.  $276.9 \text{ F g}^{-1}$ ; APVPC-1 vs. APVPC-2 vs. APVPC-3). With the increase of concentration of  $\text{H}_3\text{PO}_4$  (Fig. 2e), the specific capacitances are also progressively increased in the same way ( $1 \text{ M}$  vs.  $5 \text{ M}$  vs.  $10 \text{ M}$  vs.  $16 \text{ M}$ ,  $506$  vs.  $53.8$  vs.  $475$  vs.  $966.4 \text{ F g}^{-1}$ ; APVPC-4 vs. APVPC-5 vs. APVPC-6 vs. APVPC-2). When the concentration of  $\text{H}_3\text{PO}_4$  is relatively low, the amount of activator is inadequate to

fully activate the carbon. But with the increase of concentration of  $\text{H}_3\text{PO}_4$ , more carbon can react with the activator and further generate more pores, which eventually contribute to the increase of the specific surface area and the pore volume. In general, APVPC-2 exhibits a smaller charge-transfer resistance and larger ion diffusion than the other APVPC samples and other previously reported biomass derived porous carbon with aqueous electrolytes,<sup>59</sup> which demonstrates that despite having a moderate micropore surface area, appropriate PSD, rich active heteroatom contents and good crystallinity endow APVPC-2 with the best capacitive performance.

The long-term cycling stability is a crucial factor for electrode materials in energy storage applications. Fig. 2f demonstrates the variation of capacitance retention and GCD curves as a function of cycle number for APVPC-2 at a current density of  $10 \text{ A g}^{-1}$ . The specific capacitance decreases slowly in the first 1000 cycles and is maintained at about 95.6% of the initial specific capacitance with a slight fluctuation even after 10 000 cycles. Moreover, the 1st and 10 000th charge-discharge curves are almost fully overlapping except for a faint decay, which evidences the excellent cycling stability and the robust structure of these pomelo valve derived porous carbons.

A symmetric two-electrode cell was assembled to further evaluate the capacitive performance of APVPC-2 in  $3 \text{ M KOH}$ . The CV curves (Fig. 3a) recorded at different current densities from  $5$  to  $100 \text{ mV s}^{-1}$  have a quasi-rectangular shape, which demonstrates good electric double-layer capacitance (EDLC) behavior with fast charge/discharge processes and a good rate electrochemical performance.<sup>60</sup> Meanwhile, there are also no redox peaks/humps due to the pseudocapacitive interaction of ions with functional groups which could be observed. The pseudocapacitance behavior in porous carbon strongly depends

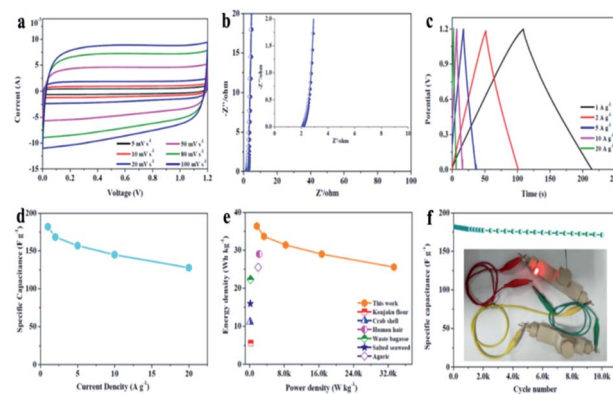


Fig. 3 (a) CV curves of the APVPC//APVPC symmetrical supercapacitor at different scan rates in the voltage window of  $0-1.2 \text{ V}$ . (b) Nyquist plots of the APVPC//APVPC symmetrical supercapacitor before and after recycling. (c) Galvanostatic charge/discharge curves of the APVPC//APVPC symmetrical supercapacitor at different current densities. (d) Specific capacitances of the as-assembled APVPC symmetrical supercapacitor at different current densities. (e) Ragone plots of the APVPC symmetrical supercapacitor and other previously reported carbon-based symmetric supercapacitors. (f) Cycle performance of the APVPC symmetrical supercapacitor at a current density of  $10 \text{ A g}^{-1}$  (inset: photograph of a lit LED bulb).





on the nature of the electrolytes (e.g.: type and pH) and the effect is almost negligible in neutral and non-aqueous electrolytes.<sup>61</sup> The Nyquist plot for APVPC-2 (Fig. 3b) is almost vertical in the low-frequency region, revealing easy electrolyte diffusion to the surface and good capacitive behavior. Simultaneously, the internal cell resistance (real  $Z'$  axis from Nyquist plots) is 2.146  $\Omega$  (6.17 kHz) for the supercapacitor. The GC curves (Fig. 3c) exhibit perfect symmetric triangular shapes at different current densities, confirming a facile, smooth adsorption/desorption process and further good capacitive characteristics. During the measurement, the change of capacitance from 181.95 to 127.78 A g<sup>-1</sup> accompanies the current density variation from 1 to 20 A g<sup>-1</sup> (Fig. 3d), which reveals a good rate performance with a capacitance retention of 70.22%. The attractive rate capability can be ascribed to the matching between the pore and the electrolyte 3 M KOH. The results are also superior to those for reported carbon-based capacitors,<sup>62,63</sup> illustrating that the electrolyte ions could efficiently, rapidly and fully penetrate into the electrochemically active interface of the APVPC-2 carbon material even at a high current density (20 A g<sup>-1</sup>), which is due to its 3D hierarchically interconnected porous structure. Compared with other energy storage systems, the Ragone plot shows an energy density of 36.39 W h kg<sup>-1</sup> at a low power density of 1.67 kW kg<sup>-1</sup> (Fig. 3e). Even at an ultra-high power density of 33.33 kW kg<sup>-1</sup>, the supercapacitor could still function efficiently with an energy density of 25.56 W h kg<sup>-1</sup>, which surpasses those of several previously reported systems.<sup>64,65</sup> The superior energy-power output behavior effectively bridges the performance gap between batteries and capacitors. The long cycle life of the supercapacitor is evaluated using constant current charge-discharge cycles at a current density of 10 A g<sup>-1</sup> as shown in Fig. 3f. The energy loss could be calculated to be 5.1% after 10 000 cycles, indicating good cycling stability. To further demonstrate the practical application of the as-fabricated SC, a tandem device is assembled by connecting two prototype units in series. Subsequently, the device is used to light up a red light-emitting-diode (LED, the lowest working potential is 2.0 V), as depicted in the inset of Fig. 3f. This tentative test suggests the great potential of the APVPC-2 materials for high performance energy storage systems.

## 4. Conclusions

In conclusion, porous carbon materials with interconnected hierarchical pores are derived from a sustainable, low-cost, and environmentally abundant waste pomelo valve biomass precursor by the hydrothermal activation with H<sub>3</sub>PO<sub>4</sub> and simple carbonization. In addition, surface functionalities on the pore wall could be tuned using nitrogen, sulfur and phosphorus heteroatoms to achieve excellent electrochemical performance. The electrochemical performance of the porous carbon is superior to that of conventional porous carbon and graphene-based electrodes. The stability and energy-power behavior were among the best reported for any porous carbon. A maximum energy density of 36.39 W h kg<sup>-1</sup> is achieved, along with an ultra-high output power of ~25.56 W h kg<sup>-1</sup>, while retaining an energy density of ~33.33 kW kg<sup>-1</sup>. Moreover, a long-lasting stability of

10 000 cycles with an energy loss of 5.1% is also demonstrated. The superior performance contributes to a good platform for large-scale production of porous activated carbon networks to achieve high-performance and stable supercapacitors.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

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