



Cite this: *RSC Adv.*, 2019, 9, 12877

Three-dimensional “skin-framework” hybrid network as electroactive material platform for high-performance solid-state asymmetric supercapacitor†

Liaoyuan Xia,^{ab} Shaoheng Hu,^{ab} Xueqin Zhang,^{ab} Le Huang,^{ab} Yu Liao,^{ab} Yan Qing,^a Yiqiang Wu,^{*a} Wenping Jiang^a and Xihong Lu^{bc}

Three-dimensional (3D) electrode materials are ideal candidates for use in fabricating high-performance supercapacitors (SCs), owing to their unique network structure and excellent electrochemical properties. In this study, an aerogel film produced by the freeze-drying self-aggregation of multiwall carbon nanotubes (MWCNTs) and cellulose nanofibers (CNFs) served as the “skin”, and an inter-connected 3D network of nickel foam (NF) as the “framework”, for the fabrication of an MWCNT/CNF-NF (called MCN) hybrid material with a distinct “skin-framework” architecture. Considering the metrics of excellent conductivity, high wettability, binder-free and unique 3D “skin-framework” structure, the MCN hybrid material has great potential as an electroactive material platform in constructing state-of-the-art asymmetric supercapacitor (ASC) electrodes. By incorporating MCN with electroactive manganese dioxide (MnO₂) and active carbon (AC), MnO₂-MCN and AC-MCN composite electrodes with respective high areal capacitances of 1784.8 (equal to 469.7 F g⁻¹) and 868.8 mF cm⁻² (equal to 126.3 F g⁻¹) at 5 mA cm⁻² were successfully prepared. Further, both kinds of electrodes exhibited high charge/discharge ability rates and good cycle performance. Moreover, an optimally assembled MnO₂-MCN//AC-MCN solid-state ASC was reversibly charged/discharged at voltages as high as 1.8 V and possessed a remarkable volumetric capacity of 9.83 F cm⁻³ and an energy density of 4.25 mW h cm⁻³, as well as good cycle stability.

Received 15th February 2019
Accepted 21st April 2019

DOI: 10.1039/c9ra01164a

rsc.li/rsc-advances

1 Introduction

The rapid development of portable electronics has induced a demand for high-energy-density and lightweight energy storage devices.^{1–6} Solid-state asymmetric supercapacitors (ASCs), which are a new type of energy storage device, are assembled from positive and negative electrodes of different voltage windows.^{7–12} They are capable of operation over a wider voltage range, resulting in higher energy and power densities. These remarkable characteristics make them very attractive as ideal energy storage devices for portable electronics. Unlike traditional supercapacitors, a solid-state ASC is typically

composed of positive and negative electrodes and a gel electrolyte. Among these components, the performance of the electrodes is very critical to the overall performance of the ASC. Considerable effort has thus been invested in the development of high-performance positive and negative electrodes in solid-state ASCs.^{13–19}

Among the different electrode materials that have been recently developed for the fabrication of solid-state supercapacitors, those with unique 3D nanostructures have been found to be very promising, with the nanostructures providing large interfaces for ion diffusion and charge transport.^{20–24} Nickel foam (NF) has particularly attracted much attention because of its usability as a substrate/current collector and a template for preparing 3D electrodes.^{25–30} The highly porous 3D structure of NF and the large electrode–electrolyte interface imbue it with improved electrolyte impregnation and reduced ion diffusion distance. In addition, the high conductivity and good mechanical strength of NF make it an ideal current collector for a supercapacitor. Encouraged by these advantages, designers have put considerable effort into the utilization of NF in 3D electrodes.^{31–33} The material has been used as the template for the development of a number of 3D electrodes with

^aCollege of Material Science and Engineering, Central South University of Forestry and Technology, Changsha 410004, P. R. China. E-mail: wuyq0506@126.com

^bHunan Province Key Laboratory of Materials Surface & Interface Science and Technology, Central South University of Forestry and Technology, Changsha 410004, P. R. China. E-mail: xly1516@126.com

^cMOE of the Key Laboratory of Bioinorganic and Synthetic Chemistry, School of Chemistry, Sun Yat-Sen University, Guangzhou 510275, P. R. China. E-mail: luxh6@mail.sysu.edu.cn

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra01164a



excellent electrochemical properties, through methods such as hydrothermal reaction, electrochemical synthesis, and chemical vapor deposition (CVD).^{34–37} However, the fabrication processes involve many complex steps, rigorous conditions, and the use of expensive equipment, which have severely impeded the practical application of such electrodes. Furthermore, these methods involve only the deposition or growing of the substances on the NF frameworks and do not favor the uniform distribution of the substances nor enhance their loading capacity owing to the limited specific surface area and poor versatility of NF. Therefore, the use of NF for the simple and scalable fabrication of high-performance 3D electrodes remains a significant challenge.

In this work, we propose a simple, effective, and generally applicable method to rationally design and fabricate advanced 3D electrodes. Unlike conventional methods for producing 3D electrodes by chemical vapor deposition or hydrothermal reaction using an NF template/substrate, the method utilizes an MCN with a 3D “skin-framework” network as the electroactive material platform. On the one hand, NF is used as a skeletal support and affords the electrodes good mechanical properties and superb electrical conductivity. On the other hand, CNF is not only used to facilitate the uniform dispersion of untreated MWCNT but favors the formation of a stable MWCNT/CNF “skin”, thus guaranteeing the introduction of active materials. Consequently, the obtained MnO₂-MCN and AC-MCN electrodes not only possess a high areal capacity, but also an outstanding discharge rate and low AC impedance. Moreover, an as-fabricated MnO₂-MCN//AC-MCN solid-state ASC device developed with such electrodes can be reversibly charged/discharged at high voltages of up to 1.8 V and has very good high-volume energy and power densities and excellent discharge and cycle performance. Therefore, this work provides a promising strategy for the rational design and fabrication of high-performance 3D electrodes for potential application in portable electronic devices.

2 Materials and method

2.1 Chemicals and materials

All the reagents and materials used in this work were used as received without further purification. The CNFs were prepared using our previously described method with minor modifications. MWCNTs (TNIM2, d8–15 nm) were purchased from the Chengdu Institute of Organic Chemistry. AC was supplied by Fujian Xinsen Carbon Limited. Nickel foam was supplied by Changsha Lyrun Material Co., Ltd. Sodium sulfate (Na₂SO₄), polyvinyl alcohol (PVA), lithium chloride (LiCl), potassium permanganate (KMnO₄), hydrochloric acid (HCl), and anhydrous ethanol were obtained from the Sinopharm Chemical Reagent Company. Deionized water was used as prepared.

2.2 Preparation of 3D network of MCN and MN hybrid materials

The typical preparation procedure was as follows. Commercial NF measuring 16.0 cm × 13.5 cm was soaked in acetone and

ethanol for 6 h and then washed several times with deionized water. It was then dried in a vacuum at 60 °C and weighed, with the weight denoted by W_0 . The pretreated NF was subsequently immersed in a homogeneously dispersed solution of MWCNT and CNF (wt% = 0.5%) with a mass ratio of 1.5 : 1. After a specified time, the NF was removed from the solution and frozen for 24 h in a freeze-dryer at –50 °C and then vacuum dried at 4.5 Pa for 3 days. This soaking and freeze-drying cycle was repeated 3–5 times, after which the final weight of the NF was taken, with the result denoted by W_1 . The loading capacity of the MWCNT/CNF per unit area of MCN was then calculated as $(W_1 - W_0)/S_{\text{NF}}$, where S_{NF} is the area of the NF. The unit area loading capacity was determined to be 5.6 mg cm⁻².

In order to demonstrate the importance of CNF for the dispersion of MWCNTs and the structural stability of MWCNT/CNF aerogel films, we have prepared MN hybrid materials using the same procedure as MCN. Except for impregnation of MN with ultrasonically dispersed MWCNT solution (wt% = 0.5%).

2.3 Preparation of the MnO₂-MCN and AC-MCN 3D electrodes

(1) The preparation of the MnO₂-MCN electrode was typically as follows. An MCN measuring 6 cm × 6 cm was used as the platform for loading the MnO₂ electroactive material to match the AC-MCN negative electrode. Based on our previous work,²⁴ the MnO₂-MCN positive electrodes were prepared using different reaction times (4, 5, 6, and 7 h) and the optimum time was determined for 6 h (Fig. S1†). All the samples were washed using ethanol and deionized water in turn and then vacuum dried at 60 °C. They were subsequently subjected to heat treatment at 220 °C for 2 h.

(2) The preparation of the AC-MCN electrode was typically as follows. Firstly, the pretreated and weighed NF (denoted by W_0) was immersed in a homogeneously dispersed solution of MWCNT, CNF, and AC (wt% = 1.43%) with a mass ratio of 1 : 1 : 8 for a certain time. Then, the NF was removed from the solution, frozen for 24 h in a freeze-dryer at –50 °C, and then vacuum dried at 4.5 Pa for 3 days. This soaking and freeze-drying cycle was repeated 4–6 times, after which the final weight of the NF was taken, with the result denoted by W_2 . The loading capacity of the MWCNT/CNF/AC per unit area of MCN was then calculated as $(W_2 - W_0)/S_{\text{NF}}$, where S_{NF} is the area of the NF. The mass per unit area of the load AC was determined to be 6.88 mg cm⁻².

2.4 Materials characterization

Scanning electron microscopy (SEM, MIRA3) and transmission electron microscopy (TEM, JEM-2100F) were used for the morphological and microstructural characterization of the electrode materials and electrode sheets. X-ray photoelectron spectrometry (XPS) (Escalab 250Xi, Thermo Fisher Scientific) was used to analyze the chemical composition and element valence state of the electrode surfaces, while atomic absorption spectrometry (ICE3000, Thermo Fisher Scientific) was used to evaluate the loading of MnO₂ onto the electrode. The contact angles of Na₂SO₄ electrolyte (1 M) drops on the surface of the



electrode sheets were measured at 25 °C using a contact angle meter (SL150). Nitrogen gas adsorption was measured using a Micromeritics ASAP 2020M analyzer at 77 K, with all the samples degassed at 150 °C in a vacuum for 8 h.

2.5 Electrochemical measurements

An electrochemical workstation (CHI 660E) was used to perform electrochemical tests on the electrodes and other components. The electrochemical testing of the electrodes was conducted using a three-electrode system with a 1 M Na₂SO₄ solution electrolyte, platinum plate counter electrode, and saturated calomel reference electrode. The working electrode had an effective area of 1.0 cm × 1.0 cm. The voltage window of the positive electrode was 0–0.8 V, while that of the negative electrode was –1 to 0 V. To fabricate a solid-state ASC, a LiCl/PVA gel electrolyte was prepared by adding LiCl (4.24 g) and PVA (2 g) into deionized water (20 mL) and heating at 85 °C for 1 h under constant stirring. Two pieces of each of the produced AC-MCN and MnO₂-MCN electrodes were immersed into the LiCl/PVA solution for 30 min, and then taken out and assembled together with a cellulose separator sandwiched in between them. Upon completion, the assembly was dried for 1 h at 40 °C and packaged with polyester (PET) film to prevent contamination of the device during the electrochemical characterization testing process.

3 Results and discussion

Fig. 1a illustrates the preparation of the 3D MnO₂-MCN and AC-MCN electrodes. As shown in Fig. 1a, the MWCNT/CNF solution

is first introduced into the 3D NF network (Fig. 1b) by a simple pouring process. Due to the fact that CNF easily forms films, the self-assembled MWCNT/CNF aerogel film formed during freeze-drying was thus randomly stacked onto the NF-network (Fig. 1d) to form the 3D-network “skin-framework” structure of the MCN hybrid material. Benefiting from the high specific surface area and versatility of MWCNT (such as its ability to combine with the metal oxide MnO₂ and conductive polymer PPy to improve their discharge rate^{38,39}), CNF has excellent electrolyte adsorption and MWCNT dispersion ability, and NF possesses high electrical conductivity and mechanical strength. Therefore, the large-scale MCN obtained (see Fig. S2†) is promising for use as an electroactive platform for fabrication of high-performance 3D electrodes. Furthermore, the MCN electrode sheet exhibits good flexibility and bending performance (Fig. 1f). In addition, benefiting from the advantages of the MCN platform, we have successfully prepared a high-performance AC-MCN electrode using a similar strategy.

To demonstrate the advantages of MCN as an electroactive material platform, its microstructure and morphology were characterized by SEM and TEM. As shown in Fig. 1b, NF has an inter-connected 3D network, which affords MCN a unique 3D porous “skin-framework” structure in which the 3D NF network constitutes the “framework” and the MWCNT/CNF aerogel film is the “skin” (Fig. 1d). This unique structure significantly enhances the conductivity and specific surface area of the MCN platform because the MWCNTs are not only a carrier for electroactive materials (such as MnO₂) but also act as a conductive agent. In fact, the N₂ adsorption–desorption measurement

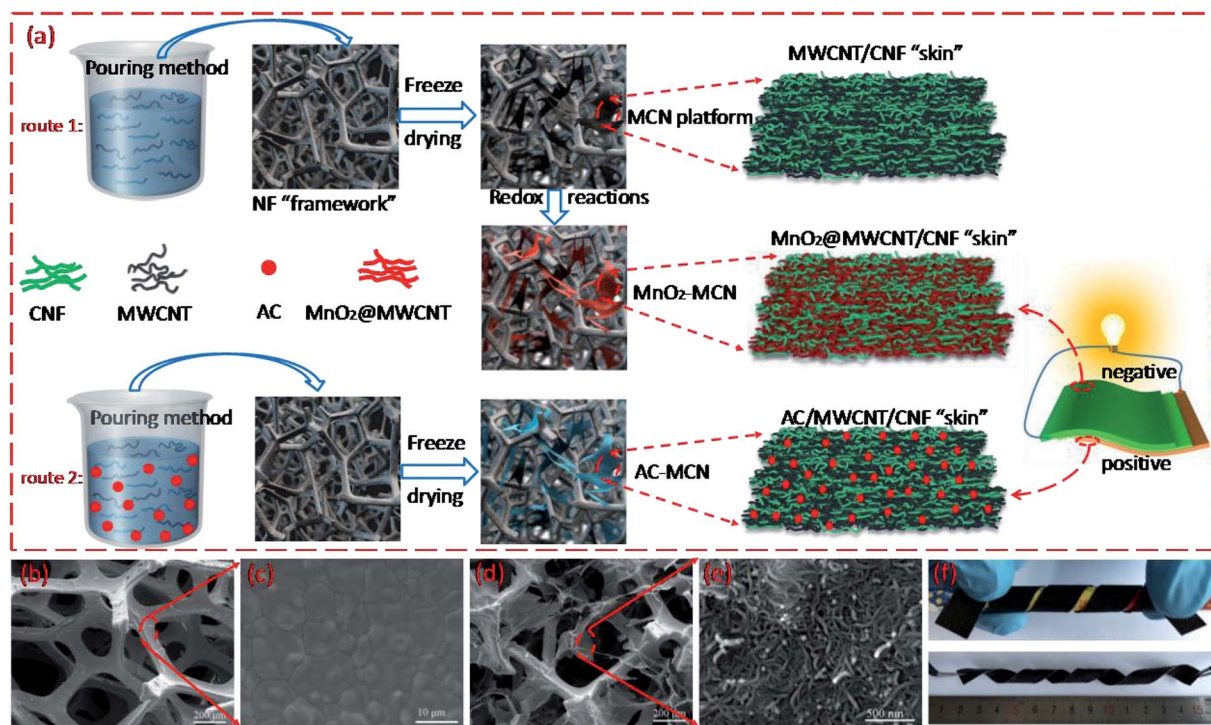


Fig. 1 (a) Illustration of the preparation of the MnO₂-MCN (route 1) and AC-MCN (route 2) electrodes, and construction of the solid-state ASC device; SEM images of the (b and c) NF and (d and e) MCN hybrid material with different magnifications; and (f) demonstration of the flexibility and bending properties of the MCN electrode sheet.



showed that the specific surface area of MCN was much larger than NF (Fig. S3†). To demonstrate the scalability of MCN as an electroactive platform for the fabrication of high-performance electrodes, MnO₂-MCN electrodes were produced by redox reactions ($4\text{MnO}_4^- + 3\text{C} + \text{H}_2\text{O} \rightarrow 4\text{MnO}_2 + \text{CO}_3^{2-} + 2\text{HCO}_3^-$).⁴⁰ As shown in Fig. 2a and b, the MnO₂@MWCNT/CNF “skin” is still entirely stacked onto the 3D “framework” of NF after redox reaction with KMnO₄ solution. Moreover, a rougher surface loaded with numerous nanoparticles can be observed in the network structure (Fig. 2c). These nanoparticles are assumed to be MnO₂ formed *in situ* at the MWCNT surface during the redox process. In addition, Fig. 2g and h show that MnO₂ nanoparticles ranging between 3 and 5 nm are loaded onto the MWCNT surface. To further demonstrate the scalability of MCN as an electroactive platform for the construction of high-performance electrodes, the AC-MCN electrode is prepared by a similar strategy (route 2). As shown in Fig. 2d and e, the AC particles are uniformly distributed in the MWCNT/CNF aerogel film. The high-magnification SEM image illustrates that the MWCNT and CNF were tightly wound on the surface of the AC particles (Fig. 2f, indicated by an arrow), providing an effective pathway for electron transport and ion diffusion in the AC electroactive materials. These results show that, compared with NF, the use of MCN as a platform for electroactive materials is more broadly applicable and offers several structural advantages for the fabrication of high-performance 3D electrodes.

As stated above, the unique 3D porous “skin-framework” network of MCN is the key to its role as a platform for electroactivity materials. This is because the homogeneously distributed untreated MWCNT in the MWCNT/CNF aerogel film is the carrier and conductive pathway of the active materials MnO₂ and AC. Hence, the dispersibility of the MWCNT and the structural stability of the MWCNT/CNF “skin” directly affect the performance of MCN as an electroactive platform for fabrication of 3D electrodes. To demonstrate the importance of CNF for the two aforementioned factors, we compared the microstructure and structural stability of the MCN platform and

MWCNT-NF hybrid materials (denoted as MN). As shown in Fig. 3a and b, the MWCNT/CNF “skin” of the MCN platform well maintains its thin film morphology before and after ultrasonic treatment in ethanol and water mixed solution, indicating excellent structural stability. Conversely, the MWCNT in MN hybrid barely forms a large-scale and film-like “skin” (Fig. 3c) and has poor structural stability. After ultrasonic treatment, the film-like “skin” obviously peels off from the NF framework (Fig. 3d). The desirable performance of MCN is because CNF not only evenly distributes untreated MWCNT, but its excellent film-forming property is also beneficial to MWCNT/CNF self-assembly to form a structurally stable aerogel film.⁴¹ However, without the auxiliary dispersion of CNF, it is difficult to uniformly disperse untreated MWCNT in a water solution, much less achieve the formation of a stable films on the 3D network of NF during freeze-drying. Moreover, after 10 minutes of ultrasound treatment for the MCN sheet, its solution remained clear and transparent (Fig. 3e, Video S1†), whereas after only 1 minute of ultrasound treatment for the MN sheet, its solution turned black (Fig. 3f, Video S1†) owing to the falling off MWCNT from the NF framework. This further demonstrates the superior structural stability of the MWCNT/CNF “skin” stacked onto the MCN platform. In addition, it is worth noting that CNF also effectively improves the wettability of the MnO₂-MCN and AC-MCN electrodes (Fig. 3g, Video S2†), thereby facilitating rapid diffusion of the electrolyte and improving the performance of the electrodes.

To further elucidate the advantages of MCN in terms of its microstructure, the porous structures of NF, MCN, and MnO₂-MCN were analyzed by N₂ adsorption. As shown in Fig. 3h, NF has an atypical type II adsorption-desorption curve. Conversely, those of MCN and MnO₂-MCN hybrids are combinations of type I and type IV, indicating the presence of micropores and mesopores.⁴² In addition, the adsorption-desorption isotherms of MCN and MnO₂-MCN have a H3 hysteresis loop at relative pressures of 0.8–1.0, suggesting the existence of macropores.⁴³ The inset of Fig. 3h shows the corresponding pore size

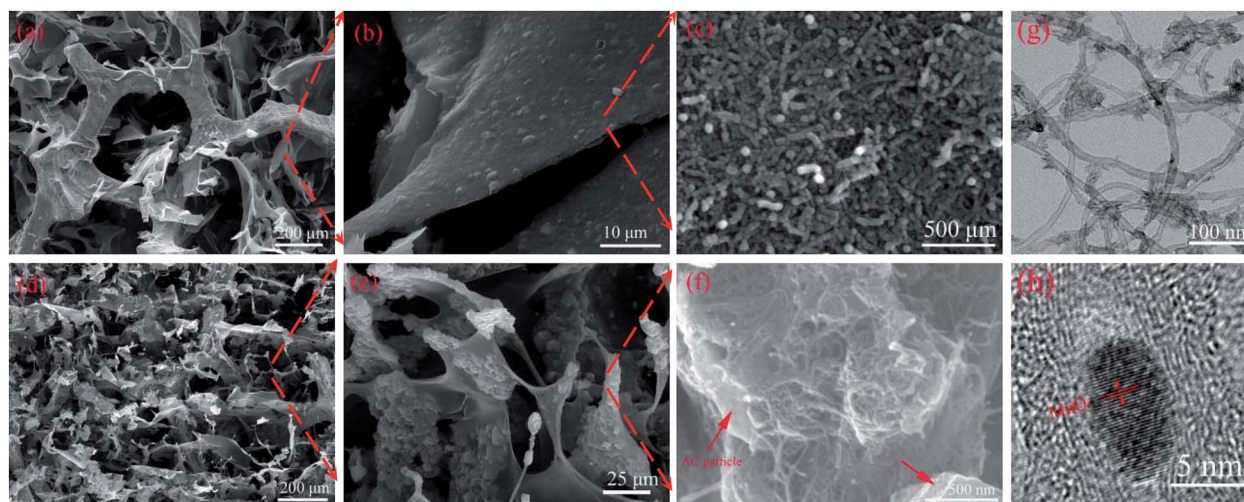


Fig. 2 SEM images of (a–c) MnO₂-MCN, (d–f) AC-MCN electrodes with different magnifications; and TEM images of (g and h) MnO₂-MCN electrode with different magnifications.



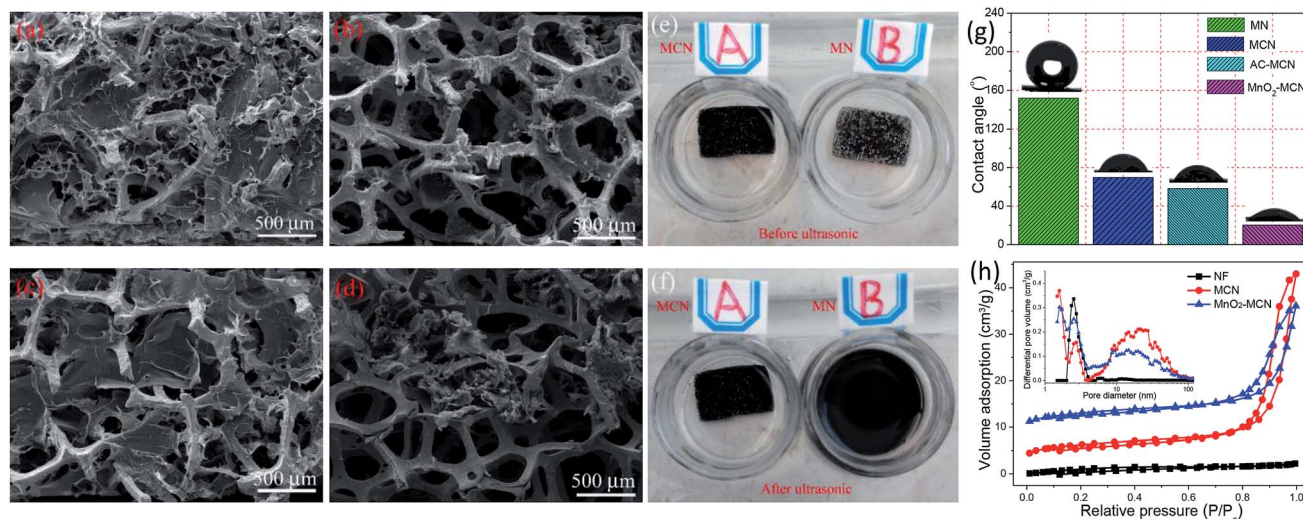


Fig. 3 SEM images of MCN (a) before and (c) after ultrasonic treatment; SEM images of MN (b) before and (d) after ultrasonic treatment; digital pictures of MCN sheet (1 cm × 1.5 cm) and MN sheet (1 cm × 1.5 cm) (e) before and (f) after ultrasonic treatment in ethanol and water mixed solution; (g) histogram and corresponding digital photos (inset) of the contact angle of the MN, MCN, AC-MCN and MnO₂-MCN electrode materials; and (h) N₂ adsorption–desorption isotherm of NF, MCN, and MnO₂-MCN and their corresponding DFT pore size distribution (inset).

distribution (PSD) curves of NF, MCN, and MnO₂-MCN obtained by the density functional theory (DFT). A characteristic peak can be observed at 2.5 nm for all the samples, primarily due to the cracks present on the NF surfaces (Fig. 1c). Conversely, MCN and MnO₂-MCN have a sharp characteristic peak at 1.6 nm, possibly due to the pores formed by the uniform distribution of MWCNT and CNF (Fig. 1e and 2c). The two materials also have multiple broad characteristic peaks between 9.3 and 34.3 nm, mainly attributed to random pores of various sizes in MCN due to the MWCNT/CNF thin-film stacking. It should be noted that macropores are not reflected in a PSD curves, although they can be verified by SEM observation (Fig. 1d and 2a). These results confirm the higher specific surface area and beneficial porous structure of MCN that enhance its use as an active material platform for the fabrication of high-performance electrodes.

XPS and EDS were used to determine the composition and chemical state of the MnO₂-MCN electrode. Fig. S4† and 4 show the full XPS spectrum of the MnO₂-MCN electrode and the corresponding Mn 2p and O 1s XPS spectra, respectively. The spectra confirm the presence of manganese oxide in the electrode. Fig. 4a reveals two characteristic peaks of Mn 2p at 653.09 and 642.39 eV, which are identical with those of Mn 2p_{1/2} and Mn 2p_{3/2} reported for Mn⁴⁺ in the literature.^{44,45} This indicates the predominant occurrence of Mn as Mn⁴⁺ in the electrode. The three characteristic peaks at 533.6, 531.9, and 530.2 eV in the fitted O 1s spectra (Fig. 4b) correspond to Mn–O, Mn–OH, and H–O–H, respectively. This agrees well with the literature^{46,47} and proves that MnO₂ is loading on the MnO₂-MCN electrode after the redox reaction process. Fig. 4c shows the distributions of Mn and O in the MnO₂-MCN electrode, as determined by EDS. It can be seen that both elements are uniformly distributed in the electrode, further proving the even loading of nano-MnO₂ formed *in situ* by redox processes when MCN is used as the electroactive material platform.

To prove the suitability of MCN for the fabrication of high-performance electrodes, a MnO₂-MCN electrode was successfully prepared by redox reaction using MCN as the platform. Fig. 5a compares the cyclic voltammogram (CV) curves of the MCN and MnO₂-MCN electrodes for a scanning rate of 100 mV s⁻¹. As expected, the MnO₂-MCN electrode exhibits a higher current density than the MCN electrode, its capacitance being mainly due to the loaded nano-MnO₂. Further, the galvanostatic charge/discharge (GCD) curves of the two electrodes for a current density of 5 mA cm⁻² (Fig. S5a†) clearly shows that the MnO₂-MCN electrode has a longer discharge time, further substantiating its excellent capacitive properties. The CV curve of the MnO₂-MCN electrode also preserves its quasi-rectangular shape for scanning rates of 5–100 mV s⁻¹ (Fig. 5b), suggesting ideal capacitive characteristics and outstanding discharge performance of the electrode.

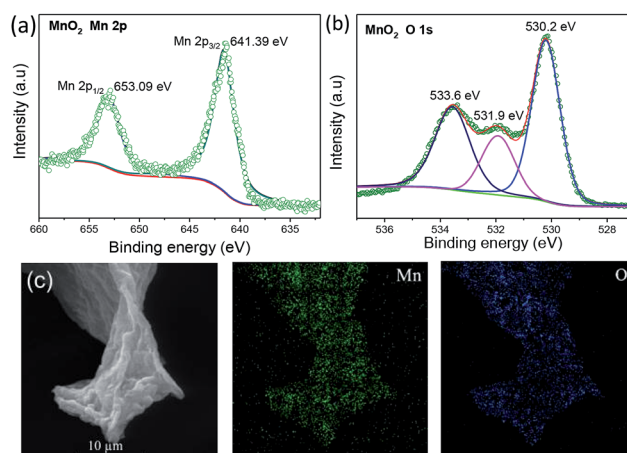


Fig. 4 XPS spectra and peak fitting of the MnO₂-MCN electrode (a) Mn and (b) O, and (c) EDS maps of Mn and O for the MnO₂-MCN electrode.



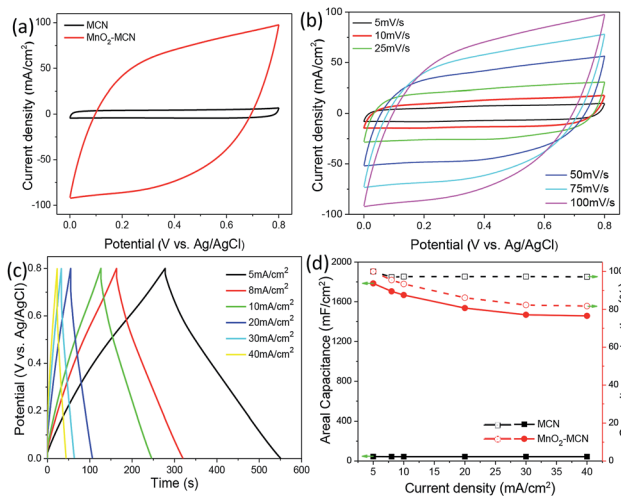


Fig. 5 (a) CV curves of the MCN and MnO₂-MCN electrodes obtained using a scanning rate of 100 mV s⁻¹; (b) CV curves of the MnO₂-MCN electrode for scanning rates of 5–100 mV s⁻¹; (c) GCD curves of the MnO₂-MCN electrode for current densities of 5–40 mA cm⁻²; and (d) areal capacitance and capacitance retention rate of the MCN and MnO₂-MCN electrodes as functions of the current density.

As shown in Fig. 5c, the GCD curves of the MnO₂-MCN electrode for different current densities all have a symmetrical and triangular shape, revealing a good capacitive behavior and reversible charging–discharging. Fig. 5d shows the relationships between the areal capacity and capacity retention of the MCN and MnO₂-MCN electrodes and the discharge current density. The areal capacity of the MnO₂-MCN electrode of 1784.8 mF cm⁻² (equal to 469.7 F g⁻¹) at a current density of 5 mA cm⁻² is obviously much higher than that of the MCN electrode, indicating a much higher discharge performance of the former. The capacity retention of the MnO₂-MCN electrode is also as high as 81.7% at a current density of 40 mA cm⁻². This is mainly because the uniformly distributed MWCNT in the MnO₂-MCN electrode functions as the MnO₂ nanoparticles carrier and effectively improves the discharge performance by affording excellent electrical conductivity;²⁴ furthermore, its high specific surface area also facilitates the even loading of MnO₂ nanoparticles. From Nyquist impedance plots (Fig. S5b and c[†]), both the MCN and MnO₂-MCN electrodes were observed to have very small impedances (~1 ohm), attributable to the high conductivity of the 3D network of NF. The MnO₂-MCN electrode maintains 92.3% of its initial capacitance after 3000 charge/discharge cycles at a current density of 10 mA cm⁻² (Fig. S5d[†]), demonstrating its good cycle performance. These results provide further evidence of the extraordinary advantages of MCN as an electroactive material platform for fabrication of high-performance 3D electrodes.

To further evaluate the feasibility of the MCN platform for the fabrication of high-performance 3D electrodes, we prepared an AC-MCN electrode by a similar strategy to match the MnO₂-MCN electrode to assemble a solid-state ASC. Fig. 6a shows CV curves of the MCN and AC-MCN electrodes collected at a scanning rate of 100 mV s⁻¹. It can be clearly seen that the current

density of the AC-MCN electrode is much higher than that of the MCN electrode. This reveals the superior electrochemical capacity of the AC-MCN electrode. The CV curve of the AC-MCN electrode (Fig. 6b) exhibits a symmetrical quasi-rectangular form when the scan rate increases from 5 to 100 mV s⁻¹, suggesting that the electrode has ideal capacitive characteristics and outstanding discharge performance. Further, the GCD curves of the MCN and AC-MCN electrodes for a current density of 5 mA cm⁻² (Fig. S6a[†]) clearly show that the AC-MCN electrode has a longer discharge time, further substantiating its excellent capacitive properties.

Fig. 6c represents the GCD curves of the AC-MCN electrode collected at various current densities from 5 to 40 mA cm⁻². All the GCD curves exhibit good symmetry, further confirming the good capacitive behavior of the AC-MCN electrode. Fig. 6d shows the relationships between the areal capacity and capacity retention rate of the AC-MCN electrode with the discharge current density. Both the areal capacity and discharge performance of the AC-MCN electrode for a given current density are clearly superior. For example, the areal capacity of the AC-MCN electrode for a current density of 5 mA cm⁻² is 868.8 mF cm⁻² (equal to 126.3 F g⁻¹), which is much higher than that of MCN (53.4 mF cm⁻²). It should also be noted that the capacity retention rate of the AC-MCN electrode is as high as 74.3% at a current density of 40 mA cm⁻². In addition, the AC-MCN electrode exhibits good electrochemical durability (Fig. S6b[†]); after 3000 charge–discharge cycles at a current density of 10 mA cm⁻², the capacitance retention remains above 89.3%. Furthermore, the AC impedance spectrum exhibits a low resistance (~1 ohm) for this electrode (Fig. S6c and d[†]), again confirming that the 3D MCN is an excellent platform for fabricating high-performance electrodes.

To assemble the solid-state ASC device, we used the MnO₂-MCN as the positive electrode and AC-MCN as the negative electrode. Prior to assembly, it was necessary to consider the

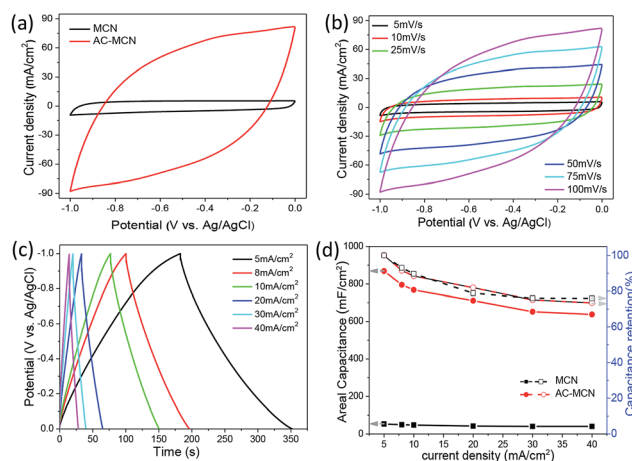


Fig. 6 (a) CV curves of the MCN and AC-MCN electrodes collected at a scanning rate of 100 mV s⁻¹; (b) CV curves of the AC-MCN electrode for scanning rates of 5–100 mV s⁻¹; (c) GCD curves of the AC-MCN electrode for current densities of 5–40 mA cm⁻²; and (d) areal capacitance and capacitance retention rate of the MCN and AC-MCN electrodes as functions of the current density.



charge balance between the anode and cathode during the assembly of the ASC. The balance can be described by the following equation:⁴⁷

$$C_s^+ \times \Delta V^+ \times S^+ = C_s^- \times \Delta V^- \times S^- \quad (1)$$

where C_s , ΔV , and S are the areal capacity, potential window, and area of an electrode, respectively. Fig. 7a shows the CV curves of the AC-MCN cathode and MnO₂-MCN anode obtained using a scanning rate of 50 mV s⁻¹. Based on eqn (1), the ratio between the charges of the positive and negative electrodes was determined to be approximately 1 : 1. Using the stable potential windows of the AC-MCN and MnO₂-MCN electrodes of -1.0 to 0.0 V and 0.0–0.8 V, respectively (Fig. 7a), the operating voltage of the as-fabricated MnO₂-MCN//AC-MCN solid-state ASC was determined to be as high as 1.8 V (Fig. 7b). Fig. S7a† shows the CV curves of the ASC device for various scanning rates and a voltage window of 1.8 V. All the curves can be observed to be symmetric and rectangle-like, indicative of good reaction reversibility and typical capacity characteristics of the ASC. This would enable the development of solid-state ASC devices with high energy densities.

Fig. 7c shows GCD curves of the MnO₂-MCN//AC-MCN ASC device for different current densities. The symmetrical and triangular shape curves demonstrate the outstanding capacity properties and rapid charging/discharging of the device. Fig. S7b† shows the GCD curves of the device for voltage windows of 1.0–1.8 V and a discharge current density of 10 mA cm⁻². All the curves are roughly symmetrical and triangular, further confirming the ASC device as an ideal capacitor. As can be seen from Fig. 7d, the MnO₂-MCN//AC-MCN ASC device has

its highest volumetric capacitance of 9.83 F cm⁻³ under a discharge current density of 5 mA cm⁻² and voltage window of 1.8 V, substantially higher than recently reported values for other ASCs such as a Ni/Co₂NiO₄-CP//Ni/AC-CP-ASC (4.48 F cm⁻³ at 4 mA cm⁻²),⁴⁸ PF/Ni/MnO₂//PF/Ni/AC-ASC (1.4 F cm⁻³ at 2.5 mA cm⁻²),⁴⁹ MnO₂//Ti-Fe₂O₃@PEDOT-ASC (1.15 F cm⁻³ at 1 mA cm⁻²),⁵⁰ MnO₂//Fe₂O₃-ASC (1.5 F cm⁻³ at 2 mA cm⁻²),⁵¹ and MnO₂@CNT//PPY@CNT-ASC (2.2 F cm⁻³ at 2 mA cm⁻²).⁵² When the current density increases from 5 to 40 mA cm⁻², the capacity retention of the MnO₂-MCN//AC-MCN ASC device is maintained as high as 75.4%, validating its excellent discharge performance. This is mainly because the MnO₂-MCN and AC-MCN electrodes consist of a unique multilayer structure (Fig. 7d and S8†), with the NF-formed 3D network conductive path between the multilayer structures also affording the electrodes excellent electrical conductivity. As can be seen from Fig. S7c,† the ASC retains 81.6% of its initial capacitance after 5000 charge/discharge cycles, revealing a good cycle performance. The electrochemical impedance spectra of the assembled ASC device were analyzed to evaluate the electrochemical behavior. As shown in Fig. 7e, the Nyquist impedance plot of the MnO₂-MCN//AC-MCN ASC is almost parallel to the imaginary linear axis, once again indicating ideal capacitance properties of the device. The charge-transfer resistance is also very small (~1 ohm), implying rapid charge transfer.

Fig. 7f compares the volumetric power and energy density of the assembled MnO₂-MCN//AC-MCN ASC device with those of other previously reported ASC devices.^{48–54} Notably, the present ASC exhibits a high energy density, high power density, and excellent discharge performance. For example, its volumetric energy is as high as 4.25 mW h cm⁻³ at a current density of 5 mA

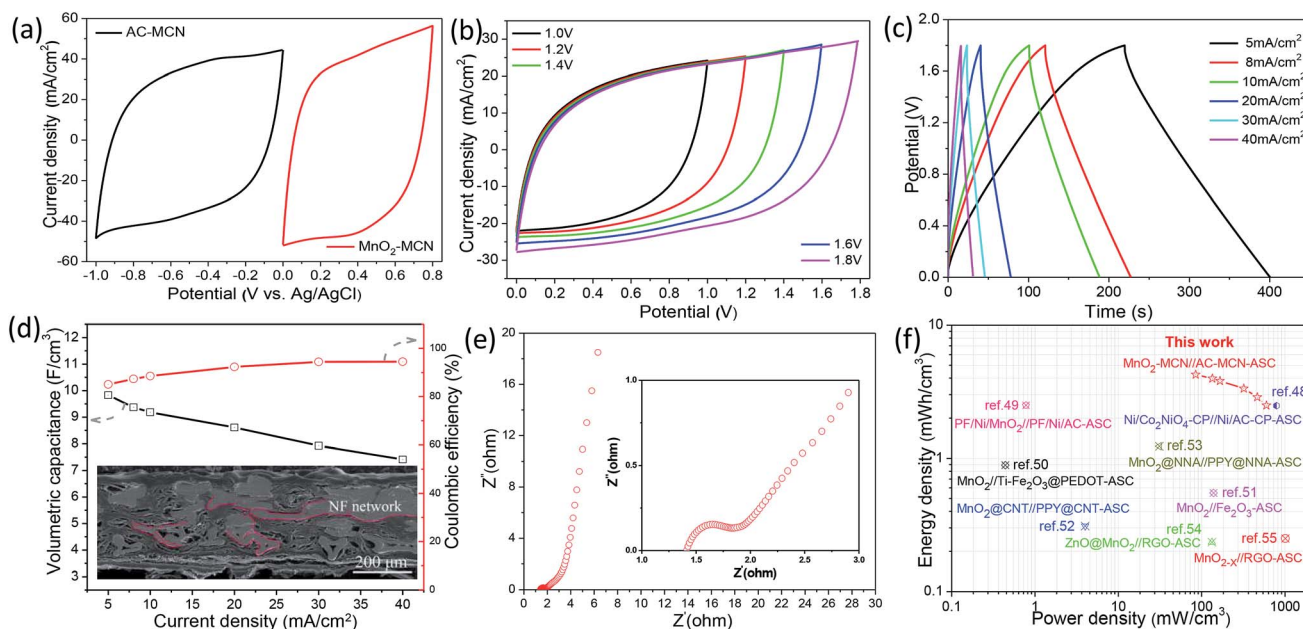


Fig. 7 (a) CV curves of the AC-MCN and MnO₂-MCN electrodes obtained using a scanning rate of 50 mV s⁻¹; (b) CV curves of the solid-state MnO₂-MCN//AC-MCN ASC device collected at a scanning rate of 50 mV s⁻¹ and different voltage windows; (c) GCD curves of the ASC device for current densities of 5–40 mA cm⁻²; (d) volumetric capacity and coulombic efficiency as a function of the current density, and (inset) cross-sectional SEM image of the ASC device; (e) Nyquist plots of the ASC device; and (f) comparison of Ragone plots of the ASC device with those of other devices in the literature.^{48–55}



cm^{-2} , higher than those of many of the other ASC devices such as the Ni/Co₂NiO₄-CP//Ni/AC-CP ASC (2.48 mW h cm^{-3} at 4 mA cm^{-2}),⁴⁸ PF/Ni/MnO₂//PF/Ni/AC ASC (0.78 mW h cm^{-3} at 2.5 mA cm^{-2}),⁴⁹ MnO₂//Ti-Fe₂O₃@PEDOT ASC (0.89 mW h cm^{-3} at 1 mA cm^{-2}),⁵⁰ MnO₂//Fe₂O₃ ASC (0.55 mW h cm^{-3} at 2 mA cm^{-2}),⁵¹ MnO₂@CNT//PPy@CNT ASC (0.31 mW h cm^{-3} at 2 mA cm^{-2}),⁵² MnO₂@NNA//PPy@NNA ASC (1.23 mW h cm^{-3} at 1 mA cm^{-2}),⁵³ ZnO@MnO₂//RGO ASC (0.234 mW h cm^{-3} at 2 mA cm^{-2}),⁵⁴ and MnO_{2-x}//RGO-ASC (0.25 mW h cm^{-3} at 2 mA cm^{-2}).⁵⁵ Even for a current density of 40 mA cm^{-2} , the volumetric energy and power density of the present ASC were still as much as 2.50 mW h cm^{-3} and 598.85 mW cm^{-3} , respectively. Two of the present ASCs (dimension: 1 cm × 1 cm) connected in series were used to continuously light up a 3.0 V LED lamp (Fig. S7c,† inset), which also demonstrates the high energy density of the as-fabricated ASC device. The above results demonstrate the very promising potential of MnO₂-MCN and AC-MCN electrodes prepared using an MCN electroactive material platform for use in the development of high-performance ASC devices.

4 Conclusions

In summary, a method for using MCN with a 3D network “skin-framework” structure as the platform of the electroactive materials of high-performance MnO₂-MCN and AC-MCN 3D electrodes was proposed in this work. Compared with conventional methods such as the use of CVD and hydrothermal reaction, the proposed method is simple, involves mild reaction conditions, effective, inexpensive, and scalable. Benefiting from the unique 3D “skin-framework” architecture of MCN, the obtained electrodes offer the following advantages: (1) superb electrical conductivity and rapid electrolyte diffusion and (2) stable “skin” and large interfacial contact area, which provide a convenient and effective platform for the electroactive materials. In addition, the as-obtained MnO₂-MCN anode and AC-MCN cathode exhibit high areal capacities and excellent electrical conductivities and rate performance. The MnO₂-MCN and AC-MCN electrodes exhibited capacitances of 1784.8 and 868.8 mF cm^{-2} , respectively, for a current density of 5 mA cm^{-2} . A 1.8 V solid-state ASC assembled using the electrodes also exhibited a very high volumetric capacity of 9.83 F cm^{-3} and maximum energy density of 4.25 mW h cm^{-3} . These findings confirm the very high potential of using electrodes based on MCN platforms for the development of high-energy ASCs for portable electronics.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

L. Y. Xia and S. H. Hu contributed equally to this work. The authors are grateful for the support of the National Natural Science Foundation of China (31530009, 31200438), National

Key R&D Program of China (2017YFD0600804), Hunan Provincial Natural Science Foundation of China (2015JJ2199).

Notes and references

- L. Dong, C. Xu, Y. Li, Z. Pan, G. Liang, E. Zhou, F. Kang and Q. H. Yang, *Adv. Mater.*, 2016, **28**, 9313–9319.
- L. Dong, C. Xu, Y. Li, C. Wu, B. Jiang, Q. Yang, E. Zhou, F. Kang and Q. H. Yang, *Adv. Mater.*, 2016, **28**, 1675–1681.
- Z. S. Wu, K. Parvez, S. Li, S. Yang, Z. Liu, S. Liu, X. Feng and K. Mullen, *Adv. Mater.*, 2015, **27**, 4054–4061.
- X. Wang, X. Lu, B. Liu, D. Chen, Y. Tong and G. Shen, *Adv. Mater.*, 2014, **26**, 4763–4782.
- H. Wu, Y. Huang, F. Xu, Y. Duan and Z. Yin, *Adv. Mater.*, 2016, **28**, 9881–9919.
- L. Wen, F. Li and H. M. Cheng, *Adv. Mater.*, 2016, **28**, 4306–4337.
- S. Liu, Y. Yin, K. S. Hui, K. N. Hui, S. C. Lee and S. C. Jun, *Adv. Sci.*, 2018, **5**, 1800733.
- D. Ghosh, J. Lim, R. Narayan and S. O. Kim, *ACS Appl. Mater. Interfaces*, 2016, **8**, 22253–22260.
- D. Yu, K. Goh, Q. Zhang, L. Wei, H. Wang, W. Jiang and Y. Chen, *Adv. Mater.*, 2014, **26**, 6790–6797.
- K. P. Annamalai, L. Liu and Y. Tao, *Adv. Mater. Interfaces*, 2017, **4**, 1700219.
- P. Wu, S. Cheng, M. Yao, L. Yang, Y. Zhu, P. Liu, O. Xing, J. Zhou, M. Wang, H. Luo and M. Liu, *Adv. Funct. Mater.*, 2017, **27**, 1702160.
- X. Lu, M. Yu, G. Wang, Y. Tong and Y. Li, *Energy Environ. Sci.*, 2014, **7**, 2160–2181.
- J.-X. Feng, S.-H. Ye, X.-F. Lu, Y.-X. Tong and G.-R. Li, *ACS Appl. Mater. Interfaces*, 2015, **7**, 11444–11451.
- C. Yang, J. Shen, C. Wang, H. Fei, H. Bao and G. Wang, *J. Mater. Chem. A*, 2014, **2**, 1458–1464.
- A. Bora, K. Mohan, S. Doley and S. K. Dolui, *ACS Appl. Mater. Interfaces*, 2018, **10**, 7996–8009.
- Z. Gao, W. Yang, J. Wang, N. Song and X. Li, *Nano Energy*, 2015, **13**, 306–317.
- Z. Pan, Y. Qiu, J. Yang, F. Ye, Y. Xu, X. Zhang, M. Liu and Y. Zhang, *Nano Energy*, 2016, **26**, 610–619.
- L. Wang, H. Yang, X. Liu, R. Zeng, M. Li, Y. Huang and X. Hu, *Angew. Chem., Int. Ed.*, 2017, **56**, 1105–1110.
- F. Zhang, T. Zhang, X. Yang, L. Zhang, K. Leng, Y. Huang and Y. Chen, *Energy Environ. Sci.*, 2013, **6**, 1623–1632.
- T. T. Nguyen, J. Balamurugan, N. H. Kim and J. H. Lee, *J. Mater. Chem. A*, 2018, **6**, 8669–8681.
- K. Ghosh, C. Y. Yue, M. M. Sk and R. K. Jena, *ACS Appl. Mater. Interfaces*, 2017, **9**, 15350–15363.
- Y. Yang, G. Ruan, C. Xiang, G. Wang and J. M. Tour, *J. Am. Chem. Soc.*, 2014, **136**, 6187–6190.
- Y. Shao, M. F. El-Kady, C.-W. Lin, G. Zhu, K. L. Marsh, J. Y. Hwang, Q. Zhang, Y. Li, H. Wang and R. B. Kaner, *Adv. Mater.*, 2016, **28**, 6719–6726.
- L. Xia, X. Li, X. Wu, L. Huang, Y. Liao, Y. Qing, Y. Wu and X. Lu, *J. Mater. Chem. A*, 2018, **6**, 17378–17388.
- X. Yang, Z. Lin, J. Zheng, Y. Huang, B. Chen, Y. Mai and X. Feng, *Nanoscale*, 2016, **8**, 8650–8657.



- 26 H. Jiang, X. Cai, Y. Qian, C. Zhang, L. Zhou, W. Liu, B. Li, L. Lai and W. Huang, *J. Mater. Chem. A*, 2017, **5**, 23727–23736.
- 27 G. Zhou, L. Li, C. Ma, S. Wang, Y. Shi, N. Koratkar, W. Ren, F. Li and H.-M. Cheng, *Nano Energy*, 2015, **11**, 356–365.
- 28 Y. He, W. Chen, X. Li, Z. Zhang, J. Fu, C. Zhao and E. Xie, *ACS Nano*, 2013, **7**, 174–182.
- 29 V. H. Nguyen and J.-J. Shim, *J. Power Sources*, 2015, **273**, 110–117.
- 30 C. Wu, J. Cai, Q. Zhang, X. Zhou, Y. Zhu, P. K. Shen and K. Zhang, *ACS Appl. Mater. Interfaces*, 2015, **7**, 26512–26521.
- 31 J. Wang, D. Chao, J. Liu, L. Li, L. Lai, J. Lin and Z. Shen, *Nano Energy*, 2014, **7**, 151–160.
- 32 D. Kong, C. Cheng, Y. Wang, J. I. Wong, Y. Yang and H. Y. Yang, *J. Mater. Chem. A*, 2015, **3**, 16150–16161.
- 33 H. Huang, Y. Tang, L. Xu, S. Tang and Y. Du, *ACS Appl. Mater. Interfaces*, 2014, **6**, 10248–10257.
- 34 X. Xiong, D. Ding, D. Chen, G. Waller, Y. Bu, Z. Wang and M. Liu, *Nano Energy*, 2015, **11**, 154–161.
- 35 K. Xu, W. Li, Q. Liu, B. Li, X. Liu, L. An, Z. Chen, R. Zou and J. Hu, *J. Mater. Chem. A*, 2014, **2**, 4795–4802.
- 36 C. Zhou, Y. Zhang, Y. Li and J. Liu, *Nano Lett.*, 2013, **13**, 2078–2085.
- 37 G. Zhu, Z. He, J. Chen, J. Zhao, X. Feng, Y. Ma, Q. Fan, L. Wang and W. Huang, *Nanoscale*, 2014, **6**, 1079–1085.
- 38 R. T. Vinny, K. Chaitra, K. Venkatesh, N. Nagaraju and N. Kathyayini, *J. Power Sources*, 2016, **309**, 212–220.
- 39 K. Liang, T. Gu, Z. Cao, X. Tang, W. Hu and B. Wei, *Nano Energy*, 2014, **9**, 245–251.
- 40 J.-G. Wang, F. Kang and B. Wei, *Prog. Mater. Sci.*, 2015, **74**, 51–124.
- 41 N. Lavoine and L. Bergström, *J. Mater. Chem. A*, 2017, **5**, 16105–16117.
- 42 L. Xia, X. Li, Y. Wu and M. Rong, *ACS Sustainable Chem. Eng.*, 2015, **3**, 1724–1731.
- 43 L. Xia, X. Li, F. Zhu, S. Hu and L. Huang, *J. Phys. Chem. C*, 2017, **121**, 20279–20286.
- 44 S. Zhu, L. Li, J. Liu, H. Wang, T. Wang, Y. Zhang, L. Zhang, R. S. Ruoff and F. Dong, *ACS Nano*, 2018, **12**, 1033–1042.
- 45 Z. Zhang, F. Xiao, L. Qian, J. Xiao, S. Wang and Y. Liu, *Adv. Energy Mater.*, 2014, **4**, 1400064.
- 46 Y. Cheng, S. Lu, H. Zhang, C. V. Varanasi and J. Liu, *Nano Lett.*, 2012, **12**, 4206–4211.
- 47 Z. Pan, M. Liu, J. Yang, Y. Qiu, W. Li, Y. Xu, X. Zhang and Y. Zhang, *Adv. Funct. Mater.*, 2017, **27**, 1701122.
- 48 J.-X. Feng, S.-H. Ye, A.-L. Wang, X.-F. Lu, Y.-X. Tong and G.-R. Li, *Adv. Funct. Mater.*, 2014, **24**, 7093–7101.
- 49 L. Zhang, P. Zhu, F. Zhou, W. Zeng, H. Su, G. Li, J. Gao, R. Sun and C. P. Wong, *ACS Nano*, 2016, **10**, 1273–1282.
- 50 C. Xu, Z. Li, C. Yang, P. Zou, B. Xie, Z. Lin, Z. Zhang, B. Li, F. Kang and C. P. Wong, *Adv. Mater.*, 2016, **28**, 4105–4110.
- 51 Y. Zeng, Y. Han, Y. Zhao, Y. Zeng, M. Yu, Y. Liu, H. Tang, Y. Tong and X. Lu, *Adv. Energy Mater.*, 2015, **5**, 1402176.
- 52 P. Yang, Y. Ding, Z. Lin, Z. Chen, Y. Li, P. Qiang, M. Ebrahimi, W. Mai, C. P. Wong and Z. L. Wang, *Nano Lett.*, 2014, **14**, 731–736.
- 53 Q. Tang, M. Chen, C. Yang, W. Wang, H. Bao and G. Wang, *ACS Appl. Mater. Interfaces*, 2015, **7**, 15303–15313.
- 54 W. Zilong, Z. Zhu, J. Qiu and S. Yang, *J. Mater. Chem. C*, 2014, **2**, 1331–1336.
- 55 T. Zhai, S. Xie, M. Yu, P. Fang, C. Liang, X. Lu and Y. Tong, *Nano Energy*, 2014, **8**, 255–263.

