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# Graphene reinforced carbon nanofiber engineering enhances Li storage performances of germanium oxide†

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The rational design of electrode materials with high power and energy densities, good operational safety, and long cycle life remains a great challenge for developing advanced battery systems. As a promising electrode material for rechargeable batteries, germanium oxide (GeO<sub>2</sub>) shows high capacity, but suffers from rapid capacity fading caused by its large volume variation during charge/discharge processes and poor rate performance owing to low intrinsic electronic conductivity. In this study, a novel one-dimensional (1D) carbon/graphene-nanocable-GeO<sub>2</sub> nanocomposite (denoted as GeO<sub>2</sub>/nanocable) is rationally designed and prepared *via* a facile electrospinning method. Specifically, amorphous carbon and graphene spontaneously construct a nanocable structure, in which graphene acts as the “core” and amorphous carbon as the “shell”, and GeO<sub>2</sub> nanoparticles are encapsulated in the nanocable. The graphene “core” promises good electrical conductivity while the amorphous carbon “shell” guarantees fast Li ions diffusion. When tested as an anode material for rechargeable lithium ion batteries, the GeO<sub>2</sub>/nanocable exhibits remarkable Li storage performance, including high reversible capacity (900 mA h g<sup>-1</sup>), high capacity retention (91% after 100 cycles), and good rate performance (595 mA h g<sup>-1</sup> at 5000 mA g<sup>-1</sup>).

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Lithium ion batteries (LIBs) require longer cycle lifetimes, and higher energy density and rate capability in order to satisfy the increasing popularity of electric vehicles (EVs) and hybrid vehicles (HEVs). Nevertheless, the current commercial LIBs using graphite anode materials are unable to meet this ever-growing demand because of their relatively low capacity (372 mA h g<sup>-1</sup>) and safety issues due to their low Li intercalation potential.<sup>1–7</sup>

GeO<sub>2</sub> is considered as a good alternative for graphite as an anode material for LIBs because of its many advantages, including a high theoretical capacity of 1125 mA h g<sup>-1</sup>, low operating voltage and rapid Li<sup>+</sup> diffusion rate.<sup>8–15</sup> In practical use, GeO<sub>2</sub> anodes suffer from fast capacity degradation and poor rate performance caused by their large volume variations during lithiation/delithiation cycles and low intrinsic electronic conductivity.<sup>6,9,11,13,16–19</sup> The hybridization of GeO<sub>2</sub> with conductive buffer materials such as graphene, amorphous

carbon, and carbon nanotubes are effective strategies to address these shortcomings.<sup>5,6,10,20–26</sup> In particular, electrospinning methods that tailor GeO<sub>2</sub> anode materials into one-dimensional (1D) carbon nanofibers have attracted the attention of many researchers, because carbon nanofibers with short Li ion diffusion pathways are recognized as good architectures for energy storage applications.<sup>27–30</sup> However, the electrochemical performances of these GeO<sub>2</sub>/carbon nanofibers are still unsatisfied because: (i) carbon nanofibers typically could not withstand the large volume change of GeO<sub>2</sub> due to their structural fragility,<sup>27</sup> thus lead to the poor cycling performance; (ii) carbon nanofibers usually exhibit relative low electronic conductivity compared to that of graphitized carbon,<sup>30</sup> therefore the rate performance of these electrodes is still not satisfactory.

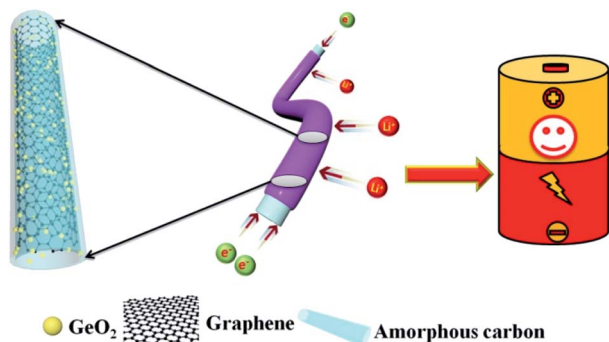
In order to overcome the above mentioned two drawbacks that widely existed in carbon nanofibers, in this work, we tailored graphene in the internal structure of carbon nanofibers to form a nanocable structure *via* a facile electrospinning method. Benefitting from the favorable mechanical properties, and electronic conductivity of graphene, the as-prepared carbon/graphene nanocable successfully mitigates the drawbacks of carbon nanofiber electrodes. As illustrated in Scheme 1, after the electrospinning and the following calcination processes, a ternary nanocomposite–amorphous carbon/graphene-nanocable-encapsulated GeO<sub>2</sub> (denoted as GeO<sub>2</sub>/nanocable) was obtained. In this unique nanocable architecture, graphene acts as the “core” and amorphous carbon as the

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Scheme 1 Schematic illustration of GeO<sub>2</sub>/nanocable.

“shell”, and simultaneously GeO<sub>2</sub> was also encapsulated into the “nanocable”. When tested as an anode material for LIBs, GeO<sub>2</sub>/nanocable exhibits enhanced cycling and rate performances compared to those of GeO<sub>2</sub>/carbon nanofibers (denoted as GeO<sub>2</sub>/CNF, prepared with the absence of graphene) electrodes.

Scanning electron microscopy (SEM) images show as-prepared products possess a 1D fiber-like morphology with a typical length on the order of 10–100 μm and an average diameter of ~300 nm (Fig. 1a and b). The microstructure of the GeO<sub>2</sub>/nanocable was further investigated by transmission electron microscopy (TEM) (Fig. 1c–e) accompanied by selective area electron diffraction (SAED). As shown in Fig. 1c, the graphene “core” was clearly embedded within an amorphous carbon “shell”, judging by the distinct contrasts in the TEM images. The “shell” has a thickness of ~100 nm while the “core” has a diameter of approximately 200 nm. Graphene enhanced

the flexibility of the GeO<sub>2</sub>/nanocable. As depicts in Fig. S1,† after bending, the structure of GeO<sub>2</sub>/nanocable could remain intact while the GeO<sub>2</sub>/CNF collapsed. The formation mechanism of the GeO<sub>2</sub>/nanocable prepared by a single-hole needle should be the conductivity difference between graphene and the electrospinning solution (PAN dissolved in DMF). Driven by a high voltage electrostatic force, graphene nanosheets with good electrical conductivity may join together to form the nanocable’s “core”, and the corresponding PAN solution forms the amorphous carbon “shell”. As shown in Fig. 1d and e, higher-magnification images show that many nanoparticles of diameter < 20 nm were attached to the “core”. The inset of Fig. 1d shows the SAED rings of GeO<sub>2</sub>, where the inner and outer diffraction rings correspond to the diffractions of the (100) and (101) planes, respectively.<sup>31</sup> Therefore, the above nanoparticles may be reasonably attributed to GeO<sub>2</sub> primary nanoparticles. Fig. 1f shows the dark field scanning transmission electron microscopy (STEM) image of GeO<sub>2</sub>/nanocable, where the bright contrast further confirms the nanocable structure of the product. Energy-dispersive spectroscopy (EDS) elemental mapping analysis was employed to investigate the elemental distribution of the GeO<sub>2</sub>/nanocable. As shown in Fig. 1g–i, the C, O, and Ge elemental maps match well with the STEM image (Fig. 1f). In Fig. 1g, as is consistent with the TEM image, the C elemental map is consisted of light red “shell” and dark red “core”. Combined with the above TEM analysis, the light red “shell” is recognized as amorphous carbon, because the texture of the amorphous carbon is the same as that obtained without graphene (as depicted in Fig. S2†). The dark red “core” is supposed as graphene based on the fact that GO is the only possible carbon source except PAN. From Fig. 1h–i, Ge and O are not uniformly distributed over the entire area of the nanocable but are concentrated in the “core” area. Because when the GO solution was mixed with Ge<sup>4+</sup>, Ge<sup>4+</sup> would be selectively bonded with the oxygenated groups by electrostatic forces due to GO nanosheets contained epoxy and hydroxyl groups on the basal planes and carboxylic acid groups.<sup>32</sup> This is another evidence that support graphene is the “core” of the nanocable.

X-ray photoelectron spectroscopy (XPS) curves of the GeO<sub>2</sub>/nanocable shown in Fig. 2a indicate the existence of Ge, C, and O elements. The corresponding high-resolution spectrum shows that there is a sharp XPS peak of Ge 3d at a binding energy at 32.8 eV, confirming the presence of Ge<sup>4+</sup> in the GeO<sub>2</sub>/nanocable (Fig. 2b).<sup>11,33</sup> Moreover, a high resolution O 1s peak is displayed in Fig. 2c at 531.8 eV, suggesting that oxygen exists in the O<sup>2-</sup> oxidation state.<sup>34,35</sup> The high-resolution C 1s spectrum shows one primary and one shoulder peak centered at 284.7 and 286.7 eV corresponding to C–C and C–N, respectively (Fig. 2d).<sup>36</sup>

Fig. 3a shows the X-ray diffraction (XRD) patterns of the GeO<sub>2</sub>/nanocable. The sharp diffraction peaks centered at 20.5°, 26.3°, and 38.2° corresponded to the (100), (101), and (102) planes of the crystalline GeO<sub>2</sub>, respectively, thereby confirming the presence of GeO<sub>2</sub>.<sup>37</sup> No carbon and graphene-related peaks were observed because of their relatively low crystallinity compared with that of GeO<sub>2</sub>.<sup>12</sup> Fig. 3b shows the Raman spectra of commercial GeO<sub>2</sub> and GeO<sub>2</sub>/nanocable. The sharp peak at 443 cm<sup>-1</sup> corresponds to the characteristic peak of GeO<sub>2</sub> (red).

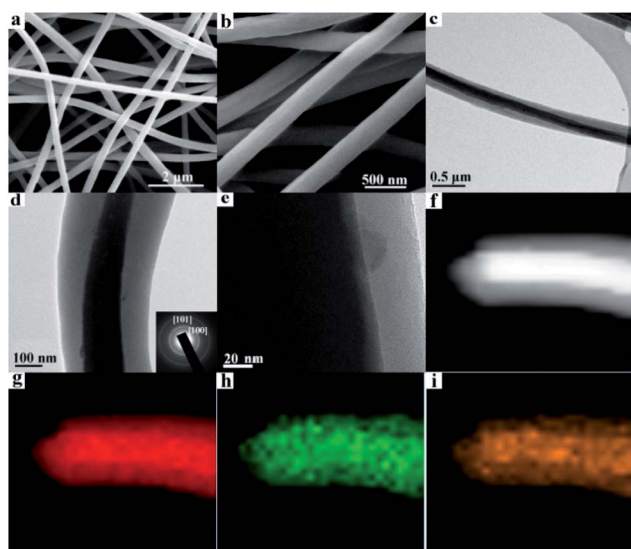


Fig. 1 (a and b) SEM images of GeO<sub>2</sub>/nanocable at low and high magnifications. (c and d) TEM and (e) HRTEM images of GeO<sub>2</sub>/nanocable, inset of (d) is the corresponding SAED patterns; (f) dark field STEM image and (g–i) EDS-elemental mapping images of a single GeO<sub>2</sub>/nanocable (images g, h and i represent C, O and Ge elements, respectively).



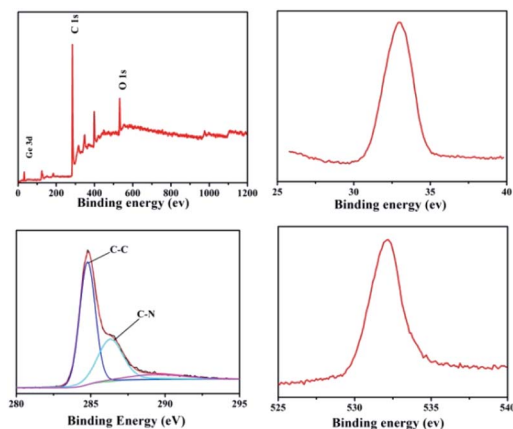


Fig. 2 XPS spectra of GeO<sub>2</sub>/nanocable: (a) the full XPS spectrum of the GeO<sub>2</sub>/nanocable; (b–d) high-resolution spectra Ge 3d, C 1s and O 1s, respectively.

The absence of GeO<sub>2</sub> peak in GeO<sub>2</sub>/nanocable (green) implies most of GeO<sub>2</sub> was beneath the amorphous carbon “shell” and its content in the “shell” was very low, this result is consistent with the above EDS mapping analysis. A 2D band, which is the characteristic band of graphene can be observed at 2600–3000 cm<sup>-1</sup> in the Raman spectra of GeO<sub>2</sub>/nanocable further confirms the existence of graphene.<sup>38</sup> Two sharp peaks at 1332 and 1590 cm<sup>-1</sup> are present in the GeO<sub>2</sub>/nanocable spectrum, which could be assigned to the defect (D) and graphitized (G) bands of carbon, respectively.<sup>39</sup> The intensity ratio of the D band is obviously higher than that of the G band, which indicates that higher amounts of disordered carbon were formed with numerous defects in the amorphous carbon layer (Raman spectra of nanomaterials primarily yield surface information). Amorphous carbon has two effects on the rate performance of LIBs. On the one hand, disordered carbon would enhance the Li

ion diffusion kinetics, thus improving the high-rate performance during charge/discharge cycles of the LIBs.<sup>40,41</sup> On the other hand, excessive amorphous carbon (or thick coating layer) would reduce the electronic conductivity of the electrode, which is harmful to its rate performance.<sup>42</sup> In the GeO<sub>2</sub>/nanocable, the graphene “core” promises the good electrical conductivity while the amorphous carbon “shell” guarantees the fast Li ions diffusion, thus the high power density of the anode material could be anticipated.

The GeO<sub>2</sub> content in the GeO<sub>2</sub>/nanocable was determined by thermal gravimetric analysis (TGA). In the GeO<sub>2</sub>/nanocable, the weight ratio of GeO<sub>2</sub> is 53.56 wt%, and the weight ratio of graphene and amorphous carbon is 46.44 wt% based on the weight loss on carbon combustion and the fact that GeO<sub>2</sub> is stable in air. The weight loss that commences at 500–600 °C could be attributed to the graphene and the amorphous carbon combustion reaction. The specific surface area of the GeO<sub>2</sub>/nanocable, which is calculated using Brunauer–Emmett–Teller (BET) measurements, is 28.3 m<sup>2</sup> g<sup>-1</sup>. The nitrogen adsorption-desorption isotherm exhibits a typical IV-type isotherm with an H3 type hysteresis loop (Fig. 3d).<sup>42</sup> These surface area values indicate that the GeO<sub>2</sub>/nanocable possesses a porous nanostructure, which may be caused by the amorphous carbon layer. According to the above structural characterization, we believe that the rationally designed GeO<sub>2</sub>/nanocable could be presented an ideal anode material for high-performance LIBs.

To systematically study the electrochemical performance of the GeO<sub>2</sub>/nanocable, various electrochemical tests including cyclic voltammetry (CV), electrochemical impedance spectroscopy (EIS) and galvanostatic charge/discharge were performed. GeO<sub>2</sub>/CNF was also tested for comparison. Initially, the Li storage mechanism of the GeO<sub>2</sub>/nanocable was investigated by using CV and the corresponding CV curves are shown in Fig. 4a. The sample was tested at a scan rate of 0.2 mV s<sup>-1</sup> from 0.0 to 3.0 V vs. Li<sup>+</sup>/Li. During the first cathodic scan, the peak at around 0.65 V arose from the decomposition of the electrolyte, the irreversible reaction between electrode and electrolyte to form a stable solid electrolyte interface (SEI) layer, and the irreversible reaction of Li and GeO<sub>2</sub> to form Li<sub>2</sub>O (GeO<sub>2</sub> + 4Li<sup>+</sup> → Ge + 2Li<sub>2</sub>O).<sup>8,14,43</sup> The sharp cathodic peak below 0.30 V corresponded to a series of Li<sub>x</sub>Ge phases.<sup>13,43</sup> During the anodic scan, the peak at around 0.35 V was caused by the dealloying reaction of Li<sub>x</sub>Ge alloys.<sup>20,44,45</sup> The broad peak located at approximately 1.15 V arose from the reoxidation of Ge to GeO<sub>2</sub>, thus result in the partial reversibility of the GeO<sub>2</sub> conversion reaction.<sup>14,15</sup> After the first cycle, the CV curves of the GeO<sub>2</sub>/nanocable overlapped well, suggesting good stability and reversibility of the GeO<sub>2</sub>/nanocable electrode for Li ions insertion and extraction.

The galvanostatic charge–discharge profiles of the GeO<sub>2</sub>/nanocable electrodes were recorded in the voltage window of 0.0–3.0 V versus Li/Li<sup>+</sup> at a current rate of 200 mA h g<sup>-1</sup> over 100 cycles (Fig. 4b). In the first discharge profile, a voltage plateau at approximately 0.4 V and a subsequent long continuous voltage drop down to 0.0 V could be observed, which match well with the CV data and are indicative of Li-alloying reactions. The GeO<sub>2</sub>/nanocable electrode displays an initial discharge/charge capacity of 1470/900 mA h g<sup>-1</sup>; the high initial irreversible

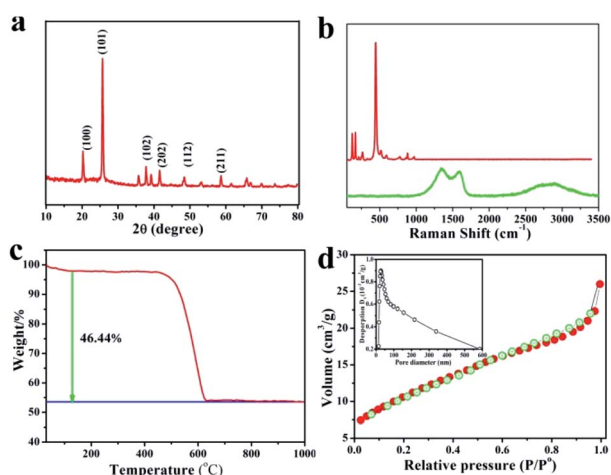


Fig. 3 (a) XRD pattern of GeO<sub>2</sub>/nanocable, (b) Raman spectra of GeO<sub>2</sub>/nanocable (green line) and commercial GeO<sub>2</sub> powder (red line), (c) TG curve of GeO<sub>2</sub>/nanocable in oxygen atmosphere, (d) nitrogen adsorption and desorption isotherms of GeO<sub>2</sub>/nanocable, inset image is the corresponding pore size distribution plots.



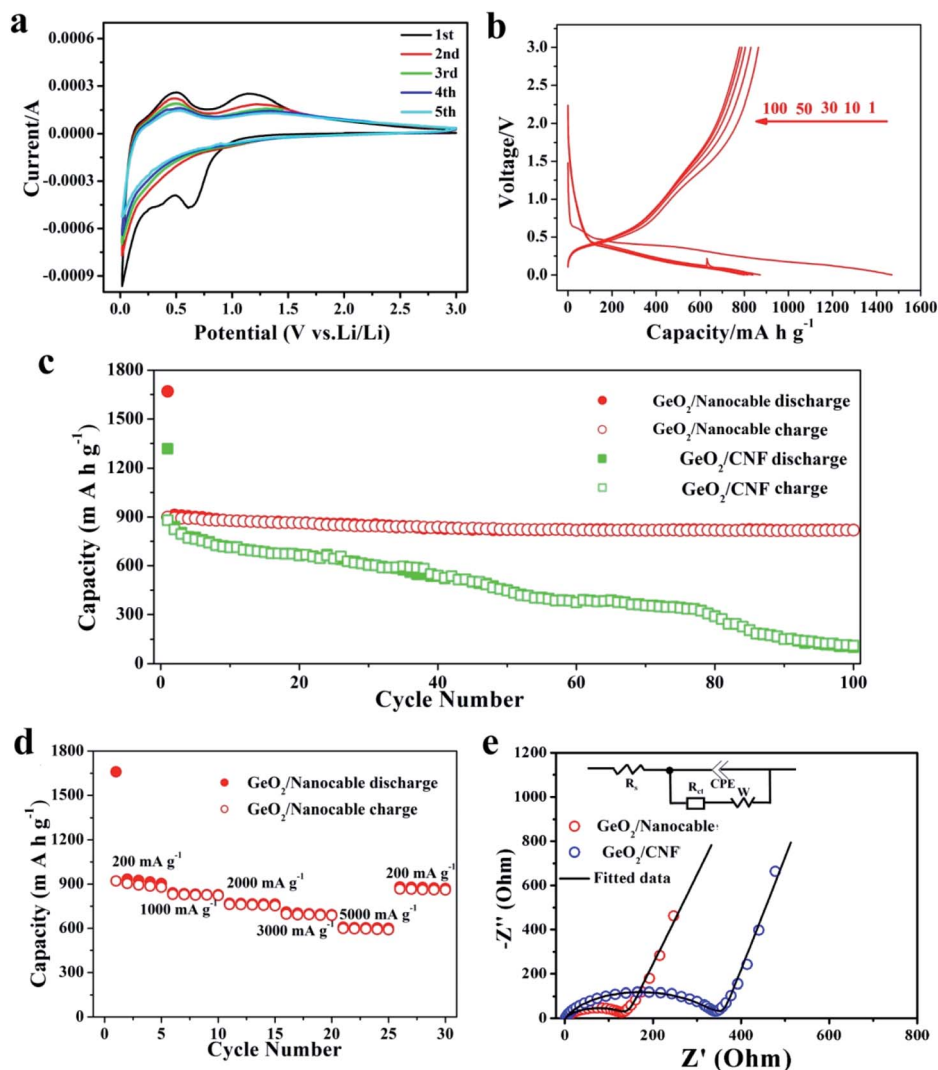


Fig. 4 (a) CV curves of the GeO<sub>2</sub>/nanocable in the voltage window 0.0–3.0 V at a scan rate of 0.1 mV s<sup>-1</sup>. (b) Charge–discharge curves of GeO<sub>2</sub>/nanocable at the 1st, 10th, 30th, 50th and 100th cycles (current: 200 mA g<sup>-1</sup>). (c) Comparison of the cycling performance of GeO<sub>2</sub>/nanocable and GeO<sub>2</sub>/CNF 200 mA h g<sup>-1</sup>. (d) Rate performance the GeO<sub>2</sub>/nanocable at different current densities. (e) EIS spectra of GeO<sub>2</sub>/nanocable and GeO<sub>2</sub>/CNF electrodes.

capacity is related to the formation of the SEI layer, electrolyte decomposition and the irreversible reaction of GeO<sub>2</sub> with Li. After the first cycle, the reversibility of the GeO<sub>2</sub>/nanocable significantly improved and the coulombic efficiency increased up to 97% after the second cycle. Fig. 4c compares the cycling performance of the GeO<sub>2</sub>/nanocable and the GeO<sub>2</sub>/CNF electrodes at a current density of 200 mA h g<sup>-1</sup>. For the GeO<sub>2</sub>/nanocable electrode, the capacity stabilized at above 819 mA h g<sup>-1</sup> after 100 cycles. The capacity loss between the 1st and 100th cycles was only 9%, thus showing the superior cyclability of GeO<sub>2</sub>/nanocable (calculated based on the reversible charge capacities). In contrast, the declining capacity plots of the GeO<sub>2</sub>/CNF electrode indicates its poor cycling performance. In fact, the GeO<sub>2</sub>/CNF electrode showed a capacity retention of only 12.5% with a final reversible capacity of 110 mA h g<sup>-1</sup>. The excellent structural strength and flexibility of graphene led to good cycling stability of the GeO<sub>2</sub>/nanocable

electrode, and this assumption could be further verified by the SEM images that obtained at the end of cycles (see ESI, Fig. S3†). Fig. S3† compares the SEM images of both electrodes after 100 cycles. From these images, it is clear that most of the GeO<sub>2</sub>/nanocables maintain their original 1D structures, while GeO<sub>2</sub>/CNF shows obvious fracture phenomena.

As shown in Fig. 4d, the rate capacities of GeO<sub>2</sub>/nanocable electrodes were also tested. The performed current increased over every 5 cycles in step from 200 mA h g<sup>-1</sup> to 5000 mA h g<sup>-1</sup> and back to 200 mA h g<sup>-1</sup> at the last 5 cycles. At the currents of 200, 1000, 2000, 3000 and 5000 mA h g<sup>-1</sup>, the corresponding reversible charge capacities were approximately 890, 825, 760, 690 and 595 mA h g<sup>-1</sup>, respectively. When the specific current was returned back to 200 mA h g<sup>-1</sup>, the capacity rose to 865 mA h g<sup>-1</sup>, which is very close to the initial charge capacity. These results demonstrate that the GeO<sub>2</sub>/nanocable electrode exhibits good tolerance to variable charge/discharge currents,



which is an important characteristic required for high-power applications. Since the rate capability is dominated by the kinetics of lithium-ion diffusion and electronic conductivity, the better electrochemical performance of the GeO<sub>2</sub>/nanocable electrode was further verified using EIS measurements with a GeO<sub>2</sub>/CNF electrode for comparison. As shown in Fig. 4e, the EIS plots consisted of a semicircle at medium to high frequency and a straight line at low frequency. The inset of Fig. 4e shows the Randles equivalent electrical circuit model of both electrodes, it can be observed that the experimental data could be well fitted using the equivalent circuit model. As is shown, the GeO<sub>2</sub>/nanocable electrode shows a considerably lower charge-transfer resistance (135 Ω) compared to that of the GeO<sub>2</sub>/CNF electrode (331 Ω) (Fig. 4e and ESI Table S1†), indicating a faster charge-transfer reaction for the GeO<sub>2</sub>/nanocable anode.<sup>46</sup> This would lead to a good rate capability of the GeO<sub>2</sub>/nanocable electrode.

## Conclusions

A novel GeO<sub>2</sub>/nanocable electrode material was successfully synthesized by a facile electrospinning method, in which graphene act as the “core” and amorphous carbon as the “shell”, and GeO<sub>2</sub> nanoparticles were encapsulated in the nanocable. As anode material for LIBs, the graphene “core” promises the good electrical conductivity while the amorphous carbon “shell” guarantees the fast Li ions diffusion. The graphene “core” could effectively alleviate volume expansion, and maintain structural stability of the GeO<sub>2</sub>/nanocable electrodes. Therefore, the GeO<sub>2</sub>/nanocable exhibited satisfactory Li storage performance including high reversible capacity (900 mA h g<sup>-1</sup>), excellent cycling performance (91% after 100 cycles) and rate performance (595 mA h g<sup>-1</sup> at 5000 mA g<sup>-1</sup>). Our work demonstrates a successful case for the preparation of GeO<sub>2</sub>/nanocable anode materials for advanced LIBs, and the proposed preparation strategy could be extended to boost other metal or oxide anodes.

## Conflicts of interest

The authors declare no competing financial interest.

## Acknowledgements

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