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Core-leaf onion-like carbon/MnO2 hybrid nano-urchins for rechargeable lithium-ion batteries

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Abstract
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Core-leaf onion-like carbon/MnO$_2$ hybrid nano-urchins for rechargeable lithium-ion batteries

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ABSTRACT

A hybrid nano-urchin structure consisting of spherical onion-like carbon and MnO$_2$ nanosheets is synthesized by a facile and environmentally-friendly hydrothermal method. Lithium-ion batteries incorporating the hybrid nano-urchin anode exhibit reversible lithium storage with superior specific capacity, enhanced rate capability, stable cycling performance, and nearly 100% Coulombic efficiency. These results demonstrate the effectiveness of designing hybrid nano-architectures with uniform and isotropic structure, high loading of electrochemically-active materials, and good conductivity for the dramatic improvement of lithium storage.
1. Introduction

Developing high-performance rechargeable lithium-ion batteries (LIBs) is among the most promising solutions to address the drastic increase in global demand of energy [1]. LIBs were introduced to the market in the 1990s by Sony and soon attracted a strong research interest due to their high energy density, stability, and no memory effect, as compared to other alternatives [2]. However, commercial LIBs mostly use graphite as the anode material, which possesses a relatively low theoretical specific capacity of \( \sim 372 \text{ mAh g}^{-1} \). This low capacity severely hampers the wide usage of LIBs in the surging consumer electronic devices and the large-scale energy applications such as hybrid electric vehicles, renewable power plants, and load levelling [3-5]. To accommodate the high-level requirements of these advanced applications, it is imperative to explore new electrode materials and novel designs for higher energy density, lower cost, flexibility, non-toxicity, and better stability.

The recent advances in nanotechnology have offered a promising route to tackle these challenges [6-9]. As compared to the bulk materials, nanostructured materials possess a large surface area with excellent electrical, optical, and mechanical properties. Nanomaterials can enhance the performance of LIBs through two approaches. The first one is by using low-dimensional carbon-based nanomaterials to provide more efficient lithiation and delithiation processes. For instance, Endo et al. firstly demonstrated the possibility of incorporating submicron vapour grown carbon fibres (VGCFs) into the anode of LIBs [10]; Hu et al. used free-standing carbon nanotube (CNT) film to fabricate flexible LIBs [11]; more recently, graphene and ultrathin graphite foams were utilized to offer higher energy capacity of LIBs [12]. The other approach is by using metal oxide nanoparticles or conductive polymers to store lithium through electrochemical redox reactions. A wide range of nanoparticles such as Si, Ge, Fe\(_3\)O\(_4\), Fe\(_2\)O\(_3\), Co\(_3\)O\(_4\), and Cu\(_2\)O have been successfully implemented to improve the
performance of LIBs. Moreover, hybrid nanostructures which combine the advantages from both carbon nanomaterials and metal oxide nanoparticles were also demonstrated [13-16].

Among various metal oxide nanoparticles, manganese oxides (MOx) which can be synthesized with different structure (e.g., crystalline or amorphous) and multiple-valent state (e.g., MnO, Mn2O3, MnO2, Mn3O4) are increasingly recognised as one of the best anode materials for LIBs [17,18]. In particular, MnO2 nanoparticles are stable, nontoxic and environmentally benign, and have a high specific capacity (~1230 mAh g⁻¹). Much effort is thus devoted to synthesize and control the structure of MnO2-based materials [19-21]. However, the performance of MnO2-based anodes remains inferior due to two main issues: (i) the pristine MnO2 has a low electrical conductivity (10⁻⁵ – 10⁻⁶ S/cm) which limits the electron transport; (ii) a large volume change of MnO2 occurs during the lithiation and delithiation processes, which leads to severe electrode pulverization, serious safety concerns and rapid loss of discharge capacity during the cycling [17,21,22].

Issue (i) can be partially mitigated by coating the MnO2 nanostructures with highly-conductive carbonaceous materials, such as single- and multi-walled CNTs, graphene, poly(3,4-ethylenedioxythiophene) (PEDOT), and mesoporous carbons [23,24]. However, issue (ii) remains largely unresolved as it is associated with the fundamental charge storage mechanism and the inherent structural properties of MnO2 nanoparticles. Indeed, the available MnO2–based LIBs often experience poor cycling performance, low stability, and even cracking, delamination, and crumbling of the anode material [17,25,26].

In this work we solve these problems by fabricating the hybrid nano-urchin structure consisting of onion-like carbon/MnO2 (OLC/MnO2) using a simple and environmentally-benign hydrothermal method. The OLC acts as both the support for MnO2 nanostructures and the conductive core for electron transport. It was chosen because of three prominent features:
the good conductivity which promotes fast electron transport, the large specific surface area which allows a high loading of the electrochemically active MnO$_2$ nanoparticles, and the isotropic structure which enhances the accessibility of nanoparticles to the electrolyte ions and accommodates the change of volume during lithium storage [27-29]. We demonstrate that LIBs incorporating the hybrid nano-urchin anode exhibit better cycling performance and rate capabilities with nearly 100% Coulombic efficiency, which are significantly improved from that of the pure MnO$_2$ or OLC anodes. Moreover, the structure of the OLC/MnO$_2$ hybrid remains distinguishable after more than 100 cycling tests, in a drastic contrast to the pure MnO$_2$ structure which degraded upon a few cycles of electrochemical operation.

2. EXPERIMENTAL

2.1 Preparation of the core-leaf OLC/MnO$_2$ hybrid nano-urchins

The synthesis procedure and the proposed nano-assembly mechanism of core-leaf OLC/MnO$_2$ hybrid nano-urchins are schematically shown in Fig. 1a. Briefly, we firstly prepared MnO$_2$ nanoparticles on the surface of OLC by stirring the mixture of 150 mg potassium permanganate (KMnO$_4$) and 50 mg OLC in 25 ml de-ionized water at 55 °C for 6 hours in a 50 mL Teflon-line stainless steel autoclave. The intermediate OLC/MnO$_2$ product was then treated in a simple hydrothermal process at 150 °C for 12 hours to transform into the nano-urchin structure. Pure MnO$_2$ was also synthesized in a hydrothermal process at 150 °C for 3 hours by mixing KMnO$_4$ with H$_2$SO$_4$ in DI water.

2.2 Morphological and structural characterizations
Morphology and structure of the samples were examined by the field-emission scanning electron microscopy (FE-SEM, Zeiss LEO 1550) and transmission electron microscopy (TEM, JEOL JEM-2100F). Compositional investigation of the samples was performed by energy-dispersive X-ray spectroscopy (EDX, JEOL JSM-5910LV). The crystal structure of samples was investigated by X-ray diffraction (XRD, Siemens D5005) with Cu Ka (λ=0.154 nm) radiation and the accelerating voltage of 40 kV. Thermal weight changes was measured by thermogravimetric analysis (TGA, Shimadzu DTG-60). Additionally, surface area analyzer (Micromeritics ASAP 242,) was used to measure the specific surface area with N₂ physisorption at 77 K.

2.3 Electrochemical measurement

The electrochemical properties were investigated by a two-electrode half-cell configuration composed of the lithium foil as the counter electrode and the active material as the working electrode. Typically, electrode slurry was prepared by mixing 80 wt% active material, 10 wt% conductive carbon black and 10 wt% polyvinylidifluoride (PVDF) binder in the N-Methylpyrrolidone (NMP) solution. The slurry was then painted onto a nickel-foam current collector, followed by drying in a vacuum oven. The mass of the active material is in the range of 3 – 4 mg cm⁻². 1M LiPF₆ solution in a 1 : 1 (v : v) mixture of ethylene-carbonate/ethyl-methyl-carbonate (EC/EMC) and Celgard 2400 membranes were employed as the electrolyte and the separator, respectively. All components listed above were assembled into a standard CR2032 button battery in a glove box filled with argon gas. The cyclic voltammetry (CV) and electrochemical impedance spectra (EIS) measurements of the half-cells were carried out on a potentiostat/galvanostat (VMP3, Bio-Logic). The scan rate of the CVs was at 0.05 mV s⁻¹ and the frequency range for the EIS spectra was varied from 0.01 Hz to 1 MHz. The galvanostatic discharging/charging test was performed in the potential
range of 0.01 – 3.0 V at various current densities from 50 to 2000 mA g\(^{-1}\) using the Neware battery tester.

3. RESULTS AND DISCUSSION

3.1 Morphology and structure of the OLC/MnO\(_2\) nano-urchins

The scanning electron microscopy (SEM) images of the pure OLC, the intermediate product, and the final OLC/MnO\(_2\) nano-urchins are shown in Fig. 1b, 1c, and 1d, respectively. One can see that the size of nanoparticles increased significantly during the hydrothermal process. Bundles of MnO\(_2\) nanosheets were observed to radiate from the OLC core (inset of Fig. 1d and Fig. S1), giving rise to the highly-porous core-leaf structure with a large surface area. Moreover, the diffraction peaks in the XRD measurements of OLC/MnO\(_2\) nano-urchins could be indexed as (001), (002), (110), (111) and (020) planes of the birnessite-type MnO\(_2\) (Fig. S2) [29]. This birnessite-type of MnO\(_2\) is characterized by a layered structure consisting of edge-shared MnO\(_6\) octahedra, which allows high mobility of the interlayer cations with fast kinetics and little structural re-arrangement [30]. Some MnO\(_2\) nanofibers were also found in the final product (Fig. 1c), which can be attributed to the transformation from birnessite-type MnO\(_2\) nanosheets into nanofibers at extended reaction time and/or high temperature [31,32]. The hybrid OLC/MnO\(_2\) was therefore architected into an intricate, three-dimensional (3D) urchin-like structure with an isotropic surface, which can effectively reduce the aggregation of Mn metal formed during the electrochemical reaction and enhance the cyclic stability, as demonstrated in detail below.

Figure 2 shows the crystalline structure and compositional analysis of the pure OLC and OLC/MnO\(_2\) nano-urchins obtained by TEM and EDX. Pure OLC had a highly-crystalline graphitic wall structure with an interplanar spacing of 0.34 nm [the (002) basal plane] and a
diameter of about 15 – 40 nm (Fig. 2a and 2b). When coated with MnO₂, the size of the core-leaf structure increased to 200 – 300 nm with the length of MnO₂ nanosheets of ~100 nm (Fig. 2c and 2d). We noted that the density of MnO₂ was much higher in the present case than that grown on hollow carbon spheres (diameter of ~200 nm) [33], making the current nanostructure promising for higher energy storage capacity. The specific surface area of pure OLC was 377 m² g⁻¹ as calculated by the Brunauer–Emmett–Teller (BET) method (Fig. S3). This large surface area of OLC might thus contribute to the high loading of MnO₂ nanoparticles.

TEM images further indicated that the interplanar spacing of birnessite-type MnO₂ nanosheets was ~0.7 nm (Fig. 2d–2e and Fig. S1), in a good agreement with others [20,34]. The multiple diffraction rings observed in the selected-area electron diffraction (SAED) pattern suggested that MnO₂ was nanocrystalline (Fig. 2f). According to Xia et al. [35], when the KMnO₄ solution is mixed with carbon nanomaterials before the hydrothermal processing, the nanocrystalline MnO₂ forms on the surface of carbon nanomaterials due to the slow redox process according to the following Eq. (1).

\[ 4\text{MnO}_4^- + 3\text{C} + \text{H}_2\text{O} \rightarrow 4\text{MnO}_2 + \text{CO}_3^{2-} + 2\text{HCO}_3^- \]  

When the solution is further treated in the hydrothermal reaction, the MnO₂ nanosheets grow from the preformed nanocrystalline due to the decomposition of KMnO₄ in water according to Eq. (2).

\[ 4\text{MnO}_4^- + 2\text{H}_2\text{O} \rightarrow 4\text{MnO}_2 + 4\text{OH}^- + 3\text{O}_2 \]  

In addition, we have previously shown that water molecules and potassium ions may diffuse into the interlayer of birnessite-type MnO₂ during the synthesis process [29]. This presence of potassium ions is beneficial for stabilizing the layered structure of MnO₂ [30]. It is estimated here from the EDX spectrum that the K/Mn ratio was about 0.23 (Fig. 2g and Table S1). Moreover, the TGA results of OLC/MnO₂ nano-urchins indicated that the water
content and OLC was about 9% and 17%, respectively (Fig. S4). The interlayer water is then calculated about 0.64 H$_2$O per chemical formula, i.e., K$_{0.23}$MnO$_2$·0.64H$_2$O [35]. The final product of the nano-urchins was thus denoted as 0.83(K$_{0.23}$MnO$_2$·0.64H$_2$O)/0.17OLC.

3.2 Lithium storage performance

Next, we fabricated a standard CR2032 button half-cell battery with the 3D OLC/MnO$_2$ nano-urchins as the working electrode and the lithium foil as the counter electrode in a glove box filled with Ar (a photo is shown as the inset of Fig. 3b). It is known that the mechanism of lithium reacting with nano-sized MnO$_2$ (also other 3d-metal oxides such as CoO$_x$ and MoO$_x$) differs from the classical lithium intercalation or alloying processes; instead, it is a reversible redox reaction process involving the formation of Li$_2$O,

\[ \text{MnO}_2 + 4\text{Li}^+ + 4e^- \leftrightarrow 2\text{Li}_2\text{O} + \text{Mn} \]  (3)

followed by the decomposition of Li$_2$O and oxidation of metallic nanoparticles [19,35].

Figure 3a shows the CV curves of OLC/MnO$_2$ core-leaf nano-urchins in the first three cycles, obtained at a scan rate of 0.05 mV/s and a potential window of 0.01 – 3 V vs Li/Li$^+$. One can see that in the first cathodic cycle there was one weak peak at 0.75 V and one main peak near 0.24 V. The former corresponded to the possible reduction of Mn$^{4+}$ to Mn$^{2+}$, while the latter was associated with the complete reduction of Mn$^{2+}$ to Mn and the formation of a solid electrolyte interface (SEI) layer at the interface of electrolyte and electrode (a polymer/gel-like film which allowed only lithium ions to diffuse) [36,37]. From the second cathodic cycle, the weak peak disappeared and the main reduction peak shifted to 0.3 V, indicating the reversible phase transformation as given in Eq. (3).

In the anodic process of Fig. 3a, one main peak was observed at around 1.3 V and one weak peak at 2.1 V, corresponding to the oxidation of metallic Mn to Mn$^{2+}$ and the
decomposition of polymer/gel film related to the SEI layer [37]. These peaks were similar to those of pure MnO₂ but different from those of pure OLC (Fig. S5a and S5b), confirming that the MnO₂ nanosheets were the main medium for lithium storage.

The galvanostatic discharging/charging measurements of OLC/MnO₂ nano-urchins at ambient temperature is shown in Fig. 3b. In the first discharging curve, there were two plateaus located at 0.75 and 0.48 V, typically ascribed to the complete reduction of Mn⁴⁺ to Mn⁰ as given in Eq. (3) [21,38]. The plateau at 0.75 V disappeared and the one at 0.48 V was shortened in the following discharging cycles, confirming the formation of metallic Mn and Li₂O matrix [21,37]. On the other hand, in the first charging curve the electrode showed a plateau in the potential range between 1.0 and 1.5 V, which was ascribed to the oxidation of Mn⁰ to Mn²⁺ [37]. These discharging/charging behaviors were consistent with the CV curves shown in Fig. 3a and were superior as compared to that of the pure MnO₂ and pure OLC (Fig. S5c–S5d) [26].

We also calculated the specific capacity to be 1278 and 854 mAh g⁻¹ in the first discharging and charging cycles, respectively. It was noted that MnO₂ has a theoretical capacity of 1230 mAh g⁻¹ [35]. The slightly larger value in our calculation could be due to the decomposition of electrolyte at a low voltage during the formation of SEI layer, which contributed extra lithium storage at the Mn/Li₂O interface [22]. Similar phenomena have also been reported for other transition metal oxide electrodes [39,40].

A more prominent feature of the OLC/MnO₂ nano-urchins is that the rate capability of LIBs incorporating this material was significantly improved as compared to that of pure MnO₂, as shown in Fig. 3c. The OLC/MnO₂ nano-urchins exhibited a capacity of 1278 mA h g⁻¹ in the first discharging cycle, which then gradually reduced to 628 mA h g⁻¹ after 10 cycles at a rate of 50 mA g⁻¹. This irreversible capacity, as mentioned above, was due to the
decomposition reactions of electrolyte and the formation of SEI layer [21]. With the increase of discharging/charging rate, however, the irreversibility disappeared and the battery could deliver a reversible capacity of 404, 263, 178 and 102 mAh g$^{-1}$ at rates of 200, 500, 1000, and 2000 mA g$^{-1}$, respectively. Remarkably, the capacity was observed to restore to ~600 mAh g$^{-1}$ and remained stable when the current density returned to 50 mA g$^{-1}$ after 50 cycles (Fig. 3c). This rate capacity of hybrid OLC/MnO$_2$ nano-urchins was superior to that of the pure MnO$_2$ taken separately (e.g., <25 mAh g$^{-1}$ for pure MnO$_2$ at a rate of 1000 mA g$^{-1}$). It was also better than the layer-by-layer graphene-MnO$_2$ nanotube film (~520 mAh g$^{-1}$ after continuous 32 cycles at various current densities) [21], and was comparable to the MnO$_2$/polyaniline anode (~210 mAh g$^{-1}$ at a current density of 1500 mA g$^{-1}$ after 50 cycles) [41].

Furthermore, Fig. 3d shows the electrochemical impedance spectroscopy (EIS) measurements of all three anode materials. The EIS curves presented in Nyquist plots could be fitted to the equivalent circuit (inset of Fig. 3d), where $R_s$ is the electrolyte resistance, $R_f$ and $Q_1$ are the SEI layer resistance and the constant phase element (CPE), respectively; $R_{ct}$ and $Q_2$ are the charge transfer resistance and the related double-layer capacitor, respectively; and $Q_3$ is a CPE corresponding to the line in the low frequency. The fitting results are summarized in Table S2, which indicated that the charge-transfer resistance of the OLC/MnO$_2$ nano-urchins (~140 $\Omega$) had been greatly reduced from that of pure MnO$_2$ (~250 $\Omega$) [42].

### 3.3 Cycling stability

We also compared the cycling performance of pure OLC, pure MnO$_2$, and hybrid OLC/MnO$_2$ nano-urchins, as shown in Fig. 4a. While the first two materials displayed a relatively stable cycling performance, an interesting feature was observed for the OLC/MnO$_2$ nano-urchins. Specifically, the capacity decayed during the first 20 cycles, flattened for the next few cycles,
and then increased to 541 mAh g\(^{-1}\) at the 100\(^{th}\) cycle. A similar trend was also observed for other metal oxide electrodes but with a much smaller increment than the current observation [39,43]. Although the exact mechanism of such increment still remains unclear, we attributed it to the enhanced conversion reaction of MnO\(_2\) nanoflowers experienced on the electrode. After certain cycles, it is known that the available active sites for lithium ion reaction could increase due to the volume change of the OLC/MnO\(_2\) nano-urchins [17]. This reasonable rise in accessible active sites could thus result in a larger capacity. As a high loading of the MnO\(_2\) nanosheets were anchored uniformly on the surface of OLC, the observed high value of the increment became quite reasonable.

Finally, we plotted the Coulombic efficiency of OLC/MnO\(_2\) nano-urchins at the rate of 200 mA g\(^{-1}\). As shown in Fig. 4b, the Coulombic efficiency increased notably to near 100% after several cycles and remained stable throughout the whole experiments. The high Coulombic efficiency not only indicated a reversible electrolyte decomposition and a complete de-conversion reactions, but also implied the high stability of LIBs [26]. This near-100% Coulombic efficiency also excluded the formation of additional SEI layers during the cycling [17]. In conjunction with the improved stability, our microscopic analysis confirmed that the OLC/MnO\(_2\) nano-urchins preserved discernible shape after 100 cycles (inset of Fig. 4b); whereas pure MnO\(_2\) had completely lost its features and became pulverized (Fig. S6).

### 3.4 Unique features of OLC

It is known that the lithium storage is critically determined by the transport of lithium ions and electrons at the electrode/electrolyte interface, the kinetics of surface incorporation of lithium ions into the electrochemically active materials, and the specific charge storage mechanism [22]. An ideal electrode architecture should possess a good conductivity to effectively transport electrons to the electrochemically active nanomaterials and a porous (but
not tortuous) structure to enhance the lithium ion diffusion [21,25,44]. Zhang et al. used an open 3D interpenetrating nickel network to anchor the MnO$_2$ nanoparticles and obtained ultrafast charge transport and intercalation/de-intercalation of active species [45]. Similarly, the porous and open structure of OLC in the present case could possibly contribute: (a) a large surface area for incorporating a high loading of the electrochemically active MnO$_2$ nanosheets (Fig. 1d); (b) a reduced charge transfer resistance for rapid ion and electron transport without sacrificing the energy density (Fig. 2d); (c) a uniform and isotropic structure that enabled the delivery of lithium ions to the active material and better accommodated the volume change during the charge/discharge cycles (Fig. 3c); and (d) an electrochemically stable core structure (Fig. 4b). The preparation method is also simple and scale as compared to other methods such as sputtering, high-temperature carbonization, or layer-by-layer assembly, as the hybrid OLC/MnO$_2$ preserved its nanostructural feature after device fabrication. We therefore believe that the 3D isotropic nano-urchins could be highly promising for high-performance rechargeable LIBs.

4. CONCLUSION

In conclusion, we have assembled the OLC/MnO$_2$ nano-urchins which acted as the stable anode material in LIBs. The electrode showed key electrochemical properties including the improved specific capacity (i.e., from ~260 mAh g$^{-1}$ of the pure MnO$_2$ to ~630 mAh g$^{-1}$ at the current density of 50 mA g$^{-1}$), the increased rate capability at various current densities, the stable cycling performance without significant degradation of the nanostructures, and nearly 100% Coulombic efficiency. These improved performance for rechargeable LIBs arose from the unique nanostructure which facilitated fast ion and electron transport and stabilized the MnO$_2$ nanoparticles during charge storage. Our proven strategy of using OLC as the core
material is cost-effective and generic, which may be applied to other high-capacity transition metal oxides also lead to the *greener and more sustainable development* of energy storage devices for advanced applications.

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Figure captions:

**Fig. 1** - (a) Schematic diagram of the synthesis procedure of the core-leaf OLC/MnO$_2$ hybrid nano-urchins. Scanning electron micrographs of (b) pure OLC, (c) intermediate product, and (d) final hybrid nano-urchins. Inset in (d) is the magnified image of a single urchin-like nano-architecture.

**Fig. 2** - (a) Low- and (b) high-resolution TEM images of pure OLC, and (c) – (e) OLC/MnO$_2$ nano-urchins. Inset in (c) illustrates the intricate, 3D core-leaf architecture. (f) Electron diffraction pattern and (g) EDX spectrum of the OLC/MnO$_2$ structures.

**Fig. 3** - (a) Cyclic voltammetry curves of the OLC/MnO$_2$ nano-urchin electrode in the first three cycles at a scan rate of 0.05 mV s$^{-1}$ in a potential range of 0.01 – 3 V vs Li/Li$^+$. (b) Galvanostatic discharging/charging curves of the OLC/MnO$_2$ nano-urchin electrode at a current density of 50 mA g$^{-1}$ for the first 10 cycles. Inset in (b) is the photo of CR2032 button battery assembled in this work. (c) Rate capabilities of the pure MnO$_2$, pure OLC, and OLC/MnO$_2$ nano-urchin electrodes. (d) EIS spectra and fittings of the pure MnO$_2$, pure OLC and OLC/MnO$_2$ nano-urchin electrodes. Inset in (d) shows the equivalent circuit model used for the fittings.

**Fig. 4** - (a) Cyclic performance of the pure MnO$_2$, pure OLC and OLC/MnO$_2$ nano-urchin electrodes at 200 mA g$^{-1}$. (b) Coulombic efficiency of the OLC/MnO$_2$ nano-urchin electrode at 200 mA g$^{-1}$. Inset in (b) is the SEM image of OLC/MnO$_2$ nano-urchin electrode after 100 cycles.
Figures:

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