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Self-assembling few-layer MoS$_2$ nanosheets on CNT backbone for high-rate and long-life lithium-ion batteries

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We demonstrate the self-assembly of few-layer MoS$_2$ nanosheets on CNT backbone via a facile hydrothermal reaction with the subsequent annealing process. In this structure, the few-layer MoS$_2$ nanosheets with controllable content are alternately and vertically grown on the surface of CNTs, forming a three-dimensional hierarchical nanostructure. The optimized MoS$_2$/CNTs hybrids could be applied as a fascinating anode material for high-rate and long cycle life lithium ion batteries (LIBs). Compared with the commercial MoS$_2$ (716 mAh g$^{-1}$), the as-prepared MoS$_2$/CNTs hybrids exhibit a much higher specific capacity of 1293 mAh g$^{-1}$ at 200 mA g$^{-1}$ with remarkably enhanced rate capability (888 mAh g$^{-1}$ even at 3200 mA g$^{-1}$). More significantly, we find that MoS$_2$/CNTs hybrids show no capacity fading after 200 cycles at 400 mA g$^{-1}$. As for MoS$_2$-based anode materials, such overwhelming electrochemical performance endows the present MoS$_2$/CNTs hybrids huge potential for developing LIBs.

Introduction

Lithium-ion batteries (LIBs) have now become the predominant power source for a wide range of portable electronic devices. In recent years, the development of electric vehicles and hybrid electric vehicles has triggered an ever-increasing demand for LIBs with higher power density and long cycle life.\textsuperscript{1,2} Their performances are strongly dependent on the choice of anode and cathode materials. As for anode materials, graphite is widely used as commercial anode materials in view of its natural abundance and good structural stability.\textsuperscript{3-6} However, it suffers from a relatively low theoretical capacity of 372 mAh g$^{-1}$. Therefore, it is crucial to search alternative anode materials with higher capacity and long cycle life for the development of LIBs.

As a typical layered transition metal sulfide, MoS$_2$ has received intense interest as a promising electrode material for LIBs because of its graphite-like structure.\textsuperscript{7-9} The layered structure and the weak van der Waals forces between MoS$_2$ layers facilitate reversible Li$^+$ intercalation and extraction.\textsuperscript{10} However, like the graphene, the freshly synthesized MoS$_2$ layers have a tendency to aggregate during practical applications, even in the drying process, greatly reducing the electrochemical active sites. Another weakness of MoS$_2$ is its poor electrical conductivity. Both the two disadvantages make its rate capability and cycling stability unsatisfactory. To solve these problems, an effective approach is to hybridize MoS$_2$ with advanced carbon materials.\textsuperscript{11-15} For example, layered MoS$_2$/graphene composites\textsuperscript{14} and MoS$_2$/amorphous carbon composites\textsuperscript{15} have been synthesized as LIBs anode materials, exhibiting an improved specific capacity with good rate and cycling performances. Notably, for the hybrid of MoS$_2$ and CNTs, the MoS$_2$ layers prefer to confine to the CNTs surface, leading to the formation of tubular MoS$_2$ layers with high crystallinity.\textsuperscript{16-18} In this regard, a high loading mass will lower the utilization of MoS$_2$ active material while a low loading mass will result in low capacity based on the MoS$_2$/CNTs hybrids. If the MoS$_2$ nanosheets can be uniformly dispersed on CNTs, which will induce the coupling effect between them, which will result in remarkable enhancement of electrochemical performance.

In the present work, we demonstrate a simple route for realizing the self-assembly of few-layer MoS$_2$ nanosheets on CNT backbone, in which the few-layer MoS$_2$ nanosheets are alternately grown on the surface of CNT, forming a three-dimensional hierarchical nanostructure. The content of MoS$_2$ can be easily controlled simply by tuning the molybdate content. When evaluated as anode materials for LIBs, the optimized MoS$_2$/CNTs hybrids indicate remarkably enhanced reversible capacity (1293 mAh g$^{-1}$ at current density of 200 mA g$^{-1}$) with excellent rate and cycling performances.

Experimental

Synthesis of the MoS$_2$/CNT Hybrids.

20 mg of CNTs was dispersed in a mixed solution with 15 ml water, 15 ml ethanol and 2 ml oleic acid containing 1.6 g sodium oleate, 0.6 g Na$_2$MoO$_4$ and 0.8 g L-cysteine by ultra-sonication for 60 min. After that, the solution was put into a 50 mL Teflon-lined stainless steel autoclave and maintained at 180 °C for 24 h. The precipitates were filtered, washing with water and ethanol several times, and dried in a vacuum at 80 °C. Afterward, the dried samples were loaded into the tube furnace and calcined in Ar atmosphere at 550 °C for 120 min with a ramp of 2 °C min$^{-1}$.

Characterization.
Structure and morphology of the as-prepared samples were characterized by X-ray diffraction (RIGAK, D/MAX 2550 VB/PC, Japan), field emission scanning electron microscopy (Hitachi FE-S4800), transmission electron microscopy (TEM; JEOL, JEM-2100F). Thermogravimetric analysis (NETZSCHSTA409PC) was carried out with a heating rate of 10 °C min⁻¹ under flowing air. Fourier transform infrared (FTIR) spectra were measured by using a Nicolet 5700 spectrophotometer, in the range of 400 to 4000 cm⁻¹ with a resolution of 4 cm⁻¹. N₂ adsorption/desorption was determined by Brunauer-Emmet-Teller (BET) measurements using an ASAP-2020 surface area analyzer.

Electrochemical Measurements.

LIB performance was determined using CR2016 type coin cells assembled in an argon-filled glove box. The working electrode was prepared by mixing the active material, carbon black (Super-P-Li), and a polymer binder (poly(vinylidenedifluoride), PVDF, Aldrich) at a weight ratio of 7:2:1. A polypropylene film (Celgard-2400) was used as a separator. Li foil was used as the counter electrode. The electrolyte was a 1 M LiPF₆ solution in a 50:50 (w/w) mixture of ethylene carbonate (EC) and diethyl carbonate (DMC). The galvanostatic charge and discharge experiment was performed with a battery tester LAND-CT2001A in the voltage range of 0.01 V – 3.0 V at room temperature. The impedance spectra were recorded by applying a sine wave with amplitude of 5.0 mV over the frequency range from 100 kHz to 0.01 Hz.

Results and discussion

The few-layer MoS₂ nanosheets assembled on CNT backbone, forming a three-dimensional hierarchical nanostructure, which has been realized by a simple hydrothermal reaction of sodium molybdate and L-Cysteine in the present of CNTs with subsequent annealing in Ar at 550 °C. The morphology of the products was characterized by both FESEM (Figure S1, ESI) and TEM (Figure 1a-c). As shown in low-magnification TEM image (Figure 1a), the uniform morphology of the MoS₂ nanosheets grown on CNTs have diameters of ~ 100 nm, while the diameter of CNTs is about ~ 20 nm. High-magnification TEM image (Figure 1b) reveals the MoS₂ nanosheets are interconnected and vertically distributed on the surface of CNTs, forming a very intriguing three-dimensional hierarchical nanostructure. By tuning the molybdate content, the content of MoS₂ can be easily controlled without changing their morphology. More interestingly, the two layered spacing can be measured to be about 1.0 nm, which is much larger than the value of the reported MoS₂ (0.64 nm). The data is in good agreement with XRD results, as shown in Figure 2 (black line). It can be found that the (002) reflection disappears while a clear broad peak at ~ 8.4° (marked by 1#) and a poor broad peak at ~ 16.7° (marked by 2#) appear. The interlayer distance of peak 1# can be calculated to ~ 1.0 nm according to the Bragg equation, which is the same as the TEM observation. Such large interlayer distance may be attributed to the presence of oleic acids on the surface of single-layer MoS₂. The insertion of oleic acids into the layer of MoS₂ has been confirmed by FTIR analysis as shown Figure S2. Prior to heat treatment, the fresh MoS₂/CNTs clearly display a keen peak at 2902 cm⁻¹, which is assigned to C-H stretching vibration of CH₂ and CH₃ in oleic acids. The peak at 1706 cm⁻¹ belongs to the C=O stretching vibration of COOH. The peak at 1399 cm⁻¹ belongs to the vibration of -CH=CH-. The peak at 1040 cm⁻¹ attributed to the vibration of C-O. The peak at 766 cm⁻¹ to the δ_C-H (bending vibration), which is characteristic of -CH₂ in long-chain alkanes. All the peaks disappeared after the calcination as a result of the removal of oleic acids as shown in Figure S2 (red line). In view of the strong confined effects of MoS₂ interlayer from their self-assembly process, it is hard to remove the surfactant oleic acids only by washing. In addition, the crystallinity of MoS₂ is also very poor from the SAED pattern in inset of Figure 1b. To totally remove the residual and improve the crystallinity, an annealing process was performed. Here, we chose 550 °C as the annealing temperature considering that too high temperature will result in the formation of tubular MoS₂ on the surface of CNTs (TEM image, Figure S3 in ESI). The TEM image is shown in Figure 1d. It can be observed that the morphology has been well-maintained. Figure 1e shows the HRTEM of the interface between the CNT and MoS₂ layer. It can be observed that the few-layers MoS₂ nanosheets are interconnected and directly grown on the CNT.
The interlayer distance is about 0.64 nm. The corresponding SAED pattern (inset in Figure 1e) shows the obvious diffraction rings, indicating a high crystallinity of the products. These results are further confirmed by XRD measurement, as shown in Figure 2 (red line). The XRD pattern of the annealed MoS$_2$/CNTs hybrids displays the distinct (002), (100), (103) and (110) diffraction peaks of 2H-MoS$_2$ (JCPDS 37-1492). Furthermore, the annealing MoS$_2$/CNTs hybrids also possess a high BET surface area of 45.0 m$^2$ g$^{-1}$ with a bimodal mesopore size distribution (Figure S4 in ESI), which is important for achieving high energy density and power density for LIBs.

Fig. 3 TG curves of MoS$_2$/CNTs hybrids with different MoS$_2$ content, labeled as sample a-c.

To optimize the composition, the annealed MoS$_2$/CNTs hybrids with different MoS$_2$ content have been synthesized, which are determined by TG analysis. As shown in Figure 3, the weight loss was measured to be 76%, 68% and 62% for sample a-c in order. Each TG curve obviously shows two weight losses. The first weight loss occurs at about 380 °C, caused by the oxidation of MoO$_3$ after TG measurement, the MoS$_2$ content of sample a-c could be estimated to be 85%, 75% and 69%, respectively. Their electrochemical performances were preliminarily evaluated by assembling them into coin-type 2016 cells, respectively. The relationship of the annealed MoS$_2$/CNTs hybrids with different MoS$_2$ content and their electrochemical performances has been investigated in Figure S5 in ESI. It can be seen that, with the increase of MoS$_2$ content, a better electrochemical performance can be obtained. In this work, a maximum MoS$_2$ content can reach as high as ~ 85%, which then has been further evaluated in detail.

Their structure and morphology of the hybrids with 85% MoS$_2$ content have already been characterized in detail before. For convenient discussion, the corresponding hybrids are labelled as MoS$_2$/CNTs hybrids in the subsequent text. Figure 4a shows the cyclic voltammograms (CVs) of the MoS$_2$/CNTs hybrids within a potential range of 0.01 V - 3 V. As shown in Figure 4a, two peaks at ~ 0.9 V and ~ 0.45 V are observed in the 1st cathodic sweep. The peak at ~ 0.9 V is attributed to the intercalation of Li ion into MoS$_2$ lattice to form Li$_x$MoS$_2$. The other peak at ~ 0.45 V corresponds to the decomposition reaction of Li$_x$MoS$_2$ to Mo and Li$_2$S. In addition, another poor peak at 1.6 V can be observed, which could be attributed to the reduction reaction of the oxygen-containing functional groups from CNTs. The strong peak at 2.4 V can be assigned to the delithiation of Li$_x$MoS$_2$. In the subsequent 2nd and 3rd cathodic sweeps, two peaks are observed at 1.8 V and 1.2 V, respectively, mainly due to the following two reactions: 2Li$^+$ + S $\rightarrow$ Li$_2$S and MoS$_2$ + xLi$^+$ + xe$^-$ $\rightarrow$ Li$_x$MoS$_2$. Fig 4b exhibits the initial three discharge–charge profiles in the potential range of 0.01 V - 3 V at current density of 200 mA g$^{-1}$. The initial discharge and charge capacities can reach 1617 mAh g$^{-1}$ and 1226 mAh g$^{-1}$, respectively, showing a remarkably enhanced Columbic efficiency of 75.8% thanks to the unique nanostructure assembled by few-layered MoS$_2$ nanosheets on CNTs. In the next two discharge and charge processes, the discharge capacity can still reach 1310 mAh g$^{-1}$ and 1296 mAh g$^{-1}$ with Coulombic efficiency as high as 95% and 96%, respectively, demonstrating a high reversible capacity and excellent cycling stability.

Fig. 4 (a) CV curves at a scan rate of 0.2 mV s$^{-1}$ for the initial 3 cycles, (b) charge-discharge curves at 200 mA g$^{-1}$ for the initial 3 cycles of the MoS$_2$/CNTs hybrids, (c) rate capabilities of the MoS$_2$/CNTs hybrids and the commercial MoS$_2$, respectively, (d) cycling behavior and Columbic efficiency of the MoS$_2$/CNTs hybrids at a current density of 400 mA g$^{-1}$. Their structure and morphology of the hybrids with 85% MoS$_2$ content have already been characterized in detail before. For convenient discussion, the corresponding hybrids are labelled as MoS$_2$/CNTs hybrids in the subsequent text. Figure 4a shows the cyclic voltammograms (CVs) of the MoS$_2$/CNTs hybrids within a potential range of 0.01 V - 3 V. As shown in Figure 4a, two peaks at ~ 0.9 V and ~ 0.45 V are observed in the 1st cathodic sweep. The peak at ~ 0.9 V is attributed to the intercalation of Li ion into MoS$_2$ lattice to form Li$_x$MoS$_2$. The other peak at ~ 0.45 V corresponds to the decomposition reaction of Li$_x$MoS$_2$ to Mo and Li$_2$S. In addition, another poor peak at 1.6 V can be observed, which could be attributed to the reduction reaction of the oxygen-containing functional groups from CNTs. The strong peak at 2.4 V can be assigned to the delithiation of Li$_x$MoS$_2$. In the subsequent 2nd and 3rd cathodic sweeps, two peaks are observed at 1.8 V and 1.2 V, respectively, mainly due to the following two reactions: 2Li$^+$ + S $\rightarrow$ Li$_2$S and MoS$_2$ + xLi$^+$ + xe$^-$ $\rightarrow$ Li$_x$MoS$_2$. Fig 4b exhibits the initial three discharge–charge profiles in the potential range of 0.01 V - 3 V at current density of 200 mA g$^{-1}$. The initial discharge and charge capacities can reach 1617 mAh g$^{-1}$ and 1226 mAh g$^{-1}$, respectively, showing a remarkably enhanced Columbic efficiency of 75.8% thanks to the unique nanostructure assembled by few-layered MoS$_2$ nanosheets on CNTs. In the next two discharge and charge processes, the discharge capacity can still reach 1310 mAh g$^{-1}$ and 1296 mAh g$^{-1}$ with Coulombic efficiency as high as 95% and 96%, respectively, demonstrating a high reversible capacity and excellent cycling stability.

Fig. 5 (a) Scheme illustration of the diffusion of electron and Li ion for the as-prepared MoS$_2$/CNTs hybrids, (b) Nyquist plots of the MoS$_2$/CNTs hybrids and the commercial MoS$_2$, respectively. The rate capability of the MoS$_2$/CNTs hybrids was further evaluated, as shown in Figure 4c. The average discharge capacities are 1293 mAh g$^{-1}$, 1203 mAh g$^{-1}$, 1092 mAh g$^{-1}$, 983 mAh g$^{-1}$ and 888 mAh g$^{-1}$ at current densities of 200 mA g$^{-1}$, 400 mA g$^{-1}$, 800 mA g$^{-1}$, 1600 mA g$^{-1}$ and 3200 mA g$^{-1}$, respectively. After the rapid charge and discharge at 3200 mA g$^{-1}$, a mean capacity of 1294 mAh g$^{-1}$ can be recovered when the current
density returns back to 200 mA g⁻¹. For comparison, the commercial MoS₂ was also tested under the same condition, showing much lower capacity of 716 mAh g⁻¹ at 200 mA g⁻¹, with poor rate performance (the capacity of only 192 mAh g⁻¹ at 3200 mA g⁻¹). Such high specific capacity and rate capability are superior or comparable at least to the best results reported for MoS₂-based electrode materials.²⁻¹²,²⁴⁻²⁷ Very recently, Yang et al.²⁴ reported the synthesis of hierarchical MoS₂/polyaniline nanowires which showed an intriguing specific capacity of 1062.7 mAh g⁻¹ at 200 mA g⁻¹ with ~ 30% capacity retention at 1000 mA g⁻¹, but still lower than our samples (1293 mAh g⁻¹ at 200 mA g⁻¹ with 888 mAh g⁻¹ capacity retention even at 3200 mA g⁻¹). After testing the rate performance, the MoS₂/CNTs hybrids subsequently continue to be evaluated at a current density of 400 mA g⁻¹ for another 200 cycles (Figure 4d). The hybrids show no capacity fading in the whole cycling process and deliver high specific capacity of ~ 1200 mAh g⁻¹ with a Coulombic efficiency of ~ 100%. The outstanding cycling stability would overwhelm the MoS₂-based anode materials in the literature, such as three-dimensional tubular architectures assembled by single-layered MoS₂ (73.8% capacity retention after 50 cycles),²⁸ hierarchical MoS₂/polyaniline nanowires (89.6% capacity retention after 50 cycles)²⁴ and MoS₂/amorphous carbon composites (95% capacity retention after 100 cycles).¹⁵ In a previous work, Chen et al.¹² reported the synthesis layered MoS₂/graphene composites, showing almost no capacity loss (1187 mAh g⁻¹ at 100 mA g⁻¹) after 100 cycles (1200 mAh g⁻¹ at 400 mA g⁻¹ after 200 cycles for our samples). Such excellent electrochemical performance is mainly attributed to the unique hierarchical nanostructure. As shown in Figure 5a, the introduction of CNTs builds high-speed conductive channel for MoS₂ nanosheets, greatly boosting the rapid electron transfer during Li insertion/extraction. To verify this viewpoint, the electrochemical impedance spectra of the MoS₂/CNTs hybrids and the commercial MoS₂ were performed. As shown in Figure 5b, the MoS₂/CNTs hybrids demonstrate a much lower resistance (~ 183.5 Ω) than the commercial MoS₂ (~ 541.6 Ω). On the other hand, the few-layer MoS₂ nanosheets were firmly and alternately assembled on the CNTs, which provided high structural sufficient electrochemical active sites and therefore resulting in high stability and meanwhile created amounts of porous configuration. The Li ion from the surrounding of MoS₂/CNTs have significantly improved contact with the Li accommodate layers, ensuring specific capacity and rate performance.

Conclusions
In conclusion, we have successfully realized the self-assembly of few-layer MoS₂ nanosheets on CNT backbone via a facile hydrothermal reaction with subsequent annealing process. In this structure, the few-layer MoS₂ nanosheets were alternately and vertically grown on the surface of CNTs forming a three-dimensional hierarchical nanostructure. The MoS₂ content can be easily controlled with a maximum content of 85%. Such MoS₂/CNTs hybrids could be applied as an intriguing anode material for the development of LIBs with high rate capability and long cycle life. Compared with the commercial MoS₂ (716 mAh g⁻¹ at 200 mA g⁻¹), the as-prepared MoS₂/CNTs hybrids demonstrated a much higher specific capacity of 1293 mAh g⁻¹ at 200 mA g⁻¹ with remarkably enhanced rate capability (888 mAh g⁻¹ even at 3200 mA g⁻¹). More significantly, they also possess a very high cycling stability, i.e. almost no capacity loss after over 200 cycles at 400 mA g⁻¹. Such overwhelming electrochemical performance endows the MoS₂/CNTs hybrids huge potential as an anode material for LIBs.

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Notes and references
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Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI.
⁵ X. Cao, Y. Shi, W. Shi, X. Rui, Q. Yan , J. Kong , H. Zhang, Small 2013, 9, 3433-3438.