MnO₂ Nanorods Intercalating Graphene Oxide/Polyaniline Ternary Composites for Robust High-Performance Supercapacitors

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New ternary composites of MnO₂ nanorods, polyaniline (PANI) and graphene oxide (GO) have been prepared by a two-step process. The 100 nm-long MnO₂ nanorods with a diameter ~20 nm are conformably coated with PANI layers and fastened between GO layers. The MnO₂ nanorods incorporated ternary composites electrode exhibits significantly increased specific capacitance than PANI/GO binary composite in supercapacitors. The ternary composite with 70% MnO₂ exhibits a highest specific capacitance reaching 512 F/g and outstanding cycling performance, with ~97% capacitance retained over 5000 cycles. The ternary composite approach offers an effective solution to enhance the device performance of metal-oxide based supercapacitors for long cycling applications.

Supercapacitor or electrochemical capacitors, featuring high power density, long cycle life, fast charge and discharge rates⁵⁷, have attracted tremendous attention in energy storage. During the last decades, different kinds of materials have been studied as possible supercapacitor electrodes, namely, carbonaceous materials, conducting polymers and transition metal oxides⁸¹⁰. Among them, transition-metal oxides such as RuO₂¹²⁻¹³, NiO¹⁴⁻¹⁵, and MnO₂¹⁶⁻²⁰, are the most widely used electrode materials thanks to their large capacitance, high surface area and relatively good electrical conductivity. Compared to other transition-metal oxides, MnO₂ is generally regarded as the most promising one for the next generation of supercapacitors because of its ideal capacitor performances, good cycle stability, low cost and environmental friendliness²¹,²². However, the poor electrical conductivity and densely packed structure limit its application in the development of high-performance supercapacitors²³. For this purpose, some efforts have been made to develop nanostructured MnO₂/carbonaceous materials (ca. graphene, carbon nanotube or carbon nanofibers) based binary composites²⁴,²⁵ or their ternary composites with conducting polymers²⁶⁻³¹. The carbonaceous materials or conductive polymers act as not only 3-dimensional (3-D) scaffold but conductive wrappings to enhance the specific capacitance and cycle stability of the electrodes fabricated with these composites. Our recent studies have shown that conductive polymer coatings prepared via in-situ redox process can greatly improve the conductivity and specific area of the resulted composites³²,³³.

As one of the most appealing carbon materials, graphene features superior electrical conductivity, excellent mechanical flexibility, and high thermal and chemical stability⁵⁴,⁵⁵. Compared with pristine graphene, graphene oxide (GO) has good dispersion in many solvents, particularly in water, and this is important for solution-processing and further derivatization⁶⁶⁻⁷⁷. Though some losses in electrical conductivity when converting from graphite, GO has demonstrated to be a potential substrate to improve the conductivity and stability of MnO₂²⁸. To enhance the performance of MnO₂-graphene composite electrodes, conductive materials were further incorporated to form ternary composites composites to achieved significantly increased specific capacitance and cycling-stability from the ternary composites based supercapacitors²⁶⁻³¹. By depositing 5–10 nm MnO₂/polyaniline (PANI) nanoparticles onto sulfonated graphene nanosheets, the composite (sGMOPANI) based two-electrode cell showed an electrochemical capacitance of 276 F/g at 1 A/g current density⁷⁸. The ternary composites of graphene/MnO₂ nanostructures with carbon nanotube or poly(3,4-ethylenedioxythiophene):poly(styrenesulfonate) showed ~20% or 45% increased capacitance than graphene/MnO₂ composites, with the highest value as
The MnO$_2$ crystalline phase forms in the synthesis, which is in good percentage of MnO$_2$ in the composites is about 20%, 46% and 70%, meaning that mass fraction.

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We report herein the successful fabrication of graphene oxide-polyaniline-MnO$_2$ ternary composites, where MnO$_2$ nanorods were incorporated to intercalate PANI/solution-exfoliated GO nanosheets. MnO$_2$ nanorods were thus fastened onto GO sheets via PANI layer (Fig. 1). The PANI layer has several advantages: (1) to anchor MnO$_2$ nanorods onto the surface of GO, (2) to prevent Go sheets and MnO$_2$ nanorods from stacking/aggregating; (3) to improve the charge transfer between GO and MnO$_2$; and (4) to prevent MnO$_2$ from falling off GO sheets during cycling electrochemistry. The composites exhibited greatly improved capacitive performance than GO-MnO$_2$ and GO-PANI composites. The ternary composites with 70 wt.% MnO$_2$ delivered a highest specific capacitance of 512 F/g at 0.25 A/g current density, with the capacitance retention of as high as 97% observed after 5100 cycles.

Results
Preparation and characterization of ternary composites. The ternary composites were prepared with a two-step protocol as shown in Fig. 1. After a first step of coating solution-exfoliated GO nanosheets with polyaniline, MnO$_2$ nanorods were allowed to self-assemble onto the GO sheets and then wrapped by in-situ prepared polyaniline coatings in the second step. The MnO$_2$ (in weight ratio) content in the ternary composites was determined with thermogravimetric (TGA) analysis (Figure 2). Hence, the as-prepared MnO$_2$-PANI-GO composites (GOPM) are designated as GOPM-20, GOPM-46 and GOPM-70, meaning that mass percentage of MnO$_2$ in the composites is about 20%, 46% and 70%, respectively.

Typical X-ray diffraction (XRD) pattern of as-prepared hierarchical porous GOPM-46 and free MnO$_2$, PANI and GO are shown in Fig. 3a. It can be seen that the XRD pattern of GOPM is similar to those of free MnO$_2$ and PANI, indicating that the MnO$_2$ particles have been well immobilized by PANI onto GO substrate. All of the major reflections [2θ = 21.2° (120), 28.0° (310), 37.5° (131), 42.5° (300), 59.0° (160) and 69.7° (003)] in the XRD pattern can be indexed to the orthorhombic phase of γ-MnO$_2$ with lattice constants a = 6.36 Å, b = 10.15 Å and c = 4.09 Å (JCPDS 14-644), confirming that γ-MnO$_2$ crystalline phase forms in the synthesis, which is in good agreement with the reported patterns for γ-MnO$_2$. The broad peak centered at about 25.3°, attributing to the diffraction of (110) plane of the as-formed PANI crystalline phase, suggests the successful synthesis of PANI. However, the sharp peak at 2θ = 11.0°, corresponding to its (002) plane for pure GO, almost disappeared, which is attributed to the increase in inter-space distance of graphene layer due to intercalation of PANI/MnO$_2$ between GO sheets.

The Raman spectra of GOPM and pristine components are shown in Fig. 3b. It can be clearly seen that both GOPM-46 and MnO$_2$ samples feature a sharp peak at 640 cm$^{-1}$, corresponding to the Mn-O vibration perpendicular to the direction of the MnO$_6$ octahedral double chains of MnO$_2$. The Raman-active peaks at 1350 cm$^{-1}$ and 1600 cm$^{-1}$, corresponding to the in-plane bond-stretching motion of C sp$^2$ atoms (G band) and the breathing modes of or of benzenoid rings of GO (D-band), were significantly suppressed in intensity for GOPM-46 sample. The typical peaks for PANI also weakened in GOPM, mainly due to its low content in the hybrid composite.

To confirm the XRD and Raman results, the compositions and the valence states of GOPM were further characterized with XPS and the results are shown in Fig. 3c-f. The peak ~284.6 eV (for C 1s) originating from the graphitic sp$^2$ carbon atoms and the peak at 531.9 eV corresponding to O1s in C–OH bond are observed for GO (Fig. 3c). The existence of C, N, O, Mn in GOPM can be confirmed from Fig. 3d. For GOPM, the XPS peaks of N 1s (Fig. 3e) are further decomposed into three Gaussian peaks with binding energies of 398.6 (-N=), 399.7 (-NH-) and 400.1 (-N$^+$-). The peak at 400.1 eV is assigned to the quinoid amine and nitrogen cationic radical (N$^+$), while the one at 398.6 eV (-N=) is due to benzenoid amine. The Mn 2p spectrum is analyzed in Fig. 3f. Both Mn 2p$_{3/2}$ peak at 634.0 eV and Mn 2p$_{1/2}$ peak centering at 654.5 eV are clearly observed, which are in good agreement with the energy splitting of the standard spectrum of MnO$_2$. The peak-to-peak separation between Mn 2p$_{3/2}$ and Mn 2p$_{1/2}$ level is 11.5 eV, which is approximately the same value as that literature for MnO$_2$. The morphologies of dry MnO$_2$, GO-PANI and GOPM-46 are observed with FESEM and TEM (Fig. 4.5). Pristine MnO$_2$ nanorods
with a length of 200 nm grow into nanospheres with diameters ranging from 300 to 600 nm (Fig. 4a). The in-situ prepared PANI self-assembled on GO sheets like a broccoli composed numerous of small flower buds with a diameter of 50 nm (Fig. 4b). When MnO₂ nanorods were introduced to GO-PANI system, GOPM hybrid composites feature a porous yet densely packed structure (Fig. 4c), with GO layers intercalated by porous interpenetrating networks of PANI-wrapped MnO₂ nanorods. PANI functions as both protecting coating for MnO₂ nanorods to avoid aggregation and adhesive to fasten GO sheets and MnO₂ nanorods into densely packed structure. Exfoliated GO sheets feature transparent wrinkled tulle in TEM image (Fig. 4d), while PANI formed a compact layer on top of GO sheets for GO-PANI composites.

A close look at the TEM images of GOPM (Fig. 4f and Fig. 5c,d), one would clearly observed the existence of MnO₂ nanorods, featuring a length less than 100 nm and a diameter ~20 nm. With the content of MnO₂ nanorods increasing from 46% to 70% in the composites, the dimensions of MnO₂ nanorods was observed to increase by 30% (Fig. 5d). Nevertheless, the intimate contact between GO sheets and PANI wrapped MnO₂ nanorods with different lengths can be clearly observed from both SEM and TEM images, which is important for improving electrical conductivity. A close look at the SEM image of ternary composites, GOPM-70 boasts increased porosity than GOPM-46, which can be confirmed by BET measurement. The GOPM-70 and GOPM-46 exhibit a surface area of 91.37 and 73.65 m²/g, respectively, much higher than that for GO-PANI (~36.92 m²/g). The increased surface area can be explained with the constitution of the ternary composites. Since PANI has the equivalent mass with GO, more MnO₂ nanorods were uncovered by PANI with the increase of MnO₂ content. More porous structure was therefore obtained to present higher surface area before heavy aggregation of MnO₂ occurs.

**Capacitive performance of ternary composite.** The capacitive performance of GO, MnO₂, GO-PANI, GO-MnO₂ and GOPM were evaluated by cycle voltammetry (CV) and galvanostatic...
charge/discharge techniques in 1 M Na$_2$SO$_4$ solutions. Fig. 6a shows the CV curves of GOPM-46 at different scan rates. At low scan rates ($\leq 50$ mV/s), GOPM-46 exhibits slight redox peaks, suggesting the pseudocapacitance feature of GOPM. The asymmetrical CV curves can be attributed to the combined double-layer and pseudocapacitive contributions to the total capacitance. The high CV currents indicate the high conductivity and low internal resistance for GOPM as the electrode material. Fig. 6b shows CV curves of GO, MnO$_2$, GO-PANI, GO-MnO$_2$ and GOPM at 50 mV s$^{-1}$, where GOPM-46 exhibits the highest capacitance than other electrodes. This can be attributed to the unique porous structure of hybrid GOPM, which effectively prevent the self-aggregation of GO sheets and MnO$_2$ nanorods. More importantly, the hybrid GOPM with PANI-coated MnO$_2$ nanorods intercalated GO nanosheets affords higher surface area to provide better conductive paths for fast electron transportation.

The galvanostatic charge/discharge curves of GOPM in 1 M Na$_2$SO$_4$ solution were carried out at a current density of 0.25, 0.5, 1, 2 and 4 A/g. As illustrated in Fig. 7a, all the curves exhibit an equilateral triangle shape, indicating high reversibility of the hybrid materials during charge/discharge process. The charging/discharging process took longer time at lower current density, which is attributed to the sufficient insertion or release of Na$^+$ during the charging/discharging process. The specific capacitance (C) of the electrode can be calculated according to equation: $C = \frac{I t}{V m}$, where C is the specific capacitance (F/g), I is the charge-discharge current (A), $t$ is the discharge time (s), V is the potential window (V), m is the mass of active material in the working electrode (g). The galvanostatic charge/discharge curves of MnO$_2$, GO-MnO$_2$, GO-PANI, GOPM at 0.25 A/g current density are compared in Fig. 7b. The specific capacitance of GOPM (409 F/g) is much higher than that of MnO$_2$ (150 F/g), GO-MnO$_2$ (162 F/g) and GO-PANI (20 F/g) at the same current density. Same trend is observed for GOPM when further increasing current density in the range of 0.25~4 A/g (Fig. 7c), indicating the robustness of the as-prepared hybrid GOPM as electrode materials.

Figure 5 | FE-SEM image of (a) GOPM-46 and (b) GOPM-70, TEM image of (c) GOPM-46 and (d) GOPM-70.

Figure 6 | (a) CV curves of GOPM-46 at scan rate of 10, 20, 30, 40, 50 and 100 mV s$^{-1}$; (b) CV curves of GO, MnO$_2$, GO-PANI, GO-MnO$_2$ and GOPM-46 at scan rate of 50 mV s$^{-1}$. 
The specific capacitance of GOPM is found to increase with the content of MnO$_2$ in the composite, as shown in Fig. 7c. The specific capacitance of GOPM-70 (412 F/g), GOPM-46 (306 F/g) and GOPM-20 (252 F/g) is about 1.2 times higher than that of MnO$_2$ (130 F/g) and GO-MnO$_2$ (156 F/g) at the current density of 1 A/g. Further increase of MnO$_2$ content over 90%, however, leads to decreased capacitive performance, e.g., the GOPM with 92% MnO$_2$ delivered a specific capacitance of 382 F/g at 1 A/g current density. The specific capacitance of 412 F/g at 1 A/g for GOPM-70 is much better than graphene/MnO$_2$ (78 wt.% MnO$_2$) binary composite under similar testing conditions. This improvement may be explained with the contribution of conductive PANI in facilitating charge transport and energy storage. Compared with the ternary composite sGMOPANI, our composite with MnO$_2$ nanorods exhibited 11% improved capacitance in supercapacitors at same current density of 1 A/g. The specific capacitance of GOPM decreases with current density, with ~62% capacitance retained for GOPM-70 when the current density increased from 0.25 A/g to 4 A/g.

The electrochemical stability of GOPM-70 nanocomposites was investigated at 4 A/g current density. As shown in Fig. 6d, the capacitance of GOPM electrode retained about 97% of the highest capacitance even after continuous galvanostatic charge/discharge process for 5100 cycles, indicating a good cycling ability of the hybrid composites. The support carbon matrix GO allowed the strong deposition of PANI-protecting MnO$_2$ nanorods on the surfaces of GO, which enhanced the mechanical strength of composite materials, resulting in the long charge/discharge ability. Interestingly, the specific capacitance showed slight increase in the first 150 cycles before decreased slowly in the later cycles, which may be explained with the insufficient contact of nanocomposites with Na$_2$SO$_4$ aqueous solution at the beginning of electrochemical measurement.

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The electrochemical properties of GOPM were further evaluated with electrochemical impedance spectroscopy (EIS). The impedance spectra of composite electrodes before and after 3000 cycles were measured in the frequency range of 100 kHz–0.1 Hz at open circuit potential with an AC perturbation of 5 mV (Fig. 8). Theoretically, an ideal Nyquist impedance plot features a semicircle over the high frequency region and a linear part in the low frequency range. The larger semicircle observed for the electrode corresponds to higher interfacial charge-transfer resistance ($R_{ct}$) for the layer on the electrode, attributed to the poor electrical conductivity of the materials. And the straight line of the Nyquist plot corresponds to the line ($R_s$) resulting from ion diffusion/transport, i.e., the more vertical line is indicative of an electrode more close to an ideal capacitor. The equivalent series resistance (ESR) of pristine MnO$_2$, GO-PANI, GOPM-46 and GOPM-70 obtained from the intersection of the Nyquist plot at the x-axis is 2.6, 0.7, 1.8, and 1.6 Ω, respectively. Considering the similar morphology of GO-PANI and GOPM, the difference in ESR of electrodes can be attributed to the different conductivities of electrode materials. The smaller ESR of GOPM than GO-PANI suggests the increased charge transfer resistance in the presence of MnO$_2$ nanorods as core. The high resistance of ion transfer in GO-PANI would be attributed to high charge density, resulting in low capacitance. In contrast, GOPM-46 exhibited a short diffusion path length of ions in the electrolyte, which could be seen from the low resistance of the capacitative part on the Nyquist plot. This may be explained by the structure of the composite: the formation of GOPM results in the surface charge of GO being compensated by the negative charge from both Cl$^-$ doped PANI and MnO$_2$ nanorods, leading to a lower resistance of ion transfer. Moreover, the presence of GO-PANI with high electrical conductivity resulted in a lower charge transfer. After 3000 cycles, the calculated ESR for...
**GO (0.1 g) dispersion in water (100 mL) was ultrasonicated (250 W, 220 V) for 1 h** during the charge/discharge cycling.

**Discussion**

By using a quick and facile two-step process, MnO$_2$ nanorods were successfully coated by in-situ prepared PANI and intercalated into GO sheets. The as-prepared ternary composites with different loading of MnO$_2$ were evaluated as electrode materials for supercapacitors. The ternary composites presented highly porous nanostructure with significantly increased surface area (91.57 and 73.65 m$^2$/g, respectively, for GOPM-70 and GOPM-46). With MnO$_2$ nanorods to facilitate the charge transport in energy storage application, the ternary composites exhibited increased specific capacitance with the loading of MnO$_2$ before a maximum reached at 70 wt.% MnO$_2$. The GOPM-70 electrode exhibits a highest specific capacitance of 512 F/g at 0.25 A/g current density. With PANI conductive coating, the electrode showed high electrochemical stability in cycling, with ~97% capacitance retention observed after 5100 cycles. Importantly, this design strategy may be employed for the fabrication of robust high-performance electrode with tailor-made hybrid composites composed of cabonaceous materials, conducting polymers and metal oxides.

**Methods**

**Preparation of MnO$_2$ nanorods**. Absorbent cotton (200 mg) was added to KMnO$_4$ solution (40 mM, 200 mL) with simultaneous vigorous stirring and ultrasonic irradiation for 20 min. The mixture suspension was then heated at 100°C for 24 h. A dark brown precipitate was obtained. The precipitates were collected by centrifugation and washed with deionized water and ethanol for 4 times. The title product was dispersed in hydrochloric acid (1 M) for use.

**Preparation of MnO$_2$/PANI/GOPM ternary hybrid nanostructures.** Freshly prepared GO(0.1 g) dispersion in water (100 mL) was ultrasonicated (250 W, 220 V) for 1 h to get an exfoliated yellow brown GO suspension. Aniline (0.12 g) was slowly added into the suspension and the stable GO/aniline suspension was obtained after stirring violently. A mixture of concentrated hydrochloric acid, ammonium persulfate (APS) in distilled water (10 mL) was then slowly added to the suspension under stirring. The molar ratio of aniline, hydrochloric acid and APS was 1:1:1. When the reaction was conducted in ice bath for 10 minutes, MnO$_2$ nanorods (0.6 g) dispersion in hydrochloric acid (1 M, 50 mL) was added to the reaction system together with second addition of aniline (0.12 g). The reaction was stirred for 12 h. Finally, the composite was filtered and rinsed with distilled water and ethanol in sequence to afford the title product (0.65 g) as a deep green solid.

**Characterization.** Thermogravimetric analysis (TGA) was conducted on a TA instrument TGA/SDTA851 at 20°C/min heating rate under nitrogen. Raman spectra were performed on a Renishaw laser confocal Raman spectrometer employing a 514 nm laser beam. The XRD patterns were recorded on an X-ray diffractometer (D8 Advance, Bruker, Germany) by using Cu Kα radiation (λ=1.54 Å) at 40 kV and 30 mA. XPS analyses were operated on a Thermo ESCALAB 250 instrument. Specific surface area of composites was measured on a BET (NOVA 1000, Quanta Chrome, America) under nitrogen atmosphere at 77 K. FE-SEM analyses were observed with a Hitachi HS4800 FESEM at an accelerating voltage of 15 kV. TEM analyses were taken with a JOEL JEM-2100 microscope.

The electrochemical properties of the composite were investigated on a CHI660D electrochemical workstation (Shanghai, China) with conventional three-electrode system at room temperature. To prepare the working electrodes, the polytetrafluoroethylene with acetylene black (15 wt%) in ethylene solution were added to as-prepared composites to produce a homogeneous paste before pressed onto nickel foam current collectors. The electrodes were then dried under vacuum at 60°C for 24 h. The electrochemical performance of composites was investigated with standard CV, galvanostatic charge–discharge and EIS technique in 1 M Na$_2$SO$_4$ solution. CV measurements were performed in voltage ranging from 0 V to 0.8 V at a scan rate of 10, 30, 50 and 100 mV/s, respectively. Charge-discharge processes were carried out galvanostatically at 0.25–4 A/g current density in 0–1 V voltage range.

**Figure 8** Nyquist impedance plots of (a) MnO$_2$, GO-PANI and GOPM composites before cycle and (b) GOPM after 3000 cycles.

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**Author contributions**

T.W. conceived the work. H.G., Z.L., L.Y. and Z.S. performed the experiments. H.G., K.E., T.J. and T.W. analyzed the data and prepared the manuscript.

**Additional information**

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