Three-dimensional Porous Graphene/Polyaniline Hybrids for High Performance Supercapacitor Electrodes

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Abstract:
Graphene-based composites took extensive attraction as electrodes for supercapacitors these years. Three-dimensional cross-linking porous graphene (3D rGO-m) was obtained by KOH activation to graphene modified by 1,2,4-triaminobenzene. 3D porous graphene/polyaniline hybrids (3D rGO-m/PANI) was prepared by the in-situ chemical oxidative polymerization. The rGO-m are reconstructed from 2D to 3D porous structure after KOH activation. The PANI nanorod arrays are successfully decorated on the surface of the 3D porous graphene sheets. The specific capacitance of the 3D rGO-m/PANI hybrids reach 985 F/g at 0.5 A/g. The capacitance retention of 3D rGO-m/PANI maintains 90% of its initial capacity after 1000 cycles, while rGO-m/PANI only keeps 83% of its initial capacity, the cycling stability of both hybrids are higher than that of pure PANI (69%).

Keywords: three dimension; porous graphene; polyaniline; hybrid materials; supercapacitor

1. Introduction

Compared to conventional capacitors and Li ion batteries, supercapacitors with outstanding charge/ discharge rate and high power capability have attracted considerable research, and will become a promising energy storage device [1]. Electrodes, as the key materials in energy storage devices, determine the performance of the supercapacitors, such as capability, delivery rates and cycling stability [2].

Among carbon based materials, such as active carbon, carbon nanotube, graphene is a highly promising electrode material in Electric double layer capacitor (EDLC) due to its high electrical conductivity, high specific surface area, high charge mobility, favorable flexibility, and good mechanical strength [3]. However, graphene sheets are easy to aggregate due to the strong $\pi-\pi$ interaction. The aggregation of graphene makes the channels among the sheets so narrow that the electrolyte ions are unable to access [4]. Compared to 2D graphene sheets, 3D porous graphene structure can provide larger specific surface area, which will contribute larger contact interfaces between electrode and electrolyte, and provide three-dimensional paths for ionic or electronic transport as well, so 3D porous structure graphene is considered to have excellent electrochemical performance.

Polyaniline (PANI) with low cost, easy synthesis and high pseudocapacitance has become a promising pseudo supercapacitor electrode material. PANI with highly-ordered nanowire arrays are beneficial to facilitate ion transport by shorten transport paths [5]. However, the inevitable volume expansion and contraction during change/discharge will result in mechanical degradation and low cycling stability, which constraint its practical application [6]. So, it is put forward that combining PANI with graphene or other inorganic materials such as metals or metal oxides as scaffold or bracket to confine the volume changes of PANI, hybrid materials will possess both high capacitance and good cycling stability [7-11].

In this work, we proposed an effective method to prepare 3D porous graphene/PANI hybrid materials by KOH activation and in-situ chemical oxidative polymerization methods. Graphene was modified by 1,2,4-triaminobenzene, the modifier serves as molecular spacer that decreases greatly the stacking of the graphene sheets. Graphene modified (rGO-m) is of thin sheets and crumpled morphology. KOH activation was used to etch pores on the rGO-m sheets to construct 3D porous structural graphene (3D rGO-m), and then 3D porous...
structure rGO-m/PANI hybrid materials was obtained by in-situ polymerization. The microstructure, morphology and electrochemical performance of the 3D rGO-m/PANI hybrid materials were characterized and investigated. The 3D rGO-m/PANI hybrid materials showed high specific capacitance and good rate capability and cycling stability.

2. Experimental

2.1 Materials

Graphite powder (30 µm) was from Nanjing XFNANO Materials Tech Co., Ltd. Aniline (An, AR), ammonium persulfate (APS, AR) were from Tianjin Damao Chemical Agent Company. Sodium carbonate anhydrous (Na2CO3, AR) and potassium hydroxide (KOH, AR) were from Tianjin Bodi Chemical Engineering Corporation. 1,2,4-triaminobenzene dihydrochloride (96%) was from Alfa Aesar. All reagents were used directly without any treatment.

2.2 Preparation of 3D porous rGO-m/PANI hybrids

2.2a 3D rGO-m: Graphene oxide (GO) was prepared by a modified Hummers method according to the reference [12]. Graphene modified by 1,2,4-triaminobenzene (rGO-m) was prepared according to the reference [13]. rGO-m were soaked in aqueous KOH solution with the mass ratio of KOH:rGO-m = 7 for 24 h, and then heat treated at 500 °C for 5 h. After activation, samples were sequentially washed by acetic acid and deionized water and further dried at 60 °C for 12 h. Samples obtained are named as 3D rGO-m. In comparison, the rGO was synthesized via the same procedure as rGO-m but without 1,2,4-triaminobenzene.

2.2b 3D porous rGO-m/PANI hybrids: The 3D rGO-m/PANI hybrids were synthesized by in-situ chemical oxidative polymerization method with aniline as a monomer in the presence of 3D rGO-m. The procedures were as followed: 3D rGO-m was ultrasonically dispersed in 1 M HCl for 0.5 h, and aniline and polyvinyl alcohol with 1:0.000075 molar ratio were added into 3D rGO-m suspension and performed another 0.5 h sonication and then decreased the temperature to 0 °C. A certain amount of APS was dropped in and kept stirring at 0 °C for 8 h. The hybrid materials were filtered and washed repetitively with DI water and ethanol, until the filtrate became colorless. Then the restructured of graphene fragments occurs. Restructuring among carbon fragments is due to the dangling bonds resulted from KOH activation. In the activation process, the etching of rGO-m matrix via redox reaction between C and KOH leaves defects and planar pores, and produces graphene fragments with a lot of C dangling bonds. Thus, the restructure of graphene fragments occurs.

Pure PANI is random relatively long rod with 400 nm thickness in the low side. The rGO-m with crumpled morphology suggests that the amino groups of 1,2,4-triaminobenzene are chemically bonded with graphene, which results in the morphology change of the rGO-m sheets. Typical FE-SEM images of activated graphene samples of 3D rGO-m are shown in figure 1(c), it can be seen clearly that the sample activated by KOH is reconstructed from 2D to 3D structure, and highly porous morphology can be observed as well. Restructuring among carbon fragments is due to the dangling bonds resulted from KOH activation. In the activation process, the etching of rGO-m matrix via redox reaction between C and KOH leaves defects and planar pores, and produces graphene fragments with a lot of C dangling bonds. Thus, the restructure of graphene fragments occurs.

2.3 Materials Characterization

The electrodes were prepared by active materials with 10% binder (polytetrafluoroethylene, PTFE). The two-electrode system was assembled with a cellulose paper as separator, carbon fibre paper as current collector and 1M H2SO4 aqueous as electrolyte.

X-ray diffraction (XRD, PANalytical) was analyzed with Cu Ka radiation (λ = 0.15418 nm) from 5° to 60°. Fourier transform infrared spectrometer (FT-IR, Bruker Equnox 55) was recorded at room temperature from 400 cm⁻¹ to 4000 cm⁻¹ at a resolution of 2 cm⁻¹, KBr powder pellet. The microstructure was observed by the Field Emission Scanning Electron Microscope (FE-SEM, Hitachi S4800).

All the electrochemical performance was studied on a PARSTAT 2237 electrochemical workstation. Cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS) and galvanostatic charge-discharge (GCD) measurements were carried out in a two-electrode system. CV and GCD measurements were conducted within the potential range of 0–0.8 V. CV curves were tested at a scan rate of 5, 10, 25, 50 and 100 mVs. GCD curves were set in the range from 0.01 Hz-100 kHz at different current densities.

The specific capacitance (Cs) of the active electrode material was determined from the GCD curves using the equation:

\[ C_s = \frac{4\Delta t}{m\Delta V} \]  

Where, \( I \) is discharge current (A), \( \Delta t \) is the discharging time (s), m is the total mass of active electrode material (g), and \( \Delta V \) is the voltage drop (V) upon discharging (excluding the IR drop). The energy density E was calculated by

\[ E = \frac{C_s \Delta V^2}{8} \]  

where the specific capacitance (Cs) was derived from GCD measurement. The same \( \Delta V \) was used as that in Cs calculation from GCD. The power density P was calculated by

\[ P = \frac{E}{\Delta t} \]  

where \( E \) referred to the energy density, \( \Delta t \) was the same as that in calculation of E.

3 Results and Discussion

3.1 Microstructure and morphology

After a hydrothermal reaction, rGO without modified appears flat and inter-stacked sheets under a FE-SEM (figure 1(a)). The rGO-m appears crumpled sheets (figure 1(b)). Some sheets are translucent, indicating that their thickness in the low side. The rGO-m with crumpled morphology suggests that the amino groups of 1,2,4-triaminobenzene are chemically bonded with graphene, which results in the morphology change of the rGO-m sheets. Typical FE-SEM images of activated graphene samples of 3D rGO-m are shown in figure 1(c), it can be seen clearly that the sample activated by KOH is reconstructed from 2D to 3D structure, and highly porous morphology can be observed as well. Restructuring among carbon fragments is due to the dangling bonds resulted from KOH activation. In the activation process, the etching of rGO-m matrix via redox reaction between C and KOH leaves defects and planar pores, and produces graphene fragments with a lot of C dangling bonds. Thus, the restructure of graphene fragments occurs.
in length and 60 nm in diameter (figure 2(a)). The FE-SEM images of rGO-m/PANI shows the structure of rGO-m supported PANI nanorod arrays with 100 nm in length and 50 nm in diameter (figure 2(b)), and the rGO-m sheets are obviously visible. 3D rGO-m/PANI shows the similar structure with rGO-m supported PANI nanorod arrays, but owing to the 3D cross-linking structure of rGO-m after KOH activation, the hybrids of rGO-m/PANI is also cross-linked (figure 2(c)), the morphology of nanorod array and cross-linking sheets is favor for the fast ion or electron transport and expected to increase the specific capacitance, which will be illustrates later.

Figure 1. SEM images of (a) rGO, (b) rGO-m and (c) 3D rGO-m.

Figure 2. SEM images of (a) PANI, (b) rGO-m/PANI and (c) 3D rGO-m/PANI.

Figure 3 represents the XRD patterns of GO, rGO-m, PANI, rGO-m/PANI and 3D-rGO-m/PANI. A broad and intense peak appears at $2\theta = 11.0^\circ$ attributed to the (001) plane basal diffraction of GO, with an inter-layer spacing of 0.80 nm in the layer-like GO sheets. After modification, the major peak of rGO-m is shifted to 26.4$^\circ$ (0.34 nm), it is indicated that rGO-m has poor stacking order, and some single layer rGO were formed by completely exfoliation, and fewer layer rGO and/or even loosely stacked layers exist as well [14].

As for rGO-m/PANI and 3D rGO/PANI hybrids, four broad peaks centered at $2\theta = 9.6^\circ$, 15.1$^\circ$, 20.3$^\circ$ and 25.2$^\circ$ can be observed, which are corresponded to (001), (011), (020) and (200) planes of PANI in an emeraldine salt form, respectively [15]. XRD patterns of rGO-m/PANI and 3D rGO/PANI hybrids are similar with PANI, and the peaks of rGO at around 26.4$^\circ$ and around 43.1$^\circ$ still can be observed in rGO-m/PANI and 3D rGO/PANI hybrids. This indicates that rGO is fully contacted with PANI and PANI is successfully coated on the rGO sheets with π-stacking between themselves [16].

Figure 4 gives the FT-IR spectra of GO, rGO, rGO-m, PANI, rGO-m/PANI and 3D-rGO-m/PANI. The spectrum of GO shows the absorption peak of carbonyl C=O ($\nu_{C=O}$ at 1722 cm$^{-1}$), benzenoid C=C ($\nu_{C=C}$ at 1574 cm$^{-1}$), epoxy C—O—C ($\nu_{C-O-C}$ at 1044 cm$^{-1}$), hydroxyl C—OH($\nu_{C-OH}$ at 1224 cm$^{-1}$) and O—H ($\nu_{O-H}$ at 3427 cm$^{-1}$) [17-19]. After reduction, the O—H peak at 3427 cm$^{-1}$ still can be observed, while the carboxyl peak at 1722 cm$^{-1}$ is unobvious. 1631 cm$^{-1}$ is corresponding to the bending vibration of C=C. 1631 cm$^{-1}$ peak in rGO-m is obviously stronger than that in rGO, and C—O—C peak around 1044 cm$^{-1}$ is adverse, and a new peak at 1405 cm$^{-1}$ appeared in the spectrum of rGO-m, which is due to the C=N stretching vibration in benzenoid rings, which further indicating that rGO-m is reduced more completely under alkaline condition. The rGO-m shows the N—H wagging vibration at 830 cm$^{-1}$ of secondary amine group [20], and two new peaks 1270 and 1127 cm$^{-1}$ in the spectrum of rGO-m are correponed to the stretching vibration of C—N and C=, respectively [21], demonstrating the amino functionalization result of rGO-m [48,49]. Compared with rGO-m, the new peak 1572 cm$^{-1}$ in the spectra of rGO-m/PANI and 3D-rGO-m/PANI is attributed to the C=C stretching vibration in the quinoid, and the stretching of C=N at 1127 cm$^{-1}$ is obviously intense and broad, which is due to the stretching vibration of C=N$^-$ in the polarized PANI structure, indicating the oxidation state existence in PANI [22]. 3D-rGO-m/PANI shows less absorption peaks of oxygen-containing functional group than rGO-m/PANI since the KOH activation at high temperature removed some of oxygen-containing functional group in rGO-m. All the above peaks observed in the spectrum of 3D rGO-m/PANI suggest that the PANI net-works are successfully decorated on graphene sheets.
3.2 Electrochemical performance

Figure 5(a) shows the CV curves of the 3D rGO-m/PANI hybrid at a scan rate of 5, 10, 25, 50 and 100 mV/s in a 1 M H₂SO₄ solution, respectively. 3D-rGO-m/PANI hybrid presents redox peaks, which are corresponding to the structural conversions of emeraldine/pernigraniline and leucoemeraldine/ emeraldine. These results indicate good pseudo-capacitance characteristics. The mirror-like and symmetrical image in 0.5 A/g galvanostatic charge-discharge (GCD) curves indicates that the reversible redox reaction proceed in 3D rGO-m/PANI, rGO-m/PANI and PANI (figure 5(b)). The specific capacitance of 3D rGO-m/PANI calculated from GCD curves is 985 F/g at 0.5 A/g, which is higher than that of rGO-m/PANI (633 F/g) and PANI (405 F/g).

Figure 5(c) depicts the Nyquist plots of 3D rGO-m/PANI and rGO-m/PANI hybrids. The obvious semicircle at the high frequency region can be seen. The straight line at the low frequency region is almost parallel to the vertical axis, which reflects the good capacity owing to the fast and reversible redox of PANI nanorod array. The first intersect

Figure 3. XRD patterns of GO, rGO, rGO-m, PANI, rGO-m/PANI and 3D rGO-m/PANI.

Figure 4. FT-IR spectra of GO, rGO, rGO-m, PANI, rGO-m/PANI and 3D rGO-m/PANI.

Figure 5. (a) CV curves of 3D rGO-m/PANI, (b) GCD curves at 0.5 A/g, (c) Nyquist plots and (d) cycling stability of 3D rGO-m/PANI, rGO-m/PANI and PANI.
of the Nyquist curve to the Z′ axis represents equivalent series resistance (R_s), and the diameter of the semicircle represents charge transfer resistance (R_{ct}) at the electrode/electrolyte interface. It can be seen that the 3D-rGO-m/PANI has lowest R_{ct} (0.3 Ω) and R_s (0.07 Ω) than that of rGO-m/PANI (R_{ct}=0.7 Ω, R_s=0.08 Ω) and PANI (R_{ct}=1.0 Ω, R_s=0.09 Ω). The low R_{ct} and R_s mean good electrical conductivity and fast charge transfer at the electrode and electrolyte interface, which can explain the high specific capacitance of 3D-rGO-m/PANI at a high current density. 3D rGO-m/PANI presents the energy density of 40.53 Wh/kg at the power density of 0.66 kW/kg.

Figure 5(d) is the cycling stability of the cell based on the 3D-rGO-m/PANI and rGO-m/PANI hybrids at 1 A/g for 1000 cycles. The capacitance of 3D-rGO-m/PANI after 1000 cycles still reaches 90% of its original one, while rGO-m/PANI only maintains 83% of its original one, the cycling stability of both hybrids are higher than that of pure PANI (69%), so it is concluded that the cycling stability of 3D-rGO-m/PANI has been enhanced after KOH activation to rGO. The improved cycling stability could be attributed to two aspects: (1) the three-dimensional cross-linking structure of 3D-rGO-m/PANI is favorable for the ion transport through a 3D continuous channel; (2) the electrode’s volume and structure changes during the charge-discharge cycling are controlled to lower degree by 3D porous structure.

4. Conclusions

In summary, 3D cross-linking porous graphene was prepared by KOH activation to the 1,2,4-triaminobenzene modified graphene precursor (rGO-m). After in-situ chemical oxidative polymerization, 3D porous rGO-m/PANI hybrids were obtained. The resulted 3D porous rGO-m/PANI hybrids possess PANI nanorod array on the 3D porous rGO-m. The 3D cross-linking porous structure provides high specific surface area, short and fast electrolyte diffusion pathways. The ion-diffusion and charge-transfer resistances of the 3D porous rGO-m/PANI hybrids are decreased. The specific capacitance is enhanced markedly compared to inactivated rGO-m/PANI hybrids and pure PANI. The maximum specific capacitance is 985 F/g at 0.5 A/g. Therefore, 3D porous rGO-m/PANI hybrids maybe become promising electrode materials for application in supercapacitors.

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