Enhanced electrochemical performance of MnO$_2$ nanoparticles: Graphene aerogels as conductive substrates and capacitance contributors

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**Capacitance calculation:**

The specific capacitance ($C_g, \text{ F g}^{-1}$) was calculated in the three-electrode system according to the following equations:

$$C_g = \frac{I \times \Delta t}{m \times \Delta V}$$

Where $I$ is the constant discharge current (A), $\Delta t$ is the discharge time (s), $\Delta V$ is the voltage window (V) during the discharge process, $m$ is the mass of the active material on the working electrode (g).

In the two-electrode asymmetric supercapacitor system, the specific capacitance was calculated based on the mass loading of the active materials both in electrodes. The specific capacity/capacitance is calculated based on the mass of $\text{MnO}_2/\text{NGA}$ for $\text{MnO}_2/\text{NGA}$ electrode, the mass of $\text{MnO}_2$ for $\text{MnO}_2$ electrode and the mass of NGA for NGA electrode, respectively.
First of all, we observe that NGA can absorb about 8 mL KMnO$_4$ (0.1 mol L$^{-1}$) in the scope of 24 h. We compare the weight of the final products from different synthetic procedures as illustrated in Fig. S1c. Procedure A: The NGA, KMnO$_4$ and MnSO$_4$ (8 mL 0.1 mol L$^{-1}$ KMnO$_4$ + 32 mL 0.0625 mol L$^{-1}$ MnSO$_4$) are mixed and heated in the sealed Teflon-lined autoclave for 12 h at 120 °C. Procedure B: The NGA (after soaking 8 mL 0.1 mol L$^{-1}$ KMnO$_4$ without freeze-drying) and the MnSO$_4$ solution (32 mL 0.0625 mol L$^{-1}$ MnSO$_4$) are mixed and heated in the sealed Teflon-lined autoclave for 12 h at 120 °C. Procedure C: The NGA (after soaking 8 mL 0.1 mol L$^{-1}$ KMnO$_4$ and freeze-drying) and the MnSO$_4$ solution (40 mL 0.05 mol L$^{-1}$) are mixed and heated in the sealed Teflon-lined autoclave for 12 h at 120 °C. As shown in Fig. S1c, the weight increase of the product from procedure C is the highest among these three samples, suggesting the most efficient loading of MnO$_2$. For the synthetic procedure A, the reaction between KMnO$_4$ and MnSO$_4$ starts from outside of NGA, and quite a few MnO$_2$ may form outside NGA. For procedure B, NGA without freeze-drying is fulfill with the KMnO$_4$ solution. These KMnO$_4$ will easily diffuse to the external surface of NGA, and some MnO$_2$ will still form outside NGA. For
procedure C, the KMnO$_4$ in NGA is in solid state after freeze-drying. In this case, most of the reactions between KMnO$_4$ and MnSO$_4$ happen inside the pores of NGA, and MnO$_2$ species form mainly inside NGA, leading to the highest MnO$_2$ content among these three synthetic procedures. If we compare the procedure B and C, during which the reactions between MnSO$_4$ and KMnO$_4$ mostly would happen inside NGA, the only difference is that whether there is solvent inside NGA. We think that with the freeze-drying process in procedure C, the KMnO$_4$ solid would be “confined” inside but not diffuse outside NGA, leading to larger amount of the reactions happen inside NGA. That is the reason why the product from procedure C has higher content of MnO$_2$ than from procedure B, and the freeze-drying process would indeed enhance the loading content of MnO$_2$.

SEM image and the electrochemical performance of the product without freeze-drying are shown in **Fig. S1d**. As shown in the SEM image, there is less MnO$_2$ formed inside the NGA in the product from procedure B than that from procedure C. The GCD cures of product from procedure B shows the specific capacity of 216.6 mAh g$^{-1}$ at 0.1 A g$^{-1}$, which is lower than the specific capacity of MnO$_2$/NGA obtained from procedure C (275.8 mAh g$^{-1}$ at 0.1 A g$^{-1}$).
Fig. S2. (a) XPS spectra of the MnO$_2$/NGA (the inset illustrates the narrow spectra of Mn 2P peaks). (b) XPS spectra of the NGA before cycle and after cycles as electrode for Zn supercapacitor. (c) SEM image of NGA. (d) SEM image of MnO$_2$/NGA.

Table S1. C, O, N and Mn contents (atomic concentration %) in MnO$_2$/NGA from XPS.

<table>
<thead>
<tr>
<th>Samples</th>
<th>C</th>
<th>O</th>
<th>N</th>
<th>Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td>MnO$_2$/NGA</td>
<td>60.72</td>
<td>26.41</td>
<td>1.42</td>
<td>11.45</td>
</tr>
</tbody>
</table>
Table S2. C, O, N, Zn and Mn contents (atomic concentration %) in NGA before and after charging/discharging cycles

<table>
<thead>
<tr>
<th>Samples</th>
<th>C</th>
<th>O</th>
<th>N</th>
<th>Zn</th>
<th>Mn</th>
</tr>
</thead>
<tbody>
<tr>
<td>NGA before cycle</td>
<td>81.52</td>
<td>15.24</td>
<td>3.18</td>
<td>0.06</td>
<td>0</td>
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<tr>
<td>NGA after cycles</td>
<td>80.32</td>
<td>15.97</td>
<td>3.43</td>
<td>0.12</td>
<td>0.16</td>
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</tbody>
</table>

Fig. S3. (a) CV curves of the MnO₂/NGA at 0.1 mV s⁻¹ (the first and the 10th cycle). (b) Cycling performance at the current density of 0.5 A g⁻¹ with the inset as the GCD curves at the initial stage and after 100 cycles.
**Fig. S4.** (a) Long-term cycling performances of the MnO$_2$ at 3 A g$^{-1}$. (b) Comparison of the Ragone plot of the MnO$_2$/NGA-based battery with some other cathode materials for ZIB.

**Fig. S5.** (a) CV curves of the NGA at different scan rate from 5 mV s$^{-1}$ to 2000 mV s$^{-1}$. (b, c) GCD curves of the NGA at different current densities from 0.1 A g$^{-1}$ to 3 A g$^{-1}$. (d) CV curves of the MnO$_2$ at different scan rate from 5 mV s$^{-1}$ to 500 mV s$^{-1}$. (e, f) GCD curves of the MnO$_2$ at different current densities from 0.1 A g$^{-1}$ to 3 A g$^{-1}$. 
Fig. S6. (a) Cycling stability of the MnO$_2$/NGA at a current density of 3 A g$^{-1}$ with the GCD curves after 5000 cycles as the inset. (b) Nyquist plots of experimental impedance data for the MnO$_2$/NGA in the frequency range of 100 kHz – 0.1 Hz before and after 5000 cycles with inset of the enlarged EIS.
Fig. S7. Capacitive performance of the asymmetric supercapacitor (porous carbon/MnO$_2$/NGA). (a) CV curves with different voltage windows at a scan rate of 30 mV s$^{-1}$. (b) CV curves with a voltage window of 1.5 V at different scan rates. (c) GCD curves at different current densities. (d) Cycling performance at a current density of 1 A g$^{-1}$ with the inset as the GCD curves at the initial stage and after 500 cycles.