High-rate-capability asymmetric supercapacitor device based on lily-like Co$_3$O$_4$ nanostructures assembled using nanowires†

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Following the growing demand for high-efficient energy storage systems, high-current-density supercapacitor devices are needed; however, their development is still a challenge. An electrode material with high electronic conductivity and abundant ion channels is an important area of research. Herein, hierarchical lily-like Co$_3$O$_4$ nanostructures constituted using nanowires with a large L/D ratio on Ni foam were in situ synthesized using a simple solvent-thermal method and subsequent sintering treatment. The as-prepared Co$_3$O$_4$ precursor was composed of nanowires with a ~50 nm average diameter and ~70 L/D ratio. The product morphology was well maintained after thermal treatment, and the product was transformed to cubic Co$_3$O$_4$ during this process. Co$_3$O$_4$/Ni exhibited excellent rate capability and cycling performance, which showed a specific capacitance of 1600 F g$^{-1}$ even at a relatively high 10 A g$^{-1}$ current density that barely declined after 5000 cycles. Furthermore, the assembled asymmetric supercapacitor device using the as-prepared Co$_3$O$_4$/Ni and activated carbon showed a high specific capacitance at a high current density (108.1 F g$^{-1}$ at a current density of 0.5 A g$^{-1}$, 96.4 F g$^{-1}$ at 10 A g$^{-1}$ and 77.9 F g$^{-1}$ even at 50 A g$^{-1}$) and displayed a long-term cycling stability performance (10 000 cycles) with a high energy density (34 W h kg$^{-1}$) at a high power density of 1963 W kg$^{-1}$. The superior performance of lily-like Co$_3$O$_4$ nanostructures on Ni foam is proposed to be attributed to benign electronic conductivity and the presence of abundant ion transport channels. Therefore, these experimental results suggest the potential application of cobalt-based oxides in a high-current-density electrochemical system.

Introduction

Following the over-exploitation of fossil fuels with rapid economic development, energy crisis is becoming increasingly serious. Therefore, clean sustainable energy or high-efficiency energy storage/conversion system is urgent to be developed in order to improve the energy utilization efficiency. For example, following the pressing demand for mobile devices and electric mobiles with fast-charge and high energy storage properties, there is an urgency to develop a high-efficiency electrochemical storage device. To date, lithium/sodium ion batteries, lithium–sulfur batteries, supercapacitors and so on, have received considerable attention. Among them, supercapacitors possess unique advantages of fast charge/discharge, environment friendliness and a long lifetime. Significant efforts have been devoted in developing high-energy-density, long cycling life supercapacitors. Compared with electric double-layer capacitors, pseudocapacitors exhibit higher specific capacitance due to fast redox reactions on the surface of electrode materials. However, the cycling life and rate performance of the pseudocapacitors is unsatisfactory. To meet the urgent demand in high-efficiency electrical equipment, improving the long-term cycling life and excellent rate performance of supercapacitors is imperative. The electrode material is the key part of a supercapacitor device and has a decisive effect on the supercapacitor performance. Therefore, much effort has been made on the design and exploitation of electrode materials with higher specific capacitance and long cycling life. However, a high-current-density supercapacitor device is needed to be developed, which is still a challenge.

The ideal electrode material for a supercapacitor probably possesses the properties of high electronic conductivity and abundant ion channels. In recent years, the research in electrode materials for supercapacitors has mainly concentrated on carbon, conducting polymers, metal oxides, metal sulfides, and so on. Among them, metal oxides, for
example Co3O4,29 NiO39 MnO2,31 are considered as important candidate electrode materials in supercapacitors due to their high specific capacitance. However, inferior electronic conductivity of metal oxides greatly hinders electron transport and thus weakens the rate and cycling performance of supercapacitors. Therefore, compositing the conductive material (e.g., metal or carbon materials) with these electrode materials is probably an effective approach to overcome this problem.24,39,40 Nickel foam possesses fine electronic conductivity and ductility, which can be considered as an ideal supporter in electrochemical power sources.41

The size and morphology of electrode materials greatly determine the exposed active surface, which show a significant effect on the electrochemical behavior of the supercapacitor device.2 A large specific surface area can accelerate surface redox reactions and enhance the contact between the active material and electrolyte. By decreasing the size, the electrode materials can expose more active surface, thus enhancing the specific surface area and shortening the ion channel, to finally improve the electrochemical performance. The morphology of electrode materials is also an important factor that influences electrochemical behavior. Due to the unique structure and high number of exposed active sites, one-dimensional nanostructures (e.g., nanowires, nanorods, nanotubes) have been devoted large attention in catalysts and energy storage/conversion fields. For example, carbon nanotubes, Co3O4 nanorods, MnO2 nanotubes, and so on have been applied in the electrode material of lithium ion batteries or supercapacitors.42–47 As mentioned previously, electrode materials with high specific capacitance and long term cycling for high performance supercapacitors should be equipped with high specific area and excellent electronic conductivity. Therefore, constructing the composite nanostructure of one-dimensional metal oxide with a conductive metal support is probably an effective approach to design an advanced supercapacitor device. For example, one-dimensional Co3O4 nanostructures showed preferable electrochemical properties in previous studies.48–52 To date, large attention has been drawn towards using different methods for the synthesis of one dimensional Co3O4 nanostructures with excellent electrochemical properties (e.g. hydro-thermal method).53,54 However, high-current-density supercapacitor devices based on cobalt-based materials are rarely reported and remain a current challenge.

In this study, hierarchical lily-like Co3O4 microflowers on Ni foam, which are assembled using nanowires, were fabricated via a simple solvent-thermal method and subsequent sintering treatment. Systematic characterizations were conducted to understand the formation mechanism and electrochemical behavior of the as-prepared product. The electrochemical performance (specific capacitance, long cycling life, energy density, power density) were evaluated in three-electrode and two-electrode systems assembled using the as-prepared Co3O4 on Ni foam and activated carbon (Co3O4/Ni//AC). In addition, practical application was evaluated through choosing the commercial light-emitting diodes (LEDs) as a load model.

**Experimental**

**Synthesis of Co3O4 nanostructures**

All chemical reagents were of analytical grade and used without further purification. Co(NO3)2-6H2O (0.5 mmol) and urea (CO(NH2)2) (10 mmol) were dissolved in a mixed solvent (4 mL ethanol, 16 mL H2O). After stirring on a magnetic stirring apparatus for 20 min, Ni foam was put into the abovementioned mixed solution. Then, the mixed suspension was transferred into a Teflon-lined stainless steel autoclave and heated at 150 °C for 15 h. The products (Co-precursor) were washed with deionized water and ethanol and further dried at 80 °C overnight. Finally, Co-precursor was heated at 350 °C for 2 h in an Ar atmosphere to obtain a Co3O4 sample on Ni foam. In addition, for comparison, Co3O4 powder was obtained from a similar method to the abovementioned method for Co3O4 sample on Ni foam except for the absence of Ni foam. The mass loading of Co3O4 on Ni foam was calculated through a difference method. The mass loading of Co3O4 for further measurement is about 2 mg (~2 mg cm−2).

**Instrument and characterization**

To realize the physical and chemical properties of the Co3O4 sample, some characterizations were carried out. The morphology of product was checked with a Zeiss Merlin Compact field emission scanning electron microscope (FESEM) equipped with an energy-dispersive X-ray spectroscopy (EDX) system. The microstructure and composition of the as-prepared Co3O4 sample were detected via transmission electron microscopy (TEM, JEOL, JEM-2100) and high-resolution TEM (HR-TEM, JEOL JEM-2010F). We examined the phase via X-ray diffraction (XRD) on a Bruker D8 Advance X-ray powder diffractometer using Cu-Kα irradiation at a scan rate of 4° min−1.

**Electrochemical properties measurement**

All the electrochemical measurements were carried out using a galvanostatic charge/discharge tester (CT2001A) and electrochemical workstation (CHI 660E) in both standard three-electrode mode and two-electrode mode. For the two-electrode mode, the working electrode was the as-prepared Co3O4 product on Ni foam, and the counter electrode was activated carbon (AC) on Ni foam (in which the active carbon, acetylene black, polyvinylidene fluoride (PVDF) with an 8 : 1 : 1 mass ratio). The working electrode and counter electrode can be separated with a piece of nonwoven separator in 1 M KOH solution. For the three-electrode mode, the reference electrode was a Hg/HgO electrode, and the other conditions (for example, working electrode and electrolyte) were similar to the two-electrode mode. Furthermore, the specific capacitance (C), energy density (E) and power density (P) could be calculated from the charge/discharge curve as follows:

\[ C = \frac{I \Delta t}{m \Delta V} \left( \text{F g}^{-1} \right) \] (1)
where \( I \) represents constant discharge current, \( \Delta t \) represents discharge time, \( m \) represents the mass of active material, and \( \Delta V \) represents width of the potential window.

\[
E = \frac{1}{2}(C\Delta V^2) \\
P = \frac{E}{\Delta t}
\]

The electrochemical impedance spectroscopy (EIS) data were recorded using a CHI660E electrochemical workstation (Chenhua, Shanghai, China) in the 0.01 Hz and 100 kHz frequency range. In addition, the commercial red light-emitting diodes (LEDs) as a load model were lightened by connecting two Co3O4/Ni//AC supercapacitor devices in series, which were charged at a current density of 5 A g\(^{-1}\).

**Results and discussion**

**Synthesis and characterization**

The Co3O4 product was fabricated through sintering treatment of the as-prepared precursor (Co-precursor), which was prepared via a simple solvent-thermal method. Fig. 1 shows the SEM images of the as-prepared Co-precursor and Co3O4 product under different magnifications. Shown from Fig. 1a, the obtained Co-precursor has lily-like nanostructures composed of one-dimensional nanowires. The petal was assembled using nanowires with different length/diameter (L/D) ratios, and some nanowires even could be curved due to the large L/D ratio. After calculation, the L/D ratio of the petal was about 70. The pistil was assembled using shorter nanorods. From the high-resolution SEM image in Fig. 1b, the diameter of nanowire or nanorod was about 50 nm. After thermal treatment, the morphology of product was well maintained from the precursor, except that the nanowire changed from smooth to rough. The one-dimensional nanowires could be observed to consist of nanoparticles based on high-resolution SEM imaging of the Co3O4 product, which is shown in Fig. 1d. The rough one-dimensional nanowires, which appeared like a necklace, probably possessed a high exposure to surface and sites, which facilitated the electron transfer.

**Electrochemical performance**

The electrochemical properties of the as-prepared Co3O4/Ni product, as a supercapacitor electrode material, were evaluated in three- and two-electrode systems. Fig. 4 demonstrates the cyclic voltammetry (CV) curves tested at different scan rates and charge–discharge performance of as-prepared product in the three-electrode system. From the cyclic voltammetry (CV) curves of Co3O4/Ni at different scan rates with 1 M KOH electrolyte at
a potential sweep window 0–0.6 V (Fig. 4a), redox peaks can be clearly observed, indicating pseudocapacitive behavior of cobalt oxides due to the redox reaction between Co^{2+} and Co^{3+}. Fig. 4b exhibits CV curves of sample after 2000 cycles, which are similar to the original CV curves, indicating the cyclic stability of as-prepared Co_{3}O_{4} electrode. Fig. 4c displays the charge-discharge curves and Fig. 4d shows the rate capability of as-prepared Co_{3}O_{4}/Ni. The discharge time at the current densities of 0.5, 1, 2, 5, 10 A g^{-1} is 1847, 905, 445, 177, 88 s, respectively. The corresponding specific capacitance of Co_{3}O_{4}/Ni was 1679, 1645, 1618, 1609, and 1600 F g^{-1} at the current densities
The shape of curves is deviated from the ideal rectangular of 0.5, 1, 2, 5 and 10 A g\(^{-1}\), respectively, indicating excellent discharge performance at a high current density. This excellent rate capability demonstrated superior discharge performance at high current density, which has potential application in commercial supercapacitors. In addition, the specific capacitance barely declined after 5000 cycles at a current density of 10 A g\(^{-1}\) (Fig. 4e). Compared with those in previous studies (Table 1), the specific capacitance and cycling performance at high current density is notably improved.\(^{48–52,55–57}\)

The asymmetric supercapacitor device was further assembled with the as-prepared Co\(_3\)O\(_4\)/Ni and active carbon (Co\(_3\)O\(_4\)/Ni/AC). The electrochemical performance was evaluated and the results are shown in Fig. 5. The cyclic voltammetric (CV) curves of Fig. 5a were tested at a scan rate of 2–100 mV s\(^{-1}\) with 1 M KOH electrolyte at a potential sweep window of 0–1.6 V. The shape of curves is deviated from the ideal rectangular voltammogram due to the pseudocapacitive behavior of cobalt oxides. Based on that, pure Ni foam has little contribution to the capacitance, and the pseudocapacitive behavior is attributed to the as-prepared cobalt oxides. Fig. 5b and c give the discharge curves of Co\(_3\)O\(_4\)/Ni//AC device. When the current density was 0.5 A g\(^{-1}\), the discharge time of Co\(_3\)O\(_4\)/Ni//AC device was high, 1477 s (the specific capacitance was 108.1 F g\(^{-1}\)). When the current density increased to 10 A g\(^{-1}\), the discharge time was 76 s (the specific capacitance was 96.4 F g\(^{-1}\)). Fig. 5d exhibits the rate capacity of the Co\(_3\)O\(_4\)/Ni//AC device. When the current density was increased to 20 A g\(^{-1}\) and even 50 A g\(^{-1}\), the specific capacitances were still maintained at 86 F g\(^{-1}\) and 78 F g\(^{-1}\), indicating the potential application in fast charge–discharge devices. Fig. 5e exhibits the cycling performance and coulombic efficiency of Co\(_3\)O\(_4\)/Ni//AC device and Co\(_3\)O\(_4\) powder//AC device. Compared with Co\(_3\)O\(_4\) powder, Co\(_3\)O\(_4\)/Ni//AC device possessed higher cycling capacity and coulombic efficiency. As shown in Fig. 5e, the specific capacitance of the Co\(_3\)O\(_4\)/Ni//AC device was faultlessly maintained at a high current density of 10 A g\(^{-1}\) after 10 000 cycles. It is proposed that the Ni substrate and binder-free equipment method increased the electron conductivity of Co\(_3\)O\(_4\).

We further conducted electrochemical impedance spectroscopy (EIS) measurement and the result is shown in Fig. 6a, where the impedance of Co\(_3\)O\(_4\)/Ni decreased evidently. In addition, the relationship between energy density (E) and power density (P) is represented in Fig. 6b. As is known, Co\(_3\)O\(_4\)/Ni//AC supercapacitor device density of 1973 W kg\(^{-1}\) also can be as

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**Table 1** The electrochemical supercapacitor performance in previous studies\(^a\)

<table>
<thead>
<tr>
<th>References</th>
<th>Material</th>
<th>(C^f) (F g(^{-1}))</th>
<th>(\varepsilon^d) (W h kg(^{-1}))</th>
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<tr>
<td>Ref. 48</td>
<td>Co(_3)O(_4)/MnO(_2)</td>
<td>560 (0.2 A g(^{-1}))</td>
<td>17.7 (158 W kg(^{-1}))</td>
</tr>
<tr>
<td>Ref. 49</td>
<td>Co(_3)O(_4)/rGO/CNT</td>
<td>790 (1 A g(^{-1}))</td>
<td>19.6 (7250 W kg(^{-1}))</td>
</tr>
<tr>
<td>Ref. 50</td>
<td>Co(_3)O(_4)/graphene</td>
<td>494.2 (7.2 A g(^{-1}))</td>
<td>—</td>
</tr>
<tr>
<td>Ref. 56</td>
<td>Co(_3)O(_4)</td>
<td>378 (1 A g(^{-1}))</td>
<td>—</td>
</tr>
<tr>
<td>Ref. 57</td>
<td>3D Co(_3)O(_4)/rGO</td>
<td>660 (0.5 A g(^{-1}))</td>
<td>40.65 (340 W kg(^{-1}))</td>
</tr>
<tr>
<td>This work</td>
<td>Co(_3)O(_4)/Ni foam</td>
<td>1600 (10 A g(^{-1}))</td>
<td>34 (1963 W kg(^{-1}))</td>
</tr>
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</table>

\(^{a}\) Note: \(C^f\) represents the specific capacitance in a three-electrode mode, \(\varepsilon^d\) represents the energy density in a two-electrode mode.
high as 34 Wh kg$^{-1}$, which displays a maximum energy density of 38 Wh kg$^{-1}$ at a power density of 100 W kg$^{-1}$. Importantly, the energy density at a power confirmed the excellent rate capability of the Co$_3$O$_4$/Ni//AC supercapacitor device. Importantly, the power density could be as high as 10 060 W kg$^{-1}$.

These results demonstrated excellent electrochemical energy storage performance of Co$_3$O$_4$/Ni electrode material and indicated commercial high-current supercapacitor device potential due to the superior rate capability. Therefore, this outstanding performance of the as-prepared binder-free Co$_3$O$_4$ electrode material is proposed to be attributed to good electronic conductivity resulting from the Ni substrate and abundant ion channels resulting from the one-dimensional structure. Moreover, practical application was evaluated through choosing the commercial light-emitting diodes (LEDs) as a load model by connecting two Co$_3$O$_4$/Ni//AC supercapacitor devices in series, and the results are shown in Fig. 6c.

Conclusion

In conclusion, using a simple solvent-thermal method and subsequent sintering treatment, lily-like Co$_3$O$_4$ nanostructures on Ni foam were successfully prepared. XRD analysis indicated high purity of the as-prepared Co$_3$O$_4$ product. SEM images showed that the as-prepared Co$_3$O$_4$ nanostructures were composed of nanowires with large $L/D$ ratio ($\sim$70) and an average diameter of $\sim$50 nm. The specific capacitance of the as-prepared Co$_3$O$_4$ nanostructures at a high current densities of 5 A g$^{-1}$, and 10 A g$^{-1}$ were 1609 F g$^{-1}$ and 1600 F g$^{-1}$, respectively, exhibiting excellent rate performance. The specific capacitance was perfectly maintained after 5000 cycles even at a high current density of 10 A g$^{-1}$ after 5000 cycles. Furthermore, the assembled asymmetric supercapacitor device using the as-prepared Co$_3$O$_4$/Ni and activated carbon exhibited a high specific capacitance (108.1 F g$^{-1}$ and 96.4 F g$^{-1}$ at current densities of 0.5 A g$^{-1}$ and 10 A g$^{-1}$, respectively) and displayed a long-term cycling performance with a high energy density (34 Wh kg$^{-1}$) at a power density of 1963 W kg$^{-1}$. Importantly, the maximum power density could be as high as 10 060 W kg$^{-1}$. Moreover, the Co$_3$O$_4$/Ni//AC device presented an excellent performance over a Co$_3$O$_4$ powder device due to the benign electronic conductivity.
resulting from the Ni substrate and abundant ion transport channels resulting from the one-dimensional structure. These results demonstrated the excellent electrochemical energy storage performance of the Co$_3$O$_4$/Ni electrode material and indicated it to be a commercial high-current supercapacitor device due to its superior rate capability.

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**Notes and references**


