Petal-like CoMoO$_4$ Clusters Grown on Carbon Cloth as a Binder-Free Electrode for Supercapacitor Application

Chuanhong Chen, Hangchun Deng, Chongshi Wang, Wenqing Luo, Dejuan Huang,* and Tianxiang Jin*

Cite This: ACS Omega, 2021, 6, 19616−19622

ACCESS Metrics & More

ABSTRACT: The development of supercapacitors with a high energy density and power density is of great importance for the promotion of energy storage technology. In this study, we designed and prepared petal-like CoMoO$_4$ clusters combined with carbon cloth as an excellent self-standing and binder-free electrode for asymmetric supercapacitors. Due to the abundant electrochemical active sites, the promising electron conduction, and ion diffusion rate, the CoMoO$_4$@carbon cloth (CoMoO$_4$@CC) electrode exhibits an excellent electrochemical performance. The results show that the CoMoO$_4$@CC material exhibits a high specific capacitance (664 F/g at a current density of 1 A/g) and an excellent cycle stability (capacitance remains at 84.0% after 1000 cycles). The assembled symmetrical supercapacitor has an energy density of 27 Wh/kg when the power density is 600 W/kg. Even at a higher power density (6022 W/kg), it still maintains a good energy density (18.4 Wh/kg).

1. INTRODUCTION

With the rapid development of the society, energy consumption and environmental pollution are increasing. The use of clean energy and renewable energy has become indispensable. The development of high-power, high-capacity alternative energy conversion/storage devices, especially environmentally friendly energy conversion/storage devices, has a special significance. Supercapacitors (SCs) are considered to be one of the most prominent and most efficient energy storage devices due to their high power density, environmental friendliness, fast charge and discharge capability, and long cycle life. In terms of the materials of electrodes, binary metal oxides (BTMOs) possess high electrochemical activity and have been widely studied to be promising electrode materials for energy storage devices. Especially, metal molybdates, which are AMoO$_4$-type compounds (where A is a divalent metal ion), have potential applications in many fields such as magnetism, photoluminescence, and catalysts. With the advantages of the high redox activity, environmental friendliness, low cost, and abundant sources, metal molybdates, such as NiMoO$_4$, CoMoO$_4$, and ZnCo$_2$O$_4$, have attracted more and more attention.

In the traditional SC manufacturing process, the active material is often combined with a current collector through the use of a polymer binder, which creates a “dead mass” in the active material, leading to a poor specific capacitance. Moreover, most reported composite electrodes are usually made by the conventional slurry coating technology, in which the binder involved undoubtedly increases the internal resistance of the electrode. To solve these problems, many researchers grew electrochemically active substances directly on the substrate to prepare binder-free electrodes.

In this work, we have developed a simple and effective method to prepare a CoMoO$_4$@CC material with a petal-like microstructure. CoMoO$_4$ is directly grown on carbon cloth to form a self-standing and binder-free electrode. The results prove that the CoMoO$_4$@CC electrode has a high specific capacity of 664 F/g at a current density of 1 A/g, and the capacitance retention rate exceeds 84% after 1000 cycles at a current density of 10 A/g. A symmetrical supercapacitor based on the CoMoO$_4$@CC material was assembled and tested for electrochemical performance. The symmetrical supercapacitor has a remarkable energy density of 27 Wh/kg when the power density is 600 W/kg.

2. RESULTS AND DISCUSSION

2.1. Materials Characterization. The morphology and nanostructure of the CoMoO$_4$@CC material were studied by...
scanning electron microscopy (SEM) and powder X-ray diffraction (XRD). As shown in Figure 1A, the surface morphology of the carbon cloth fiber is smooth before growth. Figure 1B,C shows the morphologies of CoMoO$_4$@CC materials under different magnifications. From the low-magnification image (Figure 1B), it can be seen that the carbon fibers are still woven together after growing CoMoO$_4$ nanosheets, and each carbon fiber is evenly covered by CoMoO$_4$ nanoflakes. Further observations show that there are a large number of densely undulating, highly ordered nanosheets on the surface of the carbon cloth. These nanosheets grow vertically to the fiber surface and are cross-linked (Figure 1C) to form a petal-like 3D nanoarray structure. This petal-like structure has a large specific surface area, which is beneficial to improve the ion diffusion rate. Moreover, this structure can also provide abundant active sites, which is beneficial to increase the specific capacitance. Figure 1D shows the XRD result of CoMoO$_4$. Cobalt molybdate has two crystal forms ($\alpha$ and $\beta$ forms). $\alpha$-CoMoO$_4$ can transform to $\beta$-CoMoO$_4$ when the temperature is between 330 and 410 °C.$^{23}$ In this work, the CoMoO$_4$ synthesized by a hydrothermal method has an $\alpha$ form. In the subsequent annealing process, a part of $\alpha$-CoMoO$_4$ transforms to $\beta$-CoMoO$_4$. According to XRD data, $\alpha$-CoMoO$_4$ may not transform completely due to the low annealing temperature and short annealing time. The peak at $2\theta = 33.0^\circ$ in the XRD patterns of CoMoO$_4$ is attributed to $\alpha$-CoMoO$_4$ (JCPDS card number 25-1434). The diffraction peak at $2\theta = 26.4^\circ$ corresponds to the reflection of the (002) plane of $\beta$-CoMoO$_4$ (JCPDS card number 21-0868).$^{24}$ The spectrum of CoMoO$_4$ shows wide and weak diffraction peaks, indicating that the obtained CoMoO$_4$ sample has low crystallinity. As previously reported in the literature, the poor crystallizability of the material plays a vital role in

---

**Figure 1.** SEM images: (A) bare CC, (B) CoMoO$_4$@CC material, and (C) CoMoO$_4$@CC material enlarged view; (D) CoMoO$_4$@CC XRD image.

**Figure 2.** SEM image of CoMoO$_4$@CC and the corresponding element mapping for C, O, Co, and Mo.
improving the electrochemical performance of supercapacitors.25

The X-ray energy spectrum (EDX) mapping of the SEM spectrum (Figure 2) further confirmed that CoMoO4@CC is mainly composed of Co, Mo, C, and O elements, which are uniformly distributed throughout the carbon cloth nanowire structure. The above results confirm that CoMoO4 is uniformly grown and covered on the carbon cloth surface.

To research the chemical composition and valence state of CoMoO4@CC, XPS analysis was performed. As shown in Figure 3A, C, O, Mo, and Co elements were detected in the XPS spectrum. The Co 2p spectrum in Figure 3B shows two main peaks, which are the characteristic peaks of Co2+ in CoMoO4@CC. The binding energy of Co 2p3/2 is 781.3 eV and the binding energy of Co 2p1/2 is 797.1 eV, indicating the oxidation state of Co2+.26,27 Figure 3C shows the Mo 3d spectrum; the peaks are located at 231.7 and 234.8 eV, which belong to Mo 3d5/2 and Mo 3d3/2, respectively. They are also the characteristic peaks of Mo6+ in CoMoO4@CC, indicating the oxidation state of Mo6+.28,29 Figure 3D shows the XPS spectrum of O 1s. The peaks at 532.5 and 531.2 eV correspond to metal–oxygen bonds and oxygen defects, respectively.29 These results indicate that CoMoO4@CC has been successfully prepared, which is consistent with the results of XRD and SEM-EDS.

2.2. Electrochemical Performance. A standard three-electrode system was used to conduct electrochemical tests on carbon cloth and CoMoO4@CC electrode materials in 3 M KOH solution. Figure 4A shows the cyclic voltammetry (CV) curve of bare carbon cloth and CoMoO4@CC electrode at a scan rate of 20 mV/s. Obviously, the current density and the surrounding area of pure CC are much lower than those of the CoMoO4@CC electrode, which indicates that the capacitance of carbon cloth is almost negligible. For the CV curve of the CoMoO4@CC electrode, it can be seen that there are obvious redox peaks in the range of −0.1 to 0.6 V. The redox peak originates from the faradic reaction related to the OH− ion-mediated Co3+/Co2+ ion pair in the alkaline electrolyte.
suggesting that its capacitance is provided by the faradic redox reaction.30

To further understand the charge and ion transfer mechanism of this material, electrochemical impedance spectroscopy (EIS) was used. The Nyquist curves of bare carbon cloth and CoMoO₄@CC electrode material are shown in Figure 4B. In the high-frequency region, the diameter of the semicircle represents the charge transfer resistance (R<sub>ct</sub>), and the real-axis intercept represents the equivalent series resistance (ESR).31 Both carbon cloth and CoMoO₄@CC electrode materials show an inconspicuous semicircle and a small real-axis intercept, which means that the charge transfer resistance and equivalent series resistance are low. It can be seen from the illustration that the x-axis intercept of the CoMoO₄@CC electrode material is smaller than that of bare carbon cloth, which indicates the higher electron conductivity of the CoMoO₄@CC electrode. In the low-frequency region, the linear slope of the CoMoO₄@CC electrode is closer to the y axis than that of the carbon cloth electrode, which indicates that the electrode has a faster ion diffusion rate. These excellent properties can be ascribed to the petal-like structure of the CoMoO₄ clusters. This structure can increase the contact area between the electrolyte and the active material, shorten the ion transport channel, and improve the charge transport efficiency.

Figure 5A displays the galvanostatic charge–discharge (GCD) curves of carbon cloth and CoMoO₄@CC electrodes at a current density of 1 A/g. Obviously, the discharge time of bare CC is much shorter than that of CoMoO₄@CC electrode, which indicates that the specific capacitance of pure CC is almost negligible. This result is consistent with its CV test results. Figure 5B shows the GCD curves of the CoMoO₄@CC electrode at current densities ranging from 1 to 10 A/g. The GCD curves at all current densities are nonlinear and well symmetrical, indicating that most of the capacitance of this material is pseudocapacitance. Moreover, the corresponding Coulomb efficiency of CoMoO₄@CC can reach 96.7%, indicating excellent reversibility. The specific capacitance values are calculated according to the discharge curve, and the results are shown in Figure 5C. The value of Cs decreases with the increase in the current density. This is because less ions can reach the electrode surface as the current density increases.32 When the current densities are 1, 2, 3, 5, and 10 A/g, the specific capacitances of CoMoO₄@CC are 664, 572, 480, 466, and 426 F/g, respectively. Even if the current density is increased by 10 times, it can still maintain a high specific capacitance. Figure 5D shows the results of the cycle stability test at a current density of 10 A/g. The capacity retention rate can reach 84% after 1000 cycles. The sharp decrease in the specific capacitance in the initial 400 charge/discharge cycles can be explain as the presence of the electrochemically active but unstable oxygen groups.33 The illustration in Figure 5D shows the comparison of the CV curve between the first cycle and the 1000th cycle. It can be seen that the CV curve is almost unchanged after 1000 cycles. These data indicate that the CoMoO₄@CC material has an excellent cyclic stability.

2.3. Symmetrical Supercapacitor Performance Test. To further investigate the actual performance of CoMoO₄@CC, we assembled a symmetrical supercapacitor with two identical CoMoO₄@CC electrodes. Figure 6A shows the CV curve of the CoMoO₄@CC∥CoMoO₄@CC supercapacitor at a scan rate of 10 mV/s in different voltage windows. The supercapacitor shows a typical quasi-rectangular CV curve in the voltage range of 0–1.2 V, which indicates that it has good cycle reversibility in a broad voltage window of 0–1.2 V. The GCD curves of the CoMoO₄@CC∥CoMoO₄@CC supercapacitor are shown in Figure 6B. The shape of the GCD
curves is similar to an isosceles triangle, confirming that the device has good reversibility in the entire potential range.

According to the discharge time and total mass of the electrode materials, the specific capacitance at different current densities is calculated and recorded in Figure 6C. At a current density of 1 A/g, the specific capacitance of the device is 67.5 F/g. When the current density is increased to 10 A/g, the capacitance is 46 F/g, which retains 68.1% of its initial value. Figure 6D shows the Ragone plot of the supercapacitor of this work and other Mo-based supercapacitors. It is worth noting that the maximum energy density of the CoMoO$_4$@CC∥CoMoO$_4$@CC device can reach 27 Wh/kg when the power density is 600 W/kg, superior to those of other reported materials for supercapacitors, for instance, MoO$_3$/Ag (15.51 Wh/kg, 348.97 W/kg), $\alpha$-MoO$_3$ (30 F/g, 80% retention, 100 cycle number), MoS$_2$/C (71 F/g, 81% retention, 2000 cycle number), CoMoO$_4$/C (10.94 F/g, 99.8% retention, 200 cycle number), CoMoO$_4$@CC (664 F/g, 84% retention, 1000 cycle number).

3. EXPERIMENTAL SECTION

3.1. Chemicals. Ammonium molybdate ((NH$_4$)$_6$Mo$_7$O$_{24}$·4H$_2$O, AR), cobalt chloride (CoCl$_2$·6H$_2$O), nitric acid (HNO$_3$, 65%), sulfuric acid (H$_2$SO$_4$, AR), and polyvinylpyrrolidone (PVP, AR grade) were purchased from Aladdin Co., Ltd. (China). Urea (CO(NH$_2$)$_2$) and ethanol (C$_2$H$_5$OH) were purchased from Sinopharm Reagent. Potassium hydroxide (KOH) was purchased from Xilong Science, and carbon cloth (CC) was purchased from Suzhou Yilongsheng Energy Technology Co., Ltd. Deionized (DI) water was used throughout the experiment. All reagents are analytically pure and no further purification is required.

3.2. Experimental Steps.

3.2.1. Experimental Plan Design. Using CoCl$_2$·6H$_2$O and (NH$_4$)$_6$Mo$_7$O$_{24}$ as raw materials and carbon cloth as the substrate, the CoMoO$_4$@CC material was synthesized through the hydrothermal reaction and annealing process.

3.2.2. Carbon Cloth Pretreatment. A mixed solution of 10% nitric acid and 10% sulfuric acid (3 nitrate:1 sulfur) was used to pretreat the carbon cloth at 120 °C for 3 h to improve the hydrophilicity. Then, it was washed with acetone, ethanol, and deionized water ultrasonically for 30 min to remove the grease and dirt on the surface. Finally, it was vacuum-dried at 60 °C for 12 h.

3.2.3. Preparation of CoMoO$_4$@CC. Dissolve 720 mg of CoCl$_2$·6H$_2$O, 520 mg of (NH$_4$)$_6$Mo$_7$O$_{24}$, 480 mg of CO(NH$_2$)$_2$, and 100 mg of PVP in 40 mL of deionized water. Table 1. Comparison Data of the Performance of the CoMoO$_4$@CC Electrode Material with References

| Electrode material | Current density | Specific capacitance | Capacity retention | Cycle number |
|--------------------|----------------|---------------------|--------------------|--------------|
| MoS$_2$/C          | 0.2 A g$^{-1}$ | 201 F g$^{-1}$     | 94.1%              | 1000         |
| MoO$_3$/Ag         | 1 A g$^{-1}$  | 225 F g$^{-1}$     | 71.1%              | 10,000       |
| $\alpha$-MoO$_3$   | 5 mV s$^{-1}$ | 30 F g$^{-1}$      | 80%                | 100          |
| $\alpha$-MoO$_3$   | 0.6 mA cm$^{-2}$ | 73.6 F g$^{-1}$ | 94.7%              | 2000         |
| CoMoO$_4$/PVP      | 2 mA cm$^{-2}$ | 71 F g$^{-1}$      | 81%                | 2000         |
| CoMoO$_4$/C        | 1 A g$^{-1}$  | 109.4 F g$^{-1}$   | 99.8%              | 200          |
| CoMoO$_4$@CC (this work) | 1 A g$^{-1}$ | 664 F g$^{-1}$   | 84%                | 1000         |

Table 1. Comparison Data of the Performance of the CoMoO$_4$@CC Electrode Material with References

Figure 6. Electrochemical performance of the CoMoO$_4$@CC∥CoMoO$_4$@CC symmetrical supercapacitor in 1 M Na$_2$SO$_4$ neutral electrolyte: (A) CV curves of different voltage ranges when the sweep speed is 10 mV/s; (B) GCD curves at various current densities; (C) specific capacitance under different current densities; (D) Ragone curve diagram.
After stirring for 20 min, the solution was transferred to a 100 mL autoclave and the pretreated carbon cloth (2 × 2 cm) was added to the kettle, then sealed, and hydrothermally reacted at 160 °C for 12 h. After the autoclave was cooled to room temperature, the samples were collected, washed several times with water and ethanol, and dried at 60 °C. Finally, the CoMoO₄@CC sample was heated to 350 °C at a heating rate of 5 °C min⁻¹ and annealed in a N₂ environment for 3 h.

### 3.3. Materials Characterization
Field emission scanning electron microscopy (FESEM) was performed on a Nova Nano SEM 450 scanning electron microscope. The XPS measurement was performed on a Thermo Fisher X-ray photoelectron spectrometer equipped with Al radiation as a probe (Kr, radiation), the chamber pressure was 5 × 10⁻⁹ Torr, and the diameter of the analysis spot was 400 μm. The crystal structure of the sample was studied by XRD (D8 Advance, Germany).

### 3.4. Electrochemical Tests
Whole electrochemical tests were carried out on a CH1660E workstation (Shanghai Chenhua) in 3 M KOH aqueous solution. A three-electrode system was used to investigate the electrochemical performance of the samples. A saturated calomel electrode and Pt wire served as the reference and counter electrodes, respectively.

A two-electrode system was used to study the electrochemical properties of the symmetric supercapacitor constructed by assembling two similar CoMoO₄@CC electrodes. The specific capacitances (Cs) of the individual electrode and the two-electrode cell were calculated from the galvanostatic discharge curves by eq 1. The specific energy density (Ec) and specific power density (Pc) were calculated according to eqs 2 and 3, respectively:

\[
Cs = \frac{I \Delta t}{m \Delta U}
\]  
(1)

\[
Ec = \frac{Cs \Delta U^2}{2 \times 3.6}
\]  
(2)

\[
Pc = \frac{Ec \times 3600}{\Delta t}
\]  
(3)

where I is the discharge current, Δt is the discharge time, m is the mass of the active material in a three-electrode system in a two-electrode cell, m is the total active mass of both electrodes, and ΔU is the voltage window.

### 4. CONCLUSIONS
In this work, the petal-like CoMoO₄ clusters have been directly synthesized on the carbon cloth substrate by a simple hydrothermal method. Electrochemical studies show that the CoMoO₄@CC electrode material has an excellent pseudocapacitance performance. The CoMoO₄@CC material displays a high specific capacitance (664 F/g at a current density of 1 A/g) and an excellent cycle stability (capacitance remains at 84% after 1000 cycles). The assembled symmetrical supercapacitor has an energy density of 27 Wh/kg when the power density is 600 W/kg. Moreover, the energy density can still remain at 18.4 Wh/kg at a high power density of 6022 W/kg. This convenient method provides an effective way to enhance the electrochemical performance of supercapacitors and shows broad application prospects in future energy storage systems.

---

**AUTHOR INFORMATION**

**Corresponding Authors**

Dejuan Huang — School of Chemistry, Biology, and Materials Science, East China University of Technology, Nanchang 330013 Jiangxi, China; Email: djhuang@ecut.edu.cn

Tianxiang Jin — School of Chemistry, Biology, and Materials Science, East China University of Technology, Nanchang 330013 Jiangxi, China; Email: tianxiangjin1988@gmail.com

**Authors**

Chuanhong Chen — School of Chemistry, Biology, and Materials Science, East China University of Technology, Nanchang 330013 Jiangxi, China

Hangchun Deng — School of Chemistry, Biology, and Materials Science, East China University of Technology, Nanchang 330013 Jiangxi, China

Chongshi Wang — College of Engineering, Department of Civil, Architectural & Environmental Engineering, Drexel University, Philadelphia, Pennsylvania 19104, United States

Wenqing Luo — School of Chemistry, Biology, and Materials Science, East China University of Technology, Nanchang 330013 Jiangxi, China

Complete contact information is available at: https://pubs.acs.org/10.1021/acsomega.1c02166

**Notes**

The authors declare no competing financial interest.

**ACKNOWLEDGMENTS**

This work was supported by the Foundation of Jiangxi Educational Committee under grant no. GJJ160565, East China University of Technology Research Foundation for Advanced Talents no. DHBK2016110, and National Natural Science Foundation of China (41867063 and 41562021).

**REFERENCES**

1. Zhu, M.; Shao, Q.; Pi, Y.; Guo, J.; Huang, B.; Qian, Y.; Huang, X. Ultrathin Vein-Like Iridium–Tin Nanowires with Abundant Oxidized Tin as High-Performance Ethanol Oxidation Electrocatalysts. Small 2017, 13 (36), 1701295.

2. Zhu, M.; Shao, Q.; Qian, Y.; Huang, X. Superior Overall Water Splitting Electrocatalysis in Acidic Conditions Enabled by Bimetallic Ir-Ag Nanotubes. Nano Energy 2019, 56, 330–337.

3. Zhu, T.; Ding, J.; Shao, Q.; Qian, Y.; Huang, X.; Se-Codoped MoS₂ Nanosheets as Accelerated Electrocatalysts for Hydrogen Evolution. ChemCatChem 2019, 11 (2), 689–692.

4. Chen, J.; Lin, C.; Zhang, M.; Jin, T.; Qian, Y. Constructing Nitrogen, Selenium Co-Doped Graphene Aerogel Electrode Materials for Synergistically Enhanced Capacitive Performance. ChemElectroChem 2020, 7 (15), 3311–3318.

5. Yang, L.; Lu, X.; Wang, S.; Wang, J.; Guan, X.; Guan, X.; Wang, G. Designed synthesis of nickel–cobalt-based electrode materials for high-performance solid-state hybrid supercapacitors. Nanoscale 2020, 12, 1921–1938.

6. Gao, M.; Wang, W.-K.; Rong, Q.; Jiang, J.; Zhang, Y.-J.; Yu, H.-Q. Porous ZnO-coated Co₃O₄ nanorod as a high-energy-density supercapacitor material. ACS Appl. Mater. Inter. 2018, 10, 23163–23173.

7. Qu, G.; Sun, P.; Xiang, G.; Yin, J.; Wei, Q.; Wang, C.; Xu, X. Moss-like nickel-cobalt phosphate nanostructures for highly flexible all-solid-state hybrid supercapacitors with excellent electrochemical performances. Appl. Mater. Today 2020, 20, 100713.

8. Yang, L.; Yang, Y.; Wang, S.; Guan, X.; Guan, X.; Wang, G. Multi-Heteroatom-Doped Carbon Materials for Solid-State Hybrid...
Supercapacitors with a Superhigh Cycling Performance. Energy Fuels 2020, 34, 5032–5043.

(9) Wang, J.; Yang, L.; Fu, Y.; Yin, P.; Guan, X.; Wang, G. Delicate control of crystallographic Cu2O derived Ni–Co amorphous double hydroxide nanocages for high-performance hybrid supercapacitors: an experimental and computational investigation. Nanoscale 2021, 13, 8562–8574.

(10) Fu, W.; Wang, Y.; Han, W.; Zhang, Z.; Zha, H.; Xia, E. Construction of hierarchical ZnCo2O4 @ NiCo2S2 (OH)4 core/shell nanowire arrays for high-performance supercapacitors. J. Mater. Chem. A 2016, 4, 173–182.

(11) Zhou, M.; Li, F.; Shen, X.; Xia, W.; He, H.; Zeng, X. One-pot construction of three dimensional CoMoO4/CoO hybrid nanostructures and their application in supercapacitors. J. Mater. Chem. A 2015, 3, 21201–21210.

(12) Hall, P.; Mirzaeian, M.; Fletcher, S.; Sillars, F. B.; Rennie, A. J. R.; Shitta-Bey, G. O.; Wilson, G.; Cruden, A.; Carter, R. Energy storage in electrochemical capacitors: designing functional materials to improve performance. Energy Environ. Sci. 2010, 3, 1238–1251.

(13) Rodrigues, J.; Chaturvedi, S.; Hansen, J.; Albornoz, A.; Brito, J. L. Electronic properties and phase transformations in CoMoO4 and NiMoO4: XANES and time-resolved synchrotron XRD studies. J. Phys. Chem. B 1998, 102, 1347–1355.

(14) Cai, D.; Wang, D.; Liu, B.; Wang, Y.; Liu, Y.; Wang, L.; Li, H.; Huang, H.; Li, Q.; Wang, T. Comparison of the electrochemical performance of NiMoO4 nanorods and hierarchical nanospheres for supercapacitor applications. ACS Appl. Mater. Inter. 2013, 5, 12905–12910.

(15) Peng, S.; Li, L.; Wu, H. B.; Madhavi, S.; Lou, X. W. D. Controlled growth of NiMoO4 nanosheet and nanorod arrays on various conductive substrates as advanced electrodes for asymmetric supercapacitors. Adv. Energy Mater. 2015, 5, 1401172.

(16) Cao, Y.; An, L.; Liao, L.; Liu, X.; Ji, T.; Zou, R.; Yang, J.; Qin, Z.; Hu, J. Hierarchical core/shell structures of ZnO nanorods@CoMoO4 nanolaminate used as a high-performance electrode for supercapacitors. RSC Adv. 2016, 6, 3020–3024.

(17) Yu, X.; Lu, B.; Xu, Z. Super long-life supercapacitors based on the construction of nanohoneycomb-like strongly coupled CoMoO4 3D graphene hybrid electrodes. Adv. Mater. 2014, 26, 1044–1051.

(18) Wu, H.; Lou, Z.; Yang, H.; Shen, G. A flexible spiral-type supercapacitor based on ZnCo2O4 nanorod electrodes. Nanoscale 2015, 7, 1921.

(19) Liu, B.; Liu, B.; Wang, Q.; Wang, X.; Xiang, Q.; Chen, D.; Shen, G. New energy storage option: toward ZnCo2O4 nanorods/nickel foam architectures for high-performance supercapacitors. ACS Appl. Mater. Inter. 2013, 5, 10011–10017.

(20) Wang, G.; Zhang, L.; Zhang, J. A review of electrode materials for electrochemical supercapacitors. Chem. Soc. Rev. 2012, 41, 797–828.

(21) Liu, J.; Jiang, J.; Cheng, C.; Li, H.; Zhang, J.; Gong, H.; Fan, H. J. CoO4nanowire@ MnO2 ultrathin nanosheet core/shell arrays: a new class of high-performance pseudocapacitive materials. Adv. Mater. 2011, 23, 2076–2081.

(22) Ling, J.; Zou, H.; Yang, W.; Chen, W.; Lei, K.; Chen, S. Facile fabrication of polyaniline/molybdenum trioxide/activated carbon cloth composite for supercapacitors. J. Energy Storage 2018, 20, 92–100.

(23) Rico, J. L.; Ávalos-Borja, M.; Barrera, A.; Hargreaves, J. S. J. Template-free synthesis of CoMoO4 rods and their characterization. Mater. Res. Bull. 2013, 48, 4614–4617.

(24) Xu, X.; Shen, J.; Li, N.; Ye, M. Microwave-assisted synthesis of graphine/CoMoO4 nanocomposites with enhanced supercapacitor performance. J. Alloys Compd. 2014, 616, 58–65.

(25) Liu, M.-C.; Kong, L.-B.; Kang, L.; Li, X.; Walsh, F. C.; Xing, M.; Lu, C.; Ma, X.-J.; Luo, Y. Synthesis and characterization of MnV2O7 (M = Ni or Co) based nanostructures: a new family of high performance pseudocapacitive materials. J. Mater. Chem. A 2014, 2, 4919–4926.

(26) Nti, F.; Anang, D.; Han, J. I. Facilely synthesized NiMoO4/CoMoO4 nanorods as electrode material for high performance supercapacitor. J. Alloys Compd. 2018, 742, 342–350.

(27) Li, Z.; Zhao, D.; Xu, C.; Ning, J.; Zhong, Y.; Zhang, Z.; Wang, Y.; Hu, Y. Reduced CoNi5S2 nanosheets with enhanced conductivity for high-performance supercapacitors. Electrochim. Acta 2018, 278, 33–41.

(28) Ghosh, D.; Giri, S.; Das, C. K. Synthesis, characterization and electrochemical performance of graphene decorated with 1D NiMoO4–NiO nanorods. Nanoscale 2013, 5, 10428–10437.

(29) Chen, C.; Wang, S.; Luo, X.; Gao, W.; Huang, G.; Zeng, Y.; Zhu, Z. Reduced ZnCo2O4@NiMoO4 nanotubes heterostructure electrodes with modulating oxygen vacancies for enhanced aqueous asymmetric supercapacitors. J. Power Sources 2019, 409, 112–122.

(30) Senthilkumar, B.; Meyrick, D.; Lee, T.-S.; Selvan, R. K. Synthesis and improved electrochemical performances of nano β-NiMoO4 nanocomposites for asymmetric supercapacitors. RSC Adv. 2013, 3, 16542–16548.

(31) Li, H.; Zhang, X.; Ding, R.; Qi, L.; Wang, H. Facile synthesis of mesoporous MnO2 microspheres for high performance AC/MnO2 aqueous hybrid supercapacitors. Electrochim. Acta 2013, 108, 497–505.

(32) Nasini, U.; Bairi, V.; Ramasahayam, S.; Bourdo, S.; Viswanathan, T.; Shakh, A. U. Phosphorous and nitrogen dual heteroatom doped mesoporous carbon synthesized via microwave method for supercapacitor application. J. Power Sources 2014, 250, 257–264.

(33) Wang, C.; Zhou, Y.; Sun, L.; Wan, P.; Zhang, X.; Qiu, J. Sustainable synthesis of phosphorous- and nitrogen-doped porous carbons with tunable surface properties for supercapacitors. J. Power Sources 2013, 239, 81–88.

(34) Fan, L.-Q.; Liu, G.-J.; Zhang, C.-Y.; Wu, J.-H.; Wei, Y.-L. Facile one-step hydrothermal preparation of molybdenum disulfide/carbon composite for use in supercapacitor. Int. J. Hydrogen Energy 2015, 40, 10150–10157.

(35) Zhou, C.; Wang, Q.; Yan, X.; Wang, J.; Jiang, W.; D. F.; Yuan, X.; Jiang, H.; Zhu, Y. H.; Cheng, X. N. A facile route to synthesize Ag decorated CoMoO4 nanocomposites with symmetric supercapacitor. Cerem. Int. 2020, 46, 15385–15391.

(36) Shakir, I.; Shahid, M.; Yang, H. W.; Kang, D. J. Structural and electrochemical characterization of α-MoO3 nanorod-based electrochemical energy storage devices. Electrochem. Acta 2010, 56, 376–380.

(37) Prakash, N.; Dhananjaya, M.; Narayana, A.; Shaik, D.; Rosaiha, P.; Hussain, O. High performance one dimensional α-MoO3 nanorods for supercapacitor applications. Ceram. Int. 2018, 44, 9967–9975.

(38) Ramkumar, R.; Sundaram, M. A biopolymer gel-decorated cobalt molybdate nanowires: effective grafting polymer cross linked system with an organic acid for better energy storage. New J. Chem. 2016, 40, 2863–2877.

(39) Li, J.; Zhao, C.; Yang, Y.; Li, C.; Hollenkamp, T.; Burke, N.; Hu, Z.; Tendeelo, G. V.; Chen, W. Synthesis of monodispersed CoMoO4 nanoclusters on the ordered mesoporous carbons for environment-friendly supercapacitors. J. Alloy. Compd. 2019, 810, 151841–151849.

(40) Pujari, R. B.; Lokhande, V. C.; Kumbar, V. S.; Chodanker, N. R.; Lokhande, C. D. Hexagonal microrods architecuted MoO3 thin film for supercapacitor application. J. Mater. Sci. 2016, 27, 3312.

(41) Sari, F. N. I.; Ting, J.-M. High performance asymmetric supercapacitor having novel 3D networked polypyrrole nanotube/N-doped graphene negative electrode and core-shell MoO3/PPy supported MoS2 positive electrode. Electrochim. Acta 2019, 320, 134533.

(42) Deng, H.; Huang, J.; Hu, Z.; Chen, X.; Huang, D.; Jin, T. Fabrication of a Three-Dimensionally Networked MoO3/PPy/rGO Composite for a High-Performance Symmetric Supercapacitor. ACS Omega 2021, 6, 9426–9432.