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Cobalt vanadium oxide thin nanoplates:
primary electrochemical capacitor
applicationYoujuan Zhang¹, Yuanying Liu¹, Jing Chen¹, Qifei Guo¹, Ting Wang¹ & Huan Pang^{1,2}¹Key Laboratory for Clearer Energy and Functional Materials of Henan Province, College of Chemistry and Chemical Engineering, Anyang Normal University, Anyang, 455000 Henan, P.R. China, ²State Key Laboratory of Coordination Chemistry, Nanjing University, Nanjing, 210093, Jiangsu, P.R. China.Received
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14 July 2014Correspondence and
requests for materials
should be addressed to
H.P. (huanpangchem@
hotmail.com)

Co₃V₂O₈ thin nanoplates are firstly described as a kind of electrode material for supercapacitors. More importantly, from electrochemical measurements, the obtained Co₃V₂O₈ nanoplate electrode shows a good specific capacitance (0.5 A g⁻¹, 739 F g⁻¹) and cycling stability (704 F g⁻¹ retained after 2000 cycles). This study essentially offers a new kind of metal vanadium oxides as electrochemical active material for the development of supercapacitors.

Supercapacitors, an advanced safe electrochemical energy storage device, can hold higher power density, longer cycling life and shorter charge/discharge time^{1–9}. The device stores energy on the basis of either charge accumulation (electric double layer capacitors) or fast reversible Faradaic reactions (pseudocapacitors) on the surface. Considerable efforts have been devoted to the development and characterization of new electrode materials with lower cost and improved performance.

Transition metal oxides can provide higher energy density in comparison to the electrical double layer capacitors with carbon-based active materials. And recently, binary metal oxides have been demonstrated better performance than single-component oxides due to their feasible oxidation states and high electrical conductivity^{8,10–19}. Transition metal vanadium oxides and vanadates have been widely investigated as possible active materials for rechargeable lithium batteries^{20–27}. Very recently, Wang et al have reported the successful preparation of cobalt vanadium oxide (Co₃V₂O₈) nanostructures which exhibit outstanding reversible capacity and excellent rate performance for lithium storage²⁶. However, the exploitation of Co₃V₂O₈ nanostructures for supercapacitors has few developments.

This work describes the first investigation of Co₃V₂O₈ nanoplates for supercapacitors. We provide a facile and hydrothermal method to synthesize Co₃V₂O₈ nanoplates by the modified method of Wang et al²⁶. Interestingly, the obtained Co₃V₂O₈ nanoplate electrode shows a good specific capacitance (0.5 A g⁻¹, 739 F g⁻¹) and cycling stability (704 F g⁻¹ retained after 2000 cycles) for supercapacitors.

Results

XRD patterns of as-prepared samples were performed to reveal the phase of the as-prepared samples in Fig. 1a. All intense peaks shown in Fig. 1a can be well indexed to cubic Co₃V₂O₈ (JCPDS No.16-0675). The peaks are strong, which indicates the good crystallinity of the as-prepared samples. Good crystallinity of as-prepared Co₃V₂O₈ might be good to improve the cycle life of electrodes for its crystal structure is stable and it is not easy to be destroyed during the electrochemical process. The EDS result in Fig. 1b demonstrates that the nanoplates are composed of the elements of Co, O, and V.

The morphology of as-prepared Co₃V₂O₈ is examined by field emission scanning electron microscopy (FESEM), atomic force microscope (AFM) and transmission electron microscopy (TEM). A typical low-magnification FESEM image is shown in Fig. 2a and the morphology of samples is thin nanoplates in 2D microscale with a diameter of 150 ~ 200 nm. The thickness of the thin nanosheet is ~40 nm in Fig. 2b. AFM measurement is further used to confirm the thickness of the single nanoplate in Fig. 2c. It can be seen that the accurate thickness is 42.7 nm in the inset of Fig. 2c, which is consistent with the FESEM result in Fig. 2b. In addition, there are nearly no pores seen from the TEM image in Fig. 2d, e. The selected area electron diffraction (SAED) pattern is shown in the

Figure 1 | a) XRD patterns of as-prepared samples and its corresponding JCPDS card No 16-0675; b) corresponding EDS patterns.

inset of Fig. 2e, which demonstrates the good crystalline nature of the product. The measured distance of neighboring lattice fringes in Fig. 2f is ~ 0.59 nm, corresponding well to the (110) lattice spacing of $\text{Co}_3\text{V}_2\text{O}_8$.

Discussion

The CVs of $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplate electrodes (a mass loading of ~ 5 mg) in 3.0 M KOH at different scan rates ($5 \sim 100$ mV s^{-1}) are shown in Fig. 3a. As is seen in Fig. 3a, the shapes are different from that of electric double-layer capacitance, suggesting that the capacity mainly results from pseudocapacitive capacitance. And it is clear that the Faradaic pseudocapacitive property of $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplates is based on the surface redox mechanism of Co^{2+} to Co^{3+} on the surface. Chronopotentiometry (CP) curves at different current densities are shown in Fig. 3b. Different charge–discharge current densities and times can be clearly seen. As increasing of current densities, short charge–discharge times have been obtained.

The relationship between the specific capacitances calculated by CP curves and current densities is given in Fig. 3c. Based on the curve, $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplate electrodes show a good specific capacitance and reach up to 739 F g^{-1} at the current density of 0.5 A g^{-1} , and 516 F g^{-1} even at 4.0 A g^{-1} . The specific capacitance of $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplates is better than that of our previous cobalt based phosphite nanomaterials, for example, $\text{NH}_4\text{CoPO}_4 \cdot \text{H}_2\text{O}$ microflowers (< 340 F g^{-1} at 1.5 A g^{-1})²⁸, cobalt phosphite microarchitectures (< 312 F g^{-1} at 1.5 A g^{-1})²⁹. However, some materials show larger values than that of ours^{30,31}.

The relationship of the specific capacitance against cycling number of $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplate electrodes is shown in Fig. 3d, which shows its good specific capacitance retention at 0.5 A g^{-1} . After 350 continuous charge–discharge cycles, $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplate electrodes almost retain the same specific capacitance as its initial value. More importantly, $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplate electrodes still retain more than 95.3% of their specific capacitance after 2000 continuous charge–discharge cycles. After the testing of the cycle life, we have measured the morphology and X-ray diffraction of the electrode materials in Supplementary Information (SI) Fig. 1, 2. There are few changes of $\text{Co}_3\text{V}_2\text{O}_8$ thin nanoplates after 2000 charge–discharge cycles in SI Fig. 1, and the good crystallinity of as-prepared $\text{Co}_3\text{V}_2\text{O}_8$ has also been maintained. What's more, the nanoplate structure might offer a stable structure for ion intercalation/extraction, which might improve the cycle life of the electrode.

Figure 2 | a, b) FESEM images of as-prepared $\text{Co}_3\text{V}_2\text{O}_8$; c) Corresponding AFM image; d, e) TEM images (inset-corresponding SAED pattern); f) Corresponding HRTEM image (inset-the large magnification image).

Figure 3 | a) Cyclic voltammetry within a 0.0–0.42 V range at a scan rate 5–100 mV s⁻¹ were performed on the Co₃V₂O₈ thin nanoplate electrode in 3.0 M KOH at room temperature, b) The galvanostatic charge–discharge curves during current densities, 0.5–4.0 A g⁻¹, c) Specific capacitances derived from the discharging curves at the current density of 0.5–4.0 A g⁻¹, and d) Charge/discharge cycling test at 0.5 A g⁻¹.

We have tried to propose the possible reaction mechanism of Co₃V₂O₈ in equation (1). To identify the exact electrical conductivity of electrodes, we have measured electrochemical impedance spectra (EIS) of Co₃V₂O₈ thin nanoplate electrodes at room temperature in the frequency range 0.01–10⁵ Hz at different potentials in SI Fig. 4. The impedance plot at 0.42 V in 3 M KOH exhibits a line along the imaginary axis (*Z''*). At a potential of 0, 0.60 or 0.70 V, the linear characteristic becomes weaker. Thus, the proper potential range, over which electrode can generate desirable capacitance, is below 0.42 V. Fig. 4 shows the EIS of Co₃V₂O₈ thin nanoplate electrodes at room temperature and its calculated curve by the ZSimpWin software. An equivalent circuit used to fit the impedance curve is given in the inset of Fig. 4a, which is similar to the circuit employed for the working electrode of supercapacitors. The EIS data can be fitted by a bulk solution resistance *R_s*, a charge-transfer *R_{ct}* and a pseudocapacitive element *C_p* from the redox process of electrode materials, and a CPE to account for the double-layer capacitance. The charge-transfer resistance *R_{ct}* is calculated by the ZSimpWin software. From the calculated results, *R_{ct}* of Co₃V₂O₈ thin nanoplate electrodes is 5.3 Ω. It clearly demonstrates the reduced charge-transfer resistance of the Co₃V₂O₈ thin nanoplate electrode. The nanoplate surface-interface character might decrease the polarization of the electrode, and thus increase the capacity. What's more, the phase angles for impedance plots of Co₃V₂O₈ thin nanoplate electrodes and its calculated curve by ZSimpWin software are observed in Fig. 4b. These phase angles are near 80° in the low frequencies clearly, which means that the Co₃V₂O₈ thin nanoplates allow ions or electrolyte transfer to occur

quickly. Compared with the charge-transfer resistance of other Co₃V₂O₈ nano/micro structures in SI Fig. 3, 4, and SI Table 1, it is seen that the Co₃V₂O₈ thin nanoplate can reduce the charge-transfer resistance more effectively than other Co₃V₂O₈ nano/micro structures.

In summary, Co₃V₂O₈ thin nanoplates have been successfully synthesized to show the novel nanoplate surface-interface feature, which plays a key role in the ion intercalation/extraction and electrolyte access. The electrochemical investigation of Co₃V₂O₈ thin nanoplates is interesting work, which firstly illustrates Co₃V₂O₈ thin nanoplates can be applied as an electroactive material for supercapacitors. Despite the specific capacitance of Co₃V₂O₈ thin nanoplates are not much larger than some famous materials, it might direct a new generation of nanodevices by exploring the electrochemical characteristics of novel nano/micromaterials.

Methods

In a typical synthesis, 187 mg of NH₄VO₃ was dissolved into 32 mL deionized water at 80 °C. Under severely stirring, 40 mg NaOH was added to the ammonium metavanadate solution. Then, 76 mg Co(OAc)₂·4H₂O was added to the solution under stirring. After continuous stirring for 10 mins, the resulting yellow precursor suspension was transferred into a 40 mL Teflon-lined autoclave and maintained at 120 °C for 12 hours. After reaction, the brown yellow precipitates were obtained. The precipitates were thoroughly washed with distilled water and ethanol to remove ions possibly remaining in the final products, and dried at room temperature in the air.

Characterizations. The morphology of as-prepared samples was observed by a JEOL JSM-6701F field-emission scanning electron microscope (FE-SEM) at an acceleration voltage of 5.0 kV. The phase analyses of the samples were performed by X-ray diffraction (XRD) on a Rigaku-Ultima III with Cu K_α radiation (λ = 1.5418 Å). Transmission electron microscopy (TEM) images were captured on the JEM-2100 instrument microscopy at an acceleration voltage of 200 kV. Atomic force microscopy (AFM) images were measured at CSPM4000 (Benyuan, Beijing).

Electrochemical measurement. Electrochemical study on Co₃V₂O₈ thin nanoplates electrodes was carried out on a CHI 660D electrochemical working station (Shanghai

Figure 4 | The EIS of the electrodes at room temperature and its corresponding calculated curve, in inset of a) The equivalent circuit; and b) The phase angles for impedance plots and its corresponding calculated curve.

Chenhua Instrument, Inc.). All electrochemical performances were carried out respectively in a conventional three-electrode system equipped with a platinum electrode and a saturated calomel electrode (SCE) as counter and reference electrodes. Before electrochemical measurement, we have purged out O_2 from the solution by the inert gas-Ar. The working electrode was made by mixing active materials ($Co_3V_2O_8$ thin nanoplates), acetylene black, and PTFE (polytetrafluoroethylene) at a weight ratio of 80 : 15 : 5, coating on a piece of foamed nickel foam of about 1 cm^2 , and pressed it to be a thin foil at the pressure of 5.0 MPa. The electrolyte was 3.0 M KOH solution. Electrochemical impedance spectroscopy measurements of all the samples were conducted at 0.42 V in the frequency range from 100 kHz to 0.01 Hz with the AC voltage amplitude of 5 mV by using PARSTAT2273.

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Author contributions

Y.J.Z., Y.Y.L., J.C., Q.F.G., T.W. and H.P. conceived and designed the experiments., Y.J.Z., J.C. and H.P. analyzed the measurements. H.P. wrote the manuscript in collaboration with all the authors.

Additional information

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