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Synchronous Defect and Interface Engineering of NiMoO$_4$ Nanowire Arrays for High-Performance Supercapacitors

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Abstract: Developing high-performance electrode materials is in high demand for the development of supercapacitors. Herein, defect and interface engineering has been simultaneously realized in NiMoO$_4$ nanowire arrays (NWAs) using a simple sucrose coating followed by an annealing process. The resultant hierarchical oxygen-deficient NiMoO$_4$@C NWAs (denoted as “NiMoO$_4$-$x$@C”) are grown directly on conductive ferronickel foam substrates. This composite affords direct electrical contact with the substrates and directional electron transport, as well as short ionic diffusion pathways. Furthermore, the coating of the amorphous carbon shell and the introduction of oxygen vacancies effectively enhance the electrical conductivity of NiMoO$_4$. In addition, the coated carbon layer improves the structural stability of the NiMoO$_4$ in the whole charging and discharging process, significantly enhancing the cycling stability of the electrode. Consequently, the NiMoO$_4$-$x$@C electrode delivers a high areal capacitance of 2.24 F cm$^{-2}$ (1720 F g$^{-1}$) at a current density of 1 mA cm$^{-2}$ and superior cycling stability of 84.5% retention after 6000 cycles at 20 mA cm$^{-2}$. Furthermore, an asymmetric super-capacitor device (ASC) has been constructed with NiMoO$_4$-$x$@C as the positive electrode and activated carbon (AC) as the negative electrode. The as-assembled ASC device shows excellent electrochemical performance with a high energy density of 51.6 W h kg$^{-1}$ at a power density of 203.95 W kg$^{-1}$. Moreover, the NiMoO$_4$@C ASC device manifests remarkable cyclability with 84.5% of capacitance retention over 6000 cycles. The results demonstrate that the NiMoO$_4$-$x$@C composite is a promising material for electrochemical energy storage. This work can give new insights on the design and development of novel functional electrode materials via defect and interface engineering through simple yet effective chemical routes.

Keywords: NiMoO$_4$ nanowire arrays; oxygen vacancies; core-shell electrode structure; asymmetric supercapacitors

1. Introduction

Several alternative energy technologies have been under development globally in a great effort to mitigate the energy and environmental challenges faced and in accordance with the current “carbon neutral” policies. Supercapacitors (SCs), also known as electrochemical capacitors (ECs), have been considered as one of the most promising energy storage devices due to their unique characteristics of high power density (>10 kW/kg), fast charging and discharging capability (within a few seconds), long lifespan (over 100,000 cycles), and good operational safety. SCs have been widely applied in some important fields, including smart electric grids, memory back-ups, (hybrid) electric vehicles, and aerospace crafts. Although SCs have the advantages of high power density and very long calendar lives, their further application is still hindered by their limited energy density.
Therefore, it is crucial to develop advanced high-performance SCs with higher energy densities without severely compromising the power density and cyclability.

Recently, asymmetric supercapacitors (ASCs) have been regarded as promising due to their enhanced energy density. ASCs commonly combine pseudocapacitive materials (as positive electrode) and electric double-layer capacitive materials (as negative electrode), making use of the much higher specific capacitance derived from the pseudocapacitive electrode materials and a wider potential window during operation. In this sense, the electrochemical performances of ASCs are largely influenced by the structures and properties of the pseudocapacitive electrode materials. Thus, the development of high energy density ASCs heavily relies on the rational selection and design, as well as delicate fabrication of advanced electrode materials.

The use of transition metal oxides (TMOs) [1], hydroxides [2], sulfides [3], selenides [4], carbides [5], and their composites with conductive carbon and polymers [6], as possible electrode materials for ASCs, has been attempted. Particularly, TMOs, such as NiO [7], MoO$_3$ [8], Co$_3$O$_4$ [9], and MnO$_2$ [10], have received considerable attention due to their low cost, convenience of synthesis, environmental friendliness, and relatively high capacitance. Compared to binary TMOs, ternary TMOs materials containing two different metal cations, such as NiCo$_2$O$_4$ [11], ZnCo$_2$O$_4$ [12,13], CoMoO$_4$ [14], MnMoO$_4$ [15], and NiMoO$_4$ [16], as well as some solid solutions [17], exhibit higher electrochemical activity due to the rich redox reactions stemming from the multiple oxidation states of the transition-metal components. It has been proposed that NiMoO$_4$ has a good reversible capacitance and electrochemical characteristics for SCs/ASCs applications because of the electrochemical active Ni cation and improved electronic conductivity from Mo cation [16,18].

However, the practical application of NiMoO$_4$ electrodes in SCs and ASCs is still hampered by their poor electronic conductivity, insufficient ionic transport and diffusion, and structural instability during long-term cycling [19]. Therefore, the controllable fabrication of NiMoO$_4$ with desired nano- and microstructures and rational structural engineering is highly desired but remains challenging.

Various nanostructured NiMoO$_4$ materials, such as nanosheets and nanorods arrays [20], nanotubes [21], hollow nanorods [22], mesoporous nanospheres [23], nanoparticles, and quantum dots [24], have been designed to boost the electrochemical performances of the NiMoO$_4$ electrodes via increased exposed surface for ion adsorption and insertion, shortened path distances for ion transport and diffusion, and improved electrolyte impregnation and permeation. Specifically, various low-dimensional NiMoO$_4$ nanostructures directly grown on conductive substrates (e.g., Ni/Cu foams [25,26], graphene [27], and carbon substrates [28,29]) are particularly preferred for directional electron transport with reduced charge carrier scattering at grain boundaries and easy integration into flexible devices with some specific applications.

To overcome the poor electronic conductivity of pristine NiMoO$_4$, various NiMoO$_4$/carbon composites have been synthesized by hybridizing NiMoO$_4$ nanostructures with graphene [30,31], carbon nanotubes [32], conducting polymers [33], and porous carbon architectures [34,35] Alternatively, intentional doping of NiMoO$_4$ with several kinds of heteroatoms such as Mn [36,37], P [38], Zn [39], Ce [40], or the creation of oxygen vacancies [14,41–43] in the lattice have recently been reported. In addition, NiMoO$_4$ has also been coupled with other metal oxides [44–49] or sulfides [50–52] to form heterostructure electrodes for supercapacitors with improved electrochemical performances.

However, monotonous strategy sometimes has a limited contribution for the overall electrochemical performance improvement of NiMoO$_4$ materials. In addition, some reported approaches for hybridization or doping of NiMoO$_4$ involve multiple and complex chemical and physical processes that are not economically or environmentally friendly. Thus, the rational design and the design of a NiMoO$_4$-based composite electrode for high-performance supercapacitors remains a challenge.

In this work, we report the simultaneous defect and interface engineering of NiMoO$_4$ nanowires arrays (NWAs) using a simple and effective sucrose coating followed by a
thermal treatment approach. In this process, an amorphous carbon shell was uniformly coated on the NiMoO$_4$ surface, effectively improving the electronic transport and structural integrity of the NiMoO$_4$ during electrochemical cycling. Additionally, oxygen-vacancy defects were incorporated into the NiMoO$_4$ during the carbonization process, further enhancing the electronic conductivity of NiMoO$_4$ and redox activity in the NiMoO$_4$ electrode surface. As expected, the resultant NiMoO$_{4-x}@C$ composite exhibited a higher specific capacitance than that of the pristine NiMoO$_4$ NWAs. Furthermore, an asymmetric supercapacitor (ASC) was assembled with the NiMoO$_{4-x}@C$ as positive electrode and activated carbon (AC) as negative electrode, delivering a remarkably high energy density of 51.6 W h kg$^{-1}$ at a power density of 203 W kg$^{-1}$ and an excellent cycling stability with a retention of 84.5% after 6000 cycles under a high current density of 10 A g$^{-1}$.

2. Results and Discussion

The synthesis route of oxygen-deficient NiMoO$_4$@carbon nanowire arrays (NiMoO$_{4-x}@C$) is schematically shown in Figure 1. The preparation process mainly involved three critical steps. Firstly, a NiMoO$_4$ nanowire arrays (NWAs) precursor (NiMoO$_4$·xH$_2$O NWAs, light green) was directly deposited on a ferronickel foam by a hydrothermal reaction process (Step 1, Figure 1). Secondly, the NiMoO$_4$ NWAs precursor was transferred into NiMoO$_4$ NWAs by annealing in Ar to remove crystal H$_2$O and improve crystallinity (Step 2, Figure 1). Finally, the as-obtained NiMoO$_4$ NWAs were immersed in a sucrose solution, followed by drying and annealing in an Ar atmosphere to fabricate oxygen-deficient NiMoO$_4$@carbon NWAs (NiMoO$_{4-x}@C$) (Step 3, Figure 1).

![Figure 1. Schematic illustration of the synthesis process of oxygen-deficient NiMoO$_4$@carbon nanowire arrays (NWAs) (denoted as “NiMoO$_{4-x}@C$”). Step 1, growth of the NiMoO$_4$ NWAs precursor directly on a ferronickel foam substrate using a hydrothermal process; Step 2, conversion of the NiMoO$_4$ NWAs precursor into the NiMoO$_4$ NWAs via annealing in Ar; Step 3, fabrication of NiMoO$_{4-x}@C$ composite by sucrose coating followed by annealing in Ar.](image)

The crystal structures of NiMoO$_{4-x}@C$ and neat NiMoO$_4$ NWAs samples were characterized by X-ray diffraction (XRD) analysis as depicted in Figure 2. The two strongest diffraction peaks, located at ca. 45° and 52°, were from the ferronickel foam substrate. The NiMoO$_{4-x}@C$ and NiMoO$_4$ NWAs samples showed similar diffraction peak shapes and locations. The diffraction peaks located at 14.3°, 24.0°, 25.4°, 28.9°, 32.6°, 37.1°, 38.7°, 41.3°, and 47.4° corresponded to the (110), (021), (−112), (220), (022), (−113), (−132), (040), and (−204) crystal planes of orthorhombic NiMoO$_4$ (JCPDS card No. 86-0361) [37]. Compared to the NiMoO$_4$ NWAs samples, the NiMoO$_{4-x}@C$ sample exhibited a slightly lower diffraction peak intensity possibly due to the covering of carbon on the NiMoO$_4$ surface as well
as the reduced crystallinity of NiMoO$_4$ with increased structural defects. In addition, no characteristic peaks for carbon phases were noted, suggesting the amorphous nature of the carbon species in the NiMoO$_{4-x}$@C sample. The formation of amorphous carbon could be partially ascribed to the lower low annealing temperature (400 °C herein).

![Raman spectra of pristine NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C samples.](image)

**Figure 3.** Raman spectra of NiMoO$_4$ NWAs (black) and NiMoO$_{4-x}$@C (black) and pristine NiMoO$_4$ NWAs (red) samples. The inset (top left) shows the structural model of NiMoO$_4$ crystal, whereas the green, blue, and red balls represent the Ni, Mo, and O atoms, respectively. The two strongest diffraction peaks in the square regions of dotted line are from the ferronickel foam substrate.

Raman spectra of pristine NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C samples are illustrated in Figure 3. The bands at 961 cm$^{-1}$ and 913 cm$^{-1}$ corresponded to the symmetric and asymmetric stretching modes of Mo=O bonds, while the band at 706 cm$^{-1}$ could be ascribed to the stretching mode of Ni/Mo-O bonds of the orthorhombic α-NiMoO$_4$ phase [53]. In addition, two bands ascribed to the presence of carbon species were identified. The band at around 1360 cm$^{-1}$ could be attributed to the D band from defects and disorders in the amorphous carbon layers, while the other band at around 1590 cm$^{-1}$ was related to the G band related to the vibration of sp$^2$-bonded carbon atoms [54]. This result implied the successful deposition of amorphous carbon layer on the surface of the NiMoO$_4$ NWAs.

![Raman spectra of NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C samples.](image)

**Figure 3.** Raman spectra of NiMoO$_{4-x}$@C (black) and pristine NiMoO$_4$ NWAs (red) samples.

The morphologies of the NiMoO$_4$ samples were firstly observed using scanning electron microscopy (SEM). From Figure 4a, the pristine NiMoO$_4$ NWAs sample was
composed of oriented nanowires (NWs) with a smooth surface. In addition, the NiMoO$_4$ NWAs have relatively uniform diameters of ~300 nm, on average, and lengths of several micrometers. After the coating of the carbon, the surface of the NiMoO$_{4-x}$@C sample became obviously coarse as shown in Figure 4b. The element composition analyses using energy-dispersive X-ray spectra (EDS) analysis indicated the existence of Ni, Mo, O, C, Fe, and Al elements in the NiMoO$_{4-x}$@C sample (Figure S1, Supporting Information). Note that the Fe and Al signals mainly stemmed from the ferronickel foam substrate and the sample holder, respectively.

**Figure 4.** SEM images of (a) neat NiMoO$_4$ NWAs and (b) NiMoO$_{4-x}$@C samples.

The microstructures of the pure NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C samples were further investigated by transmission electron microscope (TEM) and high-resolution TEM (HRTEM) as shown in Figure 5. The TEM image (Figure 5a) revealed that the pure NiMoO$_4$ nanowire had a smooth surface with a diameter of about 200 nm. A selected-area electron diffraction pattern (SAED) pattern (inset of Figure 5a) taken from this nanowire depicted a clear two-dimensional dot pattern, suggesting its single-crystalline structure in nature. Two diffraction spots, as marked by white circles, could be indexed to the (220) and (−222) crystal facets of orthorhombic NiMoO$_4$. From Figure 5b, the crystal plane with a lattice spacing of 2.73 Å in the HRTEM micrograph corresponded to the (−222) planes of NiMoO$_4$ [49,52]. In contrast, the TEM image in Figure 5c indicated that a layer of amorphous carbon film with a thickness of ca. 20–50 nm had been coated on the NiMoO$_4$ nanowire’s surface, confirming the core-shell structure of the NiMoO$_{4-x}$@C composite sample with different brightness contrasts of NiMoO$_4$ and carbon. The deposition of amorphous carbon on the surface of NiMoO$_4$ can be further confirmed by HRTEM micrograph as shown in Figure 5d.

**Figure 5.** TEM images (a,c) and HRTEM micrographs (b,d) of pure NiMoO$_4$ NWAs (a,b) and NiMoO$_{4-x}$@C (c,d) samples. The insets in (a,c) are corresponding SAED patterns taken from a single nanowire.
Next, the chemical composition and valence states of element on the surface of NiMoO₄ NWAs and NiMoO₄₋ₓ@C samples were identified by X-ray photoelectron spectroscopy (XPS, Figure 6). In the high-resolution Ni 2p spectrum of the pristine NiMoO₄ NWAs sample (Figure 6a), two main peaks were observed at binding energies (BEs) of 873.4 eV and 856.3 eV with a spin-orbital splitting energy of 17.1 eV, corresponding to the Ni 2p₁/₂ and Ni 2p₃/₂ of Ni²⁺ in NiMoO₄ lattice [55]. In addition, two satellite peaks with BEs of 877.3 eV and 860.6 eV were noted for Ni²⁺. In Figure 6b, similar peak locations and separations can also be observed in the Ni 2p spectra of NiMoO₄₋ₓ@C sample, indicating the coating of carbon had little effect on the chemical valence states of the Ni component in NiMoO₄. The Mo 6d spectrum of pure NiMoO₄ and NiMoO₄₋ₓ@C samples are shown in Figure 6c,d. Evidently both samples had two strong bands with BEs located at 236.0 eV and 232.9 eV, which could be assigned to Mo 3d₃/₂ and Mo 3d₅/₂ of Mo⁶⁺ cations in the NiMoO₄ lattice [56]. In addition, another pair of doublets was noted for the NiMoO₄₋ₓ@C sample, verifying the existence of Mo⁴⁺ in the NiMoO₄₋ₓ@C composite [41,42,57] possibly produced during the amorphous carbon coating process. From Figure 6e, the O 1s spectrum of the NiMoO₄ NWs sample was deconvoluted into three bands. The band centered at 530.1 eV was assigned to the lattice oxygen with O-Ni/O-Mo bonds, while the bands located at 531.3 eV and 532.9 eV correspond to the O-C and O=C bonds from moisture adsorbed on surface [58]. For the NiMoO₄₋ₓ@C sample, another band could be noted at 532 eV, suggesting the presence of oxygen vacancies (Vₒ) [59] at the NiMoO₄ surface (Figure 6f). The formation of and Mo⁴⁺ and Vₒ can tune the electronic structures and electrochemical properties of the NiMoO₄₋ₓ@C composite sample.

![Figure 6. High-resolution XPS spectra of (a,c,e) pristine NiMoO₄ and (b,d,f) NiMoO₄₋ₓ@C samples; (a,b) Ni 2p, (c,d) Mo 3d, (e,f) O 1s.](image-url)
Then, the effects of pyrolysis temperatures (from 200–800 °C) during the carbon coating of the morphologies and microstructures of the NiMoO$_4$/carbon composites were investigated. It is noted that some aggregates of residual sucrose were observed after annealing at 200 °C (Figure S2a), suggesting the carbonization of sucrose was incomplete under a lower temperature. This SEM result also coincides well with the thermogravimetric (TGA) and the differential scanning calorimetry (DSC) analyses (Figure S3), where the thermal decomposition process of sucrose mainly occurs between 223 and 389 °C. With the increase of annealing temperature, sucrose was decomposed, and the carbonization process occurred accompanied by the release of some gases (e.g., CO, CO$_2$). At a higher temperature, the generated reductive gases (e.g., CO) reacted with NiMoO$_4$ and generated some oxygen vacancies on the NiMoO$_4$ surface via abstracting some surface oxygen atoms. In contrast, well-defined nanowires were obtained for the samples prepared after annealing at 400 and 600 °C, respectively (Figure S2b,c). However, the nanowire structure was destroyed when the pyrolysis temperature was increased to 800 °C (Figure S2d), which might have been caused by the large inner stain in the NiMoO$_4$ NWAs or at the NiMoO$_4$–@C interface. Thus, the standard annealing temperature was chosen as 400 °C.

To evaluate the electrochemical performance of NiMoO$_4$ NWAs and NiMoO$_4$–@C samples, electrochemical measurements were tested by a three-electrode system with 2 M KOH electrolyte (Figure 7). Figure 7a shows the CV curves of NiMoO$_4$ NWAs and NiMoO$_4$–@C samples at a scan rate of 20 mV s$^{-1}$ with a potential window of 0 to 0.7 V. Overall, the NiMoO$_4$–@C sample had a larger integral area than that of the NiMoO$_4$ NWAs sample, indicating a significant increase of capacitance after carbon deposition and introduction of oxygen vacancies. Meanwhile, the CV curves of the two samples exhibited typical oxidation peaks, demonstrating typical pseudocapacitive charge storage characteristics. In addition, the CV curves of four different pyrolysis temperatures of NiMoO$_4$–@C samples are revealed in Figure S4. The sample collected at 400 °C shows the highest capacitance which is consistent with the result of SEM in Figure S2. The sample collected at 800 °C exhibited an unsatisfactory performance due to its collapsed morphology. Figure 7b shows the GCD curves of the two samples. It revealed that the discharge time of the NiMoO$_4$–@C sample was almost twice as much as that of the NiMoO$_4$ NWAs sample at a current density of 1 A cm$^{-2}$. Figure 7c shows the electrochemical impedance spectroscopy (EIS) of NiMoO$_4$ NWAs and NiMoO$_4$–@C. The direct impedance and charge transfer resistance of the NiMoO$_4$–@C sample had a larger integral area than that of the NiMoO$_4$ NWAs sample. The remarkably reduced size of the semicircle for the NiMoO$_4$–@C indicated an improved charge transfer kinetics due to enhanced electrical conductivity provided by the carbon shell and oxygen vacancy defects. In addition, the NiMoO$_4$–@C exhibited the steepest slope in the low-frequency region, clearly indicating the lowest Warburg impedance and, hence, the highest K-ion diffusion capability at the interface between the electrode and electrolyte. Through the AC EIS, we added the corresponding equivalent circuit diagram in Figure 7c.

The true impedance of capacitor can be estimated using the following Equation (1):

$$Z_{\text{real}} = R_{\Omega} + \frac{-j\omega C \left( R_{\text{ct}} + W \cdot j\omega^{1/2} \right)}{R_{\text{ct}} + W \cdot j\omega^{1/2} - j(\omega C)}$$  \hspace{1cm} (1)

Figure 7d shows the capacitance of NiMoO$_4$ NWAs and NiMoO$_4$–@C samples calculated from different current densities. After coating the carbon layer, the capacitance of the NiMoO$_4$–@C sample was greatly increased. The areal capacitance can be calculated as high as 2.24 F cm$^{-2}$ (1720 F g$^{-1}$) at a current density of 1 mA cm$^{-2}$. In contrast, the NiMoO$_4$ NWAs electrode only demonstrated a specific capacitance of 1.206 F cm$^{-2}$ (927 F g$^{-1}$) at the same current density. The cycling performances of NiMoO$_4$ NWAs and NiMoO$_4$–@C samples are presented in Figure 7e. The capacitance retention of NiMoO$_4$–@C is 84.5% at 20 mA cm$^{-2}$ after 6000 cycles, which is considerably better than that of the NiMoO$_4$ NWAs sample (63.1% after 6000 cycles). To illustrate the difference of the cycling process between NiMoO$_4$ NWAs and NiMoO$_4$–@C samples, we also obtained the SEM results after cycling.
as shown in Figure S5. It is evident that the NiMoO$_{4-x}$@C sample still held some nanorod structures under the protection of amorphous carbon shell. Instead, NiMoO$_4$ NWAs were aggregated after 10,000 cycles, with unsatisfactory cycle abilities. In addition, we made a comparison of the C$_s$ and cycling stability of this work with some previously reported NiMoO$_4$-based electrodes materials as summarized in Table S1.

![Figure 7](image_url)

Figure 7. (a) CV curves of the NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C at a scan rate of 20 mV s$^{-1}$. (b) GCD curves and (c) EIS spectra of NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C and its corresponding equivalent fitting circuit. (d) GCD test of NiMoO$_{4-x}$@C at different current densities. (e) Cycle test of NiMoO$_4$ NWAs and NiMoO$_{4-x}$@C at 20 mA cm$^{-2}$ over 6000 cycles.

First-principles density functional theory (DFT) simulations were next adopted to further probe the structure–performance relationship of the NiMoO$_{4-x}$@C composite electrode in supercapacitors. The optimized geometry configurations of pristine and oxygen-deficient NiMoO$_4$ (110) surface slabs are shown in Figure S6. The pristine NiMoO$_4$ (110) surface was flat and composed of fivefold Ni and Mo atoms and twofold O atoms (Figure S6a). The defective NiMoO$_4$ (110) surface can be produced after eliminating one surface O atom, leaving one threefold Ni and Mo atoms nearby (Figure S6b). The resultant oxygen-deficient NiMoO$_4$ (110) plane retains flat. Next, the adsorption behavior of OH group on the pristine NiMoO$_4$ (110) surface was first investigated. As shown in Figure 8, the OH can be adsorbed on the top of either the surface of the Mo atom (Figure 8a,b) or the Ni atom (Figure 8c,d), yielding an adsorption energy ($E_{ads}$) of $-0.50$ and $-5.97$ eV, respectively. Evidently, the adsorption of OH on the Ni site was much stronger than that on the Mo site, which is consistent with the fact that the Ni in NiMoO$_4$ is electrochemically active for pseudocapacitive charge storage process based on Faradic reactions. The chemisorption of OH on Ni and Mo sites of the NiMoO$_4$ (110) surface was further verified by the interfacial charge transfer from the charge density difference contours (Figure S7).
The adsorption of OH adsorbed on the oxygen-deficient NiMoO₄ (110) surface was evaluated. Specifically, the adsorption on the Ni and Mo sites with lower coordination due to the removal of surface O was considered. Interestingly, it is noted that the OH group was preferred to be adsorbed at the vicinity of the oxygen-vacancy position (Figure 9a), leading to an $E_{ads}$ of $-3.63$ eV and a charge transfer at the OH/NiMoO₄ interface (Figure 9b). This result suggests that the presence of surface oxygen vacancies offers more active sites for OH adsorption, concentration, and subsequent redox reactions for enhanced pseudocapacitive charge storage.

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Based on the above experimental data and theoretical simulations, the significantly improved electrochemical performances of the NiMoO₄−x@C composite can be mainly attributed to the following points: (i) the deposition of amorphous carbon shell effectively enhances the electron transport of NiMoO₄ nanowires and charge transfer at NiMoO₄/C.
heterointerface; (ii) the deposited carbon layer also improves the structural integrity of the NiMoO$_4$ nanowire arrays during long-term electrochemical cycling; (iii) the creation of oxygen vacancies in NiMoO$_4$ accompanied by the coating of the carbon further enhances the electronic conductivity of the NiMoO$_4$ electrode and creates more active sites for pseudocapacitive charge storage. Therefore, the synergy of defect and interface engineering of NiMoO$_4$ NWAs realized by carbon deposition effectively improves the overall electrochemical performance of the resultant NiMoO$_{4-x}@C$ composite electrode in supercapacitors.

To further assess the practical application potential of a NiMoO$_{4-x}@C$ sample, an asymmetric supercapacitor device (NiMoO$_{4-x}@C$/AC) was assembled with the NiMoO$_{4-x}@C$ as a positive electrode and activated carbon (AC) as a negative electrode. Before testing the ASC device, we performed the CV measurements of NiMoO$_{4-x}@C$ and AC electrodes in a three-electrode system at a scan rate of 5 mV s$^{-1}$ to estimate the suitable operating voltage range (Figure 10a). The maximum operating voltage of NiMoO$_{4-x}@C$/AC ASC was determined to be 1.6 V. The CV curves of NiMoO$_{4-x}@C$/AC ASC at different scan rates are shown in Figure 10b. All the curves display obvious redox peaks, indicating the main contribution from pseudocapacitance. An increased separation of redox peak position can be noted along with the increase of scan rates due to the increased polarization. Figure 10c shows the GCD curves of the NiMoO$_{4-x}@C$/AC ASC at different current densities while Figure 10d shows the specific capacitance calculated from different current densities. The overall capacitance of the ASC was calculated to be 1.01 F cm$^{-2}$ (156.25 F g$^{-1}$) at 1 mA cm$^{-2}$. Furthermore, the NiMoO$_{4-x}@C$/AC ASC device has demonstrated a good capacitance retention of 83.6% after 6000 cycles at 20 mA cm$^{-2}$ (Figure 10e). As a result, the ASC device can power a yellow LED (inset of Figure 10e), showing its potential in practical applications. In addition, the electrochemical performances of our NiMoO$_{4-x}@C$/AC ASC device are also superior or comparable to some recently reported NiMoO$_4$-based electrode materials for ASCs as summarized in Table S1.

![Figure 10](image-url)Figure 10. Electrochemical performance of the asymmetric NiMoO$_{4-x}@C$/AC supercapacitor. (a) CV curves of activated carbon (AC) and NiMoO$_{4-x}@C$ electrodes from −1.0−0 V and 0−0.6 V, respectively
at 5 m V s\(^{-1}\); (b) CV curves; (c) GCD curves; (d) areal capacitance and specific capacitance; (e) cyclic stability of NiMoO\(_{4-x}\)@C/AC.

3. Conclusions

In summary, synchronous defect and interface engineering was implemented in NiMoO\(_4\) material via the formation of oxygen vacancies and the coating of the carbon on NiMoO\(_4\) nanowire arrays through a simple hydrothermal method paired with sucrose pyrolysis. During this process, an amorphous carbon layer was homogeneously deposited on the surface of NiMoO\(_4\) nanowires and oxygen vacancies were created on the NiMoO\(_4\) surface during the carbonization of sucrose. The deposited carbon layer and formed oxygen vacancies in NiMoO\(_4\) boosted the electronic conductivity of NiMoO\(_4\) nanowires. In addition, the coated carbon layer also improved the structural integrity of the NiMoO\(_4\) electrode during long-term operation in supercapacitors. Consequently, the resultant NiMoO\(_{4-x}\)@C heterostructure electrode achieved a high specific capacitance of 2.24 F cm\(^{-2}\) (1720 F g\(^{-1}\)) and maintained a good capacitance retention of about 83.6% after 6000 cycles at 20 mA cm\(^{-2}\). In addition, the as-assembled NiMoO\(_{4-x}\)@C//activated carbon asymmetric supercapacitor device manifested a high energy density of 51.6 W h kg\(^{-1}\) at a high power density of 203.95 W kg\(^{-1}\), indicating that NiMoO\(_{4-x}\)@C composite is a suitable electrode material for supercapacitor applications. The proposed synergistic defect and interface engineering strategy herein can be extended for the design and development of other novel composite electrode materials for applications in electrochemical energy storage and conversion.

Supplementary Materials: The following supporting information can be downloaded at https://www.mdpi.com/article/10.3390/nano12071094/s1, Figure S1: The EDS analysis of NiMoO\(_{4-x}\)@C composite and NiMoO\(_4\) NWAs, Figure S2: SEM images of the NiMoO\(_{4-x}\)@C composite prepared under different annealing temperature: (a) 200 °C, (b) 400 °C, (c) 600 °C and (d) 800 °C, Figure S3: Thermogravimetric (TG) and differential scanning calorimetry (DSC) analysis of sucrose in Ar atmosphere at a ramping rate of 10 °C min\(^{-1}\), Figure S4: (a) CV and (b) GCD of NiMoO\(_{4-x}\)@C samples prepared under different annealing temperatures, Figure S5: SEM images of the (a) NiMoO\(_{4-x}\)@C and (b) NiMoO\(_4\) NWAs electrodes after 10,000 cycles at 20 mA cm\(^{-2}\), Figure S6: Optimized geometry structures of the (a) pristine NiMoO\(_4\) (110) surface and (b) oxygen-deficient NiMoO\(_4\) (110) surface obtained by removing one surface O atom (green). The grey, light pink and red balls represent the Ni, Mo and O atoms, respectively [60–63], Figure S7: Charge density difference contours of after adsorption of one HO molecule on (a) Mo- and (b) Ni-side of NiMoO\(_4\) (110) surface, respectively. The yellow and cyan colors denote the electron gain and loss, respectively. The isosurface level is 0.001 e bohr\(^{-3}\), Table S1: Brief comparison of electrochemical performance of current work with recently relevant literature [20,22,30,55,64–72].

Author Contributions: Conceptualization, C.Q. and P.W.; methodology, P.W. and X.D.; software, R.Z. and H.-E.W.; validation, P.W., X.D. and R.Z.; formal analysis, P.W., X.D. and C.Q.; investigation, P.W. and X.D.; resources, C.Q., H.-E.W. and Y.L.; data curation, P.W., C.Q. and H.-E.W.; writing—original draft preparation, P.W.; writing—review and editing, C.Q., T.Z. and H.-E.W.; visualization, P.W. and R.Z.; supervision, C.Q. and H.-E.W.; project administration, C.Q. and Y.L.; funding acquisition, C.Q. and H.-E.W. All authors have read and agreed to the published version of the manuscript.

Funding: This work is supported by the National Natural Science Foundation of China (Grant No. 52162037) and Yunnan Fundamental Research Projects (grant No. 2019FD113 & 202101AT070120).

Data Availability Statement: All data included in this study are available upon request by contact with the corresponding author.

Conflicts of Interest: The authors declare no conflict of interest.
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