Design of Hierarchical NiCo$_2$O$_4$ Nanocages with Excellent Electrocatalytic Dynamic for Enhanced Methanol Oxidation

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Abstract: Although sheet-like materials have good electrochemical properties, they still suffer from agglomeration problems during the electrocatalytic process. Integrating two-dimensional building blocks into a hollow cage-like structure is considered as an effective way to prevent agglomeration. In this work, the hierarchical NiCo$_2$O$_4$ nanocages were successfully synthesized via coordinated etching and precipitation method combined with a post-annealing process. The nanocages are constructed through the interaction of two-dimensional NiCo$_2$O$_4$ nanosheets, forming a three-dimensional hollow hierarchical architecture. The three-dimensional supporting cavity effectively prevents the aggregation of NiCo$_2$O$_4$ nanosheets and the hollow porous feature provides amounts of channels for mass transport and electron transfer. As an electrocatalytic electrode for methanol, the NiCo$_2$O$_4$ nanocages-modified glassy carbon electrode exhibits a lower overpotential of 0.29 V than those of NiO nanocages (0.38 V) and Co$_3$O$_4$ nanocages (0.34 V) modified glassy carbon electrodes. The low overpotential is attributed to the prominent electrocatalytic dynamic issued from the three-dimensional hollow porous architecture and two-dimensional hierarchical feature of NiCo$_2$O$_4$ building blocks. Furthermore, the hollow porous structure provides sufficient interspace for accommodation of structural strain and volume change, leading to improved cycling stability. The NiCo$_2$O$_4$ nanocages-modified glassy carbon electrode still maintains 80% of its original value after 1000 consecutive cycles. The results demonstrate that the NiCo$_2$O$_4$ nanocages could have potential applications in the field of direct methanol fuel cells due to the synergy between two-dimensional hierarchical feature and three-dimensional hollow structure.

Keywords: hierarchical hollow nanocages; NiCo$_2$O$_4$; coordinated etching and precipitation; methanol oxidation; fuel cell

1. Introduction

The ever-worsening energy and global warming issues have triggered significant research efforts in the design and development of advanced energy devices. Direct methanol fuel cells (DMFCs) have exhibited great commercialization potential credited to high energy density, low cost, easy storage and low pollutant emissions [1,2]. Generally, the performance of DMFCs was mainly related to the activity of methanol electrocatalyst [3]. Traditionally, platinum group precious metals (Pt, Ru, and Pd, etc.) were always employed as electrocatalysts for methanol. Although high electrocatalytic activity was achieved, the precious metals still suffer from high cost and low working stability [4–6]. In this regard, the design of Pt-free catalysts was considered as the best alternative to solve the problems.
Transition metal oxides (TMOs) were recognized as ideal substitutions for noble metals due to their high-active redox sites, low cost and high physicochemical stability. Over the last decade, significant efforts on TMOs have been made to obtain high-performance Pt-free electrocatalysts for methanol [7]. Generally, the nanomaterials in conventional forms of aggregated particles generally have no significant advantages in electrocatalysis. Inspired by kinetics, quantities of TMOs with different microstructures were constructed to improveelectrocatalytic kinetics and high electrocatalytic activity was obtained. Thereinto, two-dimensional (2D) nanosheets were demonstrated as ideal structure in electrocatalysis due to the unique physicochemical properties issued from high structural and morphologic anisotropies [8]. However, agglomeration of 2D nanosheets was easy to occur in electrocatalytic reactions because of the large lateral specific surface areas, leading to the decrease of active sites and diffusion channels.

Integrating amounts of 2D nanosheets into three-dimensional (3D) hierarchical nanocages provided an efficient way to obtain highly active structures. The hierarchical nanocages effectively prevented the aggregation of 2D building blocks and afforded large specific surface areas, which provided sufficient active sites for electrooxidation of methanol [9]. Meanwhile, the pores formed by the interaction of nanosheets not only provided diffusion channels for methanol and intermediate products, but also relieved the volume change and structural strain during electrocatalysis, resulting in excellent stability [10]. Further, the 2D feature of building blocks and porous thin shell of hierarchical nanocages accelerated both the collected and transfer efficiency of catalytic electrons during electrocatalysis, leading to high electrocatalytic activity. Therefore, highly active and stable methanol electrocatalysts can be acquired through the design of hierarchical porous hollow nanocages.

NiCo$_2$O$_4$ possesses bimetallic active sites (Co$^{2+}$/Co$^{3+}$ and Ni$^{2+}$/Ni$^{3+}$) and excellent conductivity, exhibiting potential applications in the field of methanol oxidization [11]. In this report, NiCo$_2$O$_4$ nanocages (NCs) were prepared by coordinated etching and precipitation (CEP) method combined with a post-annealing process. As an electrode for methanol electrooxidation, NiCo$_2$O$_4$ NCs-modified glassy carbon electrode (GCE) exhibited low overpotential, high current density and excellent stability.

2. Materials and Methods

2.1. Reagents

NaOH, CuCl$_2$·2H$_2$O, NiCl$_2$·6H$_2$O, CoCl$_2$·6H$_2$O, Na$_2$S$_2$O$_3$·5H$_2$O, KOH, methanol and polyvinylpyrrolidone (PVP, $M_w$ = 40,000) were purchased from Chengdu Kelong chemical co. LTD (Chengdu, China) without further purification. L-ascorbic acid (AA) were purchased from Sigma-Aldrich (St. Louis, MO, USA) without further purification.

2.2. Preparation of NiCo$_2$O$_4$ NCs

Cu$_2$O templates were firstly prepared according to our previous work [12]. Simply, 10 mL NaOH solution (2 M) was added into 100 mL of CuCl$_2$·2H$_2$O (0.01 M) and stirred at 55 °C for 30 min. Then, 10 mL of AA (0.6 M) was added. After 3 h of reaction, Cu$_2$O cubes were collected and dried in vacuum.

A total of 10 mg cubic Cu$_2$O, 1mg NiCl$_2$·6H$_2$O and 2 mg CoCl$_2$·6H$_2$O were dispersed into 10 mL ethanol/water (1:1), and then, 0.33 g PVP was added and stirred for 30 min. Afterwards, 4 mL Na$_2$S$_2$O$_3$·5H$_2$O solution (1 M) was slowly dropped at room temperature. After 3 h, hydroxide precursors were collected and dried in vacuum. Finally, the precursors were calcined using a tube furnace at 400 °C in air for 2 h with a heating rate of 1 °C min$^{-1}$. Co$_3$O$_4$ NCs and NiO NCs were respectively prepared as contrast samples using CoCl$_2$·6H$_2$O and NiCl$_2$·6H$_2$O only in the CEP process.

2.3. Electrochemical Measurements

Cyclic voltammetry (CV), chronamperometry and electrochemical impedance spectroscopy (EIS) were performed in 1 M KOH solution on CH1760E A191018 electrochemical workstation at room temperature. A three-electrode system was used with Ag/AgCl...
(saturated with KCl) and platinum disk (Φ = 2 mm) as the reference and counter electrodes, respectively. The Co₃O₄ NCs, NiO NCs and NiCo₂O₄ NCs-modified glassy carbon electrodes (GCE, Φ = 3 mm) were applied as working electrodes. Typically, GCE was carefully polished with 3 μm, 0.5 μm and 0.05 μm alumina powders, respectively. Then, 5 μL of the prepared sample suspension (1 mg mL⁻¹ in 0.1% Nafion solution) was measured with a pipette and dropped onto the surface of GCE, and then dried naturally.

2.4. Materials Characterization

The microstructures and morphologies of the samples were observed by field emission electron microscope (FESEM, SU8020) and high-resolution transmission electron microscope (HRTEM, FEI F20). The crystal structure and elemental composition were recorded by X-ray powder diffractometer (XRD, Rigaku D/Max-2400 using Cu-Kα radiation λ = 1.54 Å). The chemical state was determined by X-ray photoelectron spectroscopy (XPS, ESCALAB 250Xi) using a 500 μm X-ray spot (energy resolution 0.4 eV). The Brunauer–Emmett–Teller (BET, Belsort-max) was applied to analyze the specific surface area and pore structure.

3. Results and Discussion

3.1. Characterization

As shown in Figure 1a, Co²⁺ and Ni²⁺ were firstly adsorbed on the surface of Cu₂O in the ultrasonic process. The CEP process occurred on the surface of Cu₂O once S₂O₅²⁻ was added [13]. Cu₂O reacts with S₂O₅²⁻ and H₂O to form a soluble [Cu₂(S₂O₅)²⁻]₀⁻O⁻⁻ complex and abundant OH⁻ (reaction (1)). The part-hydrolyzation of S₂O₅²⁻ also facilitated the supply of OH⁻ (reaction (2)) [14]. Reactions (1) and (2) concurrently pushed reaction (3) forward, facilitating the formation of Ni-Co hydroxide precursor. The diffusion of S₂O₅²⁻ from the surface into the interior of the shell directly affected the rate of Cu₂O etching, while the transport of OH⁻ from internal to external sites promoted the growth of Ni-Co hydroxide precursor [15]. The two reaction processes coordinated together to achieve dynamic balance to promote the formation of hollow structure. In order to confirm the formation mechanism, the precipitate prepared at 0, 10, 20, 30, and 180 min was collected and observed by TEM (Figure 1b). With the introduction of S₂O₅²⁻, Cu₂O was gradually etched into the polyhedral structure due to the higher diffusion intensity of ions at the corners [15,16]. After the reaction lasted for 3 h, Cu₂O completely disappeared and hierarchical porous nanocages were obtained (Figure S1). Finally, NiCo₂O₄ NCs were obtained through the annealing of Ni-Co hydroxide precursor (reaction (4)). As observed in Figure 1c, the color of the reaction system gradually became yellow and the light green precipitates generated at the same time. The fading was attributed to the etching of Cu₂O, while the green precipitates were correlated to the formation of Ni-Co hydroxide precursor.

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\begin{align*}
\text{Cu}_2\text{O} + \text{S}_2\text{O}_5^{2-} + \text{H}_2\text{O} &\rightarrow [\text{Cu}_2(\text{S}_2\text{O}_5^{2-})_x]^{2-2x} + 2\text{OH}^- \quad (1) \\
\text{S}_2\text{O}_5^{2-} + \text{H}_2\text{O} &\leftrightarrow \text{HS}_2\text{O}_5^- + \text{OH}^- \quad (2) \\
\text{Ni}^{2+} + 2\text{Co}^{2+} + 6\text{OH}^- + 1/2\text{O}_2 &\rightarrow (\text{NiCo}_2\text{O}_2)(\text{OH})_4 + \text{H}_2\text{O} \quad (3) \\
(\text{NiCo}_2\text{O}_2)(\text{OH})_4 &\rightarrow \text{NiCo}_2\text{O}_4 + 2\text{H}_2\text{O} \quad (4)
\end{align*}
\]

As shown in Figure 2a, the strong peaks from the templates at 30°, 37°, 42°, 62°, 74° and 78° matched well with PDF#77-0199 of cubic Cu₂O. As observed in Figure 2b, no significant diffraction peaks were observed in the precursor, revealing poor crystallinity of Ni-Co hydroxide precursor. After calcination, the crystallinity of materials was obviously improved and the diffraction peaks at 36°, 43°, 64°, 75° and 77° were well indexed to the (111), (200), (220), (311) and (400) crystal planes of face-centered cubic NiCo₂O₄. XRD results clearly demonstrated the formation of high purity NiCo₂O₄ product. Furthermore, XPS measurements were performed to obtain detailed information on the elements and oxidation state of prepared NiCo₂O₄. As shown in Figure 2c, the survey spectrum displayed
a series of strong peaks related to Ni, Co, O and C species, indicating the main chemical elements of the NiCo$_2$O$_4$. In Figure 2d, two states of Co$^{2+}$ and Co$^{3+}$ were clearly observed according to Gaussian fitting. Specifically, the fitting peaks at 779.3 eV and 794.3 eV were ascribed to Co$^{3+}$. Another two fitting peaks at 781.0 eV and 795.8 eV were ascribed to Co$^{2+}$ [17]. Analogously, the Ni 2p spectra included two kinds of nickel species of Ni$^{2+}$ and Ni$^{3+}$ in Figure 2e. The fitting peaks at 854.0 eV and 871.7 eV were ascribed to Ni$^{2+}$, while the fitting peaks at 855.9 eV and 873.9 eV were related to Ni$^{3+}$ [18]. As shown in Figure 2f, the fine spectrum of O 1s displayed three peaks originated from M-O-M, C-O=C and O=C. The fitting peak of M-O-M at 528.5 eV was the typical metal-oxygen bond [19]. C-O=C at a binding energy of 530.5 eV corresponded to the high number of defect sites containing low oxygen coordination [20]. O=C at a binding energy of 531.7 eV could be ascribed to the multiplicity of physisorbed water at and within the surface [17,21]. The results of XPS demonstrated a mixed valence containing Co$^{2+}$, Co$^{3+}$, Ni$^{2+}$ and Ni$^{3+}$, which was consistent with previous reports [22]. The complex electronic states of Ni$^{2+}$/Ni$^{3+}$ and Co$^{2+}$/Co$^{3+}$ could afford enough active sites for methanol oxidation, which may be one of the important factors contributing to the high electrocatalytic performance.

Figure 1. (a) Schematic illustration of the proposed growth mechanism of NiCo$_2$O$_4$ NCs. (b) TEM images of the products monitored at different reaction times. (c) Optical photographs of the suspension at different reaction time after addition of etchant.

The surface morphologies of the CuO$_2$ templates, NiO NCs and Co$_3$O$_4$ NCs were examined and displayed in Figure 3. As shown in Figure 3a, the Cu$_2$O templates were uniformly dispersed, which was conducive to the adsorption of Ni$^{2+}$ and Co$^{2+}$. The surface of Cu$_2$O was smooth and the edge size was about 500 nm (Figure 3b). As observed in Figure 3c, the NiO cube was composed of a large number of stacked nanoparticles, and the NiO cube had nearly the same size compared to Cu$_2$O. As shown in Figure 3d, the NiO cube displayed a hollow structure with a wall thickness of about 80 nm. Similarly, the Co$_3$O$_4$ cube also displayed a hollow cubic feature (Figure 3e). However, the Co$_3$O$_4$ NCs mainly consisted of a large number of stacked nanosheets, forming a network structure (Figure 3f).
As observed in Figure 4a, the uniformly distributed Ni-Co hydroxide precursor accurately replicated the cubic structure of Cu$_2$O templates and had an edge size of 500 nm. As shown in Figure 4b, the surface of Ni-Co hydroxide precursor was composed of a large number of interacted nanosheets and formed a network structure. In addition, the precursor displayed a cage-like structure and the shell thickness was about 200 nm. After calcination, the nanosheets on the surface became thicker, more compact and the thickness of the shell reduced to about 100 nm (Figure 4c). Notably, the crinkly nanosheets structure was clearly investigated in Figure 4d and the result was consistent with the SEM image. The SAED pattern in the insert of Figure 4d exhibited well-defined rings, revealing the polycrystalline nature of NiCo$_2$O$_4$ NCs. As observed in Figure 4e,f, the lattice spacing of 0.241 nm, 0.209 nm, 0.245 nm and 0.205 nm corresponded to the (111), (200), (311) and (400) planes of spinel NiCo$_2$O$_4$, respectively. The results were consistent with the XRD analysis and a previous report [22]. On the basis of the above discussion, NiCo$_2$O$_4$ NCs were constructed by the combination of the CEP method and post calcination. The highly
porous structure provided sufficient active sites and mass transport channels, which are beneficial for electrocatalytic kinetics, leading to high electrocatalytic activity [23,24].

![Image of SEM and TEM images](Figure 4. (a) TEM images of Ni-Co hydroxide precursor and (b, c) SEM of prepared Ni-Co hydroxide precursor and NiCo2O4 NCs; (d-f) HRTEM of prepared NiCo2O4 NCs the inset shows SAED.)

### 3.2. Electrocatalytic Activity of NiCo2O4 NCs/GCE towards Methanol

The electrocatalytic activity of NiCo2O4 NCs/GCE and the contrast samples was detailedly evaluated by CV and EIS. Figure 5a shows the CV curves of the three electrodes in 1 M KOH in the absence of methanol. The distinct pairs of redox peaks were observed in all the three CV curves. The redox peaks of NiO NCs/GCE corresponded to the reversible transition of Ni ions, such as Ni2+/Ni3+ [25]. Similarly, the redox peaks of Co3O4 NCs/GCE were attributed to the transition between Co2+/Co3+ or Co3+/Co4+ [26]. The CV curve of NiCo2O4 NCs/GCE exhibited a much larger enclosed area than those of the Co3O4 NCs/GCE and NiO NCs/GCE, respectively. In addition, the onset potential towards methanol oxidation on the NiCo2O4 NCs/GCE was about 0.29 V (Figure 5c), which was lower than those of Co3O4 NCs/GCE (0.34 V, Figure S3) and NiO NCs/GCE (0.38 V, Figure S4), revealing higher electrocatalytic activity. As shown in Figure 5c, the EIS was carried out in 1 M KOH containing 0.5 M methanol and the equivalent circuit is displayed in the insert. In the circuit, $R_s$, $C$, $R_{ct}$ and $Z_w$ were the internal resistance, redox capacitance, charge transfer resistance and Warburg resistance, respectively [28,29]. Notably, the $R_s$ value of NiCo2O4 NCs/GCE (4.4 kΩ) was obviously lower than those of Co3O4 NCs/GCE (10.8 kΩ and NiO NCs/GCE (18.3 kΩ), indicating fast electron transfer rate within the electrode or at the electrode/electrolyte interface. The lower charge transfer resistance was related to the anisotropic feature of building blocks and relatively high conductivity of NiCo2O4. At low frequencies, the NiCo2O4 NCs/GCE displayed larger $Z_w$ than Co3O4 NCs/GCE and NiO NCs/GCE, revealing lower ion diffusion resistance. The lower ion diffusion resistance might be attributed to ample diffusion channels afforded by the interacted NiCo2O4 nanosheets. In order to support the kinetics analysis of EIS, the surfaces area and porosity of NiCo2O4 NCs were tested.
by BET. In Figure 5d, the curve presents a H₃-type hysteric loop in the range of 0.45–1.0, indicating a typical mesoporous characteristic [30,31]. The mean pore size of NiCo₂O₄ NCs/GCE was around 9 nm, which was ideal for the diffusion of methanol [32]. Moreover, the specific surface area and pore volume were 38.3 m² g⁻¹ and 0.2 cm³ g⁻¹, respectively, which were both higher than those of the precursor (30.0 m² g⁻¹, 0.1 cm³ g⁻¹, Figure S5). The large specific surface area provided abundant active sites for methanol catalysis, and the appropriate pore volume provided ordered diffusion channels for rapid transport [33]. In short, NiCo₂O₄ NCs/GCE exhibited rich redox active sites and transmission channels, leading to excellent electrocatalytic activity.

![Figure 5. CV curves of NiCo₂O₄ NCs/GCE, Co₃O₄ NCs/GCE and NiO NCs/GCE in 1 M KOH (a) without methanol and (b) with 0.5 M methanol at 50 mV s⁻¹; (c) EIS plots of NiCo₂O₄ NCs/GCE, Co₃O₄ NCs/GCE and NiO NCs/GCE in 1 M KOH with 0.5 M methanol; (d) N₂ adsorption–desorption isotherms of the NiCo₂O₄ NCs/GCE.](image)

The chronoamperometry is an effective tool to investigate electrochemical stability of the electrocatalyst. As shown in Figure 6a, the electrochemical stability of the NiCo₂O₄ NCs/GCE, Co₃O₄ NCs/GCE and NiO NCs/GCE for methanol oxidation at 0.45 V was investigated. Notably, the NiCo₂O₄ NCs/GCE displayed largest electrocatalytic current towards 0.5 M methanol. The current of NiCo₂O₄ NCs/GCE displayed a decrease at the initial stage due to poisoning of the intermediates, and then kept a relatively steady value until 1100 s [34,35]. The final current still maintained 85% of its original value, which was three times of the Co₃O₄ NCs/GCE and thirteen times of the NiO NCs/GCE. The CV tests were carried out for 1000 cycles to further investigate the stability of NiCo₂O₄ NCs/GCE. The maximum current density presented an 8% decrease at the 500th cycle, and maintained 80% of the initial value after 1000 cycles. The hierarchical porous structure provided sufficient interspaces for accommodation of volume change and structural strain during electrocatalysis, resulting in excellent long-term stability towards methanol.
Co3O4 NCs/GCE and NiO NCs/GCE in 0.5 M methanol; (d) The CV stability test of NiCo2O4 NCs/GCE in 0.5 M methanol at a scanning rate of 50 mV s^{-1}.

4. Conclusions

In summary, the NiCo2O4 NCs were successfully synthesized through the CEP method combined with a post-annealing process. The designed NiCo2O4 NCs were constructed through the interaction between NiCo2O4 NSs and formed a hierarchical cage-like structure. As a catalytic electrode for methanol oxidation, the NiCo2O4 NCs/GCE exhibited high electrocatalytic activity in terms of low onset potential (0.29 V) and excellent long-term stability (80% after 1000 cycles). It is demonstrated that the NiCo2O4 NCs/GCE was an ideal electrode for DMFCs and the design of hollow hierarchical structure was an effective method to obtain highly active 2D electrocatalysts.

Supplementary Materials: The Supporting Information is available free of charge on the MDPI Publications website at https://www.mdpi.com/article/10.3390/nano1102667/s1, Figure S1: XPS survey of NiCo2O4, Figure S2: CV curves of NiCo2O4 NCs/GCE in 1 M KOH without methanol and with 0.5 M methanol at 50 mV s^{-1}, Figure S3: CV curves of Co3O4 NCs/GCE in 1 M KOH without methanol and with 0.5 M methanol at 50 mV s^{-1}, Figure S4: CV curves of NiO NCs/GCE in 1 M KOH without methanol and with 0.5 M methanol at 50 mV s^{-1}, Figure S5: N2 adsorption-desorption isotherms of the Ni-Co hydroxide precursors.

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References

1. Joghee, P.; Malik, J.N.; Pylypenko, S.; O’Hayre, R. A review on direct methanol fuel cells—In the perspective of energy and sustainability. MRS Energy Sustain. 2015, 2, 3. [CrossRef]
2. Bangyang, J.; Tang, H.; Pan, M. Well-ordered sulfonated silica electrolyte with high proton conductivity and enhanced selectivity at elevated temperature for DMFC. Int. J. Hydrog. Energy 2012, 37, 4612–4618. [CrossRef]
3. Gong, L.; Yang, Z.; Li, K.; Xing, W.; Liu, C.; Ge, J. Recent development of methanol electrooxidation catalysts for direct methanol fuel cell. J. Energy Chem. 2018, 27, 1618–1628. [CrossRef]
33. Lv, X.; Zhu, Y.; Jiang, H.; Yang, X.; Liu, Y.; Su, Y.; Huang, J.; Yao, Y.; Li, C. Hollow mesoporous NiCo$_2$O$_4$ nanocages as efficient electrocatalysts for oxygen evolution reaction. *Dalton Trans.* **2015**, *44*, 4148–4154. [CrossRef] [PubMed]

34. Tong, X.; Qin, Y.; Guo, X.; Moutanabbir, O.; Ao, X.; Pippel, E.; Zhang, L.; Knez, M. Enhanced Catalytic Activity for Methanol Electro-oxidation of Uniformly Dispersed Nickel Oxide Nanoparticles-Carbon Nanotube Hybrid Materials. *Small* **2012**, *8*, 3390–3395. [CrossRef] [PubMed]

35. Amin, R.; Hameed, R.A.; El-Khatib, K.; Youssef, M.E.; Elzatahry, A. Pt–NiO/C anode electrocatalysts for direct methanol fuel cells. *Electrochim. Acta* **2012**, *59*, 499–508. [CrossRef]