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One-step synthesis of Ni(OH)₂/MWCNT nanocomposites for constructing a nonenzymatic hydroquinone/O₂ fuel cell

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In this work, a H-type hydroquinone/O₂ fuel cell was assembled and shows high energy density in neutral phosphate buffer solution at moderate temperature. The anodic material, Ni(OH)₂/MWCNTs, was synthesized by a one-step hydrothermal synthesis method to oxidize hydroquinone. The cathode material, Pt/MWCNTs, was obtained by an electrodeposition method, and shows great oxygen reduction reaction (ORR) activity. The properties and the morphology of Ni(OH)₂/MWCNT nanocomposites were characterized by TEM, XPS, EDS-mapping and electrochemical methods, like cyclic voltammetry (CV) and electrochemical impedance spectroscopy (EIS). The results show that Ni(OH)₂/MWCNTs can effectively oxidize hydroquinone and play a dominant role in enhancing the fuel cell performance. The nonenzymatic fuel cell possesses a high power density of 0.24 mW cm⁻² at a cell potential of 0.49 V.

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1. Introduction

Environmental pollution problems, such as air, soil and water pollution caused by the consumption of fossil energy are becoming more and more serious. Therefore, it is imperative to develop some efficient and green energies to substitute for fossil fuels. In recent years, biofuel cells (BFC), which can directly convert chemical energy to electric power by utilizing biocatalysts, have shown some unique advantages compared to other electric power facilities, such as low cost, excellent stability and high power output.^{1–3} It has been reported that ethanol and glucose have been catalyzed and oxidized as feedstock for conventional biofuel cells.^{4,5} In modern industrial production, phenol organic compounds are considered as important pollutants in medicines, foods and the fine chemical industry. Hydroquinone (HQ, 1,4-dihydroxybenzene), as one of the phenolic compounds, plays an important role in cosmetics, pesticides, food antioxidants and other fields.⁶ However, HQ is also a serious environmental pollutant due to its toxicity to humans and difficulty in degradation, which can cause fatigue, headache and tachycardia in humans, and even lead to acute myeloid leukemia at high levels.^{7–9} Therefore, it is very important to develop some green and effective methods for disposing HQ.

As one of transition metal hydroxides, Ni(OH)₂ has the advantages of high stability, low toxicity and inexpensive,¹⁰ and

is widely used in the fields of supercapacitors, sensors and lithium ion batteries.^{11,12} Generally, Ni(OH)₂ exists in two forms of α -Ni(OH)₂ and β -Ni(OH)₂. The β -Ni(OH)₂ possesses a hexagonal arrangement of brucite-type layers ordered along the axis *C*, correspondingly, the other structure is randomly stacked along the *C* axis.¹³ It has been reported that there were many successful methods for the synthesis of Ni(OH)₂, including solvent heat,¹⁴ ultrasonic,¹⁵ hydrothermal,¹⁶ co-precipitation¹⁷ and so on. In recent years, multi-walled carbon nanotubes (MWCNTs) are very popular and widely used in biosensors,¹⁷ supercapacitors,¹⁸ biofuel cells¹⁹ and other fields because of their small size, large specific surface area, high conductivity and chemical stability. Mostly, MWCNTs are often combined with transition metal oxides for enhancing the features in application.^{20–22} For example, MWCNTs and ZnO-Co₃O₄ are serve as electrode materials to construct the non-enzyme electrochemical sensors for high-sensitivity detection of tanshinol.²³ And SnS₂/MWCNTs composite was used as the modified material to prepare a chemically modified electrode (CME), which was used for the rapid detection of H₂O₂ and produce a strong current response.²⁴ Although the preparation of Ni(OH)₂/MWCNTs has been reported in some works, like supercapacitors,²⁵ and sensors,²⁶ the application of Ni(OH)₂ or Ni(OH)₂/MWCNTs nanomaterials in hydroquinone biofuel cells is still a new question worth exploring, which might also catalyze and degrade HQ in solution.

Hence, Ni(OH)₂/MWCNTs nanocomposites were synthesized by one-step hydrothermal method. The electrochemical properties and the morphology of nanocomposites were characterized by TEM, EDS-mapping and other electrochemical methods, like cyclic voltammetry and electrochemical impedance

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spectroscopy. The effects of HQ concentrations and scanning rates on peak current were also investigated. As a result, the nanocomposite exhibits higher electrochemistry properties than single $\text{Ni}(\text{OH})_2$ or MWCNTs prepared in this work, an H-type hydroquinone/ O_2 fuel cell was assembled and shows high energy density (0.24 mW cm^{-2}). Which demonstrated that $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanocomposites possess great application prospect as a potential anode material in HQ biofuel cell.

2. Experimental

2.1 Materials

Multiwall carbon nanotubes (MWCNTs) were purchased by Shenzhen Nanotech Port Co. Ltd., $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$, NaOH, hydroquinone, $\text{K}_3[\text{Fe}(\text{CN})_6]$, $\text{K}_4[\text{Fe}(\text{CN})_6]$ were purchased from Shanghai reagents factory (China). All reagents were analysis reagent (A.R.), and used as received without further purification. Phosphate buffered saline (PBS) solution were made of standard storage solution of $0.10 \text{ mol L}^{-1} \text{ Na}_2\text{HPO}_4$ and $0.10 \text{ mol L}^{-1} \text{ NaH}_2\text{PO}_4$. All solutions were prepared using deionized water.

2.2 Characterization

Electrochemical measurements were tested on CHI760E Electrochemical Workstation (Shanghai Chenhua Instrument Co.,

Ltd., China) and with rotating disk electrode (RDE) (PINE AFMSRCE, USA) at room temperature. A conventional three-electrode cell was used, containing a glassy carbon electrode (GCE, 3.0 mm in diameter) as the working electrode, platinum wire as the counter electrode and the standard calomel electrode (SCE) as reference electrode. Transmission electron microscopy (TEM) were tested on JEM-2100 with an accelerating voltage of 200 kV. X-ray diffraction (XRD) was performed on a Bruker D8 advance A25 at 40 kV and 40 mA for $\text{Cu K}\alpha$ with a step size of 0.01° in 2θ . X-ray photoelectron spectroscopy (XPS) analysis was measured on a PHI5000V Versaprobe system using monochromatic $\text{Al K}\alpha$ radiation, and all bindings energies were calibrated by the C 1s peak at 284.7 eV.

2.3 Synthesis of $\text{Ni}(\text{OH})_2$ and $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanocomposites

Typically, 2 g of MWCNTs was added in a 150 mL round bottle flask with a concentrated sulfuric acid (98%, 112.5 mL) and nitric acid (68%, 37.5 mL) under ultrasonic processing. The resulting black powder was collected by centrifugation and washed by deionized water until the pH of the filtrate was approximately neutral. Then, MWCNTs were dried in a vacuum oven for 24 h at 80°C .

The $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanocomposites were synthesized by hydrothermal method. The specific steps were as follows: 5 mg of MWCNTs was dispersed in 15 mL deionized water and

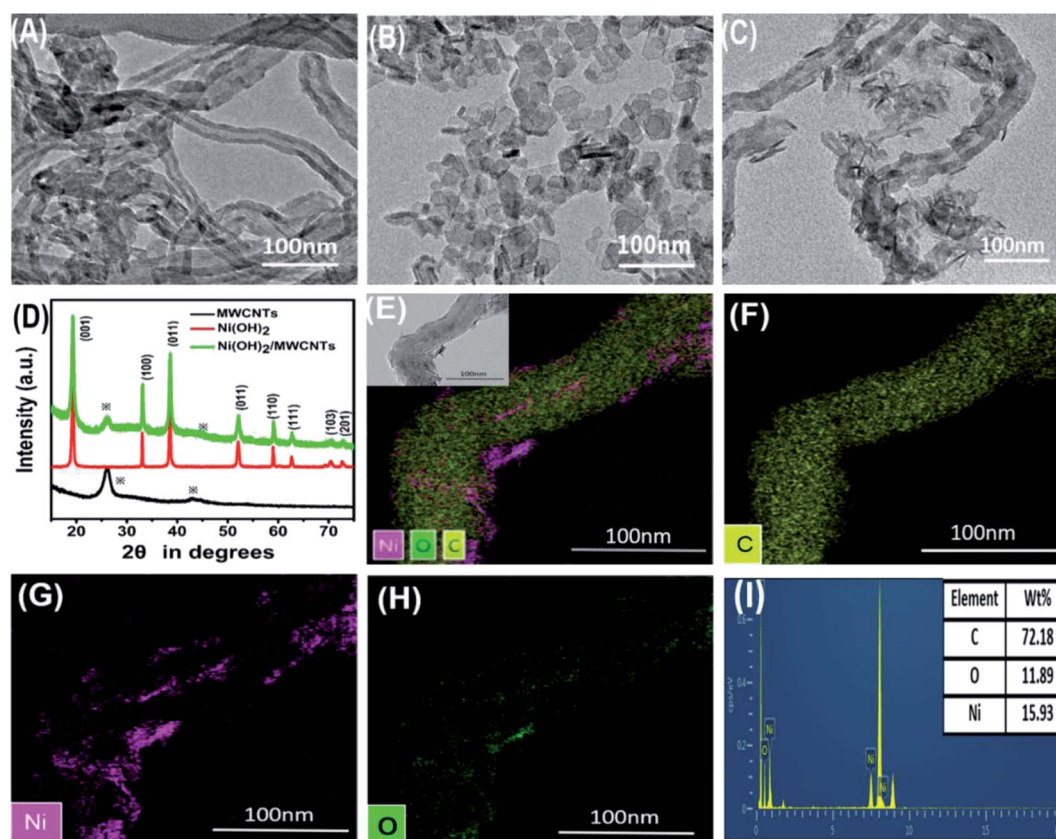


Fig. 1 TEM images of MWCNTs (A), $\text{Ni}(\text{OH})_2$ (B) and $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ (C); XRD patterns of the $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ (top), $\text{Ni}(\text{OH})_2$ (middle) and MWCNTs (bottom) (D); TEM images elemental mapping of the $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanocomposites are show in (E), (F), (G) and (H), respectively. And EDS spectra of $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ (I).



treated under ultrasonication for 30 min. Then, 100 μL of $\text{NiCl}_2 \cdot 6\text{H}_2\text{O}$ aqueous solution (0.1 M) was injected in the suspension. The solution pH was then adjusted to 12 by adding NaOH solution (0.5 M). After stirring vigorously for 1 h, the solution was transferred into a 25 mL steel autoclave, sealed and maintained at 180 $^\circ\text{C}$ for 24 h. The obtained products were centrifuged, washed with ethanol and deionized water respectively, finally dried at 80 $^\circ\text{C}$ for 10 h. For the sake of comparison, pure $\text{Ni}(\text{OH})_2$ was also synthesized by the same process without the addition of MWCNTs.

2.4 Preparation of modified electrodes

The working electrode was modified by MWCNTs, $\text{Ni}(\text{OH})_2$, or $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanocomposites. Typically the GCE ($\phi = 3.0$ mm) was polished with 1.0, 0.3, 0.05 μm of alumina slurry, washed with deionized water and absolute ethanol, and finally dried in air. An amount of MWCNTs, $\text{Ni}(\text{OH})_2$ and $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ powder was dispersed in 1 mL of water/ethanol solution (1/1 v/v), respectively, followed by ultrasonication for 30 min. Furthermore, GCEs were conducted by 5 μL of ultrasonicated mixture and 2 μL 0.05% Nafion solution, and finally dried in air at room temperature to obtain the MWCNTs/GCE, $\text{Ni}(\text{OH})_2/\text{GCE}$, $\text{Ni}(\text{OH})_2/\text{MWCNTs}/\text{GCE}$. To the Pt/MWCNTs/GCE, 5 μL of MWCNTs mixture was deposited on GCE, and the Pt film was carried out by CVs in 0.5 mol L^{-1} H_2SO_4 solution containing 1.8×10^{-3} mol L^{-1} H_2PtCl_6 with a potential

scanning from 0.40 to -0.30 V at 100 mV s^{-1} for 30 cycles. The obtained electrode is recorded as Pt/MWCNTs/GCE.

2.5 Electrochemical measurements

Electron transfer properties of the composites modified electrodes were assessed by cyclic voltammetry and electrochemical impedance spectroscopy in 0.1 mol L^{-1} KCl solution, which contains 5.0 mmol L^{-1} $[\text{Fe}(\text{CN})_6]^{3-/4-}$ redox pair. EIS measurements were tested in the frequency range from 0.10 Hz to 100 kHz at an open circuit potential (E_{ocp}) with 10 mV as the amplitude potential.

2.6 Non-enzymatic biofuel cells design

The H-type dual chambered fuel cell was constructed with $\text{Ni}(\text{OH})_2/\text{MWCNTs}/\text{GCE}$ as anode and Pt/MWCNTs/GCE as cathode, respectively. And the Nafion membrane was used to separate the anode and cathode compartments, which was injected into 0.1 mol L^{-1} PBS (pH 7.4) containing 4 mmol L^{-1} HQ as anode solution and 0.1 mol L^{-1} PBS (pH 7.4) saturated with O_2 as cathode solution. Linear sweep voltammetry (LSV) is subsequently used for obtaining the polarization curves of the fuel cell.

3. Results and discussion

3.1 Characterization of the synthesized nanocomposites

The synthesis of $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ was prepared *via* a one-step method without assistance of surfactants in aqueous phase.

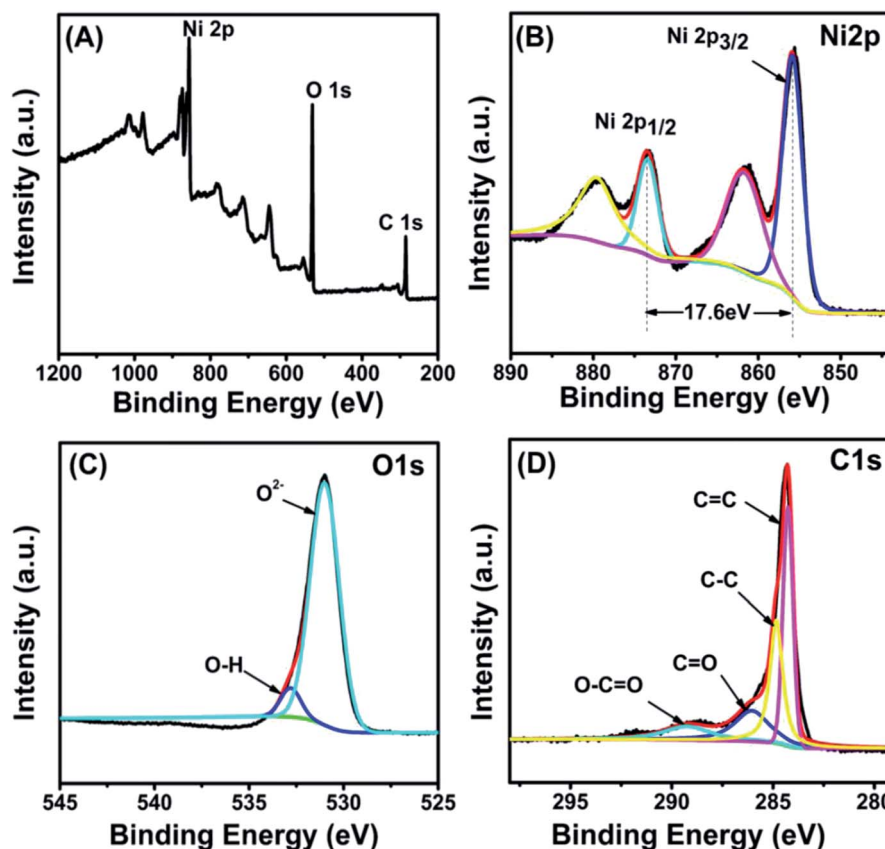


Fig. 2 XPS spectra of nanocomposite: survey spectrum (A), curve fitting spectrum of Ni 2p (B), O 1s (C), C 1s (D).

First, carbon nanotube is used as a upholder to anchor the $\text{Ni}(\text{OH})_2$. According to the literature,²⁷ $\text{Ni}(\text{OH})_2$ is usually positively charged, which is electrophilic with carbon nanotube. It can be supposed that the interaction between $\text{Ni}(\text{OH})_2$ and nanotube is electrostatic interactions.

Transmission electron microscopy was performed to investigate the morphology of the pure MWCNTs, $\text{Ni}(\text{OH})_2$ and $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanomaterials. TEM image (Fig. 1(A)) shows that the surface of bare MWCNTs is relatively smooth with an average diameter of approximately 40 nm. As observed in Fig. 1(B), the size of near hexagonal $\text{Ni}(\text{OH})_2$ nanoplates is about 10–30 nm with the thickness of 1.5–3.0 nm from the vertically aligned nanoplates. A typical TEM image (Fig. 1(C)) clearly demonstrates the successful growth of $\text{Ni}(\text{OH})_2$ nanoplates on MWCNTs. The XRD patterns in Fig. 1(D) are indexed to the hexagonal phase of typical- $\text{Ni}(\text{OH})_2$ (JCPDS-14-0117) and the crystalline MWCNTs, respectively. The sharp diffraction peaks at $2\theta = 19.258, 33.064, 38.541, 39.098, 52.100, 59.052, 60.240, 62.276, 69.346, 70.478, 72.737, 73.129$ and 82.608 correspond to (001), (100), (101), (002), (102), (110), (003), (111), (200), (103), (201), (112) and (202) crystal planes, respectively. No peaks referring to the other impurities can be observed in XRD patterns, which indicates the formation of pure $\text{Ni}(\text{OH})_2$ phase. For the further analysis, the elemental dispersive X-ray (EDX) spectroscopy of $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ composites were also

measured. TEM combined with element mapping based on EDX further confirmed the presence of Ni, O and C elements in the $\text{Ni}(\text{OH})_2/\text{MWCNTs}$. Fig. 1(E–H) shows that TEM-EDS concentration profiles of C, Ni, and O measured in the $\text{Ni}(\text{OH})_2$ also indicated the uniform coating of $\text{Ni}(\text{OH})_2$ on MWCNTs. In addition, it can be clearly seen from Fig. 1(I) that mass fractions of Ni, O and C elements are 15.93%, 11.89% and 72.18%, respectively.

XPS measurements (Fig. 2(A)) were also employed to analyze the chemical states of all elements except H in the $\text{Ni}(\text{OH})_2/\text{MWCNTs}$ nanomaterials, confirming the presence of Ni, O, and C in the composite. The high-resolution spectrum of Ni 2p in Fig. 2(B) showed two main peaks at 855.8 eV and 873.4 eV with a spin-energy separation of 17.60 eV, corresponding to Ni 2p_{3/2} and Ni 2p_{1/2}, respectively, which is typical characteristics of $\text{Ni}(\text{OH})_2$ phase.²⁸ Ni^{2+} introduced in the reaction was not converted into other substances. For oxygen element, the O 1s core level in Fig. 2(C) can be deconvoluted into two contributions at 531.1 eV and 532.9 eV, corresponding to O^{2-} and O–H in $\text{Ni}(\text{OH})_2$, respectively. In Fig. 2(D), binding energies observed at 284.74 eV, 284.84 eV, 287.6 eV and 289.25 eV correspond to C=C, C–C, C=O and O=C–O, respectively, which correspond to graphite like sp² carbon atoms C=C, C–C and carbon in carboxyl or ester groups (O=C–O) in the pretreated MWCNTs.

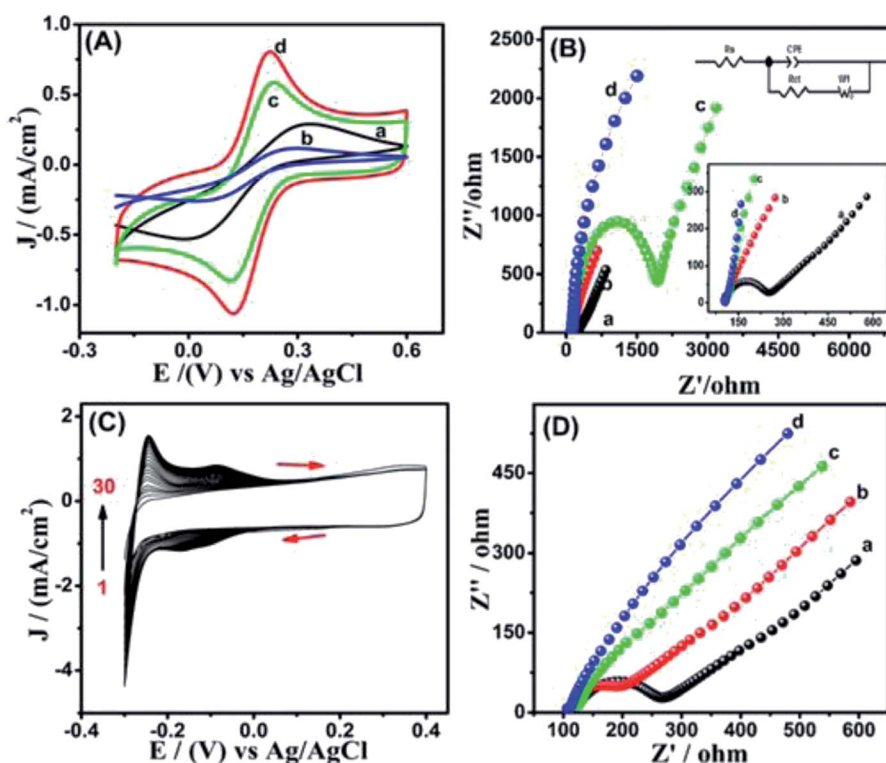


Fig. 3 (A) CV spectra of different anodic modified electrode in the 10 mmol L^{−1} $[\text{Fe}(\text{CN})_6]^{3-/4-}$ containing 0.1 mol L^{−1} KCl solution at the scan rate of 100 mV s^{−1}; (B) EIS of modified electrodes with different capping agents, bare GCE (a), $\text{Ni}(\text{OH})_2/\text{GCE}$ (b), MWCNTs/GCE (c) and $\text{Ni}(\text{OH})_2/\text{MWCNTs}/\text{GCE}$ (d), in 5.0 mmol L^{−1} $[\text{Fe}(\text{CN})_6]^{3-/4-}$ mixture containing 0.1 mol L^{−1} KCl, frequency range, 0.1–10⁵ Hz. (C) CV spectra for electrodeposition process of the film obtained from a solution mixture containing 1.8×10^{-3} mol L^{−1} H_2PtCl_6 and 0.5 mol L^{−1} H_2SO_4 between +0.4 V and −0.3 V (vs. SCE) at 100 mV s^{−1}; (D) EIS in 5.0 mmol L^{−1} $[\text{Fe}(\text{CN})_6]^{3-/4-}$ mixture containing 0.1 mol L^{−1} KCl at bare GCE (a), Pt/GCE (b), MWCNTs/GCE (c), Pt/MWCNTs/GCE (d), respectively.



All the above results indicated that β -Ni(OH)₂ nanoplates have been successfully coated on MWCNTs.

3.2 Electrochemical characterization

Cyclic voltammetry and electrochemical impedance spectroscopy are two kinds of convenient methods to study the electrochemical properties of various modified electrodes. Fig. 3(A) shows the CVs of different modified electrodes, from which the oxidation and reduction peaks of [Fe(CN)₆]^{3-/4-} can be clearly observed at bare GCE (curve a). When Ni(OH)₂ is fixed on the surface of the electrode (curve b), the peak current decreases because of the poor conductivity of Ni(OH)₂. As for MWCNTs/GCE (curve c), the peak current increases sharply due to its large specific surface area and excellent electrocatalytic performance. In conclusion, Ni(OH)₂/MWCNTs/GCE (curve d) electrode showed higher peak current than those of other electrodes, which can be attributed to the synergistic effects of Ni(OH)₂ nanoparticles and MWCNTs. Several studies have shown that the presence of oxygen species on carbon materials' surface, such as carboxyl and hydroxyl groups, can enhance the dispersion of the metal oxide, thus increasing the performance of the resulting composite materials.²⁹ Generally, the EIS consists of a semicircle in the high-frequency region and a straight line in the low-frequency region, corresponding to the charge transfer resistance and ion diffusion resistance in the electrode, respectively.³⁰ In Fig. 3(B), it is worth mentioning that the Nyquist plot for the MWCNTs electrode did not exhibit a semicircle in the high-frequency region, due to its low charge

transfer resistance (curve c). The Ni(OH)₂ electrode displays a much larger radius of semicircle in the high-frequency region compared to that of the bare electrode (curve b), indicating its higher faradaic charge transfer resistance. Besides, the Ni(OH)₂/MWCNTs electrode exhibits a more vertical line than that of the Ni(OH)₂ electrode and MWCNTs electrode in the low-frequency region (curve d), illustrating its better abilities to transfer electrons and lower diffusion resistance for ions. In conclusion, the carbon material can significantly enhance the electrical conductivity of the Ni(OH)₂, thus leading to significant improvement of the electrochemical performance.

Fig. 3(C) shows the CVs of the electrodeposition process of Pt film on MWCNTs/GCE. With the increase of cycle number from 1 to 30, all the redox peak current show positive deviation or negative deviation correspondingly, indicating that platinum has been successfully deposited on MWCNTs/GCE. Fig. 3(D) shows the EIS results on bare GCE (a), Pt/GCE (b), MWCNTs/GCE (c), and Pt/MWCNTs/GCE (d). Clearly, the diameter of the semicircle (curve d) is smaller and the slope of the straight line is higher than those of other electrodes, indicating that the surface of the electrode forms a layer of Pt, which can be used as an electron transfer medium to enhance the electron transfer.

3.3 Electrochemical behavior of HQ on modified electrodes

To analyze electrocatalytic redox phenomena of different electrodes to HQ oxidation, a comparative study was conducted by CV in 0.1 mol L⁻¹ PBS with range from -0.4–0.6 V with a scanning rate of 50 mV s⁻¹. There are no distinct redox peaks on

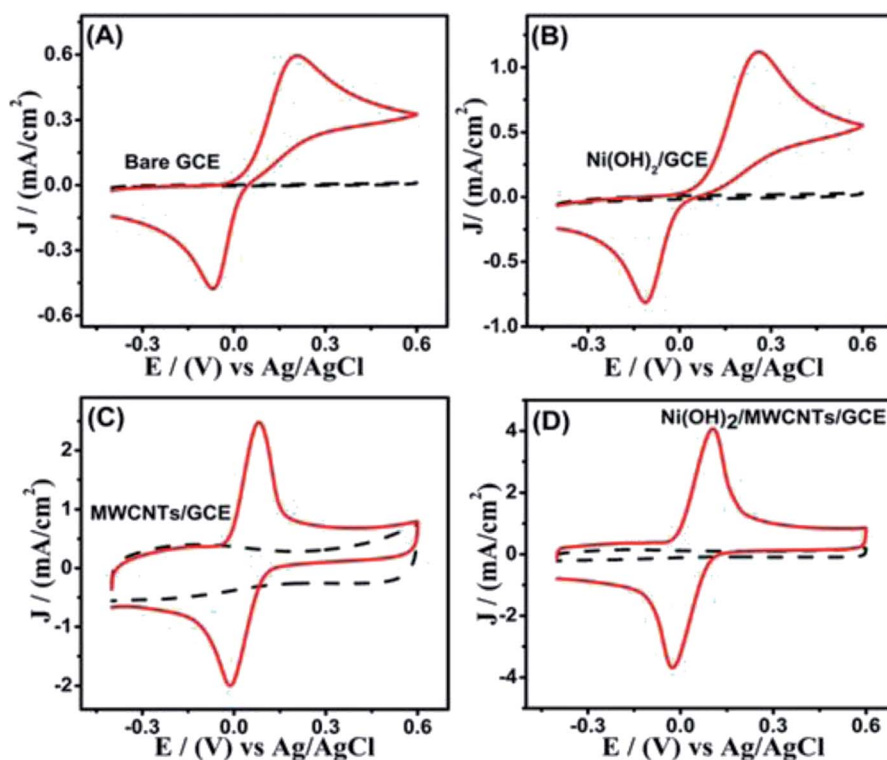


Fig. 4 CVs of bare GCE (A), Ni(OH)₂/GCE (B), MWCNTs/GCE (C), and Ni(OH)₂/MWCNTs/GCE (D) in absence (black dash line) and presence (red solid line) of 4 mmol L⁻¹ HQ in 0.1 mol L⁻¹ PBS (pH 7.4). Scan rate is 50 mV s⁻¹.

bare GCE, Ni(OH)₂/GCE, MWCNTs/GCE, and Ni(OH)₂/MWCNTs/GCE in 0.1 mol L⁻¹ PBS without addition of HQ (black dash line) in Fig. 4(A–D). However, it can be seen from Fig. 4 that all of the electrodes show huge change in the redox current density upon addition of HQ, indicating the better electrocatalytic properties of the nanomaterials for HQ oxidation. It can be concluded that the electrochemical response of Ni(OH)₂/MWCNTs/GCE electrode is intensively higher than those of bare GCE, Ni(OH)₂/GCE, and MWCNTs/GCE electrodes.

Fig. 5(A) shows the relationship of scan rate to the peak currents, a gradual shift of the peak potential in CV curves (positive shift for anodic and negative shift for cathodic peak) is detected in the range of 20–140 mV s⁻¹ and the scan rate is linear to 1/2 of the peak current in Fig. 5(B). It is also indicated that the whole electrochemical reaction is a kinetic limitation. In Fig. 5(C), with the increasing HQ concentration, the same phenomenon of anodic and cathodic peak current in CVs also occur, and these high current densities were linear with both flow rate and HQ concentration in the range from 0.05 to 20 mmol L⁻¹ in Fig. 5(D). It is obvious that Ni(OH)₂/MWCNTs nanocomposite owning more outstanding catalytic performance.

3.4 Electrocatalytic behavior of Pt/MWCNTs as cathode material

Square wave voltammetry (SWV) and linear sweep voltammetry (LSV) was carried out to study the oxygen reduction reaction

(ORR) activity. It can be seen from Fig. 6(A) that Pt/MWCNTs/GCE (d) electrode shows a higher oxygen reduction peak than those of bare GCE (a), Pt/GCE (b) and MWCNTs/GCE (c) electrodes, which indicate that Pt/MWCNTs is an excellent catalyst and can be used to further study the ORR activity. LSV measurements at 1600 rpm in Fig. 6(B), which can further prove that the ORR activity of Pt/MWCNTs is best with highest half-wave potential and diffusion limiting current density. The diffusion limiting current density of the Pt/GCE, MWCNTs/GCE and Pt/MWCNTs/GCE is 3.76, 2.79 and 5.88 mA cm⁻², respectively. And the half-wave potential is 0.788, 0.651 and 0.809 V, respectively.

In fact, platinum itself has a good catalytic effect on the reduction reaction of oxygen. However, in the presence of carbon tubes, the composites exhibited better catalytic properties. As mentioned above, the synergistic effect of MWCNTs composite with various nanomaterials often can significantly improve the catalytic performance of materials.

3.5 Performance of the fuel cell

The fuel cell test results suggest that the Ni(OH)₂/MWCNTs nanomaterials have excellent catalytic activity to HQ, while Pt/MWCNTs have ORR activity when worked as a cathode. Hence, an H-type electronical device was constructed with Ni(OH)₂/MWCNTs and Pt/MWCNTs as anode and cathode materials, respectively.

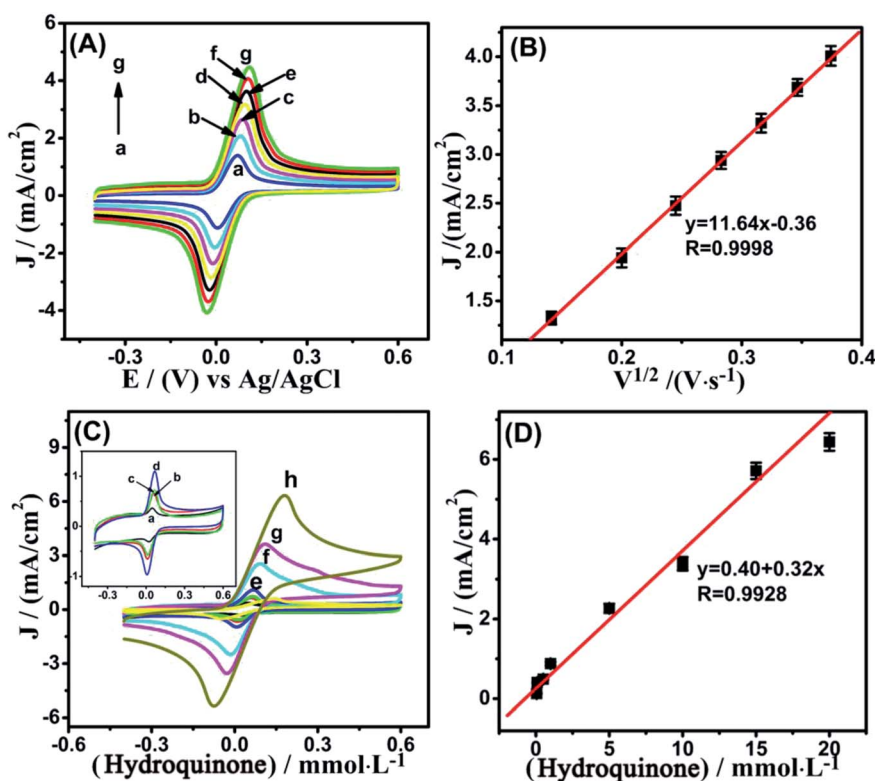


Fig. 5 (A) CV spectra of Ni(OH)₂/MWCNTs/GCE in the 0.1 mol L⁻¹ PBS (pH 7.4) containing 4.0 mmol L⁻¹ HQ at different scan rates; (B) plots of the anodic peak current density (mA cm⁻²) with the scan rates^{1/2}; (C) CV spectra of Ni(OH)₂/MWCNTs/GCE in the 0.1 mol L⁻¹ PBS (pH 7.4) at different concentrations of HQ with a scan rate of 50 mV s⁻¹; (D) plots of the anodic peak current density (mA cm⁻²) with different concentration of HQ.



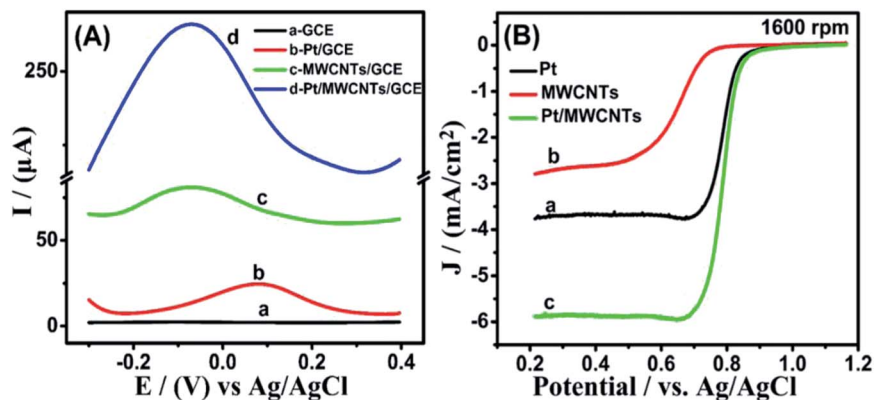
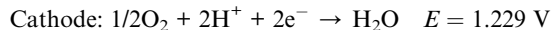
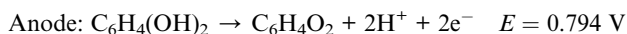


Fig. 6 (A) SWV spectra of bare GCE (a), Pt/GCE (b), MWCNTs/GCE (c), Pt/MWCNTs/GCE (d) in 0.1 mol L⁻¹ of PBS (pH 7.4) solutions saturated with O₂; (B) LSV of Pt, MWCNTs and Pt/MWCNTs in 0.1 mol L⁻¹ of KOH solutions saturated with O₂.

The detailed mechanisms of whole reactions, illustrated in Fig. 7(A), indicate that after the electrons and protons are produced from HQ, the electrons transfer to the cathode connected with external circuit, and the protons transferred through Nafion membrane. On the anode, HQ is correspondingly oxidized into *P*-benzoquinone (*P*-BQ), whereas oxygen is reduced on the cathode, thus completing the redox reaction for BFC operation. The overall reaction mechanisms in HQ fuel cell are represented as follows:



The polarization and power density curves of HQ fuel cell are shown in Fig. 7(B). It is obvious that the performance of the fuel cell constructed by the composite material (curve c and c1) is significantly better than that of the single Ni(OH)₂ (curve a and a1), or MWCNTs (curve b and b1). As can be read, a peak power

density of 0.24 mW cm⁻² at an open circuit voltage (*V*_{oc}) of around 0.49 V was obtained (curve c and c1). Most strikingly, the value of *P*_{max} of the Ni(OH)₂/MWCNTs based biofuel cell is distinctly higher than the previously reported enzyme-based BFCs^{31,32} or precious metal-based BFCs.^{33,34} Furthermore, the electrocatalytic performance of Ni(OH)₂/MWCNTs is better than single Ni(OH)₂ or MWCNTs catalysts, which indicates that the prepared composites is more suitable as an anode catalyst for biofuel cells.

4. Conclusion

A non-enzyme MWCNTs-based hydroquinone fuel cell has been assembled and shows high power density in neutral solution. Ni(OH)₂/MWCNTs was successfully prepared by hydrothermal method for the anodic material and could effectively oxidize hydroquinone in neutral PBS. It is apparent that the Ni(OH)₂/MWCNTs anodic catalyst played a dominant role in enhancing the catalytic activity and performance of HQ fuel cell. Accordingly, a biofuel cell was constructed with Ni(OH)₂/MWCNTs and Pt/MWCNTs as anode and cathode materials, respectively. As

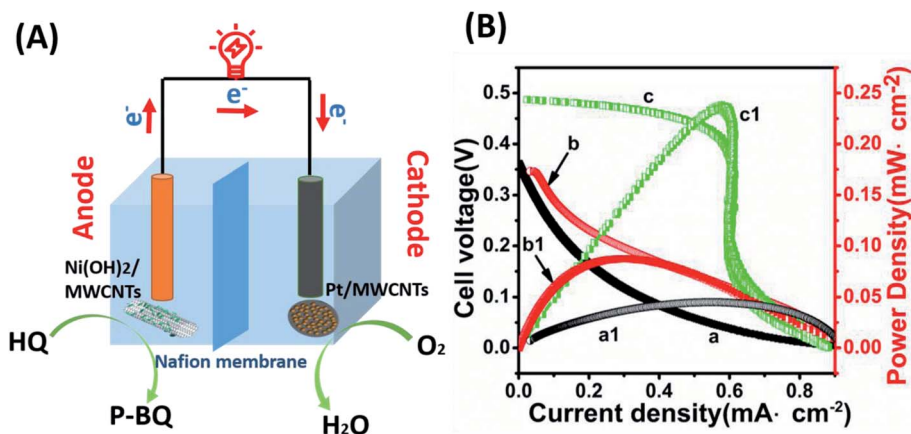


Fig. 7 (A) Schematic diagram of HQ fuel cell; (B) polarization (a, b and c) and power density curves (a1, b1 and c1) of HQ fuel cell. Anode: Ni(OH)₂/GCE (a and a1), MWCNTs/GCE (b and b1) and Ni(OH)₂/MWCNTs/GCE (c and c1) in 0.1 mol L⁻¹ PBS (pH 7.4) containing 4 mmol L⁻¹ HQ. Cathode: Pt/MWCNTs/GCE in 0.1 mol L⁻¹ PBS (pH 7.4) filled with O₂.



a result, it showed a high power density of 0.24 mW cm^{-2} at cell potential of 0.49 V, which can provide an experimental basis for the degradation of phenolic pollutants and simultaneously produce electric energy in real life in the future.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

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References

- 1 M. Nishizawa, *J. Power Sources*, 2019, **408**, 1–6.
- 2 K. Herkendell, A. Stemmer and R. Tel-Vered, *Nano Res.*, 2019, **12**, 767–775.
- 3 V. B. Oliveira, J. P. Pereira and A. M. F. R. Pinto, *Energy*, 2017, **133**, 652–665.
- 4 X. Wang, J. H. Kim and Y. B. Choi, *Korean J. Chem. Eng.*, 2019, **36**, 1172–1183.
- 5 J. H. Franco, K. J. Klunder and V. Russell, *Electrochim. Acta*, 2020, **331**, 135254.
- 6 C. Ma, N. He and Y. Zhao, *Appl. Biochem. Biotechnol.*, 2019, **189**, 1291–1303.
- 7 S. L. Allen and F. P. Zamborini, *Langmuir*, 2019, **35**, 2137–2145.
- 8 R. Cabrera-Alonso, E. Guevara and M. Ramírez-Elías, *Skin. Res. Technol.*, 2019, **25**, 20–24.
- 9 G. Ma, H. Xu and M. Wu, *Microchim. Acta*, 2019, **186**, 689.
- 10 J. J. William, I. M. Babu and G. Muralidharan, *Mater. Chem. Phys.*, 2019, **224**, 357–368.
- 11 Z. Liu, Y. Zhang and J. Feng, *Sens. Actuators, B*, 2019, **287**, 551–556.
- 12 Y. Liu, C. Yan and G. Wang, *ACS Appl. Mater. Interfaces*, 2019, **11**, 9984–9993.
- 13 X. Wang, P. Sebastian and A. C. Millan, *J. New Mater. Electrochem. Syst.*, 2005, **8**, 101–108.
- 14 Z. Zhai, Q. Liu and Y. Zhu, *J. Alloys Compd.*, 2019, **775**, 1316–1323.
- 15 V. Boychuk, V. Kotsyubynsky and B. Rachiy, *Mater. Today: Proc.*, 2019, **6**, 106–115.
- 16 S. Kettaf, O. Guellati and A. Harat, *SN Appl. Sci.*, 2019, **1**, 34.
- 17 L. A. Saghatforoush, M. Hasanzadeh and S. Sanati, *Bull. Korean Chem. Soc.*, 2012, **33**, 2613–2618.
- 18 J. Acharya, B. G. S. Raj and T. H. Ko, *Int. J. Hydrogen Energy*, 2020, **45**, 3073–3085.
- 19 J. Ji, J. Woo and Y. Chung, *Chem. Eng. J.*, 2020, **381**, 122679.
- 20 M. A. Ashraf, Z. Liu and D. Zheng, *Phys. E*, 2020, **116**, 113723.
- 21 K. Zhao, S. Li and M. Huang, *Chem. Eng. J.*, 2019, **358**, 924–935.
- 22 Y. Prabhu, B. Sreedhar and U. Pal, *Mater. Today: Proc.*, 2019, **8**, 419–426.
- 23 C. Zhang, J. Ren and Y. Xing, *Mater. Sci. Eng., C*, 2020, **108**, 110214.
- 24 Z. Duan, C. Huang and X. Yang, *Anal. Bioanal. Chem.*, 2020, **9**, 1–10.
- 25 M. Shi, Y. Li and H. Cui, *Micro Nano Lett.*, 2019, **14**, 1151–1156.
- 26 K. Khoshnevisan, H. Maleki and E. Honarvarfard, *Microchim. Acta*, 2019, **186**, 49.
- 27 D. Zhu, C. Xia, Z. Yang, T. Yang, Q. Li and R. Liu, *Electrochim. Acta*, 2020, **334**, 135591.
- 28 B. Zhang, J. Liu and J. Wang, *Nano Energy*, 2017, **37**, 74–80.
- 29 M. J. Sampaio and R. R. Bacsá, *J. Catal.*, 2015, **331**, 172–180.
- 30 S. Yang, G. Li and D. Wang, *Sens. Actuators, B*, 2017, **238**, 588–595.
- 31 H. Sakamoto, A. Koto and E. I. Takamura, *J. Nanosci. Nanotechnol.*, 2019, **19**, 3551–3557.
- 32 K. Kuroishi, T. Doi and K. Hoshi, *IEICE Trans. Electron.*, 2019, **102**, 159–163.
- 33 G. Slaughter and T. Kulkarni, *Microelectron. Eng.*, 2016, **149**, 92–96.
- 34 L. Yang, Y. Zhang and M. Chu, *Biosens. Bioelectron.*, 2014, **52**, 105–110.

