

Comparative study on various morphologies of WS₂ for electrochemical hydrogen evolution reaction

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Abstract

Nowadays, the advancement of cost-effective and efficient electrocatalysts holds significant importance in facilitating the progress of high-performance electrochemical hydrogen evolution reaction (HER). Tungsten disulfides (WS₂) are well recognized as highly promising electrocatalysts for HER due to their exceptional catalytic performance, outstanding stability, and comparatively affordable price. In this study, we used ammonium metatungstate hydrate and thioacetamide as precursors to fabricate WS₂ nanoflowers (NFs) by hydrothermal method, whereas WS₂-flakes were prepared from ammonium tetrathiotungstate through the thermolysis technique. These structures were evaluated for HER electrocatalytic activity in acidic media. The results revealed that WS₂-NFs exhibit higher catalytic activity than WS₂-flakes, with an overpotential of 325 mV to obtain a current density of 10 mA cm⁻² and a Tafel slope of 92.3 mV dec⁻¹. This performance was attributed to the larger electrochemically active surface area of WS₂-NFs compared to WS₂-flakes materials. The results of this study could bring the basis for further studies on the synthesis of WS₂-based catalytic materials for HER.

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Keywords

WS₂, electrocatalysts, hydrogen evolution reaction, nanoflowers, morphology

1 Introduction

The utilization of fossil fuels for energy generation has resulted in the emission of a substantial quantity of carbon dioxide (CO₂), leading to the phenomenon of climate change and environmental contamination. Hence, the pursuit of environmentally friendly and sustainable fuel alternatives is imperative on a global scale. Hydrogen is often regarded as a highly promising fuel source for future energy demands. Hydrogen exhibits superior energy density in comparison to conventional sources such as coal, methane, and gasoline [1-3]. Furthermore, the combustion of hydrogen yields water as a by-product, rendering it highly ecologically sustainable. At present, hydrogen can be generated by various methodologies, including coal gasification, methane gas reforming, and water

electrolysis [4-8]. Nevertheless, the initial two techniques release CO₂ during the hydrogen production process, exerting a detrimental impact on the environment. The electrochemical hydrogen evolution reaction (HER) process is often regarded as the most promising approach for hydrogen production because of its ability to meet the criteria of environmentally friendliness, efficiency, and sustainability. To date, the most optimal performance in the field of HER has been achieved by employing precious metal catalysts, namely platinum (Pt), palladium (Pd), and rhodium (Rh). [9-12]. However, the scarcity and high cost of these materials have prompted extensive research efforts, both experimental and theoretical, aimed at discovering alternative materials that are more affordable yet possess superior performance characteristics. For instance, transition metal sulfides,



selenides, and oxides have been investigated for HER. [13-15]. Among these electrocatalysts, transition metals sulfides are promising materials for hydrogen production by electrochemical method. It is worth noting that WS₂-based materials have been extensively examined as prospective alternatives to costly metal-based catalysts for the process of HER owing to their exceptional stability and superior performance. Several strategies have been suggested in previous literature to enhance the catalytic performance of WS₂, including doping, utilization of carbon-based support materials, and manipulation of the phase structure. For example, Sun et al. introduced nitrogen doping onto WS₂ nanosheets. The outcomes of their research demonstrated favorable results, as they achieved an overpotential for water splitting that was below 100 mV [16]. Lei et al. successfully synthesized WS₂/C composites that exhibit exceptional catalytic properties. These enhanced HER activities can be attributed to the higher conductivity and active surface area associated with the incorporation of carbon in the composite material [17]. Also, Lukowski and colleagues created phase 1T WS₂ from phase 2H-WS₂ with very high efficiency for HER [18]. Although WS₂-based catalysts for HER have been published, studies on the relationship between the morphological structures and catalytic activities of this material have not been systematically studied. In this study, various morphologies, including WS₂ nanoflowers (NFs) and WS₂- flakes forms were fabricated and evaluated for their catalytic activity for hydrogen production.

2 Materials and methods

2.1 Materials and chemicals

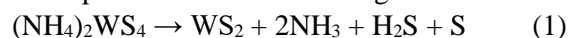
Ammonium metatungstate hydrate ($\geq 85\%$ WO₃ basis), thioacetamide (98 %), ammonium tetrathiotungstate (99.9 %), Nafion (5 %), và sulfuric acid (98 %) were provided by Sigma Aldrich.

2.2 Preparation of WS₂ with various morphologies

Preparation of WS₂-NFs. The aforementioned structure was fabricated via the hydrothermal technique. Compared to Hasani's study, WS₂-NFs were synthesized at a lower temperature and shorter reaction time [19-21]. Initially, a quantity of 8 g of thioacetamide was agitated within a glass vial that contained 40 mL of deionized water for 60 minutes. Subsequently, 8 g of ammonium metatungstate hydrate

was introduced into the mixture, followed by continuous stirring for 60 minutes. The mixture was placed into a Teflon autoclave reactor and subjected to a thermal treatment at a temperature of 180 °C for a period of 20 hours. After the reaction was finished, the reactor was subjected to a cooling process at 25 °C. Next, the solid was carefully separated and subsequently rinsed using centrifugation, repeating the process three times using deionized water. Finally, the WS₂ catalyst underwent a drying process at 100 °C in a vacuum environment for 12 hours, resulting in the production of the final powdered product.

Preparation of WS₂-flakes. This structure is produced by pyrolysis according to the following procedure [22]. A total of 8 g of ammonium tetrathiotungstate were placed in a ceramic crucible before being transferred to an inert gas furnace and heated to 900 °C for 2 hours. Following the end of the reaction, the furnace was then cooled down to ambient temperature. The obtained solid product was then isolated and subjected to a thorough washing process using centrifugation. This washing procedure involved three times of rinsing with deionized water. Finally, the WS₂ material was dried at 100 °C under vacuum for 12 hours. The formation of WS₂ was presented in the following:



2.3 Material characterization

Scanning electron microscopy (SEM) was conducted to identify various morphologies of WS₂ materials on Zeiss Sigma 300 equipment. The crystal structure of WS₂ materials was analyzed by the X-ray diffraction (XRD) method on equipment of the Bruker D8-Advance.

2.4 Catalytic activities evaluation

The electrocatalytic activities of the WS₂-NFs and WS₂-flakes were evaluated on a conventional electrochemical system in 0.5 M H₂SO₄ acidic medium. All electrochemical measurements were conducted on a potentiostat (Ivium 5612, Netherlands). The reference electrode used in the experiment was the saturated calomel electrode (SCE). The carbon rod was employed as the counter electrode. The fabrication of the working electrode is carried out under the following technique. The catalyst suspension was formed by combining 10 mg of WS₂ material with 5 mL of deionized water and 0.5 mL of Nafion, then the mixture was sonicated for 60 minutes at 25 °C to obtain the catalytic ink, after which 1 mL of this ink was coated to the surface of the glassy carbon

electrode and parched at 90 °C for 30 minutes to create a working electrode. During the electrochemical experiments, linear sweep voltammetry was recorded at a scan rate of 2 mV s⁻¹. All obtained potentials were converted by the formula, $E_{RHE} = E_0 + E_{SCE} + 0.059 \text{ pH}$. Electrochemical impedance spectroscopy (EIS) was implemented in a frequency range from 0.1 to 100,000 Hz. The stability of WS₂ was evaluated by chronoamperometry during 10 hours and cyclic voltammetry (CV) with 2000 cycles.

3 Results and discussions

The crystal structure of the WS₂ material was examined by XRD method as shown in Figure 1. WS₂-flakes catalysts have diffraction peaks at 14.3°, 32.7°, 39.5°, and 58.4° are designated the (002), (100), (103), and (110) crystal facets of the 2H-WS₂ hexagonal phase structure [19, 23, 24]. Moreover, the high-intensity peak of (002) revealed that WS₂-flakes were stacked together with a highly ordered packing, whereas WS₂-NFs displayed low-intensity peaks with the peak of the (002) crystal plane changed to diffraction angle position $2\theta \sim 13.8^\circ$. This result revealed that the number of stacks of (002) crystal planes was decreased for the WS₂-NFs sample. Besides, no strange peaks indicate the high purity of the generated WS₂ catalysts. Figure 2 illustrated the SEM images of the different morphologies of the WS₂ catalyst. Figure 2a displays the nanoflower morphology of the WS₂ material with a smaller size than the WS₂-flakes catalyst (Figure 2b). This result implies that WS₂-NFs give a higher active surface area than WS₂-flakes.

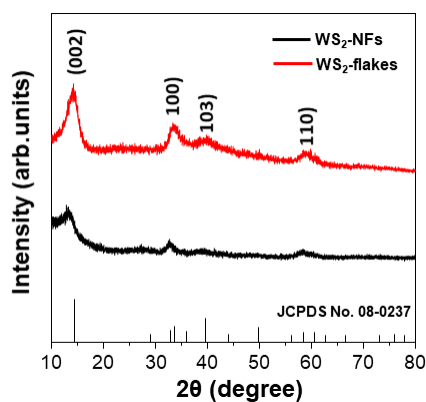


Figure 1 XRD patterns of WS₂-NFs and WS₂-flakes

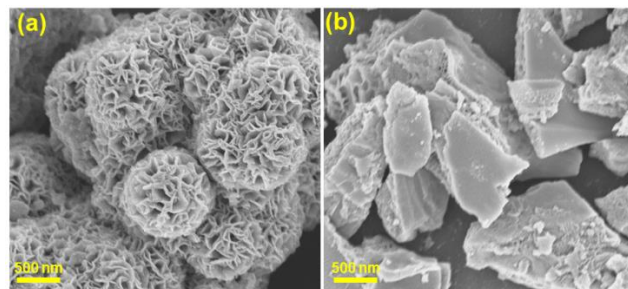
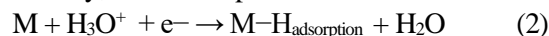
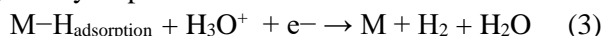


Figure 2 SEM of (a) WS₂-NFs and (b) WS₂-flakes

The HER electrocatalytic activity of the WS₂ structural morphologies was evaluated in an acidic medium. The polarization curves are shown in Figure 3a. This graph shows that the WS₂-NFs have a lower voltage requirement than the WS₂-flakes at 10 mA cm⁻². In particular, overpotential values are 325 mV and 397 mV for WS₂-NFs and WS₂-flakes, respectively. This result revealed that using a WS₂-flower catalyst would require lower energy to generate hydrogen by water electrolysis. Furthermore, the Tafel slope of the WS₂-NFs is 92.3 mV dec⁻¹ which is lower than that of the flakes WS₂ with a value of 98.1 mV dec⁻¹ (Figure 3b). This demonstrates that the WS₂-NFs catalyst has faster HER reaction kinetics than the WS₂-flakes catalyst. Because the values of the Tafel slope are in the range of 40-120 mV dec⁻¹. The HER mechanism of WS₂ catalysts follows the Volmer-Heyrovsky pathway [25]. Specifically, Volmer step:



Heyrovsky step:



M represents active sites on the surface of WS₂ materials. To confirm the reaction kinetics at the electrode and solution surfaces, impedance analysis was carried out at a specific potential of 200 mV, as shown in Figure 3c. The WS₂-NFs catalyst has a smaller Nyquist semicircle than the WS₂-flakes catalyst, indicating that the WS₂-NFs have smaller charge transfer resistance (R_{ct}) than the WS₂-flakes catalyst. Notably, the R_{ct} of the WS₂-NFs is 83.7 Ω, while the R_{ct} of the WS₂-flakes is 111.7 Ω. This result confirms that the WS₂-NFs catalyst has a better charge transfer than the WS₂-flakes catalyst, revealing a faster HER kinetic. The stability of catalysts is an important factor in assessing their practical applicability. Therefore, we examined the strength of WS₂-NFs and WS₂-flakes by chronoamperometric measurement and CV. Both WS₂-NFs and WS₂-flakes catalysts showed good stability after 10 hours of

continuous hydrogen generation. A slight decrease could be due to the covering of the catalytic active sites by the bubbles of hydrogen gas, as shown in Figure 3d. Moreover, the polarization curves of materials are only

slightly changed compared to the initial curves, as displayed in Figure 3e and f. Notably, the overpotential at 10 mA cm^{-2} is increased only by 2 mV and 5 mV after 2000 cycles for WS_2 -NFs and WS_2 -flakes, respectively.

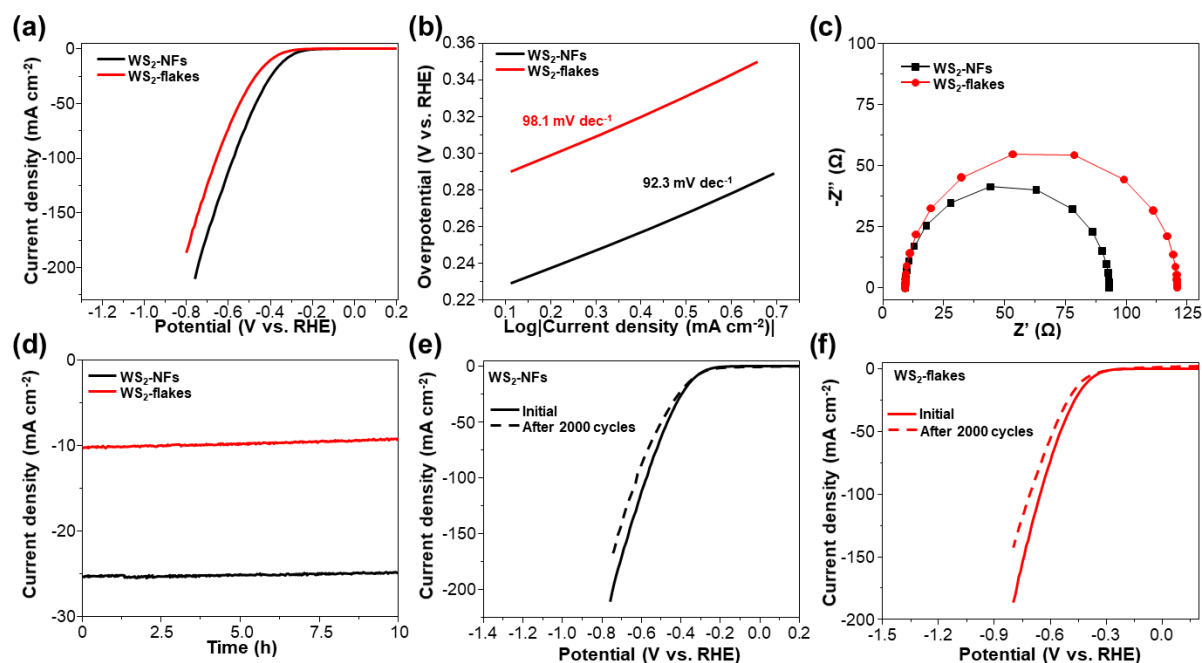
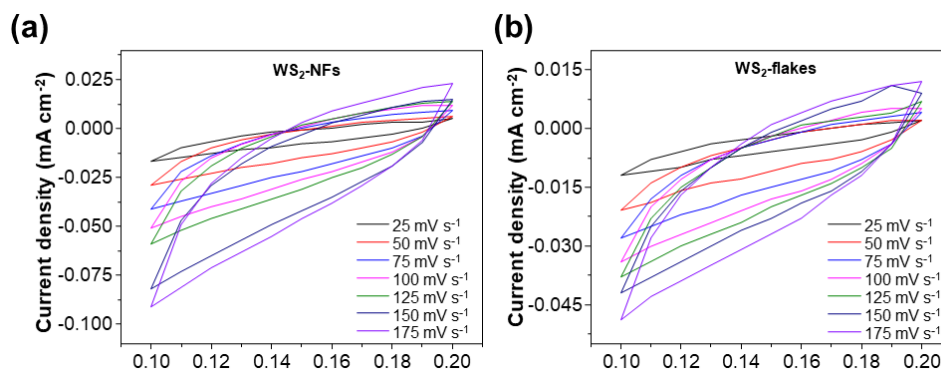


Figure 3 (a) Polarization curves of WS_2 -NFs, WS_2 -flakes, (b) Corresponding tafel slope of WS_2 -NFs, WS_2 -flakes, (c) Nyquist plots were recorded on WS_2 -NFs, WS_2 -flakes, (d) Chronoamperometric plots of WS_2 -NFs, WS_2 -flakes for 10 hours. Polarization curves were recorded initially and after 2000 CV cycles of (e) WS_2 -NFs and (f) WS_2 -flakes.

Electrochemical surface area (ESA) is an important parameter that can predict the electrocatalytic activity of materials. This value is usually calculated through the capacitance double layer (CDL), with the formula: $\text{ESA (cm}^2) = \text{CDL}/0.035$ [26]. CDL of WS_2 materials can be determined by measuring cyclic voltammetry at different scanning speeds, as exhibited in Figure 4a and b. The slope of the graph $\Delta J_{0.15}/2$ relative to the scan rate is the CDL of the material [27]. Herein, $\Delta J_{0.15}$ is the difference between anodic current density and cathodic current density at a potential of 0.15 V. The results

indicated that the CDL value of WS_2 -NFs in acidic media is 0.134 mF cm^{-2} , which is about two times higher than that of WS_2 -flakes, as shown in Figure 4c. This result implied that the WS_2 -NFs catalyst has a larger ESA than the WS_2 -flakes, as shown in Figure 4d. In particular, ESA values are 3.83 cm^2 and 2.17 cm^2 for WS_2 -NFs and WS_2 -flakes, respectively. This result could be the main reason for better HER catalytic activity of WS_2 -NFs material, as compared with WS_2 -flakes.



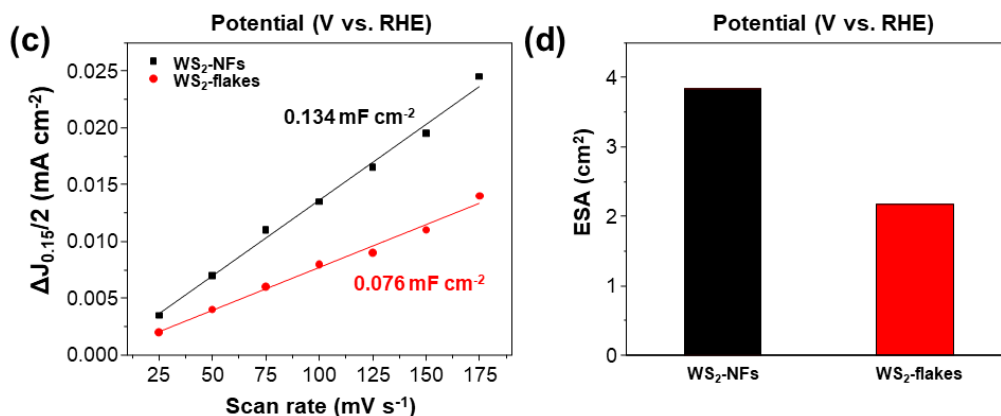


Figure 4 Cyclic voltammetry obtained in 0.5 M H₂SO₄ for (a) WS₂-NFs, (b) WS₂-flakes, (c) Extracted double-layer capacitances of WS₂-NFs, and WS₂-flakes, (d) Electrochemical surface area of WS₂-NFs, and WS₂-flakes.

The HER catalytic activities for WS₂-based electrocatalysts are provided for comparison, as shown in Table 1. The HER performance of WS₂-NFs and WS₂-flakes could be comparable with other materials such as WS₂ NF, and WS₂ NSs. This performance could be attributed to the high ESA and low Rct.

Table 1 Comparison of the HER performances of WS₂-NFs, WS₂-flakes, and against other HER electrocatalysts

Catalysts	Overpotential (mV)	Tafel slope (mV dec ⁻¹)	Reference
WS ₂ -NFs	325	92.3	This work
WS ₂ -flakes	397	98.1	This work
WS ₂ NF	400	173	[19]
WS ₂ NSs	160	72	[28]

4 Conclusion

In this study, different morphological structures of WS₂-NFs and WS₂-flakes were successfully fabricated

for HER reaction by hydrothermal and pyrolysis methods, respectively. The results of the comparison of catalytic activity revealed that WS₂-NFs have higher HER catalytic activity than WS₂-flakes. Specifically, the WS₂-NFs require only an overpotential of 325 mV to achieve a current density of 10 mA cm⁻², whereas this value is 397 mV for WS₂-flakes material. Moreover, the tafel slope of WS₂-NFs is smaller than that of WS₂-flakes, which implies a faster electrochemical reaction kinetic. These results could be explained by the fact that WS₂-NFs have a larger active surface area than WS₂-flakes catalysts. Besides, both these materials displayed outstanding stability in acidic electrolytes after 10 hours of testing.

Acknowledgments

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Nghiên cứu so sánh hoạt tính xúc tác trên các hình thái khác nhau của WS₂ cho phản ứng điện hóa tạo ra hydro

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Tóm tắt Ngày nay, sự phát triển của các chất xúc tác có độ bền cao và hiệu quả là rất quan trọng trong phản ứng tách nước tạo ra hydro (hydrogen evolution reaction – HER). Trong nghiên cứu này, ammonium metatungstate hydrate và thioacetamide được sử dụng như những tiền chất để chế tạo WS₂ hoa nano bằng phương pháp thủy nhiệt, trong khi WS₂ dạng mảnh được tạo ra từ ammonium tetrathiotungstate qua phản ứng nhiệt phân. Các cấu trúc này đã được đánh giá hoạt tính xúc tác điện hóa HER trong môi trường axit. Kết quả cho thấy WS₂ hoa nano thể hiện hoạt tính xúc tác cao hơn WS₂ dạng mảnh, với quá thế yêu cầu là 325 mV để thu được mật độ dòng 10 mA cm⁻² và một hệ số góc Tafel 92.3 mV dec⁻¹. Hiệu suất này được cho là diện tích bề mặt hoạt tính điện hóa của WS₂ hoa nano lớn hơn so với vật liệu WS₂ dạng mảnh. Kết quả nghiên cứu này tạo tiền đề cho những nghiên cứu sâu hơn về tổng hợp vật liệu xúc tác dựa trên WS₂ đối với HER.

Từ khóa WS₂, hoa nano, hydro, xúc tác điện hóa, HER