Molybdenum Carbide Anchored on N,S Co-Doped Carbon Composite Derived from Lignosulfonate as a High Performance Electrocatalyst for Hydrogen Evolution Reaction

Na Yeong Oh 1, So Young Park 1, Ji Young Hwang 1, Hyung Mo Jeong 2*, Yong Sik Kim 3,* and Duck Hyun Youn 1,*

1 Department of Chemical Engineering, Department of Integrative Engineering for Hydrogen Safety, Kangwon National University, Chuncheon 24341, Korea
2 School of Mechanical Engineering, Department of Smart Fab. Technology, Sungkyunkwan University, Suwon 16419, Korea
3 Department of Paper Science & Engineering, Kangwon National University, Chuncheon 24341, Korea
* Correspondence: yongsikk@kangwon.ac.kr (Y.S.K.); youndh@kangwon.ac.kr (D.H.Y.)

Abstract: A composite of Mo2C nanoparticles dispersed onto a nitrogen and sulfur co-doped carbon scaffold (Mo2C/N,S-C) was prepared by a simple and environmentally friendly method of one-pot annealing of MoCl5, urea, and lignosulfonate under a N2 atmosphere at 700 °C. Lignosulfonate, a by-product of the sulfite pulping process, was employed as a feedstock to fabricate the S-doped carbon scaffold and carbide simultaneously, and urea acted as a nitrogen source for N-doping to carbon. The as-prepared Mo2C/N,S-C catalyst showed high performance for the hydrogen evolution reaction (HER), with a small overpotential of 105 mV at 10 mAcm−2, and good stability for 3000 cycles. The improved HER performance of the Mo2C/N,S-C originated from the interplay between the highly active Mo2C nanoparticles and the N,S co-doped carbon scaffold with its high electrical conductivity and large surface area. Furthermore, N,S co-doping to carbon improved the hydrophilicity of the catalyst surface, thus further enhancing the HER activity.

Keywords: electrocatalyst; hydrogen evolution reaction; lignosulfonate; molybdenum carbide; N,S co-doped carbon

1. Introduction

Increasing energy consumption and environmental pollution have prompted the development of clean and renewable energy sources for the alleviation of human dependence upon exhaustible fossil fuels [1,2]. Hydrogen is recognized as an alternative energy source as it is abundant, clean, and energy-dense. Water electrolysis powered by renewable electricity is one of the most environmentally benign and sustainable hydrogen production technologies [3–5]. A hydrogen evolution reaction (HER) is the cathodic reaction of water electrolysis, wherein Pt is the most efficient electrocatalyst due to the negligible overpotential provided by the appropriate hydrogen binding energy of its surface [6]. However, the high cost and finite reserves of Pt limit large-scale application of water electrolysis. Therefore, the development of noble-metal-free electrocatalysts is of great importance [7,8].

Various molybdenum-based catalysts, including carbides (Mo2C) [9,10], sulfides (MoS2) [11,12], nitrides (MoN, Ni2Mo3N) [13,14], and phosphides (MoP) [15], have proven their good HER activity as possible replacements for Pt. In particular, molybdenum carbides are attracting tremendous attention due to their high activity and stability for HER from their Pt-like electronic structure, high electrical conductivity, and high chemical stability [16,17]. Since Vrubel and Hu revealed that commercial Mo2C possesses HER activity in both acidic and alkaline media [18], extensive research has been carried out to enhance the HER performance of Mo2C by fabricating nanostructured Mo2C catalysts and their composites with carbonaceous materials [19–26].
Next to cellulose, lignin is the second most abundant biopolymer and is generated as a by-product of the pulping process [27,28]. The lignin becomes sulfated during the sulfite pulping process, and the lignosulfonate is produced as a cross-linked polyphenolic polymer that contains sulfonic acid groups [29]. In contrast to hydrophobic lignin, lignosulfonate is an amphiphilic biopolymer and, as such, is soluble in water [30]. Small portions of produced lignosulfonate have been used as surfactants and adsorbents; however, most of it is unutilized and combusted for disposal producing carbon dioxide [31]. Thus, the effective utilization of lignosulfonate is required and inspired by its high carbon content and the presence of sulfur, we regard it as a suitable feedstock for the simultaneous fabrication of heteroatom-doped carbon and carbide.

Herein, sulfur and nitrogen co-doped carbon scaffolds decorated with molybdenum carbide nanoparticles (Mo$_2$C/N,S-C) were fabricated via the one-step pyrolysis of MoCl$_5$, urea, and lignosulfonate at 700 °C under N$_2$ flow for HER. During the synthesis, lignosulfonate plays critical roles as a source of sulfur for S-doping to carbon, and a source of carbon for the formation of amorphous carbon scaffold and carbide. At the same time, urea was employed as a nitrogen source for N-doping to carbon. Indeed, molybdenum carbide nanoparticles on S-doped carbon (Mo$_2$C/S-C) was fabricated without urea. The resultant Mo$_2$C/N,S-C catalysts exhibited excellent HER activity in alkaline solution with a low overpotential value of 105 mV at 10 mA cm$^{-2}$, which is better than Mo$_2$C/S-C and commercial Mo$_2$C (c-Mo$_2$C). Furthermore, the Mo$_2$C/N,S-C showed good stability for 3000 cycles. The enhanced HER performance of Mo$_2$C/N,S-C is due to the interplay between highly active Mo$_2$C nanoparticles and N,S-C scaffold providing high surface area and electrical conductivity. In addition, the enhanced HER performance of Mo$_2$C/N,S-C is assisted by the improved hydrophilicity of the N,S-C scaffold relative to S-C scaffold. The high electrocatalytic performance and the simple and environmentally friendly synthetic method suggest that the Mo$_2$C/N,S-C could be a promising catalyst for HER.

2. Materials and Methods

2.1. Mo$_2$C/N,S-C Synthesis

An amount of 160 mg lignosulfonate (Aldrich, Pittsburgh, PA, USA) was dissolved in 15 mL ethanol under magnetic stirring. One gram of MoCl$_5$ (Alfa aesar, Haverhill, MA, USA) was dispersed in 2.53 mL ethanol and added to the lignosulfonate solution under vigorous stirring for 30 min. Then, 109.9 mg urea was added as a nitrogen source, with a molar ratio (R) of 0.5 with respect to Mo and stirred for 1 h. After drying the solution in an oven at 100 °C for 90 min, the resultant mixture was annealed at 700 °C (3 °C min$^{-1}$ ramp) for 3 h under a N$_2$ atmosphere. As a control experiment, molybdenum carbide on a S-doped carbon (Mo$_2$C/S-C) catalyst was prepared by an identical method, except that 250 mg lignosulfonate was employed without the addition of urea. The weight contents of Mo$_2$C were measured as 35~40 wt.% for both the Mo$_2$C/N,S-C and Mo$_2$C/S-C samples.

2.2. Catalyst Characterization

Field-emission transmission electron microscopy (FE-TEM, JEOL, Akishima, Japan, JEM-2100F) and field-emission scanning electron microscopy (FE-SEM, JEOL, JSM-7900F) with energy dispersive X-ray spectrometer (EDS) were used to analyze morphologies and elemental compositions of the prepared samples. X-ray diffraction (XRD, Rigaku, Tokyo, Japan, MiniFlex 600) was conducted with Cu Kα (1.54 Å) radiation. X-ray photoelectron spectroscopy (XPS, Thermo-Scientific, Waltham, MA, USA, K-alpha) was used to investigate the chemical states of samples. The obtained binding energy values were calibrated by referencing the C 1s peak at 284.4 eV. The specific surface area and corresponding pore size distribution were investigated by measuring the N$_2$ adsorption–desorption isotherms at 77 K (Micromeritics, Norcross, GA, USA, ASAP 2020 PLUS). The contact angle measurements (Kruss, Kruss, Germany, Germany, DSA25) were conducted by loading the prepared catalysts (1 mg cm$^{-2}$) onto 1 × 1 cm carbon paper (1 wt.% wet-proofing, Toray, Tokyo, Japan, TGP-H-060).
2.3. Electrochemical Tests

All of the electrochemical measurements were conducted on a three-electrode electrochemical workstation (PAR, VersaSTAT 4) equipped with a rotating disk electrode (RDE, PINE Research) in a 1 M KOH aqueous solution. To prepare the working electrode, 20 mg of prepared catalyst was dispersed in 2 mL water and then 20 µL catalyst ink was loaded onto a glassy carbon electrode (0.19635 cm²). The reference and counter electrodes were Ag/AgCl (4 M KCl) electrode and Pt wire, respectively. All recorded potential values were converted to the reversible hydrogen electrode (RHE) with iR-compensation. Linear sweep voltammetry (LSV) curves were conducted at scan rate of 5 mV s⁻¹ with 900 rpm. Electrochemical impedance spectra (EIS) were performed in the frequency range from 100 kHz to 0.1 Hz at 105 mV (vs. RHE) overpotential with 6 mV modulation amplitude. Stability tests of prepared electrocatalysts were obtained by repeating 3000 cycles with potential range of 0.2 V to −0.2 V (vs. RHE). Electrochemical double layer capacitances (EDLC) were conducted by cyclic voltammetry (CV) from 0.1 to 0.3 V (vs RHE) at various scan rates of 20, 60, 100, 140 and 180 mV s⁻¹.

3. Results and Discussion

The synthesis of the Mo₂C/N,S-C catalyst is summarized in Scheme 1. Molybdenum chloride and urea was dissolved in ethanol solution containing the lignosulfonate and the subsequent annealing under N₂ at 700 °C yielded Mo₂C/N,S-C. During the synthesis, urea acted as a nitrogen source and lignosulfonate acted as carbon and sulfur sources. Notably, Mo₂C/S-C catalyst was generated without urea by an identical synthetic method, thereby demonstrating that that carbon sources from lignosulfonate played multiple roles in generating carbide and carbon scaffold at the same time. In addition, sulfur and nitrogen sources from lignosulfonate and urea served to S and N doping to carbon scaffold, respectively. The proposed synthetic method has the following advantages: (i) it is a simple method in which the formation of Mo₂C and the generation of amorphous carbon scaffold and N,S co-doping to the carbon scaffold are synchronously accomplished via mixing and annealing of the precursors in one pot, (ii) no toxic gases or chemicals were required for Mo₂C synthesis and N,S co-doping, (iii) the method is environmentally friendly due to the use of lignosulfonate, an industrial waste material, as a precursor, and (iv) our synthetic method produced Mo₂C nanoparticles with a size of ca. A total of 7 nm dispersed in N,S co-doped carbon scaffold (Mo₂C/N,S-C), which recorded one of the best performances for HER among the biomass-derived Mo₂C-based catalysts.

Scheme 1. Schematic illustration for the preparation of Mo₂C/N,S-C.

The SEM image of the Mo₂C/N,S-C in Figure 1a reveals the presence of amorphous carbon clusters of ~4 um in size, while the corresponding EDS elemental mapping images in Figure 1b–e reveal the almost identical distributions of molybdenum, carbon, nitrogen, and sulfur, respectively, thereby indicating that the Mo₂C nanoparticles are uniformly distributed on the N,S co-doped carbon scaffold. In the TEM image (Figure 1f), the Mo₂C nanoparticles are dispersed on amorphous carbon composite with an average particle size of 7 nm. The HR-TEM and fast Fourier transform (FFT) images are shown in Figure 1g. The lattice spacings of 0.226 and 0.237 nm are assigned to Mo₂C (101) and (002) crystalline...
planes, respectively (Figure 1g). In comparison, TEM images of Mo\textsubscript{2}C/S-C were also presented in Figure S1. The Mo\textsubscript{2}C nanoparticles are distributed on S-doped carbon scaffolds with a diameter of 16 nm. The lattice distance of 0.237 nm corresponds to the reflection of the (002) plane (Figure S1).

![Figure 1](image-url)

**Figure 1.** (a) SEM image of Mo\textsubscript{2}C/N,S-C and corresponding SEM-EDS elemental mapping for (b) Mo, (c) C, (d) N, and (e) S (scale bar = 10 um). (f) TEM and (g) HR-TEM images of Mo\textsubscript{2}C/N,S-C. Insets in (g) denote FFT images.

Figure 2a shows the XRD patterns for the Mo\textsubscript{2}C/N,S-C and Mo\textsubscript{2}C/S-C catalysts, which are consistent with their hexagonal β-Mo\textsubscript{2}C reference XRD patterns (JCPDS 00-035-0787). The peaks at 34.4°, 38.0°, 39.4°, 52.1°, 61.5°, 69.6°, 74.6°, 75.5°, 81.2° and 84.8° correspond to the (100), (002), (101), (102), (110), (103), (112), (201), (004), and (202) lattice planes, respectively. No impurity peaks were detected such as metallic molybdenum or molybdenum oxides for both catalysts.

The XPS survey spectrum for Mo\textsubscript{2}C/N,S-C (Figure S2) confirmed the existence of molybdenum, nitrogen, carbon, and sulfur elements on the catalyst surface. The amounts of N and S in Mo\textsubscript{2}C/N,S-C due to co-doping to carbon scaffold were determined to be 3.18 and 0.72 at.%, respectively, while Mo\textsubscript{2}C/S-C (Figure S3a) contained 0.49 and 0.9 at.% of N and the peaks for oxidized S was determined at 163.6/169.6 eV [35,36]. The Mo 3d and S 2p spectra of Mo\textsubscript{2}C/S-C were similar to those of Mo\textsubscript{2}C/N,S-C (Figure S3b,d), thereby suggesting similar chemical states except for the much lower content of N element (Figure S3c).
Nanomaterials 2022, 12, x FOR PEER REVIEW 5 of 10

η with (20 wt.%, E-TEK) and commercial Mo provided high surface area and electrical conductivity [35]. In addition, the co-doping of N previously reported biomass-derived Mo (BJH) method (Figure S4b).

The mesoporous texture of the Mo2C/N,S-C was revealed by the nitrogen adsorption-desorption results shown in Figure S4a, which exhibit a type IV isotherm. The Brunauer–Emmett–Teller (BET) surface area of Mo2C/N,S-C is 34.4 m² g⁻¹ and the presence of mesopores was also verified by pore size distribution (PSD) using the Barrett–Joyner–Halenda (BJH) method (Figure S4b).

Figure 3a shows polarization curves for the prepared catalysts including Pt/C (20 wt.%, E-TEK) and commercial Mo2C (c-Mo2C) in 1M KOH solution. The Pt/C exhibited the best HER activity with small overpotential of 30 mV to drive 10 mA cm⁻² (η₁₀ value) [14]. By contrast, the c-Mo2C with the η₁₀ value of 225 mV is not suitable as a HER electrocatalyst which might be due to its largely aggregated particles and low electrical conductivity [18]. In contrast, the Mo2C/S-C exhibited an improved HER activity with η₁₀ value of 141 mV than c-Mo2C, and this was significantly enhanced when the Mo2C phases are combined with the N,S co-doped carbon scaffold, achieving a η₁₀ value of 105 mV for the Mo2C/N,S-C. This improved performance is comparable to that of the previously reported biomass-derived Mo2C-based catalysts (Table S3). The coupling of Mo2C with N,S-C scaffold not only prevented agglomeration of Mo2C particles, but also provided high surface area and electrical conductivity [35]. In addition, the co-doping of N and S increases the charge and spin densities of carbon atoms compared to solely the doped carbon scaffolds, which results in a larger number of active carbon atoms [37]. Thereby, the N,S-C scaffold can further enhance the HER performance of Mo2C by modulating the catalytic activity of carbon atoms adjacent to heteroatoms [38]. In addition, previous studies have demonstrated that multi-heteroatom doping increases the surface wettability of the electrocatalysts, thereby promoting their HER activity [39–42]. Hence, the increased activity of the Mo2C/N,S-C relative to the Mo2C/S-C is further explained in terms of the surface wettability of the catalysts in Figure 4. The water contact angles of the Mo2C/N,S-C, Mo2C/S-C, and c-Mo2C are found to be 44°, 67°, and 85.3°, respectively, thereby demon-
biomass-derived Mo2C-based catalysts (Table S3). The coupling of Mo2C with N,S-C scaffold results in a larger number of active carbon atoms [37]. Thereby, the N,S-C scaffold can further enhance the HER performance of Mo2C by modulating the catalytic activity of carbon atoms adjacent to heteroatoms [38]. In addition, previous studies have demonstrated that multi-heteroatom doping increases the surface wettability of the electrocatalysts, thereby promoting their HER activity [39–42]. Hence, the increased activity of Mo2C/N,S-C relative to the Mo2C/S-C is further explained in terms of the surface wettability of the catalysts in Figure 4. The water contact angles of the Mo2C/N,S-C, Mo2C/S-C, and c-Mo2C are found to be 44°, 67°, and 85.3°, respectively, thereby demonstrating that hydrophilicity was greatly enhanced by the introduction of N,S co-doped carbon scaffold. Hydrophilicity can facilitate the HER activity by lowering the adhesion force and facilitating the detachment of gas bubbles from the catalyst surface [35,43].

The cyclic voltammogram (CV) curves of the catalysts in the non-faradaic region are presented in Figure 3c, where the charge transfer resistance (Rct) at the electrode and electrolyte interface is represented by a semicircle and is inversely proportional to the electrocatalytic activity [20,44]. The Rct value of Mo2C/N,S-C (12.4 Ω), which is smaller than that of Mo2C/S-C (35.1 Ω) and c-Mo2C (200 Ω), thereby implies rapid electron transfer and improved HER activity.

The Nyquist plots obtained from the electrochemical impedance spectroscopy (EIS) of the catalysts are presented in Figure 3c, where the charge transfer resistance (Rct) at the electrode and electrolyte interface is represented by a semicircle and is inversely proportional to the electrocatalytic activity [20,44]. The Rct value of Mo2C/N,S-C (12.4 Ω), which is smaller than that of Mo2C/S-C (35.1 Ω) and c-Mo2C (200 Ω), thereby implies rapid electron transfer and improved HER activity.

The Tafel plots, (c) Nyquist plots with an equivalent circuit, and (d) stability tests of Mo2C/N,S-C.

**Figure 3.** (a) Polarization curves of all prepared samples in 1 M KOH solution, (b) Tafel plots, (c) Nyquist plots with an equivalent circuit, and (d) stability tests of Mo2C/N,S-C.

**Figure 4.** Contact angle measurement for (a) Mo2C/N,S-C, (b) Mo2C/S-C, and (c) c-Mo2C.

Figure 3b shows the Tafel plots of the prepared catalysts, fitted to Tafel equation ($\eta = b \log|J| + a$, where $b$ is the Tafel slope and $J$ is the current density). The Tafel slope of Pt/C is 36 mV dec$^{-1}$, which is consistent with the previously reported value [20]. Meanwhile, the Tafel slope of the Mo2C/N,S-C is 56 mV dec$^{-1}$, which is smaller than that of the Mo2C/S-C (75 mV dec$^{-1}$) and the c-Mo2C (110 mV dec$^{-1}$), thereby suggesting the occurrence of the Volmer–Heyrovsky mechanism along with the faster HER kinetics in the Mo2C/N,S-C.

The Nyquist plots obtained from the electrochemical impedance spectroscopy (EIS) of the catalysts are presented in Figure 3c, where the charge transfer resistance (Rct) at the electrode and electrolyte interface is represented by a semicircle and is inversely proportional to the electrocatalytic activity [20,44]. The Rct value of Mo2C/N,S-C (12.4 Ω), which is smaller than that of Mo2C/S-C (35.1 Ω) and c-Mo2C (200 Ω), thereby implies rapid electron transfer and improved HER activity.

The cyclic voltammogram (CV) curves of the catalysts in the non-faradaic region are provided in Figure S5a–c, and the corresponding double-layer capacitance (Cdl) values
are shown in Figure S5d. The Mo$_2$C/N,S-C catalysts exhibited a higher C$_{dl}$ value of 18.02 mF cm$^{-2}$ than Mo$_2$C/S-C (5.47 mF cm$^{-2}$) and Mo$_2$C (1.34 mF cm$^{-2}$). In general, C$_{dl}$ is proportional to the contact area between catalyst and electrolyte. Accordingly, the contact area of Mo$_2$C/N,S-C is larger than other catalysts, which additionally contributes to the higher HER activity [25].

The polarization curves of the Mo$_2$C/N,S-C obtained before and after 3000 cycles between 0.2 and −0.2 V (vs. RHE) are presented in Figure 3d. This reveals the good stability of the catalyst in alkaline media, with little change in the polarization curve, and a marginal increase in the $\eta_{10}$ value from 105 to 117 mV, after 3000 cycles. Since durability is a significant factor in determining the HER performance, the highly active and durable Mo$_2$C/N,S-C has clear potential as a HER electrocatalyst.

4. Conclusions

In summary, a simple and environmentally friendly method to produce Mo$_2$C nanoparticles dispersed onto N,S co-doped carbon scaffold was designed by using the lignosulfonate, an industrial waste material. The as-prepared Mo$_2$C/N,S-C catalyst exhibited a high HER performance with a small $\eta_{10}$ value of 105 mV and a good stability for 3000 cycles. The improved HER performance resulted from a synergy between the highly active Mo$_2$C nanoparticles and N,S co-doped carbon scaffold, thus providing high electrical conductivity and large surface area. Compared to solely doped carbon scaffolds, N,S-C scaffold can further enhance the HER performance of Mo$_2$C by adjusting the catalytic activity of carbon atoms adjacent to heteroatoms. In addition, N,S co-doping to carbon modulated the hydrophilicity of the catalyst surface, thereby further enhancing the HER activity. Thus, considering this simple and environmentally friendly method, the proposed Mo$_2$C/N,S-C could be a promising HER catalyst with a high activity and stability.

Supplementary Materials: The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/nano12173047/s1, Figure S1: (a) TEM and (b) HR-TEM images for Mo$_2$C/S-C; Figure S2: XPS survey scan of Mo$_2$C/N,S-C; Figure S3: XPS spectra of Mo$_2$C/S-C. (a) Survey, (b) Mo 3d (c) N 1s, and (d) S 2p; Figure S4: (a) N$_2$ sorption isotherm and (b) pore size distribution of Mo$_2$C/N,S-C; Figure S5: CV graphs of (a) Mo$_2$C/N,S-C, (b) Mo$_2$C/S-C and (c) Mo$_2$C measured at scan rates from 20 to 180 mV s$^{-1}$ between potential range of 0.1 and 0.3 V (vs. RHE) in 1 M KOH. (d) measured capacitive currents at 0.2 V (vs. RHE) as a function of scan rate; Table S1: Element contents of lignosulfonate; Table S2: The amounts of each element for Mo$_2$C/N,S-C and Mo$_2$C/S-C; Table S3: Comparison of HER performance in alkaline media with various reported biomass-derived molybdenum carbide-based catalysts. References [45–53].

Author Contributions: Conceptualization, Y.S.K. and D.H.Y.; methodology, N.Y.O. and D.H.Y.; validation, Y.S.K. and D.H.Y.; formal analysis, N.Y.O.; investigation, N.Y.O., S.Y.P. and J.Y.H.; resources, Y.S.K. and D.H.Y.; data curation, N.Y.O., S.Y.P., J.Y.H. and H.M.J.; writing—original draft preparation, N.Y.O., S.Y.P., J.Y.H., H.M.J., Y.S.K. and D.H.Y.; writing—review and editing, N.Y.O. and D.H.Y.; visualization, N.Y.O. and D.H.Y.; supervision, D.H.Y.; project administration, Y.S.K. and D.H.Y.; funding acquisition, Y.S.K. and D.H.Y. All authors have read and agreed to the published version of the manuscript.

Funding: This study was carried out with the support of the R&D Program for Forest Science Technology (FTIS2020216B10-2222-AC01) provided by the Korea Forest Service (Korea Forestry Promotion Institute) and the National Research Foundation of Korea (NRF) grant funded by the Korean government (Ministry of Education) (2019R1I1A3A01052741). This work was also supported by the Korean Institute of Energy Technology Evaluation and Planning (KETEP) and the Ministry of Trade, Industry & Energy (MOTIE) of the Republic of Korea (No. 20224000000080).

Institutional Review Board Statement: Not applicable.

Informed Consent Statement: Not applicable.

Data Availability Statement: All data is contained within the article.
Acknowledgments: The central laboratory of Kangwon National University and Korea Basic Science Institute (Chuncheon) provided significant assistance with the TEM analyses.

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

References
1. Yu, Z.-Y.; Duan, Y.; Feng, X.-Y.; Yu, X.; Gao, M.-R.; Yu, S.-H. Clean and affordable hydrogen fuel from alkaline water splitting: Past, recent progress, and future prospects. Adv. Mater. 2021, 33, 2007100. [CrossRef]
2. Dresselhaus, M.S.; Thomas, I.L. Alternative energy technologies. Nature 2001, 414, 332–337. [CrossRef]
3. Lin, Z.; Xiao, B.; Huang, M.; Yan, L.; Wang, Z.; Huang, Y.; Shen, S.; Zhang, Q.; Gu, L.; Zhong, W. Realizing negatively charged metal atoms through controllable d-electron transfer in ternary Ir1-xRh,xSb intermetallic alloy for hydrogen evolution reaction. Adv. Energy Mater. 2022, 12, 2208553. [CrossRef]
4. Buttler, A.; Spliethoff, H. Current status of water electrolysis for energy storage, grid balancing and sector coupling via power-to-gas and power-to-liquids: A review. Renew. Sust. Energ. Rev. 2018, 82, 2440–2454. [CrossRef]
5. Kim, J.Y.; Magesh, G.; Youn, D.H.; Jang, J.-W.; Kubota, J.; Domen, K.; Lee, J.S. Single-crystalline, wormlike hematite photoanodes for efficient solar water splitting. Sci. Rep. 2013, 3, 2681. [CrossRef]
6. Sheng, W.; Myint, M.; Chen, J.G.; Yan, Y. Correlating the hydrogen evolution reaction activity in alkaline electrolytes with the hydrogen binding energy on monometallic surfaces. Energy Environ. Sci. 2013, 6, 1509–1512. [CrossRef]
7. Wang, J.; Xu, F.; Jin, H.; Chen, Y.; Wang, Y. Non-noble metal-based carbon composites in hydrogen evolution reaction: Fundamentals to applications. Adv. Mater. 2017, 29, 1605838. [CrossRef]
8. Tang, C.; Zhang, R.; Lu, W.; Wang, Z.; Liu, D.; Hao, S.; Du, G.; Asiri, A.M.; Sun, X. Energy-saving electrolytic hydrogen generation: Ni2P nanoarray as a high-performance non-noble-metal electrocatalyst. Angew. Chem. Int. Ed. 2017, 56, 842–846. [CrossRef]
9. Wu, C.; Li, J. Unique hierarchical Mo2C/C nanosheet hybrids as active electrocatalyst for hydrogen evolution reaction. ACS Appl. Mater. Interfaces 2017, 9, 41314–41322. [CrossRef]
10. Ge, C.; Jiang, P.; Cui, W.; Pu, Z.; Xing, Z.; Asiri, A.M.; Obaid, A.Y.; Sun, X.; Tian, J. Shape-controllable synthesis of Mo2C nanostructures as hydrogen evolution reaction electrocatalysts with high activity. Electrochim. Acta 2014, 134, 182–186. [CrossRef]
11. Youn, D.H.; Jang, J.-W.; Kim, J.Y.; Jang, J.S.; Choi, S.H.; Lee, J.S. Fabrication of graphene-based electrode in less than a minute through hybrid microwave annealing. Sci. Rep. 2014, 4, 5492. [CrossRef]
12. Voiry, D.; Salehi, M.; Silva, R.; Fujita, T.; Chen, M.; Asefa, T.; Shenoy, V.B.; Eda, G.; Chhowalla, M. Conducting MoS2 nanostructures as hydrogen evolution reaction catalysts for hydrogen evolution reaction. Nano Lett. 2013, 13, 6222–6227. [CrossRef]
13. Xie, J.; Li, S.; Zhang, X.; Zhang, J.; Wang, R.; Zhang, H.; Pan, B.; Xie, Y. Atomically-thin molybdenum nitride nanosheets with exposed active surface sites for efficient hydrogen production. Chem. Sci. 2014, 5, 4615–4620. [CrossRef]
14. Park, S.H.; Jo, T.H.; Lee, M.H.; Kawashima, K.; Mullins, C.B.; Lim, H.-K.; Youn, D.H. Highly active and stable nickel-molybdenum nitride (Ni2Mo3N) electrocatalyst for hydrogen evolution. J. Mater. Chem. A 2021, 9, 4945–4951. [CrossRef]
15. Lee, M.H.; Youn, D.H.; Lee, J.S. Nanostructured molybdenum phosphide/N-doped carbon nanotube-graphene composites as efficient electrocatalysts for hydrogen evolution reaction. Appl. Catal. A Gen. 2020, 594, 117451. [CrossRef]
16. Miao, M.; Pan, J.; He, T.; Yan, Y.; Xia, B.Y.; Wang, X. Molybdenum carbide-based electrocatalysts for hydrogen evolution reaction. Chem. Eur. J. 2017, 23, 10947–10961. [CrossRef]
17. Vaar, C.; Regmi, Y.N.; Leonard, B.M. Multiple phases of molybdenum carbide as electrocatalysts for the hydrogen evolution reaction. Angew. Chem. Int. Ed. 2014, 53, 6407–6410. [CrossRef]
18. Vrabel, H.; Hu, X. Molybdenum boride and carbide catalyze hydrogen evolution in both acidic and basic solutions. Angew. Chem. Int. Ed. 2012, 51, 12703–12706. [CrossRef]
19. Ma, Y.; Guan, G.; Hao, X.; Cao, J.; Abdula, A. Molybdenum carbide as alternative catalyst for hydrogen production—A review. Renew. Sust. Energ. Rev. 2017, 75, 1101–1129. [CrossRef]
20. Jing, S.; Zhang, L.; Luo, L.; Lu, J.; Yin, S.; Shen, P.K.; Tsiakaras, P. N-doped porous molybdenum carbide nanobelts as efficient catalysts for hydrogen evolution reaction. Appl. Catal. B 2018, 224, 533–540. [CrossRef]
21. Li, J.-S.; Wang, Y.; Liu, C.-H.; Li, S.-L.; Wang, Y.-G.; Dong, L.-Z.; Dai, Z.-H.; Li, Y.-F.; Lan, Y.-Q. Coupled molybdenum carbide and reduced graphene oxide electrocatalysts for efficient hydrogen evolution. Nat. Commun. 2016, 7, 11204. [CrossRef]
22. He, C.; Tao, J. Synthesis of nanostructured clean surface molybdenum carbides on graphene sheets as efficient and stable hydrogen evolution reaction catalysts. Chem. Commun. 2015, 51, 8323–8325. [CrossRef] [PubMed]
23. Lee, G.H.; Lee, M.H.; Kim, Y.; Lim, H.-K.; Youn, D.H. Facile synthesis of nanostructured molybdenum carbide/nitrogen-doped CNT-RGO composite via a modified urea glass route for efficient hydrogen evolution. J. Alloys Compd. 2019, 805, 113–119. [CrossRef]
24. Liu, Y.; Yu, G.; Li, G.D.; Sun, Y.; Asefa, T.; Chen, W.; Zou, X. Coupling Mo5C with nitrogen-rich nanocarbon leads to efficient hydrogen-evolution electrocatalytic sites. Angew. Chem. Int. Ed. 2015, 54, 10752–10757. [CrossRef]
25. Jo, H.M.; Kim, Y.; Youn, D.H. One-pot synthesis of molybdenum carbide/N-doped carbon nanotube composite using nitroltri-acetic acid for hydrogen evolution. J. Alloys Compd. 2021, 855, 157420. [CrossRef]
26. Oh, N.Y.; Lee, G.H.; Jeong, H.M.; Kim, Y.S.; Youn, D.H. Kraft lignin derived molybdenum carbide/nitrogen-doped carbon composite for efficient hydrogen evolution reaction. J. Electrochem. Soc. 2021, 168, 084511. [CrossRef]
27. Vanholme, R.; Demedts, B.; Morreel, K.; Ralph, J.; Boerjan, W. Lignin biosynthesis and structure. Plant Physiol. 2010, 153, 895–905. [CrossRef]
28. Chio, C.; Sain, M.; Qin, W. Lignin utilization: A review of lignin depolymerization from various aspects. Renew. Sust. Energ. Rev. 2019, 107, 232–249. [CrossRef]
29. Mansouri, N.-E.E.; Salvadó, J. Structural characterization of technical lignins for the production of adhesives: Application to lignosulfonate, kraft, soda-anthraquinone, organosolv and ethanol process lignins. Ind. Crops Prod. 2006, 24, 8–16. [CrossRef]
30. Kong, Q.; Yu, J.; Song, W.; Fan, X.; Li, Y.; Zhang, F.; Peng, W. Ultra-small MoCo quantum dot embedded chitosan-derived nitrogen-doped carbon nanosheets derived from peanut root nodules as highly efficient non-metal electrocatalyst for hydrogen evolution reaction. Nanoscale 2019, 12, 3047–3053. [CrossRef] [PubMed]
31. Rinaldi, R.; Jastrzebski, R.; Clough, M.T.; Ralph, J.; Kennema, M.; Bruijnincx, P.C.A.; Weckhuysen, B.M. Paving the way for lignin valorisation: Recent advances in bioengineering, biorefining and catalysis. Angew. Chem. Int. Ed. 2016, 55, 8164–8215. [CrossRef]
52. Kang, Q.; Qin, Y.; Lu, Q.; Gao, F. Waste leather-derived (Cr, N)-co-doped carbon cloth coupling with Mo2C nanoparticles as a self-supported electrode for highly active hydrogen evolution reaction performances. J. Power Sources 2020, 476, 228706. [CrossRef]

53. Chen, X.; Sun, J.; Guo, T.; Zhao, R.; Liu, L.; Liu, B.; Wang, Y.; Li, J.; Du, J. Biomass-derived carbon nanosheets coupled with MoO2/Mo2C electrocatalyst for hydrogen evolution reaction. Int. J. Hydrogen Energy 2022. [CrossRef]