

MOS₂-BASED CATALYST FOR HYDROGEN PRODUCTION IN PEM ELECTROLYSIS

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ABSTRACT

Developing low-cost electrocatalysts is key to improving proton exchange membrane (PEM) water electrolysis for sustainable hydrogen production. In the research for alternative catalyst materials, molybdenum disulfide (MoS₂) has shown great potential for the hydrogen evolution reaction (HER). This study investigates MoS₂-based catalysts synthesized via a hydrothermal method, including an unsupported variant and two supported on Ketjenblack EC carbon (KB). Physico-chemical characterization was conducted using different techniques, while electrochemical performance was evaluated in a PEM single cell through polarization, electrochemical impedance spectroscopy (EIS), and durability tests. The supported MoS₂ catalyst, incorporating KB during synthesis, exhibited superior performance with higher current densities, lower polarization resistance and higher durability, demonstrating its potential as a viable alternative to noble metal-based cathodes.

Keywords: PEM electrolyzer, non-Noble catalysts, MoS₂, Hydrogen, Proton Exchange Membrane

1. INTRODUCTION

The increasing global energy demand and the environmental impact of CO₂ emissions from fossil fuel combustion make the research for sustainable alternative energies crucial. Hydrogen is considered one of the most promising options, offering a more sustainable solution with a lower environmental impact in the context of modern energy consumption, and its combustion results exclusively in water [1]. Among the available technologies, proton exchange membrane (PEM) water electrolysis is distinguished by its efficiency and reliability, though the reliance on noble metal catalysts such as platinum and iridium limit its economic sustainability. In the context of today's scientific efforts to advance hydrogen production, there is a growing focus on developing non-precious catalysts to make water electrolysis more accessible and cost-effective [2].

Molybdenum disulfide (MoS₂) has emerged as a promising alternative due to its abundance and favorable catalytic properties, offering a viable pathway for sustainable hydrogen production.

2. MATERIALS AND METHODS

2.1. Catalyst preparation

Three MoS₂ catalysts were synthesized for cathode applications using a suitable and low-cost hydrothermal method: one unsupported and two supported on Ketjenblack EC carbon (KB). The difference between the two supported catalysts lies in the preparation: one was mechanically mixed with KB (supported-1), while in the other, KB was incorporated with the precursors (supported-2) to enhance catalytic efficiency while reducing the metal catalyst loading.

2.2. Physico-chemical characterization

To investigate the physico-chemical properties of the synthesized electrocatalysts, various characterization techniques were employed. X-ray Diffraction (XRD) was used to analyze the crystal structure and to evaluate crystallite size, while X-ray Fluorescence (XRF) was performed to determine the elemental composition of the catalysts. Transmission Electron Microscopy coupled with Energy-Dispersive X-ray Spectroscopy (TEM-EDX) allowed to know the powders morphology, structure, and elemental distribution. Additionally, Thermogravimetric Analysis (TGA) was conducted to monitor mass variations as a function of temperature under controlled atmospheric conditions.

2.3. PEM electrolyser setup

The electrochemical performances of the synthesized MoS₂ and MoS₂/C catalysts were investigated in a single PEM cell with a 5 cm² active area by polarization curves and electrochemical impedance spectroscopy (EIS) in a range of temperature between ambient to 90°C; galvanostatic durability tests were performed at an operating temperature of 80°C. The cells were assembled using a commercial IrO₂ as the anode and the three synthesized MoS₂ catalysts as the cathodes. A NAFION 212 membrane was placed between the electrodes to complete the PEM-based MEA assembly.

3. RESULTS AND DISCUSSION

The comparative analysis of the three catalysts revealed that the supported-2 MoS₂ catalyst exhibited superior electrochemical performance. In polarization measurements, it achieved higher current densities at lower voltages compared to both the unsupported and supported-1. Electrochemical impedance spectroscopy (EIS) further confirmed

lower polarization resistance at different voltage values, which suggests enhanced charge transfer efficiency; additionally, galvanostatic durability showed greater stability.

These findings confirmed the supported-2 MoS₂ as the most operative cathode catalyst for hydrogen evolution reaction in PEM electrolysis.

4. CONCLUSIONS

In this work is reported a screening of catalysts based on MoS₂ and MoS₂/C for hydrogen evolution reaction (HER) in acid environment. Non-CRM (non critical raw material) catalysts were synthesized by hydrothermal processes and characterized by physico chemical analyses, as XRD, XRF, TGA and TEM. The electrochemical characterization was carried out in terms of polarization curve, EIS at different potential value and galvanostatic durability test. The electrochemical evaluation of MoS₂ and MoS₂/C as cathodes in a PEM single cell revealed improved performance and stability for the supported MoS₂ catalysts, particularly for the one incorporating Ketjenblack EC carbon in the initial step with precursors.

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