Controllable Growth and On-Site Domain Boundary Imaging of Monolayer MoS₂ on Au foils and Its Potential Application in Hydrogen Evolution Reaction

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Abstract

Controllable synthesis of monolayer MoS_2 is the basic premise both for exploring some fundamental physical issues, and for engineering its applications in nanoelectronics, optoelectronics, etc. Herein, we report the scalable growth of domain size tunable (edge length from~ 200 nm to 80 µm), strictly monolayer MoS₂ flakes or even complete films on commercially available Au foils, via low pressure chemical vapor deposition (LPCVD) method. By introducing H₂ as carrier gas, we report the successful synthesis of large domain monolayer MoS2 triangular flakes on Au foils, with the edge length approaching to 80 µm. The growth process is proposed to be mediated by two competitive effects with H_2 acting as both a reduction promoter for efficient sulfurization of MoO₃ and an etching reagent of resulting MoS₂ flakes. By using low-energy electron microscopy/diffraction, we have further identified the crystal orientations and domain boundaries of MoS₂ flakes directly on Au foils for the first time. Of particular interesting, the nanosized triangular MoS₂ flakes on Au foils are proved to be excellent electrocatalysts for hydrogen evolution reaction (HER), featured by a rather low Tafel slope (61mV/decade) and a relative high exchange current density (38.1µA/cm²). The excellent electron coupling between MoS₂ and Au foils is considered to account for the extraordinary hydrogen evolution reaction activity. These on-site and transfer-free characterizations should shed light on the initial growth and the aggregation of MoS₂ on arbitrary substrates, further guiding the growth towards large domain flakes or monolayer films. And the synthesis of monolayer MoS₂ with introducing metal foils as substrates presents a sound proof that monolayer MoS₂ assembled on a well selected electrode can manifest comparable HER property with that of nanoparticles or few-layer MoS₂ electrocatalysts.

References

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