Promoting Alkaline Hydrogen Evolution Activity by Ordered Mesoporous Metastable *a*- MoC_{1-x} with Enhanced Water Dissociation Capability

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The sluggish kinetics of the alkaline hydrogen evolution reaction (HER) remains an important challenge for water-alkali electrolyzers. In this work, we demonstrate that metastable, face-centered-cubic a-MoC_{1-x} phase showed superior water dissociation capability and alkaline HER activity than stable, hexagonal-close-packed β -Mo₂C phase. Based on this, we prepared ordered mesoporous a-MoC_{1-x} (MMC) via a nanocasting method. In MMC structure, the a-MoC_{1-x} phase facilitates the water dissociation reaction, while the mesoporous structure enables a high dispersion of metal nanoparticles (NPs) and efficient mass transport. As a result, Pt NPs supported on MMC (Pt/MMC) catalyst showed enhanced alkaline HER activity in terms of overpotentials, Tafel slopes, mass and specific activities, and exchange current densities, compared to commercial Pt/C and Pt NPs supported on particulate a-MoC_{1-x} or β -Mo₂C. Notably, Pt/MMC showed very low

Tafel slope of 30 mV dec⁻¹, suggesting the critical role of MMC in enhancing the HER kinetics. The promotional effect of MMC was further demonstrated with an Ir/MMC catalyst.