Titanium Carbide MXenes with Different Terminations as Platinum Supports for Fuel Cells

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MXenes possess attractive hydrophilicity, metallic conductivity, and stability for application in electrocatalysis. Herein, we assess promising  $T_{i3}C_2$  as durable and active support candidates in Pt-loaded catalytic systems to drive the 4 electron-driven oxygen reduction reaction (ORR) in the advancement of proton exchange membrane fuel cells (PEMFC). Following the dealumination treatment of corresponding MAX parent materials in the presence of HF, the resulting MXene phase revealed interplanar mesoporosity. Specific electrochemical properties have drawn from the presence of distinct termination groups on the surface of  $T_{i3}C_2$ . However, the effect of these surface functional groups, which may play a important role to optimize the interaction with loaded Pt nanoparticles, has not been properly investigated to date. In this work, we direct our attention to the influence of these surface functional groups on the resulting interactions with Pt to improve the electrocatalytic activity and stability of Pt-loaded  $T_{i3}C_2$  materials toward ORR in PEMFC.