

Compositional Dependence of Hydroxylated Indium-Oxide on the Reaction Rate of CO₂/H₂ Reduction

Authors : Joel Y. Y. Loh, Geoffrey A. Ozin, Charles A. Mims, Nazir P. Kherani

Abstract : A major goal in the emerging field of solar fuels is to realize an 'artificial leaf' - a material that converts light energy in the form of solar photons into chemical energy - using CO₂ as a feedstock to generate useful chemical species. Enabling this technology will allow the greenhouse gas, CO₂, emitted from energy and manufacturing production exhaust streams to be converted into valuable solar fuels or chemical products. Indium Oxide (In₂O₃) with surface hydroxyl (OH) groups have been shown to reduce CO₂ in the presence of H₂ to CO with a reaction rate of 15 $\mu\text{mol gcat}^{-1} \text{h}^{-1}$. The likely mechanism is via a Frustrated Lewis Pair sites heterolytically splitting H₂ to be absorbed and form protonic and hydric sites that can dissociate CO₂. In this study, we investigate the dependence of oxygen composition of In₂O₃ on the CO₂ reduction rate. In₂O₃-x films on quartz fiber paper were DC sputtered with an Indium target and varying O₂/Ar plasma mixture. OH surface groups were then introduced by immersing the In₂O₃-x samples in KOH. We show that hydroxylated In₂O₃-x reduces more CO₂ than non-hydroxylated groups and that a hydroxylated and higher O₂/Ar ratio sputtered In₂O₃-x has a higher reaction rate of 45 $\mu\text{mol gcat}^{-1} \text{h}^{-1}$. We show by electrical resistivity-temperature curves that H₂ is adsorbed onto the surface of In₂O₃ whereas CO₂ itself does not affect the indium oxide surface. We also present activation and ionization energy levels of the hydroxylated In₂O₃-x under vacuum, CO₂ and H₂ atmosphere conditions.

Keywords : solar fuels, photocatalysis, indium oxide nanoparticles, carbon dioxide

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