## Tuning adsorption properties of Ga<sub>x</sub>In<sub>2-x</sub>O<sub>3</sub> catalysts for enhancement of methanol synthesis activity from CO<sub>2</sub>hydrogenation at high reaction temperature

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## ABSTRACT

Light olefins can be produced from CO<sub>2</sub> hydrogenation in a single reactor using a combination of a methanol synthesis catalyst and a methanol-to-olefin (MTO) catalyst. However, commercial methanol synthesis catalysts are active at low temperatures (200–260 °C), while MTO reaction is feasible at higher temperatures (>300 °C). Herein, we report the CO<sub>2</sub> hydrogenation to methanol at high temperatures (320–400 °C) over Ga<sub>x</sub>In<sub>2-x</sub>O<sub>3</sub> catalysts. By tuning the Ga/In ratios, phase, crystallinity, pore structure, morphology, electronic properties as well as adsorptive properties of Ga<sub>x</sub>In<sub>2-x</sub>O<sub>3</sub> catalysts can be modified. At the lowest temperature (320 °C), the pure In<sub>2</sub>O<sub>3</sub> shows the highest methanol yield. However, the maximum methanol yield declines significantly with increasing reaction temperatures. Incorporation of Ga into the In<sub>2</sub>O<sub>3</sub> crystal lattices at x = 0.4 (Ga<sub>0.4</sub>In<sub>1.6</sub>O<sub>3</sub>) maximizes the methanol yield at higher reaction temperatures of 340–360 °C. This enhancement can be attributed to an increased binding energy of adsorptive molecules with the catalyst surface to promote the hydrogenation of CO<sub>2</sub> to methanol. Further increasing Ga content (x > 0.4) leads to greatly strengthen the binding for adsorptive molecules, resulting in a lower methanol yield and the formation of methane. The surface chemisorbed oxygen is found to be a key factor determining the CO yield.

## **KEYWORDS**

CO<sub>2</sub> hydrogenation; Methanol; Indium; Gallium

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