

Supporting Information for

A highly active Au/In₂O₃-ZrO₂ catalyst for selective hydrogenation of CO₂ to methanol

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Table S1. Comparison of Au catalysts for CO₂ hydrogenation to methanol.

Catalysts	Pressure / MPa	Temperature/ K	GHSV/ cm ³ h ⁻¹ g _{cat} ⁻¹	X _{CO₂} / %	S _{MeOH} / %	STY/ g _{MeOH} h ⁻¹ g _{cat} ⁻¹	Ref.
Au/In ₂ O ₃ -ZrO ₂	5.0	523	21,000	5.4	88.2	0.27	This work
Au/In ₂ O ₃ -ZrO ₂	5.0	548	21,000	10.1	83.1	0.48	This work
Au/In ₂ O ₃ -ZrO ₂	5.0	573	21,000	14.8	70.1	0.59	This work
Au/ZrO ₂	5.0	573	21,000	26.0	13.3	0.20	This work
Au/In ₂ O ₃ -ZrO ₂ (3: 1)	5.0	573	21,000	11.3	72.9	0.47	This work
Au/In ₂ O ₃ -ZrO ₂ (6: 1)	5.0	573	21,000	14.8	66.0	0.56	This work
Au/In ₂ O ₃	5.0	573	21,000	11.7	67.8	0.47	[1]
Au/TiO ₂	0.5	513	24,000	13.1	0.6	0.0015	[2]
Au/CeO ₂	0.1	498	20,000	~1	~37.0	0.0047	[3]
Au/ZnO	5.0	513	9,000	1.0	70.0	0.014	[4]
Au/ZnO-ZrO ₂	8.0	493	3,300	2	100.0	0.019	[5]
Au/Al ₂ O ₃	0.5	493	24,000	2	3.8	0.00048	[2]

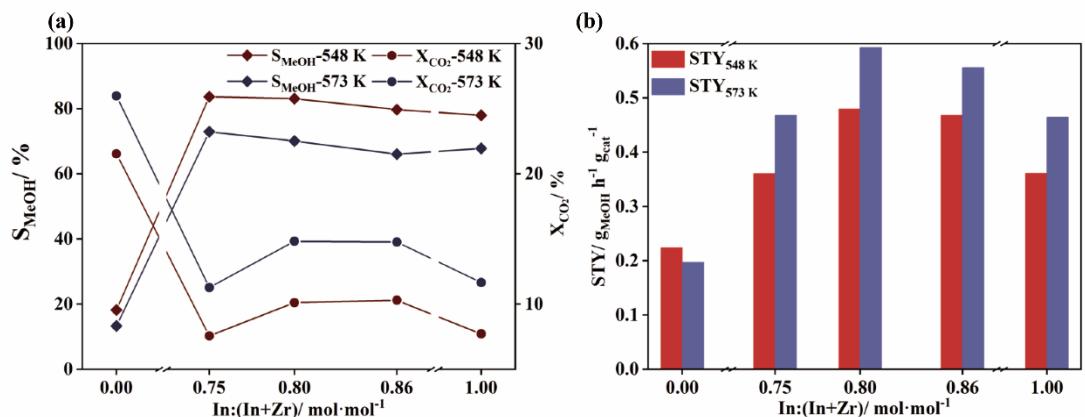


Figure S1. (a) CO_2 conversion and methanol selectivity, and (b) methanol space–time yield (STY) respect to the molar In content, which is equal to $m/(m+n)$ for the $\text{Au}/\text{In}_2\text{O}_3\text{-ZrO}_2$ ($m: n$) sample. Reaction conditions: 5 MPa, $\text{H}_2/\text{CO}_2/\text{N}_2 = 76/19/5$, GHSV = 21,000 $\text{cm}^3 \text{h}^{-1} \text{g}_{\text{cat}}^{-1}$.

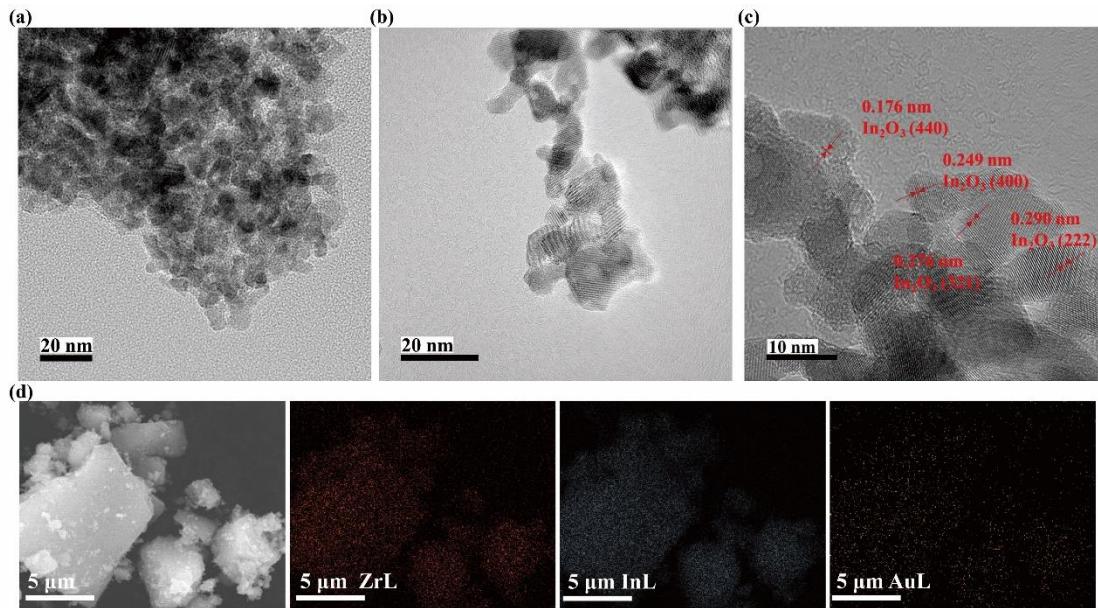


Figure S2. TEM images of Au/ In_2O_3 -ZrO₂ (a), Au/ In_2O_3 -ZrO₂-AR (b) and Au/ In_2O_3 -ZrO₂-AR (c); SEM image (d) and the corresponding EDX mapping of Au/ In_2O_3 -ZrO₂-AR.

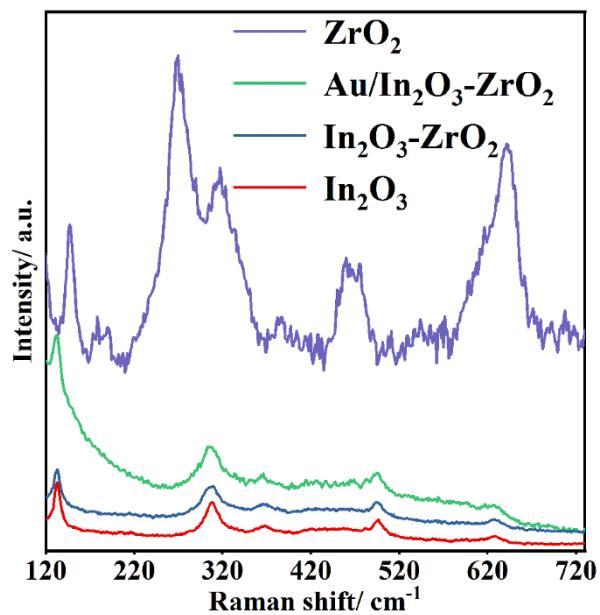


Figure S3. Visible Raman spectra of fresh ZrO₂, Au/In₂O₃-ZrO₂, In₂O₃-ZrO₂, and In₂O₃ samples.

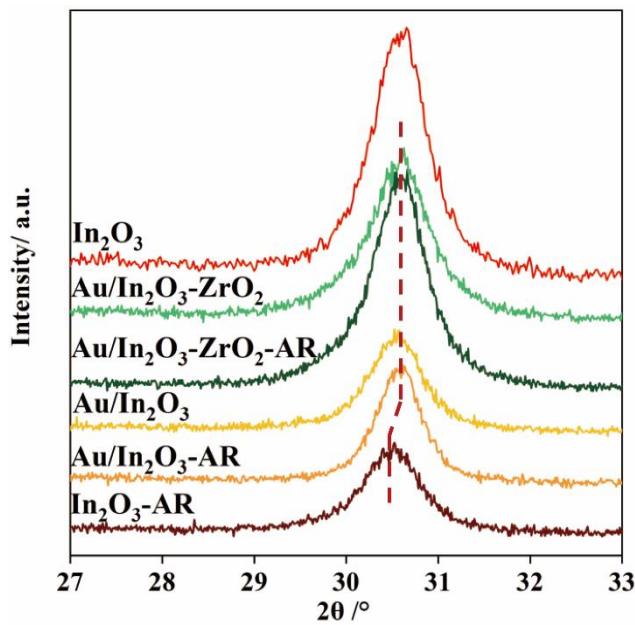


Figure S4. Enlarged XRD patterns of peaks around 32° for fresh samples and the samples after reaction.

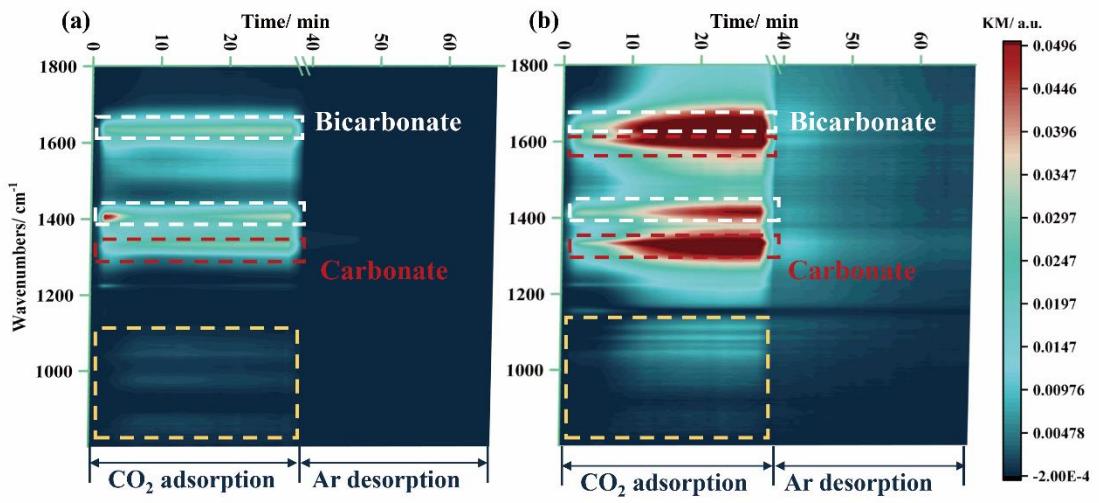


Figure S5. Time-resolved *in-situ* DRIFTS as a function of CO₂ adsorption time and Ar desorption time at 323 K after pretreatment in Ar over Au/In₂O₃ (a) and Au/In₂O₃-ZrO₂ (b). Symbol definition: bicarbonate [6] (white frame), carbonate [6-7] (red frame), CO₂ absorbed on the vacancies featuring the higher CO₂ adsorption energy [8] (yellow frame).

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