

Supplementary Material

Catalytic Conversion of High Fructose Corn Syrup to Methyl Lactate with CoO@silicalite-1

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Materials and Chemicals

Polyvinylpyrrolidone and Co(NO₃)₂·6H₂O (99%) were purchased from Aladdin Chemicals (Shanghai, China). Silica gel (30 wt% in water) and citric acid monohydrate (99.5%) were bought from Macklin reagent (Shanghai, China). Methyl lactate (98%), tetrapropylammonium hydroxide (25 wt% in water), and methanol (AR) were bought from the Sinopharm Chemical Reagent (China). Methoxyacetaldehyde dimethyl acetal (98%) was purchased from J & K Scientific (Beijing, China). Glycolaldehyde dimethyl acetal (98%) and HFCS-55 were supplied by Alfa Aesar and Qianfengxiang Food Co., Ltd (Wuhan, Hubei), respectively.

Detailed preparation of catalyst

The synthesis of c-CoO@silicalite-1 via a steam-assisted method followed the procedure of our group [1] with some modifications. Briefly, a certain amount of Co(NO₃)₂·6H₂O and 0.1 g PVP were dissolved in 30 mL water and stirred until the solution become transparent. Then, under stirring 3 g silica gel (30 wt% in water) was added and the mixture was stirred for 10 min. Then 1.5 g tetrapropylammonium hydroxide (TPAOH, 25% in water) was added into the mixture. After they were stirred at room temperature for 15 min the solution was rotary evaporated at 50 °C. After removing all the water, the dry gel was placed in the 50 mL Teflon cup and that was placed into a 100 mL Teflon-lined autoclave. A certain amount of water was added outside the Teflon cup (the mass ratio of water to dry gel is 1:5). Afterwards, the autoclave was heated to 130 °C for 3 days. The obtained solid was washed for three times with ethanol, water, and ethanol, respectively. And then the solid was dried under 60 °C for 12 h followed by calcination at 550 °C for 6 h to remove residual template. Before catalytic reaction the zeolite was only pretreated with citric acid (0.5 M, 70 °C, 100 mL/g zeolite, 2 h), then washed with water and dried for 12 h. Finally, the solid sample was calcined at 550 °C for 6 h to obtain c-CoO@silicalite-1. A series of synthesized catalysts was identified as c-Co_x, where x is the amount of Co(NO₃)₂·6H₂O (mmol) required in the synthesis of the catalyst. The pictures of the Teflon-lined autoclave and the Teflon cup were shown in Figure S1.

Figure S1

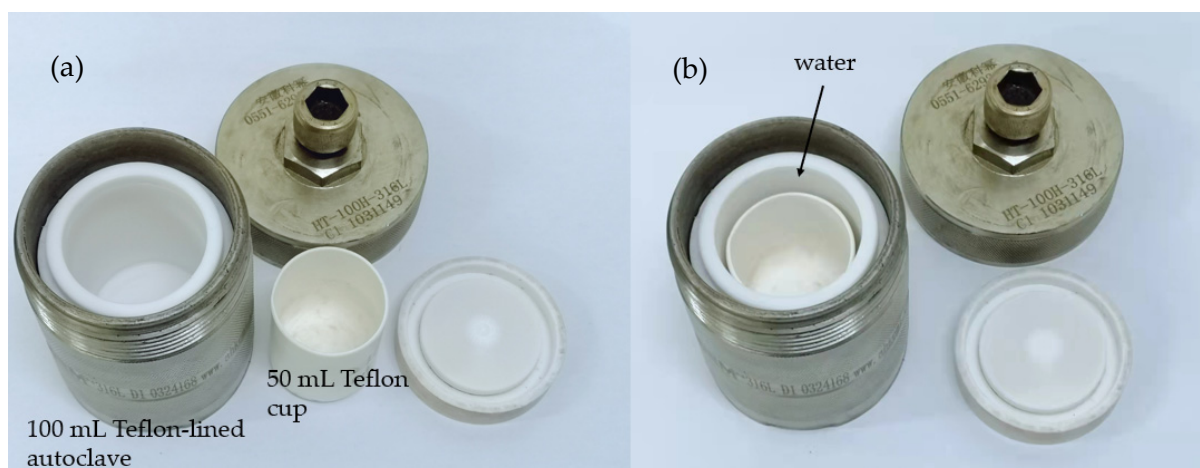


Figure S1. The Teflon-lined autoclave (a) and the Teflon cup (b).

Figure S2

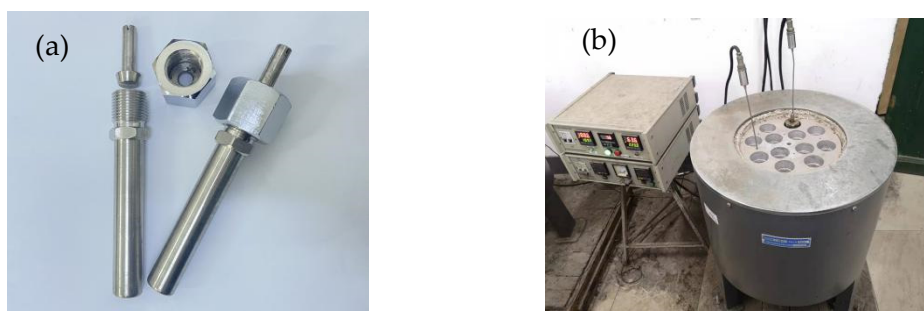


Figure S2. The photos of the stainless steel reactor (a) and heating furnace (b).

Table S1

Table S1. Composition of HFCS-55 detected by HPLC

	Content	Relative content of fructose and glucose
fructose	41.1%	55.9%
glucose	32.4%	44.1%

Figure S3

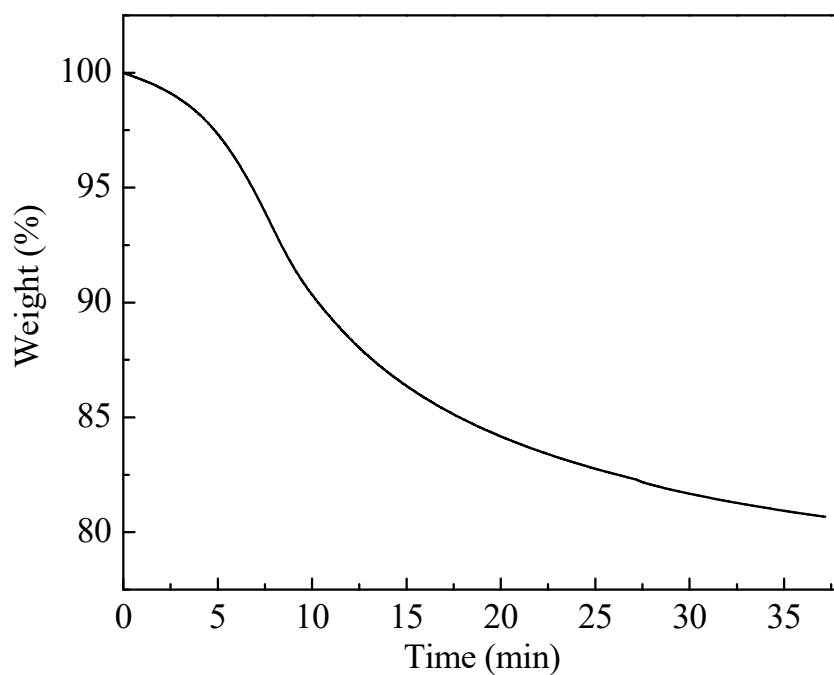


Figure S3. TG analysis of HFCS-55.

Figure S4

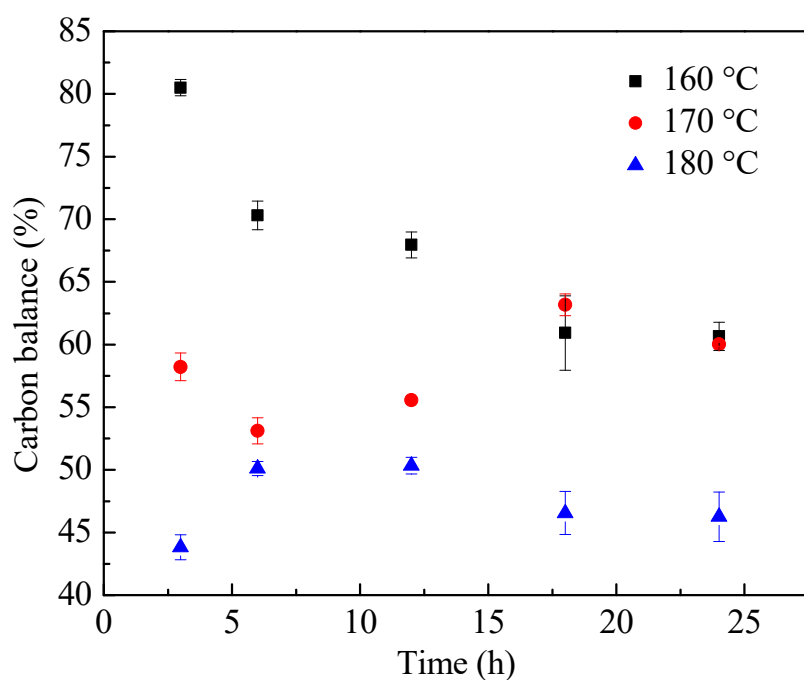


Figure S4. The carbon balance of the catalytic conversion of HFCS-55 under different reaction temperatures and time.

References

1. Yan, Y.; Zhang, Z.; Bak, S.-M.; Yao, S.; Hu, X.; Shadike, Z.; Do-Thanh, C.-L.; Zhang, F.; Chen, H.; Lyu, X.; et al. Confinement of Ultrasmall Cobalt Oxide Clusters within Silicalite-1 Crystals for Efficient Conversion of Fructose into Methyl Lactate. *ACS Catal.* **2018**, *9*, 1923–1930. <https://doi.org/10.1021/acscatal.8b03230>