

# Supporting Information

Efficient Cascade Conversion of Glucose to Levulinic Acid using Dual-Functional UiO-66 Catalyst

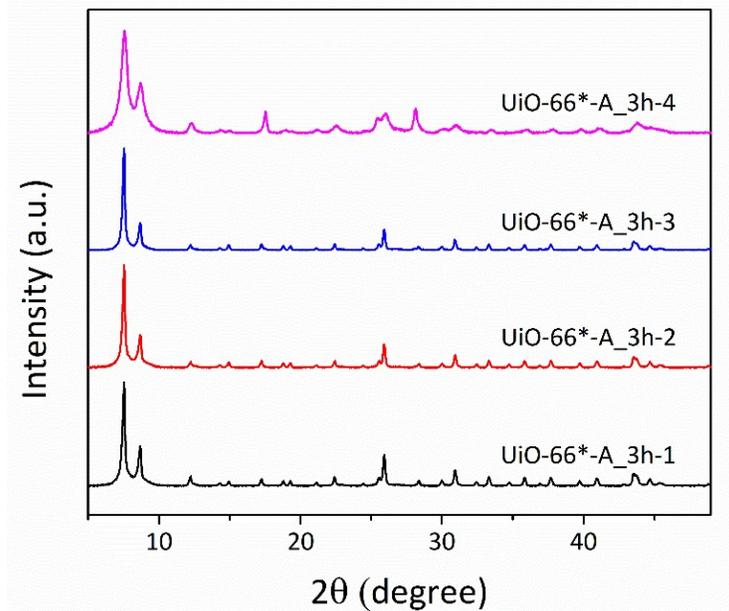
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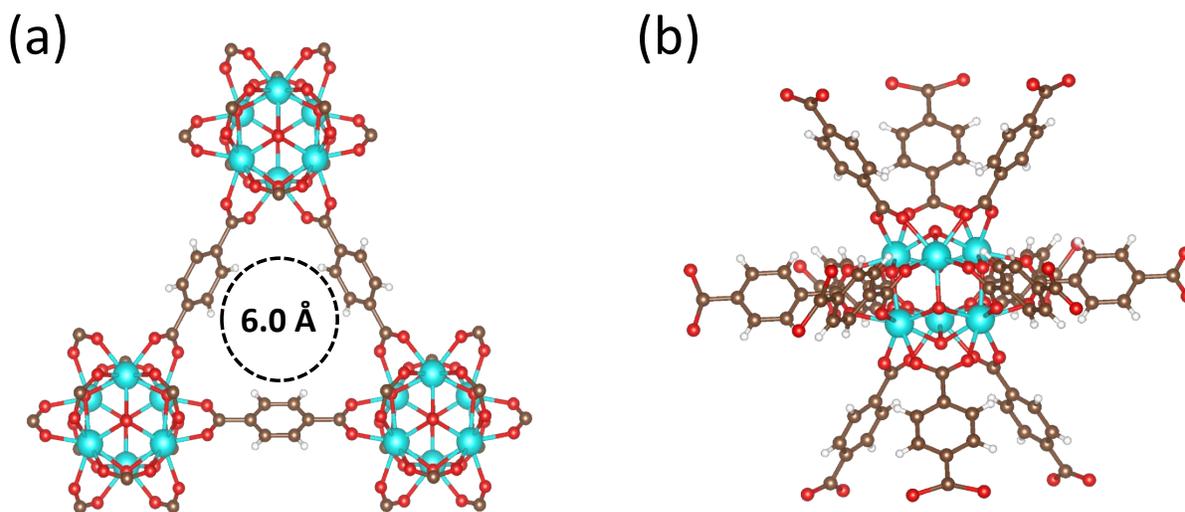
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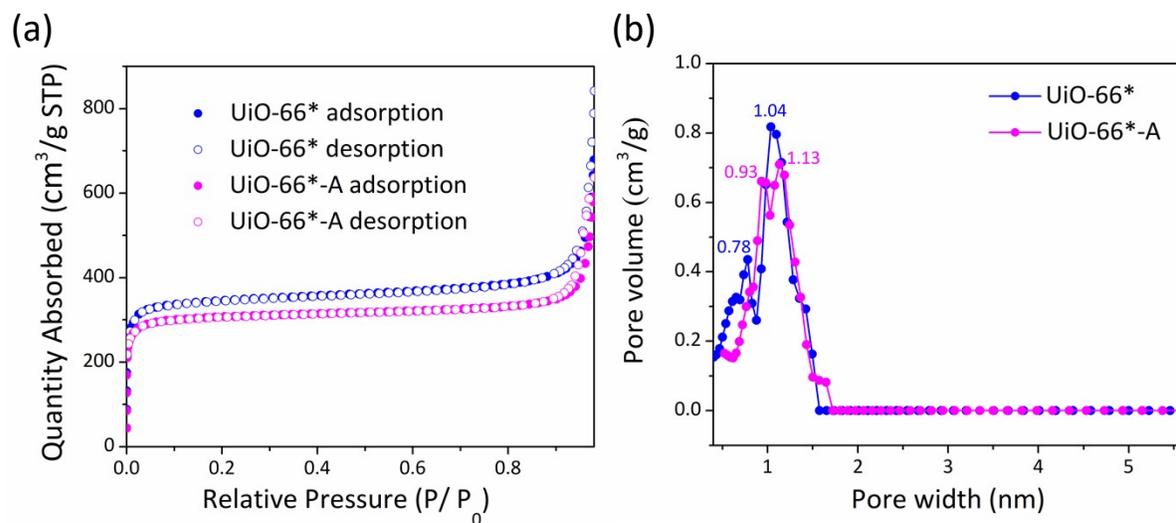
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**Figure S1.** Powder X-ray diffraction (PXRD) patterns of the UiO-66\*-A after acid treatment. The crystallinity of UiO-66\*-A remained stable up to the 3<sup>rd</sup> cycle.



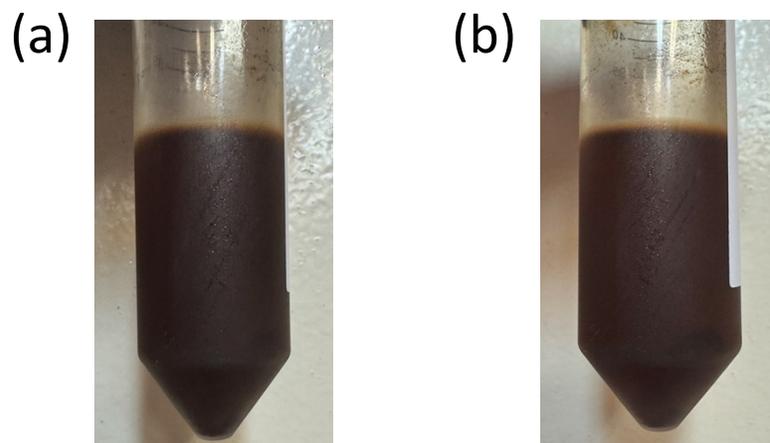
**Figure S2.** Structural illustration of UiO-66 presenting: (a) a pore window with the size of *c.a.* 6.0 Å, and (b) building unit of  $Zr_6O_4(OH)_4(BDC)_{12}$ . (Color: Zr, blue; O, red; C, brown; H, white).



**Figure S3.** (a)  $N_2$  sorption isotherms and (b) pore size distribution of the UiO-66\* and UiO-66\*-A measured at 77 K.

**Table S1.** Summary surface areas and pore volumes of UiO-66\* and UiO-66\*-A calculated from  $N_2$  sorption isotherms using Brunauer–Emmett–Teller (BET) and t-plot analysis.

Sample	Specific surface area ( $m^2 g^{-1}$ )	Total pore volume ( $cm^3 g^{-1}$ )
UiO-66*	1382.6	0.4746
UiO-66*-A	1234.3	0.4344

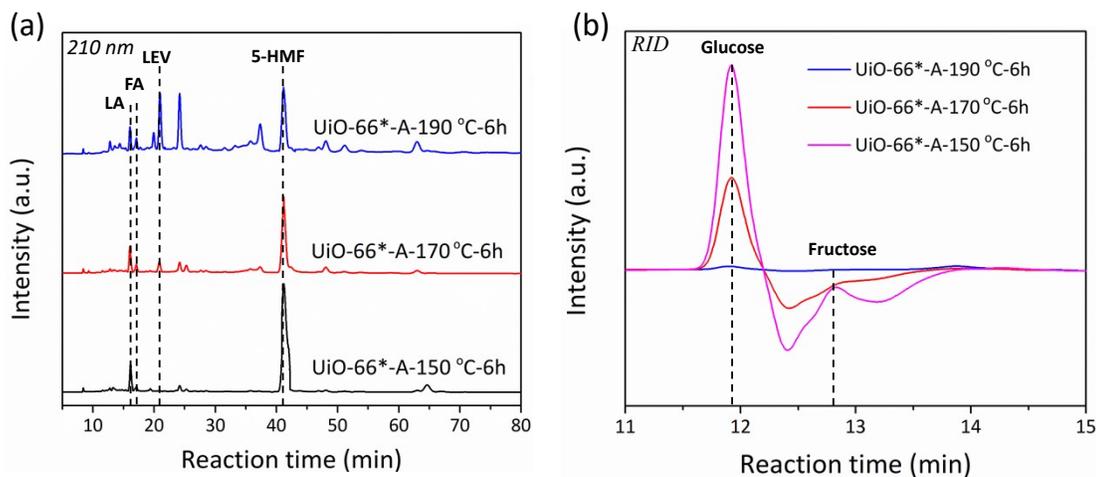


**Figure S4.** Photographs of an absence of catalyst (a)without and (b)with NaCl salt after hydrothermal reactions at 190 °C for 6 h, respectively.

**Table S2.** Conversions and yields for the catalytic conversion of glucose into LEV on UiO-66\*-A catalyst over 6 h at various reaction temperatures.

Reaction temperature (°C)	Glucose conversion (%)	Fructose (mol)	LEV (mol)	5-HMF (mol)	LA (mol)
150	64.66 ± 0.52	0.05 ± 0.00	0.01 ± 0.00	0.09 ± 0.00	0.04 ± 0.00
170	83.58 ± 0.13	0.02 ± 0.00	0.19 ± 0.01	0.07 ± 0.00	0.06 ± 0.00
190	>99.00	<i>n.d.</i> <sup>a</sup>	0.58 ± 0.01	0.06 ± 0.00	0.05 ± 0.01

<sup>a</sup>n.d.= not detected



**Figure S5.** HPLC chromatograms from (a) UV (210 nm) and (b) RID detectors of the supernatants from the reactions employing UiO-66\*-A as the catalyst at various temperatures for 6 h.

: This figure illustrates HPLC chromatograms obtained from UV (210 nm) and RID detectors of the supernatants. The chromatograms from UV detector quantify the resulting LEV, HMF, and LA from the reaction, while the results from the RID detector provide information on glucose and fructose. At a low reaction temperature of 150 °C, a prominent fructose peak is observed (0.05 mol, Table S2). As the reaction temperature increases to 170 and 190 °C, only trace amounts of fructose can be observed. These results indicate that the UiO-66\*-A catalyst efficiently converts glucose to fructose before dehydration and hydrolysis processes, which ultimately lead to LEV production.

**Table S3.** Conversions and yields of the catalytic performance of UiO-66\*-A for 190 °C at various reaction times.

Reaction time (h)	Glucose conversion (%)	LEV (mol)	5-HMF (mol)	LA (mol)
2	72.71 ± 0.59	0.28 ± 0.00	0.04 ± 0.00	0.04 ± 0.01
4	84.67 ± 0.57	0.41 ± 0.01	0.09 ± 0.01	0.08 ± 0.01
6	>99.00	0.58 ± 0.00	0.06 ± 0.00	0.05 ± 0.01
9	>99.00	0.37 ± 0.00	0.01 ± 0.00	0.03 ± 0.01

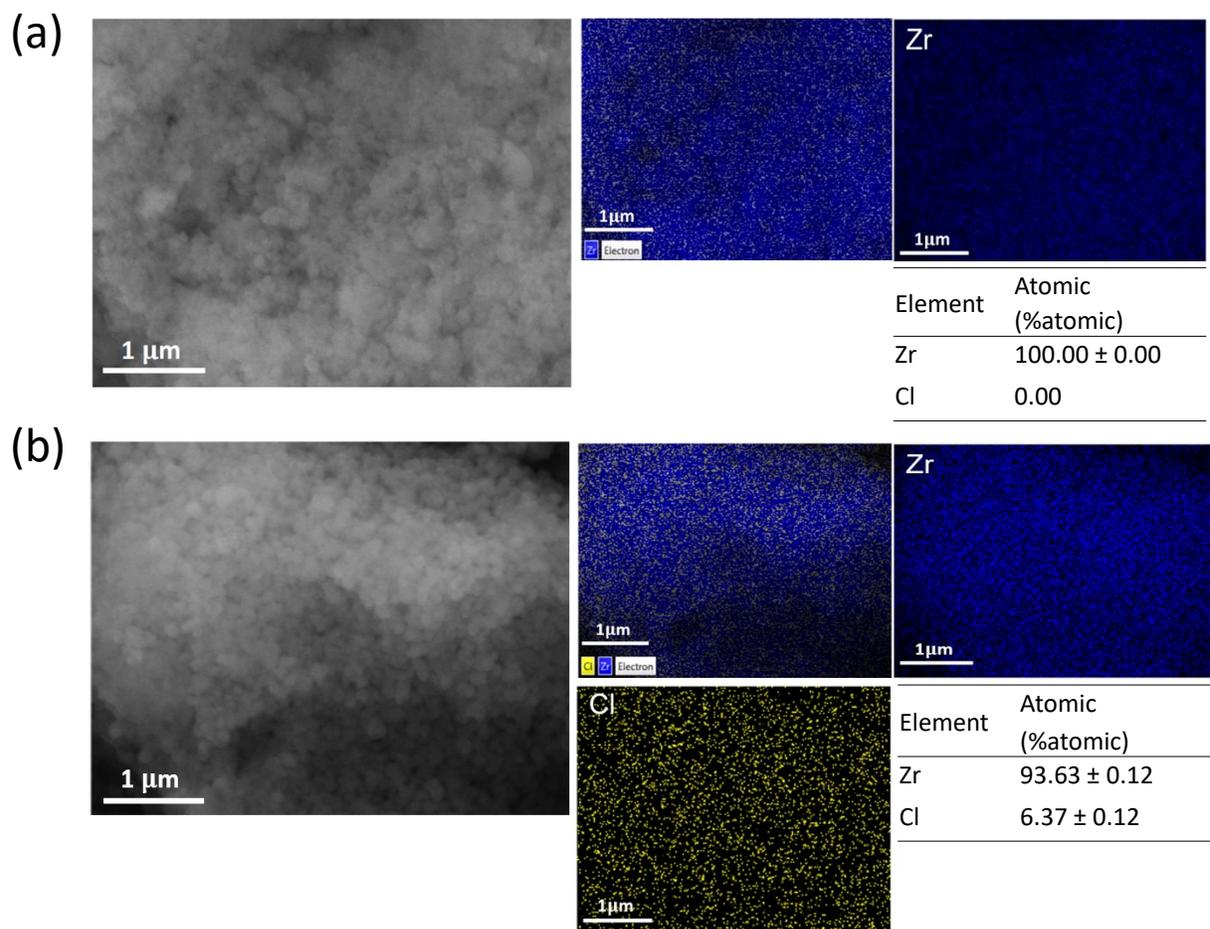
**Table S4.** Conversions and yields for catalytic conversion of glucose into LEV at the various amount of UiO-66\*-A catalyst at 190 °C for 6 h.

Amount of catalyst (g)	Glucose conversion (%)	LEV (mol)	5-HMF (mol)	LA (mol)
0.10	96.98 ± 0.14	0.46 ± 0.01	0.05 ± 0.00	0.04 ± 0.00
0.20	>99.00	0.58 ± 0.01	0.06 ± 0.00	0.05 ± 0.01
0.30	>99.00	0.60 ± 0.01	0.04 ± 0.00	0.05 ± 0.00

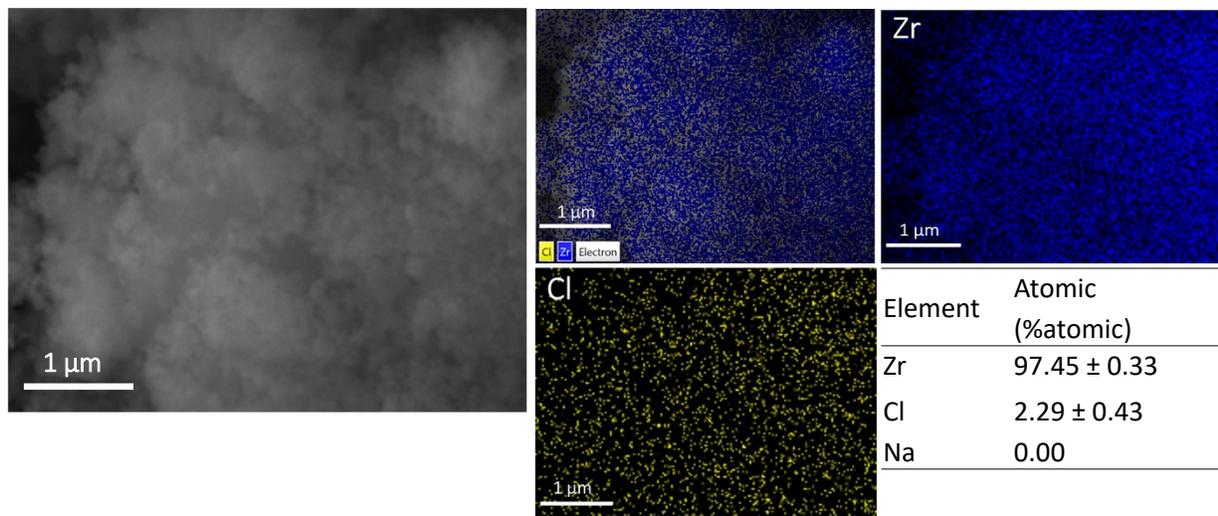
**Table S5.** Conversions and yields for catalytic conversion of glucose into LEV at the various amount of NaCl salt on UiO-66\*-A catalyst at 190 °C for 6 h.

Catalyst	<sup>a</sup> NaCl salt (w/w)	Glucose conversion (%)	LEV (mol)	5-HMF (mol)	LA (mol)
UiO-66*-A	none	>99.00	0.58 ± 0.01	0.06 ± 0.00	0.05 ± 0.01
UiO-66*-A	0.5	>99.00	0.60 ± 0.01	0.01 ± 0.00	0.04 ± 0.01
UiO-66*-A	1	>99.00	0.75 ± 0.01	0.02 ± 0.01	0.03 ± 0.01
UiO-66*-A	2	>99.00	0.83 ± 0.01	0.02 ± 0.00	0.02 ± 0.01
UiO-66*-A	3	>99.00	0.84 ± 0.00	0.02 ± 0.00	0.02 ± 0.02

<sup>a</sup>NaCl to glucose ratio (w/w).



**Figure S6.** Scanning electron microscopy-energy dispersive X-ray spectroscopy (SEM-EDS) mapping of (a) UiO-66\* and (b) UiO-66\*-A.



**Figure S7.** SEM-EDS mapping of UiO-66\*-A after the glucose conversion reaction at 190 °C for 6 h.

: SEM-EDS analysis confirmed the absence of Na in the used catalyst, likely due to the acidic final solution (pH 2-3), which promotes the formation of –COOH groups instead of –COONa.

**Table S6.** Comparisons of catalytic performance of different MOF catalysts investigated for the conversion of glucose into HMF and LEV under various conditions.

Entry	Catalyst	Reactant	Conversion (%)	Yield (mol)	Solvent	Condition	Ref.
1	PTA/MIL-101	fructose	82.00%	0.63mol HMF	DMSO	130 °C, 30 m	1
2	MIL-101(Cr)-SO <sub>3</sub> H	fructose	> 99.00%	0.90mol HMF	DMSO	120 °C, 1 h	2
3	Fe/MIL-101 (Cr)	glucose	91.50%	0.56mol HMF	DMSO/H <sub>2</sub> O	170 °C, 2 h	3
4	PO <sub>4</sub> /NU-1000	glucose	> 99.00%	0.64mol HMF	H <sub>2</sub> O/2PrOH	140 °C, 7 h	4
5	MIL-100(Fe)/Lys-PM <sub>2</sub>	glucose	98.70%	0.58mol LEV	H <sub>2</sub> O	150 °C, 9 h	5
6	UiO-66-NH-R-SO <sub>3</sub> H	glucose	> 99.00%	0.72mol LEV	H <sub>2</sub> O	170 °C, 24 h	6
7	UiO-66*-A	glucose	> 99.00%	0.83mol LEV	H <sub>2</sub> O	190 °C, 6 h	this work

: Several MOFs with Lewis and Bronsted acid attributes have been proposed for catalytic conversion of C6 sugars (*e.g.*, glucose and fructose) to 5-HMF (entries 1-4). Only a few MOF catalysts effectively convert glucose to LEV. Examples include MIL-100(Fe) loaded with an insoluble lysine-functionalized phosphotungstic acid (entry 5) and UiO-66-NH-R-SO<sub>3</sub>H (entry 6) achieving a yield of 0.58 and 0.72 LEV, respectively. In this study (entry 7), we successfully modified UiO-66 through the post-synthetic treatment method. The resulting defect-engineered UiO-66 exhibiting both Lewis and Bronsted acid characteristics enables to catalyze a multistep reaction, achieving LEV yield of 0.83 mol/mol of glucose and complete glucose conversion under hydrothermal condition in the presence of NaCl.

**Table S7.** Carbon content (%) of the UiO-66\*-A catalyst before and after the glucose conversion reaction at 190 °C for 6 h, as determined by CHNS elemental analysis.

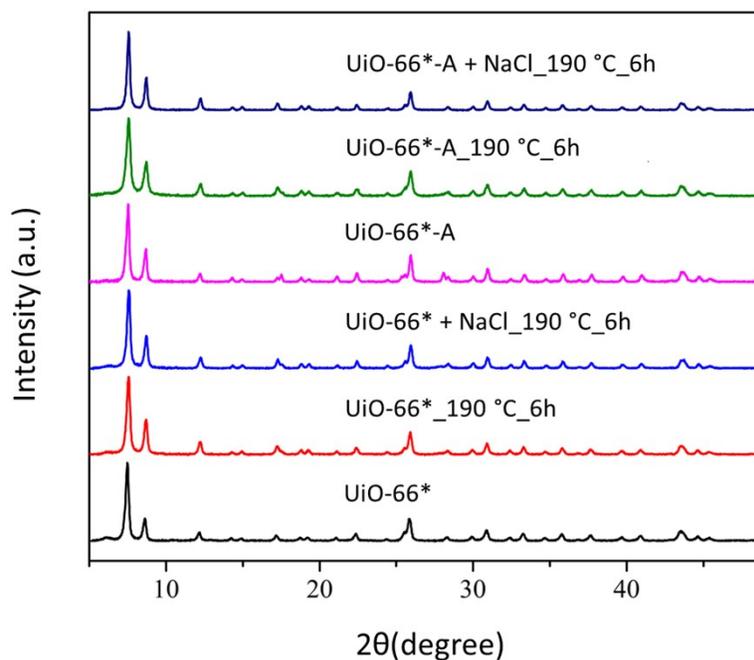
Entry	Sample	Carbon (%)
1	UiO-66*-A	22.52 ± 0.10
2	UiO-66*-A _1 <sup>st</sup>	34.84 ± 0.20
3	UiO-66*-A _2 <sup>nd</sup>	38.26 ± 0.36
4	UiO-66*-A _3 <sup>rd</sup>	40.41 ± 0.26

: The carbon content in the spent catalysts increases significantly during recycling, confirming the accumulation of additional carbonaceous substances within the framework. This accumulation hinders the diffusion of glucose into the framework, preventing it from accessing the active sites and ultimately leading to a decline in catalytic performance in subsequent runs. It is worth noting that regeneration of the spent catalyst was performed by washing it with hot methanol.

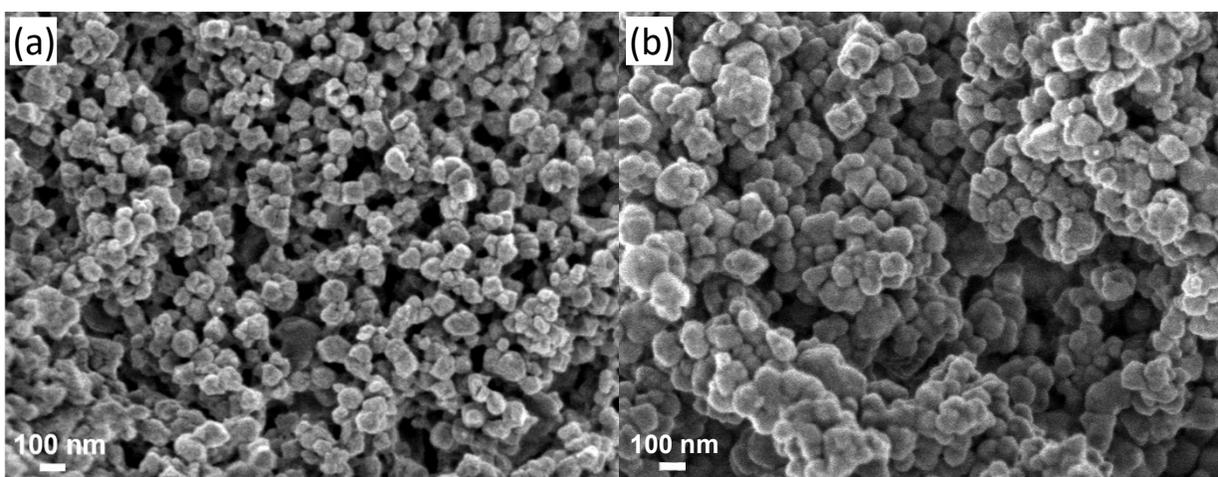
**Table S8.** Catalytic performance at different stirring speeds (300 and 400 rpm) under reaction conditions of 190 °C.

Reaction time (h)	Stirring speed (rpm)	Glucose conversion (%)	LEV (mol)	5-HMF (mol)	LA (mol)
4	300	84.67 ± 0.57	0.41 ± 0.01	0.09 ± 0.01	0.08 ± 0.01
	400	84.73 ± 0.73	0.40 ± 0.00	0.10 ± 0.00	0.08 ± 0.01
6	300	>99.00	0.58 ± 0.00	0.06 ± 0.00	0.05 ± 0.01
	400	>99.00	0.59 ± 0.00	0.06 ± 0.01	0.05 ± 0.01

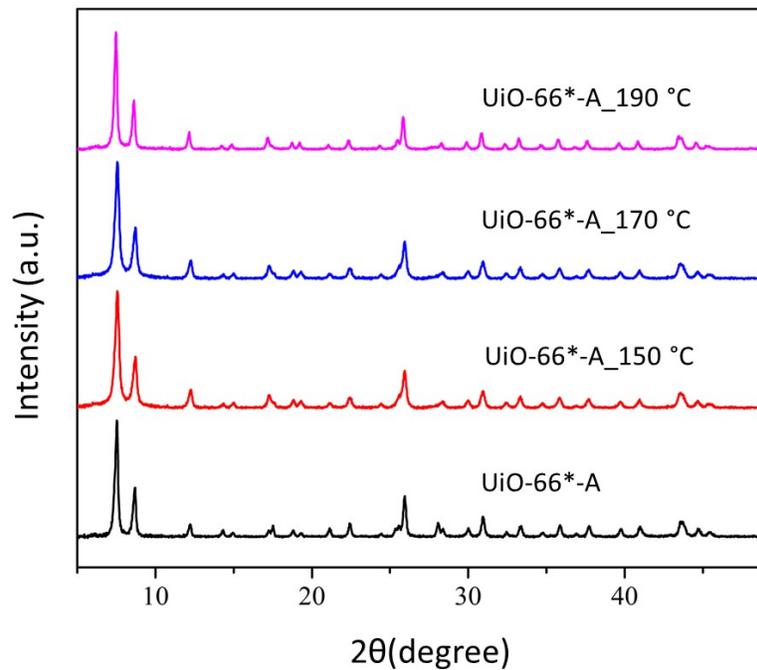
: The catalytic performance at stirring speeds of 300 and 400 rpm was comparable. This confirms that the reaction kinetics are unaffected by stirring speed.



**Figure S8.** PXRD patterns of the UiO-66\* and UiO-66\*-A before and after the glucose conversion reaction with and without NaCl at 190 °C for 6 h, respectively.

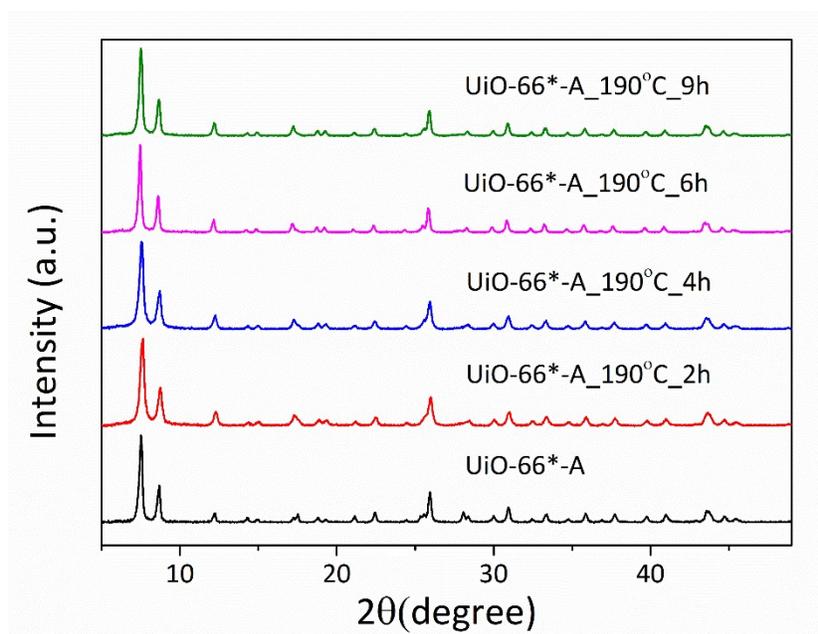


**Figure S9.** Scanning electron microscopic (SEM) images of (a) UiO-66\*-A before and (b) UiO-66\*-A after the reaction at 190 °C 6 h, respectively.



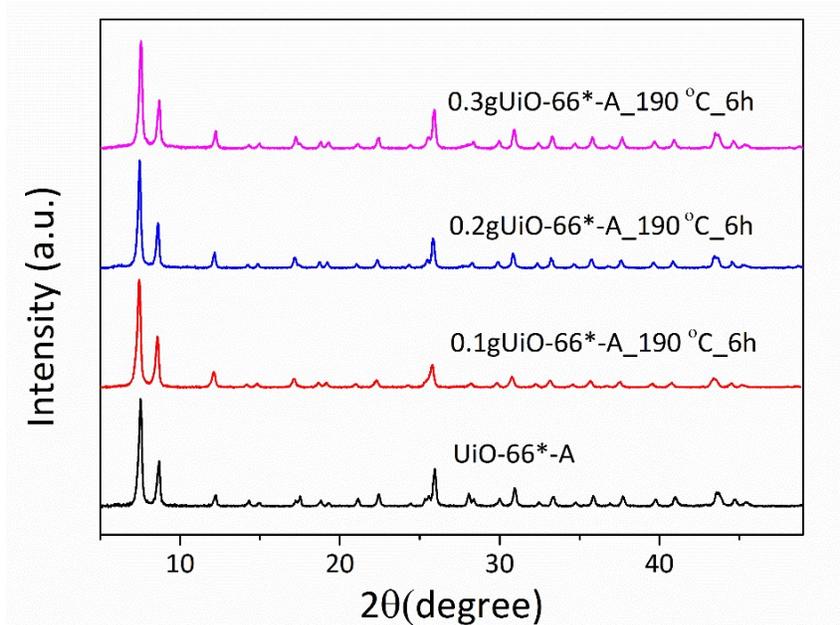
**Figure S10.** PXRD patterns of the UiO-66\*-A before and after the reaction at various temperatures for 6

h.

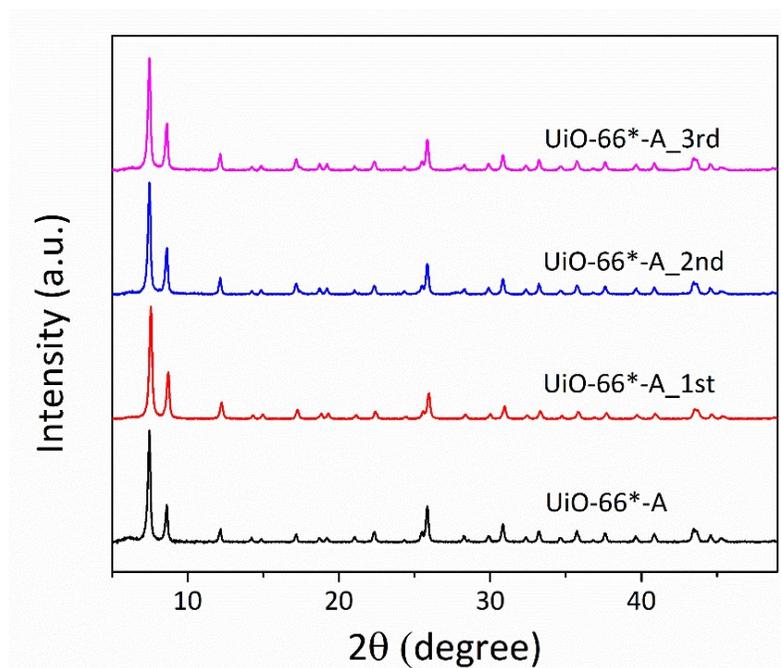


**Figure S11.** PXRD patterns of the UiO-66\*-A before and after the reactions at 190 °C for various reaction

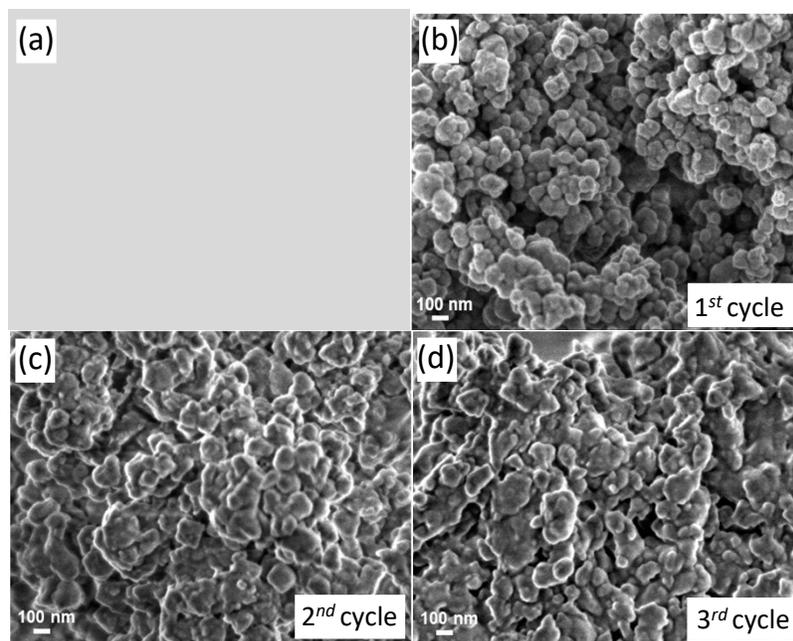
times.



**Figure S12.** PXRD patterns of the UiO-66\*-A before and after the reaction at 190 °C for 6 h at various amount of catalyst.



**Figure S13.** PXRD patterns of the UiO-66\*-A before and after the glucose conversion reaction at 190 °C for 6 h over three consecutive cycles.



**Figure S14.** SEM images of the UiO-66\*-A before and after the glucose conversion reaction at 190 °C for 6 h over 3 cycles of test.

## REFERENCES

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