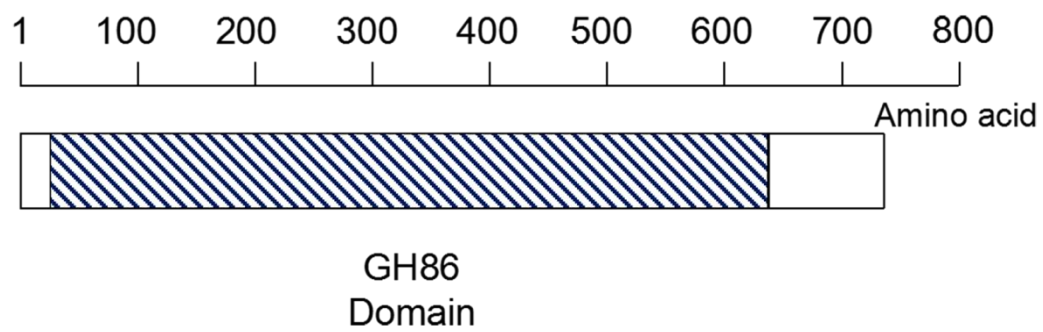


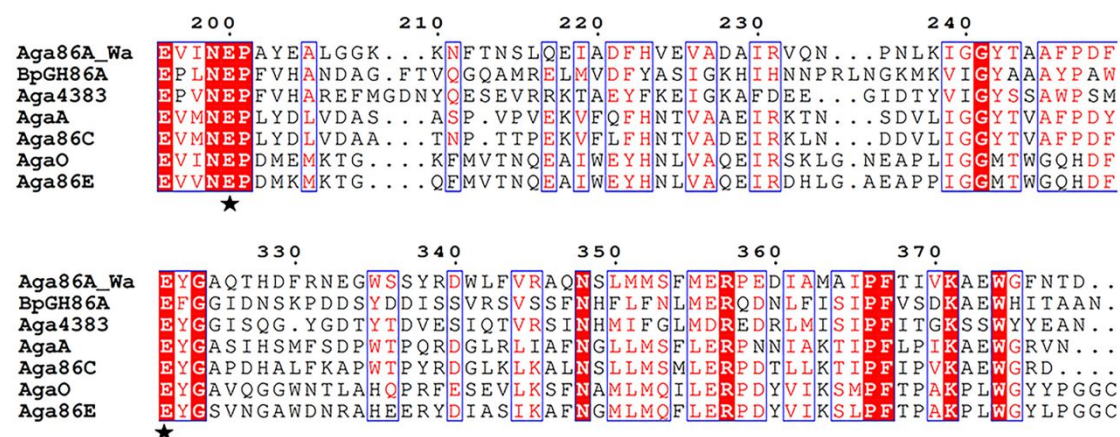
## SUPPLEMENTAL MATERIAL

**Supplementary Table 1** Percentage (%) of each oligosaccharide in products (45 U).

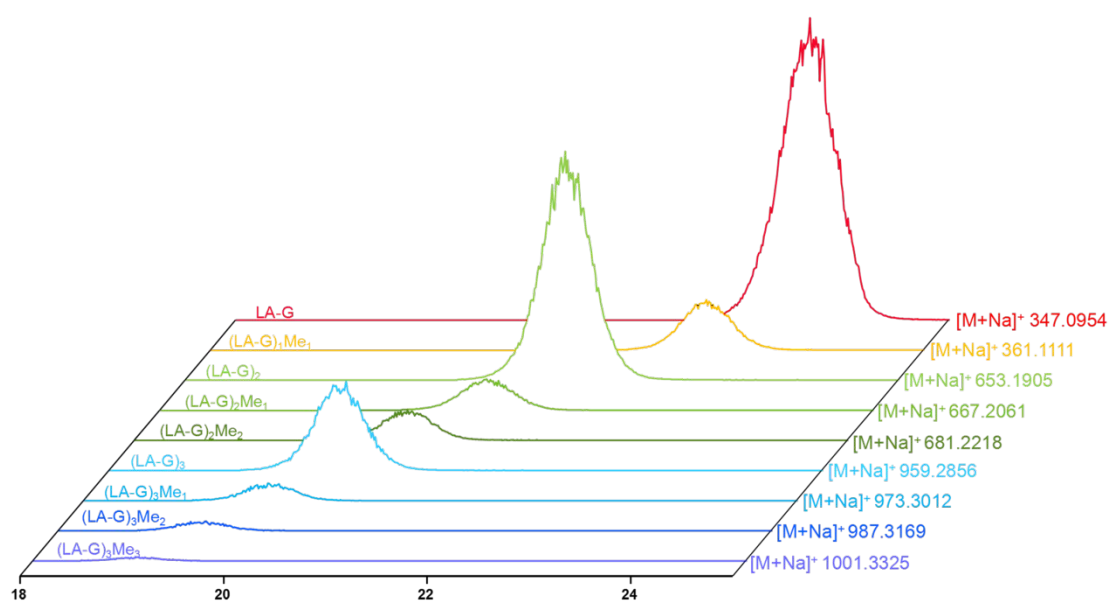
Oligosaccharide composition	Percentage (%)
(LA-G) <sub>1</sub>	40.77
(LA-G) <sub>1</sub> Me	6.27
(LA-G) <sub>2</sub>	29.22
(LA-G) <sub>2</sub> Me <sub>1</sub>	4.35
(LA-G) <sub>2</sub> Me <sub>2</sub>	4.23
(LA-G) <sub>3</sub>	11.57
(LA-G) <sub>3</sub> Me <sub>1</sub>	2.12
(LA-G) <sub>3</sub> Me <sub>2</sub>	1.08
(LA-G) <sub>3</sub> Me <sub>3</sub>	0.39



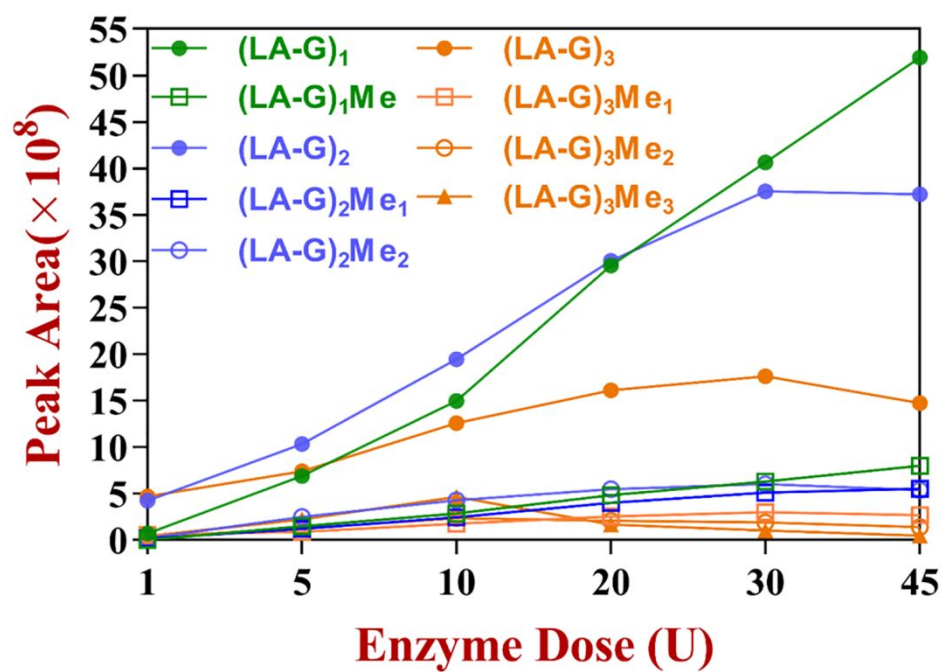
**Fig. S1** A schematic of the modular arrangement of Aga86A\_Wa. The GH86 family glycoside hydrolase catalytic domain (26-637 amino acids) was identified by the dbCAN.



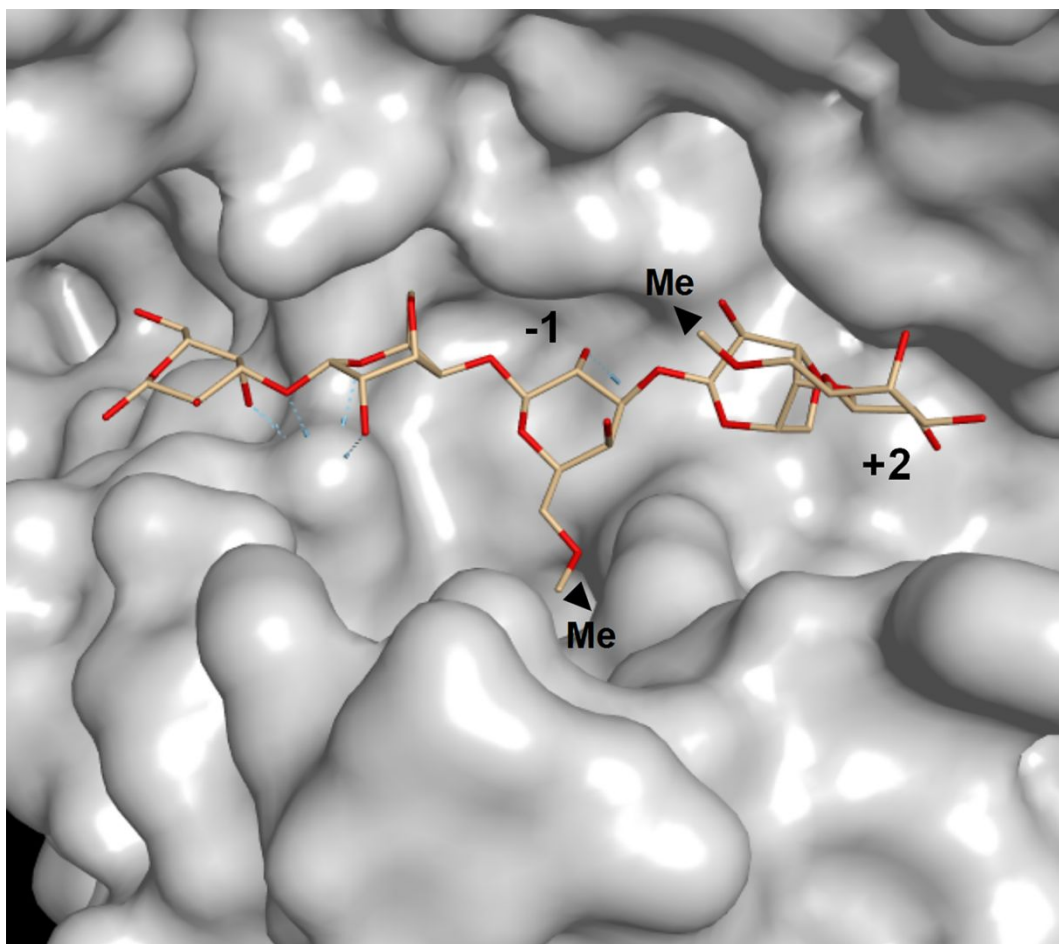
**Fig. S2** Amino acid sequence alignment of Aga86A\_Wa with characterized GH86 family enzymes. Critical catalytic residues were marked by stars.



**Fig. S3** Extracted ion chromatograms of oligosaccharides in product obtained with 45U enzyme (Fig, 5). The  $m/z$  and composition of oligosaccharides were annotated to the corresponding curve.



**Fig. S4** Products prepared by 100 mg substrate incubated with different enzyme doses of Aga86A\_Wa at 24 h.



**Fig. S5** Predicted tertiary structure of Aga86A\_Wa superimposed with the ligand in BuGH86 (5TA0). The O-6 positions of G residues at -1 and +2 subsites were modified with methyl groups.