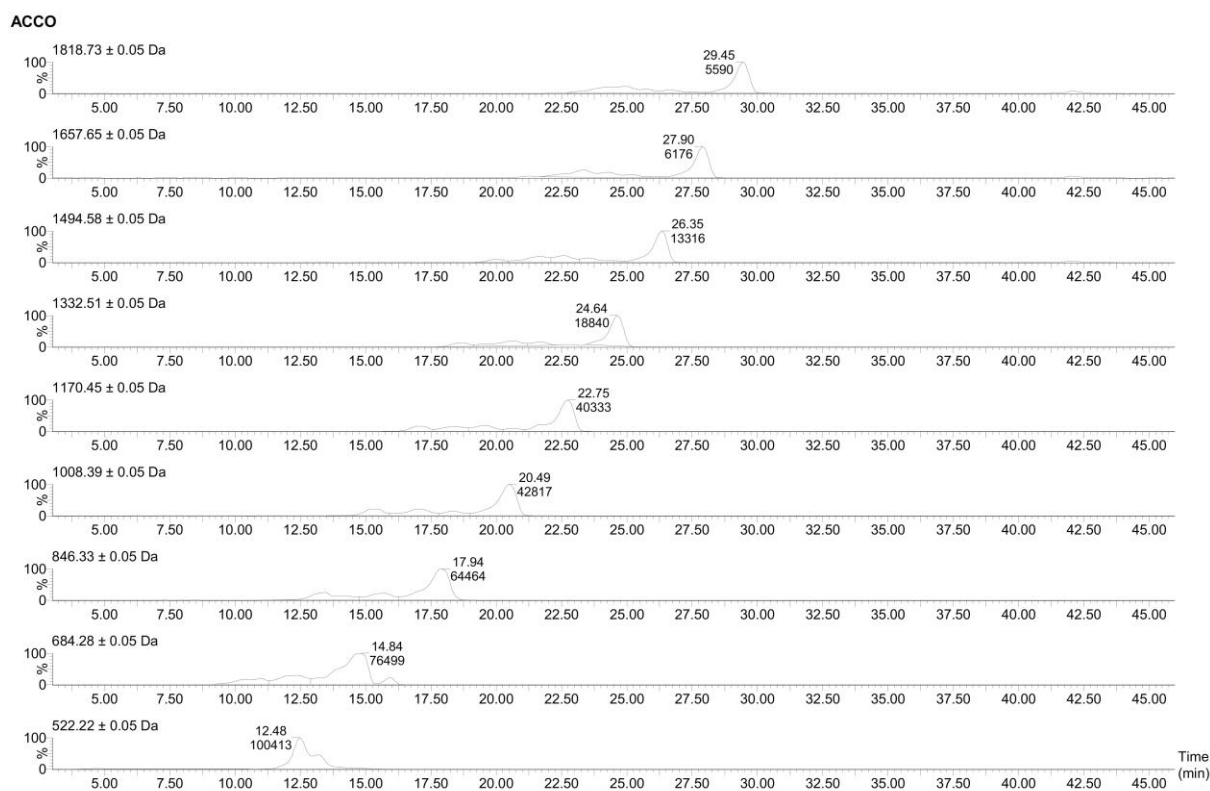


## 1 Supplementary information

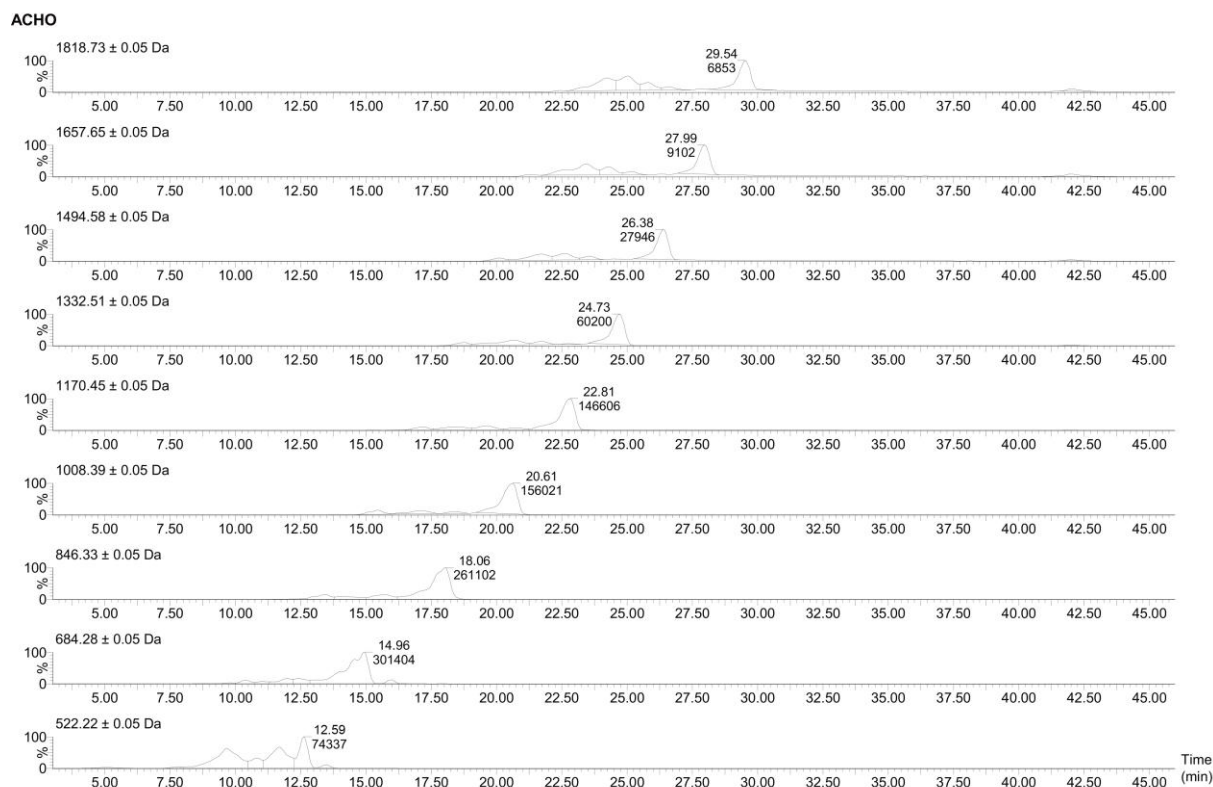
- 2 • Figure S1 and Figure S2 show the oligosaccharide composition.
- 3 • Figure S3 shows the monosaccharide composition.
- 4 • Table S1 displays the primers used in RT-PCR.
- 5 • All other text information describes supplementary methods.

### 6 1. Oligosaccharide abundance of ACCO and ACHO by UPLC-MS analysis



7

8 **Figure S1.** Ultra performance liquid chromatography of ACCO. The retention time and peak area of  
9 each oligosaccharide of ACCO were displayed.



10

11 **Figure S2.** Ultra performance liquid chromatography of ACHO. The retention time and peak area of  
 12 each oligosaccharide of ACHO were displayed.

13 **2. PMP derivatization of monosaccharides**

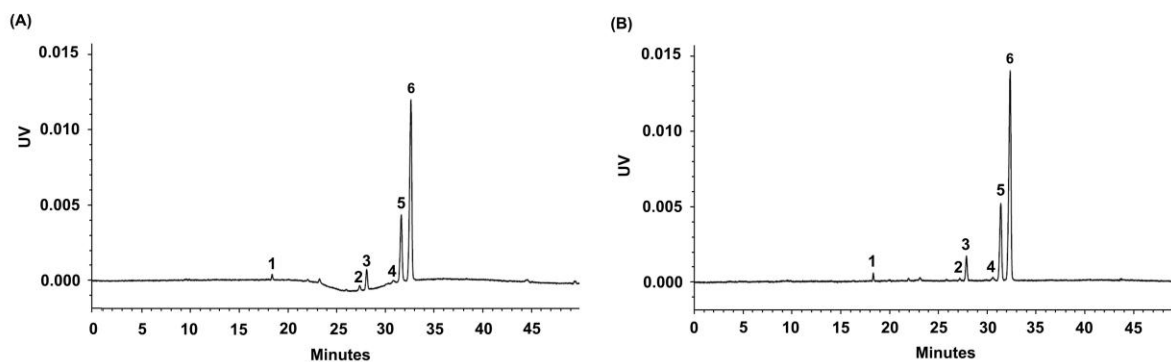
14 To prepare the PMP derivatives of monosaccharide standards, the following monosaccharides (10  
 15 mg for each) were dissolved in 5 ml of H<sub>2</sub>O and mixed well: D-(+)-arabinose, D-(+)-glucose,  
 16 D-(+)-fructose, D-(+)-galactose, D-(+)-mannose. The derivatization reaction system was composed of 40  
 17 μl of standard mixture, 600 μl of NaOH (0.3 M) and 600 μl PMP (0.5 M) in methanol. After the  
 18 incubation in water bath at 70 °C for 30 min, the reaction was stopped, cooled to room temperature and  
 19 neutralized by addition of 600 μl of HCl (0.3 M). After extracted by 1 ml of chloroform, the aqueous  
 20 layer was filtered with 0.22 μm membrane for capillary electrophoresis analysis. The PMP derivatives  
 21 of hydrolyzed ACCO or ACHO were performed as above-mentioned.

22 **3. Capillary electrophoresis analysis of PMP derivatives**

23 The PMP derivatives were separated using a P/ACE MDQ CE instrument (Beckman Coulter,  
 24 Fullerton, CA, USA) with a UV detector at a wavelength of 245 nm. The analysis was performed on an  
 25 Uncoated fused-silica capillary (Φ50 μm ×60 cm) at 26 °C. The mobile phase was a buffer solution of  
 26 sodium borate (55 mM, pH 10.55). For each monosaccharide, the calibration factor ( $f_i$ ) was calculated  
 27 based on the peak area and molecular weight. The relative abundance of monosaccharide constituents  
 28 was obtained by the ratio of  $f_i$  multiplied by their peak areas ( $A_i$ ), as shown in the following:

$$\frac{n_i}{n_j} = \frac{f_i A_i}{f_j A_j}$$

29



30  
 31 **Figure S3.** Composition analysis of ACHO and ACCO using capillary electrophoresis. A)  
 32 Monosaccharide spectrum of ACCO; B) Monosaccharide spectrum of ACHO. 1, PMP; 2, arabinose; 3,  
 33 glucose; 4, fucose; 5, galactose; 6, mannose.

34 **4. Primers for real-time PCR**

35 **Table S1. The primer sequence used for RT-PCR**

| Primer name    | Forward primer            | Reverse primer           |
|----------------|---------------------------|--------------------------|
| IL-1 $\beta$   | CGACAAAATACCTGTGGCCT      | TTCTTTGGGTATTGCTTGGG     |
| IL-6           | GAAACCGCTATGAAGTTCCTCTCTG | TGTTGGGAGTGGTATCCTCTGTGA |
| IL-8           | ATGGCTGGGATTCACCTCAA      | AAGCCTCGCGACCATTCTT      |
| TNF- $\alpha$  | AGGTCTGGGCCATAGAACT       | CCACCACGCTCTTCTGTCTAC    |
| MCP-1          | GGGATCATCTTGCTGGTGAA      | AGGTCCCTGTCATGCTTCTG     |
| $\beta$ -actin | AGGTGACAGCATTGCTTCTG      | GCTGCCTCAACACCTCAAC      |

36