

## Synthesis of Pentasaccharide and Heptasaccharide Derivatives and Their Effects on Plant Growth

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Two oligosaccharide derivatives,  $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-4)- $\alpha$ -D-ManpOMe (**1**) and  $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-4)- $\alpha$ -D-ManpOMe (**2**), have been synthesized efficiently using a convergent glycosylation strategy of 2 + 3 and 2 + 5. 1,6-Anhydro- $\beta$ -D-glucopyranose, which was prepared from cotton pyrolysis, was applied as a key synthon in the synthesis, significantly simplifying the preparation. The bioassay suggested that these two oligosaccharides can both stimulate the growth of maize cultured in liquid medium at a concentration of 3 ppm.

**KEYWORDS:** Oligosaccharide; synthesis; plant growth; maize

### INTRODUCTION

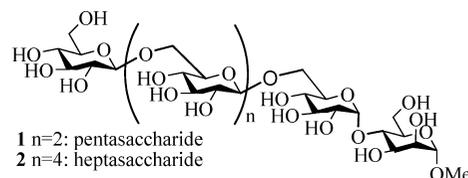
Various synthetic and naturally derived fungicides, bactericides, and antiviral agents have been used to prevent crops from infection by pathogens without deleteriously affecting their growth and ultimate harvesting. Although such compositions are effective, there have been more and more concerns from consumers regarding the potential harmful side effects of chemical agents. This, in turn, has led people to pay more attention to products of natural origin with less likely adverse side effects (1–3).

Oligosaccharides, that can exert signaling effects on plant tissues at very low concentration, are termed as oligosaccharins. Some of them are elicitors for they induce a response that may help the plant resist disease, while some have effects on plant growth and development, which are described as nontraditional plant hormones such as pectic oligosaccharins and xyloglucan oligosaccharins. Despite some oligosaccharins as plant hormones are ensured, there is still an urgent need to add flesh to our skeletal understanding of oligosaccharins, including the discovery of new biological effects (4–8). Two oligosaccharides, heptasaccharide (HS) and octasaccharide (OS), were isolated from the water extract of the rhizomes of *Paris polyphylla* var. *yunnanensis*. They were identified as linear oligomers, composed of glucose and mannose residues. The structures of compounds HS and OS were elucidated, on the basis of NMR and mass

spectra analyses, as D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)-Glc-(1-6)- $\beta$ -D-Glc-(1-4)- $\alpha$ -D-Man and D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)- $\beta$ -D-Glc-(1-6)-Glc-(1-6)- $\beta$ -D-Glc-(1-4)- $\alpha$ -D-Man, respectively. Primary bioactivity testing suggests that HS and OS may have plant growth regulatory activity in plant tissue cultures (9, 10). We hereby describe a practical method in the preparation of HS analogues, pentasaccharide  $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-4)- $\alpha$ -D-ManpOMe (**1**) and heptasaccharide  $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-6)- $\beta$ -D-Glcp-(1-4)- $\alpha$ -D-ManpOMe (**2**) (Scheme 1), and disclose our primary results on maize growth after treating with **1** and **2**.

In our ongoing project, we found an efficient method for the synthesis of  $\beta$ -D-(1 $\rightarrow$ 6)-linked gluco-oligosaccharides using levoglucoside, which was prepared from cotton pyrolysis (11) as starting material. Thus, 1,6-anhydro- $\beta$ -D-glucose (levoglucoside, **3**) was benzylated in DMF with benzyl bromide and NaH affording 1,6-anhydro-2,3,4-tri-*O*-benzyl- $\beta$ -D-glucopyranose (**4**). Treatment of **4** with 2-propanethiol and BF<sub>3</sub>·Et<sub>2</sub>O in refluxed CH<sub>2</sub>Cl<sub>2</sub> obtained isopropyl 2,3,4-tri-*O*-benzyl-1-thio- $\beta$ -D-glucopyranoside (**5**) as an  $\alpha$  product in 56% yield over two steps. The small *J*<sub>1,2</sub> value (5.5 Hz) at 5.34 ppm on the related <sup>1</sup>H NMR spectrum clearly indicated the correct structure of **5**. Conver-

Scheme 1



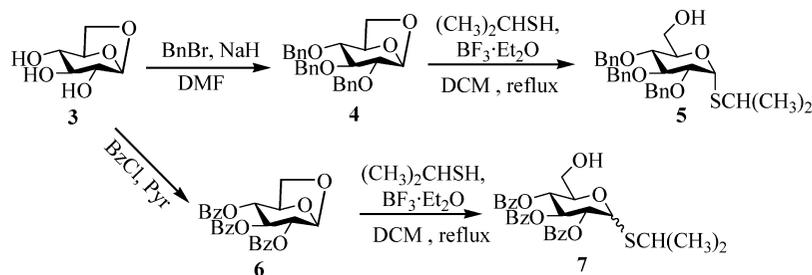
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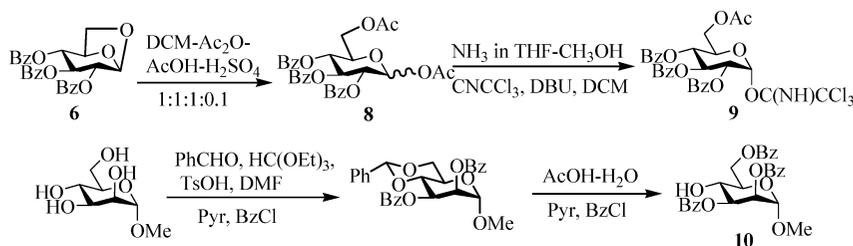
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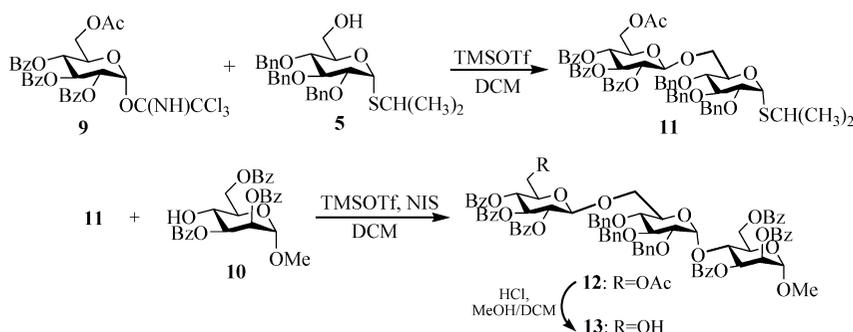
Scheme 2



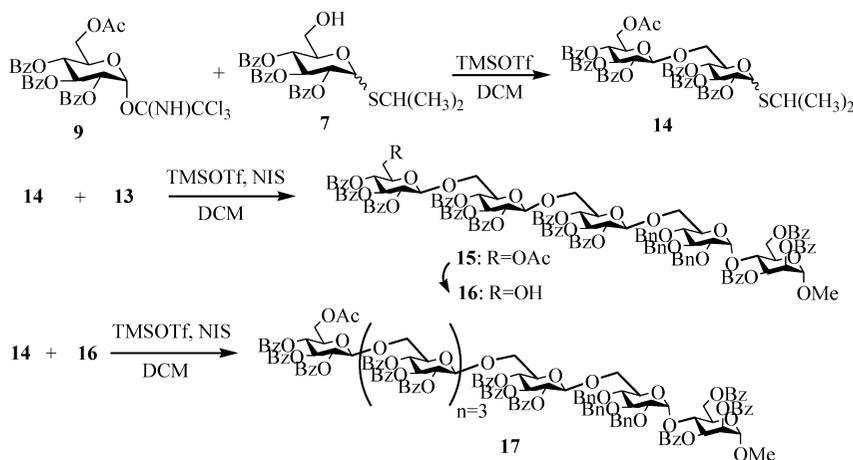
Scheme 3



Scheme 4



Scheme 5



gently, compound **3** was benzoylated with benzoyl chloride in dry pyridine affording 1,6-anhydro-2,3,4-tri-*O*-benzoyl- $\beta$ -D-glucopyranose (**6**), followed by ring opening with 2-propanethiol and  $\text{BF}_3 \cdot \text{Et}_2\text{O}$  in refluxed  $\text{CH}_2\text{Cl}_2$ , which furnished isopropyl 2,3,4-tri-*O*-benzoyl-1-thio-D-glucopyranoside (**7**) as an  $\alpha,\beta$  mixture ( $n_\alpha/n_\beta \approx 6/1$ ) in 47% yield over two steps (**Scheme 2**).

Acetolysis of **6** with  $\text{Ac}_2\text{O}$ - $\text{AcOH}$ - $\text{H}_2\text{SO}_4$  (1:1:0.1) in  $\text{CH}_2\text{Cl}_2$  gave 1,6-di-*O*-acetyl-2,3,4-tri-*O*-benzoyl- $\beta$ -D-glucopyranose (**8**), which was then transformed into glycosyl donors 6-*O*-acetyl-2,3,4-tri-*O*-benzoyl- $\alpha$ -D-glucopyranosyl trichloroacetimidate (**9**) according to published methods in 91% yield over three steps (**12**). Methyl 2,3,6-*O*-benzoyl- $\alpha$ -D-mannopyranoside (**10**) was prepared from commercial available methyl D-mannoside via

4,6-*O*-benzylideneation, benzoylation, acid hydrolysis, and regioselective 6-*O*-benzoylation in 76% yield (**13**) (**Scheme 3**).

Condensation of **5** with **9** in the presence of a catalytic amount of trimethylsilyl trifluoromethanesulfonate (TMSOTf) in  $\text{CH}_2\text{Cl}_2$  at  $-20^\circ\text{C}$  gave desired disaccharide, isopropyl 6-*O*-acetyl-2,3,4-tri-*O*-benzoyl- $\beta$ -D-glucopyranosyl-(1 $\rightarrow$ 6)-2,3,4-tri-*O*-benzoyl- $\beta$ -D-glucopyranosyl-1-thio- $\alpha$ -D-glucopyranoside (**11**), in excellent yield (98%). Coupling of disaccharide thioglycosyl donor **11** and acceptor **10** in  $\text{CH}_2\text{Cl}_2$  at  $-20^\circ\text{C}$  in the presence of cocatalyst TMSOTf and *N*-iodosuccinimide (NIS), afforded the expected  $\alpha$ -linked trisaccharide (**12**) in good yield (78%). A doublet at  $\delta$  5.44 ppm in the  $^1\text{H}$  NMR spectrum supports the newly formed  $\alpha$ -bond between glucosyl units II and III. Regioselective deacetylation

Table 1. <sup>1</sup>H-NMR Data of Compounds 5–17

comps	δ (ppm)
5	1.27 (d, 6H, J 6.7 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 2.90 (m, 1H, SCH), 5.34 (d, 1H, J 5.5 Hz, H <sub>1</sub> ), 4.62 (d, 6H, J 11 Hz, PhCH), 3.74 (dd, 1H, J 5.5 Hz, 9.8 Hz, H <sub>2</sub> ), 3.50 (dd, 1H, J 9.8 Hz, 9.1 Hz, H <sub>3</sub> ), 3.84 (t, 1H, J 9.1 Hz, H <sub>4</sub> ), 4.09 (m, 1H, H <sub>5</sub> ), 3.73 (m, 2H, H <sub>6a</sub> , H <sub>6b</sub> ), 7.25 (m, 15H, 3Ph).
7	1.23 (d, 6H, J 6.7 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 3.02 (m, 1H, CH), 6.01 (d, 1H, J 5.8 Hz, H <sub>1</sub> ), 5.44 (dd, 1H, J 5.8 Hz, 10.3 Hz, H <sub>2</sub> ), 5.48 (t, 1H, J 9.9 Hz, H <sub>3</sub> ), 6.06 (t, 1H, J 9.9 Hz, H <sub>4</sub> ), 4.46 (m, 1H, H <sub>5</sub> ), 3.73 (m, 2H, H <sub>6a</sub> , H <sub>6b</sub> ), 7.26 (m, 15H, 3Ph); 1.23 (d, 1H <sub>β</sub> , J 6.7 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 3.22 (m, 0.15H <sub>β</sub> , SCH), 4.91 (d, 0.15H <sub>β</sub> , J 10.1 Hz, H <sub>1</sub> ), 5.44 (t, 0.15H <sub>β</sub> , J 10.1 Hz, H <sub>2</sub> ), 5.44 (t, 0.15H <sub>β</sub> , J 9.9 Hz, H <sub>3</sub> ), 5.93 (t, 0.15H <sub>β</sub> , J 9.9 Hz, H <sub>4</sub> ), 3.73 (m, 0.45H <sub>β</sub> , H <sub>5</sub> , H <sub>6a</sub> , H <sub>6b</sub> ), 7.26 (m, 2.5H <sub>β</sub> , 3Ph);
9	2.07 (s, 3H, CH <sub>3</sub> ), 6.83 (d, 1H, J 3.0 Hz, H <sub>1</sub> ), 5.58 (dd, 1H, J 3.6 Hz, 10.2 Hz, H <sub>2</sub> ), 6.22 (t, 1H, J 10.0 Hz, H <sub>3</sub> ), 5.72 (t, 1H, J 10.0 Hz, H <sub>4</sub> ), 4.48 (m, 1H, H <sub>5</sub> ), 4.27 (m, 2H, H <sub>6a</sub> , H <sub>6b</sub> ), 8.7 (s, 1H, NH), 7.28 (m, 15H, 3Ph).
10	3.49 (s, 3H, CH <sub>3</sub> ), 5.61 (s, 1H, H <sub>1</sub> ), 4.93 (s, 1H, H <sub>2</sub> ), 5.60- (dd, 1H, J 3.3 Hz, 12.4 Hz, H <sub>3</sub> ), 4.26 (t, 1H, J 9.6 Hz, H <sub>4</sub> ), 4.08 (m, 1H, H <sub>5</sub> ), 4.63 (dd, 1H, H <sub>6a</sub> ), 4.87 (dd, 1H, H <sub>6b</sub> ), 7.29 (m, 15H, 3Ph).
11	1.24 (d, 6H, J 6.7 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 1.99 (s, 3H, CH <sub>3</sub> ), 2.88 (m, 1H, SCH), 4.73 (d, 1H, J 7.8 Hz, H <sub>1</sub> <sup>I</sup> ), 5.54 (dd, 1H, J 8.0 Hz, 9.6 Hz, H <sub>2</sub> <sup>I</sup> ), 5.81 (t, 1H, J 9.6 Hz, H <sub>3</sub> <sup>I</sup> ), 5.56 (t, 1H, J 9.7 Hz, H <sub>4</sub> <sup>I</sup> ), 4.19 (m, 1H, H <sub>5</sub> <sup>I</sup> ), 4.23 (m, 2H, H <sub>6a</sub> <sup>I</sup> , H <sub>6b</sub> <sup>I</sup> ), 5.36 (1H, J 5.1 Hz, H <sub>1</sub> <sup>II</sup> ), 4.85, 4.41, 4.22 (d, 6H, J 11 Hz, PhCH <sub>2</sub> O), 3.67 (dd, 1H, J 5.1 Hz, 9.5 Hz, H <sub>2</sub> <sup>II</sup> ), 3.72 (t, 1H, J 9.6 Hz, H <sub>3</sub> <sup>II</sup> ), 3.40 (t, 1H, J 9.8 Hz, H <sub>4</sub> <sup>II</sup> ), 3.90 (m, 1H, H <sub>5</sub> <sup>II</sup> ), 3.79 (m, 1H, H <sub>6a</sub> <sup>II</sup> ), 4.12 (m, 1H, H <sub>6b</sub> <sup>II</sup> ), 7.02 (m, 30H, 6Ph).
12	1.9 (s, 3H, CH <sub>3</sub> ), 3.48 (s, 3H, OCH <sub>3</sub> ), 4.55 (d, 1H, J 8.2 Hz, H <sub>1</sub> <sup>I</sup> ), 5.50 (dd, 1H, J 8.0 Hz, 9.7 Hz, H <sub>2</sub> <sup>I</sup> ), 5.77 (t, 1H, J 9.6 Hz, H <sub>3</sub> <sup>I</sup> ), 5.53 (t, 1H, J 9.7 Hz, H <sub>4</sub> <sup>I</sup> ), 5.44 (d, 1H, J 3.6 Hz, H <sub>1</sub> <sup>II</sup> ), 3.24 (dd, 1H, J 3.6 Hz, 9.9 Hz, H <sub>2</sub> <sup>II</sup> ), 3.79 (t, 1H, J 9.6 Hz, H <sub>3</sub> <sup>II</sup> ), 3.33 (t, 1H, J 9.5 Hz, H <sub>4</sub> <sup>II</sup> ), 3.66 (m, 1H, H <sub>6a</sub> <sup>II</sup> ), 3.78 (m, 2H, H <sub>5</sub> <sup>II</sup> , H <sub>6b</sub> <sup>II</sup> ), 4.89 (d, 1H, J 1.4 Hz, H <sub>1</sub> <sup>III</sup> ), 5.67 (dd, 1H, J 1.9 Hz, 2.8 Hz, H <sub>2</sub> <sup>III</sup> ), 5.89 (dd, 1H, J 9.6 Hz, 3.2 Hz, H <sub>3</sub> <sup>III</sup> ), 4.70 (t, 1H, J 9.7 Hz, H <sub>4</sub> <sup>III</sup> ), 4.03 (m, 8H, H <sub>5</sub> <sup>III</sup> , H <sub>5</sub> <sup>III</sup> , H <sub>6a</sub> <sup>I</sup> , H <sub>6b</sub> <sup>I</sup> , 4PhCH <sub>2</sub> O), 4.51 (d, 1H, J 11 Hz, PhCH <sub>2</sub> O), 4.66 (d, 1H, J 11 Hz, PhCH <sub>2</sub> O), 4.55 (m, 1H, H <sub>6b</sub> <sup>III</sup> ), 4.89 (m, 1H, H <sub>6a</sub> <sup>III</sup> ), 6.93 (m, 45H, 9Ph).
14	1.18, 1.05 (d, 3H, J 7.8 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 1.07, 1.15 (d, 0.8H <sub>β</sub> , J 6.9 Hz, (CH <sub>3</sub> ) <sub>2</sub> ), 1.93 (s, 3H, CH <sub>3</sub> ), 1.93 (s, 0.8H <sub>β</sub> , COCH <sub>3</sub> ), 2.86 (m, 1H, SCH), 3.03 (m, 0.26H <sub>β</sub> , SCH), 5.78 (d, 1H, J 6.1 Hz, H <sub>1</sub> <sup>I</sup> ), 4.87 (d, 1H, 7.8 Hz, H <sub>1</sub> <sup>I</sup> ), 4.93 (d, 0.25H <sub>β</sub> , J 7.9 Hz, H <sub>1</sub> <sup>I</sup> ), 4.77 (d, 0.24H <sub>β</sub> , J 10.0 Hz, H <sub>1</sub> <sup>II</sup> ).
15	2.01 (s, 3H, COCH <sub>3</sub> ), 3.35 (s, 3H, OCH <sub>3</sub> ), 4.86 (d, 1H, J 7.8 Hz, H <sub>1</sub> <sup>I</sup> ), 5.19 (d, 1H, J 7.7 Hz, H <sub>1</sub> <sup>II</sup> ), 4.36 (d, 1H, J 8.0 Hz, H <sub>1</sub> <sup>III</sup> ), 5.02 (d, 1H, J 3.3 Hz, H <sub>1</sub> <sup>IV</sup> ), 4.92 (d, 1H, J 1.6 Hz, H <sub>1</sub> <sup>V</sup> ).
17	2.00 (s, 3H, CH <sub>3</sub> ), 3.36 (s, 3H, CH <sub>3</sub> ), 4.74 (d, 1H, J 7.7 Hz, H <sub>1</sub> <sup>I</sup> ), 4.96 (d, 1H, J 7.6 Hz, H <sub>1</sub> <sup>II</sup> ), 4.92 (d, 1H, J 8.0 Hz, H <sub>1</sub> <sup>III</sup> ), 5.10 (d, 1H, J 7.9 Hz, H <sub>1</sub> <sup>IV</sup> ), 4.32 (d, 1H, J 8.0 Hz, H <sub>1</sub> <sup>V</sup> ), 4.96 (d, 1H, J 3.5 Hz, H <sub>1</sub> <sup>VI</sup> ), 4.94 (d, 1H, J 1.7 Hz, H <sub>1</sub> <sup>VII</sup> ).

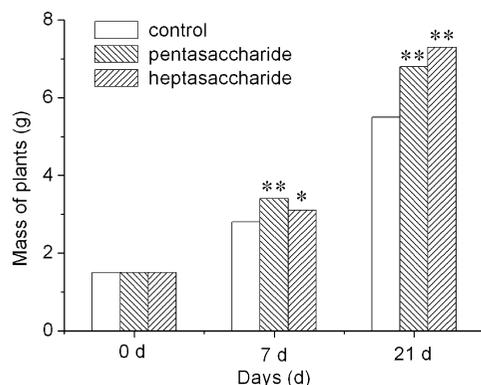
of **12** with HCl (gas) in MeOH/DCM (1:2, v/v) gave (**13**) as the trisaccharide glycosyl acceptor in good yield (89%) (**14**) (**Scheme 4**).

Condensation of **9** with **7** catalyzed by TMSOTf gave a key intermediate 6-*O*-acetyl-2,3,4-tri-*O*-benzoyl-β-D-glucopyranosyl-(1→6)-2,3,4-tri-*O*-benzoyl-1-thio-D-glucopyranoside (**14**) in 68% yield as an α,β mixture (*n*<sub>α</sub>/*n*<sub>β</sub> ≈ 4/1), which was directly used for the next reaction without further purification. Coupling of disaccharide glycosyl donor **14** with trisaccharide glycosyl acceptor **13** was promoted with TMSOTf/NIS in CH<sub>2</sub>Cl<sub>2</sub>, giving pentasaccharide (**15**) in an isolated yield of 80%. Removal of 6-*O*-deacetylation from **15** as described in the preparation of **13** afforded pentasaccharide glycosyl acceptor (**16**). Condensa-

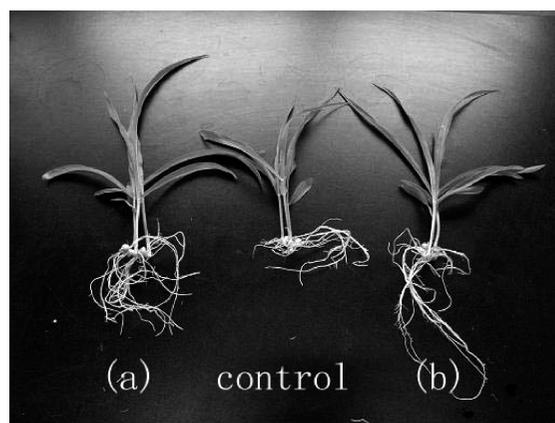
tion of **14** with **16** using the same reaction conditions as those described in the preparation of **15** obtained heptasaccharide (**17**) in 83% yield (**Scheme 5**). Global deprotection of compounds **15** and **17** (**15**–**17**) obtained pentasaccharide **1** and heptasaccharide **2**, respectively, in almost quantitative yields.

## MATERIALS AND METHODS

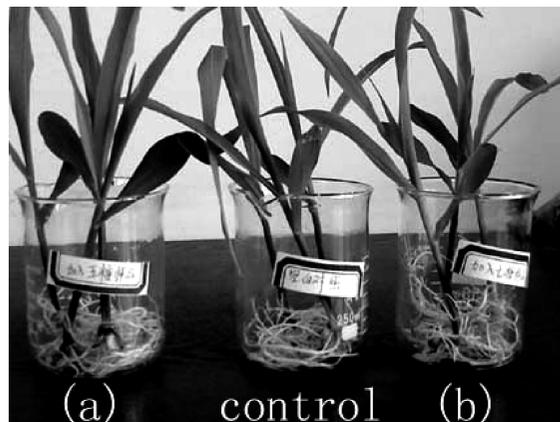
**Instruments.** <sup>1</sup>H NMR and <sup>13</sup>C NMR were recorded with a Bruker ARX 400 spectrometer for the solutions in CDCl<sub>3</sub> or D<sub>2</sub>O. The chemical shifts are given in parts per million downfield from internal Me<sub>4</sub>Si. Mass spectra were measured using a MALDI-TOF-MS with *a*-cyano-4-hydroxycinnamic acid (CCA) as matrix or recorded with a VG



**Figure 1.** Mass of maize plant after exposure to oligosaccharides for one and three weeks. The results are expressed as means (*n* = 4). The asterisk (\*) indicates a significant difference from the control group (\*, *p* > 0.05; \*\*, *p* > 0.01).

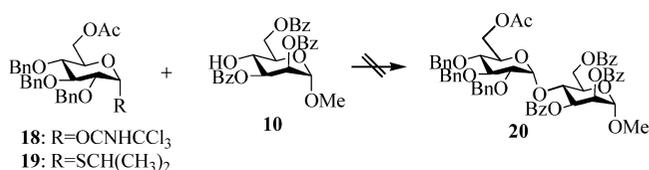


**Figure 2.** Effect on the growth of maize plants treated with oligosaccharide for one week. The control is nutrient media, (a) is nutrient media plus 3 ppm pentasaccharide, and (b) is nutrient media plus 3 ppm heptasaccharide.



**Figure 3.** Effect on the growth of maize roots treated with oligosaccharide for three weeks. The control is nutrient media, (a) is nutrient media plus 3 ppm pentasaccharide, and (b) is nutrient media plus 3 ppm heptasaccharide.

#### Scheme 6



PLATFORM mass spectrometer using the ESI(-) technique to introduce the sample. Thin-layer chromatography (TLC) was performed on silica gel HF<sub>254</sub> with detection by charring with 30% (v/v) H<sub>2</sub>SO<sub>4</sub> in MeOH or in some cases by UV detector. Column chromatography was conducted by elution of a column of silica gel (100–200 mesh) with EtOAc–petroleum ether (60–90 °C) as the eluent. The solutions were concentrated at <60 °C under reduced pressure.

**General Synthetic Procedure for Compound 5.** To a solution of levoglucose **3** (2.0 g, 12.35 mmol) in DMF (20 mL) at 0 °C was added NaH (1.77 g, 74.07 mmol), and the mixture was stirred at 0 °C for 30 min, then BnBr (5.90 mL, 49.38 mmol) was added. The mixture was stirred at rt for 2 h. The reaction mixture was diluted with cold water (50 mL) and extracted with EtOAc (150 mL). The organic phase was dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and concentrated. The residue was purified by column chromatography (4:1 petroleum–EtOAc) to give intermediate **4** (4.96 g, 93%) as a foamy solid. Solid **4** (4.5 g, 10.42 mmol) was dissolved in dry CH<sub>2</sub>Cl<sub>2</sub> (40 mL) and (CH<sub>3</sub>)<sub>2</sub>CHSH (2.83 mL, 31.25 mmol), BF<sub>3</sub>·Et<sub>2</sub>O (2.64 mL, 20.83 mmol) was added, and the mixture was refluxed for 1.5 h. The reaction mixture was neutralized with saturated aq NaHCO<sub>3</sub>, then extracted with CH<sub>2</sub>Cl<sub>2</sub>. The organic layer was dried and concentrated. Purification of the crude product by column chromatography (4:1 petroleum–EtOAc) gave **5** (2.97 g, 56%) as a foamy solid.

**General Synthetic Procedure for Compounds 2–4.** Levoglucose was benzyloated following a reported literature procedure (12), and the following step was the same as the description in the procedure for making compound **5**. Compounds **9** and **10** were prepared following reported literature procedures (12, 13).

**Synthetic Procedure for the Coupling Reaction.** Donor and acceptor were dried together under high vacuum for 1 h, then dissolved in anhydrous CH<sub>2</sub>Cl<sub>2</sub> with N<sub>2</sub> protection. TMSOTf (or TMSOTf/NIS)

was added at –20 °C, and the mixture was stirred for 20 min at –20 °C. TLC indicated that the reaction was complete. The reaction mixture was then neutralized with Et<sub>3</sub>N and concentrated under reduced pressure to dryness and purification of the crude product by column chromatography. The <sup>1</sup>H NMR of compounds **5–17** are listed in **Table 1**.

**Bioassay.** Maize seeds were immersed in 10% H<sub>2</sub>O<sub>2</sub> (wt/wt) solution for 10 min to avoid bacterium contamination and then in 0.1 M CaCl<sub>2</sub> for 30 min to improve germination. Seeds were subsequently germinated on moist filter paper in the dark at room temperature. Two days later, the pregerminated seedlings with seminal leaves were transferred to nutrient media containing 2.0 mM Ca(NO<sub>3</sub>)<sub>2</sub>·4H<sub>2</sub>O, 2.5 mM KNO<sub>3</sub>, 0.5 mM KH<sub>2</sub>PO<sub>4</sub>, 1.0 mM MgSO<sub>4</sub>·7H<sub>2</sub>O, 0.5 mM NH<sub>4</sub>NO<sub>3</sub>, 0.05 mM FeSO<sub>4</sub>·7H<sub>2</sub>O, 0.83 mg/L KI, 8.6 mg/L ZnSO<sub>4</sub>, 6.2 mg/L H<sub>3</sub>BO<sub>3</sub>, 0.25 mg/L Na<sub>2</sub>MoO<sub>4</sub>, 22.3 mg/L MnSO<sub>4</sub>, 0.025 mg/L CuSO<sub>4</sub>, and 0.025 mg/L CoCl<sub>2</sub>. After the plants had grown for 3 days, the media were then supplemented with oligosaccharides at 3 ppm. The media were exchanged every day for fresh media containing oligosaccharides. At the 7th and 21st day of plant growth, plants were wiped with tissue paper and weighed (18–20). The role of oligosaccharides in growth regulation is described in **Figures 1–3**.

## RESULTS AND DISCUSSION

**Synthesis.** Compounds **5** and **7** were synthesized efficiently using 1,6-anhydro-β-D-glucose (levoglucose), which was prepared through pyrolysis of wasted cotton in our laboratory, as the starting material. Nucleophilic attack on C-1 of protected levoglucose by 2-propanethiol, with the help of boron trifluoride etherate, significantly simplified the whole strategy as shown in **Scheme 2**. Compounds **11–17** were prepared in good yield as shown in **Schemes 4** and **5**.

Initially, we attempted to make the desired disaccharide **20** via a coupling reaction of 6-*O*-acetyl-2,3,4-tri-*O*-benzyl-α-D-glucopyranosyl trichloroacetimidate (**18**) and **10**, but failed. Only decomposed **18** and unreacted **10** were found in the reaction mixture. A condensation of isopropyl 6-*O*-acetyl-2,3,4-tri-*O*-benzyl-1-thio-α-D-glucopyranoside (**19**) and **10** was also tried, but was fruitless (**Scheme 6**). It is thus believed that active glycosyl donor does not match with the unreactive acceptor in our design strategies (21). Fortunately, a relatively inactive disaccharide donor **11** was matched with **10**; thus, we obtained the desired trisaccharide **12** under the same reaction conditions.

**Effects on the Growth of Maize Plants.** The results of growth regulation activity tests given in **Figures 1–3** show that synthetic pentasaccharide and heptasaccharide promote the growth of maize plants and also stimulate the growth of roots. After supplementing with oligosaccharides for a week, pentasaccharide increased the mass of plants by 21.4%, while heptasaccharide increased the mass of plants by 10.7%. When supplemented with oligosaccharides for 3 weeks, pentasaccharide improved by 23.6%, and heptasaccharide improved by 32.7%. We also tested the effects of different concentrations of oligosaccharides (3 ppm, 6 ppm, and 9 ppm) on the growth of plants and found that high concentration did not improve the growth rate more.

**Table 2.** Chemical Characteristics of Compounds **1** and **2**

characteristics	compd 1	compd 2
<sup>1</sup> H NMR chemical shift δ (ppm)	4.31–4.40 (H <sub>1</sub> <sup>I</sup> , H <sub>1</sub> <sup>II</sup> , H <sup>III</sup> ), 5.08 (H <sub>1</sub> <sup>IV</sup> ), 4.62 (H <sub>1</sub> <sup>V</sup> ), 3.30 (OCH <sub>3</sub> )	4.32–4.44 (H <sub>1</sub> <sup>I</sup> , H <sub>1</sub> <sup>II</sup> , H <sub>1</sub> <sup>III</sup> , H <sub>1</sub> <sup>IV</sup> , H <sub>1</sub> <sup>V</sup> ), 5.08 (H <sub>1</sub> <sup>VI</sup> ), 4.62 (H <sub>1</sub> <sup>VII</sup> ), 3.31 (OCH <sub>3</sub> )
<sup>13</sup> C NMR chemical shift δ (ppm)	102.6–103.0 (C <sub>1</sub> <sup>I</sup> , C <sub>1</sub> <sup>II</sup> , C <sub>1</sub> <sup>III</sup> ), 100.7 (C <sub>1</sub> <sup>IV</sup> ), 100.0 (C <sub>1</sub> <sup>V</sup> )	102.6–103.0 (C <sub>1</sub> <sup>I</sup> , C <sub>1</sub> <sup>II</sup> , C <sub>1</sub> <sup>III</sup> , C <sub>1</sub> <sup>IV</sup> , C <sub>1</sub> <sup>V</sup> ), 100.7 (C <sub>1</sub> <sup>VI</sup> ), 99.9 (C <sub>1</sub> <sup>VII</sup> )
ESI(+)-MS	calcd for C <sub>31</sub> H <sub>54</sub> O <sub>26</sub> : 842.29 found 860 [M + NH <sub>4</sub> ] <sup>+</sup> , 865 [M + Na] <sup>+</sup>	calcd for C <sub>43</sub> H <sub>74</sub> O <sub>36</sub> : 1166.39 found 1184 [M + NH <sub>4</sub> ] <sup>+</sup> , 1189 [M + Na] <sup>+</sup>

In this article, we first synthesized pentasaccharide and heptasaccharide derivatives **1** and **2**. At present, there have been no related reports about their plant growth activity. In the bioassay, plant auxins such as 2,4-dichlorophenoxyacetic acid and indole-3-acetic acid were used for positive control, but in our experimental conditions, 2,4-dichlorophenoxyacetic acid inhibited maize growth at a concentration of 3 ppm, and indole-3-acetic acid did not have any effects on the growth of maize at 3 ppm. The effect of glucose on maize growth was also tested, but it did not have any effect at this concentration. Compared with the known plant auxins, synthesized oligosaccharides may have different mechanisms affecting the growth of maize.

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