Structure and bioactivity of the polysaccharides in medicinal plant

*Dendrobium huoshanense*

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Abstract—Detailed structures of the active polysaccharides extracted from the leaf and stem cell walls and mucilage of *Dendrobium huoshanense* are determined by using various techniques, including chromatographic, spectroscopic, chemical, and enzymatic methods. The mucilage polysaccharide exhibits specific functions in activating murine splenocytes to produce several cytokines including IFN-γ, IL-10, IL-6, and IL-1α, as well as hematopoietic growth factors GM-CSF and G-CSF. However, the deacetylated mucilage obtained from an alkaline treatment fails to induce cytokine production. The structure and bioactivity of mucilage components are validated by further fractionation. This is the first study that provides clear evidence for the structure and activity relationship of the polysaccharide in *D. huoshanense*. © 2008 Elsevier Ltd. All rights reserved.

1. Introduction

*Dendrobium huoshanense*, Orchidaceae, is a precious herbal plant in Chinese traditional medicine.† The sections of the stem were used for the treatment of salivary, stomach, and ophthalmic disorders. It has slow growth rate and excessive harvesting has left it critically endangered in China. Among the species in the polymorphic *Dendrobium* genus, most studies have focused on small molecules with proved bioactivity, which included the structural types of bibenzyl, phenanthrene, anthracene, fluorene, coumarin, flavone, cinnamate, sesquiterpene, sterol, fatty acid, and alkaloids.‡–†‡ In contrast, the polysaccharide constituents were much less investigated, even though many plant polysaccharides have already shown to trigger immuno-modulatory activities.‡–†‡ In this paper, we report the polysaccharide constituents isolated from the leaves and stems of *D. huoshanense*, and their immuno-modulatory effects in mice splenocytes.

Keywords: *Dendrobium huoshanense*; Cell wall; Mucilage; Polysaccharide; Structure; Immune response.

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2. Results and discussion

2.1. Material and separation procedure

The plant material of *D. huoshanense* was obtained from Yuen-Foong-Yu Biotech Co., Taiwan. A flowchart describing the isolation of polysaccharides from cell wall and mucilage of *D. huoshanense* is shown in Figure 1. The tissues containing non-lignified primary cell walls were collected from leaves and stems at 4 °C. The alcohol-insoluble residue (AIR) was treated with porcine pancreatic-amylase to remove starch. The de-starched cell walls were fractionated by a sequence of extraction using CDTA (cyclohexane-1,2-diamine tetraacetate), Na2CO3, 1 M KOH, and 4 M KOH, as described previously. The remaining insoluble residue was accounted for the α-cellulose fraction. The mucilage polysaccharides collected from leaves and stems did not contain any starch granule as shown by the test with potassium iodide.

2.2. Monosaccharide composition of cell wall

The cell-wall AIR preparation and individual fractions were subjected to hydrolysis with trifluoroacetic acid to release monosaccharides, which were subsequently reduced by NaBH4 and acetylated by Ac2O to give the corresponding alditol peracetates. The composition of
neutral monosaccharides correlated with that of alditol peracetates as determined by the GC–MS analysis (Table 1). The major constituents in the primary non-lignified cell walls of leaves were Ara, Man, Glc, and Gal, with small proportions of Xyl, Rha, and Fuc. On the other hand, the primary cell walls of stems contained mainly Xyl, Glc, Man, and Gal, small proportions of Ara and Rha, and a trace amount of Fuc. It was noted that a greater proportion of Xyl was found in stem (28 mol %) than in leaf (7 mol %) cell-wall hydrolysates, but smaller proportion of Ara in stem (9 mol %) than in leaf (22 mol %). The uronic acid contents in the cell-wall preparations of leaves and stems were 11 mol % and 4 mol %, respectively.

Both the CDTA and Na2CO3 fractions from leaf and stem cell-wall preparations contained large proportions of Man, Gal, and Ara. Much greater amounts of Xyl existed in the 1 M KOH fractions of leaf (28 mol %) and stem (65 mol %) than in the CDTA and Na2CO3 fractions. Glucose was the major component in the 4 M KOH fractions of leaf (28 mol %) and stem (41 mol %). Three uronic acids were found; galacturonic acid (GalA) predominated in the CDTA and Na2CO3 fractions, whereas glucuronic acid (GlcA) and 4-O-methyl glucuronic acid (MeGlcA) existed in small amounts. In the 1 and 4 M KOH fractions of leaves, the content of GlcA was higher than MeGlcA, while the opposite was observed in stems.

Table 1. Composition of neutral monosaccharides and uronic acids in cell walls and individual fractions from the leaves and stems of *D. huoshanense*

<table>
<thead>
<tr>
<th>Neutral monosaccharides</th>
<th>Neutral monosaccharides</th>
<th>Uronic acids</th>
<th>Uronic acids</th>
</tr>
</thead>
<tbody>
<tr>
<td>AIR</td>
<td>Rha</td>
<td>Fuc</td>
<td>Ara</td>
</tr>
<tr>
<td></td>
<td>89</td>
<td>3</td>
<td>3</td>
</tr>
<tr>
<td>CDTA</td>
<td>82</td>
<td>3</td>
<td>&lt;0.5</td>
</tr>
<tr>
<td>Na2CO3</td>
<td>91</td>
<td>&lt;0.5</td>
<td>—</td>
</tr>
<tr>
<td>1 M KOH</td>
<td>89</td>
<td>2</td>
<td>17</td>
</tr>
<tr>
<td>4 M KOH</td>
<td>78</td>
<td>6</td>
<td>5</td>
</tr>
<tr>
<td>Stems</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>AIR</td>
<td>96</td>
<td>1</td>
<td>&lt;0.5</td>
</tr>
<tr>
<td>CDTA</td>
<td>73</td>
<td>&lt;0.5</td>
<td>&lt;0.5</td>
</tr>
<tr>
<td>Na2CO3</td>
<td>76</td>
<td>1</td>
<td>&lt;0.5</td>
</tr>
<tr>
<td>1 M KOH</td>
<td>94</td>
<td>2</td>
<td>3</td>
</tr>
<tr>
<td>4 M KOH</td>
<td>94</td>
<td>2</td>
<td>3</td>
</tr>
</tbody>
</table>

*Composition in mol % as average of duplicate determinations.
*Total content in mol %.
*Alcohol insoluble residue.
*The fraction obtained by extraction with trans-cyclohexane-1,2-diamine-tetraacetate.
*The fraction obtained by extraction with Na2CO3 + 25 mM NaBH4.
*The fraction obtained by extraction with Na2CO3 + 25 mM NaBH4.
*The fraction obtained by extraction with 4 M KOH + 25 mM NaBH4.
GalA is characteristic of the presence of pectic polysaccharides, and the presence of both GlcA and 4-O-methyl GlcA is typical in heteroxylans. Our results indicated that a high proportion of pectic polysaccharides existed in both the CDTA and Na$_2$CO$_3$ fractions, whereas a large amount of hetroxylan existed in the KOH fractions.

2.3. Linkage analysis of the cell-wall fractions

Glycosyl linkages between monosaccharide residues provided information for the structure of polysaccharides. The glycosyl linkages of the cell-wall fractions in leaves and stems of D. huoshanense are summarized in Table 2. In the CDTA fractions of leaves and stems cell-wall preparations, mannose in the form of 4-Man$_p$ and 4,6-Man$_p$ was most abundant. Along with the high proportions of t-Gal$_f$, 4-Gal$_p$, t-Ara$_f$, and 5-Ara$_f$, the composition of CDTA fractions was characterized by galactomannans, galactans, and arabinans (where ‘t’ represents the terminal position of sugars at the non-reducing end). The small amount of 2-Rhap found in the CDTA fractions might be attributed to the presence of rhamnogalacturonans I (RG I).

The glycosyl-linkage compositions in the Na$_2$CO$_3$ fractions were similar to those in CDTA fractions, albeit with much greater proportions of 4-Galp and smaller proportions of 4-Man$_p$ and 4,6-Man$_p$. The presence of galactomannans in the pectic fractions (CDTA + Na$_2$CO$_3$ extractions) of D. huoshanense cell-wall preparations is interesting. According to a previous study, the galactomannan of Leucaena leucocephala can form complexes with divalent metal ions, for example, Co$^{2+}$, Mn$^{2+}$, Ni$^{2+}$, and Zn$^{2+}$. It was possible that the leaf and stem cell-wall preparations of D. huoshanense might contain the complexes of galactomannan–metal ions, which were subsequently extracted by CDTA and Na$_2$CO$_3$ treatments.

High proportions of t-Xyl$_p$, 4-Xyl$_p$, and t-Ara$_p$ glycosyl linkages were observed in the 1 M KOH fractions of primary cell-wall preparations of D. huoshanense. Along with the presence of GlcA and 4-O-methyl GlcA in high proportions (Table 1), trace amount of glucuronoarabinoxylans (GAXs) was observed. Unbranched 4-Xyl$_p$ linkage predominated in the GAXs of D. huoshanense. In comparison, the GAXs found in the primary cell walls of maize (Zea mays, Poaceae) and pineapple (Ananas comosus, Bromeliaceae) contain high proportions of branched 2,4-Xyl$_p$ and 3,4-Xyl$_p$ glycosyl linkages. Furthermore, the GAXs in D. huoshanense differed from that in grasses and pineapple by having no ester-linked hydroxycinnamic acid.

The presence of xyloglucans in the KOH fractions of leaves and stems was indicated by the predominant glycosyl linkages of t-Xyl$_p$, 2-Xyl$_p$, 4-Glc$_p$, and 4,6-Glc$_p$ (see Section 2.6). The presence of t-Fuc$_p$ indicated that the xyloglucans might be fucosylated. A large amount of 4-Man$_p$ glycosyl linkage might be attributable to mannans or glucomannans.

### Table 2. Glycosyl-linkage composition in the cell-wall fractions of D. huoshanense

<table>
<thead>
<tr>
<th>Glycosyl linkage</th>
<th>Leaves</th>
<th>Stems</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>CDTA</td>
<td>Na$_2$CO$_3$</td>
</tr>
<tr>
<td>2-Rhap</td>
<td>2.6</td>
<td>1.4</td>
</tr>
<tr>
<td>t-Fuc$_p$</td>
<td>1.3</td>
<td>—</td>
</tr>
<tr>
<td>t-Ara$_f$</td>
<td>9.3</td>
<td>6.7</td>
</tr>
<tr>
<td>2-Ara$_f$</td>
<td>2.3</td>
<td>2.1</td>
</tr>
<tr>
<td>3,5-Ara$_f$</td>
<td>1.9</td>
<td>5.9</td>
</tr>
<tr>
<td>5-Ara$_f$</td>
<td>5.1</td>
<td>5.7</td>
</tr>
<tr>
<td>t-Xyl$_p$</td>
<td>2.6</td>
<td>1.5</td>
</tr>
<tr>
<td>2-Xyl$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>3-Xyl$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>4-Xyl$_p$</td>
<td>1.8</td>
<td>2.2</td>
</tr>
<tr>
<td>2,4-Xyl$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>3,4-Xyl$_p$</td>
<td>—</td>
<td>3.2</td>
</tr>
<tr>
<td>t-Gal$_p$</td>
<td>8.2</td>
<td>13.5</td>
</tr>
<tr>
<td>4-Gal$_p$</td>
<td>7.0</td>
<td>23.5</td>
</tr>
<tr>
<td>6-Gal$_p$</td>
<td>4.5</td>
<td>0.7</td>
</tr>
<tr>
<td>2-Gal$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>t-Glc$_p$</td>
<td>0.6</td>
<td>1.6</td>
</tr>
<tr>
<td>2,3-Glc$_p$</td>
<td>0.5</td>
<td>—</td>
</tr>
<tr>
<td>3-Glc$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>3,4-Glc$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>4-Glc$_p$</td>
<td>7.3</td>
<td>7.4</td>
</tr>
<tr>
<td>4,6-Glc$_p$</td>
<td>—</td>
<td>—</td>
</tr>
<tr>
<td>4-Man$_p$</td>
<td>26.3</td>
<td>14.1</td>
</tr>
<tr>
<td>4,6-Man$_p$</td>
<td>18.7</td>
<td>10.5</td>
</tr>
</tbody>
</table>

*The polysaccharide sample was subjected to a sequence of permethylation, hydrolysis, reduction, and peracetylation, and the glycosyl linkage was deduced from the GC–MS analysis of the resulting methylated alditol acetates. The data in mole percentage (mol %) are average of duplicate determinations.*
2.4. Analysis of the pectic polysaccharides in CDTA fractions

The above-mentioned analyses of the uronic acid composition and glycosyl linkages indicated the presence of pectic polysaccharides in the CDTA fractions. We further investigated the structure of such pectic polysaccharides by enzymatic digestion and spectral studies. The CDTA fractions of leaf and stem cell-wall preparations were treated with polygalacturonanase to digest the pectins. The mass spectral analysis of the enzymatic digests indicated the presence of oligogalacturonides in 4–15 degrees of polymerization (DP) containing varied degrees of methyl groups. These oligogalacturonides could be derived from homogalacturonans (HGs), which consisted of the methyl ester of GalA. The pectins isolated from *Angelica actitiloba* and *Plantago major* were shown to have anti-complement effect. The plants *Noni* (*Morinda citrifolia*) and *Panax notoginseng*, which contain high level of HGA, have medicinal applications. The high content of HGA in the pectin components of *D. huoshanense* might potentially contribute to its bioactivity.

In agreement with the structural deduction, the $^1$H NMR spectrum (600 MHz) of the polysaccharides in CDTA fractions showed an intense singlet at $\delta$ 3.7 for the methyl ester (CO$_2$CH$_3$) and a broad multiplet at $\delta$ 4.3 attributable to the C-4 protons in GalA or its methyl ester. In the $^{13}$C NMR spectrum, the signals for the C-4 attributable to the C-4 protons in GalA or its methyl ester. The presence of glucomannan was confirmed by using the glucomannan assay kit (Megazyme, see Section 3.1). The de-starched mucilage polysaccharide hydrolysate was analyzed by high pH anion-exchange chromatography–pulsed amperometric detection (HPAEC–PAD) and MALDI-TOF MS (Fig. 2) to show the presence of xyloglucans (XGs). The [M+Na]$^+$ ions in the MS spectrum corresponded to XG (m/z 791, XXGG (m/z 953), XXXG (m/z 1085), XLXG/XXLG (m/z 1247), XXFG (m/z 1393), XLLG (m/z 1409), and XLFG (m/z 1555), where G represents the backbone Glc in β1,4-linkage, X represents the Xyl(1,6)Glc unit, L represents Gal(β1,2)Xyl(1,6)Glc, F represents Fuc(1,2)Gal(1,2)Xyl(1,6)Glc. The similar composition of XGs was found in the stem cell-wall preparation of *D. huoshanense*. Our results showed that the XGs of *D. huoshanense* have both XXGG and XXXG backbones with fucosylated XG oligosaccharides XXFG and XLFG. XGs with terminal Fuc and Gal side chain have been shown to induce anti-tumor activity in cell-based assays.

2.5. Analysis of the heteroxylans in 1 M KOH fractions

The 1 M KOH fraction of leaf cell-wall preparation was hydrolyzed with the enzyme *endo*(1 → 4)-β-d-xylanase, and the released oligosaccharides were analyzed by mass spectrometry. The characteristic signals at m/z 877, 1009, 1141, and 1273 correlated with the [M+Na]$^+$ ions of Pen$_5$GlcA, Pen$_6$GlcA, Pen$_7$GlcA, and Pen$_8$GlcA, whereas the signals at m/z 1023 and 1155 were attributed to the [M+Na]$^+$ ions of Pen$_5$MeGlcA and Pen$_6$MeGlcA (where Pen represents pentoside). These results were consistent with the high contents of GlcA and MeGlcA observed in the uronic acid analysis (Table 1). Similarly, enzymatic digestion and MS analysis of the 1 M KOH fraction of stem-cell wall preparation indicated the presence of the pentose oligomers of Pen$_n$GlcA (n = 5–9), Pen$_n$MeGlcA (n = 4–9), and Pen$_n$(MeGlcA)$_2$ (n = 5–11). The pentose oligomers with substitution of (MeGlcA)$_2$ disaccharide were found in stems only, but not in leaves.

We also confirmed the presence of L-arabinose at the terminal of the oligosaccharides by treatment with α-L-arabinosidase. After the enzymatic removal of arabinosyl residues, the overall mass profile for the GlcA- and MeGlcA-substituted pentose oligomers did not change, albeit all the [M+Na]$^+$ ions showed a reduction of 132 Da in comparison with the original mass signals for the sample without α-L-arabinosidase treatment. This experiment clearly indicated the presence of mono-substituted arabinosyl residue at the terminal, a characteristic of glucuronoarabinoxylans (GAXs).

2.6. Analysis of the xyloglucans in 4 M KOH fraction

The 4 M KOH fraction of leaf cell-wall preparation was subject to enzymatic degradation by *endo*(1 → 4)-β-d-glucanase. The hydrolysate was analyzed by high pH anion-exchange chromatography–pulsed amperometric detection (HPAEC–PAD) and MALDI-TOF MS (Fig. 2) to show the presence of xyloglucans (XGs). The [M+Na]$^+$ ions in the MS spectrum corresponded to XG (m/z 791), XXGG (m/z 953), XXXG (m/z 1085), XLXG/XXLG (m/z 1247), XXFG (m/z 1393), XLLG (m/z 1409), and XLFG (m/z 1555), where G represents the backbone Glc in β1,4-linkage, X represents the Xyl(1,6)Glc unit, L represents Gal(β1,2)Xyl(1,6)Glc, and F represents Fuc(1,2)Gal(1,2)Xyl(1,6)Glc. The similar composition of XGs was found in the stem cell-wall preparation of *D. huoshanense*. Our results showed that the XGs of *D. huoshanense* have both XXGG and XXXG backbones with fucosylated XG oligosaccharides XXFG and XLFG. XGs with terminal Fuc and Gal side chain have been shown to induce anti-tumor activity in cell-based assays.

2.7. Analysis of the polysaccharides in stem mucilage

The presence of glucomannan was confirmed by using glucomannan assay kit (Megazyme, see Section 3.1). The de-starched mucilage polysaccharide hydrolysate was found to contain Man (79%) and Glc (21%) as deduced from the corresponding alditol acetates in the GC–MS analyses. Accordingly, the mass spectrum of *endo*(1 → 4)-β-d-mannanase hydrolysate of the mucilage polysaccharide gave the molecular ions attributable to glucomanno-oligosaccharides of 3–9 DP with partial acetylation.

In agreement with the structural assignments of β-(1 → 4)-d-Glc p and β-(1 → 4)-d-Man p, their anomic carbons appeared at $\delta$ 102.8 (minor component) and 100.6 (major component), respectively. The degree of acetylated mannose was estimated to be 25%. In addition to the signals at $\delta_{HC}$ 2.20/2.07/1.99/1.95/1.94/1.90 for the acetyl group of 2-O-acetyl-β-(1 → 4)-d-mannose, the minor signals (≈5%) occurring at $\delta_{HC}$ 2.50/2.50 and $\delta_C$ 173.5 for the acetyl group of 2-O-acetyl-β-(1 → 4)-d-mannose, the minor signals (≈5%) occurring at $\delta_{HC}$ 2.50/2.50 and $\delta_C$ 173.5 might be attributable to 3-O-acetyl-β-(1 → 4)-d-mannose. Moreover, the mucilage polysaccharide was subject to alkaline treatment to remove acetyl groups. After dialysis, the heteronuclear single quantum coherence (HSQC) spectrum of the deacetylated polysaccharide was recorded, and the $^1$H–$^{13}$C signals were assigned according to the literature. The HSQC spectrum clearly indicated that the deacetylated mucilage polysaccharide was composed of (β1 → 4)-linked d-Man p and (β1 → 4)-linked d-Glc p (Fig. 3).
2.8. Bioactivities of the polysaccharides and the deacetylated polysaccharides from mucilage preparations

*Dendrobium huoshanense* has been reported as one of the best Chinese medicines used for cell growth and immuno-modulatory activities; however, with only little scientific evidence. We first used the MTS assay to determine the cell proliferation effects of the polysaccharide fractions from the different *D. huoshanense* structures. As shown in Table 3, mice splenocytes incubated with mucilage fractions significantly grew faster than untreated cells (154% for stem mucilage and 141% for leaf mucilage vs 100% for control cells). The effects are more obvious compared with the glucomannan from konjac. Both ivory nut mannan (15–80 DP) and mannose β-(1 → 4)-linked oligosaccharide (15 DP) were used for the cellular experiments, but no apparent bioactivity was observed. The cytokine profiling of *Dendrobium* mucilage indicated an increased expression of several cytokines including IFN-γ, IL-10, IL-6, and IL-1α. Significant amounts of hematopoietic growth factors GM-CSF and G-CSF were also induced. We further examined the dose-dependent effects of the stem mucilage polysaccharide on the G-CSF and GM-CSF induction. The stimulatory effects on the production of both factors were dose-dependent, up to 250 μg/mL (Fig. 4).

Most interestingly, we investigated whether structural modification of stem mucilage polysaccharide by deacetylation of 2-O-acetylglucomannan would affect its stimulatory effect on cytokine production. Treatment of cells with the deacetylated stem mucilage polysaccharide was
found to maintain cell proliferation, but fail to induce cytokine production (Table 3).

### 2.9. Validation of structure and bioactivity via fractionation of mucilage components

The extract of mucilage was fractionated by anion-exchange chromatography on DEAE–cellulose (NH<sub>2</sub>/C0 form) column. Elution with distilled water afforded neutral polysaccharides, and the subsequent elution with aqueous NaCl buffer gave the uronic and aldonic acid containing polysaccharides. Six fractions were obtained, and each fraction was subject to dialysis (molecular-weight cutoff = 500) to remove salt. The most potent fraction (6% of weight) in cytokines expression was further separated by size-exclusion chromatography on Sephacryl S-200 into fractions A (31%), B (13%), C (18%), D (17%), E (8%), and F (13%). The cytokines expressions of these polysaccharide fractions were measured by the RT-PCR analyses (Fig. 5). Though G-CSF was not induced by Con A (lane 2), the cytokine expression on treatment with the polysaccharides from mucilage (lane 3) and fractions B–F (lanes 5–9) were confirmed. Fraction C is more active than fraction B for IL-6, G-CSF, and M-CSF, but fraction B is good for IL-6 (a B-cell stimulatory factor).

### Table 3. Cell proliferation and cytokines induced by different polysaccharide fractions from *D. huoshanense*

<table>
<thead>
<tr>
<th>Cell proliferation (%)&lt;sup&gt;a&lt;/sup&gt;</th>
<th>GM-CSF</th>
<th>G-CSF</th>
<th>IFN-γ</th>
<th>IL-10</th>
<th>IL-1α</th>
<th>IL-6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Untreated cells</td>
<td>100</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
</tr>
<tr>
<td>Con A</td>
<td>155 ± 5</td>
<td>408 ± 58</td>
<td>46 ± 9</td>
<td>1504 ± 387</td>
<td>590 ± 75</td>
<td>79 ± 42</td>
</tr>
<tr>
<td>Glucomannan&lt;sup&gt;c&lt;/sup&gt;</td>
<td>106 ± 2</td>
<td>NA&lt;sup&gt;c&lt;/sup&gt;</td>
<td>NA&lt;sup&gt;c&lt;/sup&gt;</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>117 ± 8</td>
</tr>
<tr>
<td>Mannan&lt;sup&gt;d&lt;/sup&gt;</td>
<td>106 ± 2</td>
<td>NA&lt;sup&gt;c&lt;/sup&gt;</td>
<td>NA&lt;sup&gt;c&lt;/sup&gt;</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>92 ± 8</td>
</tr>
<tr>
<td>Leave mucilage</td>
<td>141 ± 2</td>
<td>95 ± 17</td>
<td>582 ± 56</td>
<td>1563 ± 424</td>
<td>241 ± 9</td>
<td>322 ± 48</td>
</tr>
<tr>
<td>Stem mucilage</td>
<td>154 ± 4</td>
<td>118 ± 18</td>
<td>660 ± 58</td>
<td>1465 ± 447</td>
<td>286 ± 30</td>
<td>260 ± 91</td>
</tr>
<tr>
<td>Deacetyl-mucilage</td>
<td>100 ± 1</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
<td>&lt;50</td>
</tr>
</tbody>
</table>

<sup>a</sup> Mice splenocytes (5 × 10<sup>5</sup>) were incubated with 50 μg/mL of different polysaccharide fractions for 60 h, and then subjected to MTS assay as described in Section 3.1. The OD at 492 nm of individual samples was normalized against the OD<sub>492</sub> of the untreated cells, which were defined as 100% cell proliferation.

<sup>b</sup> After cells were incubated with 50 μg/mL of indicated preparations for 48 h, the culture medium was harvested and the cytokine production was analyzed using Quantikine mouse ELISA kits.

<sup>c</sup> Glucomannan from konjac.

<sup>d</sup> Mannan from ivory nut (15–80 DP).

<sup>e</sup> Not available.

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**Figure 4.** The dose–response relationship of mucilage polysaccharide on the production of cytokines and hematopoietic growth factors. Mice splenocytes (5 × 10<sup>6</sup> cells) were stimulated with Con A (2 μg/mL) or stem mucilage polysaccharide (2–250 μg/mL). After 48 h incubation, cell culture supernatant was harvested and analyzed for (a) G-CSF and (b) GM-CSF production using the ELISA kits.

**Figure 5.** Assessment of cytokines (G-CSF, IL-10, IFN-γ, IL-6, GM-CSF, and M-CSF) by RT-PCR on treatment of mouse splenocytes with different drugs for 12 h. Lane 1, untreated; lane 2, Con A (4 μg/mL); lane 3, crude extract of *D. huoshanense* mucilage (100 μg/mL); lane 4, fraction A (100 μg/mL); lane 5, fraction B (100 μg/mL); lane 6, fraction C (100 μg/mL); lane 7, fraction D (100 μg/mL); lane 8, fraction E (100 μg/mL); and lane 9, fraction F (100 μg/mL). Glyceraldehyde-3-phosphate dehydrogenase (GAPDH) is used as an internal control.
The composition and structure for the polysaccharide of fraction B were rigorously determined by a combination of chemical, enzymatic, and spectroscopic methods. The polysaccharide of fraction B was subject to hydrolysis with trifluoroacetic acid to release the monosaccharide components, which were subsequently reduced by NaBH₄ and acetylated by Ac₂O to give the corresponding alditol peracetates. The GC–MS analysis of such alditol peracetates correlated to the polysaccharide of fraction B with the composition of mannose and glucose in a ratio of 10:1. On the other hand, the polysaccharide of fraction B was subjected to methylation, acid-catalyzed hydrolysis, reduction and acetylation to give the corresponding methylated alditol peracetates, which was determined by GC–MS analysis to reveal the glycosyl linkages. By this method, the polysaccharide of fraction B was deduced to have a backbone consisting of (1→4)-linked Manₚ and Glcₚ.

The degree of acetylation in the polysaccharide of fraction B was determined by the NMR analysis (Fig. 6). The NMR spectrum showed the characteristic anomeric protons (H-1) for unacetylated mannopyranoside (Man), 2-O-AcManₚ (Man₂₄), and 3-O-AcManₚ (Man₃₄) at δ 4.64–4.74, 4.93–4.82, and 4.74–4.81, respectively, in a ratio of 66:19:15 deduced from the integration of individual Manₚ signals in the polysaccharide of fraction B.

High performance size-exclusion chromatography (HPSEC) coupled with RI and UV detection was used to estimate the homogeneity and molecular weight of the polysaccharide in fraction B. The retention times on two different HPLC columns (Thermo BioBasic SEC-1000 and TSK-GEL G3000 columns) were measured (Fig. 7), and the molecular weight of the dominant component was estimated to be ~10 kDa by calibration with pullulan standards.

In an alternative approach, diffusion-ordered NMR spectroscopy (DOSY) experiment was used to evaluate the molecular weights of linear and slightly branched water-soluble uncharged polysaccharides. The relationship between the molecular weight (MW) of polysaccharide and self-diffusion coefficient (D) is shown in the following equation:

\[
D = 8.2 \times 10^{-9} \text{MW}^{-0.50} \text{(m}^2 \text{s}^{-1})
\]

The calibration curve of pullulan standards is established to give an exponent value of 0.50, which is generally applied to other neutral polysaccharide systems. As to fraction B, the measured D value of ~10.08 corresponded to an average molecular weight of 9.7 kDa (Fig. 8), in agreement with that deduced from the HPSEC analysis. Thus, fraction B was attributable to a glucomannan polysaccharide of ~60 DP with partial acetylation.

Nuclear Overhauser enhancement (NOE), which depends on the interplay of proximal protons, provides valuable information for assessment of molecular con-
formation and proton assignment. In general, intramolecular NOE signals were observed in the proton pair having disposition of 1,3-diaxial or vicinal equatorial–axial relationship in a pyranosyl ring. Thus, the anomeric proton (H-1) in β-D-mannoside would exert NOEs to H-2, H-3, and H-5, whereas α-D-mannoside would show only a strong NOE between H-1 and H-2. Accordingly, the polysaccharide of fraction B was determined to contain the mannoses with β-linked configuration. Thus, the H-1 resonance of unacetylated manno.pyranosyl residues, which were linked to either manno.pyranosyl or glucopyranosyl residues, was selected for irradiation to give the differential NOE spectrum (Fig. 9). The intra-residue NOEs with the H-2, H-3, and H-5 in manno.pyranoside and the inter-residue NOE with the H-3 in glucopyranoside were observed. This result was in agreement with the β-linked configuration of mannoses in the glucomannan.

The acetyl groups in the polysaccharide of fraction B were removed by alkaline treatment. The de-esterified glucomannan was then subject to enzymatic digestion.
to determine the glycosidic linkages. After digestion with endo-\(\beta\)-mannanase or \(\beta\)-mannosidase, the hydrolysate was found to contain significant amounts of mannose and oligomannosides in a series, by comparison with the standards on HPAEC–PAD analyses. In contrast, no free mannose was released by treatment of the decetylated glucomannan with \(\alpha\)-mannosidase. These enzyme assays thus confirmed the \(\beta\)-linkage of mannoses in the polysaccharide of fraction B.

The 2D-NMR spectra further provided information for the structural assignment. Due to the deshielding effect of acetyl group, three sets of H-2 in 2-O-acetylated \(\beta\)-Manp units occurred at \(\delta 5.45, 5.42,\) and 5.36, whereas two sets of H-2 in 3-O-acetylated \(\beta\)-Manp units appeared at \(\delta 5.05\) and 4.99. Correlations of the protons with carbon signals were found by correlation spectroscopy (COSY), total correlation spectroscopy (TOCSY), HSQC, heteronuclear multiple bond correlation (HMBC), and rotating-frame Overhauser effect spectra (REOSY). The anomeric proton and carbon at the reducing end of \(\alpha\)-Manp residue was readily identified at \(\delta_H 5.14\) and \(\delta_C 94.4\) in the HSQC spectrum of fraction B. The \((1 \rightarrow 4)\)-linked \(\beta\)-Glc residue was characterized by the cross-peak between \(\delta_H 4.48\) and \(\delta_C 102.8\) in the HSQC spectrum.

In addition to the intra-glycosidic correlations of proton and carbon signals, the HMBC spectrum (Fig. 10) revealed the inter-residue cross-peaks of the anomeric proton \((H-I)\) with the C-4 of the adjacent saccharide unit via the \(J\)-coupling. Thus, the HMBC spectrum manifested the existence of the disaccharide units ManGlc, ManMan, ManMan34, \(\text{Man}_{24}\text{Man}_{24}\), \(\text{Man}_{24}\text{Man}_{34}\), \(\text{Man}_{34}\text{Man}_{34}\), \(\text{Man}_{34}\text{Man}_{34}\), and GlcMan.

In the ROESY spectrum (Fig. 11), the anomeric proton (H-1) of acetylated \(\beta\)-Manp displayed three intra-residue cross-peaks \(4,45\) namely H-1/H-2, H-1/H-3, and H-1/H-5 due to their short spatial distances similar to those found in the NOE spectrum (Fig. 9). In addition, the sequential inter-residue correlations, such as ‘H-1(Man)-/H-3(Man)’ \(\delta 4.68/5.05\) and ‘H-1(Man)-/H-3(Man)’ \(\delta 4.84/4.99\), also indicated a great population of the fragments ManGlcMan and ManGlcMan.

On the basis of the afore-mentioned 1D- and 2D-NMR spectral studies and by analogy to the previous relevant reports \(4,44,45\) the \(\text{H}\) and \(\text{C}\) NMR spectral data of the polysaccharide in fraction B are summarized in Table 4. A partial structure for the acetyl-glucomannan is tentatively depicted in Figure 12 to account for the ratio of glucoside to mannoside \((\sim 1:10)\) and the degree of acetylation \((\sim 35\%)\).

2.10. Conclusion

Our study indicates that the polysaccharides in the cell wall and stem of \(D. huoshanense\) are mainly composed of monosaccharides Xyl, Ara, Man, Glc, Gal, and GaLA, along with small proportions of Rha, Fuc, GlcA, and 4-O-methyl GalA. The glycosyl linkages of these monosaccharide residues were determined to give an insight into the structure of polysaccharides. The pectic fractions from the CDTA extraction of the cell wall and stem polysaccharides were found to contain galactomannans, galactans, arabinans, and rhamnogalacturonans I, whereas heteroxylan, glucuronoxarabin-oxylans, and xyloglucans existed in the KOH fractions. Some xyloglucans are modified with terminal fucose and Gal side chain. The stem mucilage contains glucuronman in \(\beta-(1 \rightarrow 4)\)-d-Glc \(\beta-(1 \rightarrow 4)\)-d-Manp linkages with partial acetylated mannosides at the 2- and 3-position. Our study showed that the mucilage polysaccharide exhibited specific functions in murine splenocytes. The mucilage induced several cytokines, including IFN-\(\gamma\), IL-10, IL-6, and IL-1\(\alpha\), and hematopoietic growth factors GM-CSF and G-CSF. However, the deacetylated mucilage obtained from an alkaline treatment failed to induce cytokine production.

The extract of mucilage was further fractionated by chromatography on anion-exchange DEAE–cellulose and Sephacryl size-exclusion columns. The bioactive polysaccharide fraction B was determined to have an average molecular weight of \(\sim 10\) kDa, and its composition and structure were rigorously determined by a combination of chemical, enzymatic, and spectroscopic methods. This is the first study that provides clear evidence for the structure and activity relationship of the polysaccharide in \(D. huoshanense\).

3. Experimental

3.1. Materials

\(Dendrobium huoshanense\) cultivar YFY-HS1 (US Patent No. PP16,746) was obtained from Yuen-Foong-Yu Biotech Co., Ltd, Taiwan. Tamarind XG oligosaccharide standards, the konjac glucomannan, the ivory nut mannan, the polygalacturonanase from \(Aspergillus niger\) (EC 3.2.1.15), \(endo-(1 \rightarrow 4)\)-d-xylanase from \(\text{rumen microorganism}\) (EC 3.2.1.8), \(endo-(1 \rightarrow 4)\)-d-mannanase from \(A. niger\) (EC 3.2.1.78), \(endo-(1 \rightarrow 4)\)-d-glucanase from \(Trichoderma sp.\) (EC 3.2.1.4), and \(\alpha\)-l-arabinosidase from \(A. niger\) (EC 3.2.1.55), and glucomannan assay kit were purchased from Megazyme International Ireland Ltd (Wicklow, Ireland). \(46\) Pancreatic-amylase (EC 3.2.1.1) was purchased from Sigma–Aldrich (St. Louis, MO 63103, USA). XG standards the XGX, XXG, XXFG, and XLFG were kindly gift from Dr. Philip J. Harris (University of Auckland, NZ). Concanavalin A (Con A, C5275) from \(Canavalia ensiformis\) (Jack bean) and Lipopolysaccharide (LPS, L6529) were obtained from Sigma Chemical Co. (Sigma, St. Louis, MO). Complete medium for culture of splenocytes was RPMI 1640 (Gibco, Grand Island, NY) supplemented with 10% FCS (Hyclone, Logan, UT), 0.05 mM 2-mercaptoethanol, and 1% penicillin–streptomycin (Sigma). Sephacryl S-200 was purchased from GE Healthcare, DEAE–cellulose was purchased from Sigma–Aldrich, and dialysis membrane was purchased from SPECTRUM.
3.2. Histochemical analysis of leaf and stem sections

Leaf and stem sections were cut using razor blade and examined histochemically. Secondary lignified cell walls were detected using the color reagent of phloroglucinol–HCl.\textsuperscript{47} Autofluorescence of the cell walls in UV radiation was examined using sections and isolated cell walls mounted in H\textsubscript{2}O and in 0.1 M ammonium hydroxide.\textsuperscript{48} Starch in cell wall and mucilage preparations was detected using potassium iodide, and proteins were detected with Ponceau Red.

3.3. Isolation of cell-wall alcohol-insoluble residue and mucilage

The cell-wall preparations were obtained, respectively, from the leaves and stems at 4 °C. The proportion of tissues containing non-lignified primary cell walls was in-
Figure 11. ROESY spectrum of fraction B shows the intra- and inter-glycosidic correlations (indicated by quotation marks) in the glucomannan.

Table 4. $^1$H and $^{13}$C NMR chemical shifts (in $\delta$ values) for the polysaccharide in fraction B

<table>
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<tr>
<th>Sugar residues Linkage</th>
<th>1</th>
<th>2</th>
<th>3</th>
<th>4</th>
<th>5</th>
<th>6</th>
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<td>→4)-β-D-Glc-(1→</td>
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<tr>
<td>GlcMan</td>
<td>4.48</td>
<td>3.32</td>
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<td>3.65</td>
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<td>δC</td>
<td>102.8</td>
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<td>79.2</td>
<td>74.8</td>
<td>60.6</td>
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<tr>
<td>ManGlc, ManMan</td>
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<td>4.08</td>
<td>3.77</td>
<td>3.79</td>
<td>3.52</td>
<td>3.86</td>
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<td>δC</td>
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<td>70.5</td>
<td>71.9</td>
<td>77.2</td>
<td>75.5</td>
<td>61.1</td>
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<tr>
<td>ManMan$_{34}$</td>
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<td>3.75</td>
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<td>76.4</td>
<td>60.4</td>
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<tr>
<td>Man$<em>{32}$Man$</em>{24}$</td>
<td>4.87</td>
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<tr>
<td>Man$<em>{32}$Man$</em>{34}$</td>
<td>4.84</td>
<td>5.36</td>
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<td>3.50</td>
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<td>δC</td>
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<td>70.6</td>
<td>77.2</td>
<td>76.3</td>
<td>ND*</td>
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<tr>
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<td></td>
<td></td>
<td></td>
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<td></td>
</tr>
<tr>
<td>Man$_{34}$Man</td>
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<td>5.05</td>
<td>4.00</td>
<td>3.60</td>
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</tr>
<tr>
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<td>74.0</td>
<td>73.7</td>
<td>75.7</td>
<td>ND*</td>
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<td></td>
</tr>
<tr>
<td>Man$<em>{32}$Man$</em>{24}$</td>
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<td>4.99</td>
<td>4.03</td>
<td>3.55</td>
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<td>74.4</td>
<td>73.7</td>
<td>75.7</td>
<td>ND*</td>
</tr>
</tbody>
</table>

* Not determined.

Figure 12. A tentative partial structure of the polysaccharide in fraction B.
creased by cutting away the tissues that contain lignified secondary cells walls. The collected tissues (30 g) were homogenized in 50 mM Hepes–KOH buffer (pH 7.2) containing 0.05% of 2-mercaptoethanol. The homogenate was filtered through Miracloth, and washed with the same Hepes–KOH buffer. Liquid nitrogen was added and the residue ground to fine powders. Proteins released from cell were visualized by Ponceau 2R. The fine powders were then mixed with ice-cold water, sonicated (1 min), centrifuged, and the supernatant was carefully removed. This procedure was repeated for additional three times. The pellets were recovered by filtration through Miracloth, and thoroughly washed with water and 70% ethanol until the filtrate becomes clear. Residues were then washed with 100% ethanol, followed by methanol and n-pentane, and dried overnight over silica gel under vacuum.

Mucilage was collected by gently scratching the surface of defrost leaf and stem, then homogenized in 70% ethanol containing 0.05% of 2-mercaptoethanol. The homogenate was sonicated (2 min) and dialyzed against water for 24 h at 4 °C. Mucopolysaccharides were then recovered by lyophilization. Deacetylation of the mucilage polysaccharides was done by treatment with alkali. The sample was treated with 1 M NaOH in the presence of 1% NaBH4 for 1 h at ambient temperature to remove acetyl groups, and suspension was then neutralized with glacial acetic acid to pH 5. Deacetylated mucilage polysaccharides were recovered by dialysis and lyophilization.

### 3.4. Removal of contaminating starch

The dried cell-wall preparation (200 mg) was suspended in 5 mM Tris-maleate buffer (pH 6.9), and heated (85 °C, 5 min) to gelatinize the starch granules. After cooling to 40 °C, porcine pancreatic-amylase (6300 U, Sigma) in 15 mM Tris-maleate buffer (pH 6.9, 25 mL) containing 2 mM CaCl2 was added, and the suspension was incubated for 1 h at 40 °C. The mixture was centrifuged at 500g, and the supernatant was removed. The pellets were suspended with 5 mM Tris-maleate, sonicated, filtered onto nylon mesh (pore size 11 μm), and washed with Milli-Q water until filtrate become clear. A small proportion of residue was tested with a drop of KI to indicate the absence of starch. The pellets of cell walls were dried on nylon mesh using solvent exchange by washing with ethanol, methanol, and n-pentane.

### 3.5. Sequential extraction of cell-wall polymers

The de-starched cell walls (50 mg) were fractionated by a sequence of extraction using 50 mM CDTA (pH 6.5), Na2CO3 + 25 mM NaBH4, 1 M KOH + 25 mM NaBH4, and 4 M KOH + 25 mM NaBH4 as previously described. The remaining insoluble residue was accounted for the β-cellulose fraction.

### 3.6. Neutral monosaccharide composition of cell walls and mucilage

The de-starched cell-wall preparation (10 mg) or individual cell-wall fraction (5 mg) or mucilage preparation (5 mg) were hydrolyzed, respectively, with 4 M trifluoroacetic acid (TFA) at 121 °C for 1 h. The mixture was cooled, and TFA was removed under reduced pressure. The acid hydrolysate was washed with 50% aqueous methanol, and then dried in vacuo. Reduction of monosaccharides in the hydrolysate was carried out by using NaBH4 (80 mg) in MeOH at room temperature for 30 min. This procedure was repeated for 3–5 times to assure complete reduction of monosaccharides to alditos. The mixture was washed with concentrated HCl and MeOH, and then dried in vacuo. The reduction product of alditos was subjected to acetylation using acetic anhydride in pyridine (1:2) at 80 °C for 2 h, followed by incubation at 25 °C for 16 h. The aldito peracetates were extracted by chloroform/2 M HCl (2:1), and washed carefully with saturated NaHCO3. After removal of chloroform, the composition of aldito peracetates was determined by GC–MS analysis.

### 3.7. Linkage analysis of the cell-wall polysaccharides

Each cell-wall fraction (10 mg) was dissolved in DMSO (2 mL), treated with NaH for 2 min with continuous stirring, and ice-cold MeI (1 mL) was added. After stirring at room temperature for 16 h, the methylated polysaccharides in suspension were separated by addition of chloroform/water (2:1). The organic phase was washed with 10% Na2S2O3 and water, and then concentrated under reduced pressure to give the crude product of methylated polysaccharides. The IR and 1H NMR analyses indicated no presence of hydroxyl group. According to the above-described procedure, the product of methylated polysaccharides was similarly digested with TFA, reduced with NaBH4, and acetylated with Ac2O/pyridine to give methylated aldito peracetates. The composition of methylated aldito peracetates was determined by GC–MS analysis.

### 3.8. Uronic acid composition of cell walls and the cell-wall fractions

Using m-hydroxyl diphenyl as the chromogen, the total content of uronic acids in cell walls and each cell-wall fraction was determined according to the previously reported method. Galacturonic acid (GalA), glucuronic acid (GlcA), and 4-O-methylglucuronic acid (MeGlcA) in acid hydrolysates were separated and quantified by HPAEC-PAD (Dionex, Sunnyvale, CA, USA) on a CarboPAC-1 column eluted with isocratic gradient of 150 mM NaOAc containing 100 mM NaOH at a flow rate of 0.25 mL min⁻¹.

### 3.9. Enzyme hydrolysis

The polysaccharide fractions were treated with different enzymes to release oligosaccharides for further analysis. Enzymatic reactions were stopped by boiling for 2 min. The CDTA fractions (2 mg) of leaf and stem wall fractions were treated with polygalacturonase (50 U) from A. niger (EC 3.2.1.15) with 100 mM NaOAc (pH 4.0, 200 μL) for 16 h at 50 °C. The 1 M KOH fractions (2 mg) of leaf and stem wall preparations were treated with endo-(1 → 4)-β-D-xylanase (45 U, EC 3.2.1.8) with
100 mM NaOAc (pH 6.0, 200 μL) for 1 h at 60 °C. The 4 M KOH fractions (2 mg) of leaf and stem wall preparations were treated with \textit{endo}-(1 → 4)-β-D-glucanase (EC 3.2.1.4, 5 U) with 50 mM ammonium formate (pH 5.0, 200 μL) for 16 h at 37 °C. The arabinosyl residues were removed from xylo-oligosaccharides in xylanase digest (1 M KOH fraction of wall preparations) by treatment of \textit{α-L-arabinosidase from \textit{A. niger} (EC 3.2.1.55), and \textit{endo}-(1 → 4)-β-D-xylanase} digest was freeze dried, redissolved with 100 mM NaOAc (pH 6), followed by incubation with \textit{α-L-arabinosidase (10 U) at 40 °C for 24 h.}

The stem mucilage preparation (2 mg) was treated with \textit{endo}-(1 → 4)-β-D-mannanase (EC 3.2.1.78, 5 U) with 100 mM NaOAc (pH4, 200 μL) for 16 h at 40 °C. The deacetylated glucomannan of fraction B was digested with \textit{endo}-β-1,4-mannanase \textit{(A. niger, Megazyme)} in sodium acetate buffer (100 mM, pH 4.5) for 24 h at 37 °C, using 5 μL enzyme per 1 mg of deacetylated glucomannan. Digestion with β-mannosidase that was specific for \textit{β-1,4-mannosyl linkage} \textit{(C. jimi, Megazyme)} was conducted in sodium maleate buffer (100 mM, pH 6.5) for 24 h at 37 °C, using 5 μL enzyme per 1 mg of deacetylated glucomannan. The attempted digestion with \textit{α-2,3,6-mannosidase (Jack bean, Sigma)} was performed in ammonia acetate buffer (50 mM, +1 mM ZnCl₂, pH 5.5) for 24 h at 37 °C, using 5 μL enzyme per 1 mg of deacetylated glucomannan.

### 3.10. GC–MS analysis

The alditol acetates and partially methylated alditol acetates were separated by BPX-70 column (SGE Analytical Science Pty Ltd, Victoria 3134, Australia) on PolarisQ Ion Trap GC–MS/MS System (Thermo Fisher Scientific, Inc., Waltham, MA 02425, USA). Oven temperature was increased from 38 to 150 °C at 50 °C/min, then to 230 °C at 3 °C/min and to 260°C/5 min, with carrier gas (He) at a flow rate of 1 mL/min. Quantification was done by correcting peak area to mol %.

### 3.11. HPAEC–PAD analysis

XG oligosaccharides were separated and characterized by HPAEC–PAD with known standards, using a Dionex BioLC system (Dionex, Sunnyvale, CA, USA). A CarboPac PA1 analytical column (4 mm × 250 mm) and a CarboPac PA1 guard column (4 mm × 50 mm) were used. The XG oligosaccharides were separated using a linear gradient of 50 mM NaOAc + 100 mM NaOH (solvent A) to 100 mM NaOAc + 100 mM NaOH (solvent B) at 1 mL/min over 110 min. After each run, the column was washed for 10 min with solvent B, 5 min with 300 mM NaOH (solvent C), and 30 min with solvent A.

### 3.12. Mass spectrometry

The molecular weights of the sodium adducts of oligosaccharides [M+Na]⁺ were determined using a BioTOF Ultraflex II (Bruker Daltonics, Billerica, MA 01821, USA). The enzyme hydrolysate containing oligosaccharides were mixed with 10 mM of 2,5-dihydroxybenzoic acid and 10 mM NaCl in the ratio of 5:5:3. In MS mode, the spectra were accumulated in average of 2000-3000 shuts.

### 3.13. NMR spectral analysis

The NMR spectra were recorded on Bruker ADVANCE 600 MHz NMR spectrometer (Bruker BioSpin GmbH, Rheinstetten, Germany) with 5 mm Cryoprobe DCI ¹H/¹³C. Samples were dissolved in 1 mL of D₂O. Chemical shifts are given in δ values relative to HOD signal (δH 4.8 at 25 °C) or the H-1/C-1 signals of mannopyranose (δH 4.71/δC 100.6). The HSQC spectra were recorded at 50 °C to shift the HOD peak about 0.2 ppm upfield, so that the obscured signals in the (1 → 4)-linked unacetylated β-Manp residues were revealed. ROESY and TOCSY spectra were recorded, respectively, with mixing times of 100 and 70 ms as well as the relaxation delays of 3 and 2 s.

### 3.14. Determination of molecular weight by DOSY

The sample of fraction B (1 mg) was dissolved in 400 μL of D₂O, and the average molecular weight of the polysaccharide components was determined by DOSY. The values of hydrodynamic radii were determined from pullulan standards. The DOSY experiment was carried out using a stimulated echo sequence incorporating bipolar sine gradient pulses. In addition, the HOD signal was suppressed by means of presaturation. The gradient strength was logarithmically incremented in 32 steps from 2% up to 95% of the maximum gradient strength. The diffusion time was set between 100 and 700 ms. Gradient pulse and longitudinal eddy current were set to 3 and 25 ms, respectively.

### 3.15. Preparation of SpMC (murine spleen mononuclear cell)

Six-to-eight week old male Balb/c were purchased from National Laboratory Animal Center (Taipei, Taiwan) and were sacrificed by cervical dissection. Mice splenocytes were obtained by pressing spleens through a sterilized stainless steel mesh (100 mesh) and washing the passed suspension three times with HBSS (Gibco) after depleting erythrocytes with red blood cell lysis buffer (Sigma). Finally, the viable cells were counted with a hemocytometer using trypsin blue dye exclusion. Single-cell suspensions (5 × 10⁵ cells/mL) were cultured in 10% RPMI-1640 in a humidified 5% CO₂ incubator at 37 °C.

### 3.16. Cytotoxicity/proliferation assay

Mice splenocytes (5 × 10⁵ cells/well) were stimulated directly with Con A (2 μg/mL), commercial mannan analogs (konjac glucomannan and ivory nut mannan) or different polysaccharide fractions (50 μg/mL in water) in 96-well tissue culture plates for 60 h. Afterwards, the cell proliferation was examined with MTS assay using Celltiter 96® aqueous one solution cell proliferation assay kit (Promega, Madison, WI) following the manufacturer’s instructions. Briefly, 20 μL of MTS solu-
tion was added to each well. After 4 h incubation at 37°C, the absorbance at 490 nm was obtained using a plate reader (SpectraMax M2, Molecular Device, Sunnyvale, CA). The optical density at 490 nm (OD 490) for control cells was defined as 100%. Polysaccharide preparation is screened routinely with LAL test (Pyrochrome® Kit, Cape Cod Associates, Falmouth, MA) to ensure no endotoxin contamination. The endotoxin potency (EU/mg) of each polysaccharide fraction (≤ 1 x 10 EU/mg) was shown in much lower level than that of cell culture with lipopolysaccharide (LPS ≥ 1 x 10⁵ EU/mg).

3.17. Cytokine determination by ELISA

The concentrations of IL-1β, IL-6, IL-10, IFN-γ, G-CSF, and GM-CSF in the cell culture supernatants were assayed using the Quantikine mouse ELISA kits (R&D Systems, Minneapolis, MN) according to the manufacturer’s protocol. Mice splenocytes (5 x 10⁶ cells/well) were cultured in the presence of mannan, glucomannan, or different polysaccharide fractions at concentration of 50 μg/mL in 24-well tissue culture plates. Con A at a final concentration of 2 μg/mL was used as a positive control. After 48 h following stimulation, culture supernatants were collected and stored at −80 °C until ELISA analysis. In this study, all the data points are average from three independent experiments and the represented values are average ± standard deviation.

3.18. Cytokine expression inferred from the RNA analysis

Splenocytes were removed from BALB/c mice (male, 6–8 weeks) and treated with fractions A–F (100 μg/mL each) for 12 h. Both the cell pellets and supernatants were collected at the end of the incubation period. RNA was extracted from cell pellets and converted to cDNA. Products from PCR were analyzed by gel electrophoresis on 2% gel to visualize the expressed cytokines.

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References and notes
