Structural Characterization of Anti-Complementary Polysaccharides from the Leaves of Artemisia princeps
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Abstract: Structural characterizations of the anti-complementary acidic heteroglycans, AAF IIb-2 and IIb-3, obtained from the leaves of Artemisia princeps PAMP have been studied. AAF IIb-2 consists of rhamnose, xylose, arabinose, galactose, glucose and uronic acids (galacturonic acid and glucuronic acid) in the molar ratio of 7.6: 2.6: 13.0: 10.9: 3.0: 57.9, and AAF IIb-3 consists of the same sugars in the ratio of 3.9: 2.6: 24.7: 19.7: 2.6: 46.5. Methylation analysis including carboxyl-reduction and also selective enzymolysis using exo-α-L-arabinofuranosidase suggested that AAF IIb-3 has a main chain consisting of (1→4)-linked galacturonic acid and (1→2)-linked rhamnose mostly substituted at the O-4 position. AAF IIb-3 also contained arabinosyl-3,6-galactan moiety and most of the arabinose was present as an α-L-furanosyl residue in the non-reducing terminals and highly branched side chains which mostly attached to the O-3 position of (1→6)-linked galactopyranosyl residue. The basic structure of AAF IIb-2 is similar to that of AAF IIb-3, but IIb-3 has a higher arabinogalactan content than IIb-2.

Materials and Methods

Materials

The leaves of Artemisia princeps Pamp (Japanese name = Gaiyo) were purchased from Uchida Wakanakyo Co. Ltd., Tokyo, Japan. α-L-Arabinofuranosidase from Rhodotorula flava (2) was a kind gift from Dr. Naoto Shibuya (National Food Research Institute, Ministry of Agriculture, Forestry and Fisheries, Japan). Bio-gel P-2 (200—400 mesh) was obtained from Bio-Rad. Sephadex LH-20 was purchased from Pharmacia and Sep-Pak C18 cartridge from Waters Associates Inc. Spectra Pore 6 was obtained from SPECTRUM Medical Industries Inc.

General methods

The total carbohydrate and uronic acid contents were determined by the phenol-sulfuric acid method (3) and m-hydroxydiphenyl method (4), respectively. Using arabinose and galacturonic acid as the respective standards. TLC was performed on cellulose-coated plastic sheet (Merck) with EtOAc-C2H5N-H2O-H2O (5: 5: 1: 3) as the solvent system. Reducing sugars were detected with alkaline silver nitrate (5) and uronic acid with p-anisidine hydrochloride (6). GLC (Shimadzu GC-6A) was equipped with a flame ionization detector (FID) and a glass column (0.3 X 200 cm) of 3% ECNSS-M on Uniport HP at 180° C for component sugar analysis. Nitrogen was used as carrier gas at flow rate of 60 ml/min. Polysaccharide (0.1—1 mg) was hydrolyzed with 2 M TFA at 121° C for 1.5 h. The hydrolyzate containing neutral sugar and uronic acid was converted to alditol acetates by the method of Jones and Albersheim (7). The hydrolyzates were reduced to alditol and aldonic acid by the treatment with NaBH4 in 1 N NH4OH. The aldonic acids and the alditols were separated by the addition of AG 1-X8 resin (OAc° form). Unbound alditols were then acetylated with Ac2O at 121° C for 3 h. The aldonic acids were eluted from the resin in 1 N HCl, and the HCl eluate was evaporated to dryness at 40° C. converting the aldonic acid to aldonolactones. The aldonolactones were reduced with NaBH4 to the corresponding alditol, dried and acetylated. The alditol acetates were analyzed by GLC (8).

Purification of anti-complementary polysaccharides, AAF IIb-2 and IIb-3

The crude polysaccharide from the leaves of A. princeps Pamp was prepared by hot-water extraction and dialysis (1). The crude polysaccharide was purified by ion exchange chromatography on DEAE-Sepharose, affinity chromatography on Ricinus communis-agglutinin conjugated Sepharose and gel filtrations on Sephadex G-100 and Sepharose CL-4B as reported previously (1).

Methylation analysis of polysaccharides

Before per-O-methylation, AAF IIb-2 and AAF IIb-3 were passed through the column of AG 50W-X8 resin (H+ form) in order convert all of the carbohydrate groups into protonated form because this treatment permits more complete O-methylation of uronic acid containing polysaccharides (9). Then AAF IIb-2 and IIb-3 (2 mg) were methylated by the Hakomori procedure (10) and the completeness of the methylation was checked by using tripichylmethane (11). The per-O-methylated polysaccharides were purified by gel filtration on Sephadex LH-20 (12) or reversed-phase chromatography on Sep-Pak C18 cartridge (9). Per-O-methylated polysaccharides were eluted from the Sep-Pak C18 cartridge with 100% CH3CN, followed by 100% EtOH as described by Waege et al. (9).

Introduction

Artemisiae Argyi Folium, the leaves of A. princeps Pamp (Japanese name = Gaiyo) is a well known Chinese crude drug clinically used in the treatment of colic pain, vomiting and diarrhea, and irregular bleeding from uterus.

Previously, we have reported on the purification and chemical properties of anti-complementary polysaccharides, AAF IIb-2 and IIb-3 from the leaves of A. princeps Pamp (1). AAF IIb-2 and IIb-3 are suggested to be structurally related acidic heteroglycans because both polysaccharides were composed of the same component sugars but in different molar ratios (1). The structural characterizations of AAF IIb-2 and IIb-3 are studied in the present paper.

List of Abbreviations

\( C_6H_5N \) : pyidine
\( THF \) : tetrahydofuran
\( TFA \) : trifluoroacetic acid
\( ^1H-NMR \) : proton magnetic resonance
\( NaBH_4 \) : sodium borohydride
\( NaBD_4 \) : sodium borodeuteride
\( Araf \) : arabinofuranose
\( Gal \) : galactopyranose
\( Ara \) : arabinopyranose
\( Xyl \) : xylopyranose
\( Rha \) : rhamnopyranose
\( GalA \) : galacturonic acid

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The fully methylated polysaccharides were reduced with NaBD₄ in 27:73 (v/v) 95% EtOH-THF (13). Methyl-esterified hexosuronic acid residues reduced by this procedure were converted into the corresponding 6,6-dideuteriolabeled hexosyl residue (9). Per-O-methylated and reduced polysaccharides were hydrolyzed with 2 M TFA at 121°C for 1 h. The hydrolysates were reduced to the corresponding partially O-methylated alditol acetates with NaBH₄ in 1 M ammonium in 95% EtOH. The resulting partially O-methylated alditol acetates were acetylated with Ac₂O at 121°C for 3 h, and then alditol acetate derivatives were analyzed by GLC on a Hewlett Packard 5840 A gas chromatograph equipped with dual flame ionization detectors. Samples were injected into a DB-1 capillary column (30 x 0.25 mm i.d., J and W Scientific Inc. U.S.A.) with splitless injection. The flow rate of the carrier gas, helium, was 0.9 ml/min. The gas chromatograph was programmed at 150°C for 1 min, followed by a rate of 2°C/min to 210°C. All GLC flame ionization responses to partially methylated alditol acetates were corrected to mol responses as described by Sweet et al. (14). Partially methylated alditol acetates were also analyzed by GLC-MS (15). GLC-MS was performed on a Hitachi M-80 instrument equipped with OV-1 fused silica capillary column (25 m x 0.25 mm i.d., Gaskuro Kogyo Inc., Tokyo) at 145 to 205°C (1°C/min) with a split ratio of 80:1 and operated at an ionization voltage of 20 eV with an ion source temperature at 180°C. The gas chromatograph was programmed at 150°C for 1 min, followed by a rate of 2°C/min to 210°C. All GLC flame ionization responses to partially methylated alditol acetates were corrected to mol responses as described by Sweet et al. (14). Partially methylated alditol acetates were also analyzed by GLC-MS (15). 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suggested that the low recovery of the reduction product may be caused by the low efficiency of the reduction step and the loss of uronic acid residues during Hakomori-methylation by β-elimination. To obtain more detailed information about the structure of the side chains of AAF IIb-3, AAF IIb-3 was digested with exo-α-L-arabinofuranosidase from R. flava. Then the reaction products were fractionated on a column of Bio-gel P-2 (data not shown), and two carbohydrate fractions that were eluted in the void and retained volumes were obtained. Arabinose was the only sugar detected in the retained volume by TLC and GLC. The material recovered in the void volume consisted of rhamnose, arabinose, xylose, galactose, and glucose in a molar ratio of 0.35 : 0.28 : 0.16 : 1.0 : 0.1 as the neutral sugars. These results lead to the conclusion that most of arabinosyl residues in AAF IIb-3 were linked in an α-configuration in the furanose form. The 1H-NMR spectrum of exo-α-L-arabinofuranosidase digested AAF IIb-3 showed that the signal at δ = 2.0 had decreased in intensity, suggesting that a part of O-acetyl group may be linked to a part of arabinofuranosyl residues. The results of methylation analysis of the arabinofuranosidase digested AAF IIb-3 showed that the molar ratio of the terminal, (1→5)-, and (1→3,5)-linked arabinofuranosyl residues, and (1→3)- and (1→3,6)-linked galactosyl residues had decreased significantly, and that the molar ratio of the terminal (1→6)-linked galactosyl residues had increased compared to AAF II-3 (Table II). When AAF IIb-2 was digested with the same enzyme, similar results with AAF IIb-3 were obtained, but further studies of AAF IIb-2 were not possible because of the small quantity available.

Discussion

The major anti-complementary polysaccharide, AAF IIb-3, was mainly composed of arabinose and galactose while the minor one, AAF IIb-2, was mainly composed of rhamnose, xylose, arabinose, and galactose (1). Previously, a significant amount of galacturonic acid was detected in these heteroglycans by TLC and GLC. The material recovered in the void volume consisted of rhamnose, arabinose, xylose, and galactose (1). Previously, a significant amount of galacturonic acid was detected in these heteroglycans. These results lead to the conclusion that most of arabinosyl residues in AAF IIb-3 were linked in an α-configuration in the furanose form. The 1H-NMR spectrum of exo-α-L-arabinofuranosidase digested AAF IIb-3 showed that the signal at δ = 2.0 had decreased in intensity, suggesting that a part of O-acetyl group may be linked to a part of arabinofuranosyl residues. The results of methylation analysis of the arabinofuranosidase digested AAF IIb-3 showed that the molar ratio of the terminal, (1→5)-, and (1→3,5)-linked arabinofuranosyl residues, and (1→3)- and (1→3,6)-linked galactosyl residues had decreased significantly, and that the molar ratio of the terminal (1→6)-linked galactosyl residues had increased compared to AAF II-3 (Table II). When AAF IIb-2 was digested with the same enzyme, similar results with AAF IIb-3 were obtained, but further studies of AAF IIb-2 were not possible because of the small quantity available.

Table II. Methylation analysis of AAF IIb-3, carboxyl-reduced IIb-3 and IIb-3 α-L-arabinofuranosidase (α-L-Arafase) digest

<table>
<thead>
<tr>
<th>Methylated alditol acetate derivatives</th>
<th>Mol %</th>
<th>Reduced IIb-3</th>
<th>α-L-Arafase treated</th>
<th>Linkage</th>
</tr>
</thead>
<tbody>
<tr>
<td>1, 4-Ac2-2, 3, 5-Me2-arabinitol</td>
<td>19.2</td>
<td>8.7</td>
<td>6.7 (5.1) ‡</td>
<td>Araf—*</td>
</tr>
<tr>
<td>1, 5-Ac2-2, 3, 4-Me2-xylitol</td>
<td>1.6</td>
<td>6.2</td>
<td>2.1 (1.6)</td>
<td>Xyl1—*</td>
</tr>
<tr>
<td>1, 2, 5-Ac3-2, 4-Me2-rhamnitol</td>
<td>4.1</td>
<td>3.0</td>
<td>6.9 (5.2)</td>
<td>Rha1—*</td>
</tr>
<tr>
<td>1, 5-Ac2, 3, 4, 6-Me2-galactitol</td>
<td>—</td>
<td>6.2</td>
<td>(—)</td>
<td>Glc—*</td>
</tr>
<tr>
<td>1, 4, 5-Ac2-2, 3-Me2-arabinitol</td>
<td>16.0</td>
<td>7.0</td>
<td>10.2 (7.7) ‡</td>
<td>Araf1—* or —<em>Araf1—</em></td>
</tr>
<tr>
<td>1, 5-Ac2-2, 3, 4, 6-Me2-galactitol</td>
<td>15.1</td>
<td>12.9</td>
<td>25.6 (19.5) †</td>
<td>Gal1—*</td>
</tr>
<tr>
<td>1, 2, 4, 5-Ac2-3-Me2-rhamnitol</td>
<td>4.0</td>
<td>1.1</td>
<td>6.4 (4.8)</td>
<td>Rha1—*</td>
</tr>
<tr>
<td>1, 3, 4, 5-Ac2-2-Me2-arabinitol</td>
<td>14.6</td>
<td>4.2</td>
<td>3.6 (2.7) ‡</td>
<td>Gal1—*</td>
</tr>
<tr>
<td>1, 3, 5-Ac2-2, 4, 6-Me2-galactitol</td>
<td>7.4</td>
<td>3.8</td>
<td>9.2 (7.0)</td>
<td>Araf1—*</td>
</tr>
<tr>
<td>1, 4, 5-Ac2, 3, 6-Me2-galactitol</td>
<td>8.2</td>
<td>42.7 †</td>
<td>11.2 (8.5)</td>
<td>Gal1—*</td>
</tr>
<tr>
<td>1, 5, 6-Ac2-2, 3, 4-Me2-galactitol</td>
<td>4.6</td>
<td>2.3</td>
<td>12.6 (9.6) †</td>
<td>Gal1—*</td>
</tr>
<tr>
<td>1, 3, 5, 6-Ac2-2, 4-Me2-galactitol</td>
<td>5.2</td>
<td>1.9</td>
<td>5.4 (3.6) ‡</td>
<td>Gal1—*</td>
</tr>
</tbody>
</table>

*a* Methylated different IIb-3 preparation from Table I.

*a* Molar ratio in parenthesis calculated relative to the molar ratio of 1, 5-Ac2-2, 3, 4-Me2-xylitol in IIb-3.

‡ decreased, † increased.
Angelica acutiloba Kitagawa (AR-arabinogalactan IIa and IIb-1) (16, 27, 28) and the arabinogalactans from Zea shoots (29).

When AAF IIb-2 and IIb-3 were digested with exo-α-L-arabinofuranosidase, a significant amount of the arabinosyl residue still remained in the enzyme resistant polysaccharide. The results of methylation analysis of the enzyme digested AAF IIb-3 indicate that the terminal and (1→5)-linked arabinofuranosyl residue had been retained significantly compared to AAF IIb-3. These results suggest that arabinogalactan moiety of AAF IIb-3 contained a highly branched arabinan moiety, possessing attached arabinofuranosyl side chains at O-3 of some of the (1→5)-linked arabinofuranosyl residues (Fig. 1), and the enzyme may be cleaved only at outer side chains of the branching point. The structure of this branched arabinan moiety was also similar to that of AR-arabinogalactan IIb-1 (28) from A. actiloba. The rhamnogalacturonan moieties in AAF IIb-2 and IIb-3 may be present in the inner core region because when AAF IIb-3 was subjected to partial acid hydrolysis, galacturonic acid, rhamnose, and galactose were detected as major component sugars in the acid resistant polysaccharide (unpublished results). This result also suggested that the arabinogalactan side chain may be attached to the rhamnogalacturonan backbone through position 4 of (1→2)-linked rhamnose residue in AAF IIb-2 and IIb-3 (Fig. 1). AAF IIb-2 was shown to have more potent anti-complementary activity than IIb-3 (1), but both contained similar glycosidic linkages with different molar ratios.

Although rhamnogalacturonan moieties and non-reducing end groups of xylosyl and glucuronic acid residues were more predominant than the arabinogalactan moiety in AAF IIb-2, the elucidation of the structural difference between AAF IIb-2 and IIb-3 in association with the activity must await further detailed analysis.

Acknowledgements

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