BRIEF COMMUNICATIONS

GALACTOMANNANS FROM Gleditsia aquatica

R. K. Rakhmanberdyeva

UDC 547.917

We reported previously on the isolation and study of polysaccharides from seeds of eight *Gleditsia* species [1]. In the present article we report results from a study of the properties and structure of galactomannans from seeds of *G. aquatica* Desf introduced into the Republic of Uzbekistan. Aqueous extracts of ground air-dried seeds were precipitated with alcohol to produce water-soluble polysaccharide (WSPS) in 21% yield [1]. The isolated starting WSPS was inhomogeneous. A homogeneous fraction was prepared by treating an aqueous solution of WSPS (300 mL, 0.3%) dropwise with constant stirring with alcohol (150 mL). The resulting solid was separated by centrifugation, washed with alcohol, dehydrated with acetone, and dried over P_2O_5 . The yield of fraction I was 18%. The supernatant solution was treated with another portion (150 mL) of alcohol. The resulting solid was separated as above. The yield of fraction II was 72%. Adding dropwise another portion of alcohol (150 mL) produced fraction III in 0.18% yield. Paper chromatography (PC) of the hydrolysates of all fractions (*n*-butanol:pyridine:water, 6:4:3, anilinium acid phthalate developer) detected galactose and mannose. The ratios of the sugars were determined by GC as the acetates of the aldononitriles [1] and were 1:3.4, 1:4.9, and 1:2.2, respectively. Therefore, all fractions were galactomannans.

Fraction II, designated by us GMA, was investigated in more detail.

GMA was a white powder with a cream tint. It dissolved well in water to form a thick solution with relative viscosity 20.8 (c 0.25%, H_2O). The molecular weight (MW) was determined by ultracentrifugation in a MOM-3170 instrument (50,000 rpm, 20°C) for 30 min. We studied an aqueous solution (1%) of GMA. The MW of GMA was 58,000 \pm 10%. GMA was methylated by the literature method [2] to produce the permethylate with 40.8% O–CH₃ content. The completeness of the methylation was monitored by TLC [CHCl₃:(CH₃)₂CO, 9:1; conc. H_2SO_4 developer] and IR spectroscopy (lack of OH groups). Mainly 2,3,6-tri-O-Me-D-mannose and 2,3,4,6-tetra-O-Me-D-mannose in addition to 2,3,4,6-tetra-O-Me-D-galactose and 2,3-di-O-Me-mannose were identified in the hydrolysate of the permethylate by TLC (benzene:acetone, 2:1; anilinium acid phthalate developer). The last two derivatives were found in smaller quantities.

GMA was separated into fragments using partial acid hydrolysis (H_2SO_4 , 0.5 N, 90 min, 100°C). PC of the products of partial hydrolysis and authentic specimens identified galactose and mannose and oligosaccharides with R_{gal} values of 0.85 (mannobiose), 0.74 (mannotriose), 0.2 (mannotetraose), and 0.09 (mannopentaose).

Thus, a galactomannan consisting of galactose and mannose in a 1:4.9 ratio with MW 58,000 was isolated from seeds of G. aquatica. The macromolecule contains a main β -1–4-mannopyranoside chain in which the mannose units are substituted at C-6 (branching point) by single units of D-galactopyranose. GMA of G. aquatica differs from galactomannans of previously studied Gleditsia species in MW, ratio of monosaccharide units, and degree of polymerization and is the more common type of galactomannans of the families Caesalpiniceae and Fabaceae [3-6].

REFERENCES

- 1. M. R. Mirzaeva, R. K. Rakhmanberdyeva, E. L. Kristallovich, D. A. Rakhimov, and N. I. Shtonda, *Khim. Prir. Soedin.*, 27 (1998).
- 2. S. Hakomori, J. Biochem. (Tokyo), **55**, 205 (1964).

S. Yu. Yunusov Institute of the Chemistry of Plant Substances, Academy of Sciences of the Republic of Uzbekistan, Tashkent, fax (99871) 120 64 75. Translated from Khimiya Prirodnykh Soedinenii, No. 2, p. 176, March-April, 2005. Original article submitted December 27, 2004.

- 3. N. M. Mestechkina and V. D. Shcherbukhin, *Prikl. Biokhim. Mikrobiol.*, **26**, 799 (1990).
- 4. R. K. Rakhmanberdyeva, M. R. Mirzaeva, D. A. Rakhimov, and N. D. Abdullaev, Khim. Prir. Soedin., 566 (1999).
- 5. M. R. Mirzaeva, R. K. Rakhmanberdyeva, and D. A. Rakhimov, *Khim. Prir. Soedin.*, 517 (1999).
- 6. O. V. Anulov, S. F. Ponomarenko, A. V. Egorov, and V. D. Shcherbukhin, *Rastit. Resur.*, 1, 80 (2003).