

POLYSACCHARIDES OF *Eremurus*.

## XXIX. ISOLATION OF A GLUCOMANNAN

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The tuberous roots of some species of the genus *Eremurus* contain a water-soluble glucomannan, which was first isolated from *E. regelii* Vved. and has acquired the name "éremuran" [1]. The properties, structure, and dynamics of the accumulation of éremuran have been found in their general outlines and methods for its practical use have been proposed [2-6]. The preparation of a low-molecular-mass glucomannan and its use as a bioreagent in medicine formed the object of the present investigation.

The air-dry tuberous roots of *E. regelii* (1 kg) ground to 1 mm were extracted with water (1:10) with stirring at room temperature for 2 h. The extract was separated off by centrifugation and was treated with ethyl alcohol in a ratio of 1:2. The precipitate that deposited was dissolved in aqueous trichloroacetic acid and the solution was centrifuged. It was then kept at  $80 \pm 2^\circ\text{C}$  for 40 min and was treated with ethyl alcohol. The resulting precipitate was washed with increasing concentrations of alcohol (70-96%), dried in vacuum over  $\text{P}_2\text{O}_5$ , and ground to a powder. The yield of polysaccharide was 10-12% of the weight of the air-dry raw material.

The polysaccharide consisted of a white amorphous powder with a yellowish tinge, mp  $272\text{-}276^\circ\text{C}$ ,  $[\alpha]_D^{20} -31 \pm 2^\circ$  ( $c$  2;  $\text{H}_2\text{O}$ ). Soluble in water, practically insoluble in organic solvents (acetone, alcohol, chloroform, ether). The qualitative composition of the polysaccharide — glucose and mannose, 1:2, respectively — was determined by gas-liquid chromatography [7]. It was a glucomannan. Its IR spectrum revealed the following absorption bands ( $\text{cm}^{-1}$ ): 3300-3600 (OH group); 1750, 1250 (ester group); 1650 (adsorbed water); 885, 815 ( $\beta$ -glycosidic bond). The chemical shifts in the  $^{13}\text{C}$  NMR spectrum of the glucomannan are given below: (recording and interpretation of the spectrum in accordance with [8, 9]):

	C <sub>1</sub>	C <sub>2</sub>	C <sub>3</sub>	C <sub>4</sub>	C <sub>5</sub>	C <sub>6</sub>
$\beta$ -D-Man <sub>D</sub>	101.3	71.5	72.9	77.8	76.2	61.8
$\beta$ -D-Glc <sub>D</sub>	103.8	74.4	76.6	80.0	75.5	61.8

In the spectrum weak signals were detected with chemical shifts of 21.7 and 173.5 ppm relating to the  $\text{CH}_3$  and CO atoms of an ester group. The signal at 71.5 ppm showed that the glycosidic centers of 4-substituted mannopyranose residues had the  $\beta$ -configuration of the glycosidic center, and the signal at 103.8 ppm that the glucopyranose residues also had the  $\beta$ -configuration.

The glucomannan is a native acetylated polysaccharide having a linear chain the hexopyranose residues of which are linked with one another by  $\beta$ -1 $\rightarrow$ 4 bonds.

The results of biochemical investigations have shown that the glucomannan may be used in immunology and in diagnostic reagents to create a density gradient for separating the cells of peripheral blood in order to obtain a pure population of viable lymphocytes [10].

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