R. N. Tursunova, V. A. Maslennikova, and N. K. Abubakirov

UDC 547.918:547.926

Continuing an investigation of plants of the genus Cynanchum (family Asclepiadaceae), we have studied the pregnane glycosides from the roots of the Cynanchum maximoviczii Pobed. (plant collected by P. G. Gorov on the island of Kunashir of the Kurile group).

The air-dry roots (325 g) were extracted with methanol at room temperature. When the solvent was evaporated, a large amount of crystals deposited which, after recrystallization from ethanol, were identified as sucrose (yield 4% on the weight of the raw material). After the separation of the sugar and further evaporation of the solvent in vacuum, 35 g of a resinous sum of extractive substances was obtained. From part of this material (5 g), by preparative chromatography [SiO<sub>2</sub> with 5% of gypsum, chloroform—benzene—methanol (5:5:2)] we isolated in the amorphous form 0.8 g of individual pregnane glycoside (I) showing a positive xanthydrol reaction for a 2-deoxysugar. Composition of the glycoside (I):  $C_{62}H_{100}O_{23}$ ,  $[\alpha]_D^{22}$  -24.0 ± 2° (c 1,5; methanol);  $\lambda_{\rm max}^{C_2H_5OH}$  220 nm (log  $\epsilon$  3.84).

Under the action of the pancreatic juice of the snail \*Helix plectotropis\*, the glycoside (I) was hydrolyzed, forming D-glucose and a desgluco compound (II) with the composition \$\C\_{56}H\_{90}O\_{18}\$. According to analysis, glycoside (II) contained four methoxy groups. The acid hydrolysis of compound (II) with 0.1 N sulfuric acid led to the formation of a mixture of two isomeric aglycones: cynanchogenin and sibirigenin [TLC, ethyl acetate—n-hexane (1:1); ethyl acetate—chloroform (4:1)]. In the sugar moiety of the glycoside (II) we found D-cymarose and D-oleandrose [PC, toluene—butan-1-ol-water (4:1:5), TLC, chloroform methanol (9:1)]. Similar products were isolated when desglucosibiricoside D, which we have isolated previously from \*Cynanchum sibiricum Willd\*. [1], was hydrolyzed.

On the basis of the experimental facts given, glycoside (I) was identified as sibiricoside D. The identity of the samples of sibiricoside D from the two sources of raw material was also confirmed by a direct comparison of the samples.

## LITERATURE CITED

1. R. N. Tursunova, V. A. Maslennikova, and N. K. Abubakirov, Khim. Prirodn. Soedin., 171 (1975).

Institute of the Chemistry of Plant Substances, Academy of Sciences of the Uzbek SSR. Translated from Khimiya Prirodnykh Soedinenii, No. 4, p. 522, July-August, 1975. Original article submitted March 18, 1975.

<sup>© 1976</sup> Plenum Publishing Corporation, 227 West 17th Street, New York, N.Y. 10011. No part of this publication may be reproduced, stored in a retrieval system, or transmitted, in any form or by any means, electronic, mechanical, photocopying, microfilming, recording or otherwise, without written permission of the publisher. A copy of this article is available from the publisher for \$15.00.