As reported previously, from the seeds of <u>Euonymus medirossica</u> Klok. five cardenolides have been isolated – evomonoside [1], evonoloside [2], glucoevonogenin [3], glucoevonoloside [3], and evonogenin [4].

By chromatographing the combined polar cardenolides (the method for their purification has been described previously [3]) on cellulose in the toluene-butan-1-ol (1:1)/water system and on alumina (activity grade V; eluent: chloroform-methanol) we have obtained another cardiac glycoside in the amorphous, but chromatographically individual, state with $[\alpha]_D$ -38.5 ± 2° (c 2.00; methanol); with concentrated H_2SO_4 it forms the following colors changing with time: 0 min - orange-brown; 25 min - red; 1 h - violet. Under the action of an enzyme preparation from the grape snail it hydrolyzes, forming D-glucose and evomonoside (identified by paper chromatography). The assumption that this glycoside is evonoside (digitoxigenin 3-[β -D-glucosyl- β -D-glucosyl- α -L-rhamnoside] [5] was confirmed by a direct comparison of it with a sample kindly given to us by Prof. T. Reichstein. Evonoside has recently been isolated from the seeds of jute by Abubakirov et al. [6].

To determine the position of attachment of the terminal D-glucose in evonoside, the glycoside was partially hydrolyzed with 0.2 N sulfuric acid (3 h at 95-100°C). The acid was neutralized with barium carbonate, and the hydrolyzate was analyzed by paper chromatography. The following substances were detected: D-glucose, L-rhamnose, and gentiobiose. On this basis, it may be considered with high probability that the terminal D-glucose is attached to the preceding one at C_{6} ". This conclusion is also confirmed by the results of methylation (with methyl iodide in dimethylformamide, heated with silver oxide under reflux for 8 h) and by the hydrolysis of methylated evonoside. Among the hydrolysis products a monosaccharide having the same R_f values of chromatograms as 2,3,4-tri-O-methyl-D-glucose was found.

The structure of the carbohydrate component of evobioside has been established by one of us previously [7]: evobioside is digitoxigenin 3β -O-[O- β -D-glucopyranosyl-(1 \rightarrow 4)- α -L-rhamnopyranoside]. The latter may be considered as a component part of evonoside, since it was obtained by Reichstein [8] by the enzymatic hydrolysis of the triglycoside under investigation.

Thus, evonoside can be identified as digitoxigenin 3β -O-[O- β -D-glucopyranosyl-(1 \rightarrow 6)-O- β -D-glucopyranosyl-(1 \rightarrow 4)- α -L-rhamnopyranoside].

LITERATURE CITED

- 1. S. G. Kislichenko, I. F. Makarevich, and D. G. Kolesnikov, Khim. Prirodn. Soedin., 440 (1966).
- 2. S. G. Kislichenko, I. F. Makarevich, and D. G. Kolesnikov, Khim. Prirodn. Soedin., 241 (1967).
- 3. S. G. Kislichenko, I. F. Makarevich, and D. G. Kolesnikov, Khim. Prirodn. Soedin., 193 (1969).

Khar'kov Scientific-Research Institute of Pharmaceutical Chemistry. Translated from Khimiya Prirodnykh Soedinenii, No. 6, pp. 804-805, November-December, 1973. Original article submitted May 21, 1973.

© 1975 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.

- 4. S. G. Kislichenko, I. F. Makarevich, I. P. Kovalev, and D. G. Kolesnikov, Khim. Prirodn. Soedin., 386 (1969).
- 5. A. Meyrata and T. Reichstein, Pharm, Acta Helv., 23, 135 (1948).
- 6. V. A. Maslennikova, R. U. Umarova, and N. K. Abubakirov, Khim. Prirodn. Soedin., 378 (1971).
- 7. I. F. Makarevich, Khim. Prirodn. Soedin., 681 (1972).
- 8. F. Santavy and T. Reichstein, Helv. Chim. Acta, 31, 1635 (1948).