

In the present paper we give the results of a study begun previously [1-3] of the glycosidic alkaloids of *Solanum dulcamara* L. (bitter nightshade), which grows abundantly in Belorussia. The air-dry epigeal part of the plant collected in the flowering period was extracted with 2% acetic acid. The extract was treated with 25% ammonia solution and the precipitate of "crude" glycosidic alkaloids was dried and extracted with boiling methanol. The combined glycosidic alkaloids were separated by chromatography on a column of silica in the ethyl acetate-methanol (15:10) system. Two individual compounds were obtained.

After recrystallization from methanol, the first substance had the composition $C_{50}H_{83}NO_{21}$, mp 268-270°C (decomp.), $[\alpha]_D^{20} - 58^\circ$ (c 0.6; methanol). The IR spectrum had absorption bands at, cm^{-1} : 3400 (OH); 1140 (>NH); 1050 (CH_2-OH). UV spectrum of a solution of the substance in 94% sulfuric acid: λ_{max} 255, 318, 400 nm (log ϵ 4.14, 4.85, 3.87).

The composition of the second substance was $C_{45}H_{75}NO_{16}$, mp 246-248°C (decomp.) $[\alpha]_D^{20} - 39^\circ$ (c 0.38; methanol). IR spectrum, cm^{-1} : 3400 (OH), 1140 (>NH), 1050 (CH_2-OH). UV spectrum of a solution of the substance in 94% sulfuric acid: λ_{max} 267, 325, 410 nm (log ϵ 4.01, 4.31, 3.66).

The acid hydrolysis of the two substances separately gave the same aglycone, with the composition $C_{27}H_{45}NO_2$, mp 205°C (decomp.), $[\alpha]_D^{20} - 52.5^\circ$ (c 0.6; chloroform). IR spectrum, cm^{-1} : 3600 - 3460 (alcoholic OH); 2750 - 2850 (C - CH_3); 1140 (>NH). UV spectrum of a solution of the substance in 94% sulfuric acid: λ_{max} 340, 397 nm (log ϵ 3.94, 3.93). From its IR spectrum, the UV spectrum of a solution in 94% sulfuric acid, its behavior on paper and thin-layer chromatography, and by the constants given, we identified the aglycone as soladulcidine [4-6].

The analysis of the carbohydrates in an acid hydrolyzate of each glycosidic alkaloid [7] showed that the first substance contained two molecules of D-glucose and one molecule each of D-galactose and D-xylose, and the second substance contained one molecule each of D-glucose, D-galactose, and L-rhamnose.

On the basis of what has been stated above, we identified the compounds as, respectively, α - and β -soladulcines [6, 8].

LITERATURE CITED

1. B. T. Ivanchenko and E. A. Tukalo, Proceedings of the 26th Scientific Session of the Vitebsk State Medical Institute [in Russian], Minsk (1968), p. 132.
2. E. A. Tukalo and V. T. Ivanchenko, Proceedings of the 27th Scientific Session of the Vitebsk State Medical Institute [in Russian], Vitebsk (1969), p. 260.
3. E. A. Tukalo and V. T. Ivanchenko, Sb. Nauchn. Trudov Vitebskogo Gosmedinstituta, No. 13, Book 1, 53 (1969).
4. P. Tuzson and Z. Kiss, Acta Chim. Hungar., 12, 31 (1957).
5. L. Toldy, Acta Chim. Hungar., 16, 403 (1958).
6. K. Schreiber, Planta Med., 6, 93 (1958).
7. G. N. Zaitseva and T. P. Afanas'eva, Biokhimiya, 22, No. 6, 1035 (1957).
8. M. Alkemeyer and H. Sander, Chemie und Biochemie der Solanum-Alkaloide., Tagungsberichte, Deut. Akad. Landwirtschaftswiss., Berlin, No. 27, 23 (1961).

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