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GITALOXIN FROM Digitalis ciliata

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In the adsorption chromatography of the less polar glycosides of the leaves of $Digi-talis\ ciliata$ Trautv. on a column of alumina, we isolated digitoxin and digitoxigenin bisdigitoxoside [1, 2]. When the column was eluted further with chloroform—ethanol (95:5), an individual compound was obtained in the form of a grayish white powder which, after treatment with activated carbon and recrystallization from dilute methanol, formed white acicular crystals with the composition $C_{42}H_{64}O_{15}$, mp 243-245°C. On a paper chromatogram in various systems of solvents the substance was located in the region of an authentic sample of gitaloxin. The Legal, Raymond, Kedde, and Keller—Kiliani reactions were positive. In the latter case, the layer of acetic acid was colored blue and the sulfuric acid was colored crimson. After treatment with the Svendsen-Jensen reagent the bright blue fluorescence in UV light characteristic for gitaloxigenin derivatives appeared. In the UV spectrum λ_{\max}^{C2450H} 220 nm (log ϵ 4.8). The IR spectrum was also identical with that of the foxglove cardenolides [3, 4]. Under the action of alkalis it underwent saponification with the formation of gitoxin. Acid hydrolysis of the glycoside with 0.1 N H₂SO₄ gave the aglycone gitoxigenin (mp 222-224°C) and the sugar digitoxose.

The results that we obtained permitted the conclusion that the substance studied was gitaloxigenin 3-0-tridigitoxoside or gitaloxin [4, 5].

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