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C-GLYCOSIDES OF Stellaria holostea

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Continuing a search for biologically active substances from plants of the genus Stellaria L. [1, 2], we have studied the flavonoid composition of the epigeal part of Stellaria holostea L. (easterbell starwort) collected in the environs of Khar'kov in the flowering period.

Qualitative reactions and one- and two-dimensional chromatography showed the presence in the herbage of the plant under investigation of about six flavonoids. Column chromatography on polyamide sorbent led to the isolation of substance A with mp 237-241°C, $[\alpha]_D^{20} +28^\circ$ (c 0.1; ethanol), $E_{1\text{cm}}^{1\%} = 530$, R_f (15% acetic acid, asc.) 0.42; λ_{max} (in ethanol): 350, 258, 270 nm.

The IR spectrum showed absorption bands characteristic for C-glycosides (1010-1040 cm^{-1}) [3].

For exhaustive hydrolysis we used a mixture of 30% solutions of sulfuric and acetic acids [3]. Hydrolysis for 10 h gave the aglycone, D-glucose, and D-arabinose. From the results of UV spectroscopy, alkaline degradation, and a mixed melting point with an authentic sample, the aglycone was identified as luteolin. Hydrolysis in 10% ethanolic hydrochloric acid permitted the following isomerization to be observed. On acid hydrolysis, substance A gave two compounds (A → A + B) with R_f 0.42 and 0.16 (15% acetic acid). Substance B with mp 263-265°C, $[\alpha]_D^{20} +20^\circ$ (c 0.1; ethanol) gave the same products. This enabled us to state that they were luteolin C-glycosides. Spectral investigations in the UV regions of substances B revealed free 3',4',5,7-hydroxy groups.

Its chromatographic mobility on paper, the absence of a depression of the melting point of mixtures with authentic samples, and the identity of the IR spectra of these compounds permitted substances A to be identified as homoorientin and substance B as orientin [4]. This is the first time that flavonoids from Stellaria holostea have been investigated.

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