FLAVONOIDS OF Atraphaxis spinosa. I

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From the leaves of Atraphaxis spinosa L. growing in the Alma-Ata region of the Kazakh SSR, we have isolated the combined flavonoids, consisting of 14 substances, by methanol extraction.

Chromatography on Kapron of the methanolic extract yielded a flavone glycoside (I) with mp 179-181°C in the form of lemon-yellow needles (from aqueous methanol), R_f 0.40 [butan-1-ol-acetic acid-water (4:1:5)], 0.34 (2% acetic acid), $[\alpha]_D$ -167.2° (c 0.16, methanol).

The IR spectrum of substance I shows absorption bands at (cm⁻¹) 3360 (hydroxy groups), 1669 carbonyl of a γ -pyrone bound by a hydrogen bond with a 5-OH), 1600, 1580, 1510 (benzene ring), 2970, 2850 (methoxy groups), and 890, which is characteristic for the β form of the pyranose ring of a sugar [1, 2].

UV spectrum of the glycoside with additives: λ_{max} 338, 260 nm (C_2H_5OH), 410, 260 ($+C_2H_5ONa$), 340, 266 ($+CH_3COONa$), 370, 266 ($+H_3BO_3$), 390, 260 nm ($+ZrOCl_2$) [13].

UV spectrum of the aglycone: λ_{max} 342, 266 nm (C₂H₅OH), 365, 266 (+CH₃COONa + H₃BO₃), 387, 275 (+C₂H₅ONa), 365, 273 (+ZrOCl₂), 365, 268 nm (+ citric acid).

The acid and enzymatic hydrolysis of the substance gave the aglycone (68%) and L-rhamnose. The aglycone has mp 306-308°C and R_f 0.25 [$C_6H_6-CH_3COOH-H_2O$ (125:72:3)] (1), 0.75 [HCOOH-CH₃COOH-H₂O (2:10:3)] (2), and 0.62 [butan-1-ol-CH₃COOH-H₂O (4:1:5)] (3).

Microanalyses showed the presence of one methoxy group both in the glycoside and in the aglycone.

The demethylation of the aglycone [4] gave luteolin with R_f 0.53 (1), 0.66 (2), and 0.20 (3), identified by paper chromatography with a reference sample.

Thus, on the basis of chemical and spectral studies, substance I has been characterized as 7-O-methyl-4'-O- β -L-rhamnopyranosylluteolin, and we have called it spinoside.

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