NEW FLAVONE GLYCOSIDE FROM Scutellaria ramosissima

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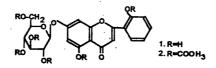
Many species of plants of the genus *Scutellaria* L. (fam. Lamiaceae) are used in folk and scientific medicines [1]. Their high pharmacological activity is due to the presence of flavonoids [2]. We are studying the flavonoids of plants of this genus growing in Central Asia.

We have previously isolated a number of flavonoids from the epigeal part and roots of *S. ramosissima* M. Pop. [3, 4]. Continuing this investigation, from an ethyl acetate fraction of an alcoholic extract, by chromatography on a column of silica gel we have isolated a new flavonoid (1) with the composition $C_{21}H_{20}O_{10}$, mp 280-281°C. Its UV spectrum (λ_{max} (ethanol) 269, 325 nm; +CH₃COONa, 268, 327) was characteristic for flavone derivatives.

According to its PMR spectrum, the substance under consideration was a glycoside and contained the signals of protons at (ppm) 3.88-4.43 (protons of the carbohydrate moiety), 5.63 (d, 6.5 Hz, H-1"), 6.59 (d, 2.0 Hz, H-6), 6.65 (d, 2.0 Hz, H-8), 6.93-7.07 (m, H-3', 5'), 7.08 (s, H-3), 7.28 (dt, 2.0 and 8.0 Hz, H-4'), 7.72 (dd, 2.0 and 8.0 Hz, H-6') and 13.48 (br.s, 5–OH).

The acid hydrolysis of flavonoid (1) yielded 2',5,7-trihydroxyflavone, $C_{15}H_{10}O_5$ (M⁺ 270) with mp 281-283°C [5] and D-glucose. The acetylation of glycoside (1) led to the hexaacetyl derivative (2) with mp 98-100°C the mass spectrum of which contained, in addition to the weak peak of the molecular ion with m/z 684, intense peaks of fragmentary ions: of a tetraacetylhexose residue with m/z 331, 211, 169, and 109, and of an aglycon with m/z 270.

The position of attachment of the carbohydrate residue to the 7-OH group of the aglycon was established by a study of the UV spectra of the glycoside (1) and its aglycon: on the addition of CH_3COONa no bathochromic shift of the absorption maxima was observed, which showed the glycosylation of the 7-OH group of the flavone. In the PMR spectrum of glycoside (1), the signal of the anomeric proton of the D-glucose residue appeared at 5.63 ppm in the form of a doublet with a SSCC of 6.5 Hz. This showed a β -glycosidic bond of the carbohydrate residue with the aglycon.



Thus, the flavone glycoside (1) had the structure of $7-\beta$ -D-glucopyranosyloxy-2',5-dihydroxyflavone.

In the roots of *S. ramosissima*, in addition to those isolated previously, we detected a flavonoid, $C_{18}H_{16}O_7$ (M⁺ 34), with mp 259°C, λ_{max} (ethanol) 270, 341 nm. On the basis of its spectral characteristics it was identified as rivularin [6] (2',5-dihydroxy-6',7,8-trimethoxyflavone).

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