

LUTEOLIN AND ITS 7-O-GLUCOSIDE FROM THE
GENUS *Trifolium*

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The epigeal part of the clover varieties *Trifolium trichocephalum* M.B., *T. canescens* Welld., *T. caucasicum* Tausch. — the section *Stenostoma* Gib. et Belli. — and *T. apertum* Bobr. — the section *Hiantia* Bobr. — family Leguminosae, collected in the flowering period in the Stavropol' and Krasnodar territories was extracted successively with chloroform and 96% ethanol. The resulting ethanolic extracts were evaporated under vacuum, and the residue was diluted with water. After a day, the chloroform-saturated aqueous extract deposited a light yellow precipitate from which two substances of flavonoid nature were isolated by fractional crystallization from aqueous methanol with subsequent chromatography on polyamide.

The first substance had the composition $C_{21}H_{20}O_{11}$, mp 256–258°C (from aqueous methanol) $[\alpha]_D^{20} - 53^\circ$ (c 0.57; methanol–pyridine, 5:1). The acetate had mp 237–240°C (from petroleum ether–chloroform, 4:1).

Acid hydrolysis with 10% sulfuric acid showed that its sugar residue was D-glucose. Under the action of the enzyme of *Aspergillus oryzae* the substance split into the aglycone and the sugar component, which shows the β configuration of the glycosidic bond. The aglycone was identified by its melting point and UV and IR spectra as luteolin. Thus, the flavonoid is luteolin 7-O- β -D-glucopyranoside [1].

The second substance, with the composition $C_{15}H_{10}O_6$, formed crystals with mp 328–330°C (from methanol). Melting point of the acetate 223–225°C (from methanol–chloroform, 4:1). From its UV and IR spectra and a mixed melting point, the substance was identified as luteolin (3',4',5,7-tetrahydroxyflavone).

The flavonoids of these species of clover have not been studied previously.

LITERATURE CITED

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