FLAVONOIDS OF Dorycnium intermedium

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The epigeal part of <u>Dorycnium</u> intermedium Ldb. collected in the environs of Tbilisi (Georgian SSR) was extracted with 80% methanol; after evaporation of the ethanol, the aqueous liquid was purified with ethyl ether, dried, and left in the cold. The yellow crystalline powder that separated out (1%) contained three flavonoids: A, B, and C. The main component was A, and substance C was present in very small amount. The separation of this material into individual flavonoids was achieved by column chromatography on polyamide sorbent and by preparative separation on a paper chromatogram.

Flavonoid A, $C_{21}H_{20}O_{12}$, mp 192-194°C, $[\alpha]_D^{20} - 157.9^\circ$ (c 0.5; ethanol). In UV light, $\lambda_{\max}^{C_2H_5OH}$ 255 nm. Acid hydrolysis with 2% H_2SO_4 gave the aglycone (yield 51%) with mp 351-358°C, which prove to be identical with myricetin [1]. L-Rhamnose was found in the carbohydrate fraction of the hydrolyzate.

On the basis of the above facts, and also IR and UV spectroscopy with complex-forming and ionizing additives, we characterized flavonoid A as 3,3',4',5,5',7-hexahydroxyflavone 3-O-rhamnoside, or myricitrin [2, 3].

Flavonoid B, $C_{27}H_{30}O_{14}$, mp 202-206°C, in UV light $\lambda_{max}^{C_2H_3OH}$ 265, 345 nm; was identified from its physicochemical constants as kaempferol 3,7-di(O- α -L-rhamnoside), or kaempferitrin [4].

Flavonoid C with mp 271-273°C, from its mobility on paper chromatography in various solvent systems and a mixed melting point, was identified as kaempferol.

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