FLAVONOIDS FROM Astragalus galegifolius

AND A. maximus

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To isolate the flavonoids, an 80% methanolic extract from 600 g of air-dry flowers of A. galegifolius was purified with chloroform after the alcohol had been distilled off. Drying the aqueous extract then gave 0.12 g of bright-yellow acicular crystals with the composition $C_{21}H_{20}O_{11}$, mp 196-198°C, UV spectrum: $\lambda_{\max}^{C_2H_5OH}$ 375, 270 nm. Acid hydrolysis formed 66% of the aglycone, kaempferol, and D-glucose; $[\alpha]_0^2 - 10^\circ$ (c 0.1; ethanol), $Kp_h[M]_0^2 - 25.5^\circ$. This value and calculations by Klyne's method showed the α configuration and the pyranose form of the sugar substituent [1]. The glycoside was readily cleaved by the enzyme of Aspergillus oryzae into kaempferol and D-glucose. By its IR spectrum, a mixed melting point, and bathochromy, the glucoside was identified as astragalin [2-4].

The combined flavonoids were obtained similarly from 700 g of the leaves and flowers of A. maximus. The ethyl acetate extracted yielded 0.15 g of yellow acicular crystals of a flavonoid with mp 192-194°C. Acid and enzymatic hydrolysis gave 79.6% of kaempferol and D-galactose. [α] $_D^{20}$ -8° (c 0.1; ethanol), Kph[M] $_D^{20}$ -20.4°. According to the molecular rotation, the galactose has the β configuration and the pyranose form [1]. In the UV spectrum ($\lambda_{max}^{C_2H_5OH}$ 359, 257 nm), bathochromic shifts showed the presence of the sugar substituent in position 3.

The results of a comparison of our experimental results with literature data indicate that the substance under investigation was kaempferol 3-O- β -D-galactoside, or trifolin [3, 4].

This is the first time that astragalin and trifolin have been isolated from the species mentioned.

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