PEUCEDANIN FROM THE ROOTS OF PEUCEDANUM LUXURIANS

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In a paper-chromatographic study of <u>P</u>. Luxurians Tamamsch, collected in Armenia between Idzhevan and Berd, we have established in it the presence of substances with the same R_f values as those in <u>P</u>. morisonii and similar species. By the method described previously [1], we have isolated from the roots of these plants a furocoumarin $C_{15}H_{16}O_4$ with mp 99-101° C (from carbon tetrachloride) which was found by mixed melting point and IR spectrum to be identical with peucedanin. Yield about 3%.

Peucadanin has previously been found in five species of Umbelliferae [1-4] belonging, like P. <u>luxurians</u>, to the section Peucedanum [5]. The presence of this substance in the section Peucedanum can be regarded as its chemical characteristic.

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FLAVONOIDS OF THE FRUIT OF SILYBUM MARIANUM

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Two flavonoid compounds have been isolated from the fruit of <u>Silybum marianum</u> (L) Gaertn., but their structure has not been definitively established at the present time [1, 2]. By means of two-dimensional paper chromatography, we have found about five compounds of a flavonoid nature in an ethanolic extract of the fruit. By using chromatography on a polyamide sorbent and cellulose, we have isolated two crystalline flavonoids, I and II, and one in the amorphous state, III.

Substance I, mp $234-237^{\circ}$ C, $[\alpha]_{D}^{22}+34.5^{\circ}$ (c 0.2; dimethylformamide; electric spectropolarimeter). UV spectrum:

 λ_{\max} 288, (325); $\lambda_{\max}^{CH_3COONa}$ 288, 327; $\lambda_{\max}^{H_3BO_8+CH_3COONa}$ 288, 328; $\lambda_{\max}^{C_2H_3ONa}$ 255, (288), 325; $\lambda_{\max}^{AlCl_3}$ (290), 315, 370 mµ; IR spectrum: 1647 (C=O, joined by a hydrogen bond to the 5-OH and the 3-OH); 1170, 1070, and 1035 (C-O-H); 3300-3440 (phenolic and alcoholic OH groups) cm⁻¹. In the solvent systems 1) ethyl acetate-benzene-acetic acid (73:24.5 :2.5) and 2) 25% acetic acid, I had Rf 0.36 and 0.57, respectively. Red orange color reaction with sodium borohydride, crimson with cyanidin, crimson with zinc + HCl, and yellow with concentrated H₂SO₄. The acetyl derivative has mp 118-120° C. In many of its properties, the substance obtained is similar to substance E₆ (silybin) previously obtained from S. marianum but differs from it in its optical activity and melting point [1, 2].

Substance II, mp 173-175°C, $[x]_D^{20} + 150^\circ$ (c 0.2; dimethylformamide). UV spectrum: λ_{max} 288, (325); $\lambda_{max}^{C_2H_6ONa}$ 245, 325; $\lambda_{max}^{CH_aCOONa}$ (288), 325; $\lambda_{max}^{H_3BO_4+CH_3COONa}$ 288 (325); $\lambda_{max}^{AlCI_5}$ (288), 315 m μ . IR spectrum: 1644 (OH in position 5 of a flavonoid); 1745 (C=O of an ester or a five-membered lactone); 3200-3300 (phenolic OH groups); 3460 (alcoholic OH groups) cm⁻¹. In system 1 R_f 0.63; in system 2, R_f 0.51. Red orange color reaction with sodium borohydride, yellow with cyanidin, yellow with zinc + HCl, and dark red with concentrated H₂SO₄. The acetate has mp 182-185°C. II is identical with substance E₅ previously isolated from S. marianum, but has a different melting point and optical activity.

Substance III. UV spectrum: $\lambda_{max} = 285$, (340); $\lambda_{max}^{C,H_5ONa} = (295)$, 325; $\lambda_{max}^{CH_5COONa} = 288$, (325); $\lambda_{max}^{H_5O_5 + CH_5COONa} = 288$, 340; $\lambda_{max}^{MCL_5} = (280)$, 315 mp . In system 1, R_f 0.79; in system 2, R_f, 0.07. Orange color reaction with sodium borohydride,

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