CHEMICAL STUDY OF THE EPIGEAL PARTS

OF THE PLANTS Ammothamnus lehmannii AND Cicer mogoltavicum

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From the roots of the plant Ammothamnus lehmannii Bge gathered close to the village of Kochka, Kenimekh region, Navoi province, Republic of Uzbekistan, we have already isolated two new flavonoids – ammothamnidin and lehmannin –, and from the epigeal part luteolin, cynaroside, quercetin, and isoquercetin, and from the roots of the plant Cicer mogoltavicum A. Kor gathered in the mountains of Kara-Tau, Dzhambul province, Republic of Kazakhstan, the isoflavonoids biochanin-A, formononetin, protensin, ononin, and trifolirhizin [1-3].

Continuing the investigations, from the chloroform fraction of an alcoholic extract of the epigeal part of A. lehmannii gathered during the flowering period in the environs of Lake Shukul', Dzhambul province, Republic of Kazakhastan, by column chromatography on silica gel chloroform—hexane elution we have isolated fractions containing two flavonoids. The rechromatography of these fractions on polyamide with elution by chloroform—hexane (3:1) led to substance (1) and with chloroform—hexane (8:2) and (9:1) to fractions containing a single flavonoid. These fractions were purified by rechromatography on a column of polyamide, and elution with chloroform—methanol (9:1) yielded substance (2).

Substance (1), with mp 112-114 °C (from acetone – hexane), $R_f 0.58$ (Silufol-254, acetone – hexane (1:1)) was identified as ammothamnidin by comparison with an authentic specimen [3].

Substance (2), with mp 102-104°C, λ_{max} (ethanol) 240, 286, 315 nm (log ε 4.05, 4.07, 3.82), was a flavonone. This fact was also shown by the signals of the H-2 and 2H-3 protons of the heterocyclic ring of the flavone nucleus, appearing in the PMR spectrum (Py-d₅) at 6.15 ppm (1H, dd, J = 11.0 and 5.0 Hz, H-2) and 2.87-3.26 ppm (5H, m, 2H-3, H-1", H-2"). On the basis of its UV, IR, and PMR spectra, and also by a direct comparison with an authentic sample, substance (2) was identified as lehmannin [4].

The chloroform fraction of an alcoholic extract of the epigeal part of *Cicer mogoltavicum* gathered in the neighborhood of Angren, Tashkent province, Republic of Uzbekistan, was chromatographed on a column of silica gel in a chloroform—hexane gradient system. Elution with chloroform—hexane (9:1) yielded fractions containing a single substance with R_f 0.26 (Silufol-254, chloroform—ethyl acetate (8:2)). After the solvent had been distilled off, the residue was recrystallized from benzene, and lustrous crystals were isolated with mp 180-181°C, M⁺ 284, λ_{max} (ethanol) 282, 287, and 311 nm. The UV spectrum was characteristic for pterocarpans. The IR spectrum revealed signals of an OH group (3480 cm⁻¹) and of aromatic C=C bonds (1506, 1043, 1020, 935 cm⁻¹). PMR spectrum (Py-d₅, δ , ppm): 3.42-3.92 (2H, m, H-6), 4.18 (1H, m, H-6a), 5.50 (1H, d, 6 Hz, H-11a), 5.85 (1H, d, 1.5 Hz, $-OCH_2O-$), 6.60 (1H, s, H-10), 6.80 (2H, br.s, H-4, H-7), 6.86 (1H, dd, 8.5 Hz and 2.0 Hz, H-2), 7.50 (1H, d, 8.5 Hz, H-1).

The spectral results and a comparison with literature information permitted the conclusion that the substance isolated belonged to the pterocarpan group and was, in fact, inermin -3-hydroxy-8,9-methylenedioxypterocarpan [5].

When the column was eluted with chloroform – hexane (8:2), yet another substance was isolated, with mp 260-262°C (from methanol), M⁺ 268. UV spectrum [λ_{max} (ethanol) 238, 250, 261, 304 nm] showed that the substance was an isoflavone. Its IR spectrum revealed signals for the absorption of hydroxy groups (3106 cm⁻¹), the carbonyl group of a γ -pyrone ring (1641 cm⁻¹), and aromatic C=C bonds (1573, 1518 cm⁻¹).

PMR spectrum (Py-d₅, δ, ppm): 3.57 (3H, s, -OCH₃), 6.99 (2H, d, 9.0 Hz, H-3, H-5), 7.72 (2H, d, 9.0 Hz, H-2, H-6), 7.07 (1H, br.s, H-8), 7.14 (1H, dd, 9.0 Hz, 2.0 Hz, H-6), 8.40 (1H, d, 9.0 Hz, H-5), 8.10 (1H, s, H-2).

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The spectral results and a comparison with literature information permitted the substance isolated to be identified as formononetin -7-hydroxy-4'-methoxyisoflavone [3].

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