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The comminuted leaves (2.0 kg) of A. turkestanica (Rgl.) Briq., family Labiatae collected in July 1970 in the flowering stage (environs of the village of Baisun, Surkhan-Dar'ya oblast) were extracted with 12 liters of methanol. The methanolic extract was concentrated, diluted with water, and treated with petroleum ether. The residual aqueous methanolic fraction, after additional removal of methanol in vacuum, was carefully extracted with ethyl acetate. The ethyl acetate extract, by chromatography on silica gel with elution by chloroform-methanol (9:1) yielded substance (I) (0.025% of the weight of the air-dry raw material) with R<sub>f</sub> 0.71 [in a thin fixed layer of silica gel, chloroform-methanol (4:1)]. Subsequent elution with chloroform-methanol (4:1) gave compound (II) (0.020%), with R<sub>f</sub> 0.50 (same conditions for chromatography).

Substance (I),  $C_{29}H_{44}O_8$  had mp 158-160°C (methanol),  $[\alpha]_D^{20} + 60.0 \pm 3.0$ ° (c 1.00; pyridine)  $\lambda_{max}^{C_2H_3OH}$  245 nm (Ig  $\epsilon$  4,05);  $\nu_{max}^{KBr}$  3450 cm<sup>-1</sup> (OH), 1752 cm<sup>-1</sup> ( $\gamma$ -lactone), 1660 cm<sup>-1</sup> (cyclohexenone). NMR spectrum in  $C_5D_5N$  at 100 MHz ( $\delta$  scale, internal standard HMDS) 0.94 ppm (3H at C-19, singlet), 1.11 ppm (3H at C-18, singlet), 1.24 (6H at C-27 and C-29, doublet, J = 6 Hz), 1.44 (3H at C-21, singlet), 6.15 ppm (1H at C-7). In the mass spectrum of (I) (MKh-1303, 160°C, energy of the ionizing electrons 40 eV), in addition to the weak molecular peak with m/e 520, peaks of ions with m/e 502, 484, 469, 466, 451, 448, 433, 430, 415, 363, 345, 327, 300, 183, 157 were found. These characteristics show that compound (I) is cyasterone [1].

Substance (II),  $C_{27}H_{44}O_7$ , was obtained with a double melting point  $152-154^{\circ}C$  and  $235-236^{\circ}C$  (from aqueous methanol) and also  $242-244^{\circ}C$  (from dry acetone)],  $[\alpha]_D^{20} + 54.5 \pm 2.5^{\circ}$  (c 0.52; methanol);  $\lambda_{\max}^{C_4H_5OH}$  243 nm (ig  $\epsilon$  4,09);  $\nu_{\max}^{KBr}$  3370-3450 (OH), 1660 cm<sup>-1</sup> (cyclohexenone). NMR spectrum (conditions analogous to those for cyasterone) (ppm): 0.93 (3H at C-19, singlet), 1.06 (3H at C-18, singlet), 1.24 (6H at C-26 and C-27, singlet), 1.44 (3H at C-21, singlet), 6.00 (1H at C-7). These facts, and also the mass spectrum of compound (II), containing the peak of the molecular ion with m/e 480 and ions with m/e 462, 444, 426, 408, 345, 344, 328, 327, 99 and 81 make it possible to consider that (II) is ecdysterone [2, 3]. The identity of (II) as ecdysterone was also confirmed by a direct comparison with an authentic sample kindly given to us by Ya. K. Yatsyuk [4]. Cyasterone and ecdysterone have been found previously in the leaves of Ajuga decumbens Thunb. ("kiranso") [5].

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