Substance (IV) — C₁₈H₁₆O₅, mp 180-183°C (ethanol). Yield 0.007%. UV spectrum, $\lambda_{\text{max}}^{\text{C}_2\text{H}_5\text{OH}}$ nm: 244 (ϵ 15,200), 276 (ϵ 13,600), 342 (ϵ 19,200).

IR spectrum, $v_{\text{max}}^{\text{KBr}}$ cm⁻¹: 3600-3400 (-OH), 1670 (>C=O), 1630, 1590 (aromatic nucleus). From its physicochemical constants and spectral characteristics, (IV) was identified as 5-hydroxy-4',7-dimethoxy-6-flavone [6].

This is the first time that arborescin and 5-hydroxy 4',7-dimethoxy-6-methylflavone have been isolated from Artemesia austriaca.

The extraction of the epigeal part of Artemesia austriaca with hot water, followed by extraction with chloroform and chromatography of the resin so obtained on a column of KSK silica gel (1:20) yielded 0.11% of arborescin and 0.05% of austricin, calculated on the airdry raw material.

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CARBONIZED CALCIUM HYDROMETASILICATE — A NEW EFFECTIVE ADSORBENT FOR THE SEPARATION OF NATURAL COMPOUNDS BY THIN-LAYER CHROMATOGRAPHY

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Carbonized calcium hydrometasilicate (KMK-20 — the numeral denotes the approximate percentage CO₂ content) is a surface-layer silicate material (the matrix being CaSiO₃ and the working surface CaCO₃; s ~ 200 m²/g, V_{pore} ~ 0.9 cm³/g, pH ~ 7.6, grain size 5-10 µm [1]). In its chromatographic activity, determined from the separation of a standard mixture of dyes (Fig. 1), KMK-20 is not inferior to adsorbents widely used in the TLC of natural compounds. Our aim was a study of the possibility of using KMK-20 for this purpose.

The results of a separation of a number of natural compounds on KSK silica gel, Silufol UV-254, and KMK-20 indicates its higher separating capacity and sensitivity. Thus, in the case of teucrins H1-H4, diterpene furolactones of the clerodane type (Fig. 2), the R_f values on KMK-20 were 13, 70, 40, and 60 (here and below, R_f 100), and on KSK silica gel they were 33, 68, 69, and 80. In the separation of the glycosides linarin and pectolinarin on KMK-20 it was possible to reveal an impurity component of the linarin which was not determined on Silufol UV-254 (Fig. 3). The aglycons of these glycosides — acacetin and pectolinarigenin — were separated on KMK-20 with R_f 35 and 83, and on Silufol UV-254 with R_f 51 and 56, respectively. A mixture of alkaloids was also readily separated on KMK-20: codeine, floripavine, fugapavine, and mecambridine — R_f 43, 55, 74, and 83, respectively (eluent: ethyl acetate—methanol—chloroform—water (7:2:1:1)). Methanolic extracts of the epigeal and radical parts of mullein were separated with the following R_f values: KMK-20 — 33, 36, 45, 51, 60, 64, 73*, 80*, 90 (epigeal part) and 51, 60, 65, 70, 75*, 81*, and 97 (radical part); Silufol UV-254 — 14, 29, 31, 34, 37, 40, 98, and 13, 29, 37, 42, and 97, respectively (* — spot detected only in TLC on KMK-20; eluent: ethyl acetate—methanol—chloroform—water (7:2:1:1)).

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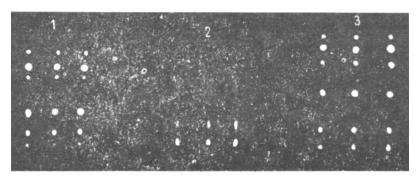


Fig. 1. Contact photograph from a chromatogram of the separation of a six-component standard mixture of Merck azo dyes: on KSK (1), Silufol UV-254 (2), and KMK-20 (3); eluent — benzene.

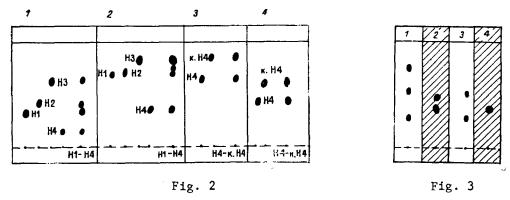


Fig. 2. Chromatograms of the separation of teucrins H1-H4, diterpene furolactones of the clerodane type, on KMK-20 (1) and KSK (2) with chloroform methanol (19:1) as the eluent, and of teucrin H4 and the ketone of teucrin H4 (K.H4) on KMK-20 (3) and KSK-4) with chloroform methanol (9:1) as the eluent.

Fig. 3. Chromatograms of the separation of linarin and pectolinarin on KMK-20 (1) and Silufol UV-254 (2) and of linarin on KMK-20 (3) and on Silufol UV-254 (4) with ethyl acetate-methanol-chloroform-water (7:2:1:1) as eluent.

Thus, the use of KMK-20 in the TLC of natural compounds can substantially facilitate the separation of substances with similar properties and provide a possibility for the reliable estimation of the purity of individual compounds. Its use in TLC does not require the performance of grinding and sieving operations; the absence of a hydroxylic coating on the surface eliminates the necessity for performing activation. The aragonite present in the carbonate layer possesses luminophoric properties and therefore eliminates the necessity for adding a luminophoric component.

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