

We have studied the chemical composition of the epigeal part of the *Pyrethrum roseum* (Adam) M. B. collected in the flowering period (July, 1978) in the region of Kislovodsk.

The comminuted raw material was extracted three times with 70% C<sub>2</sub>H<sub>5</sub>OH, the extracts were concentrated in vacuum to an aqueous residue and in this by two-dimensional chromatography using qualitative reactions [1], 16 substances of phenolic nature were detected.

Four substances were isolated from the products of the acid hydrolysis (5% H<sub>2</sub>SO<sub>4</sub>) of the dried aqueous residue by extraction with ether and chromatography on polyamide.

Substance (I), C<sub>15</sub>H<sub>10</sub>O<sub>6</sub>, mp 326–327°C. UV spectrum: λ<sub>max</sub> (C<sub>2</sub>H<sub>5</sub>OH) 350, 268, 255 nm. The IR spectra of substance (I) and of 3',4',5,7-tetrahydroxyflavone (luteolin) isolated previously from species of *Scabiosa* [2] were identical.

Substance (II), C<sub>16</sub>H<sub>12</sub>O<sub>6</sub>, mp 253–255°C. UV spectrum: λ<sub>max</sub> (C<sub>2</sub>H<sub>5</sub>OH) 345, 268, 255 nm. Isovanillic acid and phloroglucinol were found in the products of alkaline cleavage. The acetyl derivative, C<sub>22</sub>H<sub>18</sub>O<sub>9</sub>, had mp 195–197°C. On the basis of these facts, the substance was characterized as 3',5,7-trihydroxy-4'-methoxyflavone (diosmetin).

Substance (III), C<sub>15</sub>H<sub>10</sub>O<sub>7</sub>, mp 308–210°C [UV spectrum: λ<sub>max</sub> (C<sub>2</sub>H<sub>5</sub>OH) 370, 266 nm (melting point of the acetyl derivative 198–200°C)] was identified as quercetin.

Substance (IV), C<sub>15</sub>H<sub>10</sub>O<sub>6</sub>, mp 273–275°C [UV spectrum: λ<sub>max</sub> (C<sub>2</sub>H<sub>5</sub>OH) 367, 265 nm] was identified as kaempferol.

From an ethanolic extract by chromatography on polyamide we isolated a glycoside C<sub>28</sub>H<sub>32</sub>O<sub>15</sub>, mp 275–277°C. UV spectrum: λ<sub>max</sub> (C<sub>2</sub>H<sub>5</sub>OH) 342, 254, 265 nm. The UV spectra in the presence of diagnostic additives showed the presence of a free hydroxy group at C<sub>5</sub> and the absence of one at C<sub>7</sub>. From the products of quantitative acid hydrolysis we isolated diosmetin (yield of aglycone 48.7%) and a carbohydrate component which was identified as rutinose. On the basis of the results obtained and a comparative analysis with an authentic sample, the glycoside isolated was identified as diosmetin 7-rutinoside (diosmin). The aqueous extract of the freshly collected epigeal part of *P. roseum* followed by chromatography on a column containing polyamide yielded a substance C<sub>9</sub>H<sub>8</sub>O<sub>4</sub>, mp 195–198°C which was identified by UV spectroscopy, chromatography, and characteristic reactions as 3,4-dihydroxycinnamic (caffeic) acid [3].

The exhaustive extraction of a concentrated ethanolic extract with acidified diethyl ether yielded a compound C<sub>16</sub>H<sub>18</sub>O<sub>9</sub>, mp 204–207°C which was characterized by its physicochemical constants as 3-caffeoylquinic (chlorogenic) acid [3].

The study of the chemical composition of *Pyrethrum roseum* (Adam) M. B. is continuing.

## LITERATURE CITED

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