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POLYPHENOLIC COMPOUNDS OF POTATO FLOWERS

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In folk medicine, the flowers of the potato (*Solanum tuberosum* L.) have been used since antiquity as an antipyretic agent in diseases of the liver and kidneys, and in cases of constipation and tumors [1-3].

We have investigated the flowers of potatoes of the variety "Rozovyi rannii" ["early pink"] gathered in the Novo-Vodolazhskii Region of Khar'kov province.

The comminuted flowers were extracted with a fivefold volume of 80% ethanol with steeping for 24 h (three times) at room temperature. The combined extracts were evaporated in vacuum to small volume and centrifuged, and the resinous precipitate that had deposited was filtered off. After a day, an amorphous precipitate giving no reaction for flavonoids deposited from the extract obtained. Chromatography on paper in systems 1) butan-1-ol-acetic acid-water (4:1:2) and 2) 15% acetic acid showed the presence in the purified extract of not less than 9 substances of phenolic nature. In the products of the acid hydrolysis of the extract (5% ethanolic solution of hydrochloric acid, on the boiling water bath for 4 h), we found kaempferol, quercetin, myricetin, luteolin, and caffeic acid.

When the purified extract was stored in the refrigerator (5-6°C) a precipitate was formed of a substance in the form of pale yellow needles with the composition $C_{21}H_{20}O_{11}$, mp 256-258°C, $[\alpha]_D^{20} -100.0^\circ$. UV spectrum: λ_{max} 257, 268, 350 nm (substance (I)).

Chromatography on polyamide columns with elution by 30% ethanol gave a compound consisting of light yellow acicular crystals with the composition $C_{27}H_{30}O_{16}$, mp 191-193°C, $[\alpha]_D^{20} -32.0^\circ$. UV spectrum: λ_{max} 258, 264, 362 nm (substance (II)).

When using as eluent chloroform-ethanol (9:1), a substance was isolated in the form of yellow acicular crystals with the composition $C_{21}H_{20}O_{12}$, mp 229-231°C, $[\alpha]_D^{20} -33.0^\circ$. UV spectrum: λ_{max} 258, 264, 360 nm (substance (III)), and on elution with water heated to 40°C, we obtained a compound with the composition $C_{16}H_{18}O_9$, mp 202-204°C $[\alpha]_D^{20} -31.9^\circ$. UV spectrum: λ_{max} 240, 328 nm (substance (IV)).

On the basis of the results of a chemical investigation of the compounds obtained and of the products of their hydrolysis, and also of a comparison with authentic samples, substance (I) was identified as luteolin 7-O- β -D-glucopyranoside (cynaroside); (II) as quercetin 3-[O- α -L-rhamnopyranosyl-(1 \rightarrow 6)- β -D-glucopyranoside] (rutin); (III) as quercetin 3-O- β -D-glucopyranoside (isoquercitrin); and (IV) as 5-O-caffeyl-D-quinic (chlorogenic) acid.

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AN APIGENIN GLUCURONIDE FROM *Leucanthemum vulgare*

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We have previously reported the different flavonoid compositions of the ligulate and tubular flowers of *Leucanthemum vulgare* Lam. [1, 2].

Continuing a study of the chemical composition of the ligulate flowers, we have isolated another individual flavonoid — pale yellow crystals with the compositions $C_{21}H_{18}O_{11}$, mp 320–323°C (decomp.), $[\alpha]_D^{20} -92.0$ (c 0.6 pyridine–water (1:1)); molecular mass 446.4. UV spectrum $\lambda_{max}^{C_2H_5OH}$ 336, 270 nm; (+NaOAc) 336, 270 nm; (+AlCl₃) 380, 285 nm; (+AlCl₃ + HCl) 336, 270 nm; (+NaOMe) 400, 276 nm. The IR spectrum had an intense absorption band at 1725 cm⁻¹ (C=O group of an acid).

The glycoside was hydrolyzed with 10% H₂SO₄ for 5 h. This formed 68% of an aglycone with the composition C₁₅H₁₀O₅, mp 349–351°C. UV spectrum $\lambda_{max}^{C_2H_5OH}$ 336, 269 nm; (+NaOAc) 380, 274 nm; (+AlCl₃) 380, 275 nm; (+AlCl₃ + HCl) 336, 269 nm; (+NaOMe) 400, 275 nm. Phloroglucinol and p-hydroxybenzoic acid were found in the products of alkaline fusion. The aglycone was characterized as 4',5,7-trihydroxyflavone — apigenin [1–3].

The glycoside was hydrolyzed by the enzyme β-glucuronidase [4].

D-Glucuronic acid* was detected in the aqueous fractions of the acid and enzymatic hydrolysates by PC analysis.

A comparison of the physicochemical constants of the glycoside under investigation and an authentic sample of apigenin 7-O-β-D-glucuronide, together with literature information [5–7], showed their identity.

The results of the chemical study were confirmed by those of NMR spectroscopy in a comparative analysis of the aglycone, the glycoside itself, and an authentic sample.

Thus, the compound isolated has been characterized as apigenin 7-O-β-D-glucosiduronic acid.

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