FLAVONOIDS OF THE EPIGEAL PART OF Cicer mogoltavicum

A. Zh. Gafurov,^a A. Sattikulov,^a M. P. Yuldashev,^b and É. Kh. Botirov^c

UDC 547.972

Continuing an investigation of the flavonoids of plants of the genus *Cicer*, we have studied the phenolic components of *Cicer mogoltavicum* A. Kor. Earlier, inermin and formononetin had been isolated from the epigeal part of this plant [1], and formononetin, biochanin-A, pratensein, ononin, inermin, and trifolirhizin from the roots [2].

The ethyl acetate fraction of an alcoholic extract of the epigeal part of C. mogoltavicum gathered close to Angren, Tashkent oblast, was chromatographed on a column of silica gel in a chloroform—methanol gradient system.

This led to the isolation of biochanin-A and flavonoids (1-4).

Substance (1) — $C_{15}H_{10}O_6$ (H⁺ 286), mp 275-277°, λ_{max}^{EtOH} 265-370 nm, was identified as kaempferol (3,4',5,7-tetrahydroxyflavone) [3].

Substance (2) — $C_{15}H_{10}O_6$ (M⁺ 286), mp 328-331° (decomp.), λ_{max}^{EtOH} 260, 274, 356 nm, was identified as luteolin (3',4',5,7-tetrahydroxyflavone) [3, 4].

Substance (3) — $C_{21}H_{18}O_{12}$, mp 108-110°; λ_{max}^{EtOH} 266, 282, 350 nm, was identified as kaempferol 3-O-glucuronide.

Their IR spectra contained absorption bands of hydroxy groups (3250-3490 cm⁻¹), of the carbonyl of a γ -pyrone (1670-1665 cm⁻¹), and of aromatic C=C bonds. The mobilities of the substances in TLC and their IR spectra (bands of the C-O vibration of glycosides in the 1000-1100 cm⁻¹ region) showed the glycosidic nature of the compounds isolated.

The acid hydrolysis of compound (3) yielded kaempferol and D-glucuronic acid. Acetylation of substance (3) with acetic anhydride in pyridine gave a hexaacetate with the composition $C_{33}H_{30}O_{18}$ (M⁺ 714), mp 79-80°C [5].

Substance (4) — $C_{22}H_{22}O_{10}$, mp 218-220°, λ_{max}^{EtOH} 264-336 nm, which is characteristic for isoflavone derivatives, was identified as sissotrin (biochanin-A-7-O glucopyranoside).

Its PMR spectrum (in Py- d_5) showed the signals of protons at 3.73 (s, OCH₃), 4.00-4.60 (protons of the carbohydrate moiety), 5.68 (d, 6.0 Hz, H-1"), 6.60 (d, 2.0 Hz, H-6), 6.67 (d, 2.0 Hz, H-8), 6.98 (d, 9.0 Hz, H-3', H-5'), 7.66 (d, 9.0 Hz, H-2', H-6'), 8.05 (s, H-2), 13.43 ppm (br.s, 5-OH).

On acid hydrolysis, glycoside (4) gave biochanin-A (5,7-dihydroxy-4'-methoxyisoflavone, $C_{16}H_{12}O_5$, M^+ 284, mp 213-214°C) and D-glucose.

The acetylation of substance (4) with acetic anhydride in pyridine gave a pentaacetate with the composition $C_{32}H_{32}O_{15}$, mp 204-206°C, the mass spectrum of which contained, in addition to the peak of the molecular ion with m/z 972, intense peaks of fragmentary ions of acetylated tetraacetylglucose with m/z 331, 329, 271, and 169 [6].

This is the first time that compounds (1-4) have been isolated from Cicer mogoltavicum.

a) Samarkand Agricultural Institute, 703003, Samarkand, ul. M. Ulugbeka, 77. b) Institute of the Chemistry of Plant Substances, Academy of Sciences of the Republic of Uzbekistan, Tashkent, fax (3712) 40 64 75. c) Namangan Institute of Industrial Technology. Translated from Khimiya Prirodnykh Soedinenii, No. 4, pp. 642-643, July-August, 1997. Original article submitted February 10, 1997.

REFERENCES

- 1. A. Zh. Gafurov, A. Sattikulov, É. Kh. Batirov, and V. M. Malikov, Khim. Prir. Soedin., 321 (1995).
- 2. S. S. Yusopova, É. Kh. Batirov, F. Kiyamitdinova, and V. M. Malikov, Khim. Prir. Soedin., 639 (1986).
- 3. T. J. Mabry, C. R. Markham, and M. B. Thomas, The Systematic Identification of Flavonoids, Springer, New York (1970).
- 4. Sh. B. Abdullaev, A. Sattikulov, É. Kh. Batirov, Yu. V. Kurbatov, and V. M. Malikov, Khim. Prir. Soedin., 104 (1983).
- 5. J. B. Harborne and C. A. Williams, Flavones and Flavone Glycosides, in: The Flavonoids, Chapman and Hall, London (1975), p. 376.
- 6. A. L. Kazakov, Khim. Prir. Soedin., 274 (1973).