STUDIES OF COMPOUND OF DERIVATIVES OF HYDROXYCINNAMIC ACIDS AND COUMARINS IN CRANBERRY’S LEAVES BY HPLC

There was studied the chemical composition of coumarins and derivatives of hydroxycinnamic acid of cranberry’s leaves. In the cranberry’s leaves there were revealed 2 coumarins – umbelliferon, skopoletin; 4 phenolcarbonic and hydroxycinnamic acids – coffee, rosemary, p-coumarinic, chlorogenic and set their contents.

Key words: cranberry; leaves; coumarin; HPLC

INTRODUCTION
Decoction of cranberry’s leaves is used in the traditional and official medicine as a highly effective drug for the treatment of kidney and urinary tract. But this pharmaceutical form has several disadvantages: difficulty in preparation, lack of standardization, the inability to long-term storage, incomplete extraction of biologically active substances from medicinal plants. So it’s important to create a new standardized drug which will be based on biologically active substances of cranberry’s leaves [4].

The main group of substances, which has pharmacological effect in a cranberry’s leaves, are phenolic compounds. Earlier we reported on qualitative and quantitative determination of the chemical compound some classes of BAS in the leaves and extracts of leaves of cranberry: simple phenols hydroxycinnamic acid derivatives, the amount of flavonoids, the amount of phenolic compounds and organic acids [2, 3, 4]. Continuing research BAS cranberry’s leaves and their products, we noticed that the individual content of hydroxycinnamic acid derivatives and coumarins in cranberry’s leaves was not investigated. The aim was to investigate the composition of coumarins and derivatives of hydroxycinnamic acid cranberry’s leaves.

MATERIALS AND METHODS
The object of the study was the cranberry’s leaves, which was bought at the pharmacy (series 0715, a manufacturer by “Phyto Svit”).

Preparing samples for analysis of plant material. The exact weight of the sample material was crushed carefully, than was placed in a round bottom flask of 100 ml, it was extracted with 50 ml of 60 % methanol solution for 15 min in a boiling water bath with reverse refrigerator while stirring. Then the sample is treated with ultrasound for 10 min, filtered, quantitatively transferred to a volumetric flask of 100 ml, dilute to the mark with 60 % methanol [5].

Preliminary analysis of hydroxycinnamic acids was performed by TLC in the system anhydrous formic P – water P – acetate P (8 : 8 : 84) compared with authentic samples. There were identified chlorogenic and caffeic acids, which had a blue fluorescence under UV light.

To separate the amount of phenolic compounds into individual components it was used HPLC on chromatograph Agilent 1200 3 D LC System Technologies (USA), which is equipped with a flow vacuum degasser G1322A, four channel pump gradient of low pressure G1311A, autosampler (automatic injector) G1329A, thermostat of columns G 1316A, diodnomatrychnyy G1315S and refractometric G1362A detectors.

Method of determination. There was complied reversed-phase chromatography, used chromatography column SupelcoDiscovery C18 measure 250 × 4.6 mm with sorbent: modified silica gel by oktadetsyl groups, which has a diameter of 5 mkm. As mobile phase it was used: solvent A, which is 95 % of the mix of mobile phase – 0.005 N orthophosphoric acid and 5 % solvent B – acetonitrile. Mode of chromatography: maximum feed rate of the mobile phase is 0.7 ml/min, eluent working pressure is 10000-12000 kPa; column thermostat temperature is 25 °C; sample volume injected is 5-20 ml, chromatography time is 50 min. Mode of elution is gradient: 0 min 95 % solvent “A”, 5 % solvent “B”; 8 min 92 % solvent “A” 8 % solvent “B”; 15 min 90 % solvent “A”, 10 % solvent “B”; 40 min 60 % solvent “A”, 40 % solvent “B”; 41-42 min 25 % solvent “A”, 25 % solvent “B”; 43-50 min 95 % solvent “A”, 5 % solvent “B”; scan time is 0.6 seconds, the detection range is 190-400 nm.

To select the optimal analytical detection waves it has been studied ultraviolet spectra of existing standards of hydroxycinnamic acids. Based on these data detection of...
RESULTS AND DISCUSSION

In the cranberry’s leaves there was revealed phenolic compounds: coumarin – umbelliferon (0.02 %), skopoletin (0.05 %); phenolcarboxylic and hydroxycinnamic acids – coffee (0.03 %), rosemary (0.1 %), n-coumarinic (0.009 %), chlorogenic (0.16 %) (Fig.).

CONCLUSIONS

There was studied the qualitative and quantitative composition of hydroxycinnamic acids and coumarins of cranberry’s leaves. Umbelliferon content was 0.02 %, skopoletin – 0.05 %; coffee acid – 0.03 %; rosemary acid – 0.1 %; n-coumarinic – 0.009 %; chlorogenic acid – 0.16 %. The results indicate the perspectives of development new drugs which will be based on biologically active substances of cranberry’s leaves and need for better studying of the phenolic compounds of this type of material. The data can be used to standardize the cranberry’s leaves.

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ИССЛЕДОВАНИЕ СОСТАВА ПРОИЗВОДНЫХ ГИДРОКСИКОРИЧНОЙ КИСЛОТЫ И КУМАРИНОВ В ЛИСТЬЯХ БРУСНИКИ МЕТОДОМ ВЭЖХ

Исследован химический состав кумаринов и производных гидроксикоричной кислоты листьев брусники обыкновенной. В листьях брусники обнаружены 2 кумарина – умбеллиферон, скополетин; 4 фенолкарбоновые и гидроксикоричные кислоты – кофейная, розмариновая, п-кумаровая, хлорогеновая и установлено их содержание.

Ключевые слова: брусника; листья; кумарины; ВЭЖХ

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ДОСЛІДЖЕННЯ СКЛАДУ ПОХІДНИХ ГІДРОКСИКОРИЧНОЇ КИСЛОТИ ТА КУМАРИНІВ У ЛИСТІ БРУСНИЦІ МЕТОДОМ ВЕРХ

Досліджено хімічний склад кумаринів та похідних гідроксикоричної кислоти листів брусниці звичайної. В листі брусниці виявлені 2 кумарини – умбеллиферон, скополетин; 4 фенолкарбонові та гідроксикоричні кислоти – канова, розмаринова, п-кумарова, хлорогенова та встановлено їх вміст.

Ключові слова: брусниця; листя; кумарини; ВЕРХ