

UDC 615.322 577.127

DOI: 10.15587/2519-4852.2021.242761

## CHROMATO-MASS-SPECTROMETRIC RESEARCH IN *SALVIA GRANDIFLORA* L., *SALVIA PRATENSIS* L. AND *SALVIA VERTICILLATA* L. ABOVEGROUND ORGANS

Mykhailo Myha, Oleh Koshovyi, Eugene Karpun, Alla Kovalyova, Olga Mala, Volodymyr Parchenko, Olexandr Panasenکو, Vera Bunyatyan, Sergiy Kovalenko

The genus *Salvia* L. has more than 900 species distributed throughout the globe. 21 species are growing in Ukraine. All species of this genus have essential oils. *Salvia officinalis* and *Salvia sclarea* have been used in the culture and are widely used in medical practice. The chemical composition of other species of sage and the possibility of their use in pharmaceutical and medical practice are almost not studied. Taking into account the results of chemotaxonomic studies of species of the flora genus of Ukraine, their prevalence and prospects for introduction into the culture, for further studies were selected raw materials of *S. grandiflora*, *S. pratensis* and *S. verticillata*.

**The aim.** The aim of the study was to conduct a chromat-mass spectrometric study of the aboveground organs of *S. grandiflora* L., *S. pratensis* L. and *S. verticillata* L. to establish the prospects for the use of raw materials of these species in medical and pharmaceutical practice.

**Materials and methods.** The objects of the study were leaves of *S. officinalis*, leaves, stems and flowers of *S. grandiflora*, *S. pratensis* and *S. verticillata*, which were harvested on the basis of the botanical garden of Ivan Franko National University of Lviv. The research of volatile substances in the objects of the research was carried out by the method of GC-MS on the basis of the Department of Natural Sciences for Foreign Students and Toxicological Chemistry of Zaporizhia State Medical University.

**Results.** As a result of the study, 243 substances were found in the objects of the study, of which 149 were identified. 77 substances were found in the leaves of *S. officinalis*, 80, 26 and 63 substances in the leaves, stems and flowers of *S. grandiflora*, respectively, in the leaves, stems and flowers of *S. pratensis* – 28, 30 and 48 substances, respectively, in leaves, stems and flowers of *S. verticillata* – 39, 22 and 39 substances, respectively. Dominant compounds among substances of terpenoid nature are: cyclofenchene, camphene, 1,8-cineole,  $\alpha$ -thujone,  $\beta$ -thujone, camphor borneol, caryophyllene, humulene, viridiflorol, sabinene, pyranone,  $\beta$ -pinene, phytol, kolavenol,  $\beta$ -copaen, loliolide, pseudolimonene and spatulenol. Among the dominant substances, 8 were detected for the first time in these species: cyclofenchene, viridiflorol, sabinene, pyranone, phytol, kolavenol, loliolide and pseudolimonene.

**Conclusions.** The leaves of *S. officinalis*, leaves, stems and flowers of *S. grandiflora*, *S. pratensis* and *S. verticillata* of the flora of Ukraine were studied by chromat-mass spectrometric method. As a result of the study, 243 substances were identified, of which 149 were identified. Promising raw materials containing terpene compounds for *S. grandiflora* there are leaves, and for *S. pratensis* and *S. verticillata* – flowers, so they are promising agents for introduction into pharmaceutical practice

**Keywords:** *Salvia*, leaves, flowers, stems, terpenes, chromat-mass spectrometry

### How to cite:

Myha, M., Koshovyi, O., Karpun, E., Kovalyova, A., Mala, O., Parchenko, V., Panasenکو, O., Bunyatyan, V., Kovalenko, S. (2021). Chromato-mass-spectrometric research in *Salvia grandiflora* L., *Salvia pratensis* L. and *Salvia verticillata* L. aboveground organs. ScienceRise: Pharmaceutical Science, 5 (33), 32–40. doi: <http://doi.org/10.15587/2519-4852.2021.242761>

© The Author(s) 2021

This is an open access article under the Creative Commons CC BY license hydrate

### 1. Introduction

The genus *Salvia* L. has more than 900 species distributed throughout the globe, of which 21 species are represented in the Ukrainian flora. All species of this genus have essential oils. Sage and sage entered the culture [1, 2]. The chemical composition of other species of sage and the possibility of their use in pharmaceutical and medical practice are almost not studied.

Biologically active substances of the genus *Salvia* L. are characterized by a wide range of pharmaco-

logical activity, in particular: antioxidant activity [3, 4], improvement of memory and cognitive functions [5, 6], hypoglycemic and insulin-like activity [6–8], ability to inhibit angiogenesis and proliferation [9, 10], lowering cholesterol [10, 11], reducing symptoms during menopause and menopause [12, 13], antibacterial [14, 15], antidiarrheal [16, 17] and antispasmodic activity [18–20]. This determines the relevance of the study of the genus *Salvia* for the development of new drugs based on them.

Based on the literature data and the results of our own research, a chemotaxonomic study of 17 species of the genus Sage (*Salvia* L.): *S. officinalis*, *S. sclarea*, *S. grandiflora*, *S. scabiosifolia*, *S. glutinosa*, *S. aethiopsis*, *S. pratensis*, *S. stepposa*, *S. Sibthorpii*, *S. illuminata*, *S. nemorosa*, *S. pendula*, *S. sylvestris*, *S. nutans*, *S. austriaca*, *S. verticillata* and *S. cernua* to identify promising species and establish the possibility of creating new drugs based on them [21]. In the list of these species, 60 compounds of phenolic nature and 185 compounds of terpenoid nature were found [21, 22]. The chemical profile of the genus is characterized by phenolic compounds: caffeic and rosmarinic acids [3], flavonoids: cynaroside, kosmosiin, hispidulin and cyrsimaritin [23–25] and terpenoids:  $\alpha$ - and  $\beta$ -pinene, derivatives of camphor (camphene, camphor and borneol) [26], p-cymene, 1,8-cineole and limonene [27, 28]. In addition, the amino acid and monosaccharide composition of sage leaves was studied [29]. Taking into account the results of chemotaxonomic

studies and the prospects of their introduction into the culture, the raw materials of *S. grandiflora*, *S. pratensis* and *S. verticillata* were selected for further studies.

**The aim of the research** was to conduct a chromat-mass spectrometric study of the aboveground organs of *S. grandiflora* L., *S. pratensis* L. and *S. verticillata* L. to establish the prospects for the use of raw materials of these species in medical and pharmaceutical practice.

## 2. Planning (methodology) of the research

Taking into account the results of chemotaxonomic studies of sage flora of Ukraine [21], which showed the prospects of using *S. grandiflora*, *S. pratensis* and *S. verticillata* in medical and pharmaceutical practice, it is planned to conduct chromat-mass spectrometric study of aboveground organs of these species, to identify promising species raw materials and the possibility of its further use in pharmacy (Fig. 1).

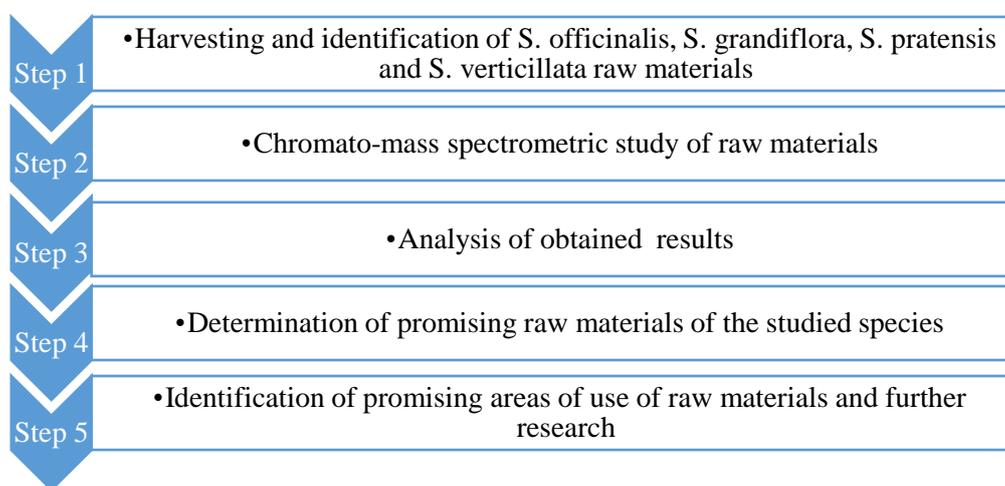


Fig. 1. Scheme of research of *S. grandiflora*, *S. pratensis* and *S. verticillata* raw materials

## 3. Materials and methods

The objects of the study were leaves of *S. officinalis*, leaves, stems and flowers of *S. grandiflora*, *S. pratensis* and *S. verticillata*, which were harvested on the basis of the Botanical Garden of Ivan Franko National University of Lviv (Lviv, 44 Cheremshyny Street, 79014), under the guidance of a senior researcher, PhD Skibitska Maria Ivanovna. The studied examples of *Salvia* genus were identified by PhD Olga Mala from the National University of Pharmacy using a special botanical catalogue [30]. The identified samples corresponded to the moisture content of SPhU [31, 32]. The voucher specimens are stored at the Department of Pharmacognosy, The National University of Pharmacy, Kharkiv, Ukraine (No. 1015–1029).

Studies of volatile substances in the objects of the research were carried out by GC-MS method [33] on the basis of the Department of Natural Sciences for Foreign Students and Toxicological Chemistry of Zaporizhia State Medical University according to the method: a sample of 2.0 (exact sample) of plant material was placed in a test tube (volume of which 20 ml), poured 10 ml of methanol (Sigma-Aldrich) and left for one day. The extract was filtered using a syringe filter, the resulting filtrate was chromatographed [34, 35].

Agilent 7890B gas chromatograph with 5977B mass spectrometric detector and Gerstel CIS 4 chilled injection system. DB-5ms chromatographic column 30 m x 250  $\mu\text{m}$  x 0.25  $\mu\text{m}$  long. The speed of the carrier gas (helium) is 1.3 ml/min. The injection volume is 0.5  $\mu\text{l}$ . Pulse injection with flow division in the ratio 1:5 was used. The temperature of the sampling unit is 200°C  $\rightarrow$  12°C/s  $\rightarrow$  265°C. Thermostat temperature: programmable, 70°C (1 min delay)  $\rightarrow$  10°C/min  $\rightarrow$  270°C (4 min delay). The total chromatography time is 25 minutes. GC/MS interface temperature – 275°C; ion sources – 230°C; quadrupole mass analyzer – 150°C. Type of ionization: electron impact at an electron energy of 70 eV. Scanned mass range: 30–700 m/z. The NIST14 mass spectrum library was used to identify the components. Quantitative determination of the content of substances in the raw material was performed in comparison with a standard sample of menthol [36, 37].

## 4. Results of the research

Determination of the qualitative composition and quantitative content of volatile substances in the study objects was performed by GC-MS method using an Agilent 7890B chromatograph. The results of the study of volatile substances in the aboveground organs

of the studied species of the genus *Salvia* are given in Table 1. In Fig. 2–5 are shown typical chromatograms

obtained in the determination of volatile compounds of the objects of the study.

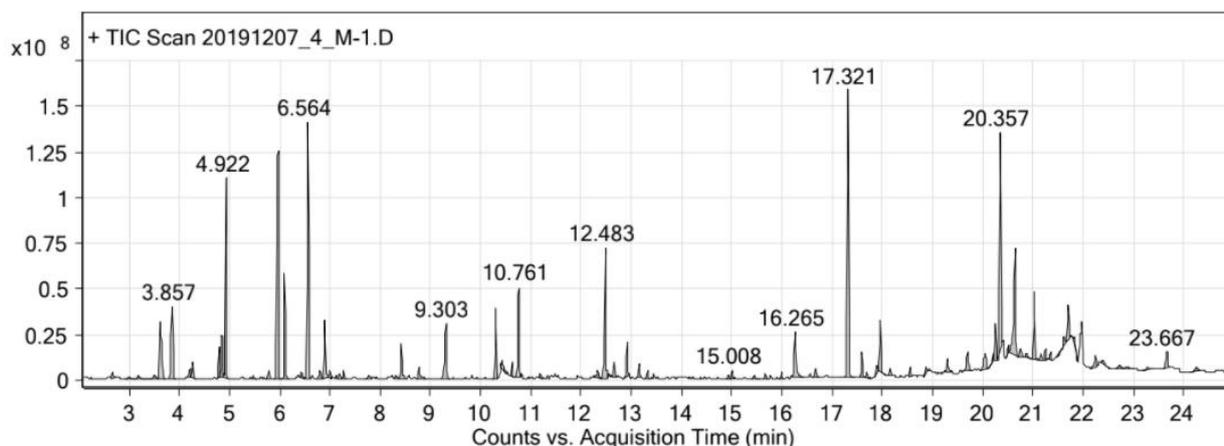


Fig. 2. Typical chromatogram of volatile compounds of *S. officinalis* leaves

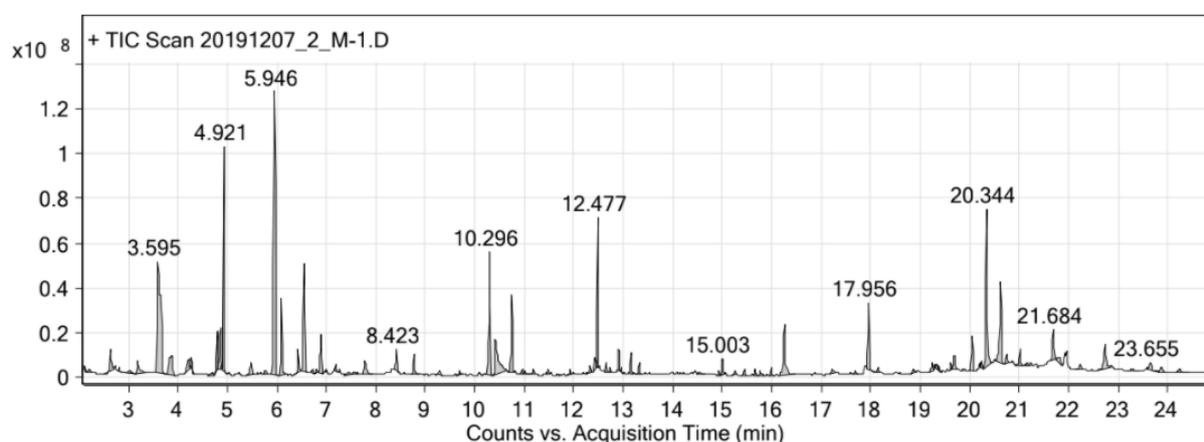


Fig. 3. Typical chromatogram of volatile compounds of *S. grandiflora* leaves

Table 1  
Content of volatile compounds in *S. officinalis* leaves; leaves, stems and flowers of *S. grandiflora*, *S. pratensis* and *S. verticillata*

1	Name of compound	Quantitative content, mg/100 g									
		<i>Salvia officinalis</i>	<i>Salvia grandiflora</i>			<i>Salvia pratensis</i>			<i>Salvia verticillata</i>		
		Leaves	Leaves	Stem	Flow-ers	Leaves	Stem	Flow-ers	Leav-es	Stem	Flow-ers
2		3	4	5	6	7	8	9	10	11	12
1	Diepoxybutane	0	1.47	0	0	0	0	0	0	0	0
2	Hydroxyacetone-	0	0	0	0	0	1.16	0.96	0	0	0
3	Methyl acetate	0	0	0	1	0	0	0	0	1.06	0
4	Ethylene glycol, monoacetate	0	0	0	0	1.69	0	0	0	0	0
5	Glycerin	0	2.24	0	0	3.47	0	3.13	0	0	0
6	Glyceraldehyde	0	0	2.34	6.48	0	4.56	0	0	10.59	2.43
7	Cyclopropane, 1,1-dimethyl-2-	0	9.23	0	0	0	0	0	0	0	0
8	Cyclohexane, 1,3-dimethyl-2-methylene-, cis-	3.78	0	0	1.29	0	0	0	0	0	0
9	Methyl 3-nitropropionate	2.05	3.79	2.24	2.98	0	2.55	2.77	0	2.76	1.90
10	Dihydroxyacetone	4.54	12.01	6.23	12.41	0	8.24	4.30	0	12.00	4.46
11	Tricyclene	3.99	0	0	0	0	0	0	0	0	0
12	$\alpha$ -pinene	0	0	0	0	0	0	0	1.84	0	0
13	Cyclofenchene	61.72	140.26	6.37	38.38	0	0	0	0	0	1.88
14	Camphene	79.72	25.77	0	3.21	0	0	0	0	0	0

Continuation Table 1

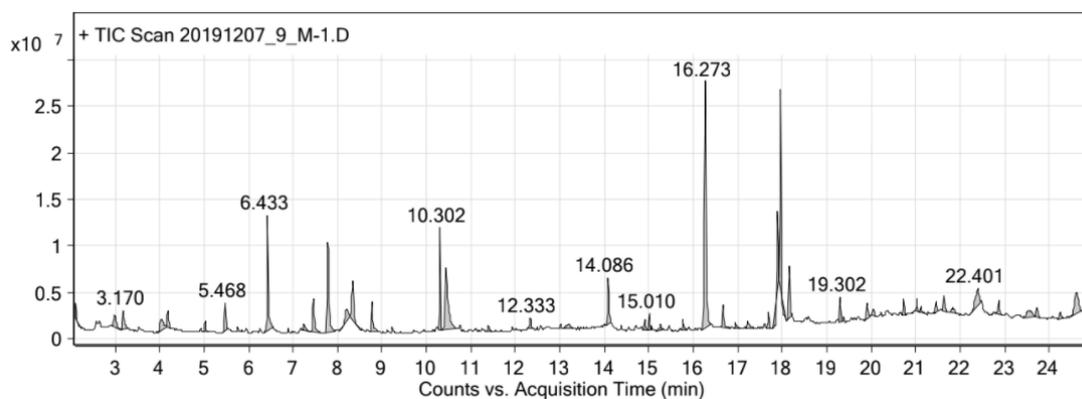
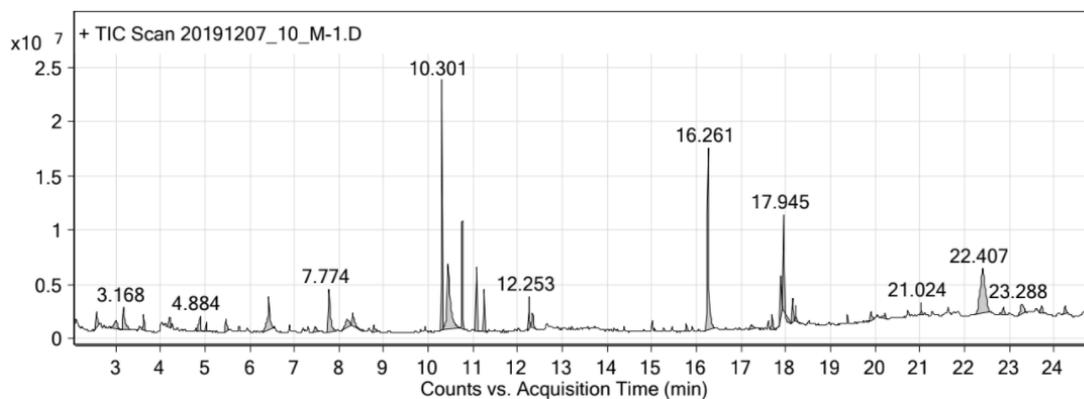
1	2	3	4	5	6	7	8	9	10	11	12
15	2,4-dihydroxy-2,5-dimethyl-3(2H)-furan-3-one	0	0	0	0	1.46	0	2.12	0	0	1.52
16	Sabinene	6.37	0	6.78	0	0	0	0	0	0	0
17	Nopinene ( $\beta$ -pinene)	0	8.75	0	10.98	0	0	0	0	0	0
18	$\beta$ -myrcene	7.31	3.94	0	1.17	0	0	0	0	0	0
19	4-carene	0	1.97	0	0	0	0	0	0	0	0.94
20	O-cymene	14.68	0	0	3.68	0	0	0	0	0	0
21	M-cymene	0	16.86	0	0	0	0	0	0	0	0
22	D-limonene	26.78	21.88	0	4.52	0	0	0	0	0	0
23	Pseudolimonene	0	0	0	0	0	0	0	0	0	2.24
24	1,8-cineole	116.64	104.05	6.45	35.67	0	0	0	0	0	0
25	Levulinic acid	0	0	0	0	0	0	1.49	0	0	0
26	Thymine	0	0	0	2.32	0	0	0	0	0	0
27	D-Alanine	2.04	7.31	0	0	0	0	4.01	0	0.92	1.47
28	Undecan	0	0	0	0	0	0	0	0.78	0	0
28	Linalool	5.42	3.21	0	0	0	0	0	0	0	0
30	$\alpha$ -thujone	171.44	186.36	36.39	40.90	0	0	0	1.69	0	0
31	$\beta$ -thujone	57.93	42.26	6.09	9.67	0	0	0	0	0	0
32	Pyranone	3.04	9.85	6.84	5.20	7.12	15.54	14.41	0	0	6.91
33	Erythritol	0	0	0	0	0	0	0	0	17.38	0
34	Camphor	189.82	76.31	2.68	0	0	0	0	0.61	0	0
35	3-pinanone	0	2.01	0	0	0	0	0	0	0	0
36	5-Caranol	5.66	0	0	0	0	0	0	0	0	0
38	Borneol	40.24	23.83	0	1.79	0	0	0	0	0	0
39	Terpinen-4-ol	3.58	0.00	0	0.89	0	0	0	0	0	0
40	$\alpha$ -terpineol	3.04	1.96	0	1.12	0	0	0	0	0	0
41	1,2,2,3-tetramethylcyclopent-3-enol	5.06	0	0	0	0	0	0	0	0	0
42	Benzofuran	0	0	0	1.87	2.92	0	5.75	0.67	0	1.07
43	Glycerol monoacetate	2.92	11.78	4.80	8.44	7.58	9.67	15.32	5.12	7.00	5.12
44	Phenylacetic acid	0	0	0	0	0	0	0	0.86	0	0
45	2-butoxyethanol	0	0	0	0	0	1.93	3.30	0	1.17	2.55
46	6-oxoheptanoic acid	0	0	0	0	2.87	0	0	0	1.84	0
47	3-hydroxy-2,3-dihydromaltol	0	0	0	3.25	0	1.59	7.23	0	0	2.69
48	Bornylacetate	23.67	11.28	0	0	0	0	0	0	0	0
49	Acetophenone, 2-hydroxy-5-methyl	9.36	10.61	0	0	6.13	0	5.28	1.00	0	1.25
50	4-hydroxy-3-methylacetophenone	0	0	0	1.87	0	1.06	0	0	0	0
51	Cytramalic acid	0	0	0	0.93	0	0	0	0	0	0
52	6-vinyl-3,3-dimethyl-6-hydroxy-5-formylmethyl-bicyclo [3.2.0]-heptan-2-one	38.02	1.77	0	0	0	0	0	0	0	0
53	Chavibetol	0	0	0	0.93	0	0	0	0	0	0
54	$\alpha$ -copaen	0	2.04	0	0	0	0	0	0	0	0
55	Caryophyllene	43.43	65.96	2.39	13.79	0	0	12.09	14.63	0	25.51
56	Salicylaldehyde hydrazone	0	0	0	0	0	0	0	8.07	0	0
57	6-methyl salicylaldehyde	0	0	0	0	13.27	0	0	0	0	0
58	Acet fluoroglucin	7.01	0	0	0	0	0	0	0	0	0
59	1,5,9,9-tetramethyl-1,4,7, cycloundecatriene	0	42.61	0	4.33	0	0	0	6.29	0	11.40
60	Humulene	62.58	0	0	0	0	0	0	0	0	0
61	$\gamma$ -Muroloene	0	2.31	0	0	0	0	0	0	0	0
62	Ethanone, 1-(3-hydroxy-4-methoxyphenyl)	0	1.58	0	0	0	0	0	0	0	0
63	(-)- $\beta$ -copaen	0	0	0	0	0	0	0	33.42	0	0
64	Methyl arachidonate	0	2.90	0	0	0	0	0	0	0	0
65	Ledene	2.79	0	0	0	0	0	0	0	0	0
67	9,10-dihydroxystearate	0	0	0	0	0	0	0.87	0	0	0
68	$\Delta$ -cardinene	0	1.58	0	0	0	0	0	0	0	0
70	Dihydroactinolide	0	0	0	0	0	0	0	1.54	0	0
71	Acetoveratron	0	2.06	0	0	0	0	0	0	0	0
72	Spatulenol	0	0	0	0	0	0	0	1.65	0	2.61
73	3,5-dihydroxy-6-(hydroxymethyl)oxan-2-one	0	0	0	0	0	2.02	0	0	0	0
74	Caryophyllene oxide	4.57	3.17	0	0	0	0	0	1.59	0	1.64
75	1-heptatriacotanol	0	0	0	0	0	0	1.33	0	0	0
76	3-hydroxy-5,6-epoxy- $\beta$ -ionone	0	0	0	0	0	0	0	0.60	0	0
77	Globulol/Ledol/Viridiflorol	87.35	4.44	7.57	23.47	0	0	0	0	0	0

Continuation Table 1

1	2	3	4	5	6	7	8	9	10	11	12
78	$\alpha$ -gurjunene	0	60.80	0	0	0	0	0	0	0	0
79	Humulene	2.34	0	0	0	0	0	0	0	0	0
80	(1R,3E,7E,11R)-1,5,5,8-tetramethyl-12-oxabicyclo [9.1.0]dodeca-3,7-diene	9.13	4.06	0	0	0	0	0	0	0	0
81	Tabanone	0	1.75	0	0	0	0	0	0	0	0
82	Diepicedren-1-oxide	17.35	9.67	0	1.44	0	0	0	0	0	0
83	11,11-dimethyl-4,8-dimethylenebicyclo-[7.2.0]undecan-3-ol	0	1.43	0	0	0	0	0	0	0	0
84	trans-longipinocarveol	4.34	7.99	0	0	0	0	0	0	0	0
85	Isoaromadendrene epoxide	6.22	0	0	1.70	0	0	0	0	0	0
88	(1R,7S,E)-7-isopropyl-4,10-dimethylenecyclodec-5-enol	0	5.06	0	0	0	0	0	2.91	0	0
89	Loliolide	0	0	0	0	0	0	0	2.66	0	0
90	Phytol acetate	3.76	11.28	0	2.28	6.04	1.69	1.54	6.63	0	0
91	Phytone	0	0	0	0	0	0	0	0.68	0	0
92	Phthalic acid	0	0	0	0	0	0	0	1.99	0	0
93	4,4,8-trimethyltricyclo[6.3.1.0(1,5)]dodecane-2,9-diol	3.34	2.30	0	0	0	0	0	0	0	0
94	Hexadecanoic acid methyl ester	0	0	0	0	0	0	0	0.70	2.12	0
95	Biformen	3.59	3.33	0	0	0	0	0	0	0	0
96	Palmitoleic acid	35.04	37.86	10.30	13.30	62.67	22.65	40.73	41.51	7.58	21.14
97	trans-2-palmitoleic acid	5.34	0	0	0	0	0	3.13	0	0	0
98	13-epimanol	169.76	0	0	0	0	0	0	0	0	0
99	8,11-octadecadienoic acid methyl ester	0	0	0	0	0	0	0	0	1.14	0
100	6,9,12-octadecatrienic acid methyl ester	0	0	0	0	0	0	0	0	1.18	0
101	Glyceryl linoleate	0	0	0	0	0	0	0	0.83	0	0
102	Phytol	3.05	1.91	0	0	8.12	0	1.51	4.75	0	1.27
103	9,12-octadecadienoic acid (Z,Z)	3.53	0	0	2.42	0	1.66	3.74	0	1.59	3.40
104	9,12,15-octadecatrienic acid, (Z,Z,Z)	29.38	32.34	0	6.79	96.30	12.19	16.39	26.27	3.33	7.24
105	Ascorbyl stearate	0	0	0	1.17	0	1.41	0	0	0	0
106	Stearic acid	3.99	2.35	0	0	4.16	0.00	6.36	3.11	0	2.00
107	Dibutyl sebacate	0	0	0	0	0	1.13	0	0	1.04	1.11
108	Oleic acid	5.09	0	4.63	0	0	0	0	0	0	0
109	Kolavenol	7.84	1.55	0	0	0	0	2.77	0	0	0
110	1,4-dimethyl-8-isopropylidetricyclo[5.3.0.0(4.10)]decane	12.36	1.88	0	0	0	0	0	0	0	0
111	Dehydroabietane	0	1.64	0	0	0	0	0	0	0	0
112	Podocarpa,8,11,13-tetraene-3one,14-isopropyl-1,13-dimethoxy	13.53	8.00	0	5.27	0	0	0	0	0	0
113	12-O-methylcarnosol	14.26	17.98	3.96	5.67	0	0	0	0	0	0
114	11-decyldocasane	0	0	0	0	0	0	1.61	0	0	0
115	Isocarnosol	19.62	0	0	0	0	0	0	0	0	0
116	Galantamine	181.55	89.06	16.26	30.93	0	0	0	0	0	0
117	Equiline derivat	0	0	0	0	1.80	1.20	0	0	0	0
118	1,2,3,4-tetrahydro-1,4-ethanoanthracene,9,10-dimethoxy-	3.56	0	0	0	0	0	0	0	0	0
119	1- (4-methyl [1,1 ' : 4', 1 " ] terphenyl-4"yl) ethanone	0	52.18	0	0	0	0	0	0	0	0
120	Atrovenetrin	0	3.69	0	0	0	0	0	0	0	0
121	Squalene	0	0	0	2.91	0	0	0	0	0	0
122	Octadecane, 3-ethyl-5-(2-ethylbutyl)-	0	0	0	0	0	0	3.81	0.59	0	0
126	1,2-dipalmitin	0	0	0	0	1.52	0	0	0	0	0
128	Ethyl isocholate	0	0	0	0	1.88	0	0	3.71	0	0
129	Dibenz[a,c]cyclohexane, 2,4,7-trimethoxy-	4.22	0	0	0	0	0	0	0	0	0
130	Pizyferol	19.53	15.38	2.70	9.70	0	0	0	0	0	0
131	Dibenz[d, f]cycloheptanone, 2,3,9-trimethoxy-	7.56	0	0	0	0	0	0	0	0	0
132	$\gamma$ -sitosterol	2.20	0	11.92	14.81	36.78	18.51	5.56	0	13.31	18.68
133	Germanicol	0	0	0	0	0	1.45	0	0	0	0
134	12-O-Methylcarnosol	3.16	20.31	0	2.57	0	0	0	0	0	0
135	Dibenz[a, c]cycloheptane, 2,3,7-trimethoxy-	2.55	0	0	0	0	0	0	0	0	0

Continuation Table 1

1	2	3	4	5	6	7	8	9	10	11	12
136	6a, 14a-methanopicene, perhydro-1,2,4a, 6b, 9,9,12a-hexamethyl-10-hydroxy-	0	0	0	0	0	20.02	0	0	5.37	0
137	Salvigenin	0	0	5.85	0	0	0	0	0	0	2.87
138	$\alpha$ -amyrin	0	0	0	0	0	7.95	0	0	0	0
140	$\alpha$ -tocopherol	0	0	9.57	0	11.39	0	0	0	0	0
141	9-octylhexacosane	0	0	0	0	0	0	1.58	0	0	0
142	Supraene	4.26	3.09	0	0	2.65	0	0	0	0	1.11
143	2,6,6,9,2',6',6',9'-octamethyl-[8,8']bi[tricyclo[5.4.0.0(2,9)]undecyl]	0	0	0	0	0	9.97	0	0	0	0
144	Betulinaldehyde	0	0	0	0	0	7.13	0	0	0	0
148	3-methoxyergost-8(14)-ene	0	0	16.77	0	0	0	0	0	0	0
149	Ursolic aldehyde	0	0	0	16.45	0	0	0	0	0	0
Unidentified substances	pcs.	15	19	4	21	8	8	21	9	5	12
	mg/100 g	152.04	86.08	24.86	48.09	29.17	8.73	56.39	22.34	14.29	34.79

Fig. 4. Typical chromatogram of volatile compounds of *S. pratensis* flowersFig. 5. Typical chromatogram of volatile compounds of *S. verticillata* flowers

As a result of the study, 243 substances were found in the objects of the study, of which 149 were identified.

### 5. Discussion of the results

77 substances were found in the leaves of *S. officinalis*, the dominant ones were cyclofenchene, camphene, 1,8-cineole,  $\alpha$ -thujone,  $\beta$ -thujone, camphor borneol, caryophyllene, humulene, viridiflorol and 13-epimanool.

In the leaves of *S. grandiflora* 80 substances were found, the dominant ones were cyclofenchene, 1,8-cineole,  $\alpha$ -thujone,  $\beta$ -thujone, camphor borneol, caryophyllene and viridiflorol. 26 substances were found in the stem of *S. grandiflora*, the dominant ones were cyclofenchene, sabinene, 1,8-cineole,  $\alpha$ - and  $\beta$ -thujone and pyranone. 63 substances were found in *S. grandiflora* flowers, the

dominant ones being cyclofenchene,  $\beta$ -pinene, 1,8-cineole,  $\alpha$ -thujone, caryophyllene and viridiflorol.

In the leaves of *S. pratensis* 28 substances were found, pyranone and phytol were dominant. 30 substances were found in the stem of *S. pratensis*, pyranone and  $\alpha$ -amyrin were dominant. 48 substances were found in *S. pratensis* flowers, pyranone, caryophyllene, phytol and kolavenol were dominant.

In the leaves of *S. verticillata* 39 substances were found, the dominant ones were caryophyllene,  $\beta$ -copaen, loliolide and phytol. 22 substances were found in the stem of *S. verticillata*. In *S. verticillata* flowers 39 substances were found, the dominant ones were cyclofenchene, pseudolimone, pyranone, caryophyllene, spatulenol and phytol.

Dominant compounds among substances of terpenoid nature are: cyclofenchene, camphene, 1,8-cineole,  $\alpha$ -

thujone,  $\beta$ -thujone, camphor borneol, caryophyllene, humulene, viridiflorol, sabinene, pyranone,  $\beta$ -pinene, phytol, kolavenol,  $\beta$ -pinene, loliolide, pseudolimonene and spatulenol. Among the dominant substances, 8 were detected for the first time in these species: cyclofenchene, viridiflorol, sabinene, pyranone, phytol, kolavenol, loliolide and pseudolimonene.

The raw materials of all three studied species *S. grandiflora*, *S. pratensis* and *S. verticillata* are rich in caryophyllene, so they could be potential sources of it. The widespread plant volatile beta-caryophyllene (BCP) was recently identified as a natural selective agonist of the peripherally expressed cannabinoid receptor 2 (CB2). Number of studies have shown that CB2 is critically involved in the modulation of inflammatory and neuropathic pain responses. Thus, the natural plant product BCP may be highly effective in the treatment of long lasting, debilitating pain states [38]. More importantly, BCP inhibited EtOH-CPP acquisition and exacerbated LORR duration. Interestingly, these effects were abrogated when mice were pre-injected with a selective CB2 receptor antagonist, AM630. Overall, the CB2 receptor system appears to be involved in alcohol dependence and sensitivity and may represent a potential pharmacological target for the treatment of alcoholism [39]. Moreover, oral administration of BCP effectively rescued  $\beta$ -cells by mitigating hyperglycemia through enhancing insulin release and also averted oxidative/inflammatory stress in pancreatic tissue of diabetic rats. The efficacy of BCP was compared with glibenclamide, a standard antidiabetic drug [40]. The characterization tests indicated that  $\beta$ CP were efficiently incorporated into  $\beta$ CD. The oral treatment with  $\beta$ CP- $\beta$ CD, at all doses tested, produced a significant ( $p < 0.05$ ) reduction on mechanical hyperalgesia and a significant ( $p < 0.05$ ) increase in muscle withdrawal thresholds, without produce any alteration in force. In addition,  $\beta$ CP- $\beta$ CD was able to significantly ( $p < 0.05$ ) decrease Fos expression in the superficial dorsal horn. Thus,  $\beta$ CP- $\beta$ CD attenuates the non-inflammatory chronic muscle pain in mice and inhibits the Fos expression in the lumbar spinal cord [41].

In addition, all three studied species *S. grandiflora*, *S. pratensis* and *S. verticillata* are rich in phytol and its derivatives, so they could be potential sources of it. In the pharma-medico viewpoint, PYT and its derivatives have been evident to have antimicrobial, cytotoxic, antitumorous, antimutagenic, anti-teratogenic, antibiotic-chemotherapeutic, antidiabetic, lipid lowering, antispasmodic, anticonvulsant, antinociceptive, antioxidant, anti-inflammatory, anxiolytic, antidepressant, immunoadjuvancy, hair growth facilitator, hair fall defense and antidandruff activities. Otherwise, the important biometabolite of PYT is phytanic acid (PA). Evidence shows PA to have cytotoxic, anticancer, antidiabetic, lipid lowering and antiteratogenic activities. In addition, it may be considered as an important biomarker for some diseases such as Refsum's Disease (RD), Sjögren Larsson syndrome (SLS), rhizomelic chondrodysplasia punctata (RZCP), chronic polyneuropathy (CP), Zellweger's disease hyperpipecolic academia (ZDHA) and related diseases. Thus, phytol may be considered as a new drug candidate [42, 43].

The raw materials of the studied species *S. grandiflora*, *S. pratensis* and *S. verticillata* are rich in  $\gamma$ -Sitosterol, so they could be potential sources of it. The study demon-

strates in vitro results, which support the ethnomedical use of  $\gamma$ -Sitosterol against cancer. Experimental results of this study suggest that  $\gamma$ -Sitosterol exerts potential anticancer activity through the growth inhibition, cell cycle arrest and the apoptosis on cancer cells [44].  $\gamma$ -sitosterol was screened for its antidiabetic property in streptozotocin (STZ) induced diabetic rats. Furthermore  $\gamma$ -sitosterol showed antihyperlipidemic activity as evidenced by significant decrease in serum total cholesterol, triglycerides and very low density lipoprotein-cholesterol levels coupled with elevation of high density lipoprotein-cholesterol levels in treated rats. A significant decrease in the activities of alanine aminotransaminase, aspartate aminotransaminase, alkaline phosphatase and acid phosphatase in  $\gamma$ -sitosterol treated rats when compared to diabetic control rats indicated its protective role against liver damage.  $\gamma$ -Sitosterol increased insulin secretion in response to glucose. Immunohistochemical study of pancreas also confirmed the biochemical findings. These results indicated that  $\gamma$ -sitosterol possessed antihyperglycemic activity [45].

Given the qualitative composition and quantitative content of terpene substances, promising raw materials for *S. grandiflora* are leaves, and for *S. pratensis* and *S. verticillata* – flowers, so they are promising agents for introduction into pharmaceutical practice.

**Study limitations.** Not all substances could be identified during the GC MS study of raw materials, as they were not available in the library that was used.

**The prospects for the further research.** To create new drugs, it is advisable to conduct a more thorough phytochemical study of raw materials of the studied species of sage, to standardize these raw materials and obtain galenic drugs based on it.

## 6. Conclusions

The leaves of *S. officinalis*, leaves, stems and flowers of *S. grandiflora*, *S. pratensis* and *S. verticillata* of the flora of Ukraine were studied by chromat-mass spectrometric method. As a result of the study, 243 substances were identified, of which 149 were identified. The dominant compounds among terpenoid substances are: cyclofenchene, camphene, 1,8-cineole,  $\alpha$ -thujone,  $\beta$ -thujone, camphor borneol, caryophyllene, humulene, viridiflorol, sabinene, pyranone,  $\beta$ -pinene, phytol, kolavenol,  $\beta$ -copaen, loliolide, pseudolimonene and spatulenol. Among the dominant substances, 8 were detected for the first time in these species: cyclofenchene, viridiflorol, sabinene, pyranone, phytol, kolavenol, loliolide and pseudolimonene.

A promising raw material containing terpene compounds for *S. grandiflora* are leaves, and for *S. pratensis* and *S. verticillata* – flowers, so they are promising agents for introduction into pharmaceutical practice.

## Conflict of interests.

Authors declare there is no conflict of interest.

## Funding.

The research was funded by the Ministry of Health Care of Ukraine at the expense of the State Budget in the framework No. 2301020 “Scientific and scientific-technical activity in the field of health protection” on the topic “Modern approaches to the creation of new medicines for a correction of metabolic syndrome”.

## References

1. Komarov, V. L. (1954). Shalfei– Salvia. Flora SSSR. Moscow-Leningrad, 21, 244–374.
2. Komarov, V. L. (1991). Rastitelnye resursy SSSR: TSvetkovye rasteniia, ikh khimicheskii sostav, ispolzovanie. Saint Petersburg: Nauka, 200.
3. Kamatou, G. P. P., Viljoen, A. M., Steenkamp, P. (2010). Antioxidant, antiinflammatory activities and HPLC analysis of South African Salvia species. Food Chemistry, 119 (2), 684–688. doi: <http://doi.org/10.1016/j.foodchem.2009.07.010>
4. Shanaida, M., Hudz, N., Korzeniowska, K., Wiczorek, P. P. (2018). Antioxidant activity of essential oils obtained from aerial part of some Lamiaceae species. International Journal of Green Pharmacy, 12 (3), 200–204.
5. Eidi, M., Eidi, A., Bahar, M. (2006). Effects of Salvia officinalis L. (sage) leaves on memory retention and its interaction with the cholinergic system in rats. Nutrition, 22 (3), 321–326. doi: <http://doi.org/10.1016/j.nut.2005.06.010>
6. Tildesley, N., Kennedy, D., Perry, E., Ballard, C., Wesnes, K., Scholey, A. (2005). Positive modulation of mood and cognitive performance following administration of acute doses of Salvia lavandulaefolia essential oil to healthy young volunteers. Physiology & Behavior, 83 (5), 699–709. doi: <http://doi.org/10.1016/j.physbeh.2004.09.010>
7. Mahdizadeh, R., Moein, S., Soltani, N., Malekzadeh, K., Moein, M. (2018). Study of molecular mechanism of Salvia species in prevention of diabetes. International Journal of Pharmaceutical Sciences and Research, 9, 4512–4521. doi: [http://doi.org/10.13040/ijpsr.0975-8232.9\(11\).4512-21](http://doi.org/10.13040/ijpsr.0975-8232.9(11).4512-21)
8. Eidi, M., Eidi, A., Zamanizadeh, H. (2005). Effect of Salvia officinalis L. leaves on serum glucose and insulin in healthy and streptozotocin-induced diabetic rats. Journal of Ethnopharmacology, 100 (3), 310–313. doi: <http://doi.org/10.1016/j.jep.2005.03.008>
9. Bjarnason, I., Scarpignato, C., Holmgren, E., Olszewski, M., Rainsford, K. D., Lanås, A. (2018). Mechanisms of Damage to the Gastrointestinal Tract From Nonsteroidal Anti-Inflammatory Drugs. Gastroenterology, 154 (3), 500–514. doi: <http://doi.org/10.1053/j.gastro.2017.10.049>
10. Ghorbani, A., Esmailizadeh, M. (2017). Pharmacological properties of Salvia officinalis and its components. Journal of Traditional and Complementary Medicine, 7 (4), 433–440. doi: <http://doi.org/10.1016/j.jtcm.2016.12.014>
11. Zagayko, A. L., Kolisnyk, T. Y., Chumak, O. I., Ruban, O. A., Koshovyi, O. M. (2018). Evaluation of anti-obesity and lipid-lowering properties of Vaccinium myrtillus leaves powder extract in a hamster model. Journal of Basic and Clinical Physiology and Pharmacology, 29 (6), 697–703. doi: <http://doi.org/10.1515/jbcpp-2017-0161>
12. Nizhenkovska, I. V., Tsurkan, O. O., Sedko, K. V. (2014). Shavlīia likarska – suchasni aspekty zastosuvannia (Ohliad literatury). Fitoterapiia. Chasopys, 2, 58–61.
13. Raja, R. R. (2012). Medicinally Potential Plants of Labiatae (Lamiaceae) Family: An Overview. Research Journal of Medicinal Plant, 6 (3), 203–213. doi: <http://doi.org/10.3923/rjmp.2012.203.213>
14. Koshovyi, O. N., Vovk, G. V., Akhmedov, E. Yu., Komissarenko, A. N. (2015). The study of the chemical composition and pharmacological activity of Salvia officinalis leaves extracts getting by complex processing. Azerbaijan Pharmaceutical and Pharmacotherapy Journal, 15 (1), 30–34.
15. Verkhovodova, Yu., Kireyev, I., Koshovyi, O., Myha, M., Osolodchenko, T. (2020). The effect of common sage extracts on the intestinal microbiota in experimental infectious colitis. Georgian Medical News, 4 (301), 165–170.
16. Nikavar, B., Abolhasani, L., Izadpanah, H. (2008). Alpha-amylase inhibitory activities of six salvia species. Iranian Journal of Pharmaceutical Research, 7, 297–303. doi: <http://doi.org/10.22073/IJPR.2010.779>
17. Khan, A., Rehman, N., AlKharfy, K. M., Gilani, A.-H. (2011). Antidiarrheal and antispasmodic activities of Salvia officinalis are mediated through activation of K<sup>+</sup> channels. Bangladesh Journal of Pharmacology, 6 (2), 111–116. doi: <http://doi.org/10.3329/bjp.v6i2.9156>
18. Hamidpour, M., Hamidpour, R., Hamidpour, S., & Shahlari, M. (2014). Chemistry, Pharmacology, and Medicinal Property of Sage (Salvia) to Prevent and Cure Illnesses such as Obesity, Diabetes, Depression, Dementia, Lupus, Autism, Heart Disease, and Cancer. Journal of Traditional and Complementary Medicine, 4 (2), 82–88. doi: <http://doi.org/10.4103/2225-4110.130373>
19. Mashkovskii, M. D. (2010). Lekarstvennye sredstva. Moscow: OOO «Izd-vo Novaia Volna», 1216.
20. Kovalenko, V. N. (2020). Compendium 2020 – Medicines. Kyiv: MORION, 2700.
21. Koshovyi, O., Raal, A., Kovaleva, A., Myha, M., Iliina, T., Borodina, N., Komissarenko, A. (2020). The phytochemical and chemotaxonomic study of Salvia spp. growing in Ukraine. Journal of Applied Biology & Biotechnology, 8 (3), 29–36. doi: <http://doi.org/10.7324/jabb.2020.80306>
22. Semenchenko, O. M., Tsurkan, O. O., Korablova, O. A., Burmaka, O. V. (2013). Determination of volatile compounds of essential oils of different species of genus of Salvia by chromatography-mass spectrometric method. Farmatsevtichnyi zhurnal, 1, 62–65.
23. Jasiccka-Misiak, I., Poliwoda, A., Petecka, M., Buslovych, O., Shlyapnikov, V. A., Wiczorek, P. P. (2018). Antioxidant Phenolic Compounds in Salvia officinalis L. and Salvia sclarea L. Ecological Chemistry and Engineering S, 25 (1), 133–142. doi: <http://doi.org/10.1515/eces-2018-0009>
24. Kharazian, N. (2013). Identification of flavonoids in leaves of seven wild growing Salvia L. (Lamiaceae) species from Iran. Progress in Biological Sciences, 3 (2), 81–98. doi: <http://doi.org/10.22059/PBS.2013.35842>
25. Myha, M., Koshovyi, O., Gamulya, O., Iliina, T., Borodina, N., Vlasova, I. (2020). Phytochemical study of Salvia grandiflora and Salvia officinalis leaves for establishing prospects for use in medical and pharmaceutical practice. ScienceRise: Pharmaceutical Science, 1 (23), 23–28. doi: <http://doi.org/10.15587/2519-4852.2020.197299>
26. Rami, K., Zheng Guo, L. (2011). Antimicrobial activity of essential oil of Salvia officinalis L. collected in Syria. African Journal of Biotechnology, 10 (42), 8397–8402. doi: <http://doi.org/10.5897/ajb10.2615>
27. Khedher, M. R., Khedher, S. B., Chaieb, I., Tounsi, S., Hammami, M. (2017). Chemical composition and biological activities of Salvia officinalis essential oil from Tunisia. EXCLI Journal, 16, 160–173. doi: <http://doi.org/10.17179/excli2016-832>
28. Gericke, S., Lübken, T., Wolf, D., Kaiser, M., Hannig, C., Speer, K. (2018). Identification of New Compounds from Sage Flowers (Salvia officinalis L.) as Markers for Quality Control and the Influence of the Manufacturing Technology on the Chemical Composition and Antibacterial Activity of Sage Flower Extracts. Journal of Agricultural and Food Chemistry, 66 (8), 1843–1853. doi: <http://doi.org/10.1021/acs.jafc.8b00581>
29. Koshevoi, O. N. (2011). Amino-acid and monosaccharide compositions of Salvia officinalis leaves. Chemistry of Natural Compounds, 47 (3), 492–493. doi: <http://doi.org/10.1007/s10600-011-9976-3>
30. Dobrochaeva, D. N., Kotox, M. I., Prokudin, Y. N., Barbarich, A. I. (1999). Opredelitel vysshikh rastenii Ukrainy. Kyiv: Naukova dumka.
31. Koshevoy, N., Zabolotny, A., Koshevaya, I., Kostenko, E. M., Rozhnova, T. (2019). Research of moisture-meter device for bulk and liquid materials. 29th International Scientific Symposium Metrology and Metrology, MMA. doi: <http://doi.org/10.1109/2019.8935983>
32. Korobiichuk, I., Bezvesilna, O., Kachniar, M., Koshovij, M., Kvasnikov, V. (2018). Methods and Ways of Piezoelectric Accelerometers Fastening on the Objects of Research. Acta Physica Polonica A, 133 (4), 1112–1115. doi: <http://doi.org/10.12693/aphyspola.133.1112>
33. Derzhavna Farmakopeia Ukrainy. Vol. 1. (2015). Kharkiv: Derzhavne pidpriemstvo «Ukrainskyi naukovyi farmatsevtichnyi tsentr yakosti likarskykh zasobiv», 1128.

34. Krivoruchko, E., Markin, A., Samoilova, V. A., Ilina, T., Koshovyi, O. (2018). Research in the chemical composition of the bark of sorbus aucuparia. *Ceska a Slovenska Farmacie*, 67 (3), 113–115.
35. Starchenko, G., Hrytsyk, A., Raal, A., Koshovyi, O. (2020). Phytochemical Profile and Pharmacological Activities of Water and Hydroethanolic Dry Extracts of *Calluna vulgaris* (L.) Hull. *Herb. Plants*, 9 (6), 751. doi: <http://doi.org/10.3390/plants9060751>
36. Shinkovenko, I. L., Kashpur, N. V., Ilyina, T. V., Kovalyova, A. M., Goryacha, O. V. et. al. (2018). The immunomodulatory activity of the extracts and complexes of biologically active compounds of *Galium verum* L. herb. *Ceska a Slovenska Farmacie*, 67 (1), 25–29.
37. Koshovyi, O., Raal, A., Kireyev, I., Tryshchuk, N., Ilina, T., Romanenko, Y. et. al. (2021). Phytochemical and Psychotropic Research of Motherwort (*Leonurus cardiaca* L.) Modified Dry Extracts. *Plants*, 10 (2), 230. doi: <http://doi.org/10.3390/plants10020230>
38. Klauke, A.-L., Racz, I., Pradier, B., Markert, A., Zimmer, A. M., Gertsch, J., Zimmer, A. (2014). The cannabinoid CB2 receptor-selective phytocannabinoid beta-caryophyllene exerts analgesic effects in mouse models of inflammatory and neuropathic pain. *European Neuropsychopharmacology*, 24 (4), 608–620. doi: <http://doi.org/10.1016/j.euroneuro.2013.10.008>
39. Al Mansouri, S., Ojha, S., Al Maamari, E., Al Ameri, M., Nurulain, S. M., Bahi, A. (2014). The cannabinoid receptor 2 agonist,  $\beta$ -caryophyllene, reduced voluntary alcohol intake and attenuated ethanol-induced place preference and sensitivity in mice. *Pharmacology Biochemistry and Behavior*, 124, 260–268. doi: <http://doi.org/10.1016/j.pbb.2014.06.025>
40. Basha, R. H., Sankaranarayanan, C. (2016).  $\beta$ -Caryophyllene, a natural sesquiterpene lactone attenuates hyperglycemia mediated oxidative and inflammatory stress in experimental diabetic rats. *Chemico-Biological Interactions*, 245, 50–58. doi: <http://doi.org/10.1016/j.cbi.2015.12.019>
41. Quintans-Júnior, L. J., Araújo, A. A. S., Brito, R. G., Santos, P. L., Quintans, J. S. S., Menezes, P. P. et. al. (2016).  $\beta$ -caryophyllene, a dietary cannabinoid, complexed with  $\beta$ -cyclodextrin produced anti-hyperalgesic effect involving the inhibition of Fos expression in superficial dorsal horn. *Life Sciences*, 149, 34–41. doi: <http://doi.org/10.1016/j.lfs.2016.02.049>
42. Islam, M. T., de Alencar, M. V. O. B., da Conceição Machado, K., da Conceição Machado, K., de Carvalho Melo-Cavalcante, A. A., de Sousa, D. P., de Freitas, R. M. (2015). Phytol in a pharma-medico-stance. *Chemico-Biological Interactions*, 240, 60–73. doi: <http://doi.org/10.1016/j.cbi.2015.07.010>
43. Islam, M. T., Ali, E. S., Uddin, S. J., Shaw, S., Islam, M. A., Ahmed, M. I. et. al. (2018). Phytol: A review of biomedical activities. *Food and Chemical Toxicology*, 121, 82–94. doi: <http://doi.org/10.1016/j.fct.2018.08.032>
44. Sundarraj, S., Thangam, R., Sreevani, V., Kaveri, K., Gunasekaran, P., Achiraman, S., Kannan, S. (2012).  $\gamma$ -Sitosterol from *Acacia nilotica* L. induces G2/M cell cycle arrest and apoptosis through c-Myc suppression in MCF-7 and A549 cells. *Journal of Ethnopharmacology*, 141 (3), 803–809. doi: <http://doi.org/10.1016/j.jep.2012.03.014>
45. Balamurugan, R., Duraipandiyar, V., Ignacimuthu, S. (2011). Antidiabetic activity of  $\gamma$ -sitosterol isolated from *Lippia nodiflora* L. in streptozotocin induced diabetic rats. *European Journal of Pharmacology*, 667 (1-3), 410–418. doi: <http://doi.org/10.1016/j.ejphar.2011.05.025>

*Received date 16.07.2021*

*Accepted date 08.10.2021*

*Published date 29.10.2021*

**Mykhailo Myha**, Postgraduate Student, Department of Pharmacognosy, National University of Pharmacy, Pushkinska str., 53, Kharkiv, Ukraine, 61002

**Oleh Koshovyi\***, Doctor of Pharmaceutical Sciences, Professor, Head of Department, Department of Pharmacognosy, National University of Pharmacy, Pushkinska str., 53, Kharkiv, Ukraine, 61002

**Yevhen Karpun**, Assistant, Department of Natural Sciences for Foreign Students and Toxicological Chemistry, Zaporizhzhia State Medical University, Maiakovskoho ave., 26, Zaporizhzhia, Ukraine, 69035

**Alla Kovaleva**, Doctor of Pharmaceutical Sciences, Professor, Department of Pharmacognosy, National University of Pharmacy, Pushkinska str., 53, Kharkiv, Ukraine, 61002

**Olga Mala**, PhD, Associate Professor, Department of Botany, National University of Pharmacy, Pushkinska str., 53, Kharkiv, Ukraine, 61002

**Volodymyr Parchenko**, Doctor of Pharmaceutical Sciences, Professor Department of Natural Sciences for Foreign Students and Toxicological Chemistry, Zaporizhzhia State Medical University, Maiakovskoho ave., 26, Zaporizhzhia, Ukraine, 69035

**Oleksandr Panasenکو**, Doctor of Pharmaceutical Sciences, Head of Department, Department of Natural Sciences for Foreign Students and Toxicological Chemistry, Zaporizhzhia State Medical University, Maiakovskoho ave., 26, Zaporizhzhia, Ukraine, 69035

**Vera Bunyatyan**, PhD, Associate Professor, I. M. Sechenov First Moscow State Medical University of the Ministry of Health of the Russian Federation, Trubetskaya str., 8, Moscow, Russian Federation, 119991

**Sergiy Kovalenko**, Doctor of Chemical Sciences, Professor, Department of Organic Chemistry, V. N. Karazin Kharkiv National University, Svobody sq., 4, Kharkiv, Ukraine, 61022

*\*Corresponding author: Oleh Koshovyi, e-mail: [oleh.koshovyi@gmail.com](mailto:oleh.koshovyi@gmail.com)*