

# Fatty Acids, Sterols, and Triterpenes of the Fruits of 8 *Heracleum* Taxa

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## Abstract

Fatty acids (FAs), sterols, and triterpenes of dichloromethane extracts of the fruits of 8 *Heracleum* L. taxa (Apiaceae) from southeastern Europe were investigated by gas chromatography with flame ionization detector and gas chromatography with mass spectrometry. In order to analyze the FAs, their volatile methyl esters were obtained by saponification and subsequent esterification of oily supernatants of the fruit extracts. Dominant was petroselinic acid (42.8%-56.5%), followed by linoleic (20.3%-33.3%) and oleic acids (12.3%-13.7%). Sterols and triterpenes were analyzed as volatile derivatives obtained by the silanization of residual unsaponifiable fractions. Among them, the most abundant was  $\beta$ -sitosterol (44.9%-56.9%), followed by stigmaterol (15.7%-25.0%),  $\Delta^7$ -stigmastenol (6.6%-12.5%), and campesterol (5.2%-8.1%). The quantity of petroselinic acid was also determined by the external standard method (298.8-433.4 mg/g of oily supernatant). The obtained results show that the investigated plants are potential valuable sources of the compounds utilized in different industries.

## Keywords

*Heracleum* taxa, Apiaceae, fruits, GC-FID and GC-MS, fatty acids, sterols, triterpenes

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Plants belonging to the Apiaceae family are widely distributed in temperate climate regions, and the fruits of some of these species (eg, *Coriandrum* L. and *Carum* L. spp.) are valued as sources for the extraction of many important oleochemicals, which can be used in the chemical, pharmaceutical, cosmetic, food, and other industries.<sup>1,2</sup> In this regard, plants from the genus *Heracleum* L. (cow parsnips, hogweeds) are insufficiently investigated. The previous investigations were mainly focused on the composition of their essential oils, some of which exhibited significant bioactivities (eg, antibacterial, antifungal, cytostatic, anticonvulsive, anti-inflammatory, and analgesic, in some cases comparable with or even better than the effects of the reference drugs).<sup>3-8</sup>

In the focus of the present study are 8 *Heracleum* taxa collected in southeastern Europe: *Heracleum sphondylium* L. (HSPH), *Heracleum sibiricum* L. (HSIB), *Heracleum montanum* Schleich. ex Gaudin (HMON), *Heracleum ternatum* Velen. (HTER), *Heracleum pyrenaicum* subsp. *pollinianum* (Bertol.) F. Pedrotti & Pignatti (HPOL), *H. pyrenaicum* subsp. *orsinii* (Guss.) F. Pedrotti & Pignatti (HORS), and *Heracleum verticillatum* Pančić (HVER), all belonging to the *H. sphondylium* group,<sup>9</sup> as well as *Heracleum orphanidis* Boiss. (HORP). Taxa of the *H. sphondylium* group can grow up to 3.5 m high, with umbels circa 20 cm wide, producing an abundance of fruits. *Heracleum orphanidis* has a stem up

to 50 cm high and umbels circa 5 cm in diameter.<sup>10</sup> Recently, we reported on the furanocoumarins of crystalline precipitates from dichloromethane extracts of the fruits of these plants.<sup>11</sup> In this study, we investigated the fatty acids (FAs), sterols, and triterpenes of oily supernatants of these *Heracleum* fruit dichloromethane extracts.

Oily supernatants of dichloromethane extracts of the fruits of the investigated *Heracleum* taxa were subjected to saponification and subsequent esterification to obtain volatile fatty acid methyl esters (FAME). The identified FAs, including their quantities determined by the peak area normalization method, are presented in Table 1. The investigated oily supernatants had both qualitatively and quantitatively very similar FA compositions. The dominant were monounsaturated (57.8%-70.3%). Each

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**Table 1.** Fatty Acid Composition of Oily Supernatants of Dichloromethane Extracts of Investigated *Heracleum* Fruits (%).

No.	R <sup>c</sup>	FA <sup>b</sup>	HSPH	HSIB	HMON	HTER	HPOL	HORS	HVER	HORP
1	10.66	Caprylic acid	C8:0	0.1 ± 0.0 (ab)	tr (a)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)	0.1 ± 0.0 (a)	-	-
2	12.46	Capric acid	C10:0	tr (a)	tr (a)	tr (a)	tr (a)	tr (a)	-	-
3	15.13	Lauric acid	C12:0	0.1 ± 0.0 (ab)	tr (a)	tr (a)	tr (a)	0.1 ± 0.0 (a)	tr (a)	0.1 ± 0.0 (a)
4	18.40	Myristic acid	C14:0	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)	0.1 ± 0.0 (ab)	tr (a)	0.1 ± 0.0 (a)
5	20.10	Pentadecanoic acid	C15:0	0.1 ± 0.0 (a)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)	0.1 ± 0.0 (a)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)
6	21.81	Palmitic acid	C16:0	5.1 ± 0.1 (f)	6.4 ± 0.0 (f)	5.1 ± 0.2 (g)	5.3 ± 0.1 (f)	5.4 ± 0.3 (e)	4.4 ± 0.0 (g)	5.5 ± 0.4 (e)
7	22.95	Palmitoleic acid	C16:1n7c	0.2 ± 0.1 (ab)	0.2 ± 0.0 (ab)	0.1 ± 0.0 (ab)	0.2 ± 0.0 (a)	0.2 ± 0.0 (ab)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)
8	23.47	Heptadecanoic acid	C17:0	0.1 ± 0.1 (a)	0.1 ± 0.0 (ab)	tr (ab)	0.1 ± 0.0 (a)	tr (a)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)
9	25.08	Stearic acid	C18:0	1.8 ± 0.1 (e)	1.5 ± 0.0 (e)	1.7 ± 0.0 (f)	1.4 ± 0.0 (e)	1.5 ± 0.0 (d)	1.1 ± 0.0 (f)	1.6 ± 0.1 (d)
10	25.97	Petroselinic acid	C18:1n12c	43.5 ± 1.1 (i)	44.6 ± 0.0 (i)	42.8 ± 0.3 (i)	51.3 ± 0.7 (i)	49.5 ± 2.9 (h)	52.4 ± 0.5 (j)	56.5 ± 3.7 (h)
11	26.04	Oleic acid	C18:1n9c	13.3 ± 0.4 (g)	12.7 ± 0.1 (g)	13.6 ± 0.1 (g)	12.6 ± 0.3 (g)	12.3 ± 0.7 (f)	13.7 ± 0.1 (h)	12.6 ± 0.8 (f)
12	26.15	cis-Vaccenic acid	C18:1n7c	1.0 ± 0.0 (d)	1.3 ± 0.0 (d)	0.9 ± 0.0 (d)	1.0 ± 0.0 (d)	0.9 ± 0.0 (c)	1.0 ± 0.0 (ef)	0.8 ± 0.0 (bc)
13	27.38	Linoleic acid	C18:2n6c	32.1 ± 0.8 (h)	30.0 ± 0.0 (h)	33.3 ± 0.5 (h)	27.4 ± 1.2 (i)	27.7 ± 1.5 (g)	25.3 ± 0.3 (i)	20.3 ± 1.2 (g)
14	28.11	Arachidic acid	C20:0	0.7 ± 0.1 (c)	0.7 ± 0.2 (c)	0.4 ± 0.1 (c)	0.6 ± 0.0 (c)	0.5 ± 0.0 (b)	0.3 ± 0.1 (d)	0.4 ± 0.1 (ab)
15	28.88	α-Linolenic acid	C18:3n3	1.1 ± 0.0 (d)	1.0 ± 0.0 (d)	1.2 ± 0.1 (de)	0.8 ± 0.0 (d)	0.9 ± 0.0 (c)	0.9 ± 0.0 (e)	1.0 ± 0.0 (cd)
16	29.00	cis-11-Eicosenoic acid	C20:1n9c	0.5 ± 0.0 (bc)	0.6 ± 0.0 (c)	0.3 ± 0.0 (bc)	0.3 ± 0.0 (b)	0.3 ± 0.1 (ab)	0.3 ± 0.0 (cd)	0.4 ± 0.1 (ab)
17	30.98	Behenic acid	C22:0	0.2 ± 0.0 (ab)	0.3 ± 0.1 (b)	0.2 ± 0.0 (abc)	0.3 ± 0.1 (b)	0.2 ± 0.0 (ab)	0.3 ± 0.0 (cd)	0.3 ± 0.1 (a)
18	33.85	Lignocenic acid	C24:0	0.2 ± 0.0 (ab)	0.2 ± 0.0 (ab)	0.1 ± 0.0 (ab)	0.1 ± 0.0 (a)	0.1 ± 0.0 (ab)	0.2 ± 0.1 (bc)	0.2 ± 0.0 (a)
		SFA		8.3	9.6	7.8	8.1	8.2	6.4	8.3
		MUFA		58.6	59.4	57.8	63.5	63.2	67.4	70.3
		PUFA		33.2	31.1	34.5	28.4	28.6	26.2	21.4
		Total identified		100.0	100.0	100.0	100.0	100.0	100.0	100.0

FAs, fatty acids; HMON, *Heracleum montanum* Schleich, ex Gaudin; HORS, *Heracleum orphanidis* Boiss.; HORS, *Heracleum pyrenaicum* subsp. *orsinii* (Guss.) F. Pedrotti & Pignatti; HPOL, *Heracleum pyrenaicum* subsp. *pollinianum* (Bertol.) F. Pedrotti & Pignatti; HSIB, *Heracleum sibiricum* L.; HSPH, *Heracleum sphondylium* L.; HTER, *Heracleum ternatum* Velen.; HVER, *Heracleum verticillatum* Panché; MUFA, monounsaturated fatty acid; PUFA, polyunsaturated fatty acid; Rt, retention times; SFA, saturated fatty acid.

<sup>a</sup>Retention times on HP-88 column (min).

<sup>b</sup>Investigated as fatty acid methyl esters.

<sup>c</sup>Relative area percentages of the compounds obtained from flame ionization detector area percent data (expressed as the means of 3 determinations ± standard deviations) or tr—trace (<0.1%).

<sup>d</sup>Significant differences ( $P < 0.05$ ) between the quantities of fatty acids in an oily supernatant are indicated by different letters in brackets.

supernatant contained significant ( $P < 0.05$ ) quantities of petroselinic (42.8%-56.5%) and oleic (12.3%-13.7%) acids, as well as of polyunsaturated linoleic acid (LA) (20.3%-33.3%). The amount of petroselinic acid was also determined by the external standard method and was  $348.1 \pm 0.5$  mg/g of HSPH,  $298.8 \pm 0.7$  mg/g of HSIB,  $340.1 \pm 1.7$  mg/g of HMON,  $374.2 \pm 10.0$  mg/g of HTER,  $355.4 \pm 12.2$  mg/g of HPOL,  $375.8 \pm 7.1$  mg/g of HORS,  $433.4 \pm 1.1$  mg/g of HVER, and  $420.0 \pm 10.6$  mg/g of HORP oily supernatant. Our study is in agreement with the one of Kleiman and Spencer,<sup>12</sup> where petroselinic (47.4%-56.3%), linoleic (24.6%-31.7%), and oleic (12.2%-14.4%) acids were the dominant ones in the fruit fatty oils of several *Heracleum* taxa, including *H. sphondylium*, *H. sibiricum*, *H. montanum*, and *H. pyrenaicum* subsp. *orsinii* collected in either southeastern Europe or Turkey. In addition, in the present research, 12 more FAs were identified, providing a more comprehensive insight into their FA compositions. A similar qualitative and quantitative FA composition was determined of the fatty oils and/or heptane extracts of the fruits of *Heracleum candicans* Wall. ex DC., *Heracleum lanatum* Michx., *Heracleum pinnatum* C.B. Clarke, *Heracleum platytaenium* Boiss., *Heracleum trachyloma* Fisch. & C.A. Mey., and *Heracleum crenatifolium* Boiss.<sup>12,13</sup>

The chemosystematic significance of petroselinic acid is well known.<sup>13</sup> Namely, it is the dominant FA in the fruits of most genera of Apiaceae.<sup>12,13</sup> For example, this FA was the most abundant in the total lipid extracts of the fruits of caraway, *Carum carvi* L. (29.46%-31.12%),<sup>2</sup> and coriander, *Coriandrum sativum* L. (76.65%).<sup>14</sup> Petroselinic acid is utilized in different industries. In the chemical industry, it is subjected to oxidative cleavage in order to obtain a mixture of adipic and lauric acids. Adipic acid can be used for a nylon polymer synthesis, whereas lauric acid is utilized as a raw material for softeners, emulsifiers, detergents, and soaps.<sup>2</sup> Petroselinic acid also synthesizes new sphorolipids, environmentally friendly biosurfactants.<sup>15</sup> In cosmetic formulations, this FA acts as a moisturizing and anti-aging agent, and in those containing  $\alpha$ -hydroxy acids, it is added as a skin-irritation reducing agent.<sup>15-18</sup> Petroselinic acid is also a potentially valuable raw material for the pharmaceutical and food industries. Namely, it was previously revealed that this FA possesses anti-inflammatory properties, since it inhibits the production of arachidonic acid metabolites and/or reduces the formation of intracellular adhesion molecules.<sup>19</sup> Additionally, the lysis of triglycerides with the incorporated petroselinic acid by pancreatic lipases occurs at much lower efficacy than the lipolysis of oleic acid rich triglycerides. Thus, it could be possibly used in low-fat diets.<sup>15,20</sup> Among the polyunsaturated FAs in the investigated *Heracleum* extracts, diunsaturated LA and triunsaturated  $\alpha$ -linolenic acid (ALA) were identified. LA and ALA are the only known essential FAs,

ie, they cannot be synthesized in the human body. Because of their effects on lipoprotein concentration, membrane fluidity, function of membrane enzymes and receptors, modulation of eicosanoid production, regulation of blood pressure, and metabolism of minerals, essential FAs have antiatherogenic and antithrombotic properties.<sup>21</sup>

Sterols and triterpenes of the oily supernatants of the investigated *Heracleum* fruit dichloromethane extracts were analyzed as their volatile trimethylsilyl derivatives obtained by the silylation of unsaponifiable fractions. The proportion of total sterols and triterpenes in the unsaponifiable fraction was  $73.3\% \pm 1.0\%$  in HSPH,  $71.9\% \pm 0.5\%$  in HSIB,  $65.3\% \pm 1.8\%$  in HMON,  $57.4\% \pm 0.8\%$  in HTER,  $62.5\% \pm 0.9\%$  in HPOL,  $62.0\% \pm 1.0\%$  in HORS,  $66.6\% \pm 1.1\%$  in HVER, and  $66.4\% \pm 0.7\%$  in HORP. The investigated unsaponifiable fractions had identical qualitative and very similar quantitative sterol and triterpene compositions (Table 2). Among them, the dominant were phytosterols (87.2%-92.5%). Namely, significant ( $P < 0.05$ ) quantities of  $\beta$ -sitosterol (44.9%-56.9%), stigmasterol (15.7%-25.0%),  $\Delta^7$ -stigmasterol (6.6%-12.5%), and campesterol (5.2%-8.1%) were determined. The health promoting benefits of phytosterols are well known. They can reduce the total and LDL cholesterol, decrease the risk of certain forms of cancer, and improve the treatment of prostate disorders. Furthermore, some phytosterols are important raw materials for the semisynthesis of various drugs with steroid structure.<sup>14,22</sup> Previously,  $\beta$ -sitosterol was isolated from the light-petroleum extract of *H. sphondylium* fruits<sup>23</sup> and from the ethanol extract of *H. pyrenaicum* Lam. aerial parts,<sup>24</sup> as well as from different extracts of various plant parts of some other *Heracleum* species, such as the petroleum fraction of the ethanol extract of *H. canescens* Lindl. roots.<sup>25</sup> On the other hand, stigmasterol was not identified in the investigated taxa previously, but was isolated from several other *Heracleum* species, eg, from the petroleum ether extract of *H. platytaenium* aerial parts.<sup>26</sup> The presence of other sterols, as well as the only identified triterpene,  $\alpha$ -amyrin (0.8%-6.0%), was reported for the first time in *Heracleum* taxa in this work. Triterpene alcohols are regularly identified as the constituents of unsaponifiable fractions of fatty oils, and  $\alpha$ -amyrin was previously detected, for example, in olive oil, *Olea oleum* and in corn oil, *Maydis oleum*.<sup>27,28</sup>

In this research, FAs, sterols, and triterpenes of the fruits of *H. ternatum*, *H. pyrenaicum* subsp. *pollinianum*, *H. verticillatum*, and *H. orphanidis* were investigated for the first time, while in the case of *H. sphondylium*, *H. sibiricum*, *H. montanum*, and *H. pyrenaicum* subsp. *orsinii*, the data for the composition of these metabolites were significantly complemented. Additionally, our results indicate that the fruits of the investigated 8 *Heracleum* taxa represent valuable natural sources for the extraction of certain oleochemicals, eg, petroselinic acid, preserving resources of other Apiaceae

**Table 2.** Sterol and Triterpene Composition of Unsaponifiable Fractions of Oily Supernatants of Dichloromethane Extracts of Investigated *Heracleum* Fruits (%).

No.	Rt <sup>a</sup>	Rt <sup>b</sup>	Constituent <sup>c</sup>	HSPH	HSIB	HMON	HTER	HPOL	HORS	HVER	HORP
1	78.34	3204	Ergostatetraenol	0.5 ± 0.0 <sup>d</sup> (bc) <sup>e</sup>	0.8 ± 0.0 (b)	0.7 ± 0.0 (a)	tr (a)	tr (a)	0.4 ± 0.0 (b)	0.1 ± 0.0 (a)	tr (a)
2	78.53	3214	Ergosterol	0.3 ± 0.0 (ab)	tr (a)	0.6 ± 0.0 (a)	tr (a)	tr (a)	tr (a)	0.1 ± 0.0 (a)	tr (a)
3	79.63	3269	Campesterol	7.0 ± 0.2 (e)	6.5 ± 0.1 (c)	8.1 ± 0.1 (b)	5.7 ± 0.1 (d)	5.2 ± 0.1 (d)	5.6 ± 0.0 (d)	6.9 ± 0.1 (e)	5.8 ± 0.1 (d)
4	80.30	3303	Stigmasterol	2.13 ± 0.1 (g)	20.4 ± 0.1 (e)	19.8 ± 0.2 (c)	20.1 ± 0.0 (f)	18.7 ± 0.1 (f)	15.7 ± 0.1 (g)	25.0 ± 0.2 (g)	19.1 ± 0.1 (f)
5	80.70	3321	$\Delta^7$ , <sup>22</sup> -Ergostadienol	0.5 ± 0.1 (bc)	1.2 ± 0.1 (b)	0.8 ± 0.1 (a)	0.1 ± 0.0 (a)	0.2 ± 0.0 (a)	0.4 ± 0.1 (b)	0.4 ± 0.1 (b)	0.2 ± 0.1 (a)
6	80.80	3328	$\Delta^7$ -Campesterol	0.1 ± 0.0 (a)	0.1 ± 0.1 (a)	0.1 ± 0.0 (a)	tr (a)	tr (a)	tr (a)	0.4 ± 0.0 (b)	0.1 ± 0.0 (a)
7	81.59	3366	$\beta$ -Sitosterol	48.9 ± 0.1 (h)	54.2 ± 0.4 (f)	56.0 ± 3.4 (d)	55.2 ± 0.2 (g)	56.9 ± 0.3 (g)	54.5 ± 0.2 (h)	44.9 ± 0.1 (h)	53.6 ± 0.1 (g)
8	82.32	3403	$\alpha$ -Amyrin	1.2 ± 0.1 (d)	0.8 ± 0.1 (b)	1.9 ± 0.1 (a)	2.6 ± 0.0 (c)	2.1 ± 0.1 (c)	6.0 ± 0.1 (e)	1.0 ± 0.1 (c)	3.1 ± 0.0 (c)
9	82.65	3419	$\Delta^7$ -Stigmasterol	9.3 ± 0.2 (f)	8.1 ± 0.2 (d)	7.7 ± 0.3 (b)	6.6 ± 0.1 (e)	8.2 ± 0.1 (e)	11.2 ± 0.2 (f)	12.5 ± 0.1 (f)	11.9 ± 0.2 (e)
10	82.95	3432	$\Delta^7$ -Avenasterol	0.7 ± 0.1 (c)	0.8 ± 0.1 (b)	0.9 ± 0.1 (a)	0.6 ± 0.0 (b)	0.9 ± 0.1 (b)	0.9 ± 0.1 (c)	2.0 ± 0.1 (d)	0.7 ± 0.0 (b)
Total identified sterols and triterpenes				89.9	93.0	96.5	90.8	92.3	94.8	93.3	94.5

HMON; *Heracleum montanum* Schleich. ex Gaudin; HORP; *Heracleum orphanidis* Boiss.; HORS; *Heracleum pyrenaicum* subsp. *orsinii* (Guss.) F. Pedrotti & Pignatti; HPOL; *Heracleum pyrenaicum* subsp. *pollinianum* (Bertol.) F. Pedrotti & Pignatti; HSIB; *Heracleum sibiricum* L.; HSPH; *Heracleum sphondylium* L.; HTER; *Heracleum ternatum* Velen.; HVER; *Heracleum verticillatum* Pančić; RI, retention indices; Rt, retention times.

<sup>a</sup>Retention times on HP-5MS column (min).

<sup>b</sup>Retention indices on HP-5MS column relative to C8-C40 *n*-alkanes.

<sup>c</sup>Investigated as trimethylsilyl derivatives.

<sup>d</sup>Relative area percentages of the compounds obtained from flame ionization detector area percent data (expressed as the means of 3 determinations ± standard deviations) or tr—trace (<0.1%).

<sup>e</sup>Significant differences ( $P < 0.05$ ) between the quantities of sterols/triterpene in an unsaponifiable fraction are indicated by different letters in brackets.

fruits, such as those of *Carum* and *Coriandrum* spp., for their utilization in the pharmaceutical and food industries, and for cookery.

## Experimental

### Chemicals

Bis-(trimethylsilyl)-trifluoroacetamide (BSTFA), *cis*-6-octadecenoic acid methyl ester (petroselinic acid methyl ester) (10 mg/mL in heptane), *cis*-11-vaccenic acid methyl ester (10 mg/mL in heptane), Supelco 37 component FAME mix (in dichloromethane), ergosterol (10 mg/mL in chloroform), and methanol (HPLC grade,  $\geq 99.9\%$ ) were purchased from Sigma-Aldrich (St. Louis, MO, United States),  $\beta$ -sitosterol, stigmaterol, and dichloromethane (HPLC grade) from Carlo Erba (Val-de-Reuil, France), and a homolog series of *n*-alkanes ( $C_8$ - $C_{40}$ ) from Fluka (Buchs, Switzerland). All other reagents were of p.a. quality.

### Plant Material

The fruits of the investigated *Heracleum* taxa were collected from the wild in southeastern Europe. The plants were identified by curator/botanist of the Natural History Museum (Belgrade), Dr Marjan Niketić. Voucher specimens have been deposited in the Herbarium of the Natural History Museum, Belgrade (BEO). Collection localities and dates, as well as voucher numbers, are shown in Table 3.

### Extraction and Sample Preparation

The fruits were air-dried, powdered, and extracted twice with dichloromethane at room temperature (maceration for 3 and 2 days, drug/solvent 1:10 w/v). The solvent was evaporated under reduced pressure, and after standing at 4°C for 24 hours, the extracts were filtered to separate oily supernatants from furanocoumarin-rich crystalline precipitates (this procedure was repeated twice). Saponification of the oily

supernatants (1 g) was achieved by 50% potassium hydroxide (5 mL)/ethanol (30 mL) at 90°C for 60 minutes. Unsaponifiable residues were removed using light petroleum, and the soap-rich polar fractions were treated with hydrochloric acid to obtain free FAs, which were then collected using diethyl ether. After evaporation of the solvent, FAs were esterified by 98% sulfuric acid (1 mL)/methanol (150 mL, purity  $\geq 99.9\%$ ) at 80°C for 60 minutes to obtain volatile FAME, which were collected using light petroleum. In order to analyze sterols and triterpenes, residual unsaponifiable fractions (500  $\mu$ L of 10 mg/ml solution in dichloromethane) were treated with BSTFA (50  $\mu$ L) and held at 60°C for 45 minutes to obtain volatile trimethylsilyl derivatives. The samples were analyzed within 6 hours after derivatization. The content of oily supernatants in the fruits of the investigated *Heracleum* taxa, as well as of FAME and unsaponifiable fractions in the oily supernatants, is presented in Table 4.

### Gas Chromatography with Flame Ionization Detector and Gas Chromatography with Mass Spectrometry Analysis

The analysis of FAME was performed on an Agilent 6890N Gas chromatograph (GC, Agilent Technologies, Palo Alto, CA, United States) equipped with a split/splitless injector (260°C), a flame ionization detector (FID), and a capillary column (Agilent J&W HP-88, 100 m  $\times$  0.25 mm, 0.20  $\mu$ m film thickness), and coupled with an Agilent 5975C mass selective detector (MSD) operating in the electron ionization (EI) mode at 70 eV. The carrier gas was He at a flow rate of 1.2 mL/min. The oven temperature was initially held at 140°C for 5 minutes, then increased linearly from 140 to 240°C at 4°C/min, and finally held at 240°C for 10 minutes. The FID and MSD transfer line temperatures were 260°C and 250°C, respectively. The split ratio was 1:25 and the injected volume 1  $\mu$ L of 1% solution of FAME in dichloromethane. The identification of the FAME was

**Table 3.** Collection Localities and Dates, and Voucher Numbers of Investigated *Heracleum* Taxa.

Taxa	Collection locality	Collection date	Voucher no.
HSPH	Litija (Slovenia)	September 2015	20150704/01 BEO
HSIB	Niš (Serbia)	September 2014	20140717/01 BEO
HMON	Kamnik-Savinja Alps (Slovenia)	September 2015	20150707/01 BEO
HTER	Mt Durmitor (Montenegro)	August 2013	20130807/14 BEO
HPOL	Mt Jablanica (North Macedonia)	August 2009	20090801/32 BEO
HORS	Mt Durmitor (Montenegro)	August 2011	20110804/01 BEO
HVER	Mts Stara Planina (Serbia)	August 2013	20140722/02 BEO
HORP	Mt Baba Planina (North Macedonia)	July 2012	20120706/01 BEO

HMON, *Heracleum montanum* Schleich. ex Gaudin; HORP, *Heracleum orphanidis* Boiss.; HORS, *Heracleum pyrenaicum* subsp. *orsinii* (Guss.) F. Pedrotti & Pignatti; HPOL, *Heracleum pyrenaicum* subsp. *pollinianum* (Bertol.) F. Pedrotti & Pignatti; HSIB, *Heracleum sibiricum* L.; HSPH, *Heracleum sphondylium* L.; HTER, *Heracleum ternatum* Velen.; HVER, *Heracleum verticillatum* Pančić.



**Table 4.** The Content of Oily Supernatants in the Fruits of Investigated *Heracleum* Taxa, as well as of Fatty Acid Methyl Ester and Unsaponifiable Fractions in the Oily Supernatants, % (w/w).

Content	HSPH	HSIB	HMON	HTER	HPOL	HORS	HVER	HORP
Oily supernatant	9.30	8.46	9.65	11.73	11.85	10.57	10.23	9.72
FAME	84.5	84.8	84.5	85.6	74.8	80.2	83.8	84.4
Unsaponifiable fraction	1.34	2.19	2.61	1.73	1.90	1.36	1.57	1.77

HMON, *Heracleum montanum* Schleich. ex Gaudin; HORP, *Heracleum orphanidis* Boiss.; HORS, *Heracleum pyrenaicum* subsp. *orsinii* (Guss.) F. Pedrotti & Pignatti; HPOL, *Heracleum pyrenaicum* subsp. *pollinianum* (Bertol.) F. Pedrotti & Pignatti; HSIB, *Heracleum sibiricum* L.; HSPH, *Heracleum sphondylium* L.; HTER, *Heracleum ternatum* Velen.; HVER, *Heracleum verticillatum* Pančić.

based on the comparison of their retention times (Rt) and mass spectra with those of representative standards ran under the same chromatographic conditions, ie, Supelco 37 Component FAME Mix, petroselinic acid methyl ester, and *cis*-11-vaccenic acid methyl ester. Relative percentages of the compounds were calculated based on the peak areas from the FID data. Additionally, the quantity of the most abundant FA, petroselinic acid, was determined by the external standard method, ie, by the construction of a calibration curve of petroselinic acid methyl ester (concentration range 0.08-10.00 mg/mL;  $y = 15\,288\,881\,008.70x + 736\,572.53$ ;  $r^2 = 0.9990$ ). Gas chromatography with flame ionization detector (GC-FID) and gas chromatography with mass spectrometry (GC-MS) analysis of the unsaponifiable fractions was performed on an Agilent 7890A GC equipped with 5975C (inert XL EI/CI) MSD and a FID detector connected by a capillary flow technology two-way splitter with make-up (250 °C). A HP-5MS capillary column (Agilent, 30 m × 0.25 mm, 0.25 μm film thickness) was used. The temperature of the GC oven was programmed from 60°C to 315°C at 3°C/min and held at 315°C for 15 minutes. He was used as carrier gas at 1.3 mL/min. The split ratio was 1:10 and the injection volume was 1 μL of a previously prepared solution of trimethylsilyl derivatives. The FID and MSD transfer line temperatures were 300°C and 315°C, respectively. The MS data were acquired in EI mode at 70 eV. The identification of the compounds was based on the comparison of their retention indices (RI), Rt, and mass spectra with those of commercially available standards (β-sitosterol, stigmasterol, and ergosterol), as well as to NIST/NBS 05, Wiley libraries 8th edition and NIST Chemistry WebBook.<sup>29</sup> The linear RIs were determined in relation to a homolog series of *n*-alkanes (C<sub>8</sub>-C<sub>40</sub>) run under the same operating conditions. Relative percentages of the compounds were calculated based on the peak areas from the FID data.

### Statistical Analysis

Determinations were carried out in triplicate. The results are expressed as mean values ± standard deviation and analyzed by one-way analysis of variance, followed by Tukey's post hoc test. Values of *P* below 0.05 were considered to indicate

significant differences. The analysis was carried out by Statistical Package for the Social Sciences (SPSS) version 23.0.

### Declaration of Conflicting Interests

The author(s) declared no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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### References

- Sriti J, Wannan WA, Talou T, Mhamdi B, Hamdaoui G, Marzouk B. Lipid, fatty acid and tocol distribution of coriander fruit's different parts. *Ind Crops Prod.* 2010;31(2):294-300.
- Laribi B, Kouki K, Bettaieb T, Mougou A, Marzouk B. Essential oils and fatty acids composition of Tunisian, German and Egyptian caraway (*Carum carvi* L.) seed ecotypes: A comparative study. *Ind Crops Prod.* 2013;41:312-318.
- Ušjak LJ, Petrović SD, Drobac MM, et al. Chemical composition, antimicrobial and cytotoxic activity of *Heracleum verticillatum* Pančić and *H. ternatum* Velen. (Apiaceae) essential oils. *Chem Biodivers.* 2016;13(4):466-476.
- Ušjak L, Petrović S, Drobac M, et al. Chemical composition and bioactivity of the essential oils of *Heracleum pyrenaicum* subsp. *pollinianum* and *Heracleum orphanidis*. *Nat Prod Commun.* 2016;11(4):529-534.
- Ušjak L, Petrović S, Drobac M, et al. Essential oils of three cow parsnips - composition and activity against nosocomial and foodborne pathogens and food contaminants. *Food Funct.* 2017;8(1):278-290.
- Ušjak L, Petrović S, Drobac M, et al. Edible wild plant *Heracleum pyrenaicum* subsp. *orsinii* as a potential new source of bioactive essential oils. *J Food Sci Technol.* 2017;54(8):2193-2202.
- Tosun F, Kizilay C, Erol K, Kilic F, Kurkcuoglu M, Baser KHC. Anticonvulsant activity of furanocoumarins and the

- essential oil obtained from the fruits of *Heracleum crenatifolium*. *Food Chem.* 2008;107(3):990-993.
8. Hajhashemi V, Sajjadi SE, Heshmati M. Anti-inflammatory and analgesic properties of *Heracleum persicum* essential oil and hydroalcoholic extract in animal models. *J Ethnopharmacol.* 2009;124(3):475-480.
  9. Tonascia N. Biosystematische Untersuchungen an *Heracleum sphondylium* s.l. in der Schweiz. *Berichte des Geobotanischen Institutes der E.T.H.* 1992;58:101-120.
  10. Brummitt RK. *Heracleum* L. In: Tutin TG, Heywood VH, Burges VH, et al, eds. *Flora Europaea*. Cambridge UK: Cambridge University Press; 1968:Vol. 2. 364-366.
  11. Ušjak LJ, Drobac MM, Niketić MS, Petrović SD. Chemosystematic significance of essential oil constituents and furanocoumarins of underground parts and fruits of nine *Heracleum* L. taxa from Southeastern Europe. *Chem Biodivers.* 2018;15(12):e1800412.
  12. Kleiman R, Spencer GF. Search for new industrial oils: XVI. Umbelliflorae-seed oils rich in petroselinic acid. *J Am Oil Chem Soc.* 1982;59(1):29-38.
  13. Bağcı E. Fatty acids and tocochromanol patterns of some Turkish Apiaceae (Umbelliferae) plants; a chemotaxonomic approach. *Acta Botanica Gallica.* 2007;154(2):143-151.
  14. Sriti J, Wannas WA, Talou T, Mhamdi B, Cerny M, Marzouk B. Lipid profiles of Tunisian coriander (*Coriandrum sativum*) seed. *J Am Oil Chem Soc.* 2010;87(4):395-400.
  15. Delbeke EIP, Everaert J, Uitterhaegen E, et al. Petroselinic acid purification and its use for the fermentation of new sophorolipids. *AMB Express.* 2016;6(1):28-37.
  16. Alaluf S, Hu H-L, Green MR, et al. Cosmetic use of petroselinic acid. *EP.* 1999;1(013):178.
  17. Barrett KE, Green MR, HL H, Parmar P, Powell JR, Rawlings AV. *Skin care composition*. U.S. Patent No. 6,455,057. U.S. Patent and Trademark Office, Washington, DC; 2002.
  18. Weinkauff R, Santhanam U, Palanker LR, Januario TE, Brinker A. *Petroselinic acid as an anti-irritant in compositions containing alpha-hydroxy acids*. U.S. Patent No. 6,022,896. U.S. Patent and Trademark Office, Washington, DC; 2000.
  19. Alaluf S, Green MR, Powell JR, et al. *Petroselinic acid and its use in food*. U.S. Patent No. 6,365,175. U.S. Patent and Trademark Office, Washington, DC; 2002.
  20. Heimermann WH, Holman RT, Gordon DT, Kowalyshyn DE, Jensen RG. Effect of double bond position in octadecenoates upon hydrolysis by pancreatic lipase. *Lipids.* 1973;8(1):45-47.
  21. Tvrzicka E, Kremmyda L-S, Stankova B, Zak A, Ales Z. Fatty acids as biocompounds: their role in human metabolism, health and disease – a review. Part 1: classification, dietary sources and biological functions. *Biomedical Papers.* 2011;155(2):117-130.
  22. Rabrenović BB, Dimić EB, Novaković MM, Tešević VV, Basić ZN. The most important bioactive components of cold pressed oil from different pumpkin (*Cucurbita pepo* L.) seeds. *LWT - Food Sci Technol.* 2014;55(2):521-527.
  23. Lawrie W, McLean J, El Garby Younes M, Younes MEG. Constituents of the seeds of *Heracleum sphondylium*. *Phytochemistry.* 1968;7(11):2065-2066.
  24. Gonzalez AG, Barroso JT, López Dorta H, Luis JR, Rodríguez Luis F. Componentes de Umbelíferas XX. Componentes del *Heracleum pyrenaicum* Lam. *Anales de Química.* 1978;74:832-834.
  25. Razdan TK, Kachroo V, Harkar S, Koul GL. Furanocoumarins from *Heracleum canescens*. *Phytochemistry.* 1982;21(4):923-927.
  26. Dincel D, Hatipoğlu SD, Gören AC, Topçü G. Anticholinesterase furocoumarins from *Heracleum platytaenium*, a species endemic to the Ida Mountains. *Turkish J Chem.* 2013;37:675-683.
  27. Gelmini F, Ruscica M, Macchi C, et al. Unsaponifiable fraction of unripe fruits of *Olea europaea*: an interesting source of anti-inflammatory constituents. *Planta Med.* 2016;82(3):273-278.
  28. Lercker G, Rodriguez-Estrada MT. Chromatographic analysis of unsaponifiable compounds of olive oils and fat-containing foods. *J Chromatogr A.* 2000;881(1-2):105-129.
  29. National Institute of Standards and Technology (NIST). Chemistry WebBook. (2018) <http://webbook.nist.gov/chemistry>. (accessed 13 Nov 2018).