ESSENTIAL OILS OF THREE ASIATIC ARTEMISIA SPECIES

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Abstract—The steam-distilled essential oils from the aerial parts of three Asiatic Artemisia, A. glabella, A. rupestris and A. persica, were analysed by GC/MS. In all about 100 compounds were identified; davanone is one of the main components of A. persica.

INTRODUCTION

For several years we have been studying wild and cultivated Artemisia species, Compositae, Anthemideae [1-5]. The objective of the present study was to determine the identities of the volatile components of three Asiatic Artemisia, A. glabella Kar. et Kir., A. rupestris L. and A. persica Boiss., which are native to the mountains and steppes of Central and Western Asia. A. rupestris is known

also in Europe (e.g. in the Baltic region and Germany) [6]. Seeds of these three *Artemisiae* were brought to Italy and plants were cultivated in the Botanical Garden of Turin university.

Numerous reports on essential oils of other Artemisia species, especially on those used in the flavour industry and for medicinal purposes, have been published; however, very little is known about the volatile components of these

Table 1. Constituents of the essential oil of Artemisia glabella

Compound	% of oil	Compound	% of oil
α-Thujene	Trace	Carvacrol	2.2
α-Pinene	1.0	Benzyl butyrate	0.4
Camphene	0.2	1-p-Menthen-9-ol	0.7
β-Pinene	0.2	Eugenol	0.2
Myrcene	Trace	α-Copaene	1.4
α-Phellandrene	0.3	Methyleugenol	4.6
α-Terpinene	0.1	3,7-Dimethyloctyl acetate	0.5
p-Cymene	0.6	β-Caryophyllene	1.6
1,8-Cineole	8.6	β-Sesquiphellandrene	0.3
γ-Terpinene	0.3	β-Cubebene	1.1
Terpinolene	0.2	Sabinyl propionate	4.3
Camphor	1.1	β-Farnesene	1.0
trans-Pinocarveol	0.1	M, 204	1.5
Pinocarvone	0.1	Sabinyl butyrate	8.2
Benzyl acetate	Trace	M, 204	2.4
Terpinen-4-ol	2.1	Farnesol	1.3
α-Terpineol	2.5	1-Dodecanol	1.1
Myrtenol	0.2	M _r 220	1.4
Phellandral	0.3	Nerolidol	1.8
cis-Piperitol	0.7	Chamazulene	0.3
Cuminic aldehyde	1.0	iso-Butyl phthalate	0.4
p-Isopropil-phenol	0.5	M, 214	6.1
p-Menth-2-en-7-ol	1.6	Hexadecanoic acid	0.8
1,5- <i>p</i> -Menthadien-7-ol	4.5	Phytol	1.1
			72.0

Table 2. Constituents of the essential oil of Artemisia rupestris

Compound	% of oil	Compound	% of oil
α-Pinene	1.1	Artemone	0.3
Camphene	Trace	Citronellyl i-valerate	0.7
β-Pinene	0.4	Linalyl 3-methyl-butyrate	1.0
Myrcene	1.6	Davanone	0.7
p-Cymene	Trace	Guaiol	1.2
Limonene	0.1	M, 222	6.2
1,8-Cineole	0.2	M, 220	2.3
2-Methyl-butyl-2-methyl-butyrate	0.4	Bazzanenol	2.0
Borneol	0.1	M, 220	1.2
Terpinen-4-ol	0.3	Tetradecanoic acid	1.7
α-Terpineol	0.2	iso-Butyl phthalate	1.5
Citronellol	0.1	4.6.10-Trimethyl-2-pentadecanone	1.1
Bornyl acetate	0.3	1-Hexadecanol	18.1
α-Terpinyl acetate	0.8	Pentadecanoic acid	1.1
Citronellyl acetate	0.9	Hexadecanoic acid	11.2
B-Elemene	3.9	1-Octadecanol	0.9
β-Sesquiphellandrene	0.7	Phytol	0.8
α-Guaiene	2.5	Linoleic acid	2.5
β-Chamigrene	1.8		
			70.2

Table 3. Constituents of the essential oil of Artemisia persica

Compound	% of oil	Compound	% of oil
2-Methyl-1-butanol	0,1	iso-Amyl tigliate	1.4
Camphene	3.3	α-Terpinyl acetate	1.6
Sabinene	1.8	Unknown	2.0
α-Terpinene	0.6	β-Elemene	0.8
p-Cymene	5.4	trans-β-Farnesene	0.4
1,8-Cineole	26.6	β-Phenylethyl-2-methylbutyrate	0.6
cis-3-Hexenyl butyrate	0.6	β-Phenylethyl n-valerate	0.4
trans-3-Hexenyl butyrate	0.8	Artemone	6.7
2-Methylbutyl-2-methylbutyrate	0.9	β-Phenylethyl tigliate	1.0
n-Amyl-3-methylbutyrate	0.4	Davanone	15.0
trans-Pinocarveol	0.4	Hydroxydavanone	1.2
δ -Terpineol	0.5	Hydroxydavanone (isomer)	0.9
Terpinen-4-ol	0.8	iso-Butyl phthalate	0.6
Myrtenal	0.4	Hexadecanoic acid	0.4
α-Terpineol	0.7		
Myrtenol	0.3		
			77.4

three Artemisiae [7, 8]. In a paper on monoterpene components of the genus Artemisia Stangl and Greger [9] analyzed three different samples of A. persica; the main components were artemisia ketone for the first sample; artemisia ketone, thujyl alcohol and camphor for the second and β -thujone for the third sample. Also the presence of sesquiterpene coumarin ethers in A. persica has been reported [10, 11].

RESULTS AND DISCUSSION

The steam-distilled essential oils were analyzed by GC/MS. The yield in essential oil was 0.34% of the dried plant for A. glabella, 0.24% for A. rupestris and 0.40% for

A. persica. The identification of each component was made by comparison of the mass spectra with a collection of literature spectra or with those of authentic samples, and was also checked against the R_f value.

43 components were identified in A. glabella oil (totalling 72.0%), 33 in A. rupestris (70.2%) and 29 in A. persica (77.4%) (Tables 1-3). The components not identified were reported as 'unknown' or only with M_r (when known), if they were more abundant than 1% of the oil. iso-Butyl phthalate which is present in little amounts in the three oils is probably an impurity. Chamazulene is responsible for the blue colour of the A. glabella oil and is probably an artefact formed during distillation, as well as β -elemene.

The major constituents of the essential oil of A. persica

are 1,8 cineole (26.6%) and davanone (15.0%). Davanone is reported to occur in one of the chemotypes of *Tanacetum vulgare* [12], in *A. pallens* [13, 14] and in *A. rehan* [15]. The presence in the oil of *A. persica* of a compound (6.7%) with a mass spectrum similar to that of davanone, but with a shorter GC R_1 , which is in agreement with the data published by Naegely *et al.* [16] and the cooccurrence of artemone and davanone in the oil of *A. pallens* [13, 14] and of two davanone isomers (one probably being artemone) in *A. rehan* [15], led us to believe that this compound might be artemone.

EXPERIMENTAL

Artemisia rupestris and A. glabella seeds were imported in 1979 from the Botanical Garden of the Kazakistan-S.S.R. Sciences Academy, 470032 Karaganda, U.S.S.R.; A. persica in 1980, from the Botanical Garden of the Uzbekistan-S.S.R. Sciences Academy, Ul. Dshachan Abidovi 272, Taschkent, Uzbekistan-S.S.R., U.S.S.R. Plants were cultivated in the Botanical Garden of Turin university: a voucher is deposited in the 'herbarium'. From each plant we had at our disposal 25 g of leaves and buds.

A previous distillation of 1 g from each sample was performed on a micro-scale apparatus, already described elsewhere [17]. The whole sample (ca 24 g) was then distilled for 2 hr in a modified Marcusson apparatus [18].

Essential oil (1.0 μ l) dil 1/2500 in hexane was injected, using the on column technique, into a glass capillary column, PS 255, 25 m \times 0.32 mm i.d., film thickness 0.3 μ m, crosslinked. Temp. were prog from 50° to 200° at 3°/min. Carrier gas was H₂, flow rate 3 ml/min, with FID.

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