

ESSENTIAL OIL OF *Perovskia scrophulariifolia**

Kh. R. Nuriddinov,^a K. Kh. Khodzimatov,^b Kh. N. Aripov,^a
T. Ozek,^c B. Demirchakmak,^c and K. H. C. Basher^c

The herbal parts of Perovskia scrophulariifolia (Labiatae) collected from Yakkabag in the Kashkadarya region of Uzbekistan in August, 1993 yielded 0.54% oil by water distillation. The oil was analyzed by GC/MS. Seventy-one components were characterized representing 95% of the oil with 1,8-cineole (11%), caryophyllene oxide (10%), camphor (9%), humulene epoxide II (7.9%), bornyl acetate (7.8%), and p-cymene (5.7%) as major constituents.

The genus *Perovskia* (Labiatae) is represented in the flora of the former USSR by six species which are distributed in Central Asia, North Iran, Beluchistan, Afghanistan, West Tibet, and West India [1].

Constituents of the essential oil of the aerial parts of *P. scrophulariifolia* growing in Turkmenistan were reported as borneol, camphene, geraniol, linalool, β -pinene, sabinene, terpinene, and terpinolene without quantitative indication [2]. In a separate study, 0.2-2.1% essential oil was obtained from herbal parts of the plants growing in Tyanshan, Fergana, Pamir-Alay, and Zarafshan mountains. Major constituents found in the oils were as follows: 1,8-cineole (15-19%), camphene (3.7-6.2%), δ -3-carene (4-6.5%), linalool (3-7.2%), linalylacetate (7.9-8.4%), humulene (5-7.7%), α -betulenol (4-6.7%), β -caryophyllene (7-13%), and α -pinene (8-12%) [3].

We have analyzed the essential oil of *Perovskia scrophulariifolia* Bunge occurring in Uzbekistan. The hydrodistilled essential oil was analyzed by GC/MS and the results are given in Table 1. Seventy-one components were characterized representing 95% of the oil. 1,8-Cineole, caryophyllene oxide, camphor, humulene epoxide II, bornyl acetate, and p-cymene were found as major constituents.

EXPERIMENTAL

The plant material was collected from Yakkabag in Kashkadarya region of Uzbekistan in August, 1993. The herbal parts were subjected to hydrodistillation for 3 h using a Clevenger-type apparatus to produce essential oil in 0.54% yield.

The essential oils were analyzed by GC/MS using a Hewlett-Packard GCD system. An Innowax FSC column (60 m \times 0.25 mm \varnothing) was used with helium as carrier gas. GC oven temperature was kept at 60° for 10 min and programmed to 220°C at a rate of 4°C/min and then kept constant at 220°C for 10 min. Split ratio was adjusted at 50:1. The injector temperature was 250°C. MS were taken at 70 eV. Mass range was from m/z 10 to 425. A library search was carried out using the Wiley GC/MS Library and the TBAM Library of Essential Oil Constituents [4-9]. Relative percentage amounts of the separated compounds were calculated from total ion chromatograms by the computerized integrator.

*Materials presented at the 2nd International Conference of Natural Compounds.

^aInstitute of the Chemistry of Plant Substances, Academy of Sciences, Tashkent, Uzbekistan.

^bInstitute of Botany, Academy of Sciences, Tashkent, Uzbekistan.

^cCentre (TBAM), 26470 Eskisehir, Turkey.

TABLE 1. Composition of the Essential Oil of *Perovskia scrophulariifolia*

Compounds	Amo- unts. %	Compounds	Amo- unts. %	Compounds	Amo- unts. %
α -pinene	1.43	p-cymen-8-ol	0.32	thujol	0.30
α -thujene	0.10	α -calacorene I	0.14	α -muurolene	0.79
camphene	1.25	piperitenone	0.97	α -selinene	0.16
β -pinene	1.44	cubebol	0.08	isocaryophyllene oxide	1.75
sabinene	0.09	cis-jasmone	0.14	caryophyllene oxide	9.98
δ -3-carene	1.22	α -calacorene II	0.07	humulene epoxide I	0.54
myrcene	0.18	α -gurjunene	0.30	humulene epoxide II	7.85
limonene	2.79	linalool	0.54	elemenal*	0.45
1,8-cineole	11.0	cis-sabinenehydrate	0.07	1-epi-cubanol	0.10
p-cymene	5.71	linalylacetate	0.27	viridiflorol	0.57
α -p-dimethyl styrene	0.01	pinocarvone	0.09	cumin alcohol	0.13
α -cubebene	0.19	bornyl acetate	7.82	cis-p-menth-3-en-1,2-diol	0.09
trans- sabinenehydrate	0.15	β -caryophyllene	0.72	T-cadinol	0.42
α -copaene	0.69	terpinen-4-ol	0.54	T-muurolol	0.20
camphor	9.10	cis-p-mentha-2,8-dien- 1-ol	0.11	α -bisabolol	0.91
carvone	0.12	myrtenal	0.11	carvacrol	0.15
geranylacetate	1.49	aromadendrene	0.29	α -eudesmol	0.22
δ -cadinene	0.22	trans-pinocarveol	-0.29	α -cadinol	0.77
γ -cadinene	1.32	δ -terpineol	0.48	caryophylladienol*	0.84
selina-3,7(11)- dien	0.29	α -humulene	1.80	calacorene alcohol I*	0.26
myrtenol	0.32	cryptone	0.44	calacorene alcohol II*	0.22
trans-carveol	0.16	γ -muurolene	0.48	caryophyllenol II	0.62
calamenene	0.43	α -terpinylacetate	4.56	epi-13-manool	2.57
m-cymen-8-ol	0.24	borneol	2.81		

*Tentative identification by GC/MS data alone.

REFERENCES

1. B. K. Shishkin, Flora of the USSR (Flora SSSR) XXI, AN SSSSR, Moscow-Leningrad (1954); Israel Program for Scientific Translations, Jerusalem (1977), p. 267.
2. M. O. Karryev, Mater. Yubileinoi Respub. Nauch. Konf. Farm. Posvyashennoy 50-letiyu Obraz. SSSR, Tashkent, SSSR (1972), pp. 62-63.
3. A. D. Dembitski, Izv. An Kaz SSR, Ser. Khim., 4, 4-10 (1984).
4. A. A. Swigar and R. M. Silverstein, Monoterpenes: Infrared, Mass, $^1\text{H-NMR}$ and $^{13}\text{C-NMR}$ Spectra and Kovats Indices, Aldrich Chemical Co., Milwaukee, WI (1981).
5. W. Jennings and T. Shibamoto, Qualitative Analysis of Flavor and Fragrance Volatiles by Glass Capillary Gas Chromatography, Academic Press, London (1980).
6. Eight Peak Index of Mass Spectra, 3rd Edition, The Royal Society of Chemistry, London (1986), Vols. 1-7.
7. F. W. McLafferty and D. B. Stauffer, The Wiley/NBS Registry of Mass Spectral Data, John Wiley and Sons, New York (1988), Vols. 1-7.
8. R. P. Adams, Identification of Essential Oils by Ion Trap Mass Spectroscopy, Academic Press (1989).
9. R. P. Adams, Identification of Essential Oil Components by Gas Chromatography/Mass Spectroscopy, Allured (1995).