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RESEARCH NOTE

Essential Oils from Leaves and Inflorescence of *Ocimum basilicum* var. *purpurascens* Benth. from Northeastern Brazil

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Abstract

Essential oils from the leaves and inflorescences of *Ocimum basilicum* L. var *purpurascens* Benth. were obtained by steam distillation and using microwave distillation. Oils were investigated by GC and GC/MS. The main constituent in the leaf oils and inflorescence oil was linalool (39.3-79.6% and 81.5%).

Key Word Index

Ocimum basilicum var *purpurascens*, Labiateae, essential oil composition, linalool.

Plant Name

Ocimum basilicum var *purpurascens* Benth., local name : mangericão roxo.

Source

The plant material was collected in the Medicinal Plant Garden of the Universidade Federal do Ceará, Brazil. The botanical material was identified in the Kew Royal Garden, London, England and voucher specimens have been deposited in the Prisco Bezerra Herbarium (Ceará-Brazil) under number 18777.

Plant Part

Fresh leaves were collected in October 1996. Steam distillation produced an oil (UFC code- F2546) in 0.5% yield, and distillation using microwave oven (1) produced 0.1% yield (leaves) and only traces with inflorescences.

Previous Work

Chemical composition of the leaf oil of *O. basilicum* var *purpurascens* collected in Jammu-Tawi from tissue culture techniques (2) and in India (3) was reported in the literature. Methyl cinnamate was the

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Table I. Percentage composition of the leaf and inflorescence oils of the *Ocimum basilicum* var. *purpuracens*

Compounds	Retention			Method of identification
	index	Leaf ^a	Leaf ^b inflorescence ^b	
α -pinene	927	0.6		RRT, MS
camphene	935	0.4		RRT, MS
β -pinene	958	1.3		RRT, MS
p-cymene	1001	0.5		RRT, MS
1,8-cineole	1004	5.0	2.4	RRT, MS
limonene	1008	1.0		RRT, MS
γ -terpinene	1038	0.3		RRT, MS
cis-linalool oxide [†]	1044		2.3	RRT, MS
trans-linalool oxide [†]	1060		3.2	RRT, MS
linalool	1091	39.3	79.6	81.5
camphor	1107	0.9	2.0	RRT, MS
borneol	1139	0.6		RRT, MS
terpinen-4-ol	1155	1.8	4.3	RRT, MS
α -terpineol	1167		2.6	RRT, MS
methyl chavicol	1187	1.9		RRT, MS
octyl acetate	1204	2.3		RRT, MS
bornyl acetate	1271	0.9		RRT, MS
δ -elemene	1343	3.4		RRT, MS
α -copaene	1377	0.6		RRT, MS
β -bourbonene	1383	0.5		RRT, MS
β -cubebene	1384	2.0		RRT, MS
β -elemene	1393	0.8		1.9
β -caryophyllene	1416	0.8		RRT, MS
trans- α -bergamotene	1442	4.0		2.5
α -humulene	1449	1.6		RRT, MS
bicyclogermacrene	1491	0.7		RRT, MS
germacrene A	1499	1.0		RRT, MS
δ -guaiene	1503	0.4		RRT, MS
γ -cadinene	1512	7.7		RRT, MS
humulene epoxide II	1586	1.5		RRT, MS
1-epi-cubenol	1600	2.3		RRT, MS
T-cadinol	1612			7.0
α -muurolol	1627	11.0		RRT, MS
β -eudesmol	1630	0.5		RRT, MS
Total		95.6	96.4	92.9

^a = steam distillation; ^b = microwave distillation; [†] = furanoid form

major component identified. A recent review of basil oil from different origins was also published (4).

Present Work

Oil analysis were performed on a Hewlett-Packard 5971 GC/MS instrument employing the following conditions: Column: Dimethylpolysiloxane DB-1 fused silica capillary column (30 m x 0.25 mm, film thickness 0.1 μ m); carrier gas: Helium (1 mL/min); injector temperature: 250°C; detector temperature: 200°C; column temperature: 35°-180°C at 4°C/min then 180°-250°C at 10°C/min; mass spectra: electronic impact 70 eV. Compounds were identified by mass spectral library search (5-7) and retention indices (8). The oil constituents are presented in Table I. The extraction using microwave oven produced linalool

in highest yield. The chemotype growing in Northeastern Brazil is very different to the chemotype growing in India, and is very similar to the *O. basilicum* of Taiwanese origin (9).

Acknowledgments

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