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Article

Lavandula stoechas subsp. *luisieri* (Rozeira) Rozeira: Variability of Chemical Composition of Essential Oil in Wild Populations

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Abstract

Lavandula stoechas subsp. *luisieri* (Rozeira) Rozeira is an aromatic shrub endemic to the south-west of the Iberian Peninsula. It is distinguished by being the only species of lavender that contains necrodol derivatives in its essential oil. This study aims to evaluate the diversity of the chemical composition of *L. stoechas* subsp. *luisieri* essential oil and how it differs from essential oils of other lavender species with which it shares its habitat and with which it can hybridize. *The variability in the chemical composition of L. stoechas subsp. luisieri essential oil has been evaluated across 66 populations distributed among 14 areas in southwestern Iberian Peninsula. The main compounds present in the essential oil of Lavandula stoechas subsp. luisieri are trans- α -necrodyl acetate (20.68% $\pm$ 4.17%), 1,8-cineole (7.79% $\pm$ 7.14%) and trans- α -necrodol (8.66% $\pm$ 2.18%). Other compounds may occasionally be present in percentages greater than 6.00%, such as α -cadinol, linalool, lavandulyl acetate, fenchone and camphor. Two chemotypes have been identified in the essential oil of L. stoechas subsp. luisieri: 1) trans- α -necrodyl acetate – 1,8-cineole, and 2) trans- α -necrodyl acetate, with little or no presence of 1,8-cineole. Furthermore, the absence or very low percentage of camphor (0.16 – 7.61%) and fenchone (0.00 – 7.39%) has been confirmed as a unique characteristic of this essential oil.*

Keywords: essential oil; *Lavandula stoechas* subsp. *luisieri*; necrodol derivatives; Spanish Lavander

1. Introduction

The genus *Lavandula* L. (Lamiaceae) comprises approximately 40 species, as well as a variety of infraspecific taxa and hybrids (around 75 taxa). These species are, generally, endemics (with a narrow distribution), that are primarily distributed in the Mediterranean region, encompassing from north Atlantic Islands to South Asia. There are important hotspots in Canary Islands (5 species and 7 subspecies), Morocco (9 species, 4 of those are endemic species) and South of the Arabian Peninsula and West of the Horn of Africa (18 species, 12 of them endemic) [1,2].

Several lavender species have been used by humans since ancient times. First written references about medicinal use of lavender were from first century BC, in “De Materia Medica (pp. 460-461)” from Dioscorides, where is mentioned the *stoechas* term refer to the use of lavender species attributable to *Lavandula stoechas* L. and *Lavandula vera* DC. (= *Lavandula angustifolia* L.). Their uses extend from Egyptian, Roman, Greek and Persian civilizations [3-5]. The wide use as aromatic and medicinal plant of these species has been maintained over millennia, thanks to their recognized properties (sedatives, carminatives, antidepressants and anti-inflammatory), and, during First World War, after the discovering of the antibiotic capacity of the *L. angustifolia* essential oil, the growing of its essential oil demanding contributed to the expansion of lavender crops [3,6,7]. At present, the cultivation of lavender is predominantly focused on three species, *L. angustifolia* (true lavender), *L.*

latifolia Medik. (spike lavender) and their hybrid *L. × intermedia* Emeric ex Loisel. (lavandin), which are present over several countries, among which stand out: Bulgaria, France, United Kingdom, Spain, Italy, Russia, Ukraine, Moldova, Romania, Hungary, Poland, Turkey, Morocco, United States, Australia, South Africa, and China [8-12].

In recent decades, a significant number of research projects have been conducted on the chemical composition, traditional uses and medicinal properties of essential oils from other lavender species, being especially high the number of published studies about *L. stoechas* [13,14].

Two subspecies of *L. stoechas* are recognized: *L. stoechas* subsp. *stoechas* and *L. stoechas* subsp. *luisieri* (Rozeira) Rozeira (Figure 1), which present differences both morphologically, in the chemical composition of their essential oils, and geographic distribution.

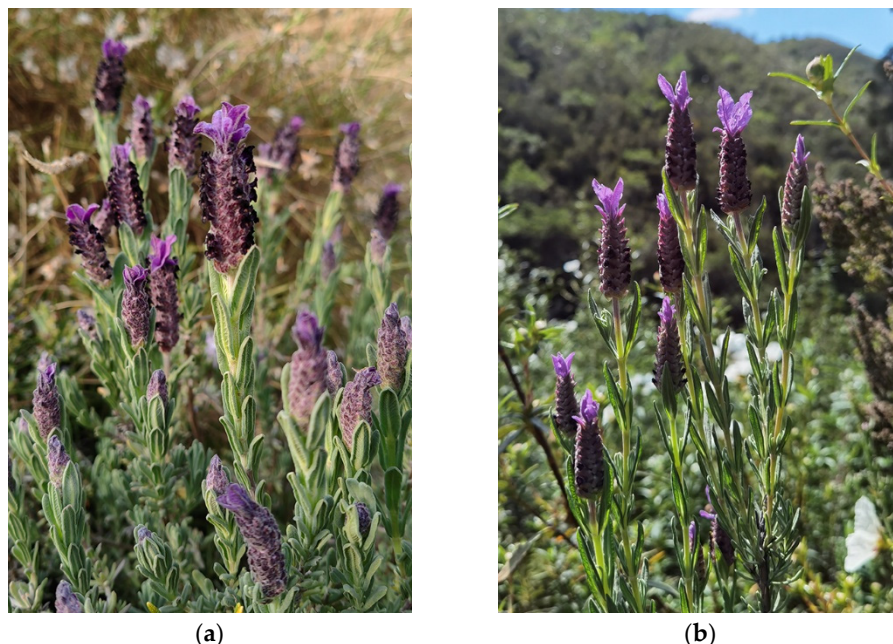


Figure 1. Photography of *L. stoechas* subspecies: (a) *L. stoechas* subsp. *stoechas*; (b) *L. stoechas* subsp. *luisieri*.

L. stoechas subsp. *stoechas* (French lavender) is an aromatic shrub native to the coastal areas of the Mediterranean region, penetrating inland where topography allows and at altitude where the climate is suitable, on coastal and internal mountain ranges, at altitudes between sea level and 1000(-1350) m (Figure 2a) [1].

Lavandula stoechas subsp. *luisieri* (Spanish lavender) is an endemic shrub to the SW of the Iberian Peninsula. It inhabits a variety of environments, including cliffs and coastal dunes, as well as inland mountain ranges, at altitudes below 900-950 m a. s. l. (Figure 2b) It is associated with pioneer plant formations of Mediterranean scrublands and with the shrub stratum of the understory in more or less dense forests of oaks (*Quercus* L. spp.) and pines (*Pinus* L. spp.) [1,15,16].

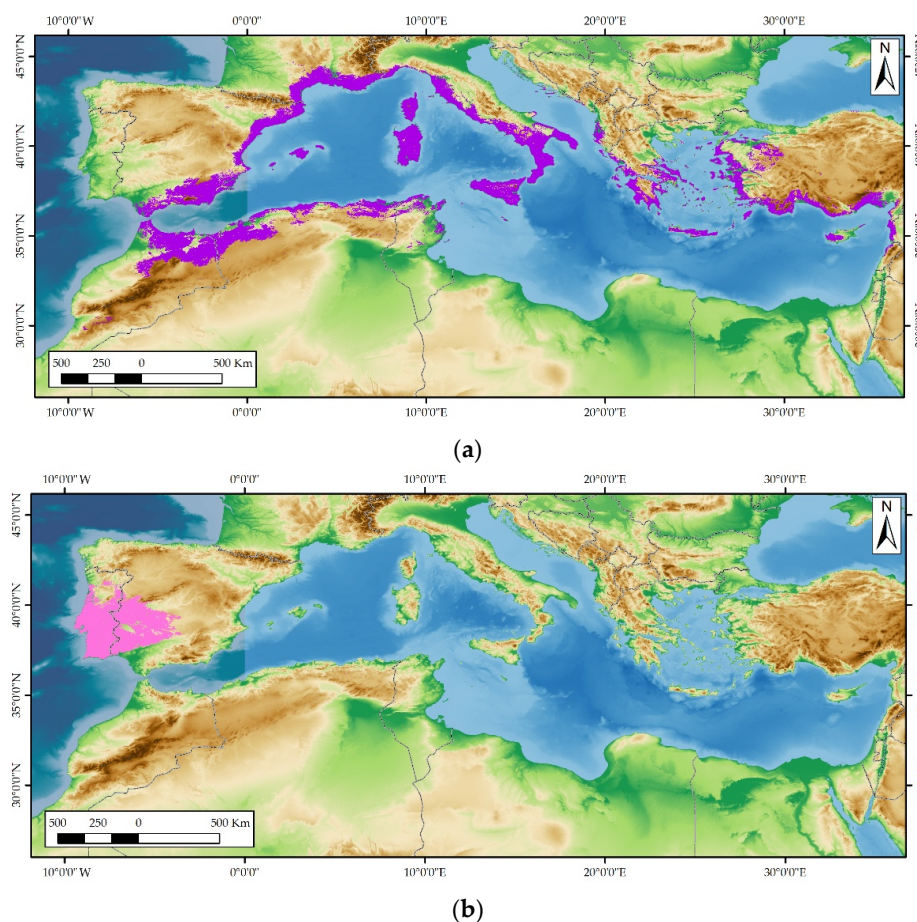


Figure 2. Distribution maps of *L. stoechas* subspecies: (a) *L. stoechas* subsp. *stoechas*; (b) *L. stoechas* subsp. *luisieri*.

(a) Morphological differences: *L. stoechas* subsp. *stoechas* is characterized by ascending stems up to 70 cm high. The inflorescence peduncle is up to 2(3) cm long and no longer than the spike itself. The indumentum of the fertile bracts is woolly (scattered to dense) with simple hairs. *L. stoechas* subsp. *luisieri* has erected stems up to 150(200) cm in height, an inflorescence peduncle reaching up to 6 cm, and the indumentum of the fertile bracts velvety, denser on the venation, with short and branched hairs [1,15,16].

(b) Chemical composition of *L. stoechas* subsp. *stoechas* essential oil has as main components fenchone [17-34], camphor [35-45], and 1,8-cineole [46], and rarely other majority components as linaol [47,48] and pulegone [49]. Instead, *L. stoechas* subsp. *luisieri* has an essential oil rich in necrodol derivatives, *trans*- α -necrodyl acetate mainly, accompanied by other components such as 1,8-cineole, lavandulyl acetate, lavandulol, viridiflorol, fenchone, and camphor [50]. However, these essential oils do not show a stable chemical composition in their whole natural distribution area and can be observed wide ranges of variation in their major components [13,50].

L. stoechas subsp. *luisieri* essential oil differs from the rest of lavender essential oils because it is the only one that contains necrodol derivatives (*trans*- α -necrodyl acetate, *cis*- α -necrodyl acetate, *trans*- α -necrodol, and *cis*- α -necrodol) [13,50]. First studies developed about chemical composition over this essential oil indicated that 1,8-cineole (0.10-43.20%) and *trans*- α -necrodyl acetate (6.30-28.40%) are the main components and camphor and fenchone have very low percentages of presence (0.40-6.30% and 0.00-5.40% respectively) [51-53]. Subsequent studies on the potential applications of the essential oil revealed the presence of camphor as the primary component (42.90–60.30%), with *trans*- α -necrodyl acetate having a residual presence (0.00–12.39%). [54-56]; and in other research, the majority component has been identified as fenchone, with a percentage of up to 36.77%. [56,57].

The presence of *trans-α-necrodyl acetate* and other *necrodol* derivatives renders the *L. stoechas* subsp. *luisieri* essential oil particularly distinctive, thereby evoking the interest of the cosmetic and pharmaceutical industries. Several studies give to it good antifungal and antibacterial activity [59-63], moderate antioxidant capacity [55,63-65], good anti-inflammatory activity [66-69] and high inhibition capacity of *proBACE-1* (enzyme related with Alzheimer disease) [69], good effects over trypanosomatid diseases [70-71], and over infections made by *Leishmania* species (Protozoal) [72]. Finally, its antifeedant activity is subject to variation, exhibiting low activity against *Hyalomma lusitanicum* [56], and between low and high against *Spodoptera littoralis*, *Leptinotarsa decemlineata*, *Myzus persicae*, and *Rhopalosiphum padi*, among others [58,73-75].

The divergent properties exhibited by this species (e.g. anti-inflammatory and antioxidant properties) in conjunction with the diversity observed in the chemical composition of its essential oil, the morphological similarity with *L. stoechas* subsp. *stoechas*, and the high hybridization capacity of *L. stoechas* subsp. *luisieri* with other lavender species (*L. pedunculata* (Mill.) Cav. and *L. viridis* L'Hér.) necessitate a comprehensive evaluation of the morphological variability and chemical composition of its essential oil throughout its natural distribution area. The objective of the present study is twofold: firstly, to ascertain the presence or absence of diverse chemotypes in *L. stoechas* subsp. *luisieri*, and secondly, to identify the chemical components that contribute to its economic value.

2. Results

A total of 66 populations of *L. stoechas* subsp. *luisieri* were analyzed. The distribution of these populations is primarily concentrated in inland mountainous regions, with elevations ranging up to 912 meters above sea level (m.a.s.l.) and are seldom observed in coastal areas or alluvial plains, with elevations ranging from 36 to 192 meters above sea level (m.a.s.l.) (see Figure 3, Table 1 and Supplementary Table S1).

Table 1. List of sampling areas for *L. stoechas* subsp. *luisieri*.

Zone	Type	Country	N	IDs	Elev.
Alconera	MR	ES	1	1	608
Atlantic coast	CZ	PT	7	2-8	36-112
Guadiana International	AP	PT	2	9-10	192-194
Montes de Toledo	MR	ES	10	11-20	515-824
Rio Tejo	AP	PT	2	21-22	204-373
Serra da Gardunha	MR	PT	5	23-27	425-617
Serra de Alvelos	MR	PT	5	28-32	311-829
Serra de Monchique	MR	PT	7	33-39	104-764
Serra de São Mamede	MR	PT-ES	2	40-41	362-526
Sierra de San Pedro	MR	ES	5	42-46	362-500
Sierra de Aracena y Picos de Aroche	MR	ES	9	47-55	260-709
Sierra Morena de Sevilla	MR	ES	8	56-63	296-709
Vegas Bajas	AP	ES	1	64	179
Villuercas	MR	ES	2	65-66	593-912

Note. N: number of populations collected per zone, IDs: identification number assigned to each population, Elev.: elevation(m.a.s.l.), Type (MR: mountain range, CZ: coastal zone; AP: alluvial plain), Country (ES: Spain, PT: Portugal).

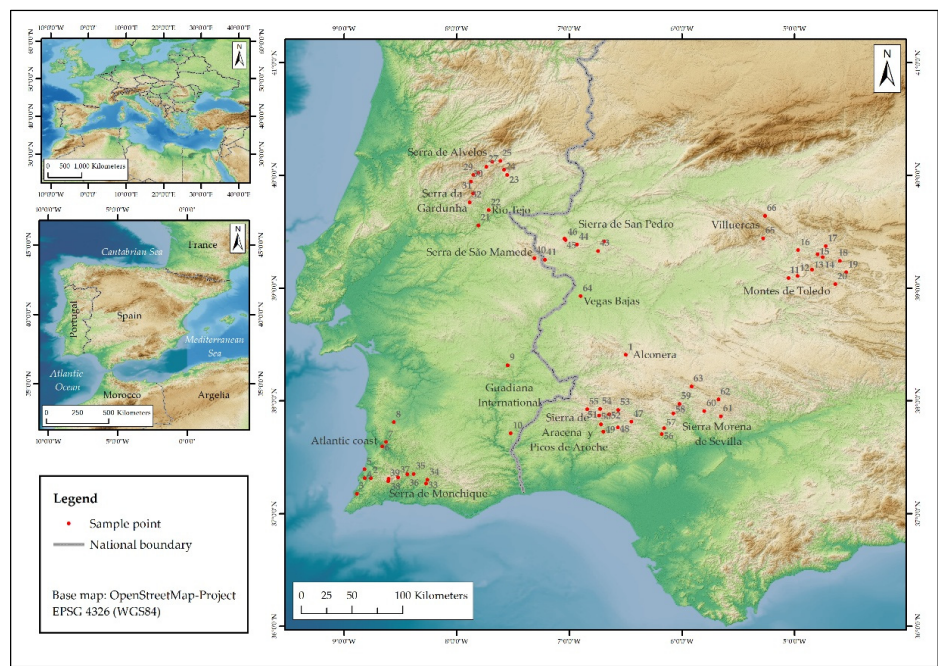


Figure 3. Distribution of *L. stoechas* subsp. *luisieri* sampling points utilized in this study.

2.1. Essential Oil Yield

The mean essential oil (EO) yield of the *L. stoechas* subsp. *luisieri* populations obtained is 3.08 ± 1.42 g/kg (grams of EO/kg fresh sample), with the lowest yield (1.08 g/kg) and highest yield (8.17 g/kg) located in Serra de Alvelos (Portugal) and Montes de Toledo (Spain), respectively. With regard to the average essential oil yield, the majority of the areas studied demonstrate an average yield of 2.50–3.50 g/kg. The lowest yields were observed in the Portuguese mountain ranges of Serra de Alvelos and Serra da Gardunha, with average values of 1.46 ± 0.45 g/kg and 2.23 ± 0.60 g/kg, respectively. The lowest yield was recorded in the isolated population located in the alluvial plain of the Guadiana River (Vegas Bajas), with a value of 1.75 g/kg. The areas with the highest average yields are the Montes de Toledo Mountain range (Spain) and the Guadiana River floodplain as it passes through Portugal (Guadiana International), with values of 5.20 ± 1.30 g/kg and 4.25 ± 1.07 , respectively (Table 2 and Supplementary Table S2a-d).

Table 2. Essential oil yield (g/Kg) of *L. stoechas* subsp. *luisieri* for each zone.

Zone	N	Yield (g/kg)		
		$\mu \pm \sigma$	min.	max.
Alconera	1	2.65		
Atlantic coast	7	2.67 ± 1.24	1.42	4.43
Guadiana International	2	4.25 ± 1.07	3.46	4.97
Montes de Toledo	10	5.20 ± 1.29	3.01	8.17
Rio Tejo	2	3.31 ± 1.40	2.34	4.30
Serra da Gardunha	5	2.23 ± 0.60	1.42	2.95
Serra de Alvelos	5	1.46 ± 0.45	1.08	2.05
Serra de Monchique	7	2.62 ± 1.79	1.34	5.94
Serra de São Mamede	2	2.30 ± 0.47	1.97	2.64
Sierra de San Pedro	5	2.78 ± 0.46	2.20	3.27
Sierra de Aracena y Picos de Aroche	9	2.87 ± 0.87	1.79	4.25
Sierra Morena de Sevilla	8	3.11 ± 0.66	2.10	4.14
Vegas Bajas	1	1.75	1.75	

Villuercas	2	3.42 ± 0.82	2.80	4.03
	66	3.08 ± 1.42	1.077	8.17

Note. $\mu \pm \sigma$: plus or minus one standard deviation from the mean, min.: minimum value; max.: maximum value.

2.2. Chemical Composition

The chemical composition of *L. stoechas* subsp. *luisieri* essential oil is highly variable. A total of 159 distinct compounds were detected in the 66 samples analyzed, and their presence exhibited significant variability (Supplementary Table S2a-d). The chemical composition of the essential oil is predominantly comprised of between 80–89, 90–99, and 70–79 different compounds, in 26, 21, and 12 samples, respectively. Only two samples (belonging to populations 17 and 18) have fewer than 70 different compounds in the essential oil, and four samples have more than 100 compounds (101 compounds in population 4, and 100 compounds in populations 47, 54, and 63). Furthermore, this essential oil is characterized by a high number of unknown compounds (20 in total), some of which may account for up to 3.60–3.71% of the total composition (e.g. unknown sesquiterpenol), as well as 4–20% of unidentified compounds. The percentage of compounds identified in each essential oil sample ranges from 80.99% (population 1) to 96.70% (population 19), and the average percentage of compounds identified in the essential oil of the 66 populations studied is $88.19 \pm 3.09\%$.

A total of 24 different chemical compounds were identified among the ten compounds with the highest percentage of presence in at least one essential oil sample. (Supplementary Table S2a-d and Table 3). Of these, *trans*- α -necroeryl acetate, *trans*- α -necrodol, and lavandulyl acetate are present in all samples among the 10 major compounds. The following compounds were detected in all essential oil samples analyzed: 1,8-cineole, camphor, linalool, 5-methylene-2,3,4,4-tetrame-2-cyclopentenone, *cis*- α -necrodol, α -pinene, arbozol, eudesma-3,7(11)-diene, and *cis*- α -necroeryl acetate; however, these compounds did not always appear among the top ten. It is noteworthy that the presence of fenchone, cymene isomer, viridiflorol, α -cadinol, copaborneol and several unidentified components (monoterpene, sesquiterpene or ester) may be observed among the ten compounds, although they may not be present in the essential oil.

Table 3. Major compounds are present in the essential oil of *L. stoechas* subsp. *luisieri*.

Components	$\mu \pm \sigma$	min.	max.	N	N1	N2	N3	N4	N5
<i>trans</i> - α -Necroeryl Acetate	20.68 ± 4.17	11.11	30.50	66	60	6	0	0	0
1,8-Cineole	7.79 ± 7.14	0.04	24.24	45	6	20	7	12	21
<i>trans</i> - α -Necrodol	8.66 ± 2.18	4.32	13.57	66	0	37	22	7	0
Lavandulyl Acetate	4.74 ± 0.91	2.95	6.96	66	0	1	22	43	0
Camphor	2.01 ± 1.55	0.16	7.61	30	0	1	3	26	36
α -Cadinol	2.24 ± 2.76	0.02	10.78	13	0	1	3	9	21
Fenchone	2.23 ± 2.91	0.01	12.92	17	0	0	6	11	31
Linalool	3.25 ± 1.19	1.09	6.58	62	0	0	3	59	3
5-Methylene-2,3,4,4-tetrame-2-Cyclopentenone	2.73 ± 0.70	1.26	4.13	58	0	0	0	58	8
Cymene Isomer	2.01 ± 0.79	0.09	3.29	50	0	0	0	50	15
Viridiflorol	1.96 ± 0.81	0.01	3.43	36	0	0	0	36	28
Unknown Sesquiterpenol	2.03 ± 0.86	0.14	3.71	30	0	0	0	30	24
<i>cis</i> - α -Necrodol	1.73 ± 0.27	1.10	2.29	29	0	0	0	29	37
α -Pinene	1.73 ± 0.98	0.41	5.36	20	0	0	0	20	46
Arbozol	1.69 ± 0.38	0.84	2.57	20	0	0	0	20	46
Unknown Sesquiterpenol	1.52 ± 0.70	0.09	2.89	16	0	0	0	16	46
Unknown Mw=153	1.56 ± 0.40	0.57	2.31	13	0	0	0	13	43
Unknown Ester	1.60 ± 0.22	1.11	2.10	9	0	0	0	9	47
Eudesma-3,7(11)-diene	1.07 ± 0.59	0.12	3.24	6	0	0	0	6	60
Unknown Mw=154	1.80 ± 0.26	1.24	2.08	6	0	0	0	6	4
<i>cis</i> - α -Necroeryl Acetate	1.38 ± 0.28	0.82	1.92	5	0	0	0	5	61

Unknown Sesquiterpenol	1.38 ± 0.90	0.66	3.60	3	0	0	0	3	7
Copaborneol	1.01 ± 0.37	0.04	1.78	1	0	0	0	1	59
Unknown Sesquiterpenol	0.80 ± 0.42	0.05	2.09	1	0	0	0	1	53

Note. $\mu \pm \sigma$: plus or minus one standard deviation from the mean, min.: minimum value; max.: maximum value, N: number of occurrences in the top ten, N1: number of times in first place, N2: number of times in second place, N3: number of times in third place, N4: number of times between fourth and tenth place, N5: number of times in chemical composition outside of top ten.

Trans- α -Necrodiyl acetate was identified as the predominant compound in 60 samples and the secondary compound in the remaining 6 samples analyzed, with an average percentage of $20.68 \pm 4.17\%$. 1,8-Cineole was the predominant compound in six samples and the second most abundant in twenty samples. The mean percentage of presence was found to be $7.79 \pm 7.14\%$, with a wide range of variation observed, from traces ($< 0.05\%$) to 24.24% . *Trans- α -necrodol* is the second most abundant compound, appearing in 37 samples, with an average percentage of presence of $8.66 \pm 2.18\%$.

As illustrated in Figure 4 and outlined in the supplementary data (Supplementary Table S3a-f), the distribution of the eight primary components in the essential oil of *L. stoechas* subsp. *luisieri* is depicted by geographical area. The populations studied in the Spanish mountain ranges of Alconera, Villuercas and Sierra Morena de Sevilla generally have an essential oil that is rich in *trans- α -necrodiyl acetate* and 1,8-cineole, with 1,8-cineole being the main component in some cases. In other areas, the percentage of *trans- α -necrodiyl acetate* generally exceeds that of 1,8-cineole. This is of particular significance in the populations studied in the Serra da Gardunha, Serra de Alvelos, Serra de São Mamede, and Sierra de San Pedro Mountain ranges, and in the alluvial zone population of Vegas Bajas, where 1,8-cineole has very low percentages of presence. The mean percentage of presence in these areas ranges from $0.62 \pm 0.81\%$ in Serra de São Mamede to $2.54 \pm 1.47\%$ in Serra da Gardunha.

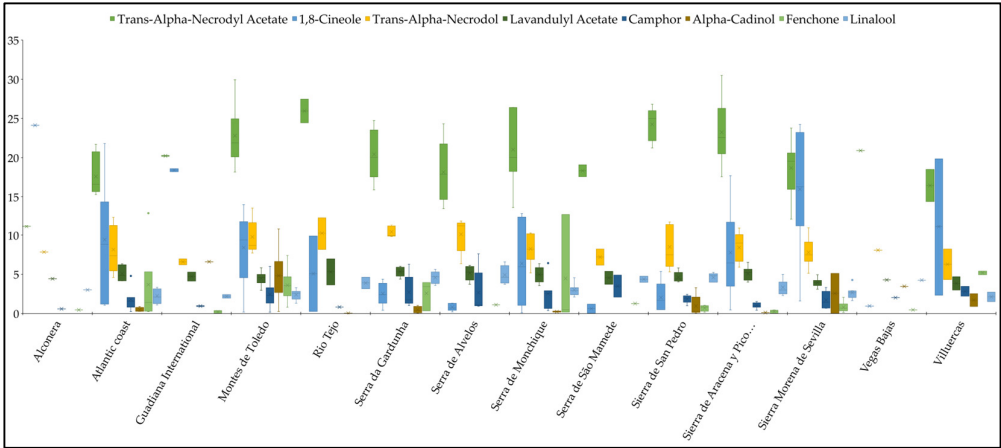


Figure 4. Box plot shows the distribution of the eight most prevalent compounds in the essential oil of *L. stoechas* subsp. *luisieri*, categorised according to zone.

In contrast, the *trans- α -necrodol* exhibited a certain degree of stability, with levels ranging from 5–12% across all areas. Of the other compounds, the presence of α -cadinol is worthy of note in the populations of the Montes de Toledo, Sierra Morena de Sevilla, Sierra de San Pedro y Villuercas, and Fenchone mountain ranges, principally in Serra de Monchique, where it can reach percentages of up to 12.76%.

2.3. Cluster Analysis

Hierarchical clustering was utilized to investigate the correlation between the chemical composition of the 66 populations of *L. stoechas* subsp. *luisieri* (Figure 5). The dendrogram obtained reveals the presence of two clusters, which are further subdivided into four subclusters at various hierarchical levels.

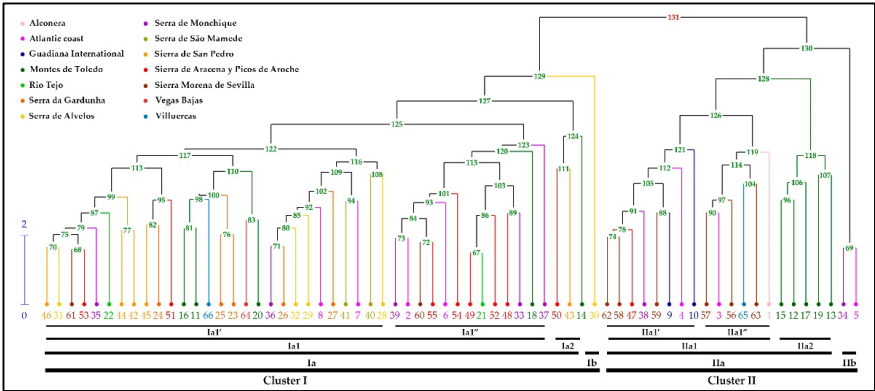


Figure 5. Hierarchical clustering of the percentage composition of the 66 samples of essential oils of *L. stoechas* subsp. *luisieri* analyzed.

The two primary groups (clusters I and II) of the dendrogram demonstrate a clear delineation of the essential oil samples according to their chemical composition (see Table 4). The primary distinctions between these groups pertain to the percentage of 1,8-cineole present, which ranges from 0.04% to 12.84% in cluster I and from 10.50% to 24.24% in cluster II, and the presence or absence of other minor compounds, including copaborneol, α -cadinol, and several unknown compounds.

Table 4. Range of values (in percentage) known compounds (which at least once are among the ten most abundant) in the essential oil of *L. stoechas* subsp. *luisieri*, grouped according to cluster analysis.

Components	Cluster I								Cluster II							
	Ia1'		Ia1''		Ia2		Ib	IIa1'		IIa1''		IIa2		IIb		
	min.	max.	min.	max.	min.	max.		min.	max.	min.	max.	min.	max.	min.	max.	
α -Pinene	0.41	3.62	0.74	2.12	1.81	2.40	4.77	0.93	1.96	1.01	3.23	1.92	5.36	1.45	1.50	
Cymene Isomer	0.09	3.29	0.17	3.16	0.26	1.37	1.91	1.85	2.76	1.80	2.36	a	2.31	2.04	2.06	
1,8-Cineole	0.04	5.63	5.38	12.84	1.35	7.56	1.33	12.42	18.60	19.84	24.24	10.50	14.01	12.86	13.06	
Fenchone	a	7.39	a	1.40	0.39	2.75	a	a	5.31	0.34	5.41	2.32	4.88	12.76	12.92	
Camphor	0.38	6.27	0.64	6.42	0.16	1.90	7.61	0.23	2.90	0.58	3.45	1.36	6.00	1.56	2.72	
Linalool	1.52	6.58	1.89	4.99	1.72	5.25	4.62	1.26	2.91	1.09	3.01	1.31	2.87	2.10	2.10	
<i>trans</i> - α -Necrodyll Acetate	15.30	26.42	17.54	28.15	26.83	30.50	13.48	16.59	20.46	11.11	19.48	18.17	26.04	13.64	15.68	
Lavandulyl Acetate	3.72	6.96	3.65	6.35	3.92	6.53	3.73	3.75	5.28	3.00	4.43	2.95	5.83	3.55	4.15	
<i>cis</i> - α -Necrodyll Acetate	0.91	1.80	1.09	1.81	1.21	1.92	0.84	0.95	1.67	0.84	1.38	1.10	1.72	0.82	1.16	
Arbozol	0.84	2.57	1.25	2.55	1.33	1.67		1.39	2.24	1.11	1.70	1.16	1.96	1.57	1.58	
5-Methylene-2,3,4,4-tetrame-2-Cyclopentenone	1.57	4.00	1.59	3.51	1.26	2.56	3.04	1.89	4.13	1.83	3.15	1.33	1.77	2.23	2.23	
<i>trans</i> - α -Necrodol	4.59	12.38	5.92	13.57	6.71	10.48	6.36	6.22	10.93	4.32	7.84	7.70	8.85	5.16	5.43	
<i>cis</i> - α -Necrodol	1.58	2.29	1.34	2.22	1.32	1.88	1.10	1.36	1.96	1.27	1.69	1.27	1.55	1.46	1.46	
Eudesma-3,7(11)-diene	0.20	2.29	0.12	2.69	0.14	0.99	3.24	0.53	1.39	0.40	1.31	0.40	2.01	1.42	1.76	
Viridiflorol	0.44	3.43	a	3.09	0.14	2.50	2.96	0.56	2.73	1.12	2.07	a	0.92	1.44	1.45	

Copaborneol	a	1.78	a	1.43	a	0.82	1.48	0.19	1.43	0.54	0.95	0.04	0.45	0.71	0.72
α-Cadinol	a	4.85	a	0.25	a	6.09	a	a	6.60	a	5.10	2.99	10.78	0.31	0.31

Note. min.: minimum value; max.: maximum value. Bold: dominant compounds for each cluster.

Cluster I is further subdivided into four subclusters, which are distributed across a range of hierarchical levels. The initial subdivision successfully isolates a sample (subcluster Ib) that is distinctly different from the others due to its elevated camphor content (7.61%), which is the highest percentage detected in all the samples analyzed in this study. Subcluster Ia is further subdivided into three groups (Ia1', Ia1'', and Ia2) that are primarily categorized based on the percentages of *trans*-α-necrodiyl acetate and 1,8-cineole present. However, there is overlap between the ranges of values due to the influence of other minor compounds that are not among the top ten (e.g. α-cadinol, camphor, fenchone).

Cluster II is further subdivided into two subclusters (IIa and IIb). Subcluster IIb is distinguished by its unique composition, with two samples of essential oil exhibiting a high concentration of fenchone, approximately 13% of the total composition. The remaining three groups contained within subcluster IIa are distinguished by a high percentage of 1,8-cineole (greater than 10%) and its relationship with the percentage of *trans*-α-necrodiyl acetate. In group IIa2, the range of 1,8-cineole is from 10.50% to 14.01%, and for *trans*-α-necrodiyl acetate, it is from 18.17% to 26.04%. In addition, α-cadinol is present in all samples, with a range of 2.99% to 10.78%. Group IIa is subdivided into two subgroups, designated IIa1' and IIa1''. Subgroup IIa1' consists of samples in which the percentage of *trans*-α-necrodiyl acetate (16.59–20.46%) exceeds that of 1,8-cineole (12.42–18.60%), while subgroup IIa1'' comprises samples in which the percentage of 1,8-cineole (19.84–24.24%) surpasses that of *trans*-α-necrodiyl acetate (11.11–19.48%).

Regarding the relationship between the chemical composition of the essential oil and the harvesting area, Figure 5 shows that there is a weak relationship between samples from the same population, except for those from Montes de Toledo, where five of the ten populations are included in subgroup IIa2 (cluster II).

3. Discussion

The populations of *L. stoechas* subsp. *luisieri* collected are predominantly located in mountainous areas (54 populations), with sporadic occurrences in more or less flat areas associated with the course of major rivers (Guadiana and Tagus rivers) (5 populations) and in areas of the Portuguese Atlantic coast (7 populations). These populations, in conjunction with those from other studies [51, 53, 57, 58, 63, 73], indicate a clear preference of the species for mountainous areas in the interior of the Iberian Peninsula (Figure 6a).

The essential oil yield obtained in this study (Table 1) ranges from 1.08 g/kg (0.11% w/w) to 8.14 g/kg (0.81% w/w), with an average yield of 3.44 ± 0.87 g/kg (0.34 ± 0.01%). In the harvests from Almadén de la Plata, located in the north of Seville and included in the study area of Sierra Morena de Sevilla, yields of 0.20% and 0.30% w/w fresh weight [53,76] were obtained. These values are comparable to, or slightly lower than, those obtained in the eight populations analyzed in the Sierra Morena de Sevilla area (average yield: 0.31 ± 0.07% w/w. As demonstrated in the various harvests conducted in central Portugal (Penamacor, Vila Velha de Ródão, Mata, and Casal da Fraga), the yields ranged from 0.07% to 0.38% w/w [57], which is analogous to the yields obtained for the mountain ranges of central Portugal studied, Serra da Gardunha and Serra de Alvelos, where the minimum and maximum values recorded were 0.11% and 0.30%, respectively (Figure 6b). Other studies indicate higher yields of up to 2.80%; however, these units are expressed in terms of dry matter volume/weight (% v/w) [51,54], and are therefore not comparable with the data obtained in our study.

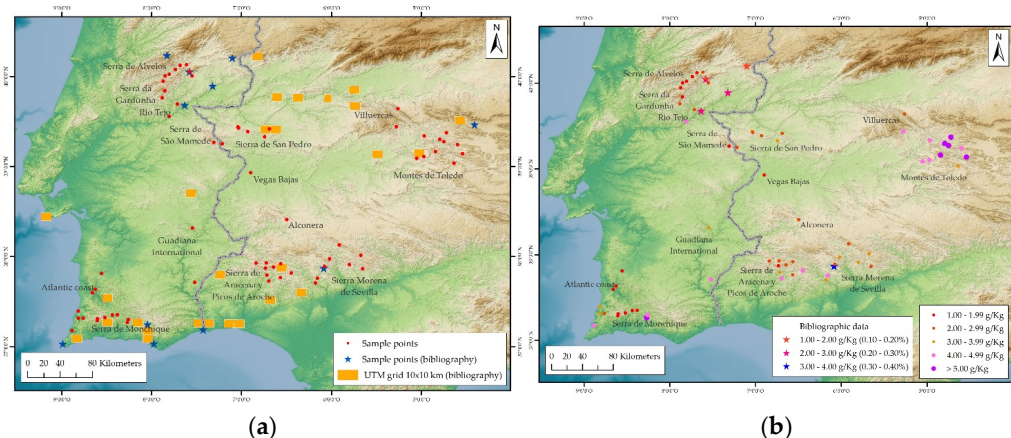


Figure 6. (a) Distribution map of sampling points for the analysis of the essential oil from *L. stoechas* subsp. *luisieri*. (b) Distribution map of essential oil yield data (stars – bibliography references and points – sampling points in this study).

The chemical composition of the essential oil of *L. stoechas* subsp. *luisieri* is distinguished by the presence of necrodol derivatives, which are absent in other lavender species. Preliminary studies undertaken by García-Vallejo [51,52] utilizing 37 specimens from 27 locations have indicated that *trans*- α -necrodyl acetate (6.30–28.80%) is the predominant compound in 24 essential oil samples, while 1,8-cineole (0.10–43.20%) is the primary compound in the remaining 13 samples. Conversely, camphor (0.40–6.40%) and fenchone (0.00–5.40%), which are major compounds found in other morphologically similar lavender species (*L. stoechas* subsp. *stoechas*, *L. pedunculata*, and *L. viridis*), are present in low concentrations. Subsequently, the characteristic profile of the essential oil of *L. stoechas* subsp. *luisieri* was confirmed in samples collected in Almadén de la Plata (in the north of Seville) [53,76], and novel unidentified compounds began to appear, such as: The following compounds have been identified: 1,1,2,3-tetramethyl-4-hydroxymethyl-2-cyclopentene, 2,3,4,5-tetramethyl-2-cyclopenten-1-one, and 2,3,5,5-tetramethyl-4-methylene-2-cyclopenten-1-one (= 5-Methylene-2,3,4,4-tetramethylcyclopent-2-enone).

The major compound in essential oil samples obtained from leaves collected in populations in northern Seville and southern Toledo (with values reaching up to 38.30%) is 2,3,5,5-tetramethyl-4-methylene-2-cyclopenten-1-one [73]. The study also highlights the presence of camphor (with percentages of 51.80% and 53.70% for essential oil samples obtained from flowers and leaves of the same individual), fenchone (with values reaching 22.0% in some samples), 1,8-cineole (with a range of presence between 0.40% and 20.60%, being the major component in three samples), and viridiflorol (with 12.10% being the major compound in a sample of essential oil obtained from flowers). In contrast, necrodol derivatives exhibit remarkably low percentages, particularly *trans*- α -necrodyl acetate, which displays an occurrence percentage of 1.10–3.70% [73].

In the course of further research conducted on samples collected in various areas of Portugal, 1,8-cineole was found to be the main component (25.70–34.30%), followed by *trans*- α -necrodyl acetate (11.30–17.50%) in three areas of the Alentejo coast. Montenegro (Faro), Salir (Loulé), and Vila Real de Santo António [63]. Conversely, in four towns in the south-east of the Beira Interior region (Vila Velha de Ródão, Mata, Penamacor, and Casal da Fraga), *trans*- α -necrodyl acetate was identified as the predominant compound in essential oil samples from leaves (25.23–49.95%) and flowers (9.40–26.99%) in three localities (Penamacor, Mata, and Casal da Fraga). Meanwhile, 1,8-cineole was detected in trace amounts (0.00–4.95%). In the sample from Vila Velha de Ródão, fenchone was identified as the predominant compound, accounting for 19.16% and 36.77% of the essential oil derived from leaves and flowers, respectively. *Trans*- α -necrodyl acetate was the second most abundant compound, contributing 19.38% and 10.27% to the essential oil extracted from leaves and flowers, respectively [57]. The Penamacor population is subsequently used in other studies, in which, together with other properties of the essential oil, the variation in chemical composition between

leaves and flowers during the different phenological stages of the plant is evaluated. The research findings indicate that the predominant compound in the sample is *trans*- α -necrodiyl acetate (1.80–26.90 per cent), with 1,8-cineole present in substantial concentrations in the leaves (13.90–16.40%), and camphor reaching 42.90% in an essential oil sample obtained from the leaves at the end of the flowering period [54,59]. The most recent populations the focus of study in Portugal are Piodão (in the central region) and Cabo São Vicente (in the Algarve), which exhibit markedly divergent chemical compositions. In Piodão, the essential oil is characterized by a high concentration of *trans*- α -necrodiyl acetate (16.00–17.40%), lavandulyl acetate (6.10–7.60%), and *trans*- α -necrodol (7.10–8.40%), while in Cabo São Vicente, the predominant compounds are 1,8-cineole (33.90%), fenchone (18.40%), and *trans*- α -necrodol (4.50%), with *trans*- α -necrodiyl acetate exhibiting a significantly lower concentration (3.20%) [58,68,69,72].

Finally, the final population studied is in Montes de Toledo (Pueblo Nuevo de Bullarque), and its essential oil has a chemical composition that is rich in camphor (74.40%), with no traces of the various necrodol derivatives and 1,8-cineole [74].

The observed percentage variation in the compounds that constitute the essential oil of *L. stoechas* subsp. *luisieri* is largely consistent with the results obtained in this study. *Trans*- α -necrodiyl acetate is typically the most prevalent compound, with concentrations reaching up to 48.22–49.45% in essential oils derived from fresh leaves collected in Mata and Penamacor, respectively [57]. These values exceed those observed in this study, where the maximum concentration of *trans*- α -necrodiyl acetate is 30.50% in an essential oil derived from fresh whole plants from a population in Sierra de Aracena and Picos de Aroche. This discrepancy in values can be attributed to the nature of the sample type employed, given that *trans*- α -necrodiyl acetate is found in higher concentrations in leaves than in flowers [57].

1,8-Cineole has been found to be present in a wide range of proportions in the essential oil of *L. stoechas* subsp. *luisieri*, ranging from very low percentages to as high as 43.20% in a population from Aznalcollar [51] and 24.24% for a population studied in this work from Sierra Morena de Sevilla. In the present study, the geographical distribution of the samples examined did not appear to be related to the percentage of 1,8-cineole present in the essential oil of *L. stoechas* subsp. *luisieri*. Samples exhibiting high or low percentages of 1,8-cineole may be observed in the same area, as has been previously reported in other studies [51,73].

Other compounds are characteristic of the essential oil of *L. stoechas* subsp. *luisieri*, including camphor and fenchone. The prevalence of both is low (i.e., less than 8.00%), and in some cases, they may even be absent [51,52]. However, in this study, fenchone exceeds 10.00% in two populations (ID numbers 5 and 34, from the Atlantic coast and Serra de Monchique areas, respectively) that appear isolated in the cluster analysis (Figure 5). In addition, other studies have identified samples exhibiting even higher values, including two samples from southern Toledo with values reaching up to 22.00% [73], a sample from Cabo São Vicente (18.20%), and a sample from Vila Velha de Ródão with values up to 36.77% [57]. Morphological analysis (unpublished data) of the populations collected in this study indicates the existence of several individuals with morphologies intermediate between *L. stoechas* subsp. *luisieri* and *L. pedunculata* (hybrids), which may be responsible for the increase in fenchone in the essential oil. A comparable phenomenon has been observed in the sample from Vila Vela de Ródão. The three samples under scrutiny have been found to contain necrodol derivatives, with concentrations ranging from 10.00 to 25.00%. Conversely, in samples from southern Toledo and Cape São Vicente, the presence of necrodol derivatives is found in minimal percentages (< 2.00%), thus prompting uncertainty regarding the authenticity of these samples as *L. stoechas* subsp. *luisieri*.

A comprehensive analysis of the extant literature pertaining to the essential oil of *L. stoechas* subsp. *luisieri* reveals the presence of two samples that are characterized by a high concentration of camphor: one from southern Toledo (with 51.80–53.70%) [73], and one from Pueblo Nuevo de Bullarque (with 74.40%) [74]. In both cases, the necrodol derivatives characteristic of the essential oil of *L. stoechas* subsp. *luisieri* are not present, so it is possible that these samples correspond to another species of *Lavandula*, possibly *L. pedunculata* subsp. *sampaioana* (Rozeira) Franco, a species

from which samples of essential oil rich in camphor (up to 84.40%) have been found in various locations in the Montes de Toledo and Sierra Morena Mountain ranges [51].

4. Materials and Methods

4.1. Collection of Plant Materials

In spring 2024 (April–May), 66 populations of *L. stoechas* subsp. *luisieri* were collected, distributed throughout much of the natural distribution area (Supplementary Table S1).

In each population (sample), 20 individuals were selected that were representative of the morphological diversity of the species existing in the population. For each individual (subsample), a fraction of the aerial part (stems, leaves, and flowers) was collected individually and stored in raffia bags at room temperature (20–25 °C) for 12–24 hours. Next, in the laboratory, each subsample was cut into small pieces, and 25 g of sample were selected at random. Finally, the 20 subsamples or individuals were mixed to obtain the final sample from each population (25 × 20 = 500 g fresh weight). All samples were frozen at -40 °C until distillation.

4.2. Essential Oil Extraction

For the extraction of essential oil, each sample (500 g of fresh, frozen material) was placed in a 5000 ml spherical flask and mixed with distilled water (temperature: 20–25°C) in a proportion of 20% (w/w). The samples were left to thaw for 10–15 minutes and then distilled.

The distillation method used was hydrodistillation in a Clevenger-type apparatus for 2 hours. The extracted essential oil was stored in an amber vial at 4 °C.

4.3. Chemical Characterization of Essential Oils

The chemical analysis of the essential oils was carried out using a combination of two gas chromatography techniques (GC-FID + GC-MS), chemical compounds were identified by CG-MS and quantified by CG-FID. The analysis was performed on Agilent 8890 GC (Agilent Technologies, Inc., Santa Clara, CA, USA) paired with the Agilent 5977B MSD (Mass Selective Detector). Polar column DB-WAX UI (60 m long, 0.25 mm diameter and 0.5 µm film thicknesses) was employed using Helium carrier gas at constant flow of 2 mL/min. Apolar column HP-5MS UI (60 m long, 0.25 mm diameter and 0.25 µm film thicknesses) was employed using Helium carrier gas at constant flow of 1 mL/min. The column temperature started at 50°C and increased to 240°C (polar column) and 285°C (apolar column).

4.4. Statistical Analysis

Statistical parameters were calculated using SPSS statistics version 23.0 (IBM, Nueva York, United States) and for the graphical representations of data, MS Excel version 2507 (Microsoft 365 for business, Microsoft, United States).

The percentage composition of the isolated essential oils was used to determine the relationship between the different samples by cluster analysis using the DARwin, version 6.0.021 (CIRAD-BIOS UMR AGAP, Montpellier, France) [77,78]. For cluster analysis, the Euclidean distance metric was selected as a measure of similarity among samples and the Unweighted Pair Group Method using Arithmetic Averages (UPGMA) was used for cluster definition.

5. Conclusions

The essential oil of *L. stoechas* subsp. *luisieri* is clearly distinguishable from any other lavender essential oil due to the presence of necrodol derivatives. These compounds can be found in four forms: *trans*- α -necrodiyl acetate, *cis*- α -necrodiyl acetate, *trans*- α -necrodol, and *cis*- α -necrodol. *Trans*- α -necrodiyl acetate is typically the most prevalent compound in the essential oil.

The presence of 1,8-Cineole has been detected in percentages ranging from trace amounts (< 0.10%) to over 25.00%, indicating a considerable degree of variability. This behavior is instrumental

in differentiating two chemotypes in the essential oil of *L. stoechas* subsp. *luisieri*: one chemotype is characterized by a high concentration of *trans*- α -necroeryl acetate and a low representation of 1,8-cineole (less than 6%), while the other chemotype is distinguished by a high concentration of both compounds, with *trans*- α -necroeryl acetate and 1,8-cineole at percentages between 15.00–25.00%, rarely higher, and with 1,8-cineole as the predominant component in some cases.

A further distinguishing characteristic of *L. stoechas* subsp. *luisieri* essential oil is the comparatively low representation of camphor and fenchone, with percentages of presence of less than 8.00%. This characteristic differentiates the essential oil of this species from other species in the *stoechas* section of the *Lavandula* genus (*L. stoechas* subsp. *stoechas*, *L. pedunculata*, *L. viridis*, and their hybrids).

Supplementary Materials: The following supporting information can be downloaded at the website of this paper posted on Preprints.org, Table S1: Checklist of collected populations of *L. stoechas* subsp. *luisieri*; Table S2a: Chemical composition of *L. stoechas* subsp. *luisieri* (Zone: 1. Alconera, 2. Atlantic coast, 3. Guadiana International, 4. Montes de Toledo); Table S2b. Chemical composition of *L. stoechas* subsp. *luisieri* (Zone: 5. Rio Tejo, 6. Serra da Gardunha, 7. Serra de Alvelos, 8. Serra de Monchique); Table S2c. Chemical composition of *L. stoechas* subsp. *luisieri* (Zone: 9. Serra de São Mamede, 10. Sierra de San Pedro, 11. Sierra de Aracena y Picos de Aroche); Table S2d. Chemical composition of *L. stoechas* subsp. *luisieri* (Zone: 12. Sierra Morena de Sevilla, 13. Vegas Bajas, 14. Villuercas); Table S3a. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 1. Alconera, 2. Atlantic coast); Table S3b. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 3. Guadiana International, 4. Montes de Toledo); Table S3c. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 5. Rio Tejo, 6. Serra da Gardunha); Table S3d. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 7. Serra de Alvelos, 8. Serra de Monchique); Table S3e. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 9. Serra de São Mamede, 10. Sierra de San Pedro); Table S3f. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 11. Sierra de Aracena y Picos de Aroche, 12. Sierra Morena de Sevilla); Table S3f. Main components of *L. stoechas* subsp. *luisieri* essential oil (Zone: 13. Vegas Bajas, 14. Villuercas).

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