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### A new species of Micromeria (Lamiaceae) from Köyceğiz (Muğla, southwest of Turkey)

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**Abstract:** A new species, *Micromeria aybalae* H.Duman & Dirmenci (Lamiaceae), *Micromeria* Benth. sect. *Micromeria*, is described from Muğla Province in southwestern Turkey. A description, taxonomic note, distribution map, habitat, and nrDNA ITS and cpDNA *trnL*-F based phylogeny are presented. The differences between the new species and its allies, *Micromeria cremnophila* Boiss. & Heldr. s.l. and *M. hispida* Benth., are discussed, and an identification key is provided for the Turkish *Micromeria*.

Key words: Labiatae, Muğla, phylogeny, Sandras Mountain, Micromeria, Turkey

#### 1. Introduction

The genus *Micromeria* Benth. belongs to the family Lamiaceae, subfamily Nepetoideae, tribe Mentheae, and subtribe Menthinae (Harley et al., 2004). *Micromeria* Benth. s.str. is currently circumscribed as monophyletic and is represented by nearly 70 species in the world (Govaerts, 1999; Brauchler et al., 2008b; Puppo and Meimberg, 2015a, 2015b). The genus is distributed from the Macaronesian-Mediterranean region to southern Africa, India, and China (Bräuchler et al., 2008b).

Micromeria s.l., a taxonomically complex genus, was divided into six sections as sect. Micromeria, sect. Pseudomelissa Benth., sect. Cymularia Boiss., sect. Pinoelentia P.Perez, sect. Xenopoma (Willd.) Benth., and sect. Hesperothymus Benth. by Morales (1993). In recent studies some changes were made to the taxonomic border of the genus Micromeria, as listed below. The species of sections Xenopoma and Hesperothymus pro parte distributed in southeastern Africa were transferred to the genus Killickia Bräuchler, Heubl & Doroszenko by Bräuchler et al. (2008a), and to Clinopodium by Cantino and Wagstaff (1998), Govaerts (1999), and Harley and Granda (2000); the species of section Pseudomelissa were transferred to Clinopodium by Bräuchler et al. (2005) and by Ryding (2006).

In Flora of Turkey (Davis, 1982), Micromeria s.l. is represented by 14 species (22 taxa), 12 of which are endemic and grouped into three sections: sect. Micromeria (7 species, 12 taxa), sect. Pseudomelissa (6 species, 9 taxa), and sect. Cymularia (1 species). After the latest taxonomic

evaluations performed by Bräuchler et al. (2005), the species of section *Pseudomelissa* were transferred to *Clinopodium*. Finally, *Micromeria* is represented in Turkey by 8 species and 13 taxa, 8 of which are endemic (Arabacı et al., 2010; Dirmenci, 2012).

In 2015, the first author collected some unusual specimens belonging to *Micromeria* sect. *Micromeria* in southwestern Turkey (Figure 1). These specimens were clearly different from other known *Micromeria* species growing in Turkey, Greece, and Cyprus. After taxonomic studies it was concluded that the collected specimens represented an undescribed species close to *Micromeria cremnophila* Boiss. & Heldr. and *M. hispida* Boiss. & Heldr. ex Benth. It is being described as a new species here.

The main purposes of this study are: 1) to identify a new species, 2) to discuss the morphological and micromorphological differences between the new species and its allies, 3) to discuss the phylogenetic relationships of the new species with other *Micromeria* species, 4) to determine the phylogenetic position of the other Turkish *Micromeria* species, 5) to evaluate the condition of Turkish *Micromeria* species according to recent studies, and 6) to make a new identification key consisting of the new species and other species in Turkey.

#### 2. Materials and methods

#### 2.1. Plant material

The plant materials of the new species were collected by the first author during the GEF-5 project, carried out in the Köyceğiz district of Muğla Province in November

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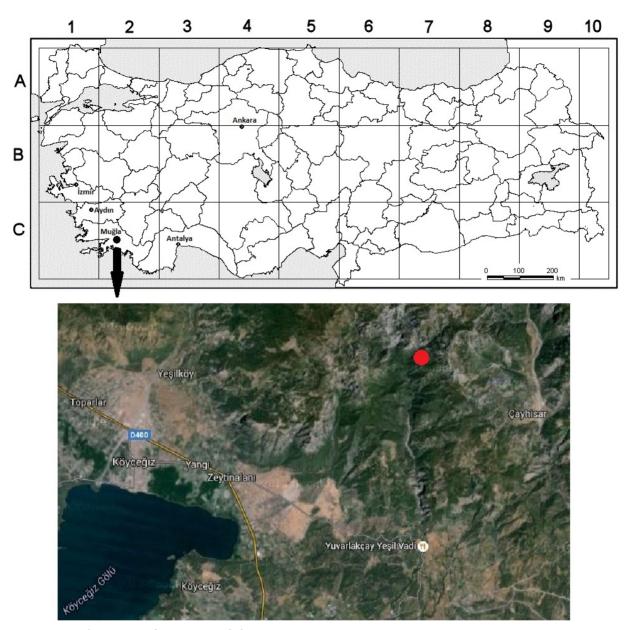


Figure 1. Distribution map of Micromeria aybalae.

2015 (Figure 1). These specimens were identified using the relevant literature (Ball and Getlifle, 1972; Baden, 1990; Davis, 1982; http://www.cretanflora.com) and compared to other known *Micromeria* species (Appendix 1) using specimens found in the following herbaria (acronyms according to Index Herbarium 2016): ANK, B, BM, E, EGE, G, GAZI, GOET, HUB, ISTF, ISTE, K, LE, MA, W, and WU (http://sweetgum.nybg.org/science/ih/).

#### 2.2. Micromorphological studies

Trichome morphology of *Micromeria cremnophila*, *M. aybalae*, and *M. hispida* was studied using tabletop scanning electron microscopy (SEM) in the Basic Sciences

Research and Applied Center of Balıkesir University. Average samples were chosen for micromorphology of stems and leaves. Trichomes were investigated and photographed using a NeoScope JCM. They were fixed on aluminum stubs using double-sided adhesive. The SEM micrographs were taken with a NeoScope JCM-5000 at an accelerating voltage of 10 kV.

#### 2.3. Genomic DNA isolation

About 0.2 g of dried leaf tissue was used for DNA isolation. Some of the specimens were collected from their natural distribution areas. These fresh leaves were dried in silica gel. In addition, some leaves from some species were taken

from herbarium materials. Samples dried in silica gel and herbarium materials were squashed using a mortar and pestle with liquid nitrogen. The GeneJET Plant Genomic DNA Purification Mini Kit (Thermo Scientific, Lithuania) was used for DNA extraction following the manufacturer's protocol. Information on the investigated samples, GenBank numbers, and voucher numbers are provided in Appendix 2 (samples used in our molecular studies are marked with an asterisk). The quality of DNA extraction was confirmed by electrophoresis on a 1% agarose gel.

#### 2.4. PCR amplification

The internal transcribed spacer (ITS) region of the nuclear ribosomal DNA (nrDNA) sequences and the trnL-F region of the chloroplast DNA (cpDNA) were used for molecular analysis of the Micromeria species. Polymerase chain reaction (PCR) amplifications of the ITS nrDNA were performed using ITS5A (5'-CCTTATCATTTAGAGGAAGGAG-3') (Stanford et al., 2000) and ITS4 (5'-TCCTCCGCTTATTGATATGC-3') (White et al., 1990) primers. The trnL-trnF (trnLeutrnPhe) cpDNA amplifications were performed using (forward) CGAAATCGGTAGACGCTACG trnL-c *trn*L-f (reverse) (Taberlet et al., 1991) and ATTTGAACTGGTGACACGAG (Taberlet et al., 1991) primers. The total volume of each PCR tube was 25 µL, comprising 2.5 µL of CoralLoad PCR Buffer (QIAGEN, Germany), 3.0 µL of Q-solution (QIAGEN), 0.25 µL of Taq DNA polymerase (QIAGEN), 1.25 μL of 5 μM ITS5A  $(1.25 \mu L \text{ of } 5 \mu M \text{ } trnL\text{-c}), 1.0 \mu L \text{ of } 5 \mu M \text{ } ITS4 (1.25 \mu L \text{ of } 1.25 \mu L \text{ of$ 5 μM trnL-f) (Sigma-Aldrich, Germany), and 0.4 μL of 20 µM dNTP solution (QIAGEN), and it was autoclaved in deionized water. During the PCR amplification, a thermal cycler machine (Techne-Prime, USA) was used for the routine amplification. Initial denaturation was performed for 5 min at 95 °C. The following 35 cycles were carried out: 1 min at 94 °C for denaturation, 1 min at 51 °C (59 °C for tnrL-F) for annealing, and 2 min at 72 °C for extension. A final extension cycle (5 min at 72 °C) followed. PCR products were checked by electrophoresis in 1% agarose gel.

# 2.5. Data analysis and editing of the ITS nrDNA and trnL-F cpDNA data

PCR products were sent to Genoks (Gene Research and Biotechnology Company, Ankara, Turkey) to be sequenced. The sequenced DNA was edited using Sequencher, version 4.9 (Gene Code Corporation, Ann Arbor, MI, USA). Clustal W (Larkin et al., 2007) was used for alignment. Some nucleotides from the 5' end of ITS1 (rpl32) and the 3' end of ITS2 (trnL-trnF) were cut to avoid doubtful base callings and redundant gaps. Finally, sequences of 660–672 nucleotides in length were produced from nrITS regions and 850–870 nucleotides in length from *trnL-trn*F cpDNA regions for the taxa studied. Polymorphic sequence regions

of the *Micromeria* species were determined. A maximum parsimony tree was obtained using PAUP\* (Swofford, 2002).

#### 3. Results

**Micromeria aybalae** H.Duman & Dirmenci **sp. nov.** (Figures 2A–2E, 3A–3C, 4A–4I)

**Type:** Turkey. C2 Muğla: Köyceğiz, Sandras Mountain, west of Çayhisar village, calcareous rocks, 980–1000 m a.s.l., 4.11.2015. *H.Duman* 10395 (holotype GAZI!, isotypes ANK!, HUB!, ISTE!).

#### 3.1. Diagnosis

Micromeria aybalae is similar to Micromeria cremnophila s.l. and M. hispida. It differs from M. cremnophila s.l. in the following aspects: stems prostrate to procumbent, slender, fragile (not ascending to erect and woody rootstock) and pilose with minutely glandular papillate (not puberulent, short hispid, or scabrid); leaves revolute only at apex (not revolute); calvx 4-4.5 mm and pilose with minutely glandular papillate (not 2-3 mm and puberulent to scabrid-pubescent); teeth 1.25-1.75 mm and 1/3-1/2 of tube (not 0.3-0.5 mm and 1/5-1/4 of tube), ciliate (not ciliate); corolla 5-6 mm (not 3-4 mm), pinkish-white (not purplish-pink). It differs from *M. hispida* in its herbaceous appearance: herbs only woody at base; stems prostrate to procumbent, branched, pilose with minutely glandular papillate (not dwarf shrub, procumbent to ascending, often branched, patent-pubescent to hispid); verticillasters 2-flowered (not 1-6-flowered); calyx pilose with minutely glandular papillate (not hispid); teeth 1/3-1/2 of tube (not 3/4 of tube); corolla pinkish-white (not purple).

#### 3.2. Description

Perennial herbs. Stems numerous, prostrate to procumbent, 5-15 cm long, slender, brittle, pilose with minutely glandular papillate thoroughly, sometimes ±hispid at the base. Leaves ovate, 4-8 × 2.5-4.5 mm, slightly revolute towards the apex, pilose with minutely glandular papillate on both surfaces, subsessile, petioles up to 0.5 mm long, attenuate to rounded at base, acute to obtuse at apex, veins invisible. Inflorescence lax, 2-flowered, to  $6 \times 2$  cm, raceme or paniculate; cymules distinctly pedunculate. Bracts similar to leaves,  $2-5 \times 1.5-3$  mm, ovate, shortly petiolate to subsessile; bracteoles 2, 1-1.5 mm long, linear, shorter than pedicel, pedicel 3-5 mm long. Calyx tubular, 4-4.5 mm long, subbilabiate, pilose with minutely glandular papillate, veins 15, visible, pilose at throat; teeth linear-lanceolate, 1.25-1.75 mm long, 1/3-1/2 of the tube, ±equal or lower ones slightly longer than upper ones, ciliate. Corolla 5-6 mm long, pinkish-white, exserted from corolla, minutely glandular papillate; upper lip emarginate, lower lip divided into 3 lobes; hairy inside of upper lip and mouth. Style slightly exserted from corolla, unequally 2-lobed. Stamens 4, included in corolla, filaments hairy at

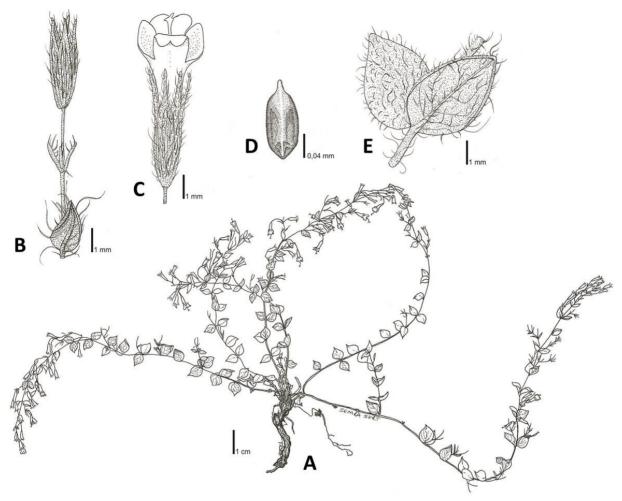


Figure 2. Micromeria aybalae: A- habit, B- calyx, C- flower, D- nutlet, E- leaves.



Figure 3. Micromeria aybalae: A, B- flowers and leaves, C- habit (photos from Dr Hayri Duman).

base. Nutlets oblong, c. 1.5 mm long, acuminate, mucro 0.3–0.4 mm, brown.

# 3.3. Habitat, geographical distribution, and conservation status

The species was found as growing only on Sandras Mountain, one of the highest mountains in southwestern Turkey, located in the northeastern part of the Köyceğiz

district of Muğla Province (Figure 1). The bedrock of Sandras Mountain mainly comprises serpentine and limestone (Doğan, 2011). This geomorphological characteristic is responsible for the high number of endemic species. Sandras Mountain presents about 118 endemic plants, 25 of which (including *M. aybalae*) grow only there (Pirhan et al., 2014), and 4 of which are under

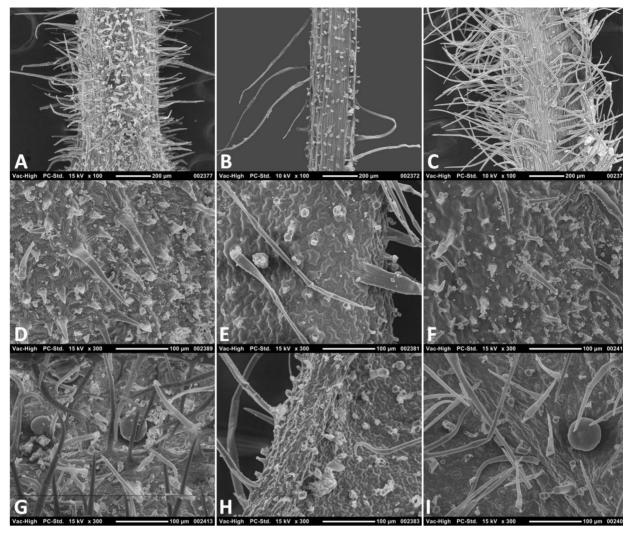


Figure 4. SEM photographs of Micromeria cremnophila (A, D, G), M. aybalae (B, E, H), and M. hispida (C, F, I).

threat on the European and global scale (Özhatay et al., 2003). Phytogeographically, Sandras Mountain is under the influence of the Eastern Mediterranean.

Micromeria aybalae was collected at an altitude of 1000 m in the northwest of Çayhisar village, on the limestone bedrock of black pine forest clearances. The species growing in the same area together with the new species are Phagnalon rupestre (L.) DC subsp. graecum Batt (Asteraceae), Inula heterolepis Boiss. (Asteraceae), Micromeria myrtifolia Boiss. & Hohen. (Lamiaceae), Teucrium *montbretii* Benth. subsp. pamphylicum P.H.Davis (Lamiaceae), and Ptilostemon chamaepeuce (L.) Less. (Asteraceae). Micromeria aybalae is known only from the type locality. The new species is endemic and an element of the East Mediterranean. The distribution area of the species is less than 10 km<sup>2</sup>. Its population in the type locality is considerably poor. The number of individuals is about 200. Moreover, its habitat is subject to rain erosion. Because the region is too hot, forest fires are often observed. As a result, the species' habitat may be destroyed, and its present distribution area might get narrower in the future. It should, therefore, be regarded as Critically Endangered (CR) under the B2abi-v, C1-2ai-ii, D criteria (IUCN, 2014).

#### 3.4. Etymology

*Micromeria aybalae* is a name given in dedication to the first author's daughter.

#### 4. Discussion

According to the latest studies, the genus *Micromeria* is represented by 9 species in Turkey (Bräuchler et al., 2008a; Arabacı et al., 2010; Dirmenci, 2012). The identification key for all the species occurring in Turkey is provided below.

#### Key to the species of *Micromeria* present in Turkey

- Annual herb, corolla resupinate (Sect. *Cymularia*) .....

   M. cymuligera

   Subshrub or perennial herbs, corolla not resupinate

- **3**. Flowers in lax verticillasters, peduncles and pedicels conspicuous ...... **5**.

- - 6. Calyx 2-3 mm; corolla 3-4 mm ...... M. cremnophila

- **8**. Calyx subbilabiate to 1/5, scabrid-pubescent; teeth lanceolate, erect to subpatent (NE of Turkey) ... **M. elliptica**
- 8. Calyx subbilabiate to 1/3–2/5, shortly pubescence to hispidulous; teeth subulate, curved (W and S of Turkey) .....

#### ...... M. graeca subsp. graeca

Micromeria aybalae is geographically isolated and morphologically clearly different from other Micromeria species growing in Turkey, Greece, and Cyprus. On the other hand, Micromeria aybalae has some morphological similarities with Micromeria cremnophila s.l. and M. hispida in sect. Micromeria (Figures 2-4). As seen as Figure 4, it can be easily distinguished from allied species morphologically and micromorphologically. It differs from M. cremnophila s.l. in the following aspects: stems pilose with minutely glandular papillate (not puberulent, short hispid, or scabrid) (Figures 4A and 4B), leaves pilose on both surfaces (not scabridulous and hispidulous) (Figures 4D, 4E, 4G, and 4H), calyx 4-4.5 mm and pilose with minutely glandular papillate (not 2-3 mm and puberulent to scabrid-pubescent), corolla 5-6 mm (not 3-4 mm) (Figures 2A-2E and Figures 3A-3C). It differs from M. hispida in appearance; herbs only woody at base, stems prostrate to procumbent, branched and pilose with minutely glandular papillate (not dwarf shrub, procumbent to ascending, often branched and patent-pubescent to hispid) (Figures 4B and 4C), leaves pilose on both surfaces (not patent pubescent to pilose) (Figures 4E, 4F, 4H, and 4I), verticillasters 2-flowered (not 1–6-flowered), calyx pilose with minutely glandular papillate (not hispid); teeth 1/3–1/2 of tube (not 3/4 of tube), and corolla pinkish-white (not purple) (Figures 2A–2E and Figures 3A–3C). Detailed differences between *Micromeria aybalae* and related species are presented in the diagnosis and in the key.

*Micromeria aybalae* is close to *M. chionistrae* Meikle, which is endemic to Cyprus, and *M. microphylla* (d'Urv.) Benth., a widespread species from the Mediterranean basin. It is distinguished from *M. chionistrae* by its perennial herbs (not subshrub and cushion-forming), verticillasters 2-flowered (not 3–8-flowered), calyx 4–4.5 mm (not 2.5–3 mm), and corolla 5–6 mm (not 3.5–4 mm). It can be distinguished from *M. microphylla* by its perennial herbs (not subshrub), verticillasters 2-flowered (not 1–7-flowered), calyx 4–4.5 mm (not 2–2.5 mm), corolla 5–6 mm (not ca. 4 mm).

According to our molecular and GenBank data results, and as seen in Figures 5 and 6, the genus Micromeria s.str. is a monophyletic genus when all the Micromeria clades are considered regarding the nrITS data (Figure 5) and the cptrnL-F data (Figure 6). On the other hand, in classical sectional classification, some of the section Pineolentia members are together with section Micromeria members (from the Canary Islands). This can be easily seen from the cptrnL-F phylogenetic tree (Figure 6). According to the data of Bräuchler et al. (2010), section Pineolentia and section Micromeria have the same clade, and this shows that they have a common ancestor. The data of Puppo et al. (2014, 2015) contributed to sections Pineolentia and Micromeria. M. pineolens shares the same clade with the Canary Islands section Micromeria specimens. Likewise, section Cymularia and Madagascarian Micromeria species do not belong to the section Micromeria clade according to Bräuchler et al. (2010). M. cymuligera (from section Cymularia) is closer to the Madagascarian group. This species (M. cymuligera) is known to grow only in Turkey, and the collected specimens are not as sufficient as mentioned before by Bräuchler et al. (2010). Our phylogenetic data and results confirm the results of Bräuchler et al. (2005, 2010).

In Flora of Turkey, the genus Micromeria is divided into three sections: Cymularia, Pseudomelissa, and Micromeria. Bräuchler et al. (2006) transferred the section Pseudomelissa to Clinopodium s.l. In this study, three Turkish species (M. mollis, M. carica, and M. dolichodonta) belonging to sect. Pseudomelissa (Davis, 1982) that differ from the other sections with larger, flat at the margin, and entire to weakly crenate leaves were also molecularly

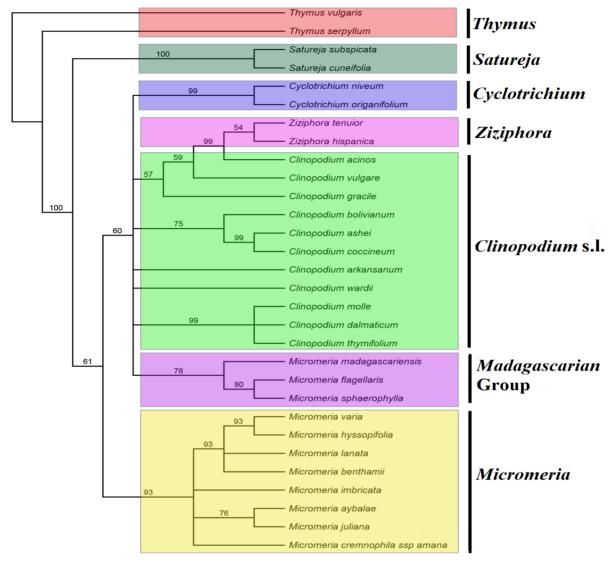


Figure 5. Maximum likelihood tree of *Micromeria* and related genera with bootstrap values (based on ITS data).

examined, and it was observed that these species belong to *Clinopodium* s.l. (Figures 5 and 6). The findings were consistent with the results of Bräuchler et al. (2005, 2010).

Sect. *Cymularia*, which is a monotypic section, is characterized by being annual, morphologically with ovateacuminate bracteoles and resupinate corolla. According to Brauchler et al. (2010), *Micromeria cymuligera* is closer to *Mentha* and *Cyclotrichum* than to *Micromeria*. In addition, our *trn*L-F data obtained from this study also showed this close relationship (Figure 6).

Sect. *Micromeria*, which contains the new species *Micromeria aybalae*, differs from the other two sections by a prominent vein at the leaves' margins, and mostly revolute and entire leaves. According to our nrDNA data (Figure 5, bootstrap value: 93) and cpDNA data (Figure 6, bootstrap value: 98), *M. aybalae* is included in Sect. *Micromeria*-real

*Micromeria*. It is seen in Figure 5 that *Micromeria aybalae* has a close relationship with *M. juliana* in the ITS region (bootstrap value: 76).

Finally, morphological differentiation was also supported using two different DNA regions. With this new species, the *Micromeria* species numbers in Turkey have increased to 9 species and 14 taxa, 9 of which are endemic to Turkey.

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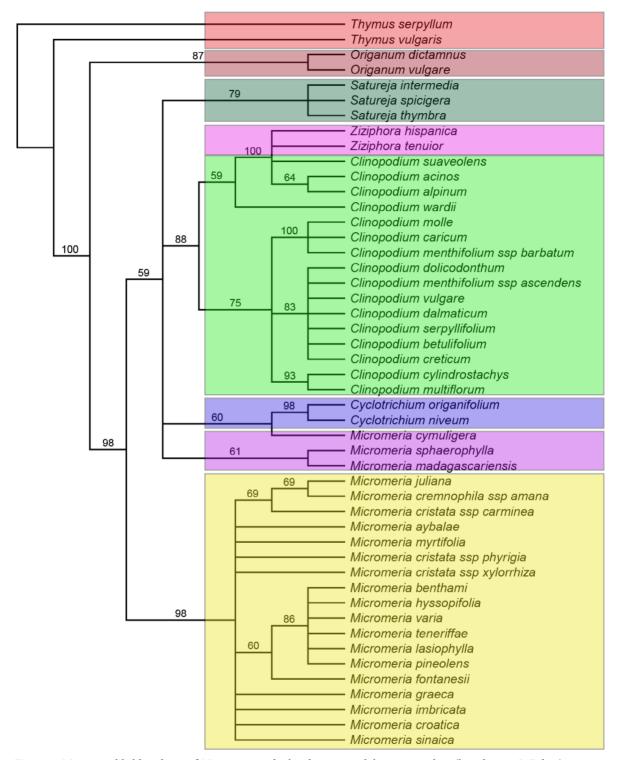


Figure 6. Maximum likelihood tree of *Micromeria* and related genera with bootstrap values (based on trnL-F data).

the specimens: ANK, B, BM, E, EGE, G, GAZI, GOET, HUB, ISTE, ISTF, K, LE, W, and WU. We also thank Infrastructure Action under the FP6 (SYNTHESYS Project GB-TAF 3087), FP7 (SYNTHESYS Project ES-TAF 264)

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#### Appendix 1. Examined specimens.

Micromeria cremnophila subsp. amana: Type. (Turkey) C6 Adana: Mons Amanus, mont de Düldül, 1500-2000 m a.s.l., 07.1911, Haradjian 3887 (E W); B5 Kayseri: Tomarza, Toklar, around Aslandaş village, 1700-1800 m a.s.l., 14.07.1989, M.Koyuncu 7198 (AEF); Bakırdağ, Kisge, at west foot of Bakır Da., 1300 m a.s.l., 27.06.1952, P.H.Davis 19217 (ANK, E, EGE, K, BM); Adana: Feke, Göksu gorge, below Himmetli village, 700-800 m, P.H.Davis 19870 (E, K, BM); Feke: between Suphandere and Belanköy, in gorge, 900 m a.s.l., 02.07.1952, P.H.Davis 19542 (BM); ibid., P.H.Davis 19565 (K, BM); ibid., P.H.Davis 19546 (ANK, K); C5 Adana: Pozanti, Alpu village, Karınca mountain, 1650 m a.s.l., 07.07.2007, T.Dirmenci 3450 & E.Akcicek (GAZI); 20 km from Saimbeyli to Feke, 800-900 m a.s.l., 11.07.2006, T.Dirmenci 3237 (GAZI); 10 km from Saimbeyli to Tufanbeyli, 1200 m a.s.l., 11.07.2007, T.Dirmenci 3462 & E.Akcicek (GAZI); C6 Osmanive: Bahce, montis amanus, regione de Düldül, 7000 ft., 07.1908, Haradjian 2407 (K); Düldül mountain, near Haruniye, 700 m a.s.l., 26.07.1949, P.H.Davis 16377 (ANK, K); Düldül mountain, Baskonus plateau, 1800 m a.s.l., 14.08.2014, T.Dirmenci 4254, T.Arabacı & T.Yazıcı (GAZI); Kahramanmaraş: Engizek Dağı, Aksu village, 1000-1100 m a.s.l., 05.07.1986, H.Duman 2083 (GAZI); subsp. anatolica: Type: (Turkey) B9 Van: 5 km north of Çatak, in crevices of boulders, 25.07.1954, P.H.Davis 23258 & O.Polunin (holo. E, iso K, BM, ANK); B6 Kayseri: Akkışla, above Ganişeyh, Hınzır mountain, 1900 m a.s.l., 22.07.1980, N.Çelik 1459 (AEF); Yalak, Binboğa mountain, 1750-2200 m a.s.l., 20.07.1992, Z.Aytaç & H.Duman 5367 (GAZI); Kahramanmaraş: Göksun, between Yesilköy and Kınıkkoz village, 1400-1600 m a.s.l., 13.06.1978, B.Yıldız 2092 (HUB); Sivas: Gürün, Gökpınar village, 1500 m a.s.l., 21.07.1987, M.Koyuncu (AEF-14434); Malatya: Darende, 1070 m a.s.l., 19.06.1954, P.H.Davis 21852

(E, K, BM); 50 km SE of Darende, 1200 m a.s.l., 05.08.1956, McNeill 440 (K); Malatya: Darende, ca. 1070 m a.s.l., 19.06.1954, P.H.Davis 21852 (ANK); between Malatya and Darende, 1 km east of Develi village, 4795 ft., 10.08.2007, T.Dirmenci 3504 & T.Arabacı (GAZI); B7 Elazığ: Harput, around castle, 17.07.1990, K.Alpınar (ISTE-62187); Erzincan: between Kemaliye and Sarıkonaklar, 1100-1200 m a.s.l., 08.06.1989, M.Koyuncu 8859 (AEF); Erzincan: between Kemaliye and Divriği, old road, the entrance of Bülent Ecevit tunnel, 15.06.2016, T.Dirmenci 4564 & A.Kahraman (GAZI). Tunceli: Munzur Da., Aksu De., above Ovacık, 1800 m a.s.l., P.H.Davis 31480 (E); B8 Muş: 15 km from Mus to Tatvan, 26.06.1983, T.Ekim 7773 (GAZI); Van: Atatürk forest, around DSİ and Radar, 2000 m a.s.l., 23.06.1988, Z.Aytaç 2330 (GAZI); B9 Van: Gürpınar, south of Hamurkesen village, 1900 m a.s.l., 15.07.2003, M. Ünal 8595 (VANF); C4 Konya: Ereğli, around Cakıllar village, 1900 m a.s.l., 20.07.1995, Z.Aytac 7169 & N.Adıgüzel (GAZI); C10 Hakkari: Cilo Da., Diz De., 1760 m a.s.l., 06.08.1954, P.H.Davis 23996 (E, K, BM).

*Micromeria hispida*: In rupestribus pr. Kainurio Chorio, (Eparhiae Mirabello) Cretae, 21.04.1846, Heldreich 1422 (isotypes: K WU P-photo); Crete: Selinaris, dans la gorge traverse par la route Iraklion-Agios Nikolaos, a proximate du monatere d'Agios Georgios, fin Avril, 1997, G.Van Buggenhout 18344 (G); Crete, Hierapetra, Felsgerölle des aphendi Kavusi, 02.08.1904, Dörfler 1054 (G-Boiss.).

*Micromeria microphylla*: Malta: "In collibus aridis insulae Melitae copiosissime", d'Urville (Lectotype: P-photo); Karpathos, in fissuris rupium calc. ad Vrondi adversus Pigadia, 17.06.1935, K.H.Rechinger 8249 (G-Boiss.).

*Micromeria chionistrae*: Cyprus, Phini 1000 m a.s.l., in cracks of bare rock, 06.06.1939, Kennedy 1495 (holotype: K).

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**Appendix 2.** GenBank information.

	GenBank Acce	GenBank Accession Numbers	
	ITS	trnL-trnF	
Thymus vulgaris L.	AY506646	GU381636	
Thhymus serpyllum L.	EU796890	AY570502	
Satureja subspicata Bartl. ex Vis.	EU823288		
Satureja cuneifolia Ten.	EU823290		
Satureja intermedia C.A.Mey.		GU381622	
Satureja spicigera (K.Koch) Boiss.		GU381623	
Satureja tymbra L.		JQ669068	
Cyclotrichium niveum (Boiss.) Manden. & Scheng.	GU381397	GU381525	
Cyclotrichium origanifolium (Labill.) Manden. & Scheng.	GU381399	GU381527	
Ziziphora tenuior L.	KP278588	GU381507	
Ziziphora hispanica L.	KP278578	GU381503	
Origanum dictamnus L.		GU381643	
Origanum vulgare L.		AY506614	
Clinopodium acinos (L.) Kuntze	JQ669074	GU381498	
Clinopodium vulgare L.	DQ667324	AY506593	
Clinopodium gracile (Benth.) Kuntze	JQ669082		
Clinopodium bolivianum (Benth.) Kuntze	DQ017564		
Clinopodium ashei (Weath.) Small	DQ667237		
Clinopodium coccineum (Nutt. ex Hook) Kuntze)	AY943485		
Clinopodium arkansanum (Benth.) Kuntze	JQ669080		
Clinoopodium wardii (C.Marquand & Airy Shaw) Bräuchler	GU381392	GU381516	
Clinopodium dalmaticum (Benth.) Bräuchler & Heubl	JQ669118	JQ669055	
Clinopodium thymifolium (Scop) Kuntze	JQ669121		
Clinopodium suaveolens (Sm.) Kuntze)		GU381499	
Clinopodium alpinum (L.) Kuntze		AY840180	
Clinopodium serpyllifolium (M.Bieb.) Kuntze		GU381535	
Clinopodium serpyllifolium (M.Bieb.) Kuntze		GU381536	
Clinopodium betulifolium (Boiss. & Balansa) Kuntze		GU381532	
Clinopodium creticum (L.) Kuntze		GU381533	
Clinopodium cylindrostachys (Epling & Játiva) Govaerts		GU381605	
Clinopodium multiflorum (Ruiz & Pav.) Kuntze		GU381602	
Micromeria madagascariensis Baker	GU381374	GU381481	
Micromeria flagellaris Baker	GU381375		
Micromeria sphaerophylla Baker	GU381376	GU381487	
Micromeria varia Benth.	GU381447	AY840203	
Micromeria hyssopifolia Webb & Berthel.	AY227142	AY506612	
Micromeria lanata (C.Sm. ex Link) Benth.	JQ669120		
Micromeria benthamii Webb & Berthel.	GU381446	AY840183	
Micromeria imbricata (Forssk.) C.Chr.	KF805105	AY840193	
Micromeria cymuligera Boiss. & Hausskn.		GU381523	

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### Appendix 2. (Continued).

Micromeria teneriffae (Poir.) Benth. ex G.Don		AY840206
Micromeria lasiophylla Webb & Berthel.		AY840205
Micromeria pineolens Svent.		AY840182
Micromeria fontanesii Pomel		AY840195
Micromeria graeca (L.) Benth. ex Rchb.		AY840198
Micromeria croatica (Pers.) Schott		GU381624
Micromeria sinaica Benth.		AY840200
Micromeria cremnophila Boiss. & Heldr. subsp. amana (Rech.f.) P.H.Davis*	KY020411	KX381823
Micromeria cristata (Hampe) Griseb. subsp. phyrigia P.H.Davis*		KX381824
Micromeria cristata subsp. carminea (P.H.Davis) P.H.Davis*		KX381825
Micromeria cristata subsp. xylorrhiza (Boiss. & Heldr. ex Benth.) P.H.Davis*		KX381826
Clinopodium serpyllifolium subsp barbatum (P.H.Davis) Bräuchler*		KY038988
Micromeria aybalae H.Duman & Dirmenci*	KX381815	KX381818
Micromeria juliana (L.) Benth. ex Rchb.*	KX381816	KX381821
Micromeria myrtifolia Boiss. & Hohen.*		KX381822
Clinopodium molle (Benth.) Kuntze*	KX381814	KX381819
Clinopodium caricum (P.H.Davis) Bräuchler & Heubl*		KX381817
Clinopodium dolicodonthum (P.H.Davis) Bräuchler & Heubl**		KX381820

<sup>\*:</sup> Isolated in this study.