

Nuevas citas de murciélago ratonero bigotudo pequeño, *Myotis alcathoe* (Vespertilionidae) en Italia

New records of the Alcathoe bat, *Myotis alcathoe* (Vespertilionidae) for Italy

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Abstract: The Alcathoe bat (*Myotis alcathoe*) is a recently described cryptic species; in Europe its distribution range is poorly known. In Italy this species has been recorded in a small number of locations in Abruzzo (central Italy) and Campania (southern Italy). Our report refers to three bats captured in a mountainous area dominated by forest habitats in the Appennino Lucano Val d'Agri Lagonegrese National Park (Basilicata region). The identification of bats captured was confirmed by molecular analysis using the technique of DNA barcoding. In this paper we present new recordings that highlight the presence of the species in other regions of southern Italy and that help define its distributional status in Europe.

Key words: *Myotis alcathoe*, cryptic species, DNA barcoding, coxI.

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The presence of cryptic species seem to be widespread among vespertilionid bats and especially in the genus *Myotis* (Jones and Barlow 2004).

Myotis alcathoe is a recently described cryptic species (von Helversen et al. 2001) belonging to the group *mystacinus*.

This small bat is difficult to identify only through morphological analysis.

Although the species can be identified using both morphological characters and genetic markers, only the later provide full confidence about it (Niermann et al. 2007).

In Europe population size and trends of *M. alcathoe* are poorly known.

In Italy the species was recorded for five localities in Abruzzo (central Italy) and Campania (southern Italy). All bats were captured in beech forests (Tereba et al. 2008, Galimberti et al. 2012).

Our records were collected as part of a survey of bats covering the entire territory of the Appennino Lucano Val d'Agri Lagonegrese National Park (south-west of the Basilicata region). We made two capture sessions on 3 and 6 August, 2012 near two small mountain lakes at 1167 and 1304 m a.s.l., respectively. Small lakes are part of a forest reserve dominated by beech (*Fagus sylvatica*) associated with other tree species, such as Lobel maple (*Acer lobelii*), sycamore (*Acer pseudoplatanus*) and Small-leaved Lime (*Tilia cordata*), fig. 1.

Bats were captured with 2.5 x 12 m mist-nets (50 denier, 38 mm mesh) placed on commuting routes and near small lakes used by bats for drinking and foraging.

The nets were deployed half an hour after sunset and kept in place for 4 hours. Forearm length and body mass of each bat trapped were measured using a digital callipers (± 0.1 mm) and a pesola digital scale (± 0.1 g). Wings were trans-illuminated to distinguish juveniles from adults (Antony 1988).



Fig. 1 – Records of *Myotis alcaethoe*.

Bats captured were subjected to a skin biopsy using a sterile punch of a 3 mm diameter from the tail membrane (uropatagium) (Worthington Wilmer and Barratt 1996).

Samples were stored in sterile tubes containing 95% ethanol for subsequent molecular analysis.

Total genomic DNA was extracted from tissue samples using 5 PRIME, ArchivePure DNA Purification Kit. A fragment of ca. 650 bp of the mitochondrial subunit 1 of cytochrome c oxidase, suitable for echolocating bats identification (Galimberti et al. 2012), was amplified for the three sampled bats using the primers VF1d 5'-TTCTCAACCAACCACAARGAYATYGG-3' and VR1d 5'-TAGACTTCTGGGTGGCCRAARAAYCA-3' from Ivanova et al. (2007).

PCR reactions were performed in 20µl reactions using ca. 1 ng of genomic DNA, 0.2µl of VF1d (0.2 mM), 0.2µl of VR1d (0.2 mM), 2µl of total dNTPs (0.2 mM), 0.1µl of (0.5U) of MasterTaq Eppendorf®, 2µl 1x Buffer including MgCl₂ at 1.5 mM and 14.5µl of water. PCR conditions were: 1 min at 94°C, followed by 5 cycles of 30 s at 94°C, 40 s at 50°C, and 1 min at 72°C, followed by 35 cycles of 30 s at 94°C, 40 s at 55°C, and 1 min at 72°C, and ending with 10 min at 72°C. The light strands were sequenced using an ABI3730XL by Macrogen Inc. Chromatographs were checked by eye and sequences were edited, when necessary, using the BioEdit sequence alignment editor (version 7.0.5.3; Hall 1999). To assess species attribution each sequence was compared using the BLAST algorithm in GenBank, where sequences

belonging to almost all the Italian echolocating bats species are available thanks to a previous study (Galimberti et al. 2012). All sequences have been deposited in GenBank (HG325822-23-24).

We caught three lactating females (FAL = 33.6, 32.9 and 33.0 mm respectively; weight = 4.7, 4.9 and 4.0) from the “*M. mystacinus* group”, two in Marsico Nuovo (altitude 1167 m a.s.l.) and one in Calvello (altitude 1304 m a.s.l.). All there caught in beech forest sites.

Molecular analysis of these samples allowed the taxonomic identification of the three bats as belonging to the species *M. alcaethoe*. The high similarity matches with reference deposited sequences excluded the misidentification with other congeners.

In central and eastern Europe, *M. alcaethoe* seems to prefer deciduous forests with old trees and streams (Niermann et al. 2007; Řehák et al. 2008; Lučan et al. 2009; Bashta et al. 2011). Our data adds further records for the presence of *M. alcaethoe* in beech forests of southern Italy (Tereba et al. 2008).

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