



ELSEVIER

Contents lists available at ScienceDirect

Data in Brief

journal homepage: www.elsevier.com/locate/dib



Data Article

Molecular data on the CO1 and beta fibrinogen gene in the evolutionary relationships of the mastiff bat (Chiroptera, Molossidae, *Molossus*)

Livia O. Loureiro^{a,*}, Burton K. Lim^b, Mark D. Engstrom^{a,b}

^a Department of Ecology and Evolution, University of Toronto, Toronto, ON, Canada M5S 3B2

^b Department of Natural History, Royal Ontario Museum, Toronto, ON, Canada M5S 2C6

ARTICLE INFO

Article history:

Received 23 January 2018

Accepted 23 April 2018

Available online 30 April 2018

Keywords:

Molossidae
Phylogenetics
Neotropics
Evolution

ABSTRACT

Molossus is one of the most diverse genera of free-tailed bats in the pantropical family Molossidae and occurs through all the Neotropics. Nevertheless, the taxonomy and phylogeny of this group is poorly understood. Here, we present the data on evolutionary relationships of *Molossus* based on DNA barcodes of COI gene from 346 specimens of *Molossus* and its sister genus *Promops* and another New World molossid *Eumops*. Of these specimens, 50 are new sequences and 296 were obtained from GenBank. In addition, the nuclear gene beta fibrinogen was sequenced from a subset of 35 specimens. These data provide the basis for further exploration and understanding of the phylogenetic relationships of the genus *Molossus* (Loureiro et al., 2018) [1].

© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Specifications table

Subject area	Biology
More specific subject area	Genetics, Molecular phylogeny

DOI of original article: <https://doi.org/10.1016/j.mambio.2018.01.008>

* Corresponding author.

E-mail address: livia.loureiro@mail.utoronto.ca (L.O. Loureiro).

<https://doi.org/10.1016/j.dib.2018.04.088>

2352-3409/© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Type of data	Figures
How data was acquired	Tissues were extracted using the DNeasy Tissue Kit (QIAGEN Inc., Valencia, California). PCR protocols followed [2–4]. The nucleotides were sequenced in an ABI 3130 (Applied Biosystems_) automated sequencer using Big Dye Terminator Cycle Sequencing methodology (Applied Biosystems_).
Data format	Analyzed
Experimental factors	Total genomic DNA was extracted from liver, heart or kidney tissue that were frozen in liquid nitrogen or preserved in ethanol.
Experimental features	Sequences were assembled in SEQUENCHER and aligned using the Muscle algorithm [5]. Phylogenetic relationships were reconstructed MEGA 6.06 [6].
Data source location	Bonaire, Dominican Republic, Ecuador, El Salvador, French Guiana, Guyana, Jamaica, Martinique, Mexico, Panama, Peru, Suriname, Venezuela, Bolivia, and Montserrat.
Data accessibility	The sequences have been deposited in the public repository of GenBank and BOLD systems (Supplementary material 1)

Value of the data

- Phylogenetic relationships within the genus remain undefined and until recently there have been few molecular studies of *Molossus* [1,11,12]. Therefore, these data combined with more genetic markers, more species of the genus, and more comprehensive geographic sampling could clarify the evolution of *Molossus*.
 - These data could help to test the homology of many morphological and ecological characters, such as echolocation calls.
 - *Molossus* is a Neotropical genus, occurring from the southeastern United States to southern Argentina, and throughout the Caribbean islands. Therefore, these data could be used in the development of biogeographic studies in the Neotropics.
 - This data could be used for comparative studies related to other genera of molossid bats to understand the evolution of the family Molossidae.
-

1. Data

Molossus is one of the most diverse and common genera of the family Molossidae, but its taxonomy and phylogenetic relationships are still poorly understood. Maximum likelihood trees for the mitochondrial CO1 gene (Fig. 1) and nuclear beta fibrinogen gene (Fig. 2) from more than 300 specimens of *Molossus* and its sister groups distributed through all the Neotropics estimate the phylogenetic relationships within the genus. We also present the specimen vouchers used in the genetic analyses for the genes COI and Beta fibrinogen, the GenBank and BOLD Systems accessions numbers, the species identification, and country where the specimens were collected (Supplementary material 1).

2. Experimental design, materials and methods

2.1. DNA sequencing

Total genomic DNA was extracted from liver, heart or kidney tissue that were frozen in liquid nitrogen or preserved in ethanol. Tissues were extracted using the DNeasy Tissue Kit (QIAGEN Inc.,

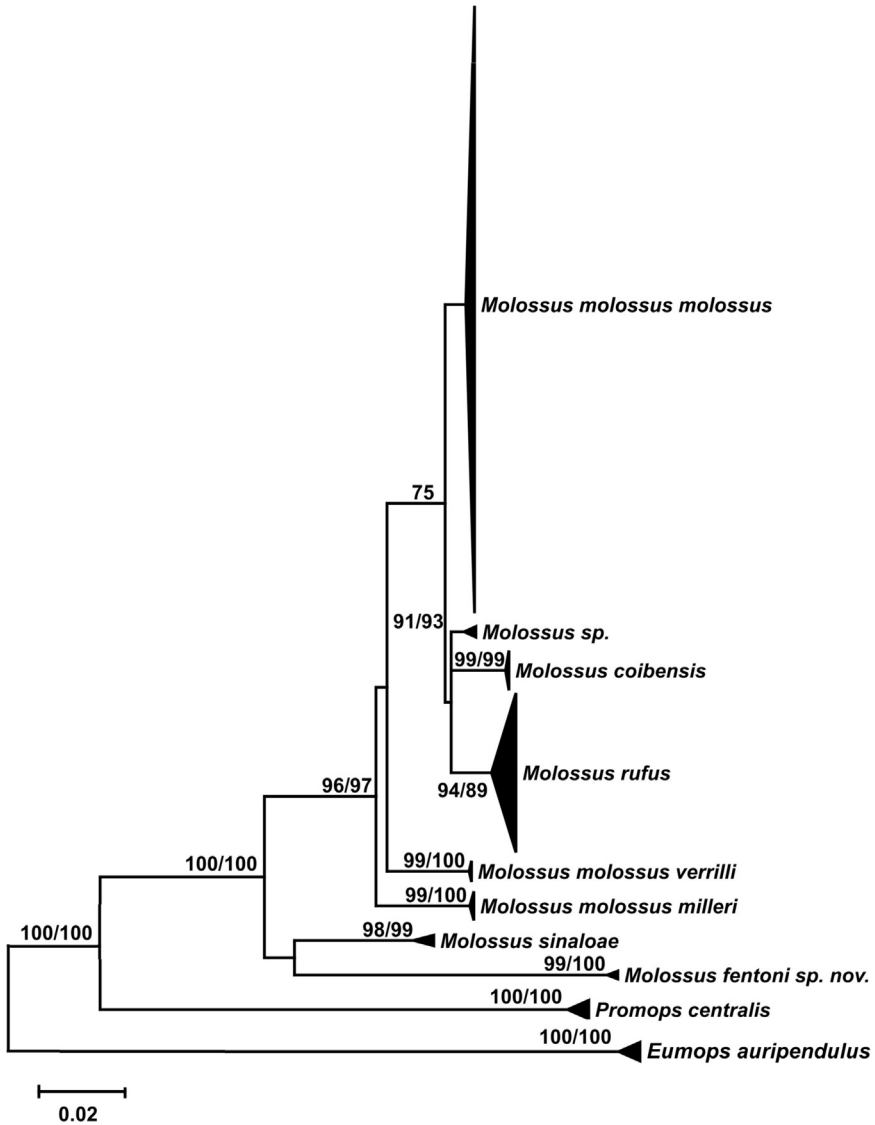


Fig. 1. Maximum likelihood tree of COI sequences of *Molossus*. Bootstrap support values (maximum likelihood/maximum parsimony) > 70% are reported for well-supported nodes. *M. m. daulensis* was recovered inside the *M. m. molossus* clade.

Valencia, California) following the manufacturer's protocol. Molecular protocols for the COI gene followed the methods outlined by Refs. [2,3] and protocols for the beta fibrinogen gene followed [4]. The nucleotides of both strands were sequenced in an ABI 3130 (Applied Biosystems_) automated sequencer using Big Dye Terminator Cycle Sequencing methodology (Applied Biosystems_). These analyses were carried out in accordance with the recommendations of the Canadian Council on Animal Care, the requirements under the Animals for Research Act (Revised Statutes of Ontario, 1990), and the Royal Ontario Museum Animal Care Policies and Guidelines for animal experiments.

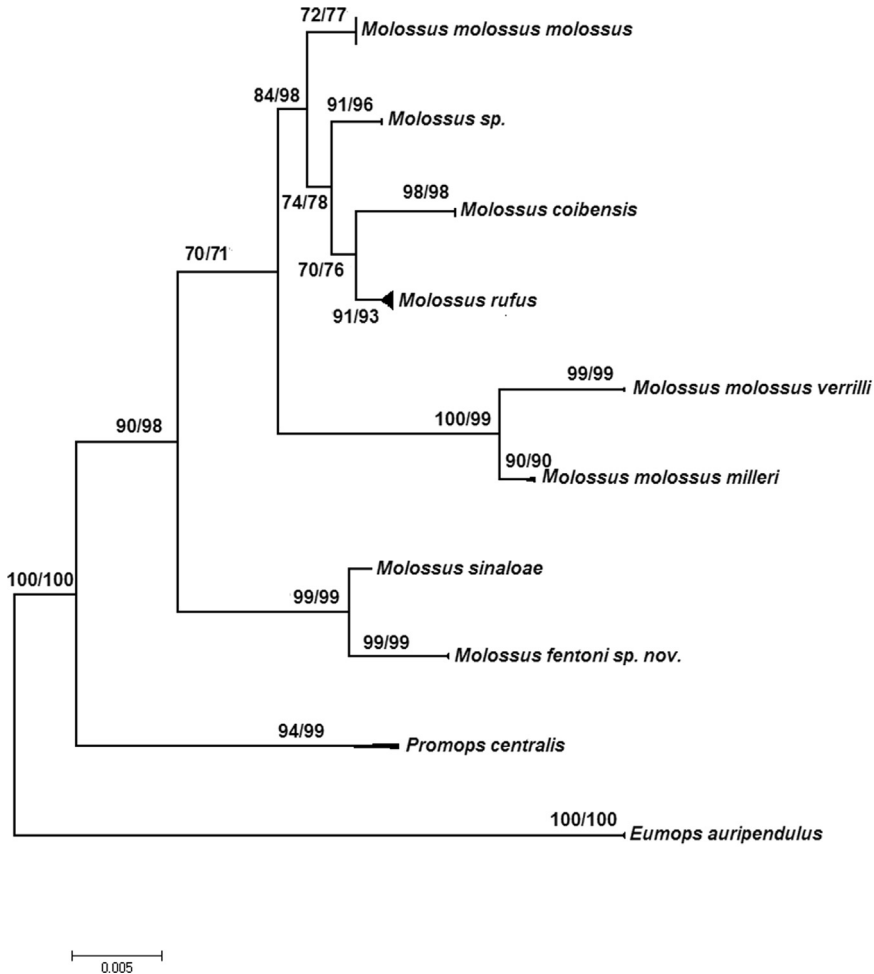


Fig. 2. Maximum likelihood tree of beta-fibrinogen sequences of *Molossus*. Bootstrap support values (maximum likelihood/maximum parsimony) > 70% are reported for well-supported nodes. *M. m.*

2.2. Phylogenetic analyses

DNA barcodes of 657 basepairs of COI were analyzed from 346 specimens of *Molossus* from across the Neotropics including Bonaire, Dominican Republic, Ecuador, El Salvador, French Guiana, Guyana, Jamaica, Martinique, Mexico, Panama, Peru, Suriname, and Venezuela. The genera *Eumops* and *Promops* were included as outgroups following [7,8]. Of these specimens, 50 are new sequences and 296 were obtained from GenBank (Supplementary material 1). Based on genetic divergence in COI, a subset of 35 specimens spanning the breadth of variation was sequenced for 764 basepairs of the nuclear gene beta fibrinogen. Sequences were assembled in SEQUENCHER and aligned using the Muscle algorithm [5] as implemented in MEGA 6.06 [6]. Phylogenetic relationships were reconstructed for each single dataset using maximum likelihood analyses with 1000 bootstrap replications as implemented in MEGA 6.06 [6] (Figs. 1 and 2). Aligned datasets were subjected to alternative models of sequence evolution in Partition Finder 1.0.1 [9] to select the best partitions and models of sequence evolution [10].

Acknowledgements

This work was supported by Coordenação de Aperfeiçoamento de Pessoal de Nível Superior (Capes) (9 99999.011880/2013–09). Neotropical fieldwork has been primarily funded by the Royal Ontario Museum with additional financial support in Ecuador by MAXUS Inc. and in Guyana by Conservation International and funding through the Academy of Natural Sciences, Philadelphia. We thank the following curators and collection support staff that provided access or loaned specimens: B. Patterson (FNMH), C.J. Conroy (MVZ), M. Campbell (MSB), Dr. B. S. Coyner (Sam Noble Museum), Dr. N. Simmons (AMNH), Dr. H.J. Garner (TTU), Dr. C. Lopez-Gonzalez (Instituto Politécnico Nacional, Mexico City), Dr. J. Juste (CSIC), and A.L. Gardner (NMNH/ USMN).

Transparency document. Supporting information

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.dib.2018.04.088>.

Appendix A. Supporting information

Supplementary data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.dib.2018.04.088>.

References

- [1] L.O. Loureiro, B.K. Lim, M.K. Engstrom, A new species of mastiff bat (Chiroptera, Molossidae, *Molossus*) from, Guyana and Ecuador. *Mamm Biol.* 90 (2018) 10–21.
- [2] A.V. Borisenko, B.K. Lim, N.V. Ivanova, R.H. Hanner, P.D.N. Hebert, DNA barcoding in surveys of small mammal communities: a field study in Suriname. *Mol. Ecol. Resour.* 8 (2008) 471–479.
- [3] E.L. Clare, B.K. Lim, M.D. Engstrom, J.L. Eger, P.D.N. Hebert, DNA barcoding of Neotropical bats: species identification and discovery within Guyana. *Mol. Ecol. Notes* 7 (2007) 184–190.
- [4] S.A. Reeder, R.D. Bradley, Phylogenetic relationships of neotomine–peromyscine rodents using DNA sequences from beta fibrinogen and cytochrome b, in: D.A. Kelt, E.P. Lessa, J.A. Salazar-Bravo, J.L. Patton (Eds.), *The Quintessential Naturalist: Honoring the Life and Legacy of Oliver P. Pearson*, University of California Publications in Zoology, 2007, pp. 883–900.
- [5] R.C. Edgar, MUSCLE: multiple sequence alignment with high accuracy and high throughput. *Nucleic Acids Res.* 32 (2004) 1792–1797.
- [6] K. Tamura, G. Stecher, D. Peterson, A. Filipiński, S. Kumar, MEGA6: molecular evolutionary genetics analysis version 6.0. *Mol. Biol. Evol.* 30 (2013) 2725–2729.
- [7] L.K. Ammerman, D.N. Lee, T.M. Tipps, First molecular phylogenetic insights into the evolution of free-tailed bats in the subfamily Molossinae (Molossidae, Chiroptera). *J. Mammal.* 93 (2012) 12–28.
- [8] R. Gregorin, A. Cirranello, Phylogeny of Molossidae Gervais (Mammalia: chiroptera) inferred by morphological data. *Cladistics* 32 (2016) 2–35.
- [9] R. Lanfear, B. Calcott, S.Y.W. Ho, S. Guindon, Partition Finder: combined selection of partitioning schemes and substitution models for phylogenetic analyses. *Mol. Biol. Evol.* 29 (2012) 1695–1701.
- [10] H. Akaike, A new look at the statistical model identification. *IEEE Trans. Autom. Control* 19 (1974) 716–723.
- [11] Y. Gager, E. Tarland, D. Lieckfeldt, M. Ménage, F. Botero-Castro, S.J. Rossiter, R.H.S. Kraus, A. Ludwig, D.K.N. Dechmann, The value of molecular vs. morphometric and acoustic information for species identification using sympatric molossid bats. *PLoS ONE* 11 (2016) e0150780.
- [12] L.L. Lindsey, L.K. Ammerman, Ammerman. Patterns of genetic diversification in a widely distributed species of bat, *Molossus molossus*. *Occas. Papers Mus. Texas Tech* 339 (2016) 1–16.