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Mitogenome of northern long-eared bat

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Bellevue University, Bellevue, NE, USA; Nebraska Cooperative Fish and Wildlife Research Unit, and School of Natural Resources, University of Nebraska-Lincoln, Lincoln, NE, USA; U.S. Geological Survey—Nebraska Cooperative Fish and Wildlife Research Unit, and School of Natural Resources, University of Nebraska-Lincoln, Lincoln, NE, USA; Department of Biology, University of Nebraska Omaha, Omaha, NE, USA; University of Nebraska-Lincoln, Lincoln, NE, USA; School of Natural Resources, and University of Nebraska State Museum, University of Nebraska-Lincoln, Lincoln, NE, USA

ABSTRACT

The complete mitogenome of the northern long-eared bat (Myotis septentrionalis) was determined to be 17,362 bp and contained 22 tRNA genes, 2 rRNA genes and one control region. The whole genome base composition was 33.8% GC. Phylogenetic analysis suggests that M. septentrionalis be positioned next to M. auriculus in the Nearctic subclade of the Myotis genus. This complete mitochondrial genome provides essential molecular markers for resolving phylogeny and future conservation efforts.

The northern long-eared bat (Myotis septentrionalis) has recently experienced drastic population declines in eastern and midwestern parts of its range because of the invasive fungal disease white-nose syndrome (WNS) (Frick et al. 2015; Langwig et al. 2015). The disease induces physiological and behavioral changes in bats during hibernation, which can result in death (Verant et al. 2014). Population declines have been so severe that M. septentrionalis was listed as threatened in the United States of America (USFWS 2015) and endangered in Canada (COSEWIC 2013). M. septentrionalis seems to be more susceptible to WNS than other closely related species, such as the little brown bat (Myotis lucifugus); however, the cause of this susceptibility has yet to be determined and may be due to genetic differences or varying environmental preferences (Frick et al. 2015; Langwig et al. 2016). Regulation of specific mitochondrial genes, including COI, ND2, ATP6 and ATP8, is crucial during the hibernation process (Hittel and Storey 2002); therefore, comparative analysis of mitochondrial genomes of hibernating bat species might offer some insight into how WNS affects species differently. Here we report the first complete mitogenome of M. septentrionalis and examine the phylogenetic position of M. septentrionalis within the genus Myotis based on complete mitogenomes.

We collected wing tissue from an adult, female M. septentrionalis on 5 July 2017 at Ponca State Park (Dixon County) in northeastern Nebraska (42.6022°N, 96.7154°W). We used sterile 2-mm disposable biopsy punches to collect two tissue plugs from the flight membrane near the leg of the bat to avoid large blood vessels, which were easily seen in the flight membrane. A representative tissue plug from an adult, female M. septentrionalis that was collected at the same site on the same evening was deposited at the University of Nebraska State Museum (catalog number UNSM ZM-31046). After tissue collection, bats were released at points of capture. Each tissue plug was stored dry in a cryogenic tube with several silica beads. Upon return from the field, tissue plugs were frozen at −80 °C until DNA extraction. Genomic mitochondrial DNA was extracted and purified from one tissue plug using the standard protocol of the Abcam Mitochondrial DNA Isolation Kit and sequenced on an Illumina NextSeq500 at the University of Nebraska Medical Center. The mitogenomic sequence was assembled and annotated using Geneious (Kearse et al. 2012).

The total length of the mitogenome was 17,362 bp (GenBank Accession No. MK547202). The mitogenome consisted of 22 tRNA genes, two rRNA genes and one control region. The whole genome base composition was 33.8% GC.

To investigate the position of M. septentrionalis within the genus Myotis, we constructed a maximum likelihood tree based on 19 complete mitochondrial genomes using MEGA 6 under the GTR + G + I model with 500 bootstrap replicates (Pattengale et al. 2010; Tamura et al. 2013). The phylogenetic tree contained two subclades, Nearctic and Neotropical (Figure 1). The position of M. septentrionalis next to M. auriculus in the Nearctic subclade of the Myotis genus corresponds with previously proposed phylogenetic relationships (Stadelmann et al. 2007).
This mitogenome establishes a basis for additional, future phylogenetic studies of this diverse genus, as well as studies on the effects of WNS on *M. septentrionalis* in comparison to other species of hibernating bats. Future studies should consider the susceptibility of *Myotis* bats to WNS in relation to their Nearctic and Neotropical subclade groupings.

**Disclosure statement**

The authors report no conflict of interest. This study was performed under the auspices of the University of Nebraska Omaha IACUC protocol # 18-072-06-FC and USFWS permit number TE79842A-1. Any use of trade, firm, or product names is for descriptive purposes only and does not imply endorsement by the U.S. Government.

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**Data availability statement**

The data that support the findings of this study are openly available in GenBank of NCBI at [https://www.ncbi.nlm.nih.gov](https://www.ncbi.nlm.nih.gov), reference number MK547202.

**References**


