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Sarah J. Gaughan, Kevin L. Pope, Jeremy A. White, Cliff A. Lemen & Patricia W. Freeman

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MITOGENOME ANNOUNCEMENT

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 storeddry 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. 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.

**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.

**KEYWORDS**

*M. septentrionalis*; mitochondrial genome; next generation sequencing
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, reference number MK547202.

References


