MITOGENOME ANNOUNCEMENT

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The complete mitochondrial genome of *Sarcophila mongolica* (Diptera: Sarcophagidae)

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ABSTRACT

Sarcophila mongolica Chao & Zhang, 1988 (Diptera: Sarcophagidae) is considered to be of ecological and medical significance. In this study, we report the mitochondrial genome (mitogenome) of *S. mongolica*. This mitogenome was composed of 15,936 bp in length (GenBank accession no. MT845211), comprising 13 protein-coding genes (PCGs), two ribosomal RNAs (rRNAs), 22 transfer RNAs (tRNAs), and a non-coding control region. The arrangement of genes was identical to that of ancestral metazoan. Nucleotide composition revealed a strong A + T bias, accounting for 75.40% (A 38.2%, G 9.7%, C 14.9%, and T 37.2%). Phylogenetic analysis indicated that *S. mongolica* was obviously separated from the other flesh flies. This mitogenome provides important genetic data for further understanding of the evolutionary relationship within Sarcophagid flies.

ARTICLE HISTORY Received 19 August 2020

Accepted 8 November 2020

Taylor & Francis

Taylor & Francis Group

KEYWORDS Mitochondrial genome; *Sarcophila mongolica*; phylogenetic analysis

Although the larvae of Sarcophila genus usually colonize on invertebrate carcasses, Sarcophila mongolica Chao & Zhang, 1988 (Diptera: Sarcophagidae) has so far been rarely reported except that the adult was found only in Inner Mongolia (Pape 1996; Xu and Zhao 1996). In this study, adult specimens were first trapped by pig liver in June 2019 from Ürümqi city (43°50′N, 87°37′E), Xinjiang province, China. Whether they are of forensic importance or just accidental visitors remain unknown. All specimens were killed by freezing, and then identified by traditional morphological keys (Xu and Zhao 1996). These specimens were deposited at -80 °C in Guo's lab (Hunan, Changsha, China) with a unique code (CSU19111977). Total DNA was extracted from thoracic muscle tissues of an adult specimen using QIANamp Micro DNA Kit (Qiangen Biotech Co., Ltd) according to the manufacture's instruction. The genome sequencing of S. mongolica was performed on an Illumina HiSeg 2500 Platform, and then de novo assembly was carried out with MITObim version 1.9 and SOAPdenovo version 2.04 (https://github.com/chrishah/ MITObim and http://soap.genomics.org.cn/soapdenovo.html)

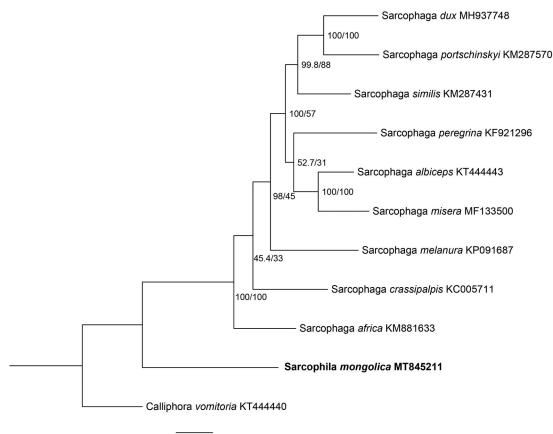
(Ren et al. 2020). Finally, the rough boundaries of all genes were initially identified by MITOS2 Web Server (http://mitos2. bioinf.uni-leipzig.de/index.py) (Ren et al. 2020).

In this study, the mitogenome of S. mongolica was 15,936 bp in length (GenBank accession no. MT845211), containing 13 protein-coding genes (PCGs), two ribosomal RNAs (rRNAs), 22 transfer RNAs (tRNAs), and a non-coding control region. The arrangement of genes was identical to that of ancestral metazoan (Cameron 2014). Nucleotide composition revealed a highly A+T bias, accounting for 75.40% (A 38.2%, G 9.7%, C 14.9%, and T 37.2%). Phylogenetic analysis of S. mongolica and other nine Sarcophagids species was constructed using maximum likelihood (ML) method based on the 13 PCGs, and Calliphora vomitoria (Diptera: Calliphoridae) was used as an outgroup (Figure 1). ML analysis was performed by IQ-TREE version 1.6.12 (Ren et al. 2020). The phylogenetic relationships indicated that the species of S. mongolica was clearly separated from the other flesh flies. This study provided important mitochondrial data for further

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Figure 1. Phylogenetic trees of S. mongolica with nine sarcophagids species based on 13 PCGs by maximum likelihood (ML) method. Calliphora vomitoria was selected as an outgroup.

studying on evolutionary relationships and species identification of flesh flies.

Disclosure statement

No potential conflict of interest was reported by the author(s).

Acknowledgments

We are grateful to Prof. Lushi Chen (Guizhou Police College) for species identification.

Funding

This study is supported by the National Natural Science Foundation of China [No. 81772026 and 82072114] and the National Natural Science Foundation of Hunan Province [No. 2020JJ4763].

Data availability statement

Mitogenome data supporting this study are openly available in GenBank at: https://www.ncbi.nlm.nih.gov/nuccore/MT845211. Associated BioProject, SRA, and BioSample accession numbers are https://dataview.ncbi.nlm.nih. gov/object/PRJNA665806, https://www.ncbi.nlm.nih.gov/sra/SRR12719806, and SAMN16268934, respectively.

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