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#### MITOGENOME ANNOUNCEMENT

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# The complete mitochondrial genome of *Artemia sinica* Cai, 1989 (Crustacea: Anostraca) using next-generation sequencing

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#### ABSTRACT

The complete mitochondrial genome of *Artemia sinica* was obtained using the next-generation sequencing (NGS) method. The mitochondrial genome is a circular molecule of 15,689 bp in length, with the typical structure of 13 protein-coding genes (PCGs), 22 transfer RNA genes (tRNAs) and 2 ribosomal RNA genes, and a non-coding control region (CR). The base composition is 31.53% A, 18.99% C, 16.50% G, and 32.98% T, with an A + T content of 64.51%. All tRNAs have a cloverleaf structure excepting *tRNA-Ser*<sub>1</sub>, that represents the D-loop structure.

#### **ARTICLE HISTORY**

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#### **KEYWORDS**

Mitogenome; brine shrimp; Artemia sinica; proteincoding genes; transfer RNA genes; ribosomal RNA genes

The genus Artemia consists of seven bisexual species and a large number of parthenogenetic populations with different ploidy degrees (Asem et al. 2010). So far, the complete mitogenome of three bisexual species of Artemia (Artemia franciscana, Artemia urmiana, and Artemia tibetiana) have already been characterized(Valverde et al. 1994; Zhang et al. 2013). In this study, we sequenced and described the complete mitochondrial genome of Artemia sinica (GenBank: MK069595). The present study is a part of a more comprehensive project of characterizing the complete mitochondrial genome of genus Artemia.

The cysts of A. sinica (ARC: 1166) were collected from Ejinor Lake (Inner Mongolia, China) and stored in the Artemia Reference Center (Ghent University, Belgium). The total DNA was extracted from a cultured adult specimen. A genomic library was established followed by paired-end  $(2 \times 150 \text{ bp})$ next-generation sequencing (10 Gb), using the Illumina HiSeq X-ten sequencing platform. Quality checks for sequencing reads were performed by FastQC (Andrews 2010) and the sequences were assembled and mapped to the reference Artemia mitochondrial genome (A. franciscana, X69067) with Spades v3.9.0 (Bankevich et al. 2012) and bowtie v2.2.9 (Langmead and Salzberg 2012). Putative tRNA genes were determined using the tRNAscan-SE2.0 (http://lowelab.ucsc. edu/tRNAscan-SE/) and ARWEN (http://130.235.46.10/ARWEN/) online software. All genes were annotated based on gene order on the reference mitochondrial map and using BLAST analysis (https://blast.ncbi.nlm.nih.gov). Additionally, to annotate PCGs and tRNAs, the position of start and stop codons,

and secondary structures and the position of anticodons were re-considered, respectively.

The complete mitogenome of *A. sinica* was 15,689 bp in length, with 13 protein-coding genes (PCGs), 22 transfer RNAs (tRNAs), 2 ribosomal RNAs (rRNAs), and a control region (CR). The overall nucleotide composition of the major strand of the *A. sinica* mitogenome was as follows: 31.53% A, 18.99% C, 16.50% G, and 32.98% T, with a total A + T content of 64.51%.

Nine tRNAs (*tRNA-lle*, *tRNA-Gln*, *tRNA-Cys*, *tRNA-Tyr*, *tRNA-Phe*, *tRNA-His*, *tRNA-Pro*, *tRNA-Leu*, and *tRNA-Val*) and four PCGs (*ND5*, *ND4*, *ND4L*, and *ND1*), as well as both rRNAs were encoded on the light strand. Just six PCGs (*ND2*, *COX1*, *ATP6*, *COX3*, *CYTB*, and *ND1*) began with the common ATG start codon. Stop codons included TAA (*ND2*, *ATP8*, *ATP6*, *ND3*, *ND4L*, *CYTB*, and *ND1*), TAG (*ND6*) and non-complete codons T (*COX1*, *COX2*, *COX3* and *ND5*, and *ND4*). The *12S rRNA* and *16S rRNA* were separated by the *tRNA-Val*.

Based on the results, the highest and lowest values of %GC tRNA composition belong to *tRNA-Lys* (46.9%) and *tRNA-Glu* (18.2%), respectively. Secondary structures of *tRNA-Ser*<sub>1</sub> showed the D-loop structure. The longest and shortest tRNAs were *tRNA-Ser*<sub>2</sub> (67 bp) and *tRNA-Ala* (59 bp), respectively.

CR was located between *12S rRNA* and *tRNA-Met*, with an A + T content of 67.58%. The longest gap and overlapping were determined between *tRNA-Gln/tRNA-Cys* (54 bp) and *tRNA-Phe/ND5* (13 bp), respectively.

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Figure 1. Phylogenetic tree showing the relationship among *A. sinica* and three other species from the *Artemia* based on maximum-likelihood (ML) approach. Numbers behind each node denote the bootstrap support values. The GenBank accession numbers are indicated on the right side of species names. *Streptocephalus sirindhornae* was used as an outgroup.

The phylogenetic relationship of *A. sinica* with members of the genus *Artemia* was determined from a concatenated dataset including the 13 PCGs and 2 rRNAs using the software MEGA 7.0.26 v. (Kumar et al. 2016) with 1000 bootstrap replicates and a GTR model (Figure 1). According to the result, *A. sinica* was placed as a clade sister to other Asian spp. All Asian species were clearly separated from American *A. franciscana*.

#### **Disclosure statement**

No potential conflict of interest was reported by the authors.

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