Characterization of the complete mitochondrial genome of *Glyptothorax minimaculatus* and phylogenetic studies of Sisoridae

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**ABSTRACT**

The entire mitochondrial genome (mitogenome) of *Glyptothorax minimaculatus* was sequenced; it spanned 16,536 bp in length and contained 13 protein-coding genes (PCGs), 2 ribosomal RNAs, and 22 transfer RNA genes. A total of 37 genes formed a typical vertebrate mitochondrial gene arrangement. The phylogenetic tree of Sisoridae based on 13 PCGs was constructed and supported that *G. minimaculatus* was closely related to *Glyptothorax sinensis*, *G. zanaensis*, *G. longinema*, *G. granosus* and *G. lanceatus*. The mitogenome of *G. minimaculatus* described in this study provided molecular evidence for its current taxonomic status and laid a groundwork for further study concerning phylogenetics within Sisoridae.

*Glyptothorax minimaculatus* is a species in the family Sisoridae (Chen 2013) and is distributed in the Yingjiang River and Longchuan River in China. The description of this species has mainly focused on current morphology, and the genetic information of this species needs to be established. In this study, we sequenced and characterized the complete mitochondrial genome of *G. minimaculatus* and analyzed its phylogenetic position within Sisoridae. This research was expected to provide evidence for the current taxonomic status and investigate the phylogenetic relationship with other species in Sisoridae.

The specimen was obtained from the Longchuan River (Yunnan Province, China) with geographic coordinates of 25.024°N, 98.677°E and deposited at the zoological Museum of Southwest University under unique voucher number LJ-SCH-2-4-202001004 (the contact person is the corresponding author). The TiANamp genomic DNA Kit (Tiangen Biochemistry Technology Co., Ltd., China) was employed for total genomic DNA extraction from the muscle tissue. After amplification and purification, the DNA libraries were constructed and then sequenced by an Illumina NovaSeq 6000 sequencing platform. Sequenced fragments were assembled to procure the complete mitogenome by using GetOrganelle version 1.7.0+ (Jin et al. 2020) and NOVOPlasty version 4.2 (Dierckxsens et al. 2016). The entire mitogenome was annotated by using MITOS2 software (Bernt et al. 2013). All animal operations in this study were approved and supervised by the Committee of Laboratory Animal Experimentation at Southwest University (Chongqing Province, China) and strictly conformed to the guidelines (protocol number [2014]25) enacted by the committee.

The entire mitochondrial genome of *G. minimaculatus* spanned 16,536 bp in length and exhibited nucleotide contents of 31.17% A, 25.87% T, 15.41% G, and 27.54% C. The mitogenome formed a closed loop with a light (L) strand and a heavy (H) strand and comprised 13 protein-coding genes (PCGs, 11,414 bp in total), 2 ribosomal RNA genes (12S and 16S rRNA, 2,605 bp in total), and 22 transfer RNA (tRNA, 1,564 bp in total) genes. Except for ND6 and 8 tRNAs (Ala, Asn, Cys, Tyr, Ser2, Glu, Pro, Gln), which were located on the L-strand, the residuals were distributed on the H-strand (Miya and Nishida 2000). Furthermore, the original replication on the L-strand and the control region (D-loop) on the H-strand were predicted.

In 13 PCGs, ATG was identified as the canonical start codon for 12 PCGs, with the particular case that the COI gene applied the GTG as its start codon. The use of stop codons was varied as follows: TAA: COI, ATP6, ATP8, ND1, ND4L and ND5; TGA: ND2, ND3 and ND6; TA-: COII; T-: COII, Cytb and ND4. The incomplete stop codons of TA- and T- are ordinarily present in fish mitochondrial genes (Lee et al. 2019). Twenty-two tRNAs distributed between rRNAs and PCGs were detected based on their respective anticodon sequences. The 12S and 16S rRNAs clamped by tRNAVal and interposed by tRNALeu(TAA) and tRNAArg were 957 bp and 1,648 bp, respectively.

A phylogenetic tree based on concatenated sequences of 13 PCGs in 20 fish species was constructed by using the neighbour-joining method with 1000 bootstrap replicates in
the software MEGA 11 (Figure 1) (Tamura et al. 2021). G. minimaculatus was clustered into the Glyptothorax branch and closely related to Glyptothorax sinensis, Glyptothorax zanaensis, Glyptothorax longinemus, Glyptothorax granosus and Glyptothorax lanceatus, with bootstrap probabilities of 100%. The position of G. minimaculatus in the NJ tree supported its current taxonomic status based on morphological features and clearly indicated that it is a valid species.

Author contributions
ZJW and XY conceived and designed the experiments. ZJW contributed the experimental reagents, materials and instruments. HRG and XY collected and identified the specimens. XY and HLG performed the experiments. XY analyzed the data and wrote the manuscript. ZJW approved the final version of the manuscript to be published. All authors agree to be accountable for all aspects of the work.

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

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Data availability statement
The genome sequence data that support the findings of this study are openly available in GenBank of NCBI under the accession no. OK329966 (https://www.ncbi.nlm.nih.gov/nuccore/OK329966). The associated BioProject, SRA, and Bio-Sample numbers are PRJNA786366, SRR17200913, and SAMN23669246, respectively.

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