

Mitochondrial DNA Part B



Resources

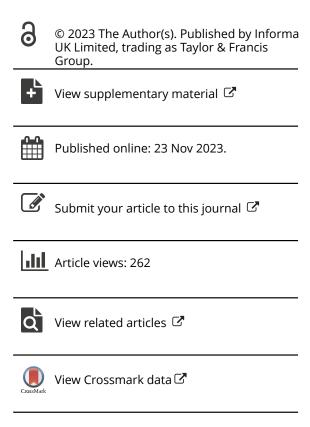
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The complete mitochondrial genome of a new geographical population of freshwater fish *Macropodus hongkongensis* (Freyhof & Herder, 2002)

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MITOGENOME ANNOUNCEMENT



The complete mitochondrial genome of a new geographical population of freshwater fish Macropodus hongkongensis (Freyhof & Herder, 2002)

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ABSTRACT

Macropodus hongkongensis (Freyhof & Herder, 2002), is sparsely distributed in Hong Kong and Guangdong provinces. Recently, a new geographical population of M. hongkongensis was discovered in the Wanquan River in the Hainan province. Therefore, this study focused on sequencing the complete mitochondrial genome of the new geographical population of Macropodus hongkongensis from the Wanquan River. The circular mtDNA molecule was 16,492 bp in size, and the overall base compositions were A (30.30%), C (24.90%), T (29.80%), and G (15.00%), with a slight bias toward A+T. The complete mitogenome encoded 13 protein-coding genes (PCGs), 22 tRNA genes, 2 rRNA genes, and a control region. Phylogenetic analysis indicated that M. hongkongensis of the Hainan Wanquan River was most closely related to the M. hongkongensis of the Gongdong population. These results provide useful genetic information for species identification and phylogenetic studies of Macropodus species.

ARTICLE HISTORY

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KEYWORDS

Macropodus hongkongensis; mitochondrial genome; phylogenetic analysis; new geographical population

Introduction

Macropodus hongkongensis, commonly known as the Hong Kong paradise fish, belonging to the genus Macropodus of the family Belontiida and is sparsely distributed in Hong Kong and Guangdong provinces (Chan et al. 2008; Wang et al. 2009). Recently, a new geographical population of M. hongkongensis was discovered in the Wanguan River in the Hainan province. The known geographical distribution area of M. hongkongensis in Hainan is limited, and its habitat consists of small streams in the intermountain thickets (Shen et al. 2021). In this study, the complete mitogenome of M. hongkongensis of the Wanguan River, a new geographical location, was determined and its phylogenetic relationship was analyzed.

Materials and methods

Specimens of adult M. hongkongensis (Figure 1) were collected from the Wanquan River, Hainan province (19°11'49.36"N, 110°47′60.59 E) and deposited in the Hainan Academy of Ocean and Fisheries (http://www.hnhky.cn/, Qingfeng Zhang, zhangqf@hnhky.cn) under the voucher number HNFF0930101. Total DNA was extracted using an Ezup DNA kit and sequenced using Illumina 6000. The genome was assembled using the GetOrganelle (Jin et al. 2020). The annotated sequence was submitted to GenBank with the accession number OQ630878.1, and the read coverage depth map is shown in Figure S1 (Supplementary material). A phylogenetic tree was constructed



Figure 1. Image of M. hongkongensis from the Hainan Wanguan River, taken by Shen Zhixin.

by software RAxML (Stamatakis 2014), using sequences from Genbank: M. hongkongensis (MN128300.1), M. erythropterus (NC_029946.1), M. opercularis (NC_025932.1), Betta apollon (LC433688.1), Trichopodus pectoralis (KY606170.1), Trichogaster trichopterus (KP100265.1), Colisa fasciata (KP301136.1), Colisa Ialia (AP006039.1), (Liu et al. 2019); Betta splendens (AB571120.1), (Yu et al. 2016); Epinephelus hexagonatus (NC_065829.1), (Wang et al. 2022); Alces alces (JN632595.1), (Hassanin et al. 2012); Centropyge eibli (KT001113.1) (Shen et al. 2016).

Results

The complete circular mitochondrial genome of M. hongkongensis of the new population was 16,492 bp in length (Figure 2)

CONTACT Zhixin Shen 🔯 shen_266@msn.com 🗗 Research Institute of Freshwater Fisheries, Hainan Academy of Ocean and Fisheries Science, Haikou, China Supplemental data for this article can be accessed online at https://doi.org/10.1080/23802359.2023.2278819.

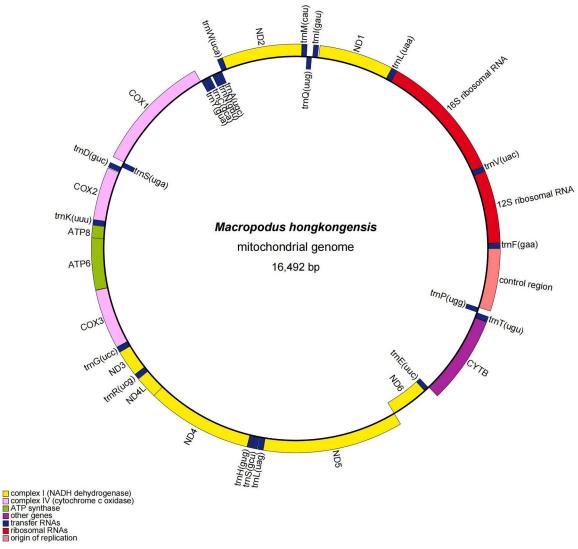


Figure 2. Mitogenome gene map of the M. hongkongensis (OQ630878.1).

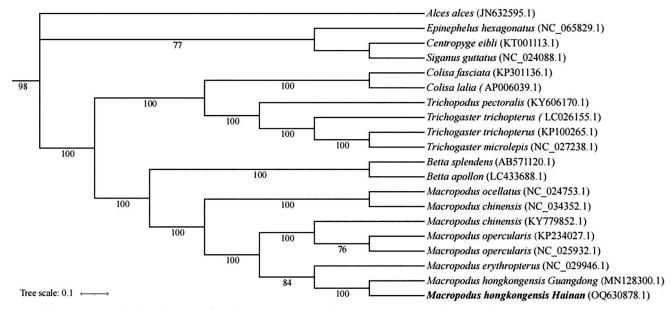


Figure 3. Phylogenetic tree of M. hongkongensis inferred by ML. The position of the Hainan population is shown in bold.

with a base composition of 30.30, 24.90, 29.80, and 15.00% of A, C, T, and G, respectively. The composition showed a slight bias toward A+T content (60.10%), which was consistent with other Macropodus fishes (Xu et al. 2016). The mitogenome contained 13, 22, and 2; protein (PCGs), tRNA and rRNA-coding genes, respectively, along with a control region. All PCGs had an ATG start codon, except COX1 (GTG) and ND6 (CTA). Four PCGs (ND1, ATP6, ATP8, and ND5) terminated with TAA, and eight PCGs (ND2, COX2, COX3, ND3, ND4L, ND4, ND6, and CYTB) ended with the incomplete stop codon TA- or T--. Except for the ND6 and the eight tRNA genes, all other genes were encoded on the heavy strand (H).

Conclusion

In the phylogenetic analysis, M. hongkongensis of the Hainan population was most closely related to the Guangdong population (Figure 3) and the nucleotide sequence divergence of 13 protein coding genes between the two population was 4.0%, smaller than the interspecific distance of Macropodus erythropterus (13.9%). These results offer molecular markers and a reference for conservation research of M. hongkongensis in different populations.

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Author contributions

Zhixin Shen designed the study and planned the experiments. Qingfeng Zhang drafted the manuscript and performed the data analysis. Fangyuan Li and Gaojun Li conceptualized and revised the manuscript. Yang Dong, Ji Wang, and Dikai Chen were responsible for sample collection, data acquisition, and revision of the manuscript. All authors agree to be accountable for all aspects of this study.

Ethical approval

This study did not require ethical approval or permission to collect samples.

Disclosure statement

No potential conflict of interest was reported by the authors.

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Data availability statement

The genome sequence data that support the findings of this study are openly available in GenBank at [https://www.ncbi.nlm.nih.gov] under the accession number: OQ630878.1. The associated **BioProject**, **SRA**, and **Bio-Sample** numbers are PRJNA955001, SRR24305076, and SAMN34163124, respectively.

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