

MITOGENOME ANNOUNCEMENT

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## The complete mitochondrial DNA sequence of Xinjiang arctic grayling *Thymallus arcticus grubei*

Yun-Guo Liu, Yao-Yao Li, Wei Meng and Ling-Xiao Liu

College of Life Sciences and technology, Xinjiang University, Urumqi, China

### ABSTRACT

*Thymallus arcticus grubei* is a specific grayling species that naturally occurs in the Xinjiang Irtysh River basin, China. The complete mitochondrial genome of The Xinjiang arctic grayling *T. arcticus grubei* was first determined (Accession number LC168675), with a total length of 16,658 nucleotides, of which 15,587 nucleotides are coding DNA, and 1071 nucleotides are non-coding DNA. It has the common feature with those of other grayling with respect to genome structure and gene arrangement. It contained 13 protein-coding genes, 2 ribosomal RNA genes, 22 transfer tRNA genes, and 1 control region (D-loop region). The complete mitochondrial genome of Xinjiang arctic grayling *T. arcticus grubei* provides an important data set for further study on its classification.

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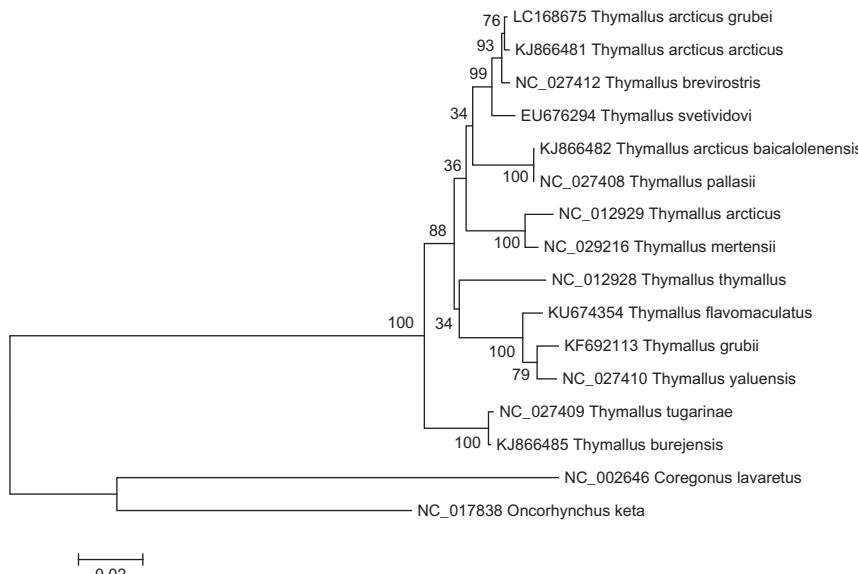
### KEYWORDS

Xinjiang arctic grayling;  
*Thymallus arcticus grubei*;  
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*Thymallus arcticus grubei*, which is classified as *Thymallus*, Salmonidae, Salmoniformes, Protacanthopterygii, Euteleostei is a specific grayling species that naturally occurs in Xinjiang Irtysh River basin, China (Liu et al. 2016). Commonly, *T. arcticus grubei* is classified as one of the subspecies of arctic grayling. However, there is still a lack of molecular data of mitochondrial DNA sequence to prove this opinion. So, it is important to obtain the whole mitogenome of *T. arcticus grubei* for the further study. The specimen of *T. arcticus grubei*, named as Thyag-01 based on the phylogenetic analysis, was collected from Irtysh River, Xinjiang, China (48°12' N, 86°40' E) in 2014 and stored in Xinjiang Fisheries Research Institute, Urumqi, China. Genomic DNA was extracted from Xinjiang arctic grayling muscle according to Liu et al. (2012, 2014). The mitochondrial genome was amplified using 25 pairs of primers, designed based on mitogenome sequence of the arctic grayling, *T. arcticus arcticus* (Accession number KJ866481) and the Mongolian grayling, *T. brevirostris* (Accession number NC\_027412). Protein-coding genes were analyzed by ORF Finder (<http://www.ncbi.nlm.nih.gov/gorf/gorf.html>) using the invertebrate mitochondrial code. The tRNA genes were identified by ARWEN (Laslett & Canback 2008) and tRNA-scan SE (Lowe & Eddy 1997).

The complete *T. arcticus grubei* mitochondrial genome (Accession number LC168675) was 16,658 nucleotides long, of which 15,587 nucleotides are coding DNA, and 1071 nucleotides are non-coding DNA. The mitochondrial genome of *T. arcticus grubei* was found to contain 13 protein-coding genes, 2 ribosomal RNA genes, 22 transfer RNA genes, and 1

control region. Its structure and composition is also comparable to other vertebrates (Yasuike et al. 2010; Liang et al. 2012; Liu et al. 2013; Ma et al. 2015; Xu et al. 2015; Shan & Liu 2015). A phylogenetic tree was constructed based on the comparison of the complete mitochondrial genome sequences with other *Thymallus* species using Neighbour-Joining method (Figure 1). On average, the AT content of the *T. arcticus grubei* mtDNA was 55.18%. To investigate the nucleotide bias, skew for a given strand was calculated as  $(A - T)/(A + T)$  or  $(G - C)/(G + C)$  (Perna & Kocher 1995). The AT and GC skews for the *T. arcticus grubei* mitochondrial genome were 0.015 and -0.225, respectively; this finding indicated that the strand that encoded genes contained more A and C than T and G, and this skew was evidence of codon usage bias. Of the 13 protein-coding genes identified in the *T. arcticus grubei* mitochondrial genome, all but *cox1* used conventional start codons, ATG. The ORFs of six protein-coding genes, *ND1*, *cox1*, *Atp8*, *Atp6*, *nad4L*, and *nad5*, ended with a TAA codon, and only one gene, *nad6*, ended with TAG, respectively. In addition, six genes, *nad2*, *cox2*, *cox3*, *nad3*, *nad4*, and *cytb*, ended with incomplete stop codon T or AT. The 12S rRNA gene was 946-bp long and was flanked by the *tRNA-phe* and *tRNA-val*, while the 16S rRNA was 1679 bp long and was flanked by *tRNA-val* and *tRNA-leu*. The intergenic region between 12S rRNA and 16S rRNA was found to be 72 bp and to contain only one tRNA (*tRNA-val*). The lengths of 22 tRNA coding genes range in size from 67 to 75 nucleotides. The control region was located between *tRNA-pro* and *tRNA-phe* with a length of 1003 bp.



**Figure 1.** A phylogenetic tree constructed based on the comparison of complete mitochondrial genome sequences of the grayling, *T. arcticus grubei* and other thirteen species of *Thymallus* genus. They are *T. thymallus* (European grayling), *T. pallasii* (East Siberian grayling), *T. arcticus* (Arctic grayling), *T. tugariniae* (Lower Amur grayling), *T. burejensis* (Bureya grayling), *T. grubii* (Amur grayling), *T. arcticus arcticus* (Arctic grayling), *T. brevirostris* (Mongolian grayling), *T. yaluensis* (Yalu grayling), *T. mertensii* (Kamchatka grayling), *T. flavomaculatus* (Yellow-spotted grayling), *T. baicalolenensis* (Baikal black grayling), and *T. svetividovi* (Upper Yenisei grayling). *Oncorhynchus keta* and *Coregonus lavaretus* are used as outgroup. Genbank accession numbers for all sequences are listed in the figure. The numbers at the nodes are bootstrap percent probability values based on 1000 replications.

## Disclosure statement

The authors report no conflicts of interest. The authors alone are responsible for the content and writing of the article.

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