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The male and female complete mitochondrial genome sequences of the Endangered freshwater mussel *Potomida littoralis* (Cuvier, 1798) (Bivalvia: Unionidae)

Elsa Froufe¹, Han Ming Gan²,³, Yin Peng Lee²,³, João Carneiro¹, Simone Varandas⁴, Amilcar Teixeira⁵, Alexandra Zieritz⁶, Ronaldo Sousa⁷, and Manuel Lopes-Lima¹

¹CIIMAR/CIMAR – Interdisciplinary Centre of Marine and Environmental Research, University of Porto, Porto, Portugal; ²School of Science, Monash University Malaysia, Selangor, Malaysia; ³Monash University Malaysia Genomics Facility, Monash University Malaysia, Petaling Jaya, Selangor, Malaysia; ⁴CITAB-UTAD – Centre for Research and Technology of Agro-Environment and Biological Sciences, University of Trás-os-Montes and Alto Douro, Vila Real, Portugal; ⁵CIMO-ESA-IPB – Mountain Research Centre, School of Agriculture, Polytechnic Institute of Bragança, Campus de Santa Apolónia, Bragança, Portugal; ⁶Faculty of Science, School of Geography, University of Nottingham Malaysia Campus, Semenyih, Selangor Darul Ehsan, Malaysia, and ⁷CBMA – Centre of Molecular and Environmental Biology, Department of Biology, University of Minho, Campos de Gualtar, Braga, Portugal

Abstract

Freshwater mussels of the family Unionidae exhibit a particular form of mitochondria inheritance called double uniparental inheritance (DUI), in which the mitochondria are inherited by both male and female parents. The (M)ale and (F)emale mitogenomes are highly divergent within species. In the present study, we determine and describe the complete M and F mitogenomes of the Endangered freshwater mussel *Potomida littoralis* (Cuvier, 1798). The complete M and F mitogenomes sequences are 16,451 bp and 15,787 bp in length, respectively. Both F and M have the same gene content: 13 protein-coding genes (PCGs), 22 transfer RNA (trn) and 2 ribosomal RNA (rrn) genes. Bayesian analyses based on the concatenated nucleotide sequences of 12 PCGs and 2 rrn genes of both genomes, including mitogenome sequences available from related species, were performed. Male and Female lineages are monophyletic within the family, but reveal distinct phylogenetic relationships.

The freshwater mussel *Potomida littoralis* (Bivalvia: Unionidae) is an Endangered freshwater mussel with a circum-Mediterranean distribution (Lopes-Lima et al., 2014). Phylogenetic studies within the family are scarce with descriptions of several subfamilies being based primarily on morphological traits (Carter et al., 2011). The inclusion of many genera in these subfamilies is still unclear, mostly due to the lack of sampling coverage and number of available markers (Bogan, 2008). Unionid bivalves, such as several other bivalve families, possess an interesting mitochondrial inheritance, called doubly uniparental inheritance (DUI), in which the male gonad tissue contains (M) mitochondria inherited from the fathers, the male somatic tissue contains (F) mitochondria inherited from the mothers, and females inherit their (F) mitochondria through ordinary single uniparental inheritance (Hoeh et al., 1996). The M and F mitochondrial lineages are highly divergent within each animal and have distinct gene order arrangements (Huang et al., 2013).

In this study, the complete M and F mitogenomes of *P. littoralis* were sequenced, assembled and annotated using an established pipeline (Gan et al., 2014). The M and F mitogenomes have been deposited in GenBank database under the accession numbers KT247375 and KT247374, respectively. The length of the F and M mitogenomes (16,451 bp and 15,787 bp, respectively) of *P. littoralis* sequenced in this study is within the expected range for each gender-specific haplotypes within Unionidae. This is mainly due to the 3' extension of COX2, common to all freshwater mussel male mitogenomes. Both haplotypes have the same gene content: 13 protein-coding genes (PCGs), 22 transfer RNA (trn) and 2 ribosomal RNA (rrn) genes involved in mitochondrial transcription and translation processes. Regarding the gene orientation, again both have the same genes (4 PCGs, 20 tRNAs and 2 rRNAs) encoded on the heavy strand and the remaining (9 PCGs and 2 rRNAs) encoded on the complementary strand.

Additional mitogenome sequences (M and F) available from related species were downloaded from GenBank. Each gene sequence was aligned using GUIDANCE (version 1.5, Penn et al., 2010) with the MAFFT multiple sequence alignment algorithm (version 7, Katoh & Standley, 2013). To build our single gene alignments, we used the following GUIDANCE parameters: score algorithm: GUIDANCE; bootstraps replicates: 100; Sequence cut-off score: 0.0 (no sequences removed); Column cut-off score: below 0.8. The final concatenated data set included 12 mitochondrial protein-coding genes (excluding the ATP8 due to its extreme length variation) and the 2 rRNA genes of both genomes. To infer the phylogenetic relationships among these sequences with Bayesian methodology, we used MrBayes v3.2.1 (Ronquist et al., 2012). The alignment (12,084 bp) was partitioned according to the best scheme determined by JmodelTest2
(GTR + I + G, HKY + I + G and HKY + G models were used). Each chain started with a randomly generated tree and ran for $1 \times 10^6$ generations with a sampling frequency of 1 tree for every 100 generations. The resultant trees, after discarding the first 25% as burn-in, were combined in a 50% majority rule consensus tree. The final BI tree was rooted at the split between Male and Female haplotypes (based on previous studies, e.g. Huang et al., 2013).

As expected, the obtained phylogenetic tree (Figure 1) revealed two main clades divided between the monophyletic M and F mitogenomes clades. However, the three sub-families (for which GenBank sequences are available) revealed distinct phylogenetic relationships. Although a smaller number of M genomes are available, relationships are very well resolved and supported only for M genomes but not for F genomes. In the M-clade, the Gonideinae and Ambleminae form a sister lineage and then sister to the Anodontinae, whereas in the F-clade, the relationships of these subfamilies are not well resolved. The newly sequenced *P. littoralis* genomes cluster inside the Gonideinae with *Pronodularia japonensis* in the M-clade, and with a clade including *P. japonensis* and *Lamprotula leai* in the F-clade.

![Figure 1. Bayesian phylogenetic tree of 20 Unionidae Male and Female mitogenomes sequences based on concatenated nucleotide sequences of 12 mitochondrial protein-coding genes (excluding the ATP8 gene) and the 2 rRNA genes. The numbers behind the species names are the GenBank accession numbers and those at the nodes indicate the posterior probability.](image)

**Declaration of interest**

The authors report that they have no conflicts of interest. The authors alone are responsible for the content and writing of the paper. This work was financially supported by the Portuguese Foundation for Science and Technology (FCT) project PTDC/AAC-AMB/ 117688/2010, and partly by the Strategic Funding UID/Multi/04423/2013 through national funds provided by FCT and European Regional Development Fund (ERDF), in the framework of the programme PT2020. JC is supported by Portuguese Foundation for Science and Technology (FCT) post-doc fellowship with reference SFRH/BPD/100912/2014. HMG and YPL are supported by the Monash University Malaysia Tropical Medicine Biology Platform.

**References**


