

Complete mitochondrial genome of the hard clam (*Mercenaria mercenaria*)

Zhi Hu^{a,b,c*}, Hao Song^{a,b*}, Cong Zhou^{a,b,c}, Zheng-Lin Yu^{a,b}, Mei-Jie Yang^{a,b,c} and Tao Zhang^{a,b,d}

^aCAS Key Laboratory of Marine Ecology and Environmental Sciences, Institute of Oceanology, Chinese Academy of Sciences, Qingdao, Shandong, China; ^bLaboratory for Marine Ecology and Environmental Science, Qingdao Pilot National Laboratory for Marine Science and Technology, Qingdao, Shandong, China; ^cUniversity of Chinese Academy of Sciences, Beijing, China; ^dCenter for Ocean Mega-Science, Chinese Academy of Sciences, Qingdao, Shandong, China

ABSTRACT

The hard clam (*Mercenaria mercenaria*) is an important economic and ecological bivalve. In this study, the mitochondrial genome was sequenced. The sequenced genome size was 18,360 bp. The nucleotide composition was asymmetric with a AT bias. Mitogenome contained 13 protein-coding genes (PCGs), 2 rRNA genes, and 22 tRNA genes. Of 13 PCGs, 3 genes (*cox3*, *nad3*, and *cox2*) had incomplete stop codons. Furthermore, phylogenetic analysis using 12 PCGs (except *atp8*) figured out that *M. mercenaria* was closely related to genus *Dosinia*. The complete mitogenome of *M. mercenaria* provides essential information for further phylogenetic and evolutionary analysis in Veneridae.

ARTICLE HISTORY

Received 2 August 2019

Accepted 13 October 2019

KEYWORDS

Mercenaria mercenaria; Veneridae; mitogenome

Veneridae presumably has more than 800 species and is a large and diverse family of Bivalvia (Mikkelsen et al. 2006), many of which are commercially important in benthic communities (Canapa et al. 2003). The hard clam (*Mercenaria mercenaria*) is a native kind of bivalve on the east coast of the US and Canada (Menzel 2010). Fu-Sui Zhang imported hard clams from America to China in 1997. Because of its strong life, it became an important cultured bivalve in China. The mitogenome has been suggested to be a good source for molecular biology because of rapid evolutionary rate and lack of recombination (He et al. 2011). In this paper, we report the complete mitochondrial genome of the hard clam *M. mercenaria* to better understand the phylogenetic position within the Veneridae family.

The adult hard clam was collected from Dongying, Shandong province, China (37.25 N 118.55 E) and then cultured by the Mashan Group Co., Ltd. in Shandong Province. During culture, the hard clam was acclimated to the seawater (25 °C, 30‰ salinity) under continuous aeration and fed with *Isochrysis galbana*. The adductor muscle was sequenced. Specimen (Collection Number: MBM286619) was deposited in the museum of Institute of Oceanology, Chinese Academy of Sciences. The complete mitogenome of *M. mercenaria* was 18,360 bp in length (GeneBank Accession: MN233789), which was within the range of genome sizes for already sequenced molluscan mitogenomes. The mitogenome size was similar to *Dosinia* clam and length differences were mostly due to the

size variations of the non-coding region (Lv et al. 2018). The overall base composition was 27.17% A, 21.62% G, 9.44% C, and 41.77% T. The GC content was 31.06%. The mitogenome contained 13 protein-coding genes (PCGs), 2 ribosomal RNA genes, 22 transfer RNA genes, and a control region. *Atp8* gene was found in the mitogenome, which has been reported as missing in several bivalve species (He et al. 2011). Six PCGs (*cox3*, *nad2*, *nad4l*, *nad4*, *atp6*, and *cox2*) used ATG as initiation codon. Four PCGs (*nad1*, *cob*, *nad3*, and *nad5*) used ATT as an initiation codon. *nad6* and *cox1* used ATA as initiation codon and *atp8* used ATC as initiation codon. Most PCGs used TAG or TAA as stop codon. Incomplete stop codons in *cox3*, *nad3*, and *cox2* were found as T(aa) and TA(a), these incomplete termination codons might be completed as TAA by post-transcriptional polyadenylation (Ojala et al. 1981).

The other 14 clams of family Veneridae mitogenome sequences in public were used in phylogenetic analysis. *Solen grandis* and *Solen strictus* were used as outgroups. We used MEGA 7 (Kumar et al. 2016) to construct the phylogenetic relationships of the *M. mercenaria* and related Veneridae by neighbour-joining method with 1000 bootstrap replicates based on the 12 PCGs (except *atp8*) (Figure 1). The resultant phylogenetic tree indicated that *M. mercenaria* was closely related to genus *Dosinia*. The complete mitogenome of *M. mercenaria* provides essential information for further phylogenetic and evolutionary analysis in Veneridae.

CONTACT Tao Zhang  tzhang@qdio.ac.cn  Institute of Oceanology, Chinese Academy of Sciences, 7 Nanhai Road, Qingdao 266071, Shandong, China

*These authors contributed equally to the work.

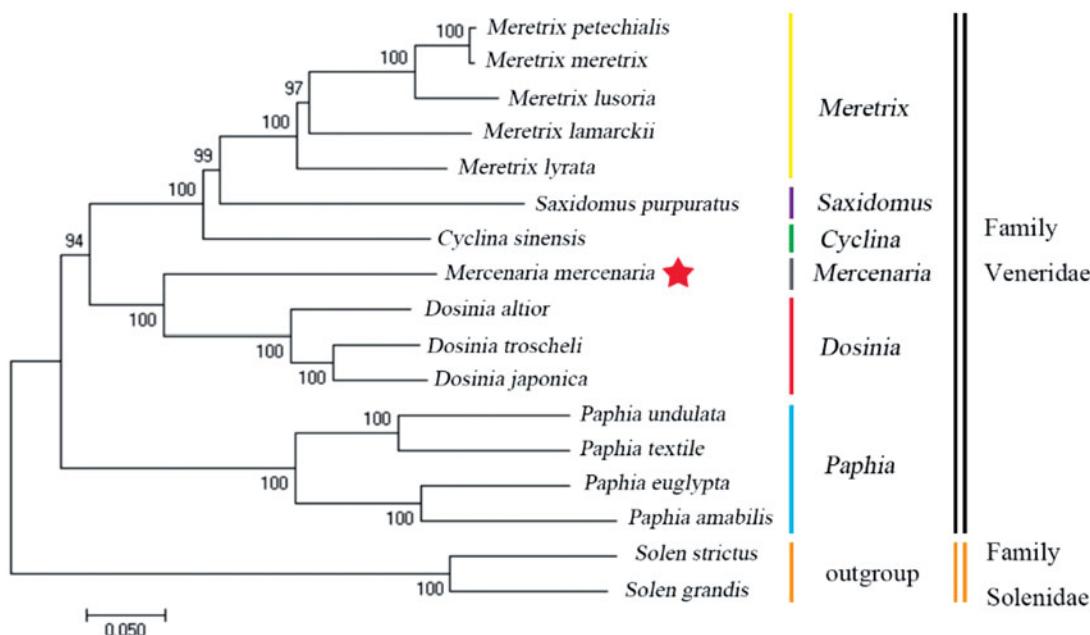


Figure 1. Phylogenetic trees based on the nucleotide sequences of 12 concatenated protein-coding genes. The numbers in the phylogenetic tree are the bootstrap probability values. Vertical stick indicates genus, double vertical sticks indicates families. The genome sequence in this study is labelled with a red star. GenBank accession numbers of the sequences were used for the tree as follows: *Meretrix petechialis* (EU145977); *Meretrix meretrix* (GQ463598); *Meretrix lusoria* (GQ903339); *Meretrix lamarckii* (GU071281); *Meretrix lyrata* (KC832317); *Saxidomus purpuratus* (KP419933); *Cyclina sinensis* (KU097333); *Mercenaria mercenaria* ★; *Dosinia altior* (MG543473); *Dosinia troscheli* (MG543474); *Dosinia japonica* (MF401432); *Paphia undulata* (JF969278); *Paphia textile* (JF969277); *Paphia euglypta* (GU269271); *Paphia amabilis* (JF969276); *Solen strictus* (JN786377); *Solen grandis* (HQ703012).

Disclosure statement

The authors declare no conflicts of interest.

Funding

This research was supported by the earmarked fund for the Modern Agro-industry Technology Research System (CARS-49), Major Applied Technology Innovation Project in Agriculture of Shandong Province [SF1405303301], supporting programme of Science and Technology Service Network Initiative in Fujian Province: the selection breeding of the hard clam, large-scale seed cultivation and high efficiency ecological culture technology in pond (2017T3014), the Industry Leading Talents Project of Taishan Scholars (Recipient: Tao Zhang), the 'Double Hundred' Blue Industry Leader Team of Yantai (Recipient: Tao Zhang), and the Creative Team Project of the Laboratory for Marine Ecology and Environmental Science, Qingdao Pilot National Laboratory for Marine Science and Technology [LMEES-CTSP-2018-1]. The funders had no role in the study design, data collection and analysis, decision to publish, or preparation of the manuscript.

References

- Canapa A, Schiaparelli S, Marota I, Barucca M. 2003. Molecular data from the 16S rRNA gene for the phylogeny of Veneridae (Mollusca: Bivalvia). *Mar Biol*. 142:1125–1130.
- He CB, Wang J, Gao XG, Song WT, Li HJ, Li YF, Liu WD, Su H. 2011. The complete mitochondrial genome of the hard clam *Meretrix meretrix*. *Mol Biol Rep*. 38:3401–3409.
- Kumar S, Stecher G, Tamura K. 2016. MEGA7: Molecular Evolutionary Genetics Analysis version 7.0 for bigger datasets. *Mol Biol Evol*. 33: 1870–1874.
- Lv CD, Li Q, Kong LF. 2018. Comparative analyses of the complete mitochondrial genomes of *Dosinia* clams and their phylogenetic position within Veneridae. *PLoS One*. 13:e0196466.
- Menzel RW. 2010. Quahog clams and their possible mariculture. *J World Aquacult Soc*. 2:21–36.
- Mikkelsen PM, Bieler R, Kappner I, Rawlings TA. 2006. Phylogeny of Veneroidea (Mollusca: Bivalvia) based on morphology and molecules. *Zool J Linn Soc*. 148:439–521.
- Ojala D, Montoya J, Attardi G. 1981. TRNA punctuation model of RNA processing in human mitochondrial. *Nature*. 290:470.