Complete mitochondrial genome of the apple snail *Pomacea diffusa* (Gastropoda, Ampullariidae) with phylogenetic consideration

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**ABSTRACT**

We present the complete mitochondrial genome of *Pomacea diffusa* in this study. The results show that the mitochondrial genome is 16,640 bp in length, which is comprised of 13 protein-coding genes, two rRNA genes, and 21 tRNA genes. The nucleotide compositions of the light strand are 39.62% of A, 30.13% of T, 16.02% of C, and 14.24% of G. Except eight tRNA genes (Glu, Gly, Trp, Cys, Tyr, Met, Thr, Val) on the light strand, the rest are encoded on the heavy strand. All the protein-coding genes start with ATC initiation codon, and two types of inferred termination codons are TAA and TAG. There are 26 intergenic spacers and two gene overlaps. The phylogenetic analysis shows that *P. diffusa* clusters with *P. canaliculata* and *P. maculata* with high bootstrap support, which is consistent with the morphological and molecular evidence.

The apple snail *Pomacea diffusa* (Blume 1857) originated in South America and was introduced to Oceania, North America and Asia with aquarium trade (Hayes et al. 2008; Horgan et al. 2014). Unlike its congeneric invasive species *P. canaliculata* (Lamarck 1822) and *P. maculata* (Perry 1810), *P. diffusa* feeds mostly on algae and has little direct impact on macrophytes (Morrison and Hay 2011). But the hybrids between *P. diffusa* and other *Pomacea* species might pose serious threats to paddy plants and gastropod fauna (Aditya and Raut 2001a, 2001b; Morrison and Hay 2011).

Considering that *P. diffusa* was often confused with *P. bridgesii* (Reeve 1856) in the morphological identification (Hayes et al. 2008; Horgan et al. 2014) and few studies focused on *P. diffusa*, we sequenced the complete mitochondrial genome of *P. diffusa* by using the next-generation sequencing (NGS) techniques for its population genetics and polymorphism studies, which also helps to understand its introducing process and seek effective management strategy. The specimen was purchased from Yuehe Pet Market in Guangzhou (23°09′ N, 113°24′ E) and confirmed as *P. diffusa* based on morphological characteristics and molecular identification (Rawlings et al. 2007; Hayes et al. 2008). The pleopod muscle of *P. diffusa* was preserved in 95% ethanol and stored at −40°C in South China Agricultural University, Guangzhou, China. The procedure referred from Green and Sambrook (2012) was carried out in the total genomic DNA extraction.

The complete mitochondrial genome of *P. diffusa* presented in this study (GenBank accession number KY008698) is 16,640 bp in length, containing 13 protein-coding genes, two ribosomal RNA genes (L-rRNA and S-rRNA), 21 transfer RNA genes (tRNA). All of them are encoded on the heavy strand except eight tRNA genes (Glu, Gly, Trp, Cys, Tyr, Met, Thr, Val). Among 13 protein-coding genes (total 11,268 bp) encoding 3743 amino acids, the maximum is ND5 with 1728 bp and the minimum is ATP8 with only 159 bp. The S-rRNA and L-rRNA genes are 950 and 1345 bp, respectively, located between the tRNA^{Glu} and tRNA^{Aeu} genes and separated by the tRNA^{Val} gene. The inner ring indicates the GC percent varying from 22.90% to 37.60%. The overall nucleotide compositions of the light strand in descending order are 39.62% of A, 30.13% of T, 16.02% of C, and 14.24% of G. The most representative base is A, and the bias against G is observed. The absence of D-loop is consistent with the Gastropoda (Zeng et al. 2015; Yang et al. 2016a, 2016b, 2016c), but, at least, one lengthy non-coding region is an essential regulatory element for the initiation of transcription and replication (Wolstenholme 1992).

All the protein-coding genes start with ATC initiation codon, and two types of inferred termination codons are TAA (ATP8, ATP6, ND1, ND6, ND4L, ND4, ND5, COX3, ND3, ND2) and TAG (COX1, COX2, CYTB). Twenty-one tRNA genes vary from 62 to 72 bp in length, and all fold into the typical cloverleaf secondary structure. There are 26 intergenic spacers (total 1684 bp) varying from 1 to 1073 bp in length and two gene overlaps (total 27 bp), the larger of which is 20 bp between the ND5 and tRNA^{The} genes. The tandem repeat...
sequences are observed in inter genetic space of tRNA$^{\text{Phe}}$ (GAA) and COX3 genes.

A phylogenetic tree is constructed using the maximum-likelihood method based on the complete mitochondrial genomes of the closely related 17 Gastropoda species to assess their actual phylogenetic relationship and evolution (Figure 1). *P. diffusa* clusters with *P. canaliculata* and *P. maculata* with high bootstrap support. The phylogenetic analysis is consistent with morphological and molecular evidence, indicating that *P. diffusa* has higher homology with *P. canaliculata* and *P. maculata* (80.08% and 80.45% nucleotide sequence identity, respectively) than other snail species (Figure 1; Rawlings et al. 2007; Hayes et al. 2008).

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**Disclosure statement**

The authors report no conflicts of interest. The authors themselves are responsible for the content and writing of the paper.

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**References**

Aditya G, Raut SK. 2001a. Food of the snail, *Pomacea bridgesi*, introduced in India. Curr Sci India. 80:919–921.

Aditya G, Raut SK. 2001b. Predation of water bug *Sphaerodema rusticum* fabricius on the snail *Pomacea bridgesi* (Reeve), introduced in India. Curr Sci India. 81:1413–1414.
Green MR, Sambrook J. 2012. Molecular cloning: a laboratory manual. 4th ed. New York (NY): Cold Spring Harbor Laboratory Press.

Hayes KA, Joshi RC, Thiengo SC, Cowie RH. 2008. Out of South America: multiple origins of non-native apple snails in Asia. Divers Distrib. 14:701–712.

Horgan FG, Stuart AM, Kudavidanage EP. 2014. Impact of invasive apple snails on the functioning and services of natural and managed wetlands. Acta Oecol. 54:90–100.

Morrison WE, Hay ME. 2011. Feeding and growth of native, invasive and non-invasive alien apple snails (Ampullariidae) in the United States: invasives eat more and grow more. Biol Invasions. 13:945–955.

Rawlings TA, Hayes KA, Cowie RH, Collins TM. 2007. The identity, distribution, and impacts of non-native apple snails in the continental United States. BMC Evol Biol. 7:97.

Wolstenholme DR. 1992. Animal mitochondrial DNA: structure and evolution. Int Rev Cytol. 141:173–216.

Yang H, Zhang JE, Deng Z, Luo H, Guo J, He S, Luo M, Zhao B. 2016a. The complete mitochondrial genome of the golden apple snail Pomacea canaliculata (Gastropoda: Ampullariidae). Mitochondrial DNA B. 1:45–47.

Yang H, Zhang JE, Guo J, Deng Z, Luo H, Luo M, Zhao B. 2016b. The complete mitochondrial genome of the giant African snail Achatina fulica (Mollusca: Achatinidae). Mitochondrial DNA A. 27:1622–1624.

Yang H, Zhang JE, Luo H, Luo M, Guo J, Deng Z, Zhao B. 2016c. The complete mitochondrial genome of the mudsnail Cipangopaludina cathayensis (Gastropoda: Viviparidae). Mitochondrial DNA A. 27:1892–1894.

Zeng T, Yin W, Xia R, Fu C, Jin B. 2015. Complete mitochondrial genome of a freshwater snail, Semisulcospira libertina (Cerithioidea: Semisulcospiridae). Mitochondrial DNA. 26:897–898.