Complete mitochondrial genome of a sea cucumber, *Euapta godeffroyi* (Echinodermata, Holothuroidea, Apodida, Synaptidae)

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**ABSTRACT**

We determined the complete mitochondrial genome sequence of a holothurian *Euapta godeffroyi* belonging to the order Apodida. The complete mitogenome of *E. godeffroyi* was 16,410 bp in length and consisted of 13 protein-coding genes (PCGs), two ribosomal RNA genes, and 22 transfer RNA genes. The orders of PCGs and rRNAs did not match those of any recorded holothurian mitogenomes. The maximum likelihood (ML) phylogenetic tree placed *E. godeffroyi* as the sister group to chirodoid species and supported the monophyly of the order Apodida.

**Keywords**

Echinodermata; sea cucumber; complete mitogenome; Synaptidae

*Euapta godeffroyi* (Semper, 1868) (Figure 1) is a common sea cucumber in coral reefs in the tropical Indo-Pacific region and shows geographically large distribution in the tropical shallow waters from the west Indian Ocean to the east Pacific Ocean (e.g., Massin 1999). Apodid holothurians including *E. godeffroyi* are a stem group within the class Holothuroidea (Kerr and Kim 2001; Miller et al. 2017). Miller et al. (2017) confirmed the monophyly of the order Apodida, and that apodid holothurians are divided into two clades in the order Apodida: (1) family Myriotrochidae and (2) other apodid holothurians. In the later clade, however, the family Synaptidae including *E. godeffroyi* did not form a monophyletic clade. A part of synaptid species, *Leptosynapta clarki* Heding, 1928, was placed in Chirodidae. Their molecular phylogenetic study of Miller et al. (2017) used six mitochondrial/nuclear gene markers, covering all common gene markers for Echinodermata, but their study included limited OTUs from apodid holothurians, and the relationships between *L. clarki* and chirodoid species lacked strong nodal support. Therefore, the order Apodida requires the phylogenetic and systematic revision using further genetic markers and OTUs, and 13 protein-coding genes of mitogenome are expected to be effective in revising the phylogenetic relationship of apodid holothurians. There has been no mitogenome record for the family Synaptidae in Apodida, and we choose *E. godeffroyi* as representative of the family. We determined the whole mitogenome sequence by a shotgun sequencing method.

The specimen was collected in 2019 from North Point, Nyaung Oo Phee Island, Myanmar, facing the Andaman Sea (10.087 N 97.963 E), at the depth of approximately 22 m. Total DNA was extracted using DNeasy Blood & Tissue Kit (QIAGEN) and processed using the Collibri\textsuperscript{TM} PS DNA Library Prep Kits for Illumina Systems (Invitrogen). Paired-end sequencing (300 cycles) was conducted using HiSeqX (Illumina) of Macrogen Japan Corp., with 150 bp read length and additional inserts of ca. 100 bp for a total of approximate 16 million reads. Assembly was performed using CLC Genomics Workbench ver. 12 (QIAGEN) with the default setting. Gene identification was made using the MITOS web server (Bernt et al. 2013). The voucher specimen is deposited in the National Museum of Nature and Science, Tsukuba, Japan (NSMT E-13912).

The mitogenome of *E. godeffroyi* (GenBank/DDJB/EMBL accession number LC704718) is 16,410 bp long and encodes 13 proteins, two rRNAs, and 22 tRNAs for a total of 37 gene products. The overall A + T content is 64.0%, which is slightly lower than that of other apodid species (Sun et al. 2021). Unlike other sea cucumber mitogenomes, *COX1*, *ND4L*, and *ND1* start with GTG codon. All other protein-coding genes (PCGs) start with the ATG start codon. Eight of PCGs stop with the termination codon TAG, other five of PCGs (*CYTB*, *ND2*, *ND3*, *ND4*, and *ATP6*) end with TAA codon. Although previously three variations of the gene orders of 13 PCGs and rRNAs were known in the class Holothuroidea (Sun et al. 2021), the gene order of *E. godeffroyi* did not match those of...
any recorded holothurian mitogenomes. The gene order of *E. godeffroyi* shows unique arrangement with COX3 following after COX1, and ND3 located next to ND6 (Figure 1).

The maximum-likelihood phylogenetic analysis (ML) based on 13 PCGs of *E. godeffroyi* and 16 other holothurians (Scouras et al. 2004; Sun et al. 2010, 2020; Fan et al. 2011; Mu et al. 2018; Takano et al. 2019; Yang et al. 2019, 2020; Liao et al. 2020; Zeng et al. 2020; Figueroa et al. 2021; three other direct submitted sequences in GenBank), and of two echinoids (Jacobs et al. 1988; another direct submitted sequence in GenBank) as outgroup, was conducted using RAxML-NG ver.1.0.2 (Kozlov et al. 2019) with bootstrap analyses of 1000 replicates (Figure 1). PartitionFinder 2.1.1 (Lanfear et al. 2017) was used to determine the best partitioning scheme and the substitution model with branch lengths linked and a greedy search algorithm (Lanfear et al. 2012). The optimal partitioning strategy and evolutionary models consisted of thirteen genes data set for ML analyses was as follows; partition 1 (COX1), partition 2 (COX2), partition 3 (COX3), partition 4 (CYTB), partition 5 (ND1), partition 6 (ND2 and ND4), partition 7 (ND3, ND4L, and ATP6), partition 8 (ND5), and partition 9 (ATP8) with GTR + I + G; partition 9 (ND6) with K81UF + I + G. Three apodids, *E. godeffroyi*, *Chiridota heheva* and *Chiridota* sp., form a monophyletic clade, and *E. godeffroyi* is a sister taxon to the other Apodida (*C. heheva* and *Chiridota* sp.) with high nodal support (Figure 1). This relationship between the three apodid species in our tree was consistent with the result of Miller et al. (2017). Although additional OTUs are needed, this mitogenome will be useful for reconstructing higher systematics of Holothuroidea phylogeny.

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Ethics statement

All applicable international, national, and/or institutional guidelines for the care and use of animals were followed.

Author’s contributions

All authors conceived the study. MMA and TF contributed to the application for sampling permits and conducted the voucher specimen collection and the DNA sample preparation in Myanmar. AO and SFH performed DNA analysis in the laboratory. All authors were involved in the interpretation of the data; the drafting of the paper, revising it critically for intellectual content; and the final approval of the version to be published; and that all authors agree to be accountable for all aspects of the work.

Disclosure statement

No potential conflict of interest was reported by the author(s).

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Data availability statement

The consensus genome sequence is openly available in GenBank of NCBI at [https://www.ncbi.nlm.nih.gov] (https://www.ncbi.nlm.nih.gov) under the accession no. LC704718. The associated BioProject, SRA, and BioSample numbers are PRJDB13488, DRR361674, and SAMD00467977 respectively. The voucher specimen is deposited at the National Museum of Nature and Science, Tsukuba, Japan (Toshihiko Fujita; fujita@kahaku.go.jp) under the catalog number NSMT E-13912.

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