

[단보, Short communication]

Full-length mitochondrial genome of the boring polychaete species, *Polydora hoplura* (Annelida) isolated from abalone, *Haliotis discus hannai* shells

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ABSTRACT

The mitochondrial genome (mitogenome) of the invasive and harmful polychaete species, *Polydora hoplura* (Annelida), was analyzed by next generation sequencing (NGS). Its mitogenome was found to be 17,597 bp in length, comprising 13 protein-coding genes, two ribosomal RNA genes, and 23 transfer RNA genes, with one additional *trnM* gene. The gene composition and order of *P. hoplura* were distinct from those of polychaetes and even differed from those of *Marenzelleria neglecta* belonging to the same family, Spionidae.

Key words: Invasive species, harmful species, mitochondrial genome, polychaete, *Polydora hoplura*

The polychaete *Polydora hoplura* (Annelida) is an invasive and harmful species that has been distributed in new regions by the anthropogenic transportation of live imported shellfish worldwide (Radashevsky and Migotto 2017). It is one of the most notorious pests that bores into mollusk shells (Leonart *et al.* 2003; Boonzaaier *et al.* 2014), and has caused serious damage to abalone and oysters on shellfish farms along the southern coast of South Korea since first reported in 2004 as *P. uncinata* (Radashevsky *et al.* 2017; Sato-Okoshi *et al.* 2012).

A voucher specimen of *P. hoplura* I-13 (NIFS Lot No. Ab-WB-2019-004) was collected from an abalone farm on the southwestern coast of South Korea (34°16'24"N, 126°62'89"E) and deposited in the

invertebrate collection of the National Institute of Fisheries Science (NIFS, Busan, South Korea). Its genomic DNA (gDNA) was extracted from a piece of excised tissue according to Asahida *et al.* (1996). For next generation sequencing (NGS), a library was constructed with the gDNA according to the manufacturer's instruction of MGIEasy Universal DNA Library Prep Set (MGI Tech Co. Ltd., Shenzhen, China). A library with an average insert size of ca. 420 bp was generated and subjected to a NGS analysis using Genetic Sequence MEISEQ-2000 (MGI Tech Co. Ltd.). Approximately 11 gigabyte (Gb) of raw data were obtained by 150 bp paired-end sequencing. The raw data were trimmed with cutadapt v1.9.1 (Martin 2015) and subjected to de novo assembly and annotation using Geneious Prime v2020.2 (Biomatters Ltd, Auckland, New Zealand). The gene composition and order of *P. poplura* were further confirmed through online MITOS WebServer (<http://mitos.bioinf.uni-leipzig.de/index.py>). The full-length mitogenomic sequence analyzed in this study was deposited in the GenBank under the accession number MW248165.

In this study, the mitogenome of *P. hoplura* was a circular molecule of 17,597 bp in total length. It

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