

Complete mitochondrial genome of *Micractinium pusillum* CCAP 231/1 (Chlorellaceae, Trebouxiophyceae)

Nam Seon Kang^{a*}, Seung-Woo Jo^{b,c*}, Jung A Lee^a, Kyeong Mi Kim^a, Hyeong Seok Jang^a, Eun Song Kim^a, Moongeun Yoon^a, Ji Won Hong^a and Ho-Sung Yoon^{b,c}

^aDepartment of Taxonomy and Systematics, National Marine Biodiversity Institute of Korea, Seocheon, Republic of Korea; ^bDepartment of Energy Science, Kyungpook National University, Daegu, Republic of Korea; ^cSchool of Life Sciences, BK21 Plus KNU Creative BioResearch Group, Kyungpook National University, Daegu, Republic of Korea

ABSTRACT

The mitochondrial genome of *Micractinium pusillum* CCAP 231/1 was completely sequenced. This mitogenome has 70,061 bp in length and consists of 62 genes including 32 protein-coding, 3 rRNA, and 27 tRNA genes. The overall GC content of the genome is 31.3%.

ARTICLE HISTORY

Received 7 November 2019
Accepted 23 November 2019

KEYWORDS

Complete mitochondrial genome; *Micractinium pusillum*; Chlorellaceae; Trebouxiophyceae

Micractinium Fresenius is a globally distributed genus in a variety of aquatic habitats. There are currently 16 taxonomically accepted species (Guiry and Guiry 2019). *Micractinium pusillum* Fresenius (Chlorellaceae, Trebouxiophyceae) is a holotype species of the genus and this species is known to be a cosmopolitan microalga commonly found in many types of freshwater environments (John et al. 2002). It is often considered as an indicator of nutrient-enriched water. In this study, the complete mitogenome of *M. pusillum* CCAP 231/1 was determined for the first time.

Micractinium pusillum CCAP 231/1 was purchased from the Culture Collection of Algae and Protozoa (CCAP). This strain was originally isolated from Wicken Lode, Cambridgeshire, England ($52^{\circ}18' 24.29''\text{N}$, $0^{\circ}16'32.86''\text{E}$). The culture was inoculated onto fresh BG-11 agar medium (UTEX, USA) containing imipenem (Sigma-Aldrich, St. Louis, USA) at a concentration of $100\text{ }\mu\text{g ml}^{-1}$ to suppress bacterial growth (Hong et al. 2015). A resulting single colony on the agar plate was then aseptically transferred to a liquid BG-11 medium and grown at 18°C under cool fluorescent light (approximately $40\text{ }\mu\text{mole m}^{-2}\text{ s}^{-1}$) in a light:dark cycle (14:10 hrs) for 4 weeks. Microalgal biomass was harvested by centrifugation at centrifugation at $2063 \times g$ (1580R; Labogene, Daejeon, Korea). Whole genomic DNA was extracted from the sample using a DNeasy Plant Mini Kit (Qiagen, Hilden, Germany) followed by preparation of a library using an MGIEasy DNA Library Prep Kit

V1 (BGI, Shenzhen, China) according to the manufacturer's instruction. Whole genome sequencing was performed using BGISEQ-500 (BGI, China) sequencer and raw data were filtered to obtain $>10\text{ Gb}$ clean data per each sample. *De novo* mitogenome assembly was carried out using NOVOPlasty v3.6 software (Dierckxsens et al. 2017). The size of the circular mitogenome produced is 70,061 bp (GenBank accession number MN649871) which is smaller than that of the previously reported *M. conductrix* mitogenomes (74,708 bp, GenBank accession number KY629619). The nucleotide composition is 34.6% A, 34.1% T, 15.2% G, and 16.0% C. The overall GC content is 31.3%. The *M. pusillum* mitogenome contains 62 genes, including 32 predicted protein-coding, 3 rRNA, and 27 tRNA genes. Among 32 genes, 19 genes were revealed as complete protein-coding genes, which all of them were started with ATG as a start codon and ended by TAA as a stop codon. It was found that there was only 1 case of gene-overlapping with a size of 85 bp and 27 tRNA genes ranged from 31 to 86 bp in length.

Phylogenetic analysis was carried out by PhyML 3.0 with 11 reported mitogenome sequences (Fan et al. 2017) belonging to the Trebouxiophyceae family and it was visualized by FigTree v1.4.4. The result showed the phylogenetic position of *M. pusillum* CCAP 231/1 (Figure 1) within this family. This new information would contribute to a better understanding of the phylogenetic relationships of the *Micractinium* species

CONTACT Ji Won Hong  jwhong@mabik.re.kr  Department of Taxonomy and Systematics, National Marine Biodiversity Institute of Korea, Seocheon, Republic of Korea; Ho-Sung Yoon  hsy@knu.ac.kr  Department of Energy Science, Kyungpook National University, Daegu, Republic of Korea.

*These authors have contributed equally to this work.

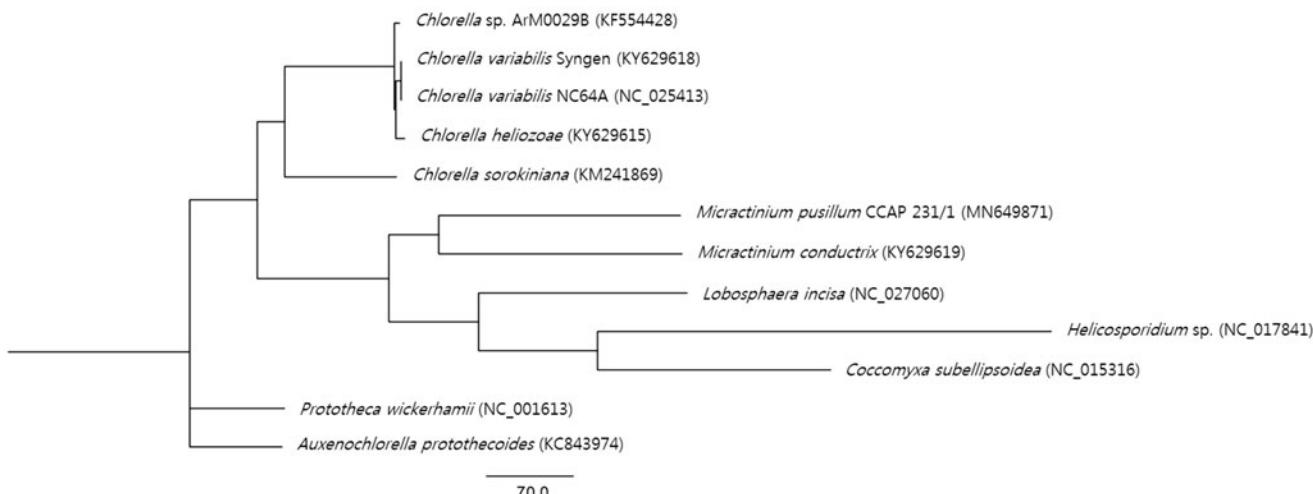


Figure 1. Maximum-likelihood phylogenetic tree of *M. pusillum* CCAP 231/1 and 11 other species. GenBank accession numbers were indicated in the parentheses.

and mitochondrial genome diversity and evolution in the Trebouxiophyceae.

Disclosure statement

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

Funding

This work was supported by the Securement, Discovery, and Basic Survey of Marine Biological Resources [2019M00100], funded by the National Marine Biodiversity Institute of Korea (MABIK). This study was also supported by a grant from the National Research Foundation of Korea [NRF-2017R1A2B4002016].

References

- Dierckxsens N, Mardulyn P, Smits G. 2017. NOVOPlasty: *de novo* assembly of organelle genomes from whole genome data. *Nucleic Acids Res.* 45(4):e18.
- Fan W, Guo W, Van Etten JL, Mower JP. 2017. Multiple origins of endosymbionts in Chlorellaceae with no reductive effects on the plastid or mitochondrial genomes. *Sci Rep.* 7(1):10101.
- Guiry MD, Guiry GM. 2019. AlgaeBase. Galway: National University of Ireland (World-wide electronic publication). <http://www.algaebase.org>.
- Hong JW, Jo SW, Cho HW, Nam SW, Shin W, Park KM, Lee Kl, Yoon HS. 2015. Phylogeny, morphology, and physiology of *Micractinium* strains isolated from shallow ephemeral freshwater in Antarctica. *Phycol Res.* 3:212–218.
- John DM, Whitton BA, Brook AJ. 2002. The freshwater algal flora of the British Isles: an identification guide to freshwater and terrestrial algae. 2nd ed. Cambridge (UK): Cambridge University Press; p. 488–490.