VIE ET MILIEU - LIFE AND ENVIRONMENT, 2018, 68 (2-3): 151-155

# SEQUENCE ANALYSIS CONFIRMS A NEW ALGAL CLASS

L. K. MEDLIN<sup>1</sup>, Y. DESDEVISES<sup>2</sup>

<sup>1</sup> Marine Biological Association of the UK, The Citadel, Plymouth PL1 2PB, UK <sup>2</sup> Sorbonne Université, CNRS, UMR 7232, Biologie Intégrative des Organismes Marins, BIOM, Observatoire Océanologique, 66650 Banyuls/Mer, France

MICROALGAL CLASS CHLOROMORUM PHYLOGENETICS ABSTRACT. – As judged by its position in an SSU rRNA tree, a new algal class in the heterokonts has been recovered from a Bayesian phylogenetic analysis of 12 heterokont algal classes. The sequences were deposited in Genbank as raphidophytes, possibly because cells of these species resemble raphidophytes. A constrained tree placing them in the raphidiophytes yielded a significantly worse tree as determined by the Shimodaira-Hasegawa test (P = 0.0001). New efforts should be made to describe this class formally because cultures do exist for these microalgae.

#### INTRODUCTION

In an earlier search for the appropriate number and type of outgroups to recover monophyletic classes in the diatoms (Medlin 2014), there were a group of sequences in the Silva database (SSUREF\_96 and SSUREF\_199\_128) that did not fall within the Raphidophyceae, although they were labeled in Genbank as being new species of Raphidophyceae (Suppl Table I). A revised taxon sampling of the heterokonts with a reduced number of diatoms for the 18S SSU rRNA gene showed that this group of sequences consistently fell into a well supported clade separated from the raphidophytes and thus would appear to be a new microalgal class in the pigmented heterokonts.

### MATERIALS AND METHODS

rRNA sequences from the Chloromorum spp. in Suppl Table I were uploaded from Genbank and aligned to the SILVA SSU rRNA sequence alignment in the ARB program Version 5.5 using maximum primary and secondary structural similarity (Ludwig et al. 2004). No filter was applied to the alignment and full length SSU sequences (2029 bases) from 59 taxa were exported for further analysis, Phylogenetic analyses were performed on the SSU alignment with RAxML (Randomized Axelerated Maximum Likelihood, Stamatakis et al. 2014) and a GTR + G model from the CIPRES online analysis portal (https://www.phylo.org/, Miller et al. 2010). Tree robustness was assessed with a bootstrap procedure with 1000 replicates. This tree was compared with a constrained phylogeny built with all putative raphidophytes forced as a monophyletic group. The comparison was performed with a SH-test (Shimodaira & Hasegawa 1999) using PAUP\* (Swofford 2003) with the FullOpt setting and 1000 bootstrap replicates in the testing procedure. The phylogenetic tree was also reconstructed by Bayesian inference with MrBayes 3.2.6 (Ronquist et al. 2012) using 4 chains of 2.106 generations, trees sampled every 1000 generations, and burnin value set to 20% of the sampled trees (400). We checked that standard deviation of the split frequencies fell below 0.01 to ensure convergence in tree search.

#### RESULTS AND DISCUSSION

The Chloromorum sequences fell into a well-supported clade (bootstrap support in % (bt) = 100, posterior probabilities pp = 1) that was sister with no support to a moderately supported larger clade (bt = 61) containing the Rhaphidophyceae, Chrysomerophyceae, Xanthophyceae, Phaeophyceae (Figs 1, 2), basically the SI clade of the pigmented heterokonts of Yang et al. (2012). This "SI" clade was collapsed into a polytomy with a posterior probability of 1 with the BI analysis (Fig. 2) and all other sister relationships were the same as the RAxML tree. Shimodaira-Hasegawa tests using a constrained tree (Fig. 3) to place these new sequences inside the rhapidophytes yielded a significantly worse tree (P = 0.0001). Based on these preliminary analyses, it is concluded that this group of sequences is not raphidophytes but belongs to a new class of microalgae with at least three species based on SSU analysis. There are twelve strains in culture at the MarbioNC Living Algal Resource Collection (Larc) at the University of North Carolina, Wilmington, which should be examined in more detail and with other genes sequenced to ascertain conclusively that they represent a new algae class with their own set of distinct morphological characters separating them from the other pigmented heterokont microalgae and how many new species are present.

This is not the first time that a new algal class has been identified first with sequence data and to be later defined taxonomically with morphological/physiological features from cultured cells. In 1995, Cavalier-Smith *et al.* revised the Ochristan (pigmented heterokont) algae and defined several new classes based on sequence analysis alone, such as the Chrysomerophyceae. Before that, the Pelago-

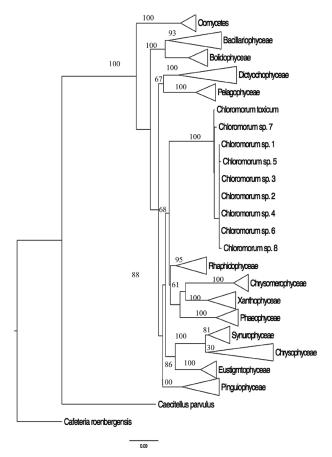


Fig. 1. – Phylogenetic analysis of the *Chloromorum* sequences using a RAxML analysis of selected members of each algal class in the Heterokonta to place them in a phylogenetic context. Bootstrap support values > 50 % are placed at each node.

phyceae were described from sequence data and little supportive morphological data (Andersen *et al.* 1993, Saunders *et al.* 1997), and after that several other microalgal classes were described, all with sequence data providing the strongest and usually the initial support for the new class: Bolidophyceae (Guillou *et al.* 1999), Pinguiophyceae (Kawachi *et al.* 2002), and Phaeothamniophyceae (Bailey *et al.* 1998). The picobiliphytes were recognized first as a new class with molecular data (Not *et al.* 2007) and it was several years and with much effort before cultures could be established to show conclusively that this was not a new algal class as originally predicted but a new heterotrophic protist phylum (Seenivasan *et al.* 2013).

Nor is it the first time that algal species first believed to be raphidophytes have been shown with molecular data to be more closely related to other groups (Bowers *et al.* 2006). In the Bowers *et al.* study, *Chattonella verreculosa* was found to be a dictyochophyte and later moved into a new genus *Pseudochattonella* (Edvardsen *et al.* 2007, Hosoi-Tanabe *et al.* 2007).

The sterols of fatty acids of *Chloromorum toxicum* have already been examined (Giner *et al.* 2008) referring to its original identification as a species of *Chattonella*, a valid member of the Raphidophyceae and compared to

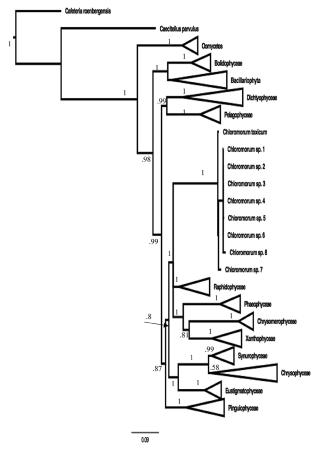


Fig. 2. – Phylogenetic analysis of the *Chloromorum* sequences using a Bayesian analysis of selected members of each algal class in the Heterokonta to place them in a phylogenetic context.

two other microalgae that had originally been described as *Chattonella* spp. Its sterols were significantly different from the other two algae and from those found in another study by Marshall *et al.* (2002) who suggested that sterols and fatty acids were conserved within a genus and thus more informative phylogenetically than pigments, which have been traditionally used to define groups of raphidophytes.

Yang et al. (2012) studied the evolution of the pigmented heterokonts and recovered basically three major clades in their analyses, which could primarily be defined by carotenoid pigments. Taxa in the SIII clade have only the diatoxanthin-diadinoxanthin cycle (D-D cycle) carotenoid cycle, as do some taxa in the SI clade, whereas the SII clade is defined by taxa with the violaxanthin-antheraxanthin cycle (V-A) cycle). Based on our phylogenetic analysis, it could be hypothesized that this new algal class contains diatoxanthin-diadinoxanthin cycle (D-D cycle), something that could easily be tested to support their placement in this major clade of the pigmented heterokonts.

The one sequence that has a name, *Chloromorum* toxicum Tomas, has been used without being validly published (Giner et al. 2008), having no Latin description or

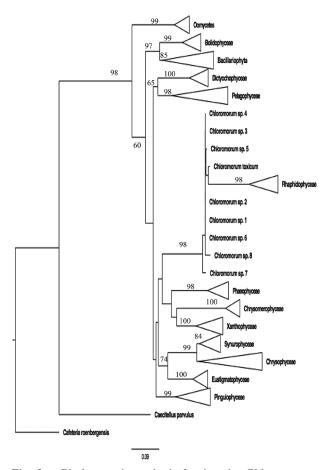


Fig. 3. – Phylogenetic analysis forcing the *Chloromorum* sequences into the Rhapidophyta for testing the validity of these sequences as members of that microalgal class. Bootstrap support values > 50 % are placed at each node.

designation of type material, and thus is a *nomen nudum*, which would make also a class name of Chloromorophyceae also illegal. Thus this species will have to be renamed and another class name linked to the new genus name for a valid description of all taxa.

#### REFERENCES

- Andersen RA, Saunders GW, Paskind MP, Sexton JP 1993. Ultrastructure and 18S rRNA gene sequence for *Pelagomonas calceolata*gen. et sp. nov. and the description of a new algal class, the Pelagophyceae classis nov. *J Phycol* 29: 701-715.
- Bailey JC, Bidigare RR, Christensen SJ, Andersen RA 1998. Phaeothamniophyceae classis nova: a new lineage of chromophytes based upon photosynthetic pigments, rbcL sequence analysis and ultrastructure. *Protist* 149: 245-263.
- Cavalier-Smith T, Chao EE, Allsopp MTEP 1995. Ribosomal RNA evidence for chloroplast loss within Heterokonta: Pedinellid relationships and a revised classification of Ochristan algae. Arch Protistenk 145: 209-220.
- Edvardsen B, Eikrem W, Shalchian-Tabrizi K, Riisberg I, Johnsen G, Naustvoll L, Throndsen J 2007. *Verrucophora farcimen* gen. et sp. nov. (Dictyochophyceae, Heterokonta) a

- bloom-forming ichthyotoxic flagellate from the Skagerrak, Norway. *J Phycol* 43: 1054-1070.
- Giner JL, Zhao H, Thomas C 2008. Sterols and fatty acids of three harmful algae previously assigned as *Chattonella*. *Phytochem* 69: 2167-2171.
- Guillou L, Chrétiennot-Dinet MJ, Medlin LK, Claustre H, Loiseaux-de Goër S, Vaulot D 1999. *Bolidomonas*: a new genus with two species belonging to a new algal class, the Bolidophyceae (Heterokonta). *J Phycol* 35: 368-381.
- Hosoi-Tanabe S, Honda D, Fukaya S, Otake I, Inagaki Y, Sako S 2007. Proposal of *Pseudochattonella verruculosa* gen. nov, comb. nov. (Dictyochophyceae) for a former raphidophycean alga *Chattonella verruculosa*, based on 18S rDNA phylogeny and ultrastructural characteristics. *Phycologia* 55.
- Kawachi M, Inouye I, Honda D, O'Kelly Ch J, Bailey JC, Bidigare RR, Andersen RA 2002. The Pinguiophyceae classis nova, a new class of photosynthetic stramenopiles whose members produce large amounts of omega-3 fatty acids. *Phycol Res* 50: 31-47.
- Ludwig W, Strunk O, Ralf Westram R, Richter L, Meier H, Yadhukumar Buchner, Lai T, Steppi S, Jobb G, Förster W, Brettske I, Gerber S, Ginhart AW, Gross O, Grumann S, Hermann S, Jost R, König A, Liss T, Lüßmann R, May M, Nonhoff B, Reichel B, Strehlow R, Stamatakis AP, Stuckmann N, Vilbig A. Lenke M, Ludwig T, Bode A, Schleifer KH 2004. ARB, a software environment for sequence data. *Nucleic Acids Res* 32: 1363-1371.
- Marshall JA, Nichols PD, Hallegraeff GM, 2002. Chemotaxonomic survey of sterols and fatty acids in six marine raphidophyte algae. *J Appl Phycol* 14: 255–265.
- Medlin LK 2014. Evolution of the Diatoms: VIII. Re-Examination of the SSU-Rrna Gene Using Multiple Outgroups and a Cladistic Analysis of Valve Features. *J Biodivers Biopros Dev* 1: 129.
- Miller MA, Pfeiffer W, Schwartz T 2010. Creating the CIPRES Science Gateway for inference of large phylogenetic trees. *In* Proceedings of the Gateway Computing Environments Workshop (GCE), 14 Nov. 2010, New Orleans, LA: 1-8.
- Not F, Valentin K, Romari K, Massana R, Vaulot D, Medlin LK 2007. Picobiliphytes: a marine picoplanktonic algal group with unknown affinities to other eukaryotes. *Science* 315: 253-255.
- Ronquist F, Teslenko M, van der Mark P, Ayres DL, Darling A, Höhna S, Larget B, Liu L, Suchard MA, Huelsenbeck JP 2012. MrBayes 3.2: efficient Bayesian phylogenetic inference and model choice across large model space. *Syst Biol* 61: 539-542.
- Saunders GW, Potter D, Andersen RA 1997. Phylogenetic affinities of the Sarcinochrysidales and Chrysomeridales (Heterokonta) based on analyses of molecular and combined data. *J Phycol* 33: 310-318.
- Seenivasan R, Sausen N, Medlin LK, Melkonian M 2013. *Picomonas judraskeda* gen. et sp. nov.: the first identified member of the *Picozoa phylum* nov., a widespread group of picoeukaryotes, formerly known as 'Picobiliphytes'. *PLoS ONE* 8: e59565.
- Shimodaira H, Hasegawa M 1999. Multiple comparisons of loglikelihoods with applications to phylogenetic inference. *Mol Biol Evol* 16: 1114-1116.
- Stamatakis A, Ludwig T, Meier H 2005. RAxML-III: a fast program for maximum likelihood-based inference of large phylogenetic trees. *Bioinformatics* 21: 456-463.
- Swofford DL 2002. PAUP\*. Phylogenetic Analysis Using Parsimony (\*) and other methods. Version 4.0b10. Sinauer Associates Sunderland MA.

U41054

Yang EC, Boo GH, Kim HJ, Cho SM, Boo SM, Andersen RA, Yoon HS 2012. Supermatrix data highlight the phylogenetic relationships of photosynthetic stramenopiles. *Protist* 163: 217-231.

Received on October 25, 2017 Accepted on January 31, 2018 Associate editor: E Magnanou

Suppl Table I. – Summary of taxa used in this analysis

Accession Number	Taxon
Bicoecea unpigmented heterok	onts
AF174364	Cafeteria roenbergensis Fench., Pat.
AY642126	Caecitellus parvulus (Greiss) Pat, Nyg. Steinb, Turl.
Oomycetes water molds	
M32705	Achlya bisexualis Cok., Couch
X54266	Lagenidium giganteum Couch
New Algal Class	
EU038275	Chloromorum toxicum Tomas
EU038278	Chloromorum sp. 1
EU038280	Chloromorum sp. 2
EU038274	Chloromorum sp. 3
EU038277	Chloromorum sp. 4
EU038279	Chloromorum sp. 5
EU038276	Chloromorum sp. 6
AY788946	Chloromorum sp. 7
EU038273	Chloromorum sp. 8
Bacillariophyceae	
M87326	Cylindrotheca closterium (Ehr.) Reim., Lewin
M87329	Rhizosolenia setigera Brightw.
M87334	Tryblionella apiculata Greg.
Bolidophyceae	
AF123596	Triparma mediterranea (Guil., ChretDin.) Ich., Santos
AF123595	Triparma pacifica (Guil., ChretDin.) Ich., Santos
Chrysophyceae	
AF123282	Chromophyton rosanoffii Woronin
AF123291	Dinobryon sociale var. americanum (Brun.) Bach.
AF123293	Ochromonas tuberculata Hibb.
Chrysomerophyceae	
AJ295822	Antarctosaccion applanatum (Gain) Del.
U78034	Giraudyopsis stellifera Dang.
Dictyochophyceae	
U14384	Apedinella radians (Lohm.) Campb.
U14385	Dictyocha speculum Ehr.
AY254857	Florenciella parvula Eik.
U14387	Pseudopedinella elastica Skuj.
Eustigmatophyceae	
U41051	Eustigmatos magus (Pet.) Hibb.

Monodus subterranea (Pet.) Hibb.

## $Suppl\ Table\ I.-Continued.$

Accession Number	Taxon
U41092	Nannochloropsis granulata Karl, Pot.
U41052	Pseudocharaciopsis minuta (Braun) Hibb.
AF045051	Vischeria helvetica (Vis., Pas.) Hibb.
Pelagophyceae	
U40257	Aureococcus anophagefferens Harg., Sieb.
U14386	Pelagococcus subviridis Norr.
U78033	Sarcinochrysis marina Geit.
Pinguiophyceae	
AF123284	Chrysochaete britannica (God.) Ros.
AF438325	Glossomastix chrysoplasta O'Kel.
AF438324	Pinguiococcus pyrenoidosus And, Pot, J. Bail.
AF438322	Polypodochrysis teissieri Magne
Phaeophyceae	
X53229	Costaria costata (C. Ag.) D. Saunders
L43062	Ectocarpus siliculosus (Dill.) Lyng.
AB011423	Fucus distichus L.
L43066	Scytosiphon lomentaria (Lyng.) Link
Rhapidophyceae	
AY788922	Chattonella antiqua (Hada) Ono
AY788944	Chattonella subsalsa Bieche.
U41649	Chattonella subsalsa
AY788931	Fibrocapsa japonica Tor., Tak.
U41650	Heterosigma akashiwo (Hada) Hada
AB217629	Pseudochattonella verruculosa (Har., Chi.) Tan, Hon, Fuk, In,, Sako
U41651	Vacuolaria virescens Cienk.
Synurophyceae	
U73228	Mallomonas caudata (lw.) lv.
U73220	Synura mammillosa Tak.
U73219	Tessellaria volvocina (Play.) Play.
Xanthophyceae	
AF083398	Bumilleriopsis filiformis Vis.
U43277	Heterothrix debilis Vis.
AF083399	Heterococcus caespitosus Vis.
AF083400	Mischococcus sphaerocephalus Vis.
U73219	Tessellaria volvocina (Play.) Play.
M55286	Tribonema aequale Pas.
AF083397	Tribonema intermixtum Geit.