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Taxonomy and nomenclature of the recently published chlamydomonad genera *Ostravamonas* and *Paludistella* (Volvocales, Chlorophyceae)

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A recently published chlamydomonad green algal genus, *Paludistella* H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 72), was found to be superfluous as it originally included the type of *Ostravamonas* Barcytë & Hodač in Barcytë *et al.* (2020: 214). Here, the taxonomic and nomenclatural status under these generic names were examined based on morphological, phylogenetic, and ITS-2 secondary structure analyses, proposing two new combinations for *P. asymmetrica* H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 76) and *P. trianguloculus* H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 76) under *Ostravamonas*.

Traditionally, *Chlamydomonas* Ehrenberg (1834: 288) (Volvocales, Chlorophyceae) includes hundreds of flagellate green microalgae (e.g., Ettl 1983), but recent molecular phylogenetic studies have revealed its polyphyly, urging its division into monophyletic genera (e.g., Pröschold *et al.* 2001, Demchenko *et al.* 2012, Nakada *et al.* 2016). In the course of the taxonomic revisions, *Ostravamonas* and *Paludistella* have been published.

Ostravamonas was first validly published online on 11 December 2019 in the *European Journal of Phycology*, and included three species, *O. chlorostellata* (E.A. Flint & H. Ettl 1966: 420) Barcytë & Hodač in Barcytë *et al.* (2020: 215) (type), *O. meslinii* (Bourrelly 1951: 258) Barcytë & Hodač in Barcytë *et al.* (2020: 217), and *O. tenuiincisa* Barcytë & Hodač in Barcytë *et al.* (2020: 217). On the same day, a manuscript proposing *Paludistella* was accepted by *Phytotaxa*, and it was published online on 5 February 2020 (Susanti *et al.* 2020). Because *Paludistella* includes *P. chlorostellata* (E.A. Flint & H. Ettl) H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 75), which was the type of *Ostravamonas*, *Paludistella* is an illegitimate superfluous name under Art. 52 of the *Shenzhen Code*. Susanti *et al.* (2020) also proposed two new species, *P. asymmetrica* and *P. trianguloculus*. While these new species were discriminated from *O. chlorostellata* (\equiv *P. chlorostellata*) and *O. meslinii* (\equiv *P. meslinii*), they were not compared with *O. tenuiincisa*, newly described in Barcytë *et al.* (2020). Here, genera and species published by Barcytë *et al.* (2020) and Susanti *et al.* (2020) were compared based on descriptions, 18S *r*DNA, and ITS-2 secondary structures, and the correct names under *Ostravamonas* are clarified.

The 18S *r*DNA sequences of *O. chlorostellata* strain SAG 12.72 (MK912143) and *O. meslinii* strain SAG 75.81 (MK912144) published by Barcytë *et al.* (2020) were identical to those published by Susanti *et al.* (2020) (MK696129 and MK696131, respectively). Therefore, these sequences were used for further analyses. The 18S *r*DNA sequences of *O. tenuiincisa* (FR865525) was appended to the 18S *r*DNA alignment generated by Susanti *et al.* (2020), manually aligned (provided in Supplemental Material), and used for phylogenetic analyses according to the methods of Susanti *et al.* (2020). The morphological descriptions of *Ostravamonas* and *Paludistella* were generally consistent with each other. In the 18S *r*DNA phylogenetic tree (Fig. 1), all species belonging to *Ostravamonas* and *Paludistella* formed a robust clade (posterior probability [PP]=1.00, bootstrap proportion [BP] in maximum likelihood analysis = 100%) within the clade *Chloromonadinia* (sensu Nakada *et al.* 2008).

Three species described in Barcytë *et al.* (2020) and Susanti *et al.* (2020) were morphologically compared based on their original descriptions. The morphological characteristics of three species, *O. tenuiincisa*, *P. asymmetrica*, and *P. trianguloculus*, were similar, with some differences. Young cells of *P. asymmetrica* were frequently asymmetrical, while no such cells were reported for *O. tenuiincisa* and *P. trianguloculus*. The stigma was string-like in *O. tenuiincisa*, oblong to

small elliptical in *P. asymmetrica*, and triangular to elliptical in *P. trianguloculus*. Zoosporangia including eight zoospores were reported for *O. tenuiincisa* but not for *P. asymmetrica* or *P. trianguloculus*. Similarly, akinetes were reported only for *P. asymmetrica*. However, these characters related to asexual reproduction can be affected by culture conditions, and their importance in species taxonomy requires further studies.

In the 18S *r*DNA phylogenetic tree (Fig. 1), *P. asymmetrica* and *O. chlorostellata* were sister to each other (PP = 1.00, BP = 98%), and they formed a clade with *P. trianguloculus* (PP = 1.00, BP = 100%) but were distantly related to *O. tenuiincisa*. The predicted secondary structure of nuclear *r*DNA ITS-2 of *O. tenuiincisa* (Barcytë *et al.* 2020) was compared with models for *P. asymmetrica* and *P. trianguloculus* (Susanti *et al.* 2020). The four helices of ITS-2 were comparable among the new species, and five compensatory and six or seven hemi-compensatory base changes, respectively, were recognized between *O. tenuiincisa* and *P. asymmetrica* or *P. trianguloculus* (Fig. 2).

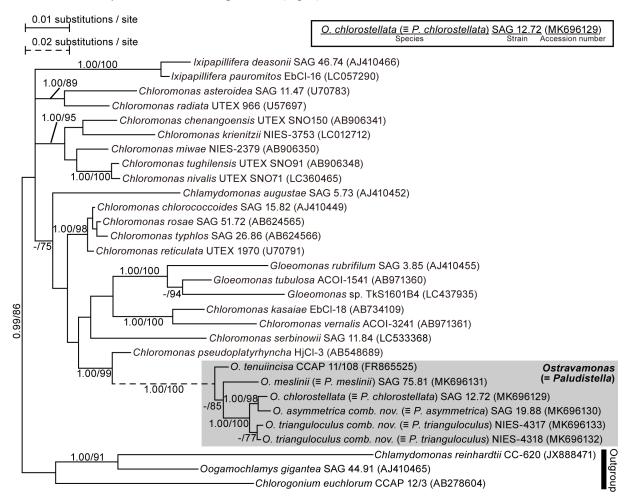


FIGURE 1. Bayesian phylogenetic tree of the clade *Chloromonadinia* based on 18S *r*DNA. Numbers on branches indicate posterior probabilities (≥ 0.95 ; left) and bootstrap percentages for the maximum likelihood analysis ($\geq 70\%$; right).

	Helix II	Helix III	Helix IV
O. tenuiincisa CCAP 11/108	5'UAGA 1111 3'AUCU	GCUAC GUGUUCAUGUUGUUAUGGAUACA UAA 	UGAA
O. asymmetrica comb. nov. (≡ P. asymmetrica) SAG 19.88	5'UUAU 1111 3'AGUA	gccaAugauuaucuuuuuca-agauguaagcaa cuguuaccaguaaaaaagu-ucuacguucuuu	UGAA 56 1111 ACUU
O. trianguloculus comb. nov. (≡ P. trianguloculus) NIES-4317	5′UCAA 1111 3′AGUU	gcca augaucaucuuuuuca - agauguaagcaa cggu uacugg - agaaaagu - ucuacguucuuu	UGAA56 ACUU
O. trianguloculus comb. nov. (≡ P. trianguloculus) NIES-4318	5′UCAA 1111 3′AGUU	gccaaugaucaucuuuuuca-agauguaagcaa caguuacugg-agaaaggu-ucuacguucuuu	UGAA IIIII ACUU

FIGURE 2. Comparisons of ITS-2 secondary structures of *Ostravamonas tenuiincisa* with *O. asymmetrica* and *O. trianguloculus* based on Barcytë *et al.* (2020) and Susanti *et al.* (2020), except where modified (*). Hyphens (-) and ellipses (...) indicate gaps and omitted regions, respectively. The positions marked by gray boxes and empty boxes respectively indicate the presence of compensatory and hemicompensatory base changes compared to the *O. tenuiincisa* sequence. Numbers are indicated on the right.

All morphological, phylogenetic, and ITS-2-structural analyses supported the synonymy of *Paludistella* with *Ostravamonas* and the independence of *O. tenuiincisa* from *P. asymmetrica* and *P. trianguloculus*. Therefore, the correct generic name for the taxon is *Ostravamonas*. Here, a species list with synonymy and new combinations for *P. asymmetrica* and *P. trianguloculus* under *Ostravamonas* are compiled.

Ostravamonas Barcytë & Hodač in Barcytë et al. (2020: 214)

Type: Ostravamonas chlorostellata (E.A. Flint & H. Ettl) Barcytë & Hodač in Barcytë et al. (2020)

Heterotypic synonym: Paludistella H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe, (2020: 72), nom. illeg. superfl.

Ostravamonas chlorostellata (E.A. Flint & H. Ettl 1966: 420) Barcytë & Hodač (2020: 215)

Type: Flint & Ettl (1966), fig. 2 (holotype); SAG 12.72 (authentic strain; permanently cryopreserved in a metabolically inactive state) (epitype; designated by Barcytë *et al.* 2020)

Homotypic synonyms: *Chlamydomonas chlorostellata* E.A. Flint & H. Ettl (basionym) (excl. var. *gracillima* H. Ettl), *Paludistella chlorostellata* (E.A. Flint & H. Ettl) H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 75)

Ostravamonas asymmetrica (H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe 2020: 76) Nakada & H. Susanti *comb. nov*.

Type: TNS-AL-58967 (specimen prepared from the strain SAG 19.88) (holotype)

Homotypic synonym: *Paludistella asymmetrica* H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe 2020: *Phytotaxa* 432: 76 (figs. 3, 6E, F; basionym)

Ostravamonas meslinii (Bourrelly 1951: 258) Barcytë & Hodač in Barcytë et al. (2020: 217)

- Type: Bourrelly (1951), pl. 3 fig. 52 (holotype); SAG 75.81 (authentic strain; permanently cryopreserved in a metabolically inactive state) (epitype; designated by Barcytë *et al.* 2020)
- Homotypic synonyms: *Chlamydomonas meslinii* Bourr. (basionym); *Paludistella meslinii* (Bourr.) H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe (2020: 73)

Ostravamonas tenuiincisa Barcytë & Hodač in Barcytë et al. (2020: 217)

Type: CCAP 11/108 (permanently cryopreserved in a metabolically inactive state) (holotype)

Ostravamonas trianguloculus (H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe 2020: 76) Nakada & H. Susanti *comb. nov*.

- Type: NIES-4318 (permanently cryopreserved in a metabolically inactive state) (holotype); NIES-4317 (permanently cryopreserved in a metabolically inactive state) (paratype); TNS-AL-58966 (specimen prepared from the strain NIES-4318) (paratype); TNS-AL-58965 (specimen prepared from the strain NIES-4317) (paratype)
- Homotypic synonym: *Paludistella trianguloculus* H. Susanti, Mas. Yoshida, Tak. Nakayama, Nakada & M.M. Watanabe 2020: *Phytotaxa* 432: 76 (figs. 4, 5, 6C, D; basionym)

Note: In Susanti et al. (2020), TNS-AL-58966 was erroneously referred as an isotype.

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Supplemental Material. Multiple sequence alignment of nuclear 18S rDNA in FASTA format.