

## *Stephanodiscus akanensis* sp. nov., a new species of extant diatom from Lake Akan, Hokkaido, Japan

Akihiro Tuji<sup>1</sup>, Ayako Kawashima<sup>2</sup>, Matthew L. Julius<sup>3</sup> and Eugene F. Stoermer<sup>4</sup>

<sup>1</sup>Department of Botany, National Science Museum, Tokyo,  
4–1–1, Amakubo, Tsukuba, 305–0005 Japan  
E-mail: tuji@kahaku.go.jp

<sup>2</sup>Hiraoka Environmental Science Laboratory, 5–15–6, Harajuku, Shiroyama-machi,  
Tsukui-gun, Kanagawa, 220–102 Japan

<sup>3</sup>Biological Sciences, St. Cloud State University, 225 Mathematics & Science Center,  
720 Fourth Avenue South, St. Cloud, MN 56301–4498, U.S.A.

<sup>4</sup>Center for Great Lakes and Aquatic Sciences, University of Michigan,  
Ann Arbor, MI 48109–1090, U.S.A.

**Abstract** The morphology of a new centric diatom species, *Stephanodiscus akanensis* Tuji, Kawashima, Julius & Stoermer is described using light and scanning electron microscopy. The species is so far only known from Lake Akan, Hokkaido, Japan. It is separated from other similar *Stephanodiscus* species by the characteristic position of the valve face fuloportulae and the numbers of rimoportulae.

**Key words:** Bacillariophyceae, Lake Akan, Centric diatoms, ultrastructure, initial valve.

### Introduction

Lake Akan is a eutrophic system in eastern Hokkaido, Japan (43°27'N, 144°07'E) and lies at an elevation of 429 m. It has a surface area of 13.3 km<sup>2</sup>, a 42 m maximum depth, and an 18 m mean depth. The lake is well known for the occurrence of *Cladophora sauteri* (Nees) Kützing colonies, which appear as macroscopic “balls” (Canter-Lund and Lund, 1995). Takayasu *et al.* (1930) first described Lake Akan’s plankton flora and fauna. Negoro and Watanabe (1977) expanded our knowledge of Lake Akan’s plankton flora by a light microscope (LM) investigation. Kawashima and Kobayasi (1993, 1994, 1995, 1996) and Kawashima and Mayama (1997, 1998, 2000, 2001, 2002) described the lake’s diatom flora in greater detail using both the LM and scanning electron microscope (SEM). In this paper we continue to expand the knowledge base about the plankton flora with the description of a new species, *Stephanodiscus akanensis*, from the system.

### Materials and Methods

*S. akanensis* is described from a near shore plankton tow collected in August 1995 from a 3 m depth. The samples were processed for LM and SEM observations following the protocols of Kawashima and Kobayasi (1993). Material for SEM observation was dried onto poly-L-lysine coated 10 mm coverslips, washed with distilled water, and sputter-coated with platinum. Observations were performed with a JOEL-6301F SEM. A single specimen on an unsputter-coated coverslip was examined with the SEM. After completing SEM observations of the coverslip, it was used to make a permanent Pleurax mount for LM observation. This LM preparation serves as the holotype for the species. Observation of the specimen in the SEM prior to its preparation as a LM slide provides the opportunity to obtain complete information about the LM and SEM morphology of a single diatom specimen, in this instance the holotype specimen. Terminology for specific morphological features follows Håkansson and Locker (1981).

### Diagnosis

*Stephanodiscus akanensis* Tuji, Kawashima, Julius & Stoermer sp. nov. (Figs. 1–5)

Valvae circulares, fronte leviter concentricè undulatae, 13–60  $\mu\text{m}$  diam. Areola in fronte limboque divergentes, cribris tholiformibus penitus instructae, fasciculos formantes. Fasciculi 7–8 per 10  $\mu\text{m}$ , prope centrum valvae uniseriati, versus limbum ex 2–3 seiebus areolarum constans. Costa extrinsecus parum elevata. Spinae in quiaequè extremitatibus interstriarum regulariter dospositae. Fulportulae frontis valvae ex primo tubo poris brevi et 2 satellitibus instructae. Fulportulae limbi per costam secundariam vel tertiariam in parte inferiori spinae extus crebrae, interne ex primo tubo brevi et 3 poris satellitibus praeditae. Rimoportulae duae, oppositae, ad partem conjunctam frontis valvae et limbi praesentes, velut tubo proxime subter spina visiviles.

Valves circular, 13–60  $\mu\text{m}$  in diameter with 7–8 fascicles in 10  $\mu\text{m}$ . Valve surface slightly concentrically undulate. Areolae form fascicles. Fascicles uniseriate near valve center becoming bi- or triseriate near valve margin. Areolae on valve face and mantle have internally domed cribra. Interfascicles slightly raised externally. Spines regularly placed at the end of each interfascicle. One or two valve face fulportulae, with a short primary pore tube and two satellite present. Mantle fulportulae located below spines externally, occurring on every second or third interfascicles. Internally mantle fulportulae have a short primary tube and three satellite pores. Two rimoportulae present opposite one another at the valve face/mantle junction, and visible externally as a short tube immediately beneath a spine.

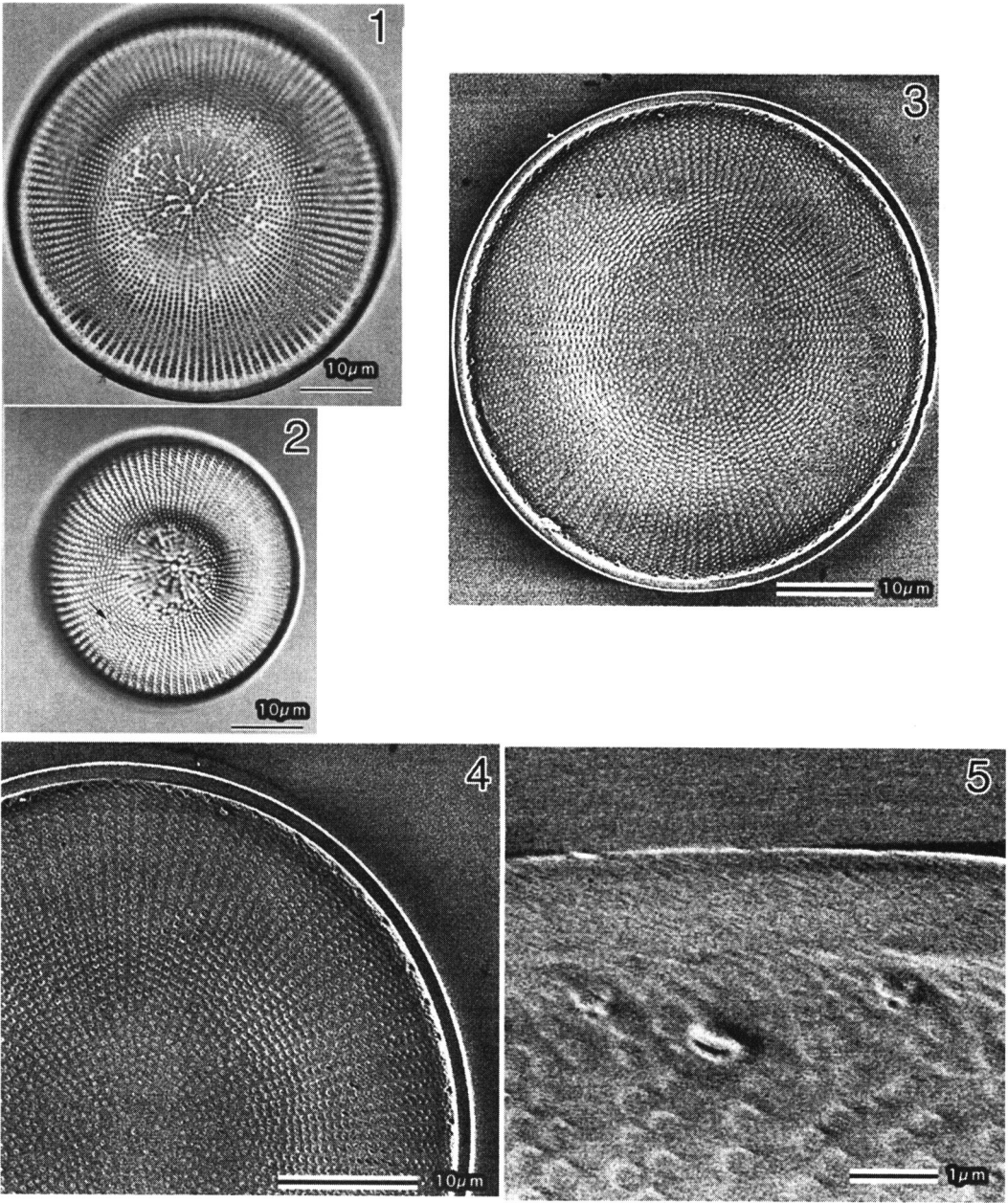
Holotype: TNS-AL-53966s, Department of Botany, National Science Museum, Tokyo, Japan.

Type Material: TNS-AL-53966m, Department of Botany, National Science Museum, Tokyo, Japan.

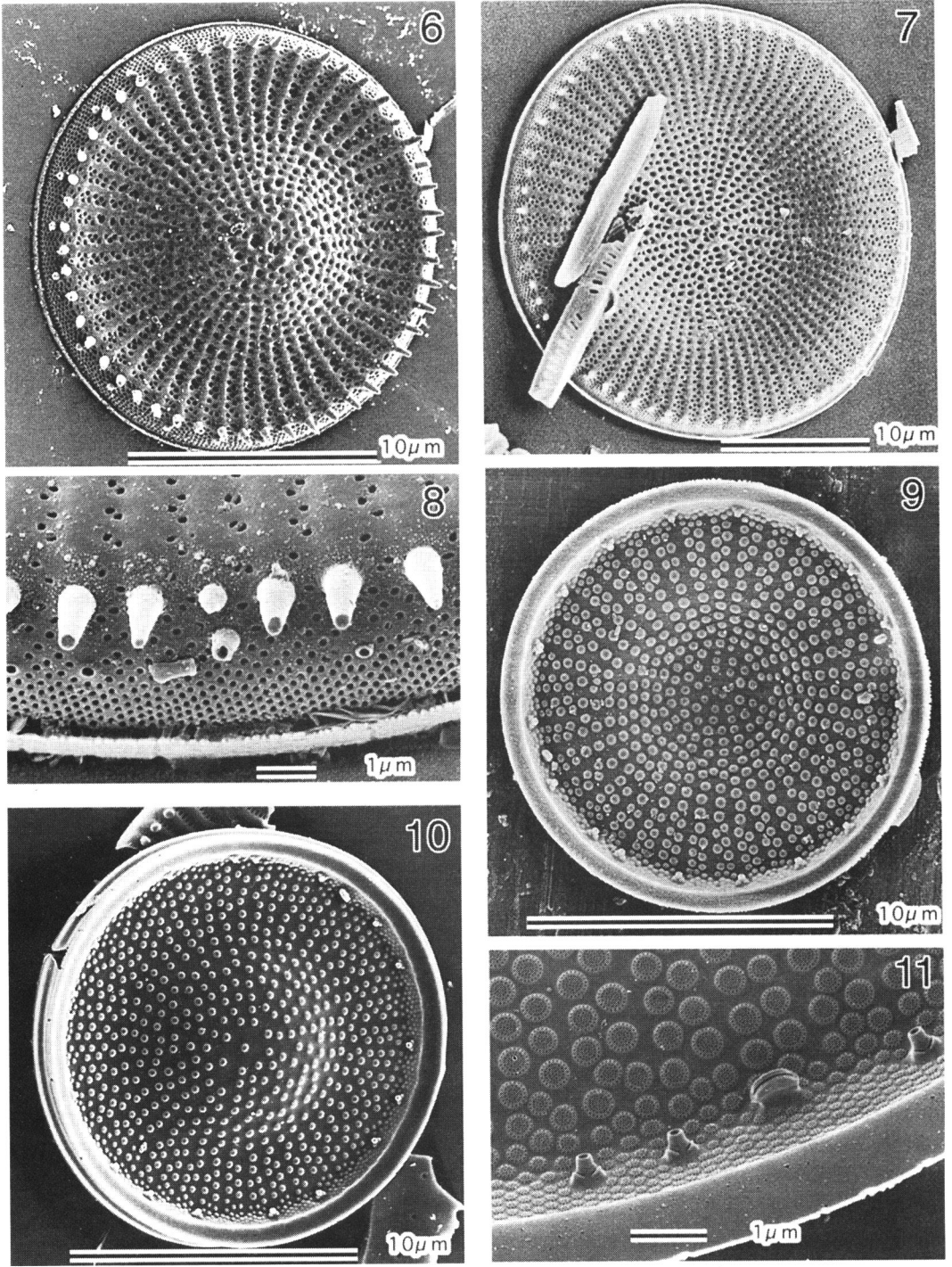
Type Locality: Lake Akan, Hokkaido, Japan.

### Observations

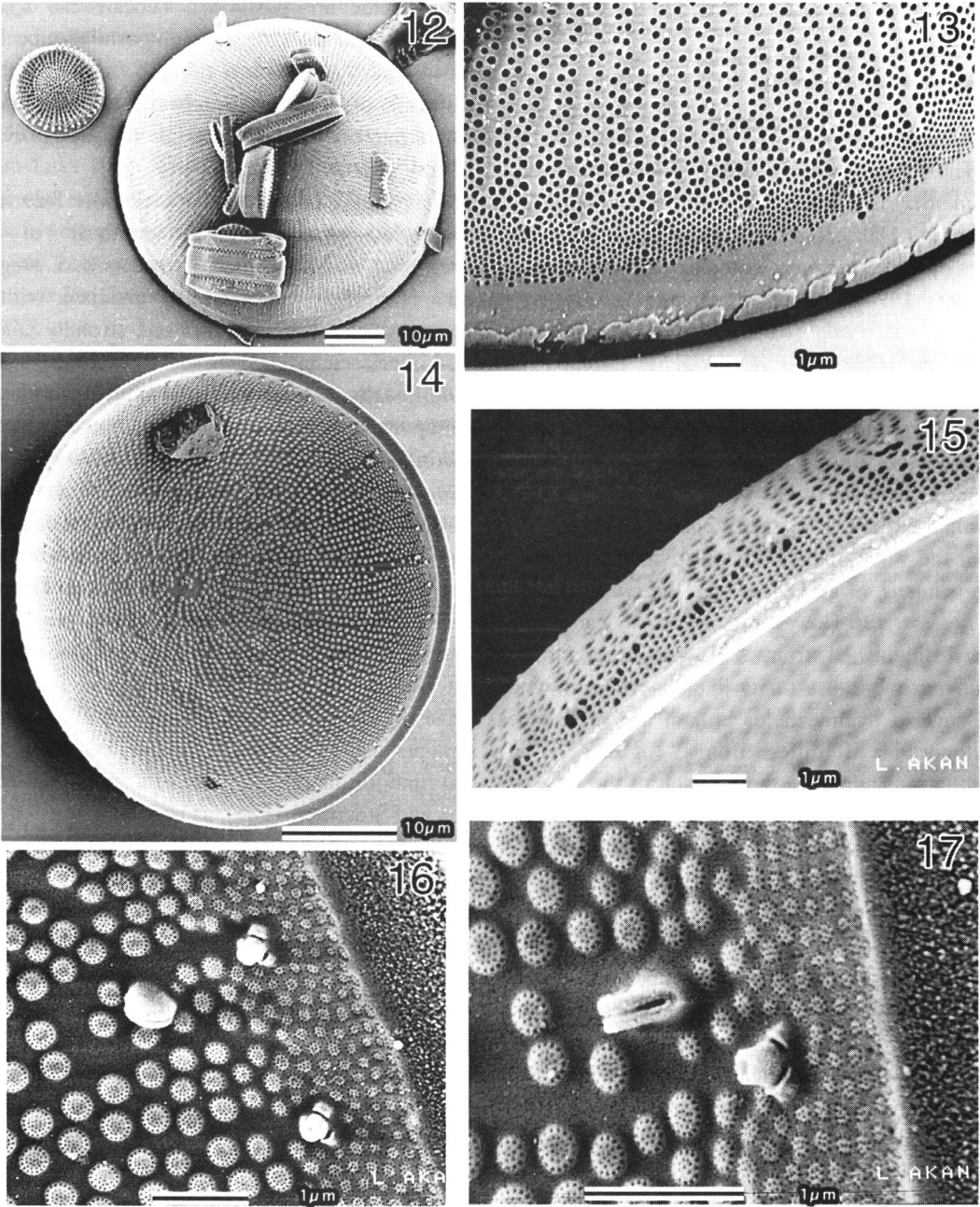
The fascicles of areolae are separated by distinct interfascicles (Figs. 1, 2). The areolae begin as single rows at the center of the valve face and increase to 2–3 rows (rarely 4) toward the valve face margin (Figs. 1–4, 6, 7). Areola density increases on the valve mantle beginning near the mantle fulportulae horizon and continuing to the valve mantle edge. The areolae pattern also changes in conjunction with the density difference, moving from a fasciculate pattern to a decussate one. Areolae density on the mantle is approximately 60 in 10  $\mu\text{m}$  and a single row parallel to the valve radius contains 4–9 areolae. Punctae on the mantle occur in vertical files, numbering 6–7 punctae per vertical file. Spines occur at each interfascicle. One (rarely 2) valve face fulportulae with two satellite pores occurs near the valve margin (Fig. 10). In some specimens this valve face process appears to be absent (Fig. 9). The number of valve face fulportulae does not appear to be cell size related. *Stephanodiscus akanensis* does not have a “heterotopic fulportulae position” as defined by Håkanson and Meyer (1994). The mantle fulportulae with 3 satellite pores are positioned on the upper valve mantle (Figs. 3, 5, 6–8). Externally, mantle fulportulae openings appear as simple pores with a slightly thickened rim, lacking a well developed external tube (Fig. 8). The two (rarely one) rimoportulae are opposite position from one another (Fig. 9). The rimoportulae occur directly underneath a spine and above the mantle fulportulae horizon (Fig. 8). The labium of the rimoportulae appears to be variable direction (Figs. 9, 11). Initial cells were observed, and range from 55–60  $\mu\text{m}$  in diameter. Spines are absent on initial valves and a broad, hyaline zone is present on the lower mantle (Figs. 12, 13), in contrast to the morphology expressed in vegetative valves. Internally, initial valves possess mantle fulportulae with 3 satellite pores (Figs. 14, 16, 17), similar to the vegetative valves. Externally, the fulportulae and rimoportulae appear as simple pores (Fig. 15), differing from the fulportulae siliceous rim



Figs. 1–5. *Stephanodiscus akanensis* sp. nov. Figs. 1, 3–5. Holotype. Figs. 1–2. LM DIC images. Figs. 3–5. SEM images using the low vacuum SEM. Figs. 1, 3 Holotype frustule without central fultoportulae. Fig. 2. Frustule with central fultoportula (arrow). Figs. 4, 5. Rimoportula and fultoportulae.



Figs. 6–11. *Stephanodiscus akanensis* sp. nov. Figs. 6–7. Exterior of whole frustule. Fig. 8. Spines and opening of rimoportula and fultoportulae. Fig. 9. Interior of a frustule without central fultoportulae. Fig. 10. Interior of a frustule with a central fultoportula. Fig. 11. Rimoportula and fultoportulae.



Figs. 12–17. *Stephanodiscus akanensis* sp. nov., Initial valves. Fig. 12. Exterior of whole initial valve and a vegetative cell. Figs. 13, 15. Opening of fuloportulae. Fig. 14. Interior of whole initial valve. Figs. 16–17. Rimoportula and fuloportulae on initial valve.

and rimoportulae tube expressed in the vegetative valve.

### Discussion

The valve face morphology of *Stephanodiscus akanensis* is similar to that of *S. oregonica* (Ehrenb.) Håk., *S. alpinus* Hust., and the members of the *S. niagarae* complex (Håkansson and Meyer 1994), which includes *S. niagarae* Ehrenb., *S. rotula* (Kütz.) Hendy, *S. neoastrea* Håk. et Hickel, *S. aegyptiacus* Ehrenb., *S. galileensis* Håk. et Ehrlich, *S. agassizensis* Håk. et Kling, and *S. heterostylus* Håk. et Meyer (Håkansson. 2002).

In *S. akanensis*, the valve face fuloportulae number either 0 or 1 (rarely 2) and are positioned near the valve margin. This also characterizes *S. aegyptiacus*, *S. galileensis*, *S. agassizensis*, and *S. heterostylus*. However, *S. akanensis* can easily be distinguished from *S. aegyptiacus* and *S. galileensis* because of its finer interfascicle and absence of hyaline fields around the valve face fuloportulae. It differs from *S. agassizensis* in having a smaller size range. Each of the preceding species always has a single rimoportula, differing from *S. akanensis* which generally bears 2 rimoportulae and only rarely exhibits a single rimoportula. *S. akanensis* differs from *S. heterostylus* in its mantle fuloportulae positioning. In *S. heterostylus* the mantle fuloportulae are always located beneath a spine and positioned near the spine base. In *S. akanensis* the mantle fuloportulae are also located beneath spines, but their distance from the spine base is greater than the positioning expressed in *S. heterostylus*. Additionally, *S. heterostylus* has a higher maximum number of valve face fuloportulae, as described by Håkansson and Meyer (1994). *S. niagarae* has a broader mantle with areolae consisting of peralvar rows of 10–12 areolae, in contrast to *S. akanensis* which possesses peralvar rows of 4–9 areolae. The rimoportulae of *S. niagarae* are positioned within the spine horizon, with the external rimoportulae tube taking the place of a spine (Theriot and Stoermer. 1984). In contrast, *S. akanensis* the

rimoportulae are positioned beneath the spine horizon, with the external rimoportulae tube located directly beneath a spine. *S. niagarae* also differs from *S. akanensis* in having a higher maximum number of valve face fuloportulae (Theriot and Stoermer. 1984).

Valves of *S. akanensis* that lack valve face fuloportulae appear similar to those of *S. oregonica*, *S. alpinus*, and *S. neoastrea*. *S. oregonica* is strongly concentrically undulate, with a lacunate central area, and raised, strongly silicified interfascicles (Håkansson and Kling. 1999). Additionally, *S. oregonica* only expresses a single rimoportula (Håkansson. 1986). These features distinguish *S. oregonica* from *S. akanensis*. *S. neoastrea* can be distinguished from *S. akanensis* through observation of rimoportulae placement. In *S. neoastrea* the rimoportulae are positioned between interfascicle (Håkansson and Hickel, 1986; Håkansson and Meyer, 1994), but in *S. akanensis* the rimoportulae are positioned on a interfascicle (Figs. 9–11). *S. akanensis* is distinguishable from *S. alpinus* through differences in rimoportulae placement. *S. alpinus* has the external rimoportula tube in place of a spine (Håkansson and Stoermer, 1984), as in *S. niagarae*, while in *S. akanensis* the external rimoportula tube occurs directly under a spine. The maximum number of valve face fuloportulae expressed in each species represents an additional difference between *S. akanensis* and *S. alpinus*. *S. alpinus* always has 0 or 1 valve face fuloportulae, while *S. akanensis* can have 2 valve face fuloportulae.

Takayasu *et al.* (1930) first described Lake Akan's plankton flora and fauna. There is no information concerning *Stephanodiscus* species in this paper. Negoro and Watanabe (1977) discussed *S. astrea* and *S. astrea* var. *minutula* being present in Lake Akan. Kawashima and Kobayasi (1993) described the diatom flora in detail, using both LM and SEM observations. They discussed an unidentified *Stephanodiscus* sp. and *Actinocyclus* sp., which they felt had been identified by earlier investigators as *S. astrea* and *S. astrea* var. *minutula*. Hustedt (1930) was widely used by many Japanese diatomists for identifica-

tion of freshwater diatom species. This work contains limited information about *Stephanodiscus* species, and its use can lead to misidentification of specific *Stephanodiscus* species. Recently, Tuji & Kociolek (2000) described two new species from Lake Biwa, *Stephanodiscus suzukii* Tuji & Kociolek and *S. pseudosuzukii* Tuji & Kociolek, which were previously identified as *S. carconensis* var. *carconensis* Grunow and *S. carconensis* var. *pusilla* Grunow by Skvortzow (1936). Tuji (2002) also re-described *Aulacoseira nipponica* (Skvortzow) Tuji from Lake Biwa, where they had been identified previously as *A. solida* (Eulenstein) Krammer. Misidentification of Japanese centric diatom taxa should continue, and the re-discovery of new species like *S. akanensis* emphasize the need for more study of freshwater centric diatom taxonomy in Japan.

### Acknowledgements

We dedicate this work in memory of the late Dr. Hiromu Kobayaim, whose observations led us to undertake this study. The authors thank Dr. Isamu Wakana, Akan Educational Board for assistance in collecting samples from Lake Akan, and Dr. Ken Katumoto for the Latin diagnosis. We also thank Dr. Yashushi Kusuoka and the Lake Biwa Museum for the use of the FE-SEM.

### References

- Håkansson, H. 1986. A taxonomic reappraisal of some *Stephanodiscus* species (Bacillariophyta). *British phycolological journal*, **8**, 89–98.
- Håkansson, H. 2002. A compilation and evaluation of species in the general *Stephanodiscus*, *Cyclostephanos* and *Cyclotella* with a new genus in the family Stephanodiscaceae. *Diatom Research*, **17**, 1–139.
- Håkansson, H. & B. Hickel. 1986. The morphology and taxonomy of the diatom *Stephanodiscus neoastraea* sp. nov. *British phycolological journal*, **21**, 39–43.
- Håkansson, H. & H. Kling. 1989. A light and electron microscope study of previously described and new *Stephanodiscus* species (Bacillariophyceae) from central and northern Canadian Lakes, with ecological notes on the species. *Diatom Research*, **4**, 239–288.
- Håkansson, H. & B. Meyer. 1994. A comparative study of species in the *Stephanodiscus niagarae*—complex and a description of *S. heterostylus* sp. nov. *Diatom Research*, **9**, 65–85.
- Håkansson, H. & E. F. Stoermer. 1984. An investigation of the morphology of *Stephanodiscus alpinus* Hust. *Bacillaria*, **7**, 159–172.
- Hickel, B. & H. Håkansson. 1993. *Stephanodiscus alpinus* in plusssee, Germany. Ecology, morphology, and taxonomy in combination with initial cells. *Diatom Research*, **8**, 89–98.
- Kawashima, A. & H. Kobayasi. 1993. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 1. Centric Diatoms. *Nat. Envir. Sci. Res.*, **6**, 41–58.
- Kawashima, A. & H. Kobayasi. 1994. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 2. *Fragilaria sensu lato*. Diatoms. *Nat. Envir. Sci. Res.*, **7**, 9–22.
- Kawashima, A. & H. Kobayasi. 1995. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 3. Araphid diatoms except for *Fragilaria sensu lato*. Diatoms. *Nat. Envir. Sci. Res.*, **8**, 35–49.
- Kawashima, A. & H. Kobayasi. 1996. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 4. Raphid diatoms: *Eunotia*, *Cocconeis*, *Achnanthes*, *Rhoicosphenia*. *Nat. Envir. Sci. Res.*, **9**, 15–32.
- Kawashima, A. & S. Mayama. 1997. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 5. Raphid diatoms: *Aneumastus*, *Craticula*, *Diatomella*, *Diploneis*, *Frustulia*, *Gyrosigma*, *Luticola*, *Neidium*, *Sellaphora*, *Stauroneis*. *Nat. Envir. Sci. Res.*, **10**, 35–52.
- Kawashima, A. & S. Mayama. 1998. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 6. Raphid diatoms: *Cavinula*, *Diadesmis*, *Geissleria*, *Hippodonta*, *Navicula*, *Placoneis*. *Nat. Envir. Sci. Res.*, **11**, 23–41.
- Kawashima, A. & S. Mayama. 2000. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 7. Raphid diatoms: *Caloneis*, *Pinnularia*. *Nat. Envir. Sci. Res.*, **13**, 67–83.
- Kawashima, A. & S. Mayama. 2001. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 8. Raphid diatoms: *Cymbella*, *Encyonema*, *Gomphoneis*, *Gomphonema*, *Gomphosphenia*, *Reimeria*. *Nat. Envir. Sci. Res.*, **14**, 89–109.
- Kawashima, A. & S. Mayama. 2002. Diatoms from Akan-ko (Lake Akan) in Hokkaido, Japan. 9. Raphid diatoms: *Amphora*, *Epithemia*, *Rhopalodia*. **15**.
- Negoro, K. & M. Watanabe, 1977. On the phytoplankton of Lake Akan. *Bull. Jap. Soc. Phycol.*, **25**, Suppl., 221–237.
- Skvortzow, B. W. 1936. Diatoms from Biwa Lake, Honshu Island, Nippon. *Philipp. J. Sci.* **61**, 253–96, 8 pls.
- Takayasu, S., H. Igarashi, & J. Sawa. 1930. Akan-ko Cyosa. *Report of Fisheries Research*, **21**, 67–70.
- Theriot, E. & K. Serieyssel. 1994. Phylogenetic systematics as a guide to understanding features and potential morphological characters of the centric diatom family

- Thalassiosiraceae. *Diatom Research*, **9**, 429–450.
- Theriot, E. & E. F. Stoermer. 1984. Principal component analysis of *Stephanodiscus*. *Bacillaria*, **7**, 37–58.
- Theriot, E. H. Håkansson, & E. F. Stoermer. 1988. Morphometric analysis of *Stephanodiscus alpinus* (Bacillariophyceae) and its morphology as an indicator of trophic status. *Phycologia*, **27**(4), 485–493.
- Tuji, A. 2002. Observations on *Aulacoseira nipponica* from Lake Biwa, Japan, and *Aulacoseira solida* from North America (Bacillariophyceae). *Phycol. Res.* **50**, 313–316.
- Tuji, A & J. P. Kociolek. 2000. Morphology and taxonomy of *Stephanodiscus suzukii* sp. nov. and *S. pseudosuzukii* sp. nov. (Bacillariophyceae) from Lake Biwa, Japan and *S. carconensis* from North America. *Phycol. Res.* **48**, 231–9.