New species of the dinoflagellate cyst genus
Svalbardella Manum, 1960, emend. from the Paleogene
and Neogene of the northern high to middle latitudes

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Received: 17 March 2020 – Revised: 27 June 2020 – Accepted: 7 July 2020 – Published: 10 September 2020

Abstract. Species of the fusiform peridiniacean dinoflagellate cyst genera Svalbardella Manum, 1960, emend. (Eocene–Oligocene) and Palaeocystodinium Alberti, 1961 (Late Cretaceous–Miocene), have been examined from the high to middle latitudes of the Northern Hemisphere: Spitsbergen, Norwegian-Greenland Sea, Labrador Sea, western North Atlantic, and the North Sea basin. The genus Svalbardella is emended to comprise species with smooth or finely ornamented surfaces and for which one or both horns are bluntly rounded. Svalbardella clausii sp. nov. has a narrow range restricted to the lowermost Chattian (close to the NP24–NP25 boundary and within Chron C9n), and it therefore appears a useful stratigraphical marker. This species has a wide distribution across the North Atlantic, having been reported from the North Sea basin, western North Atlantic, and the Labrador Sea. Svalbardella clausii sp. nov. overlaps stratigraphically with the reoccurrence interval of Svalbardella cooksoniae Manum, 1960, and spans the Oi-2b cooling maximum. Its presence may therefore be related to the establishment of cooler surface waters at this time.

Svalbardella kareniae sp. nov. has a discordant occurrence: Lower Oligocene and Lower Miocene of the Norwegian Sea at Deep Sea Drilling Project Site 338 and Ocean Drilling Program Site 643, respectively, and mid-Oligocene of the North Sea. Its distribution suggests that Svalbardella kareniae sp. nov. favours more open marine conditions.

Palaeocystodinium obesum Fensome et al., 2009, described from offshore eastern Canada where it has a highest occurrence in the Lower Oligocene, is emended to include specimens with a finely ornamented periphragm and traces of tabulation in addition to the archeopyle.

1 Introduction

The fossil dinoflagellate cyst genus Svalbardella Manum, 1960, is known from the Middle Eocene to Oligocene primarily of the Northern Hemisphere: Spitsbergen (Manum, 1960), Western Siberia (Iakovleva, 2011), Greenland (Birkenmajer et al., 2010; Nør-Hansen, 2003), the Norwegian-Greenland Sea (Eldrett et al., 2004, 2019; Śliwińska and Heilmann-Clausen, 2011), Arctic Canada (Ioannides, 1986), the North Sea basin (Heilmann-Clausen and Van Simaëys, 2005; Schiøler, 2005; Śliwińska, 2019; Van Simaëys et al., 2005), the western North Atlantic (Egger et al., 2016; Firth et al., 2013; Head and Norris, 1989), and the Paratethys (e.g. Pross et al., 2010). There are, however, a few records of Svalbardella species from the southern high latitudes (e.g. Brinkhuis et al., 2003; Van Simaëys et al., 2005).

Currently only two species are formally assigned to this genus, Svalbardella cooksoniae Manum, 1960, and Svalbardella partimtabulata Heilmann-Clausen and Van Simaëys, 2005. Svalbardella cooksoniae ranges from Chron C19r (~42.2 Ma; Eldrett et al., 2004) to Chron C9n (Cocclioni et al., 2018; Pross et al., 2010), whereas Svalbardella partimtabulata is presently known only from a narrow in-
terval within the Middle Eocene (Chron C18r–C18n; mid-Bartonian; Heilmann-Clausen and Van Smaey, 2005; Śliwińska et al., 2016; Thomsen et al., 2012) of Denmark. Notably, in the Oligocene succession of the North Sea basin, the Norwegian-Greenland Sea, and the Paratethys, *Svalbardella cooksoniae* occurs in several discrete intervals. Three of these intervals are synchronous with Oligocene cooling maxima (Śliwińska, 2019; Śliwińska and Heilmann-Clausen, 2011; Van Smaey et al., 2005). Morphologically, the genus *Svalbardella* is most similar to *Palaeocystodinium* Alberti, 1961. Common features are as follows: (i) standard peridiniacian tabulation (Fensome et al., 1993) and fusiform shape, (ii) single apical and antapical horns, (iii) cornucovate wall, (iv) an ellipsoidal endocyst, and (v) a 2a intercalary archeopyle. It is generally accepted that *Svalbardella* differs from *Palaeocystodinium* in having bluntly rounded apical and antapical horns and indications of tabulation other than the archeopyle. However, the distinction between the two genera based on these two features is not straightforward. Although expression of tabulation is not in the original diagnosis of *Svalbardella*, the description of the genus does state that “there are indications of plates and a longitudinal furrow [sulcus] in the type species” (Manum, 1960, p. 21). However, under the description of the type species, *Svalbardella cooksoniae*, “Some specimens are recognizable by their shape only, the fine ornamentation and other details not having preserved” (Manum, 1960, p. 22). Moreover, even though in the original generic description of *Palaeocystodinium* the tabulation is expressed by archeopyle alone, several species assigned to this genus show signs of additional tabulation. The cingulum and sulcus are typically observed in *Palaeocystodinium hampedenense* (Wilson, 1977) Wrenn and Hart, 1988, and sporadically in *Palaeocystodinium australinum* (Cookson, 1965) Lentin and Williams, 1976. Both species were originally assigned to *Svalbardella*. Regarding *Palaeocystodinium minor* Strauss in Strauss et al., 2001, specimens “may display very faint longitudinal parasutural ledges at the horn bases” (Strauss et al., 2001, p. 407).

Already in 1967, Evitt (1967, p. 37) considered *Palæocystodinium* to be a junior synonym of *Svalbardella*. Further, Lindgren (1984) argued that discriminating two taxa at the rank of genus simply because of slight variability in tabulation and horn shape is not reasonable (p. 186), and the author thus suggested that *Svalbardella* is the senior synonym of *Palaeocystodinium*. Wrenn and Hart (1988) opposed that notion and argued that the presence of tabulation and broad, rounded horns which are so characteristic of *Svalbardella* are not highly variable features. Furthermore, the authors observed that “horns in various species of *Palaeocystodinium* may be pointed or bluntly round, but they are still thin and taper markedly from the area of the central body to their terminations” (Wrenn and Hart, 1988, p. 361). However, in *Svalbardella partimtabulata* Heilmann-Clausen and Van Smaeys (2005) and *Svalbardella kareniae* sp. nov. (this study) the antapical horn is long and tapers gradually.

The most recent emendation of *Palaeocystodinium* restricts this genus to forms with pointed horns (Fensome et al., 2009). However, the literature reveals several distinctive morphotypes with at least one bluntly rounded horn that are assigned either to *Svalbardella* or *Palaeocystodinium* (e.g. Damassa et al., 1990; Egger et al., 2016; Poulsen et al., 1996; Schüster, 2005; Śliwińska et al., 2012) but have never been formally described as species.

In this study, we accept that the presence of at least one bluntly rounded horn is sufficiently distinctive to separate *Svalbardella* from *Palaeocystodinium*. Accordingly, *Svalbardella* is amended to include species with a single bluntly rounded horn. The present study describes two new species, *Svalbardella clausii* sp. nov. and *Svalbardella kareniae* sp. nov., from the Oligocene of the North Atlantic region, and discusses their stratigraphical and palaeoenvironmental significance. Additionally, *Palaeocystodinium obsenum* Fensome et al., 2009, is emended.

### 2 Material and methods

This research is based on samples from Spitsbergen, the Norwegian-Greenland Sea, the North Sea basin, the Labrador Sea, and western North Atlantic. The locations of those North Atlantic–Arctic sites where the genus *Svalbardella* has been reported, including the present study, are shown in Fig. 1 and listed in Table 1. Palynological slides from Sarsbukta, Spitsbergen, studied previously by Manum (1960) were reanalysed by Kasia K. Śliwińska in 2011 during a research stay at the Natural History Museum in Oslo, Norway. The preparation methods are described in Manum (1960). During this visit, palynological slides from Renardodden, Spitsbergen, studied previously by Head (1984), were also examined (and subsequently reexamined by Martin J. Head). Palynological slides from Ocean Drilling Program (ODP) Site 634 in the Norwegian Sea were loaned from the Natural History Museum in Oslo, their preparation having been described in Manum et al. (1989). Palynological slides from the offshore 11/10-1 well, central North Sea, were borrowed from the Norwegian Petroleum Directorate (NPD); the preparation methods for these are described in Śliwińska (2019). Palynological slides from the offshore Nini-I well in the eastern North Sea were also examined, as were palynological slides from the Harre-1 borehole, onshore Denmark. The preparation methods for these slides were described by Śliwińska et al. (2010, 2012). The palynology of Deep Sea Drilling Project (DSDP) sites from the northern North Atlantic and Labrador Sea was previously studied by Damassa et al. (1990). The present research also uses palynological slides from Integrated Ocean Drilling Program (IODP) sites U1411 and U1406 in the western North Atlantic, which were prepared at the Geological Survey of Denmark and Greenland (GEUS) and included in the study by Egger et al. (2016). Several new
palynological slides from IODP Hole 1411B were prepared for the present study, following the processing methods described in Egger et al. (2016).

Key taxa discussed in the text are presented in Table 1 and on Plates 1–5. The position of each of the photographed specimens is indicated by microscope coordinates (MC; the A point = 0 : 4 × 90 : 3 [XM1 × YM1] following the method described in Śliwińska, 2019) and/or an England Finder (EF) reference. The horn length is measured from the tip of the endocyst to the corresponding tip of the pericyst. The type material and the illustrated dinocysts marked with MGUH numbers are stored in the type collection of the Natural History Museum of Denmark (Øster Voldgade 5–7, DK-1350 Copenhagen K, Denmark). The type material and the illustrated dinocyst marked with a PMO number are stored in the type collection of the Natural History Museum in Oslo (Sars’ gate 1, NO-0562 Oslo, Norway).

3 Formal description of new species

Division Dinoflagellata (Bütschli, 1885) Fensome et al., 1993
Subdivision Dinokaryota Fensome et al., 1993
Class Dinophyceae Pascher, 1914
Subclass Peridiniphycideae Fensome et al., 1993
Order Peridiniales Haeckel, 1894
Family Peridiniaceae Ehrenberg, 1831
Subfamily Deflandreoideae Bujak and Davies, 1983
Genus Svalbardella Manum, 1960, emend.

Type
Manum, 1960, pl. 1, figs. 1–3; text-fig. 2, as Svalbardella cooksoniae.

Original diagnosis
Shells of planktonic microorganisms. Shape fusiform with somewhat swollen middle part and blunt ends. No appendages. Girdle approximately equatorial. Middle part of shell entirely filled by a thin-walled ellipsoidal body (Manum, 1960, p. 21).

Emended diagnosis
Cornucavate cysts with a fusiform shape, a single apical and antapical horn, and an ellipsoidal endocyst. One or both horns are conspicuously bluntly rounded. Surface smooth or bears low ornamentation. Cingulum when present is approximately equatorial. Archeopyle 2a intercalary. Other traces of tabulation may be present.

Accepted species
Svalbardella clausii sp. nov., Svalbardella cooksoniae Manum, 1960, Svalbardella kareniae sp. nov., Svalbardella partimtabulata Heilmann-Clausen and Van Simaeyns, 2005.

Remarks
The genus is emended to contain species in which one (e.g. Svalbardella kareniae sp. nov.) or both (e.g. Svalbardella cooksoniae) horns are bluntly and conspicuously rounded.

Stratigraphic range
The oldest stratigraphic appearance of the genus Svalbardella is reported from Chron C19r; late Lutetian (as Svalbardella cooksoniae) by Eldrett et al. (2004). The youngest stratigraphic appearance is reported from the Early Miocene A. granosa zone of Manum et al. (1989) in ODP Hole 643A (as Palaeocystodinium sp. 2, now Svalbardella kareniae sp. nov.).

Svalbardella clausii sp. nov.
Plate 1A–I
1990 Palaeocystodinium sp. 2; Damassa et al., fig. 5M.
2005 Palaeocystodinium sp. 2 of Damassa et al. (1990); Schröter, pl. 10, figs. 6–7.
2010 Svalbardella cooksoniae auct. non Manum, 1960; Pross et al., pl. IV, fig. 5.
2010 Svalbardella sp. 1; Śliwińska et al., pl. 1, fig. 4.

Derivation of name
After Claus Heilmann-Clausen, the PhD supervisor of Kasia K. Śliwińska, for his important contributions to the study of Cretaceous and Paleogene dinoflagellate cysts.

Diagnosis
A species of Svalbardella with rounded, short apical and antapical horns of approximately equal length. The endocyst is broad with a length approximately 1.5 times the width. The endocyst is ellipsoidal and has a smooth wall. The periphragm is nearly smooth to microrugulate or incompletely microreticulate. On more coarsely ornamented specimens, tabulation may be expressed by the cingulum and/or plates 1a, 3a, 3–5′, and/or 1–2′′ in addition to the archeoype. The archeoype is intercalary type I (2a); adcingular margin of the periarcheopyle slightly overlaps that of the endoarcheopyle (operculum free).

https://doi.org/10.5194/jm-39-139-2020 J. Micropalaeontology, 39, 139–154, 2020
Table 1. Drill sites and outcrops yielding specimens here attributed to the genus *Svalbardella* from the North Atlantic–Arctic region illustrated in Fig. 1.

| Drill site/outcrop | Taxon | Reference |
|--------------------|-------|-----------|
| Sarsbukta          | *Svalbardella cooksoniae* holotype | Manum (1960) |
| Renardodden        | *Svalbardella* sp. cf. *S. partimtabulata* | Head (1984), this study |
| DSDP 908           | *Svalbardella* sp. 1, *Svalbardella* spp. | Eldrett et al. (2019), Poulsen et al. (1996) |
| ODP 913            | *Svalbardella cooksoniae*, *Svalbardella* cf. *S. cooksoniae* | Eldrett et al. (2004), Firth (1996) |
| DSDP 338           | *Svalbardella kareniae* sp. nov. (as *Svalbardella* (*Palaeocystodinium*) spp.) | Eldrett et al. (2004), Manum (1976), Śliwińska and Heilmann-Clausen (2011) |
| ODP 643A           | *Svalbardella* sp. cf. *S. cooksoniae*, *Svalbardella* sp. 1, *Svalbardella* sp. 2 | Eldrett et al. (2004), Head and Norris (1989) |
| ODP 985            | *Svalbardella cooksoniae* | Eldrett et al. (2019), Williams and Manum (1999) |
| Hellefisk-1        | *Svalbardella* sp. cf. *S. cooksoniae* | Nøhr-Hansen et al. (2003) |
| Kap Brewster       | *Svalbardella* sp. | Birkenmayer et al. (2010) |
| 6609/11-1, 6610/2-1S, 6407/9-2, and 6407/9-5 | *Svalbardella cooksoniae* | Eidvin et al. (2007) |
| 11/10-1            | *Svalbardella cooksoniae* | Śliwińska (2019), Śliwińska and Heilmann-Clausen (2011) |
| Alma-1X            | *Svalbardella cooksoniae*, *Svalbardella* cf. *cooksoniae*, *Svalbardella clausii* sp. nov. (as *Palaeocystodinium* sp. 2 of Damassa et al., 1990) | Schiessler (2005), Śliwińska and Heilmann-Clausen (2011) |
| Nini-1             | *Svalbardella clausii* sp. nov. holotype (as *Svalbardella* sp. 1), *Svalbardella sp.*, *Svalbardella cooksoniae* | Claussn et al. (2012), Śliwińska (2019), Śliwińska et al. (2010, 2014a) |
| Mona-1             | *Svalbardella* sp. | Śliwińska and Heilmann-Clausen (2011), Van Simaeys et al. (2005) |
| Harre-1            | *Svalbardella kareniae* sp. nov. (as *Palaeocystodinium* sp. 1), *Svalbardella cooksoniae*, *Svalbardella* sp. 1 | Śliwińska et al. (2012), this study |
| Kysing-4           | *Svalbardella partimtabulata* holotype | Heilmann-Clausen and Van Simaeys (2005) |
| Kallo, Hingene-Wintham boring 42E-212 | *Svalbardella* sp. indet. | De Coninck (1996) |
| Leluchów           | *Svalbardella* sp. | Gedl (2004) |
| Umbria-Marche (incl. the Massicore borehole, as well as Contessa, Pieve d’Accinelli, and Monte Cagnero outcrop sections) | *Svalbardella cooksoniae*, *Svalbardella* sp. | Van Mourik and Brinkhuis (2005), Van Simaeys et al. (2005) |
| ODP 647A           | *Svalbardella cooksoniae*, *Svalbardella* sp. cf. *S. cooksoniae*, *Svalbardella* sp. 1, *Svalbardella* sp. 2 | Firth et al. (2013), Head and Norris (1989) |
| DSDP 112           | *Svalbardella clausii* sp. nov. (as *Palaeocystodinium* sp. 2) | Damassa et al. (1990), this study |
| Cumberland B-55    | *Svalbardella* sp. | Barss et al. (1979) |
| IODP U1405, U1406, U1411 | *Svalbardella* sp. | Egger et al. (2016) |
Figure 1. The distribution of drill sites and outcrops where Svalbardella specimens have been reported (see Table 1). Modified from Egger et al. (2016). Reprinted from Review of Palaeobotany and Palynology, 234, Egger, L. M., Śliwińska, K. K., van Peer, T. E., Liebrand, D., Lippert, P. C., Friedrich, O., Wilson, P. A., Norris, R. D., and Pross, J.; Magnetostratigraphically-calibrated dinoflagellate cyst bioevents for the uppermost Eocene to lowermost Miocene of the western North Atlantic (IODP Expedition 342, Paleogene Newfoundland sediment drifts), 159–185, Copyright (2016), with permission from Elsevier.

Holotype

Slide 158L-2; MGUH no. 33445, EF reference S23/1, MC 23 × 98.2. Ditch cutting sample from the eastern North Sea basin, Nini-1 well; depth: 1100.0 m below kelly bushing (mbKB). Dated to earliest Chattian. Slide is curated at the Geological Museum of the University of Copenhagen. Plate 1A–C.

Paratypes

Slide 27227-3; MGUH no. 33446, MC 21.3 × 105.1, EF reference K20/3. IODP Leg 342, Site U1411; Sample 342-U1411B-6H-4, 24–26 cm; depth: 43.64 m b.s.f. Offshore Newfoundland. Earliest Chattian. Slide is curated at the Geological Museum of the University of Copenhagen. Plate 1F.

Slide P-76119; MGUH no. 33447; MC 35.5 × 110.3, EF reference E36/3. DSDP Site 112, Sample 112-5-1, 23–30 cm. Western North Atlantic. Mid-Oligocene, Plate 1G.

Description

A fusiform peridiniacean species with apex and antapex, each having a single short and bluntly rounded horn. The cyst is long, wide, and cornucavate. The endocyst is ellipsoidal and has a smooth wall. The periphragm is nearly smooth to faintly or conspicuously ornamented with a microrugulate pattern that may include discrete and partially fused granules and may develop into an incomplete microreticulation. The endophragm and periphragm are appressed except at the horns. The horns are rounded to bulbous in shape. A small tapering antapical or apical protuberance up to 7 µm long may occur at the termination of a horn (marked by an arrow in Plate 1E and H). Horns are approximately equal in length, and each is generally less than 30% of the endocyst length. When the periphragm is nearly smooth, the tabulation is expressed only by the archeopyle. On more coarsely ornamented specimens, the tabulation may be expressed by the cingulum and/or plates 1a, 3a, 3–5'' and/or 1–2'' in addition to the archeopyle. The archeopyle is intercalary type I (2a), isodeltaform (operculum free). The periarcheopyle is of similar size or slightly larger than the endoarcheopyle. The adcingular margin of the periarcheopyle is up to ~1.5 µm higher than that of the endoarcheopyle (Plate 1F).

Dimensions

Holotype: endocyst length 87 µm; pericyst length 122 µm; width 57 µm; apical horn length 20 µm; antapical horn length 20 µm. Range (n = 12): endocyst length 75–98 µm (mean 85.2 µm); pericyst length 100–150 µm (mean 120.2 µm); width 52–65 µm (mean 57.5 µm); apical horn length 12–25 µm (mean 19.4 µm); antapical horn length 15–30 µm (mean 20.2 µm).
Plate 1. A scale bar of 20 µm applies to all figures except where stated. (A–I) Svalbardella clausii sp. nov. (A–C) Holotype, (A) ventral surface and (B) dorsal surface; Nini-1 well; depth: 1100.0 mbKB; slide K158L-2; MGUH no. 33445; MC: 23 × 98.2; EF: S23/1; cyst length: 122 µm; (C) tracing of (B) showing expression of tabulation around the archeopyle (operculum partially detached). (D) DSDP Site 112, Sample 112-5-1, 23–30 cm; slide P-76119; MC: 36.4 × 102; EF: O36/2; (E) DSDP Site 112, Sample 112-5-1, 23–30 cm; slide P-76119; MC: 35 × 102.2 EF: O35/1; (F) Paratype, dorsal surface; IODP Site U1411, Sample 342-U1411B-6H-4, 24–26 cm; depth: 43.64 m b.s.f.; slide 27227-3; MGUH no. 33446, MC: 21.3 × 105.1; EF: K20/3; cyst length 112 µm. (G) Paratype DSDP Site 112, dorsal surface. Sample 112-5-1, 23–30 cm; slide P-76119; MGUH no. 33447; MC: 35.5 × 110.3; EF: E36/3; cyst length: 115 µm. (H) Dorsal surface; DSDP Site 112, Sample 112-5-1, 23–30 cm; slide P-76119; MC: 43 × 97 EF: T44/1; cyst length 150 µm. (I) Dorsal surface; IODP Site U1406, Sample 342-U1406B-17H-4, 76–79 cm; depth: 148.76 m b.s.f.; slide 25266-4; MC: 28.7 × 103.3; EF: M28/4; cyst length 100 µm.

Occurrence

North Sea basin, Alma-1x and Nini-1 wells; western North Atlantic, Hole IODP U1411B (40.6 and 53.1 metres below sea floor, m b.s.f.; Chron C9n), Hole IODP U1406B (148.76 m b.s.f.; Chron C9n), and Hole IODP U1406C (141.31 and 142.82 m b.s.f.; Chron C9n); Labrador Sea, DSDP Sites 112 and 119.

Age

Existing records suggest that Svalbardella clausii sp. nov. is limited to a narrow range within the mid-Oligocene. The widest and least specific range for Svalbardella clausii sp. nov. is provided by Damassa et al. (1990; as Palaeocystodinium sp. 2), who observed the lowest occurrence (LO) somewhere in NP24 and the highest occurrence (HO) somewhere within NP25 (a maximum time span between approximately 29.5 and 23 Ma; Vandenberghe et al., 2012) in the
western North Atlantic. In the two studied sites from offshore Newfoundland, IODP Sites U1411 (Hole U1411B, 40.6 and 53.1 m b.s.f.; Chron C9n) and U1406 (Hole U1406B, 148.76 m b.s.f.; Hole U1406C, 141.31 and 142.82 m b.s.f.), the species ranges within Chron C9n (Site U1411 magnetostratigraphy after Norris et al., 2014; Site U1406 magnetostratigraphy after van Peer et al., 2017) and thus is of earliest Chattian age.

In the North Sea, this species is reported in two industrial wells (Nini-1 and Alma-1x), from which only ditch cuttings are available. Due to the high risk of caving we focus here on the highest occurrences (HOs). The HO of Svalbardella clausii sp. nov. is observed either at the Rupelian/Chattian boundary (Schiøler, 2005) or in the lowermost Chattian (Śliwińska et al., 2010, 2014a). In both wells, the HO of Svalbardella clausii sp. nov. occurs below the HO of Licracysta? semicirculata (Schistler, 2005; Śliwińska et al., 2010), which in the North Sea basin is a well-established lowermost Chattian marker occurring within Chron C8r (Śliwińska et al., 2012). The range of Svalbardella clausii sp. nov. is synchronous with the mid-Oligocene Svalbardella cooksoniae occurrence interval, which in the Nini-1 well can be correlated with the uppermost NP 24 zone to lowermost NP25 zone (Śliwińska et al., 2014a).

Comparison

Svalbardella partitabulata is significantly more slender, has a smaller endocyst, and relatively longer horns, with the antapical horn being significantly longer than the apical horn. Svalbardella cooksoniae has a slightly longer but significantly narrower endocyst and longer and more conical horns in contrast with the short, almost bulbous horns in Svalbardella clausii. Palaeocystodinium obesum is also broad and has very short horns, but they are pointed and slightly smaller in size. However, forms transitional between Palaeocystodinium obesum and Svalbardella clausii sp. nov. (Plate 5D) have been observed in the lower Chattian in the western North Atlantic (e.g. IODP Hole U1406B, Sample 342-U1406B-17-4; 76–79 cm).

Svalbardella kareniae sp. nov.  
Plate 2A–E and H–J

1972 Palaeocystodinium golzwense auct. non Alberti, 1961; Benedek, pl. 3, fig. 7.  
1976 Svalbardella (Palaeocystodinium) spp.; Manum, pl. 5, fig. 1 only.  
1989 Palaeocystodinium sp. 2; Manum et al., pl. 19, fig. 7.  
2005 Palaeocystodinium sp. 1 of Manum et al., 1989; Schistler, pl. 10, fig. 1  
2012 Palaeocystodinium sp. 1; Śliwińska et al., pl. II, fig. 7.

Derivation of name

After Karen Dybkjær, the PhD co-supervisor of Kasia K. Śliwińska, for her important contributions to the study of Jurassic and Miocene dinoflagellate cysts.

Diagnosis

A species of Svalbardella with a short and blunt apical horn and long and pointed antapical horn. The periphragm has a smooth to finely ornamented surface; the endophragm is smooth and slightly thicker than the periphragm. The antapical horn is at least 1.5 times as long as the apical horn.

Holotype

Slide 158L-2, MGUH 33448, EF reference T27/3, MC 27 × 96.7. Ditch cutting sample from the Nini-1 well, the eastern North Sea basin; depth: 1100.0 mbKB. Lowermost Chattian, from the uppermost part calcareous nannofossil zone NP24, correlated to Chron C9n (Śliwińska et al., 2010, 2014a). Slide is curated at the Geological Museum of the University of Copenhagen (Plate 2A, B).

Paratype

Slide 4879iii, PMO 235.064, EF reference M43/3. ODP Leg 104, Hole 643A, Sample 643A-40-1, 55–57 cm; Norwegian Sea; depth: 371.95 m b.s.f. Early Miocene (Manum et al., 1989). Slide is curated at the Natural History Museum in Oslo, Norway (Plate 2H).

Description

Peridiniacean cyst comprising an ellipsoidal endocyst and fusiform pericyst. The pericyst has a single apical and antapical horn. Periphragm and endophragm are closely appressed except at the horns. The periphragm is smooth to irregularly and minutely reticulate. The endophragm is smooth and slightly thicker than periphragm. The apical horn is blunt, and the termination is tapering with a rounded tip. The antapical horn is conical, pointed distally, and at least 1.5 times longer than apical horn. Tabulation expressed by archeopyle alone, which is intercalary type I (2a), isodeltaform (operculum free).

Dimensions

Holotype: endocyst length 79 µm; pericyst length 156.5 µm; width 36 µm; apical horn length 28.5 µm; antapical horn length 49 µm. Range (n = 8): endocyst length 53–79 µm (mean 69.6 µm); pericyst length 115–166 µm (mean 148.4 µm); width 28–36.5 µm (mean 33.0 µm); apical horn length 14–34 µm (mean 26.2 µm); antapical horn length 34–66 µm (mean 53.8 µm).
Plate 2. (A–E) Svalbardella kareniae sp. nov. (A–B) Holotype, (A) dorsal surface and (B) ventral surface; Nini-1; depth: 1100 m; slide 158L-2; MGUH 33448; MC: 27 × 96.7; EF: T27/3; cyst length: 156.5 µm. (C) Ventral surface and (D) dorsal surface; Harre-1 borehole; depth: 105.25 m; slide K94F-1; EF: S53/0; cyst length: 115 µm (the same specimen as photographed in Śliwińska et al., 2012; see their pl. 2.7). (E) Archeopyle is seen to the right; the Hesselbjerg outcrop section, level −11.0 m, slide 171K-1; MC: 25 × 99; EF: R25/1; cyst length: 143.5 µm. (F–G) Svalbardella cooksoniae Manum, 1960. (F) Dorsal surface with tabulation around the cingulum; ODP Site 643, Sample 643A-50-1, 30–32 cm; depth: 468.40 m b.s.f.; slide 4668iv; EF: O37/3; cyst length: 123.2 µm. (G) Dorsal surface with no tabulation in the cingular area; 11/10-1 well; depth: 823.0 m b.s.f.; MC: 47.5 × 104.5; EF: L48/3. (H–J) Svalbardella kareniae sp. nov. (H) Paratype, dorsal surface, ODP Site 643, Sample 643A-40-1, 55–57 cm; depth: 371.95 m b.s.f.; slide 4879iii; PMO: 235.064; EF: M43/3; cyst length: 161.7 µm. (I) Ventral surface, (J) dorsal surface; ODP Site 643, Sample 643A-40-1, 55–57 cm; depth: 371.95 m b.s.f.; slide 4879iii; EF: K42/0; cyst length: 161.5 µm.
Occurrence

Norwegian Sea, DSDP Leg 38, Hole 338, Sample 338-26-5, 100–102 cm – as “Svalbardella (Palaeocystodinium) spp.” in Manum, 1976, pl. 5, fig. 1 only; Norwegian Sea, ODP Leg 104, Hole 643A, Samples 42-1, 19–21 cm (390.89 m b.s.f.), to 37-3, 20–22 cm (345.50 m b.s.f.), inclusive (as Palaeocystodinium sp. 2 in Manum et al., 1989); the North Sea, Alma-1x well, recorded in ditch cutting samples between 1844.0 and 1725.2 m (as Palaeocystodinium sp. 1 of Manum et al., 1989, in Schiøler, 2005); onshore Denmark in the Viborg-1 borehole between 164.0 and 133.5 m, in the Harre-1 borehole between 113.75 and 70.75 m, and in the Hesselbjerg outcrop section from −9.0 to 7.5 m (as Palaeocystodinium sp. 1 in Śliwińska et al., 2012). In Denmark the sediments yielding Svalbardella kareniae sp. nov. belong to the Branden Clay unit (see Śliwińska et al., 2012).

Age

In the Norwegian Sea the species is observed in two Holes. In DSDP Hole 338 in Sample 338-26-5, 100–102 cm, the sample was initially dated as Late Eocene (Manum, 1976). However, more recently Eldrett and Harding (2009) suggested an Early Oligocene age for the sample. In another hole, ODP Hole 643A, the species is recorded in the Early Miocene (A. granosa zone of Manum et al., 1989, above the HO of Chiropteridium spp.; Manum et al., 1989; Śliwińska et al., 2014b). In the North Sea, the species is observed in the Rupelian and Chattian stages of the Oligocene (Schiøler, 2005; Śliwińska et al., 2012).

Comparison

Svalbardella partimbulata also has a short apical and a long antapical horn, but faint sutural lines or low ridges on the periphragm surface partially delimit the tabulation. Also the apical horn tends to be relatively longer than for Svalbardella kareniae sp. nov. and the central body shorter relative to the total length. Svalbardella cooksoniae has coarser ornamentation on the periphragm and wider horns with apical and antapical horns being of similar lengths. Svalbardella clausii sp. nov. has significantly shorter horns that are both rounded and equal in length. The shape of the antapical horn in Svalbardella kareniae sp. nov. resembles that in Palaeocystodinium golzowense.

Description

The periphragm is about 0.3µm thick and has an outer surface with low microrugulae with interspersed granules, in places forming a discontinuous micoreticulum. The endophragm is about 0.6µm thick and smooth. Archeopyle intercalary, type I (2a), periarcheopyle slightly larger than endoarcheopyle, its adcingular margin overlapping that of the endoarcheopyle by about 2µm (operculum free). The tabulation is also indicated by sutural features on the hypocyst expressed as rudimentary septa.

Dimensions

Endocyst length: 62µm; pericyst length: 151µm; width: 22µm; apical horn length: 31µm; antapical horn length: 61µm. One specimen measured.

Occurrence

A single specimen, slide No. MCP 2276 (5), Renardodden, Spitsbergen. The slide is curated at Brock University.

Age

The sample is dated palynologically as Late Eocene or Early Oligocene (Head, 1984).

Remarks

This specimen is most similar to Svalbardella partimtabulata but has a relatively longer endocyst and relatively shorter apical horn. Additionally, the boundary between the postcingular and antapical plate series (marked by an arrow in Plate 3A, B) occurs at the approximate level of the endocyst antapex, whereas in Svalbardella partimtabulata it is much lower. This may be a consequence of the relatively longer endocyst in the Spitsbergen specimen.
Plate 3. (A–F) Svalbardella sp. cf. S. partimtabulata Heilmann-Clausen and Van Simaeys, 2005. Right lateral view (A–D) at upper- through mid-foci, with margin between postcingular and antapical plate series shown by arrows (A, B), adcingular margin of periarcheopyle overlapping that of endoarcheopyle shown by arrow (D), and detail of wall surface on central body (E) and antapical horn (F); Renardodden, Spitsbergen; slide MCP 2276(5); EF: M41/2; cyst length: 151 µm. Scale bar = 2 µm in (E), (F).

Genus Palaeocystodinium Alberti, 1961

Type

Alberti, 1961, pl. 7, fig. 12, as Palaeocystodinium golzowense.

Palaeocystodinium obesum Fensome et al., 2009, emend.

Plates 4A–H and 5A–C

1972 Svalbardella cf. granulata; Benedek, p. 14, pl. 3, fig. 5a, b.
1990 Palaeocystodinium sp. 1; Damassa et al., fig. 5 G, J.
1996 Svalbardella sp. 1; Poulsen et al., pl. 3, fig. 13.
2009 Palaeocystodinium obesum Fensome et al., p. 50, pl. 8, figs. j–l.

Original description

A species of Palaeocystodinium with relatively short but pointed horns, a smooth periphragm, and a broad endocyst. Bicavate, with pericoels restricted to polar regions; wall layers appressed laterally (Fensome et al., 2009, p. 50).

Emended diagnosis

A species of Palaeocystodinium with short, pointed horns, a nearly smooth to microrugulate–microreticulate periphragm and a broad endocyst. Faint tabulation, including cingulum, may be expressed. Cornucavate, with pericoels restricted to polar regions; wall layers appressed laterally. The $2a/5'$ margin is considerably longer than the $2a/3'$ margin.

Emended description

Peridiniacean cyst comprising broad ellipsoidal endocyst and short, pointed horns. Periphragm has a nearly smooth to finely ornamented periphragm. This fine ornament varies...
from a complete microreticulation (apparently present on the holotype, Plate 5A) to a microrugulate pattern that forms an incomplete microreticulation (Plate 4A). When the periphragm is smooth or faintly ornamented, the tabulation is expressed by the archeopyle only. On more prominently ornamented specimens, tabulation may be expressed also by a cingulum, and on some specimens plates 2–3′, 1a, 3a, 2–5″, and/or 1–2‴ are distinguishable (Plate 4E–G) in addition to the archeopyle. The 2a/3′ margin is relatively long, and the 2a/5″ margin is considerably longer than the 2a/3″ margin (Plate 4A, B, D–H). Expression of tabulation is, however, a variable feature. The antapical horn may be asymmetrical owing to a “bulge”, as shown in Plate 4E–H (this study), in Benedek (1972, pl. 3 fig. 5), in Damassa et al. (1990, fig. 5J), and on the holotype itself (Plate 5A, also in Fensome et al., 2009, pl. 8, fig. 1). Archeopyle intercalary, type I (2a), isodeltaform (operculum free).
Plate 5. (A–C) *Palaeocystodinium obesum* Fensome et al., 2009, emend. (A) Holotype, with microreticulate surface ornamentation and distinctive bulge on antapical horn indicated by arrow; sidewall core sample at 751.6 m (2466 ft) in Onondaga E-84 well, Scotian Margin off eastern Canada; GSC type collection no. 130270; slide P37515; slide 01; EF: O49/2; total length: 129 µm (photo supplied by Robert A. Fensome; see Fensome et al., 2009, pl. 8, fig. l). (B) Topotype specimen; sidewall core sample at 751.6 m (2466 ft) in Onondaga E-84 well, Scotian Margin off eastern Canada; GSC type collection no. 130269; sample P37515; slide 01; EF: L51/0; total length: 94 µm (photo supplied by Robert A. Fensome; see Fensome et al., 2009, pl. 8, fig. k). (C) Topotype specimen; sidewall core sample at 751.6 m (2466 ft) in Onondaga E-84 well, Scotian Margin off eastern Canada; GSC type collection no. 130268; sample P37515; slide 01; EF: K40/4; total length: 115 µm (photo supplied by Robert A. Fensome; see Fensome et al., 2009, pl. 8, fig. j). (D) Specimen transitional between *Svalbardella clausii* sp. nov. and *Palaeocystodinium obesum*; dorsal surface; IODP Site U1406, Sample 342-U1406B-17H-4, 76–79 cm; depth: 148.76 m b.s.f.; slide 25266-4; MC: 55 \times 100; EF: Q56/1; total length: 117 µm. Figures (A)–(C) are reprinted from “Late Cretaceous and Cenozoic fossil dinoflagellates and other palynomorphs from the Scotian Margin, offshore eastern Canada” by Fensome, R. A., Williams, G. L., and MacRae, R. A.; Journal of Systematic Palaeontology, copyright 2009 © The Natural History Museum, reprinted by permission of Informa UK Limited, trading as Taylor & Francis Group, https://www.tandfonline.com on behalf of The Natural History Museum.

Dimensions

**Holotype:** endocyst length 92 µm; pericyst length 129 µm, overall width 45 µm (Fensome et al., 2009). **Range (this study):** endocyst length 69–104 µm (mean 81.9 µm); pericyst length 92–137 µm (mean 110.8 µm); width 37–56 µm (mean 49.3 µm); apical horn length 8–19 µm (mean 13.9 µm); antapical horn length 9–20 µm (mean 16.1 µm). Twelve specimens from DSDP Site 112 and IODP Hole 1411B, showing slightly wider size ranges than reported by Fensome et al. (2009) based on three specimens.

Occurrence

In addition to the Scotian Margin, which is the type locality, this species is present off Newfoundland (IODP Hole U1411B), the Greenland Sea (ODP Site 908), the Norwegian Sea (DSDP Site 338), and the North Sea (Nini-1) (Damassa et al., 1990; Poulsen et al., 1996; this study).

Age

On the Scotian Margin off Canada, from where the holotype is described, the HO is in the upper Rupelian (Fensome et al., 2009). In the North Atlantic *Palaeocystodinium obesum* (as *Palaeocystodinium sp. 1*) is reported from NP19/20 to NP24, suggesting a range from Priabonian to mid-Oligocene (Damassa et al., 1990). Off Newfoundland, this species ranges from Chron C12r to Chron C9n, although deposits older than earliest Oligocene were not investigated (this study). In the Greenland Sea and the North Sea, it is observed in the Chattian (Poulsen et al., 1996; this study).
Remarks

The holotype bears faint lineations on the hypocyst running from the central body down to the antapical horn that might represent tabulation. Re-examination of the specimen will be needed to confirm this. This species is emended to include specimens having a microrugulate–microreticulate periphragm and traces of tabulation in addition to the archeopyle.

4 Discussion

The two new species, *Svalbardella clausii* sp. nov. and *Svalbardella kareniae* sp. nov., widen the morphological diversity of the genus *Svalbardella* emend., which until, now has been limited to two formally described species: *Svalbardella cooksoniae* Manum, 1960, and *Svalbardella partimtabulata* Heilmann-Clausen and Van Simaeys, 2005.

Our new observations from the Oligocene succession of the North Sea and offshore Newfoundland require the emendation of *Palaeocystodinium obesum* Fensome et al., 2009, which now includes forms with fine ornament as well as traces of tabulation in addition to the archeopyle. This study also confirms that within both *Svalbardella* and *Palaeocystodinium*, as presently conceived, tabulation is expressed variably, even within individual species (e.g., *Svalbardella clausii* and *Palaeocystodinium obesum*) and may be related to the degree of ornamentation on the periphragm.

*Svalbardella kareniae* provides a link between the genera *Svalbardella* and *Palaeocystodinium*. If a specimen of *Svalbardella kareniae* broke along the cingulum, its antapical horn would easily be confused with that of *Palaeocystodinium golzowense*. The presence of forms transitional between *Palaeocystodinium obesum* and *Svalbardella clausii* (Plate 5D) provides an additional link. This emphasizes the close morphological similarity between *Svalbardella* and *Palaeocystodinium* and implies a close phylogenetic relationship. However, we consider it premature to synonymize these genera while the stratigraphic ranges and fine morphological details of many species are incompletely known. Future studies incorporating both light and scanning electron microscopy that document the fine ornamentation of the periphragm, details of the tabulation where exhibited, and morphology of both the endo- and periarcheopyle, augmented by morphometric studies documenting intraspecific variability—and all accomplished using finely age-calibrated samples—will be needed to understand fully the relationship between these two genera over their ranges.

*Svalbardella clausii* is observed in the northern high and middle latitudes in a very narrow time interval spanning the NP24–NP25 boundary and Chron C9n. The species has a wide spatial distribution across the North Atlantic (North Sea, western North Atlantic, and Labrador Sea). The horizon containing *Svalbardella clausii* is therefore a prime marker for the lower Chattian succession in this region. The range of this species overlaps with the timing of the reappearance of *Svalbardella cooksoniae* and spans the Oi-2b cooling maximum (Śliwińska et al., 2010; Van Simaeys et al., 2005). Therefore, the presence of *Svalbardella clausii* may be related to an establishment of colder surface waters associated with the cooling. This species notably appears not to have penetrated into the Norwegian-Greenland Sea.

*Svalbardella clausii* appears in narrow but diachronous intervals. In the North Sea basin this species is observed in the Oligocene, whereas in the Norwegian Sea it is recorded either from the Lower Oligocene (DSDP Hole 338) or Lower Miocene (ODP Hole 643A). The Branden Clay unit onshore Denmark, where *Svalbardella kareniae* is present, represents a short interval during the Oligocene when conditions were more fully marine (Friis, 1994). The Branden Clay is the only unit in the Danish Oligocene succession that yields well-preserved specimens of the planktonic foraminiferal genus *Globigerina* (Ulleberg, 1987). In DSDP Hole 338, the appearance of *Svalbardella kareniae* coincides with a change to open marine/oceanic conditions, as suggested by the dinoflagellate cyst assemblage (Eldrett and Harding, 2009). In ODP Hole 643A, the appearance of *Svalbardella kareniae* similarly corresponds to a change to more open marine conditions, as suggested by the biomarker record and dinoflagellate cyst assemblages (Śliwińska et al., 2010, 2014a). All these observations imply that *Svalbardella kareniae* was sensitive to changes in the palaeoenvironment, presumably favouring more open marine conditions.

5 Conclusions

*Svalbardella clausii* sp. nov. and *Svalbardella kareniae* sp. nov. are formally described herein, requiring the emendation of the genus *Svalbardella*, and *Svalbardella* sp. cf. *S. partimtabulata* Heilmann-Clausen and Van Simaeys, 2005, is reported from Spitsbergen. This study also emends *Palaeocystodinium obesum* Fensome et al., 2009, which now includes specimens with a finely ornamented periphragm and traces of tabulation in addition to the archeopyle.

The distribution of *Svalbardella clausii* appears to reflect an affinity for colder sea surface temperatures, and that of *Svalbardella kareniae* reflects open marine conditions. The narrow and synchronous range of *Svalbardella clausii* makes it an excellent mid-Oligocene marker for the North Atlantic region.

The morphology of *Svalbardella kareniae* and those forms transitional between *Palaeocystodinium obesum* and *Svalbardella clausii* illustrate the close phylogenetic relationship between *Svalbardella* and *Palaeocystodinium*, these two genera being separated on the presence of bluntly rounded versus pointed terminations of the horns. This study also clearly illustrates that surface ornamentation and the tabulation it may express are variable features within both genera.
Data availability. Palynological slides are stored at the Natural History Museum in Oslo, Norway (643A), GEUS (Nini-1, Harre-1 U1411, U1406), the Norwegian Petroleum Directorate (11/10-1 well), and Brock University (Renardodden).

Author contributions. KKŚ designed the project and carried out the analysis (Harre-1, Hesselbjerg, Nini-1, 11/10-1, Site U1411, Site U1406, Site 112, Site 338, and Hole 643A) together with MJH (Renardodden). Both authors wrote the paper.

Competing interests. The authors declare that they have no conflict of interest.

Acknowledgements. This research was conducted on samples provided by the International Ocean Discovery Program (IODP), which includes its predecessors, the International Ocean Drilling Program (IODP), Ocean Drilling Program (ODP), and Deep Sea Drilling Project (DSDP). Kasia K. Śliwińska is most grateful to the late Svein B. Manum for inspiring discussions and sharing his view on the morphology of Svalbardella cooksoniae, Franz-Josef Linde-mann (Natural History Museum in Oslo) for his assistance during her visit to the Museum, and Sarah Damassa for sharing her slides from DSDP Site 112 and a slide from DSDP 338. Annette Ryge and Dorthe Samuelsen (GEUS) are kindly acknowledged for preparing palynological slides from Sites U1406 and U1411. Robert Williams (NPD) took a photo shown on Plate 4A. We are grateful to the journal reviewers, Laurent Londeix and Robert A. Fensome. Further, we thank Robert A. Fensome for providing original photographs of Pulaeocystodinium obesum which are illustrated here (Plate 5A–C). The COVID-19 pandemic prevented us from personally re-examining type specimens.

Financial support. This research has been supported by the Danish Council for Independent Research (grant no. 11-107497 to Kasia K. Śliwińska) and the Natural Sciences and Engineering Research Council of Canada (to Martin J. Head).

Review statement. This paper was edited by Luke Mander and reviewed by Laurent Londeix and Robert A. Fensome.

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