

The Arctic is a critical region in the current and past global climate system. One major challenge for investigating the geological history of the Arctic is the establishment of a rigorous chronology. To get better insights into the late Neogene history of oceanographic and climatic change in the region, well-calibrated ocean drill cores are essential. Nevertheless, detailed stratigraphy from (sub)Arctic ocean drill sites remains problematic due to the generally poor preservation of calcareous microfossils in high northern-latitude oceans. Here, organic-walled palynomorphs (dinoflagellate cysts and acritarchs) can compensate for the lack of calcareous microfossils because diverse and rich palynological assemblages can be recovered. We defined three magnetostratigraphically-calibrated dinoflagellate cyst and acritarch biozones in the Upper Miocene to Upper Pliocene of Norwegian Sea Ocean Drilling Program Hole 642B (Vøring Plateau) that show potential for correlation within the Nordic Seas. It is also noted that several bioevents in the Nordic Seas are strongly diachronous with the North Atlantic, highlighting the limitations of applying North Atlantic bioevents directly to the Nordic Seas. For each of the three interval biozones (VP1 to VP3) we use the highest occurrences of acritarch and dinoflagellate cyst species ("*Veriplicidium franklinii*" of Anstey 1992, *Reticulosphaera actinocoronata*, and *Invertocysta lacrymosa*) that are relatively synchronous across the Nordic Seas and North Atlantic and thus show potential for a North Atlantic–Arctic reference stratigraphy.

## **Highlights**

A new late Neogene marine palynomorph biozonation is presented for the Norwegian Sea

Three biozones covering the uppermost Miocene to Upper Pliocene are defined

The zonation is an important step towards a regional late Neogene Arctic stratigraphy

Correlation between the Nordic Seas and North Atlantic is strongly diachronous

1Late Neogene dinoflagellate cyst and acritarch biostratigraphy for Ocean

2Drilling Program Hole 642B, Norwegian Sea

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## 22Abstract

23The Arctic is a critical region in the current and past global climate system. One  
24major challenge for investigating the geological history of the Arctic is the  
25establishment of a rigorous chronology. To get better insights into the late  
26Neogene history of oceanographic and climatic change in the region, well-  
27calibrated ocean drill cores are essential. Nevertheless, detailed stratigraphy  
28from (sub)Arctic ocean drill sites remains problematic due to the generally poor  
29preservation of calcareous microfossils in high northern-latitude oceans. Here,  
30organic-walled palynomorphs (dinoflagellate cysts and acritarchs) can  
31compensate for the lack of calcareous microfossils because diverse and rich  
32palynological assemblages can be recovered. We defined three  
33magnetostratigraphically-calibrated dinoflagellate cyst and acritarch biozones in  
34the Upper Miocene to Upper Pliocene of Norwegian Sea Ocean Drilling Program  
35Hole 642B (Vøring Plateau) that show potential for correlation within the Nordic  
36Seas. It is also noted that several bioevents in the Nordic Seas are strongly  
37diachronous with the North Atlantic, highlighting the limitations of applying  
38North Atlantic bioevents directly to the Nordic Seas. For each of the three interval  
39biozones (VP1 to VP3) we use the highest occurrences of acritarch and  
40dinoflagellate cyst species ("*Veriplicidium franklinii*" of Anstey 1992,  
41*Reticulatosphaera actinocoronata*, and *Invertocysta lacrymosa*) that are relatively  
42synchronous across the Nordic Seas and North Atlantic and thus show potential  
43for a North Atlantic–Arctic reference stratigraphy.

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## 46Keywords

47Miocene | Pliocene | palynology | biozones | Nordic Seas | North Atlantic

## 481 Introduction

49The Norwegian Sea links the high-latitude eastern North Atlantic with the Arctic  
50Ocean and is today a critical component of the high-latitude ocean circulation and  
51Arctic climate because of northward heat transport via the Norwegian Atlantic  
52Current. Circulation in the Nordic Seas was different from the present during the  
53earliest Pliocene (Jansen et al., 2000), with a modern-like circulation developing  
54around 4.5 Ma in the Early Pliocene (Knies et al., 2014; De Schepper et al., 2015).  
55In order to gain further understanding of the timing of oceanographic and  
56climatic changes in the Arctic Ocean and Nordic Seas, detailed studies of  
57chronostratigraphically well-calibrated successions are essential (e.g.  
58Mattingsdal et al., 2013). The major challenge for a detailed stratigraphy at  
59(sub)Arctic ocean drill sites remains the poor preservation of calcareous  
60microfossils, or also simply their absence due to the harsh environmental  
61conditions. Calcareous microfossils have not been recorded from the only  
62existing Neogene Arctic Ocean sediment record (ACEX), and establishment of a  
63reference stratigraphy for that region is further hampered by incomplete core  
64recovery and the generally rare presence of organic-walled microfossils (e.g.  
65Moran et al., 2006; Matthiessen et al., 2009a; 2009b). However, recently, at the  
66margin of the Arctic Ocean, on the Yermak Plateau in the Fram Strait, an  
67uppermost Miocene to Quaternary stratigraphic framework for the Atlantic–  
68Arctic gateway was successfully established based on seismic correlation,  
69magnetostratigraphy and biostratigraphy at Ocean Drilling Program (ODP) Sites  
70910, 911 and 912 (Mattingsdal et al., 2013). Based mainly on palynology and  
71magnetostratigraphy, the base of ODP Hole 911A was placed in the lowermost

72Pliocene, with an age likely around 5.2 Ma (Mattingsdal et al., 2013; Grøsfjeld et  
73al., 2014). Schreck et al. (2012) calibrated Neogene dinoflagellate cyst and  
74acritarch bioevents to magnetostratigraphy at Iceland Sea ODP Hole 907A. These  
75bioevents were successfully applied to dating a Central Arctic Late Miocene  
76interval (Stein et al., 2016), highlighting the potential of palynology in  
77contributing to dating Arctic and Nordic Seas sediments and establishing an  
78Arctic reference stratigraphy. In the Norwegian Sea, a diverse late Neogene  
79dinoflagellate cyst flora has already been documented by Mudie (1989), but a  
80detailed and up-to-date record of dinoflagellate cyst stratigraphic events is  
81currently required.

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83In the Nordic Seas, organic-walled palynomorphs (dinoflagellate cysts and  
84acritarchs) can compensate for the absence of calcareous microfossils since their  
85assemblages are known to be diverse, rich and well preserved. Furthermore, our  
86knowledge of late Neogene dinoflagellate cysts and acritarchs has increased  
87steadily since the 1990s following major taxonomic advances and the formal  
88description of numerous new dinoflagellate cyst and acritarch taxa (e.g. Head,  
891993, 1996, 1997; Head and Norris, 2003; De Schepper et al., 2004; De Schepper  
90and Head, 2008a, 2014; Schreck et al., 2012; Verhoeven et al., 2014; Versteegh  
91and Zevenboom, 1995). Dinoflagellate cysts have been successfully applied for  
92establishing a detailed late Neogene stratigraphy in the eastern North Atlantic  
93(De Schepper and Head, 2008a, 2009), Iceland Sea (Verhoeven et al., 2011;  
94Schreck et al., 2012) and the North Sea Basin (Louwye et al., 2004; De Schepper  
95et al., 2009; Dybkjær and Piasecki, 2010), suggesting that this should also be  
96achievable in the Norwegian Sea.

97

98Because of the oceanographic and climatic importance of the Norwegian Sea in  
99influencing Arctic climate, we have reinvestigated the palynology of ODP Site 642  
100with the help of advances in dinoflagellate cyst and acritarch taxonomy. ODP Site  
101642 (Figure 1) located on the Vøring Plateau is currently influenced by Atlantic  
102waters via the Norwegian Atlantic Current, yet is situated at a latitude  
103comparable to that of the Iceland Sea ODP Site 907, where Schreck et al. (2012)  
104documented Miocene to Pliocene dinoflagellate cyst assemblages. The aim of this  
105study is to establish a new, detailed palynostratigraphy for the Upper Miocene to  
106Pliocene succession in the Norwegian Sea, which may represent a benchmark  
107biozonation that can interlink the North Atlantic, Norwegian Sea, and Arctic  
108Ocean.

## 1092 **Materials and methods**

### 1102.1 ***Ocean Drilling Program Hole 642B: lithology and samples***

111ODP Hole 642B (67°13.5'N, 2°55.7'E; water depth 1268 m, core recovery of  
11297.5%) located on the outer Vøring Plateau, was drilled with an advanced  
113hydraulic piston corer in 1985 as a part of the ODP Leg 104 (Shipboard Scientific  
114Party, 1987) (Figure 1). The drill hole reached a depth of 221.1 meters below sea  
115floor (mbsf), with a total core recovery of 215.6 m, and penetrated  
116predominantly pelagic to hemipelagic sediments of Holocene through Early  
117Miocene age.

118

119The 40 samples investigated in this study were collected from the lower part of  
120lithological Unit I, Subunit IIA and the upper part of IIB (Figure 2). Unit I (0–65.7



121mbsf) consists of repeated alternations of dark, carbonate-poor glacial mud and  
122light, carbonate-rich interglacial sandy mud. Bioturbation and the presence of  
123dropstones are common throughout. Subunit IIA (65.7–90.4 mbsf) consists  
124predominantly of nannofossil oozes, with minor diatom-nannofossil oozes and  
125muds. This subunit is moderately to heavily bioturbated. Subunit IIB (90.4–107.2  
126mbsf) consists of siliceous oozes and siliceous muds, one short interval of  
127nannofossil ooze and several minor volcanic ash layers, which have been  
128moderately to heavily bioturbated.

### 1292.2 *Palynological preparation*

130Samples were first wet sieved at 150 and 63  $\mu\text{m}$  for measurements of  
131foraminiferal stable isotopes (Risebrobakken et al., 2015). Consequently, the <63  
132 $\mu\text{m}$  sediment fraction was dried and processed to extract organic-walled  
133palynomorphs at Palynological Laboratory Services Ltd (Holyhead, UK) following  
134a slightly modified procedure described in De Schepper and Head (2008b). A  
135weighed quantity of each sample (Table 1) was disintegrated and placed into a 1L  
136Tripour beaker. One *Lycopodium clavatum* spore tablet (Batch no. 483.216, n =  
13718,583  $\pm$  1,708 spores per tablet) was added prior to chemical degradation.  
138Calcium carbonate was removed by slowly adding 50% hydrochloric acid (HCl)  
139until the sample had stopped reacting. Subsequently, the beaker was topped up  
140fully with water, stirred and allowed to settle. The diluted supernatant liquid was  
141then sieved through a 10- $\mu\text{m}$  sieve cloth, and the collected residue returned to  
142the beaker. To remove silicate, 100 ml of 60% hydrofluoric acid (HF) was added  
143and left for two days with periodic stirring. The sample was then topped up with  
144water for dilution and the entire residue sieved through a 10- $\mu\text{m}$  sieve cloth and

145collected, before it was placed in a 250 ml Pyrex glass. Oxidation was carried out  
146on four samples (642B-9H5, 100–101 cm, 642B-10H2, 40–41 cm, 642B-10H2,  
147100–101 cm, 642B-10H3, 102–103 cm) before mounting by adding 50% cold  
148nitric acid (HNO<sub>3</sub>) to the residue in the beaker (in 25 ml of water) and left for  
149some time, depending on the preservation of the residue present. A short  
150ultrasonic treatment then preceded the sieving as before using water. Before  
151mounting, the residue was mixed with a 1% solution of polyvinyl alcohol (PVA) to  
152prevent clumping, and stained with Safranin-O if necessary. It was then pipetted  
153onto a 32x22 mm cover slip on a low temperature drying plate and allowed to  
154dry. Once dry, the coverslip was mounted onto the glass microscope slide using  
155glycerine jelly optical adhesive.

### 156**2.3 Counting, photography and data storage**

157The focus of this study was the dinoflagellate cyst assemblage, but also acritarchs  
158and terrestrial palynomorphs were identified. At least 250 dinoflagellate cysts  
159were counted in each slide along non-overlapping traverses at 400x  
160magnification using a transmitted light microscope (Zeiss Axiophot and  
161AxioImager.A2). During this regular count, all encountered acritarchs, spores,  
162pollen and fresh water algae were also enumerated. The remainder of the slide  
163was scanned at 200x magnification to identify rare taxa not seen during the  
164regular count. Broken palynomorphs were counted as one unit when more than  
165half of the original form was present.

166Photographs of selected dinoflagellate cysts and acritarchs were taken using a  
167Zeiss Axiocam 506 Color on a Zeiss AxioImager.A2 microscope.

168All raw data are available at <http://doi.pangaea.de/10.1594/PANGAEA.846838>.

#### 1692.4 **Taxonomy and nomenclature**

170Some taxa were grouped due to problematic taxonomy and/or limited  
171stratigraphic value. This includes most *Spiniferites* and *Achomosphaera* species  
172which were lumped as *Spiniferites/Achomosphaera* spp. indet. According to  
173Schreck and Matthiessen (2013), the *Batiacasphaera micropapillata* complex  
174comprises both micropapillate and microreticulate forms. Although the  
175distinction is not always easy to make (depending on the quality of the  
176microscope objective), we have classified the purely microreticulate forms as  
177*Batiacasphaera minuta* s.s. *Pyxidinopsis* sp. A has a wall ornament that resembles  
178closely that of *Batiacasphaera sphaerica*, but has a precingular instead of  
179antapical archeopyle. Proximate specimens with low wall ornament comparable  
180to *Batiacasphaera sphaerica*, *Batiacasphaera micropapillata* and *Pyxidinopsis* sp.  
181A where the archeopyle could not unquestionably be determined were assigned  
182to the *Batiacasphaera/Pyxidinopsis* spp. indet. group. Reworked dinoflagellate  
183cysts were identified on preservation state and known stratigraphic range  
184outside the uppermost Miocene to Pleistocene.

185Nomenclature generally follows De Schepper et al. (2004), De Schepper and Head  
186(2008b, 2014) and Fensome et al. (2008).

#### 1872.5 **Age model, errors and sample resolution**

188The paleomagnetostratigraphic study of Bleil (1989) reports a complete and  
189apparently continuous sequence of the Bruhnes and Matuyama Chrons, including  
190all major reversals as well as the Gauss/Matuyama boundary. A minor hiatus  
191occurs in the uppermost Gauss Chron (near the 8H and 9H cores boundary), but  
192the Mammoth subchron was positively identified. Also the Gilbert Chron and its

193four normal polarity subchrons were identified. The palaeomagnetic reversal  
194ages for ODP Hole 642B reported in Bleil (1989) were recalibrated to the  
195Astronomically Tuned Neogene Time Scale 2012 (ATNTS 2012; Hilgen et al.,  
1962012) (Table 2; see also De Schepper et al., 2015). By linearly correlating the  
197mid-points between two adjacent samples that record a reversal in inclination,  
198we provide an age estimate for each investigated sample and bioevent (Table 3).  
199The error on the palaeomagnetic reversals can amount to half the age difference  
200between the two samples either side of the observed reversal (Weaver and  
201Clement, 1987). The distance between two samples across a magnetic reversal  
202varies from 3 to 41 cm (average 29 cm) in the Pliocene succession of ODP 642,  
203which corresponds to an error of 1.5–21 kyr (average 15 kyr) for the age of the  
204magnetic boundaries.

205

206An accurate age for the samples in core-section 642B-8H6 has not been possible  
207to provide because they lie between a hiatus around 66 mbsf (Bleil, 1989) and  
208the Gauss/Matuyama polarity reversal (2.588 Ma). Sample 642B-8H6, 129–130  
209cm (65.69 mbsf) is younger than 3.20 Ma, and is certainly older than 2.58 Ma but  
210it may also be older than 3.04 Ma.

211

212Sample spacing varies between 5 and 195 cm, with an average of 84 cm, which  
213corresponds to age intervals of 7 to 257 kyr, with an average of 75 kyr. The  
214sampling resolution has an impact on the accuracy of the estimated ages of a  
215highest occurrence (HO), because the true HO of a species is between the highest  
216sample containing this species and the overlying sample without this species. For  
217ODP Hole 642B, this results in an age for the true HO that can be up to  $7(75)257$

218kyr younger than the reported age, or older in the case of a lowest occurrence  
219(LO).

## 2203 Results

### 2213.1 General

222Most samples contain mainly well-preserved palynomorph assemblages.  
223Exceptions are samples 642B-10H2, 40–41 cm (77.80 mbsf) and 642B-10H3,  
224102–103 cm (79.92 mbsf) where counting was hampered by a large amount of  
225amorphous material, and sample 642B-11H6, 110–111 cm (94.00 mbsf) where  
226preservation was poor and cysts showed some degradation. Dinoflagellate cysts  
227and acritarchs dominate the assemblages, but terrestrial palynomorphs (spores  
228and pollen) also occur abundantly (Table 1). The fresh water alga *Gelasinicysta*  
229*vangeelii* is present in low abundance in most samples.

230The focus of this study was marine palynomorphs, especially the dinoflagellate  
231cyst assemblage. The concentration of dinoflagellate cysts is between  $376 \pm 44$   
232and  $79,854 \pm 27,861$  dinoflagellate cysts/g dry sediment. A total of 95  
233dinoflagellate cyst taxa were recorded in the studied interval and the number of  
234dinoflagellate cyst taxa per sample is high at 18(35)53 taxa per sample. Diversity  
235(Shannon-Wiener index, Table 1) is highly variable but reveals a gradual  
236declining trend towards younger samples. Evenness remained relatively constant  
237throughout the studied interval and fluctuates between 0.55–0.78, indicating that  
238there is always one taxon (or a few taxa) dominating the samples. Dominant taxa  
239in the studied interval include the *Batiacasphaera micropapillata* complex,  
240*Habibacysta tectata*, *Nematosphaeropsis labyrinthus*, *Operculodinium? eirikianum*  
241var. *eirikianum*, cysts of *Protoceratium reticulatum*, round brown cysts and

242 *Spiniferites/Achomosphaera* spp. The overall assemblage throughout the studied  
243 interval contains typical Upper Miocene to Pliocene taxa such as *Achomosphaera*  
244 *andalousiensis* subsp. *andalousiensis*, *Amiculosphaera umbraculum*,  
245 *Bitectatodinium raedwaldii*, *Invertocysta lacrymosa*, *Operculodinium tegillatum*  
246 and *Pyxidinopsis braboi*, as well as extant taxa (e.g. *Impagidinium aculeatum*,  
247 *Lingulodinium machaerophorum*).

248

249 Acritarchs constitute a significant part of the marine palynological assemblages,  
250 and often outnumber dinoflagellate cysts. The concentration of acritarchs in the  
251 studied interval varies between  $53 \pm 21$  and  $114,498 \pm 39,840$  acritarchs/g. A  
252 total of 18 acritarch taxa was recorded and the number of acritarch taxa per  
253 sample is 1(6)10. The taxa *Cymatiosphaera? invaginata*, *Nannobarbophora*  
254 *walldalei* and small spiny acritarchs dominate the acritarch assemblages, and in  
255 the middle Zanclean (after ~4.5 Ma) also *Cymatiosphaera? icenorum* and  
256 *Lavradosphaera crista* are abundant.

### 257 **3.2 Age calibration of selected bioevents**

258 Since a solid palaeomagnetostratigraphy is available for ODP Site 642 (Bleil,  
259 1989), it is possible to provide calibrated ages for the bioevents of individual  
260 dinoflagellate cyst and acritarch species following the procedures outlined in De  
261 Schepper and Head (2008a) and/or Schreck et al. (2012). Table 3 summarises  
262 the calibrated ages for lowest, highest and highest common occurrences in ODP  
263 Site 642.

### 264**3.3 Biozonation**

265We have attempted to construct a Nordic Seas biozonation that can serve as a  
266basis for correlation with the North Atlantic and Arctic oceans. Therefore, the  
267zone boundaries have been based on bioevents of species that are relatively  
268synchronous across these regions. Three zones are defined as interval biozones,  
269the body of fossiliferous strata between two specified biohorizons (Figure 2). The  
270several bioevents occurring within these zones can be used for more local-to-  
271regional correlations in the Nordic Seas. Range charts for dinoflagellate cysts and  
272acritarchs are provided in an Appendix.

273

274The biozonation follows the *International Stratigraphic Guide* (abridged version  
275of Murphy and Salvador, 1999). The biozones are given informal names (VP1 to  
276VP3) referring to the location of ODP Site 642 on the Vøring Plateau. The  
277biohorizons used are the HO or LO of dinoflagellate cyst and acritarch taxa. Also,  
278the concepts of highest common occurrence (HCO) and highest persistent  
279occurrence (HPO) (De Schepper and Head, 2008a) are used to describe events  
280within biozones. A HCO is recorded in a sample that marks the highest sample  
281where a particular taxon is noticeably common, but this taxon occurs in lower  
282abundance above the level. A HPO marks the highest sample below which a taxon  
283is persistently recorded (i.e. in successive samples), even if the records consist of  
284few specimens only. The classification of relative abundance of the dinoflagellate  
285cysts and the acritarchs is as follows: *Rare*, 0–2.9 %, *Frequent*, 3–9.9 %, *Common*,  
28610–29.9%, *Abundant*, 30–49.9 %, and *Dominant*  $\geq$  50 %.

287

#### 288**VP1 Interval biozone**

289**Definition.** The body of strata from the base of the studied interval to the HO of  
290"*Veriplicidium franklinii*" of Anstey (1992).

291**Characteristic events.** "*Veriplicidium franklinii*" of Anstey (1992) has a HO in  
292sample 642B-10H7, 30–31 cm (85.20 mbsf) at the top of the zone (Figure 2,  
293Appendix). Additional events characterizing the top of the zone are the HO  
294"*Impagidinium densiverrucosum*" of Zevenboom and Santarelli in Zevenboom  
295(1995), the HPO of *Trinovantedinium glorianum*, and the LOs of *Operculodinium*  
296*tegitatum* and *Heteraulacacysta* sp. A of Costa and Downie (1979). *Corrudinium*  
297*devernaliae* has a LO in sample 642B-11H3, 65–66 cm (89.05 mbsf) in the middle  
298of the zone.

299**Dinoflagellate cyst association.** *Nematosphaeropsis labyrinthus*, *Operculodinium*  
300*centrocarpum* s.s., *Operculodinium?* *eirikianum* var. *eirikianum*, round brown  
301cysts, *Reticulosphaera actinocoronata*, *Selenopemphix* spp. and  
302*Spiniferites/Achomosphaera* spp. indet. are the most abundant taxa. Several  
303heterotrophic taxa (*Barssidinium*, *Lejeunecysta*, *Selenopemphix*) are recorded in  
304every sample in variable abundance (rare to common). The species *Lejeunecysta*  
305*hatterasensis* is recorded in several samples. *Cerebrocysta poulsenii* and  
306*Selenopemphix armageddonensis* are restricted to this zone only.

307**Acritarch association.** "*Veriplicidium franklinii*" of Anstey (1992) occurs in all  
308samples and is rare to common. Small spiny acritarchs and *Cymatiosphaera?*  
309*invaginata* dominate the acritarch assemblage. *Lavradosphaera crista* is rare to  
310common. *Lavradosphaera canalis*, *Lavradosphaera lucifer*, *Cymatiosphaera?*  
311*aegirii*, *Cymatiosphaera?* *icenorum* and *Nannobarbophora walldalei* are rare and  
312sporadic within the zone.



313**Reference section.** Samples 642B-11H7, 55–56 cm, to 642B-10H7, 30–31 cm  
314(94.95–85.20 mbsf).

315**Age.** Late Messinian to earliest Zanclean, from >5.89 Ma to 5.29 Ma.

316**Calibration.** In ODP Hole 642B, Zone VP1 corresponds to the middle part of the  
317planktonic foraminifer *Neogloboquadrina atlantica* sinistral Zone (Spiegler and  
318Jansen, 1989), the lower radiolarian *Antarctissa whitei* Zone (Goll and Bjørklund,  
3191989), the upper NN15/NN17 calcareous nannofossil Zone (Donnally, 1989), and  
320the upper PM3 dinoflagellate cyst Zone (Mudie, 1989) (Figure 3).

321**Correlation.** The VP1 Zone (>5.89 Ma to 5.29 Ma) partly overlaps with the  
322*Selenopemphix armageddonensis* Zone (7.6–5.0 Ma) of Dybkjær and Piasecki  
323(2010) (Figure 4), which is defined from the HO of *Hystrichosphaeropsis obscura*  
324to the HO of *Barssidinium evangelinae*. The range of *Selenopemphix*  
325*armageddonensis* is presumed to correspond approximately to the entire  
326*Selenopemphix armageddonensis* Zone (Dybkjær and Piasecki, 2010).  
327*Hystrichosphaeropsis obscura* and *Barssidinium evangelinae* were not recorded  
328in ODP Hole 642B, but *Selenopemphix armageddonensis* was last recorded in the  
329upper part of the VP1 Zone (sample 642B-11H2, 15–16 cm) (Figure 2).

330**Comments.** Anstey (1992) describes the acritarch “*Veriplicidium franklinii*” from  
331the Upper Miocene through possibly Lower Pliocene of ODP Site 645 (Baffin Bay).  
332Although only known under an informal name, this acritarch appears to have  
333correlation potential across the North Atlantic.

334“*Impagidinium densiverrucosum*” of Zevenboom and Santarelli in Zevenboom  
335(1995) is mainly known from the Middle Miocene of The Netherlands and Italy  
336(Zevenboom, 1995) and the Upper Miocene of the Danish North Sea (Dybkjær  
337and Piasecki, 2010) In our study, it has a HO at the top of the VP1 Zone (sample

338642B-10H7, 30–31 cm). It is morphologically very distinct, although often  
339broken, and is usually present in low numbers only. Therefore, the HO of the  
340more abundant "*Veriplicidium franklinii*" of Anstey (1992) was chosen as the  
341defining event for the VP1 Zone upper boundary (Figure 2). *Selenopemphix*  
342*armageddonensis* occurs sporadically in the uppermost Miocene and lowermost  
343Pliocene of the Danish North Sea, and ODP Hole 642B, and was recorded also in  
344the Lower Pliocene of Belgium (Louwye et al., 2004). The LO of *Operculodinium*  
345*tegillatum* at the top of the zone (Table 2) could be a useful local marker, although  
346it has been reported from the Upper Miocene of the Iceland Sea (Schreck et al.,  
3472012) and the North Sea Basin (Louwye, 1999; Louwye and De Schepper, 2010).  
348Dinoflagellate cysts dominate the marine palynomorph assemblage with an  
349average concentration of  $8,202 \pm 1,202$  cysts/g compared to  $5,648 \pm 867$   
350acritarchs/g.

351

### 352VP2 Interval biozone

353**Definition.** The body of strata from the HO of "*Veriplicidium franklinii*" of Anstey  
354(1992) to the HO of *Reticulosphaera actinocoronata*.

355**Characteristic events.** The top of the zone is defined by the HO of  
356*Reticulosphaera actinocoronata* in sample 642B-10H2, 40–41 cm. Also the  
357*Batiacasphaera micropapillata* complex, *Selenopemphix brevispinosa* and Cyst  
358type I of de Vernal and Mudie (1989) have their HO in the same sample (Figure 2,  
359Appendix). The range of *Operculodinium tegillatum* corresponds almost to the  
360entire zone: its LO is immediately at the boundary between the VP1 and VP2  
361Zones, whereas its HO is found in the sample immediately above the zone. The

362LOs of *Filisphaera filifera* subsp. *filifera* and *Bitectatodinium raedwaldii* are at the  
363base of the zone in sample 642B-10H6, 65–66 cm (Table 3).

364**Dinoflagellate cyst association.** *Nematosphaeropsis labyrinthus*, *Operculodinium*  
365*tegillatum*, round brown cysts and *Spiniferites/Achomosphaera* spp. are the most  
366abundant taxa. The *Batiacasphaera micropapillata* complex is frequent to  
367abundant. *Operculodinium centrocarpum* s.s. and *Operculodinium?* *eirikianum* var.  
368*eirikianum* are rare to common. *Filisphaera filifera* subsp. *filifera* is frequent in  
369one sample. *Corrudinium?* *labradori* and *Reticulosphaera actinocoronata* are  
370rare to frequent, and occur in every sample. Cyst type I of de Vernal and Mudie  
371(1989) has an acme in the uppermost part of the zone. Cysts of *Protoceratium*  
372*reticulatum* (= *Operculodinium centrocarpum* sensu Wall and Dale, 1966) are rare  
373and frequent in the upper two samples of the zone.

374**Acritarch association.** The zone corresponds to an acme of *Cymatiosphaera?*  
375*invaginata* which dominates the acritarch assemblage. Small spiny acritarchs are  
376common. *Cymatiosphaera?* *icenorum*, *Lavradosphaera crista* and  
377*Nannobarbophora walldalei* are rare to frequent throughout the zone.  
378*Lavradosphaera lucifer* has a HO in sample 642B-10H2, 100–101 cm (78.40  
379mbsf).

380**Reference section.** Samples 642B-10H6, 65–66 cm, to 642B-10H2, 40–41 cm  
381(84.05–77.80 mbsf).

382**Age.** Early Zanclean, 5.18–4.64 Ma.

383**Calibration.** In ODP Hole 642B, Zone VP2 corresponds to the middle to upper  
384part of the planktonic foraminifer *Neogloboquadrina atlantica* sinistral Zone  
385(Spiegler and Jansen, 1989), the upper *Antarctissa whitei* radiolarian Zone (Goll  
386and Bjørklund, 1989), and the uppermost NN15/NN17 calcareous nannofossil

387zone (Donnally, 1989) (Figure 3). The upper boundary of the VP2 Zone coincides  
388with the upper boundary of the *Antarctissa whitei* radiolarian Zone and  
389NN15/NN17 calcareous nannofossil Zone. This zone has no equivalent in the  
390dinoflagellate cyst zonation, as this interval was left unstudied by Mudie (1989).

391**Correlation.** In the Nordic Seas, the *Batiacasphaera micropapillata* complex  
392(including *Batiacasphaera minuta*) has a HO in the sample below the HOs of  
393*Corrudinium devernaliae* and *Operculodinium tegillatum*. In the North Atlantic,  
394the RT1, RT2 and RT3 zones (~4.0–3.71 Ma) are defined respectively by the HOs  
395of *Corrudinium devernaliae* (top RT1), *Batiacasphaera minuta* (top RT2) and  
396*Operculodinium tegillatum* (top RT3). The sequential order of the HOs is different  
397between both regions making a detailed correlation between the North Atlantic  
398and Nordic Seas biozones difficult. Based on the occurrence of the same marker  
399dinoflagellate cyst species, the RT1, RT2 and RT3 biozones (~4.0–3.71 Ma) can  
400be correlated to the upper VP2 Zone (5.18–4.64 Ma), but the correlation is  
401strongly asynchronous (Figure 4). The upper VP2 Zone correlates to the lower  
402*Melitasphaeridium choanophorum* Zone (5.0–3.60 Ma) of Dybkjær and Piasecki  
403(2010).

404**Comments.** Our study presents a refinement of the stratigraphic ranges of  
405*Reticulosphaera actinocoronata* and *Filisphaera filifera* subsp. *filifera* at ODP  
406Hole 642B. Mudie (1989) recorded the HO of *Reticulosphaera actinocoronata* in  
407Hole 642B, sample 642B-14CC, 20 cm (120.10 mbsf), but we document the HO of  
408*Reticulosphaera actinocoronata* in sample 642B-10H2, 40–41 cm (77.80 mbsf,  
4094.64 Ma), with five higher occurrences being interpreted as reworked. We  
410consider the specimens of *Reticulosphaera actinocoronata* recorded above  
41177.80 mbsf to be reworked based on the observations that (1) these are either

412single occurrences or occurrences recorded outside the regular count, (2) in the  
413same samples single occurrences of other Lower Pliocene species are recorded  
414(e.g. *Batiacasphaera micropapillata* complex, *Operculodinium tegillatum* in  
415samples at 68.3, 70.45 mbsf), or (3) they co-occur with a large amount of  
416undifferentiated reworked specimens (n=134 reworked cysts in sample 65.34  
417mbsf). Mudie (1989) also reported *Reticulosphaera actinocoronata* from Hole  
418642C, where it has a HPO in sample 642C-11H3, 56–57 cm (76.56 mbsf) at ~4.4  
419Ma (based on linear interpolation between the palaeomagnetic reversal for the  
420hole by Bleil (1989), updated to ATNTS2012 of Hilgen et al., 2012). Our new  
421observations place the range top of this species in Hole 642B closely to its range  
422top in Hole 642C.

423Mudie (1989) placed the LO of *Filisphaera filifera* in sample ODP 642B-9CC, 18  
424cm (76.25 mbsf) considerably higher than our observation of its LO in sample  
425642B-10H6, 65–66 cm (84.05 mbsf). We even recorded a single specimen of  
426*Filisphaera filifera* subsp. *pilosa* in the underlying VP1 Zone in sample 642B-  
42711H3, 65–66 cm (89.05 mbsf). The difference between our observations and  
428those of Mudie (1989) can be explained by the lower sampling resolution of 1  
429sample per 9 m in this part of the section (see discussion).

430Cyst type I of de Vernal and Mudie (1989) occurs between 4.81 to 4.64 Ma (Table  
4313), with a clear acme between 4.69 and 4.64 Ma. This record corresponds to  
432other Zanclean records in the Labrador Sea (ODP Hole 646B), western North  
433Atlantic (DSDP Hole 603C), and off West Greenland (M.J. Head, pers. comm. in  
434Louwye et al., 2004; Piasecki, 2003). In DSDP Hole 603C, this species has a  
435restricted range between 4.66 and 4.21 Ma, with an acme between 4.54 and 4.32  
436Ma (M.J. Head, unpublished data).

437Average dinoflagellate cysts concentrations ( $34,331 \pm 9392$  cysts/g) are  
438considerably higher than the VP1 Zone, but acritarchs are the dominant marine  
439palynomorph group with an average concentration of  $40,698 \pm 11,050$   
440acritarchs/g.

441

#### 442**VP3 INTERVAL BIOZONE**

443**Definition.** The body of strata from the HO of *Reticulosphaera actinocoronata* to  
444the HO of *Invertocysta lacrymosa*.

445**Characteristic events.** At the base of the zone, *Corrudinium devernaliae* and  
446*Operculodinium tegillatum* have their HOs in sample 642B-10H1, 145–146 cm  
447(77.35 mbsf). The ranges of *Ataxiodinium confusum*, *Operculodinium? eirikianum*  
448var. *crebrum*, *Impagidinium solidum*, *Melitasphaeridium* sp. A of De Schepper and  
449Head (2008b) and *Spiniferites elongatus* are restricted to this zone (Figure 2,  
450Appendix). At or just below the top of this zone, the HOs of *Bitectatodinium?*  
451*serratum*, *Melitasphaeridium choanophorum*, *Ataxiodinium zevenboomii*,  
452*Corrudinium? labradori*, *Heteraulacacysta* sp. A of Costa and Downie (1979),  
453*Operculodinium janduchenei*, *Trinovantedinium glorianum* and *Achomosphaera*  
454*andalousiensis suttonensis* are recorded.

455**Dinoflagellate cyst association.** *Filisphaera filifera*, *Nematosphaeropsis*  
456*labyrinthus*, cysts of *Protoceratium reticulatum*, and *Spiniferites/Achomosphaera*  
457spp. are the most abundant taxa. The relative abundance of cysts of  
458*Protoceratium reticulatum* increases sharply across the lower boundary of the  
459zone to reach at least 25% in most samples of this biozone, except the uppermost  
460sample which is dominated by *Habibacysta tectata* and *Filisphaera filifera* subsp.  
461*filifera*. *Operculodinium? eirikianum* var. *eirikianum* is less abundant than in Zone

462VP2. *Corrudinium? labradori* (rare), *Impagidinium pallidum* (rare to frequent),  
463*Impagidinium aculeatum* (rare) and *Impagidinium paradoxum* (rare) are  
464consistently present.

465**Acritarch association.** Acritarchs are very abundant in this zone and outnumber  
466the dinoflagellate cysts in the lower and middle part of the zone, except in the  
467lowermost sample which yielded very few acritarchs (Table 1). The assemblages  
468are dominated by small spiny acritarchs. *Lavradosphaera crista* is generally  
469common to frequent, and shows a decline in the upper part of the zone when  
470dinoflagellate cysts become more abundant. *Lavradosphaera crista* has a  
471HCO/HPO in sample 642B-9H1, 120–121 cm (67.60 mbsf). *Cymatiosphaera?*  
472*invaginata* is abundant throughout the zone. *Cymatiosphaera? icenorum* has a  
473well-expressed acme between samples 642B-10H1, 40–41 cm (76.30 mbsf) and  
474642B-9H2, 110–111 cm (69.00 mbsf).

475**Reference section.** Samples 642B-10H1, 145–146 cm, to 642B-9H1, 120–121 cm  
476(77.35–67.60 mbsf).

477**Age.** Zanclean to Piacenzian, between 4.49 to 3.27 Ma.

478**Calibration.** In ODP Hole 642B, Zone VP3 corresponds to the uppermost part of  
479the planktonic foraminifer *Neogloboquadrina atlantica* sinistral Zone (Spiegler  
480and Jansen, 1989), and the upper boundary of both zones seems to coincide  
481(Figure 3). Zone VP3 corresponds to the *Pseudodictyophimus gracilipes*  
482*tetracanthus* radiolarian Zone (Goll and Bjørklund, 1989), the lowermost  
483NN16/NN19 calcareous nannofossil Zone (Donnally, 1989), and the lowermost  
484PM2 dinoflagellate cyst Zone (Mudie, 1989).

485**Correlation.** Apparently diachronous (see section 4.3), the upper  
486*Melitasphaeridium choanophorum* Zone (5.0–3.60 Ma, Dybkjær and Piasecki,

4872010) corresponds to the VP3 Zone (Figure 4). The VP3 Zone further correlates  
488to the North Atlantic RT4 (3.71–3.15 Ma) and RT5 (3.15–2.74 Ma) Zones, but  
489reveals a strong diachroneity.

490**Comments.** Acritarchs are the dominant palynomorph group within the zone  
491with an average concentration of  $10,888 \pm 1,534$  acritarchs/g. Dinoflagellate  
492cysts occur with an average of  $6,100 \pm 919$  cysts/g.

493

#### 494**NOT ZONED**

495The four samples in the interval between 66.90 and 64.54 mbsf (642B-9H1, 50–  
49651 cm to 642B-8H6, 14–15 cm) are difficult to date and we have not established a  
497zone, due to a hiatus near the 8H and 9H cores boundary (~66 mbsf; Bleil, 1989).  
498In the second sample below the hiatus, several dinoflagellate cyst species as well  
499as the acritarch *Lavradosphaera crista* disappear from the record. These HOs are  
500part of the clear decrease in diversity that is visible in the upper part of the  
501studied interval (Table 1). Several of the species recorded in the not zoned  
502interval (e.g. *Barssidinium* spp., *Filisphaera filifera* subsp. *filifera*, *Habibacysta*  
503*tectata*, *Amiculosphaera umbraculum*) disappear from the North Atlantic record  
504during the Pleistocene (De Schepper and Head, 2009).

## 5054 Discussion

### 5064.1 Comparison with the ODP Hole 642B palynological study by Mudie (1989)

507Assisted by considerable advances in dinoflagellate cyst and acritarch taxonomy  
508since the mid-1990s, we have recorded 95 dinoflagellate cyst taxa and 18  
509acritarch taxa in the Pliocene of ODP Hole 642B compared to 52 dinoflagellate  
510cyst and 3 acritarch taxa in Mudie (1989). Only 19 taxa are in common with the



511study of Mudie (1989), however this number does not take into account the  
512several taxa left in open nomenclature by Mudie (1989).

513

514The sampling resolution of our study is considerably higher than that of Mudie  
515(1989), and was more focused on the Miocene–Pliocene. We analysed 40 samples  
516from the Pliocene–Miocene interval in ODP 642B from core-section 642B-8H6 to  
517642B-11H7 with a sample interval from 5 to 195 cm (~5,000 to ~200,000 years).  
518In comparison, Mudie (1989) investigated the entire Quaternary to Miocene  
519section with a 1 m sample resolution down to core 642B-8H6 and every core  
520catcher (~9 m) between 642B-9H to 642B-20H (i.e. the Pliocene–Miocene  
521interval).

522

523The three zones (VP1, VP2, VP3) defined in the present study compare to the PM  
524zones of Mudie (1989) as follows (Figure 3): Zone VP1 corresponds to the  
525uppermost part of Zone PM3 and our youngest Zone VP3 corresponds to the  
526lowermost part of Zone PM2. The top of zone PM3, thought to be roughly  
527corresponding to the Miocene–Pliocene boundary, was defined at the HCO of  
528*Amiculosphaera umbraculum* (Mudie, 1989). We identified, however, this species  
529throughout the studied interval, with its highest abundance near the top of the  
530studied interval. *Amiculosphaera umbraculum* is known also from the Pleistocene  
531eastern North Atlantic up to 1.44 Ma (De Schepper and Head, 2008a) and can  
532therefore not be used as a stratigraphic marker for the Miocene–Pliocene  
533transition. Nevertheless, by coincidence, the upper boundary of our VP1 Zone  
534corresponds to the upper boundary of the PM3 Zone.

535

536Mudie (1989) recorded the LO of *Tectatodinium pellitum* and *Filisphaera filifera*  
537(642B-9H-CC, 18 cm) at the base of zone PM2 although one occurrence of  
538*Filisphaera filifera* below in sample 642B-12-CC, 28 cm is shown (Fig. 5 in Mudie,  
5391989). It must be pointed out that the illustrated specimen of *Tectatodinium?* sp.  
540of Piasecki (1980) (pl. 5, fig. 2 in Mudie, 1989) is in fact *Tectatodinium pellitum*,  
541whereas the illustrated *Tectatodinium pellitum* (pl. 5, figs. 1, 5 in Mudie, 1989) is  
542likely a different taxon. This could be the result of mislabeling the plates or reflect  
543taxonomical issues. In the present study, *Tectatodinium pellitum* was recorded  
544throughout the study interval. We identified the LO of *Filisphaera filifera* subsp.  
545*filifera* in sample 642B-10H6, 65–66 cm, at the base of our VP2 Zone. Note that  
546our VP2 Zone was identified in an interval of the core that was left unstudied by  
547Mudie (1989) (Figure 3).

548

#### 549**4.2 Correlations with and implications for the (Pliocene) Danish North Sea**

##### 550 ***zonation of Dybkjær and Piasecki (2010)***

551In Denmark, a Neogene dinoflagellate cyst biozonation was established from a  
552combination of onshore and offshore well sections (Dybkjær and Piasecki, 2010).  
553Due to the absence of sections with reliable marine isotope stratigraphy or  
554magnetostratigraphy within the Danish North Sea, the ages assigned to the  
555different Danish biozones were obtained from other locations where calibrated  
556sections are available and/or from data compilations (Powell and Brinkhuis in  
557Lourens et al. 2005). The identification of index species in the Danish Neogene  
558North Sea sediments allowed the biozones to be correlated to the western North  
559Atlantic zonation of de Verteuil and Norris (1996). The Danish North Sea

560 *Selenopemphix armageddonensis* and *Melitasphaeridium choanophorum* biozones  
561 can also be recognised in the Norwegian Sea, where they broadly correspond to  
562 the VP1–VP2 and VP2–VP3 zones respectively (Figure 4).

563

564 The Upper Miocene to lowermost Pliocene *Selenopemphix armageddonensis* Zone  
565 (7.6–5.0 Ma) in the Danish North Sea is defined between the HO of  
566 *Hystriosphæropsis obscura* and the HO of *Barssidinium evangelinae* (Dybkjær  
567 and Piasecki, 2010). The range of *Selenopemphix armageddonensis* is presumed  
568 to approximate the range of the zone (Dybkjær and Piasecki, 2010). Specimens  
569 of *Selenopemphix armageddonensis* were identified in three samples of our VP1  
570 Zone, suggesting at least a partial overlap of the VP1 Zone with the  
571 *Selenopemphix armageddonensis* Zone. However, *Barssidinium evangelinae* was  
572 not identified in ODP Hole 642B.

573

574 Although sometimes problematic due to the occurrence of reworked specimens  
575 in Upper Pliocene sediments, a generally good index species for correlating  
576 Lower Pliocene deposits between the Norwegian Sea and Danish North Sea is  
577 *Reticulosphaera actinocoronata*. This species has its LO within the  
578 *Melitasphaeridium choanophorum* Zone in the Danish North Sea (Dybkjær and  
579 Piasecki, 2010) and defines the top of our VP2 Zone. It has a relatively well-  
580 established HO in several North Atlantic locations at 4.4–4.5 Ma (Louwye et al.,  
581 2004; Schreck et al., 2012) adding value to its use as an index fossil. Using the HO  
582 of *Reticulosphaera actinocoronata* to subdivide the *Melitasphaeridium*  
583 *choanophorum* Zone in the Danish North Sea could further refine the  
584 dinoflagellate cyst biozonation there.

586The top of the *Melitasphaeridium choanophorum* Zone (Dybkjær and Piasecki,  
5872010) is characterised by the HO of *Melitasphaeridium choanophorum* which  
588occurs ~3.6 Ma according to the data compilation of Powell and Brinkhuis in  
589Lourens et al. (2005). In contrast, this species is recorded in the VP3 Zone until  
590~3.3 Ma in the Norwegian Sea and has a HPO around 3.0 Ma in the North  
591Atlantic, where also occasional occurrences are recorded in the Pleistocene  
592(Head and Westphal, 1999; De Schepper and Head, 2009). In fact, this species is  
593recorded in modern sediments of the Gulf of Mexico, which may act as a refuge  
594for this warm-water species today (Limoges et al., 2013). It is thus clear that the  
595HO of *Melitasphaeridium choanophorum* is diachronous, and an age of 3.6 Ma – as  
596listed in the compilation of Powell and Brinkhuis in Lourens et al. (2005) – for  
597the upper boundary of the *Melitasphaeridium choanophorum* Zone in the Danish  
598Basin is probably too old. Especially since it occurs in younger deposits in the  
599Norwegian Sea (this study), North Atlantic (De Schepper and Head, 2009) and  
600the southern North Sea Basin (Louwye et al., 2004; De Schepper et al., 2009). In  
601addition, the HOs of *Invertocysta lacrymosa* and *Barssidinium graminosum* occur  
602just above the HO of *Melitasphaeridium choanophorum* in the Danish North Sea  
603(Dybkjær and Piasecki, 2010). Both *Invertocysta lacrymosa* and *Barssidinium*  
604*graminosum* have a last appearance at 2.74 Ma, about 250,000 years after the last  
605appearance of *Melitasphaeridium choanophorum* in the eastern North Atlantic  
606(De Schepper and Head, 2008a; 2009). Also at the Vøring Plateau both species  
607occur well into the Upper Pliocene (this study). Together, this suggests a Late  
608Pliocene age for the top of the Danish Basin *Melitasphaeridium choanophorum*

609Zone rather than an Early Pliocene age proposed by Dybkjær and Piasecki  
610(2010).

### 6114.3 **Correlations with the Iceland Sea and North Atlantic**

#### 6124.3.1 Upper Miocene correlations

613There are only few sections available for comparison that cover the Upper  
614Miocene to Pliocene in the North Atlantic realm, and these often only have  
615limited age control (e.g. Labrador Sea ODP Site 646; de Vernal and Mudie, 1989).  
616Eastern North Atlantic DSDP Hole 610A has a calibrated dinoflagellate cyst  
617zonation that covers the Lower Pliocene from ca. 4 Ma onwards to Pleistocene  
618(Figure 4) (De Schepper and Head, 2009). In Iceland Sea ODP Hole 907A, Schreck  
619et al. (2012) recorded stratigraphically-calibrated ranges of selected  
620dinoflagellate cyst species for the Miocene to Pliocene, but a biozonation was not  
621established.

622

623Correlations of the uppermost Miocene at ODP Site 642 with the Iceland Sea and  
624North Atlantic are hampered by the limited availability of marker species for this  
625time interval (e.g. Schreck et al., 2012). For example, *Cristadinium*  
626*cristatoserratum* (HO at 8.3 Ma) and the acritarch *Decahedrella martinheadii* (HO  
6276.5 Ma) are two of the Upper Miocene markers of Iceland Sea ODP Hole 907A  
628(Schreck et al., 2012), but these events are older than the studied section at ODP  
629Hole 642B.

#### 6304.3.2 Lower Pliocene correlations

631The HO of *Reticulosphaera actinocoronata* likely presents the best correlative  
632marker within the Nordic Seas and with the wider North Atlantic region in the

633 Lower Pliocene. The last appearance in the Norwegian Sea at 4.64 Ma and in the  
634 Iceland Sea at 4.45 Ma correspond favourably to the relatively well-established  
635 last appearance in several North Atlantic locations at around 4.4–4.5 Ma (reviews  
636 in Louwye et al., 2004; Schreck et al., 2012). The HO of *Reticulatosphaera*  
637 *actinocoronata* could therefore be used in the future for a Neogene North Atlantic  
638– Nordic Seas reference stratigraphy.

639

640 Applying the eastern North Atlantic zonation of De Schepper and Head (2009) to  
641 the Vøring Plateau is problematic due to the difference in stratigraphic ranges of  
642 the marker species. Whereas several Early Pliocene bioevents and biozones are  
643 near-synchronous between the Norwegian Sea and Iceland Sea, these are  
644 strongly diachronous compared to the North Atlantic (Figure 4). The Lower  
645 Pliocene North Atlantic RT1 to RT3 zones correlate to the uppermost VP2 and  
646 lowermost VP3 zones, but this correlation is strongly diachronous. The  
647 sequential disappearance of *Corrudinium devernaliae* (3.90 Ma), *Batiacasphaera*  
648 *micropapillata* complex (3.83 Ma) and *Operculodinium tegillatum* (3.71 Ma),  
649 which characterise the upper boundaries of respectively the North Atlantic RT1,  
650 RT2 and RT3 zones, is not observed in the Nordic Seas (Figure 4). Instead, in ODP  
651 Hole 642B, the HOs of *Operculodinium tegillatum* and *Corrudinium devernaliae*  
652 are observed in one sample (4.49 Ma) immediately above the HO of the  
653 *Batiacasphaera micropapillata* complex (4.64 Ma). Also, all three species have  
654 their HO (or HPO) in the Early Pliocene of Iceland Sea ODP Site 907 between  
655 4.45–4.55 Ma (Schreck et al., 2012). Thus, the HOs of *Corrudinium devernaliae*,  
656 *Batiacasphaera micropapillata* complex and *Operculodinium tegillatum* are  
657 recorded at least 500,000 years later in the North Atlantic than in the Iceland and

658 Norwegian seas. This discrepancy in timing of the bioevents between both basins  
659 can likely be attributed to the first development of the modern ocean circulation  
660 in the Nordic Seas around 4.5 Ma (De Schepper et al., 2015). The  
661 contemporaneous extinction events and major overturn in the phytoplankton  
662 assemblages across the Nordic Seas, suggest a cooling throughout the Nordic  
663 Seas followed by the development of a proto-East Greenland Current and  
664 Norwegian Atlantic Current.

665

666 Correlation of the North Atlantic RT4 Zone (De Schepper and Head, 2009) to the  
667 Norwegian Sea appears similarly problematic. The RT4 Zone is the interval  
668 between the HO of *Operculodinium tegillatum* and the HO of *Impagidinium*  
669 *solidum* in DSDP Hole 610A and straddles the Early–Late Pliocene boundary. In  
670 the Nordic Seas, it is not practical to define a biozone using the HOs of these two  
671 species. Firstly, we already established that the HO of *Operculodinium tegillatum*  
672 is recorded up to 800,000 years earlier in the Norwegian Sea than in the North  
673 Atlantic. Secondly, *Impagidinium solidum* occurs rarely in ODP Hole 642B  
674 between 3.83 and 3.59 Ma and this may not represent its total stratigraphic  
675 range. If the record at 3.59 Ma does represent its HO, then the HO of  
676 *Impagidinium solidum* is also recorded 500,000 years earlier in the Norwegian  
677 Sea (3.59 Ma) than in the North Atlantic (3.15 Ma). Since *Impagidinium solidum*  
678 has a preference for warm waters (De Schepper et al., 2011), its presence in the  
679 Nordic Seas may be restricted to the Early Pliocene, whereas it can persist longer  
680 in the warmer, more southerly eastern North Atlantic.

681

### 6824.3.3 Upper Pliocene correlations

683The stratigraphic ranges of Upper Pliocene dinoflagellate cysts are also strongly  
684diachronous between the Nordic Seas and North Atlantic. For example,  
685*Melitasphaeridium* sp. A of De Schepper and Head (2008b) and *Operculodinium?*  
686*eirikianum* var. *crebrum* have overlapping ranges near the Early–Late Pliocene  
687boundary in the Norwegian Sea (3.6–3.5 Ma). Such overlap is also encountered in  
688the North Atlantic, yet this occurs considerably later around ca. 3.2 Ma (De  
689Schepper and Head, 2008a). This observation demonstrates also a diachroneity  
690between the North Atlantic and Norwegian Sea bioevents in the Late Pliocene.

691

692A Late Pliocene major extinction event is recognised in the Norwegian Sea  
693around 3.3 Ma when four dinoflagellate cysts (*Heteraulacacysta* sp. A of Costa  
694and Downie (1979), *Invertocysta lacrymosa*, *Operculodinium janduchenei* and  
695*Melitasphaeridium choanophorum*) and one acritarch (*Lavradosphaera crista*)  
696disappear simultaneously (Figure 2, 4, Appendix). Their disappearance from ODP  
697Hole 642B at 67.60 mbsf is not related to the hiatus in the upper part of the  
698Gauss Chron of ODP Hole 642B which occurs higher around 66 mbsf (Bleil,  
6991989). These HOs, dated at 3.27 Ma, could be related to a major Northern  
700Hemisphere cooling event during MIS M2 (ca. 3.3 Ma; De Schepper et al., 2013).  
701However, after MIS M2 sea surface temperatures remain fluctuating between 0  
702and 3 °C above Holocene values (Bachem et al. 2016), suggesting that high  
703latitude cooling may not be the only cause of their disappearance. Irrespective of  
704what is causing the HOs, a strong asynchrony remains apparent with their HOs in  
705the North Atlantic (Figure 4), where these species range into younger Late  
706Pliocene sediments (< 3 Ma). It is thus possible to recognise the North Atlantic



707RT5 zone, defined between the HO of *Impagidinium solidum* (but see earlier  
708comments on its range) and HO of *Invertocysta lacrymosa*, in the Norwegian Sea  
709where it corresponds to the upper VP3 Zone (Figure 3, 4). But also here, there is  
710strong diachroneity between the different zones (RT5 Zone, 3.15–2.74 Ma vs. VP3  
711Zone, 4.49–3.27 Ma), which could be related to deteriorating climate associated  
712with the onset of the Northern Hemisphere glaciation and/or the hiatus in the  
713Norwegian Sea record. These species are not recorded from the Iceland Sea  
714where the Late Pliocene sediments are generally barren (Schreck et al., 2012).

## 7155 **Conclusions**

716Three dinoflagellate cyst interval biozones (VP1, VP2, VP3) are established for  
717the latest Miocene to Late Pliocene of Vøring Plateau ODP Hole 642B using the  
718most up-to-date taxonomy and a higher resolution compared to the pioneering  
719work of Mudie (1989). The zones and stratigraphic ranges of selected  
720dinoflagellate cysts and acritarchs are calibrated to the available  
721magnetostratigraphy (Bleil, 1989). The oldest Zone VP1 corresponds to the late  
722Messinian–earliest Zanclean, Zone VP2 is restricted to the early Zanclean and the  
723youngest Zone VP3 covers the late Zanclean to middle Piacenzian. Some of the  
724Upper Miocene Arctic (e.g. *Decahedrella martinheadii*) and North Atlantic (e.g.  
725*Barssidinium evangelinae*) marker species were missing from our studied  
726interval in ODP Hole 642B, hampering correlations with the Upper Miocene  
727biozones from the Iceland Sea and Danish Basins. The HOs of the dinoflagellate  
728cyst species *Reticulosphaera actinocoronata* and *Invertocysta lacrymosa* were  
729used to define the upper boundaries of the VP2 Zone, and the upper boundary of  
730the VP3 Zone, respectively. These two Pliocene bioevents seem to be relatively

731synchronous between the Nordic Seas and North Atlantic and may serve as the  
732basis for a future Pliocene North Atlantic–Arctic reference stratigraphy.

733

734Our dinoflagellate cyst biozonation for the Vøring Plateau ODP Hole 642B has  
735potential to be a benchmark for linking Arctic and North Atlantic zonations and  
736for dating and correlating late Neogene deposits along the Norwegian Shelf. The  
737Early Pliocene Norwegian Sea bioevents (disappearances of *Reticulatosphaera*  
738*actinocoronata*, *Operculodinium tegillatum*, *Corrudinium devernaliae*,  
739*Batiacasphaera micropapillata* complex) are observed in contemporaneous  
740Iceland Sea sediments, but these Nordic Seas events are strongly asynchronous  
741with the North Atlantic. Late Pliocene Norwegian Sea HO (e.g. HOs of  
742*Ataxiodinium confusum*, *Operculodinium? eirikianum* var. *crebrum*,  
743*Melitasphaeridium choanophorum*) are also asynchronous with the North  
744Atlantic, demonstrating that several events are only useful for local to regional  
745correlations. Because of the diachronous ranges of several species in the different  
746basins, it is clear that one must apply North Atlantic biozonation schemes with  
747caution in the Nordic Seas (e.g. Anthonissen, 2009) or if applying the Nordic Seas  
748biozonation into the North Atlantic and North Sea. To improve inter-basin  
749correlations, a well-calibrated, continuous upper Neogene record in the southern  
750Norwegian Sea – northern North Sea region would be a major asset in tying the  
751North Atlantic and Norwegian biozonations together.

752

753Finally, while we focused on the Late Miocene and Pliocene, it would be valuable  
754to re-investigate Early and Middle Miocene marine palynology (Manum et al.,  
7551989) applying the modern taxonomic concepts and biozonation schemes

756available (de Verteuil and Norris, 1996; Munsterman and Brinkhuis, 2004;  
757Dybkjær and Piasecki, 2010) to improve understanding of Neogene  
758oceanographic and climate evolution of the Arctic region.

759

## 760Taxonomic appendix

761

762"*Veriplicidium franklinii*" of Anstey (1992): This thin-walled, autophragmal  
763palynomorph with scabrate to finely granulate wall was first described in an  
764unpublished MSc thesis (Anstey, 1992). The vesicle appears elongate, likely due  
765to the typical occurrence of several mainly longitudinal folds of the autophragm.  
766On our specimens, an excystment aperture was not observed, but Anstey (1992)  
767reports an angular rupture-type aperture in the apical region based on SEM  
768analysis. Measurements based on 8 specimens: vesicle length, 27[32]34 µm;  
769width, 19[21]23 µm. These compare well with the dimensions reported by  
770Anstey (1992): length, 27[32]38 µm; width, 10[24]27 µm.

771

772*Batiacasphaera micropapillata* complex and *Batiacasphaera minuta* s.s.: We  
773follow the concept of Schreck and Matthiessen (2013) which groups all  
774(sub)spherical, transparent specimens bearing an apical archeopyle and a  
775microreticulate, vermiculoreticulate–vermiculate and rugulate ornamentation.  
776Our *Batiacasphaera micropapillata* complex likely includes specimens that can be  
777attributed to the exclusively microreticulate *Batiacasphaera minuta* s.s. (e. g. in  
778sample 642B-10H5, 115–116 cm; 642B-10H5, 11–12 cm), but when  
779*Batiacasphaera minuta* s.s. was unequivocally recognised, it was categorised as  
780such.

781

782*Batiacasphaera/Pyxidinosia* spp. indet.: This group contains all (sub)spherical,  
783transparent cysts bearing a low wall ornament (granulate, punctate to

784punctoreticulate) of which the archeopyle (precingular or apical?) could not  
785clearly be determined.

786

787*Bitectatodinium* sp. A: (Sub)spherical cysts bearing a precingular (2P) archeopyle  
788as for the genus *Bitectatodinium*. This species has a wall ornament that is  
789intermediate between the typical vermiculate ornament of *Bitectatodinium*  
790*tepikiense* and the “hairy” ornament of *Bitectatodinium raedwaldii*. The wall  
791ornament of *Bitectatodinium* sp. A is a dense pattern of pili (as for *B. raedwaldii*  
792Head, 1997) and short, straight (as for *B. tepikiense* Wilson, 1973) or irregular  
793polygonal ridges. This pattern is best observed using L-O (Lux-Obscura) analysis  
794at 1000x magnification. *Bitectatodinium* sp. A is here considered to have a  
795transitional morphology between the end-members *Bitectatodinium raedwaldii*  
796and *Bitectatodinium tepikiense*, and is therefore left in open nomenclature.

797

798“*Impagidinium densiverrucosum*” Zevenboom and Santarelli in Zevenboom  
799(1995): Rare but distinct large, (sub)spherical cyst with a dense and coarse  
800verrucate wall ornament on all plates. Relatively high and smooth crests appear  
801to delineate tabulation, likely of an *Impagidinium*. The encountered specimens  
802are often broken. Measurements based on 5 specimens: maximum inner  
803diameter, 66[78]88  $\mu\text{m}$ ; crest height, 3[3.3]3.5  $\mu\text{m}$ . Diameter measurements  
804should be treated with care since 4 specimens were broken and one was folded.  
805This taxon was listed as *Impagidinium vermiculatum* by Beck (2013).

806

807Cysts of *Protoceratium reticulatum*: Paez-Reyes and Head (2013) proposed this  
808name for specimens of *Operculodinium centrocarpum* sensu Wall and Dale, 1966.  
809Both forms with short and long processes are included here.

810

811*Pyxidinopsis* sp. A: (Sub)spherical transparent cysts with apical archeopyle,  
812bearing a low wall ornament that is granulate, punctate to punctoreticulate. This  
813taxon closely resembles *Batiacasphaera sphaerica* except for the archeopyle  
814which is precingular. When the archeopyle was not fully visible, specimens were  
815generally grouped under *Batiacasphaera/Pyxidinopsis* spp. indet., however it is  
816possible that some specimens of *Batiacasphaera sphaerica* may have been  
817classified as *Pyxidinopsis* sp. A.

818

819Cyst type I of de Vernal and Mudie (1989): spherical palynomorph, possibly a  
820dinoflagellate cyst with an apical archeopyle. Numerous thin solid, straight and  
821bent processes end distally in broad, ornamented trabeculae. Trabeculae delimit  
822large (4–6 µm diameter) and occasional small (2 µm diameter) circular to oval  
823openings. Large excystment aperture (apical archeopyle?), operculum free.

824

825*Lavradosphaera?* sp.: Questionably assigned to the genus *Lavradosphaera* due to  
826the very reduced cancellous wall, which consist of solitary thin rods that support  
827the outer wall on these specimens.

828

829*Lavradosphaera* sp. cf. *canalis*: Spherical cyst bearing U-shaped channels typical  
830of *Lavradosphaera canalis*. The specimens are assigned questionably to the  
831species because the typical cancellous wall structure for the genus

832 *Lavradosphaera* is very reduced, and appears to consist of several, solitary thin  
833 rods supporting the outer wall.

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843

844

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1021

## 1022 **FIGURE CAPTIONS**

1023 **Figure 1.** Bathymetric map of the Nordic Seas showing the location of ODP Site  
1024 642 in the Norwegian Sea and the other locations in the Iceland Sea (ODP Site  
1025 907), the North Atlantic (DSDP Site 610) and the North Sea Basin discussed in  
1026 the text.

1027

1028 **Figure 2.** ODP Hole 642B lithology, studied samples (core, section and depth  
1029 (cm) in section are indicated) and stratigraphic occurrence of selected  
1030 dinoflagellate cysts and acritarch taxa and biozones defined in this study. Grey  
1031 shading in the biozonation indicate the intervals between the biozones that were  
1032 not sampled. All raw data are available from the range charts in the appendix and  
1033 from <http://doi.pangaea.de/10.1594/PANGAEA.846838>.

1034

1035 **Figure 3.** Comparison of the new zonation at ODP Hole 642B to the  
1036 palaeomagnestratigraphy (Bleil, 1989) and previously established zonations  
1037 using dinoflagellate cysts (Mudie, 1989), planktonic foraminifers (Spiegler and  
1038 Jansen, 1989), radiolarians (Goll and Bjørklund, 1989), and calcareous  
1039 nanofossils (Donnally, 1989). Grey shading indicates intervals between the  
1040 biozones that were not sampled, i.e. the interval between the sample with the  
1041 highest occurrence of a taxon and the next sample above.

1042

1043 **Figure 4.** Early and Late Pliocene stratigraphic ranges of selected dinoflagellate  
1044 cysts in the Norwegian Sea ODP Hole 642B (this study), Iceland Sea ODP Hole  
1045 907A (Schreck et al., 2012), eastern North Atlantic DSDP Hole 610A (De  
1046 Schepper and Head, 2009; 2008a), western North Atlantic DSDP Hole 603C (M.J.

1047Head, unpublished data) and Labrador Sea ODP Hole 646B (de Vernal and Mudie,  
10481989). The biozonation of the Norwegian Sea (this study) is compared to  
1049zonations in Denmark (Dybkjær and Piasecki, 2010) and the eastern North  
1050Atlantic (De Schepper and Head, 2009). Correlations of the zonal boundaries  
1051using the marker species between the different zonations is shown. Correlation  
1052of individual taxa with the Danish zonation is approximate and does not reflect  
1053precise absolute ages derived from within the Danish North Sea basin.

1054Abbreviations used: Oteg = *O. tegillatum*, Bmin = *B. minuta*/*B. micropapillata*  
1055complex, Cdev = *C. devernaliae*, Bgr = *B. graminosum*, Ilac = *I. lacrymosa*, Mcho =  
1056*M. choanophorum*, Isol = *I. solidum*.

1057

#### 1058TABLE CAPTIONS

1059**Table 1.** Samples studied from ODP Hole 642B, dry weight of sediment  
1060processed for palynology, counts of *Lycopodium clavatum* spike added, and  
1061counts and concentrations of dinoflagellate cyst, acritarchs and terrestrial  
1062palynomorphs.

1063

1064**Table 2.** Magnetostratigraphic reversals in ODP Hole 642B identified by Bleil  
1065(1989) and updated to Geological Time Scale 2012 (Hilgen et al. 2012).

1066

1067**Table 3.** Highest (HO) and lowest (LO) occurrence of dinoflagellate cysts and  
1068acritarchs (indicated by asterisk) in ODP Hole 642B, including estimated age and  
1069error (based on sampling interval).

1070

1071

1072

1073**APPENDIX**

1074The appendix contains stratigraphic range charts containing the raw data  
1075(counts) of dinoflagellate cysts and acritarchs for ODP Hole 642B. Red shading  
1076shows taxa used for defining the biozones, green shading shows taxa with  
1077important stratigraphic ranges.



## 1078 PLATE CAPTIONS

### 1079 Plate I

1080 Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
1081 depth (cm) in section. England Finder references and slide number are also  
1082 given. All photographs are taken in transmitted light. 1,2. *Achomosphaera*  
1083 *andalousiensis andalousienis* Jan du Chene, 1977 emend. Jan du Chene and  
1084 Londeix, 1988. ODP 642B-11H-3, 65–66 cm, W45/0, slide 11J501. Uncertain  
1085 view at (1) high focus on characteristic fenestrate process tip, and at (2) mid-  
1086 focus showing the free archeopyle. 3,4. *Ataxiodinium confusum* Versteegh and  
1087 Zevenboom in Versteegh, 1995. ODP 642B-9H-2, 65–66 cm, B55/2, slide 12F475.  
1088 Oblique dorsal view at (3) high focus on archeopyle and granular wall, and at (4)  
1089 mid-focus demonstrating the dorsal and ventral connections between endo- and  
1090 periphragm. 5,6. *Ataxiodinium choane* Reid, 1974. ODP 642B-10H-1, 40–41 cm,  
1091 Y67/1, slide 12G102. Ventral view at (1) mid-focus with view on faintly granular  
1092 pericyst, and at (2) low focus on archeopyle. 7,8. *Ataxiodinium zevenboomii* Head,  
1093 1997. ODP 642B-11H-7, 15–16 cm, V64/0, slide 11J512. Dorsal/apical? view at  
1094 (7) high focus on archeopyle and dorsal/apical surface, and (8) mid-focus. 9,10.  
1095 *Amiculosphaera umbraculum* Harland, 1979. ODP 642B-9H-1, 120–121 cm,  
1096 Y38/2, slide 12F407. Dorsal view at (9) mid-focus on antapical funnel-shaped  
1097 connection between peri- and endophragm, and at (10) low focus revealing  
1098 traces of tabulation (cingulum?) on the periphragm. Note the apical boss on the  
1099 endophragm (9). 11,12. *Cerebrocysta poulsenii* de Verteuil and Norris, 1996. ODP  
1100 642B-11H-7, 55–56 cm, T31/2, slide 11J513. Oblique dorsal view at (11) high  
1101 focus on granulate wall and at (12) mid-focus. 13. *Barssidinium pliogenicum*  
1102 (Head, 1993) De Schepper and Head, 2004. ODP 642B-9H-3, 70–71 cm, G51/1,

1103slide 12G12. Vertical stack of several foci. 14,15. *Dapsilidinium pastielsii* (Davey  
1104and Williams, 1966) Bujak, Downie, Eaton and Williams, 1980. ODP 642B-9H-2,  
110565–66 cm, J48/3, slide 12F475. Antapical view of (14) high focus on granulate  
1106wall, and at (15) mid-focus. 16. *Heteraulacacysta* sp. A of Costa and Downie,  
11071979. ODP 642B-9H-3, 70–71 cm, G33/2, slide 12G12. Vertical stack of several  
1108foci. 17, 18. *Impagidinium aculeatum* (Wall, 1967) Lentin and Williams, 1981.  
1109ODP 642B-9H-3, 60–61 cm, J41/1, slide 12G10. Dorsal view at (17) high focus on  
1110archeopyle (3”), and at (18) mid-focus. 19, 20. *Impagidinium pallidum* Bujak,  
11111984. ODP 642B-9H-3, 70–71 cm, E32/3, slide 12G12. Dorsal view at (19) high  
1112focus on archeopyle (3”), illustrating the incomplete expression of the tabulation,  
1113and at (20) mid-focus.

1114

## 1115Plate II

1116Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
1117depth (cm) in section. England Finder references and slide number are also  
1118given. All photographs are taken in transmitted light. 1–3. *Batiacasphaera*  
1119*hirsuta* Stover, 1977. ODP642B-11H-3, 65–66 cm, W54/0, slide 11J501. Uncertain  
1120view at slightly different high foci (1,2) illustrating the pili (hairs) characteristic  
1121for this species and the apical archeopyle at the top, and at (3) mid-focus. 4, 8.  
1122*Invertocysta lacrymosa* Edwards, 1984. ODP 642B-9H-3, 60–61 cm, O50/1, slide  
112312G10. Dorsal view at (4) high focus, and at (8) low focus showing an apical boss  
1124on the endocyst. 5–7. *Batiacasphaera micropapillata* complex sensu Schreck and  
1125Matthiessen, 2013. ODP 642B, 11H3, 65–66 cm, M43/3, slide 11J501. Apical view  
1126at (5) high focus on apical archeopyle and microreticulate wall ornament, at (6)  
1127mid-focus, and at (7) low focus on antapical surface. 9–16. *Bitectatodinium* sp. A.

1128(9–12) ODP 642B-8H-6, 129–130 cm, F34/0, slide 12F401. Ventral view at  
1129slightly different high foci (9,10) illustrating a dense ornament of pili and short  
1130ridges (“columellate”) on the wall revealed by L(ux)–O(bscura) analysis; at (11)  
1131mid-focus and at (12) low focus on 2P archeopyle. (13–16) ODP 642B-8H-6, 129–  
1132130 cm, G35/0, slide 12F401. Dorsal view at (13,14) high focus on 2P archeopyle  
1133and dense wall ornament of pili and short ridges (“columellate”); at (15) mid-  
1134focus and at (16) low focus on ventral surface. 17–20. *Melitasphaeridium*  
1135*choanophorum* (Deflandre and Cookson 1955) Harland and Hill, 1979. ODP  
1136642B-9H-2, 65–66 cm, K48/3, slide 12F475. Oblique ventral view at (17) high  
1137focus on aculeate process tips; at (18) slightly lower high focus on granular wall;  
1138at (19) mid-focus and at (20) lower focus on 3” archeopyle.

1139

### 1140Plate III

1141Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
1142depth (cm) in section. England Finder references and slide number are also  
1143given. All photographs are taken in transmitted light. 1–3. *Corrudinium?*  
1144*labradori* Head, Norris and Mudie, 1989. ODP 642B-10H-4, 15–16 cm, X56/0,  
1145slide 11J378  
1146Dorsal view at (1) high focus on 2P archeopyle, reticulate wall ornament and  
1147cingulum expressed by single margin; at (2) mid-focus and at (3) ventral surface  
1148with expressed cingulum and possibly one margin of the sulcal area. 4,8.  
1149*Filisphaera filifera* subsp. *filifera* Bujak, 1984 emend. Head, 1994. ODP642-9H-2,  
115065–66 cm, B53/4, slide 12F475. Oblique view at (4) high focus on finely  
1151reticulate wall with smaller and larger lumina, and at (8) mid-focus illustrating  
1152the radiating septa of the luxuria. 5–7. *Corrudinium devernaliae* Head and Norris,

11532003. ODP 642B-10H-1, 145–146 cm, L55/0, slide 12G104. Oblique dorsal view  
1154at (5) high focus on archeopyle and crests that form an incomplete and large  
1155reticulum; at (6) mid-focus and at (7) low focus on ventral surface. 9–11.

1156*Habibacysta tectata* Head, Norris and Mudie, 1989. ODP 642B-8H-6, 129–130 cm,  
1157N35/1, slide 12F401. Ventral view at (9) high focus showing the characteristic  
1158wall ornament, at (10) mid-focus, and at (11) lower focus on the 3” archeopyle  
1159and dorsal surface. 12,16. “*Impagidinium densiverrucosum*” of Zevenboom and  
1160Santarelli in Zevenboom (1995). ODP 642B-11H-7, 55–56 cm, U37/2, slide  
116111J513. Dorsal view at (12) high focus on precingular archeopyle (free) and  
1162coarsely verrucate wall ornament, and at (16) mid-focus. Note the 3” precingular  
1163operculum central in the picture. 13,14. *Pyxidinosia braboi* De Schepper and  
1164Head, 2004. ODP 642B-9H-1, 120–121 cm, D52/2, slide 12F407

1165Oblique dorsal view at (13) high focus and at (14) mid-focus. 15. *Selenopemphix*  
1166*nephroides* Benedek, 1972 emend. Bujak in Bujak, Downie, Eaton and Williams,  
11671980. ODP 642B-9H-3, 60–61 cm, W39/3, slide 12G10. Vertical stack of several  
1168foci. 17,18. *Operculodinium janduchenei* Head, Norris and Mudie, 1989. ODP  
1169642B-9H-2, 65–66 cm, J46/2, slide 12F475. Dorsal view at (17) high focus and at  
1170(18) mid-focus. Processes on this specimen are somewhat long compared to  
1171originally illustrated specimens in Head, Norris and Mudie, 1989. 19,20. Cyst of  
1172*Pentapharsodinium dalei* Indelicato and Loeblich III 1986. ODP 642B-9H-6, 100–  
1173101 cm, M61/0, slide 12G98. Uncertain view at subsequently lower foci (19,20)  
1174showing the solid processes that occasionally branch.

1175

1176**Plate IV**

1177 Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
 1178 depth (cm) in section. England Finder references and slide number are also  
 1179 given. All photographs are taken in transmitted light. 1–3, 5–7. *Operculodinium?*  
 1180 *eirikianum* var. *eirikianum* Head, Norris and Mudie, 1989 emend. Head, 1997. (1–  
 1181 13) ODP 642B-9H-3, 55–56 cm, V39/0, slide 12G9. Dorsal view at (1) high focus  
 1182 on large precingular archeopyle (3”), at (2) mid-focus, and at (3) lower focus. (5–  
 1183 7) ODP 642B-11H-5, 60–61 cm, X38/2, slide 11J507. Ventral view at (5) high  
 1184 focus on ventral surface, at (6) mid-focus, and at (7) low focus on dorsal surface  
 1185 with archeopyle. 4,8. *Operculodinium?* *eirikianum* var. *crebrum* De Schepper and  
 1186 Head, 2008. ODP 642B-9H-3, 60–61 cm, O49/0, slide 12G10. Uncertain view at  
 1187 (4) high focus on microreticulate wall structure and at (8) mid-focus. Note the  
 1188 very thick luxuria [2.5–3.0 µm], consisting of radiating, non-tabular septa. 9–  
 1189 11, 12, 16. *Operculodinium tegillatum* Head, 1997. (9–11) ODP 642B-10H-1, 145–  
 1190 146 cm, L56/0, slide 12G104. Apical view at (9) high focus showing the  
 1191 precingular archeopyle (3”), at (10) mid-focus, and at (11) lower focus. (12, 16).  
 1192 ODP 642B-10H-3, 102–103 cm, S62/0, slide 12G135. Uncertain view at (12) high  
 1193 focus and at (16) mid-focus. 13, 14. *Spiniferites elongatus* Reid, 1974. ODP 642B-  
 1194 9H-2, 65–66 cm, D57/2, slide 12F475. Dorsal view at (13) high focus and at (14)  
 1195 mid-focus. 15. *Selenopemphix brevispinosa* Head, Norris and Mudie, 1989. ODP  
 1196 642B-11H-7, 15–16 cm, V35/0, slide 11J512. Vertical stack of several foci. 17–20.  
 1197 *Reticulosphaera actinocoronata* (Benedek, 1972) Bujak and Matsuoka, 1986.  
 1198 (17–18) ODP 642B-10H-2, 40–41 cm, C48/0, slide 12G105. Uncertain view at  
 1199 (17) high focus on the irregularly branching process tips and at (18) mid-focus.  
 1200 (19–20) ODP 642B-10H-3, 102–103 cm, M37/0, slide 12G135. Uncertain view at

1201(19) high focus and at (20) mid-focus. Arrow indicates possible opercular plate  
1202with a single process.

1203

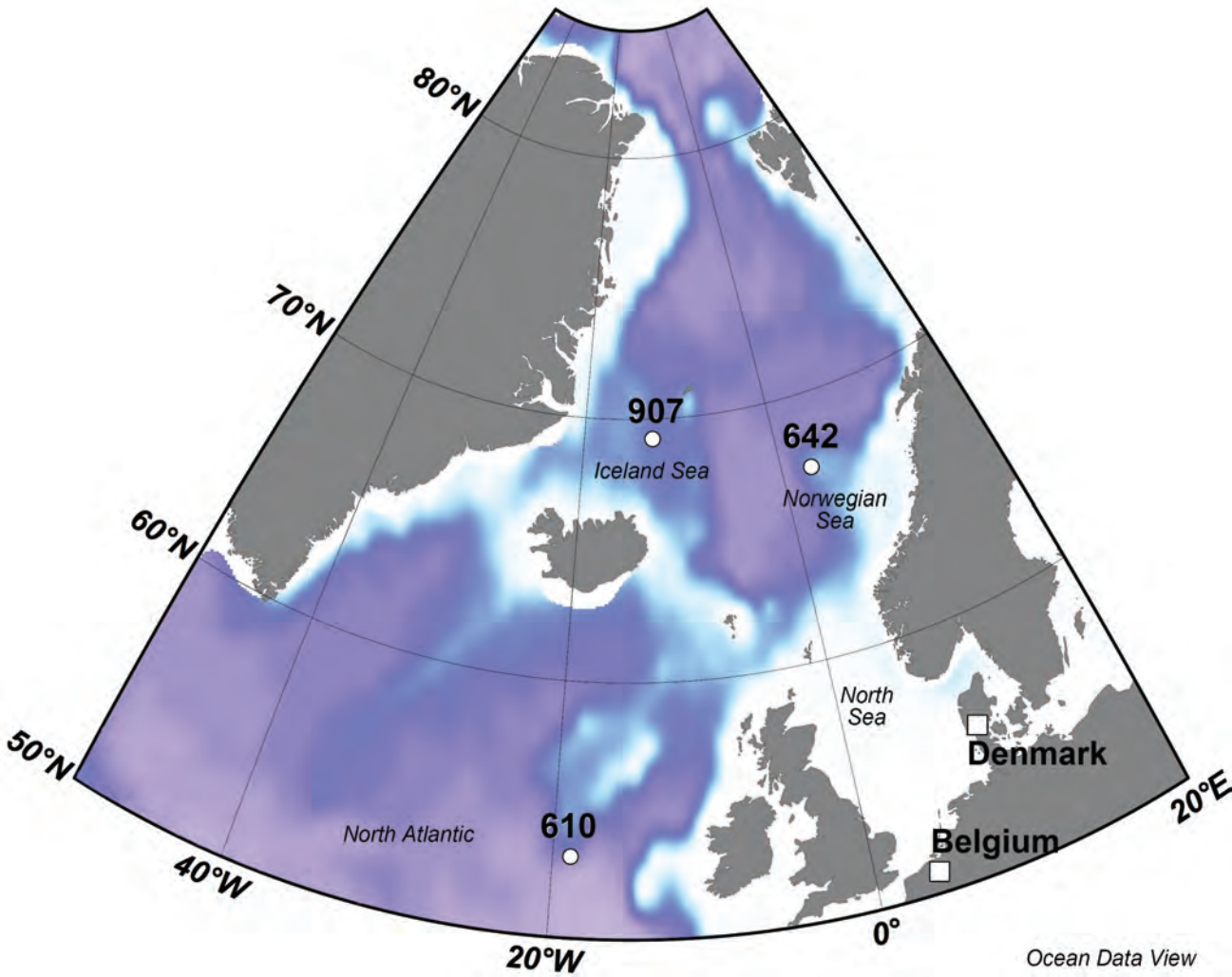
#### 1204**Plate V**

1205Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
1206depth (cm) in section. England Finder references and slide number are also  
1207given. All photographs are taken in transmitted light. 1–7. Cyst type I of de Vernal  
1208and Mudie (1989). (1–4) ODP 642B-10H-2, 40–41 cm, L40/3, slide 12G105.  
1209Uncertain view at (1) high focus on granular trabeculae forming a network with  
1210circular openings, at two slightly different mid-foci (2,3) showing numerous, thin  
1211solid processes ending distally in broad, ornamented trabeculae, and at (4) low  
1212focus on (apical?) archeopyle. (5–7) ODP 642B-10H-2, 40–41 cm, N41/0, slide  
121312G105. Uncertain view at (1) high focus, at (6) mid-focus, and at (7) low focus  
1214on (apical?) archeopyle. 8,12–16. *Lavradosphaera canalis*. (8,12) ODP 642B-11H-  
12156, 70–71 cm, W42/1, slide 11J510. Uncertain view at (8) high focus and (12) low  
1216focus. (13–16) ODP 642B-11H-6, 70–71 cm, W49/3, slide 11J510. Antapical view  
1217at (13) high focus, at two slightly different mid-foci (14,15) showing the U-  
1218shaped channels typical of *Lavradosphaera canalis*, and at (16) low focus on  
1219pylome. 9–11. "*Veriplicidium franklinii*" of Anstey (1992). ODP 642B-11H-7, 15–  
122016 cm, W32/0, slide 11J512. Uncertain view at (9) high focus, (8) mid-focus and  
1221(11) lower focus. 17–20. *Lavradosphaera?* sp.. ODP 642B-10H-3, 102–103 cm,  
1222056/3, slide 12G135. Two slightly different high foci (17,18), mid-focus (19) and  
1223low focus (20) revealing outline of pylome.

1224

#### 1225**Plate VI**

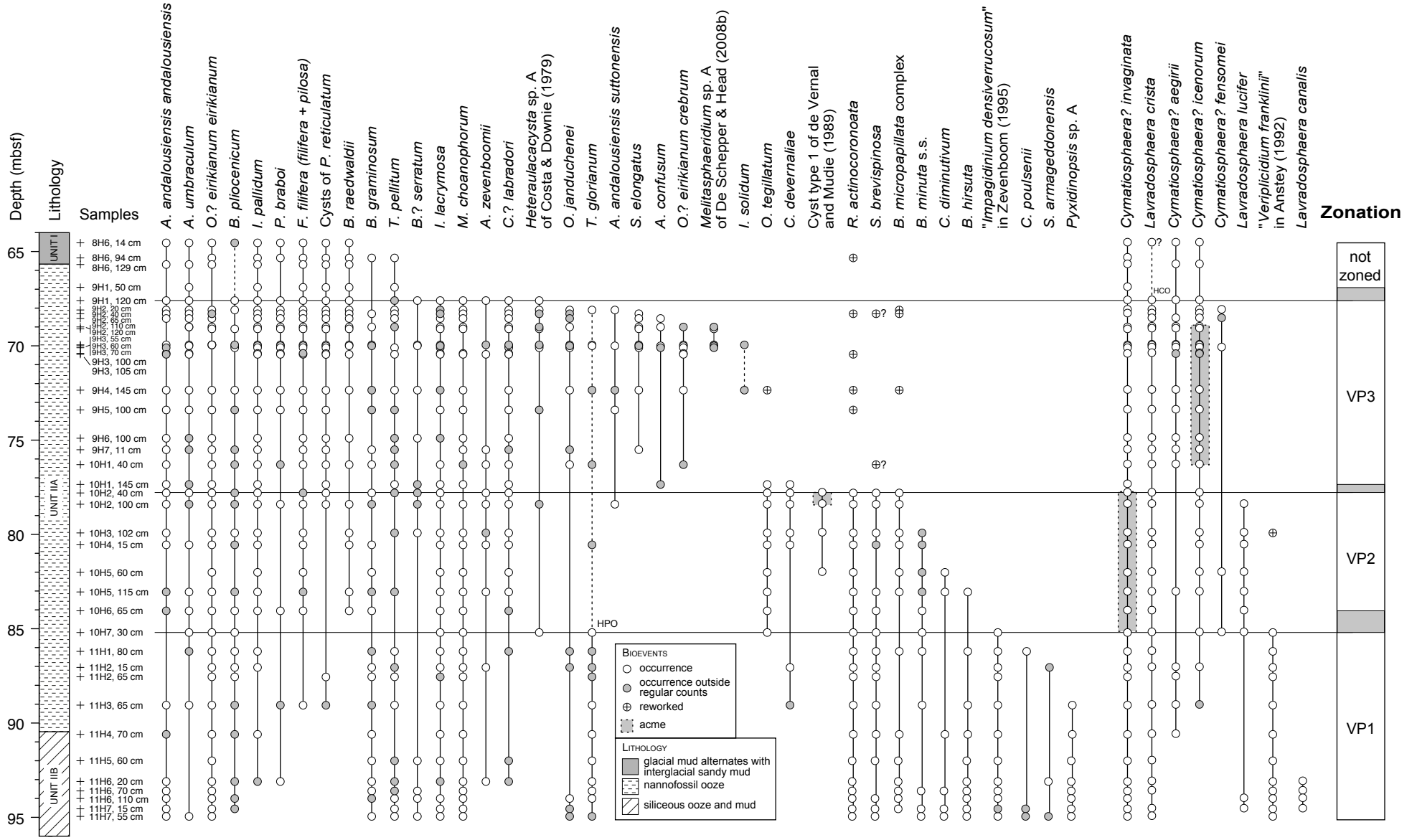
1226 Sample code follows ODP nomenclature, listing Site, Hole, Core, Section and  
1227 depth (cm) in section. England Finder references and slide number are also  
1228 given. All photographs are taken in transmitted light. 1–3. *Gelasinicysta vangeelii*  
1229 Head, 1992. ODP 642B-10H-4, 15–16 cm, X57/3, slide 11J378. High focus (1),  
1230 mid-focus (2) and low focus (3). 4. *Pediastrum*. ODP 642B-9H-3, 60–61 cm,  
1231 C45/2, slide 12G10. 5–7. *Cymatiosphaera? aegirii* De Schepper and Head, 2014.  
1232 ODP 642B-9H-3, 70–71 cm, H45/2, slide 12G12. Slightly lower high foci (5,6) on  
1233 trifurcate process tips that connect to the next process via thin, round and solid  
1234 trabeculae, and mid-focus (7). 8,12,16. *Cymatiosphaera? invaginata* Head, Norris  
1235 and Mudie, 1989. (8,12) ODP 642B-9H-3, 70–71 cm, Q49/0, slide 12G12  
1236 High focus (8) on undulating crests and mid-focus (12) showing the Y-shaped  
1237 distal ends of the crests. (16) ODP 642B-9H-3, 60–61 cm, W37/3, slide 12G10.  
1238 Mid-focus. 9–11. *Cymatiosphaera? icenorum* De Schepper and Head, 2014. ODP  
1239 642B-9H-3, 60–61 cm, W37/3, slide 12G10. Slightly different high foci (9,10)  
1240 showing a broad process typical for the species, and mid-focus (11). 13–14.  
1241 *Lavradosphaera crista* De Schepper and Head, 2008. ODP 642B-9H-3, 70–71 cm,  
1242 G46/0, slide 12G12. Apical view at (13) high focus on pylome, resembling an  
1243 apical archeopyle, and at (14) mid-focus. 15. *Nannobarbophora walldalei* Head,  
1244 1996. ODP 642B-9H-3, 60–61 cm, D44/0, slide 12G10.

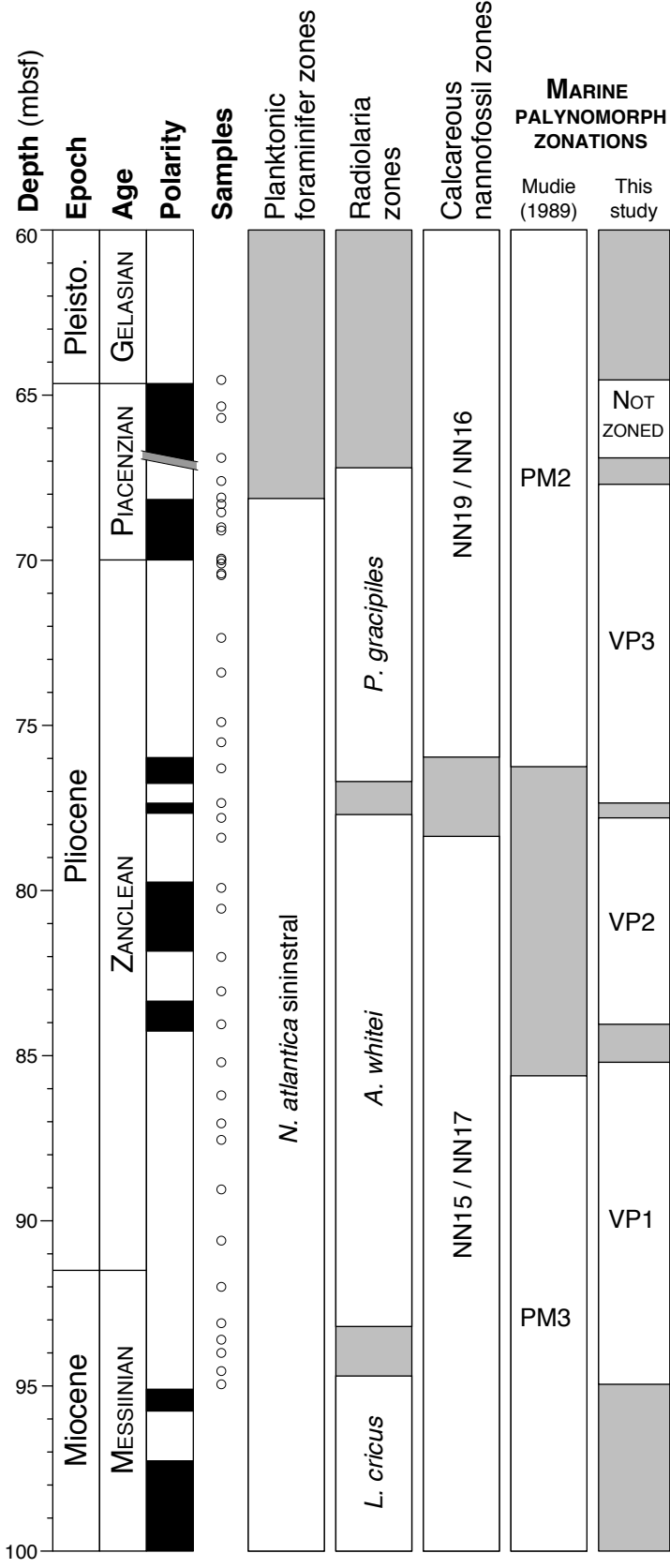




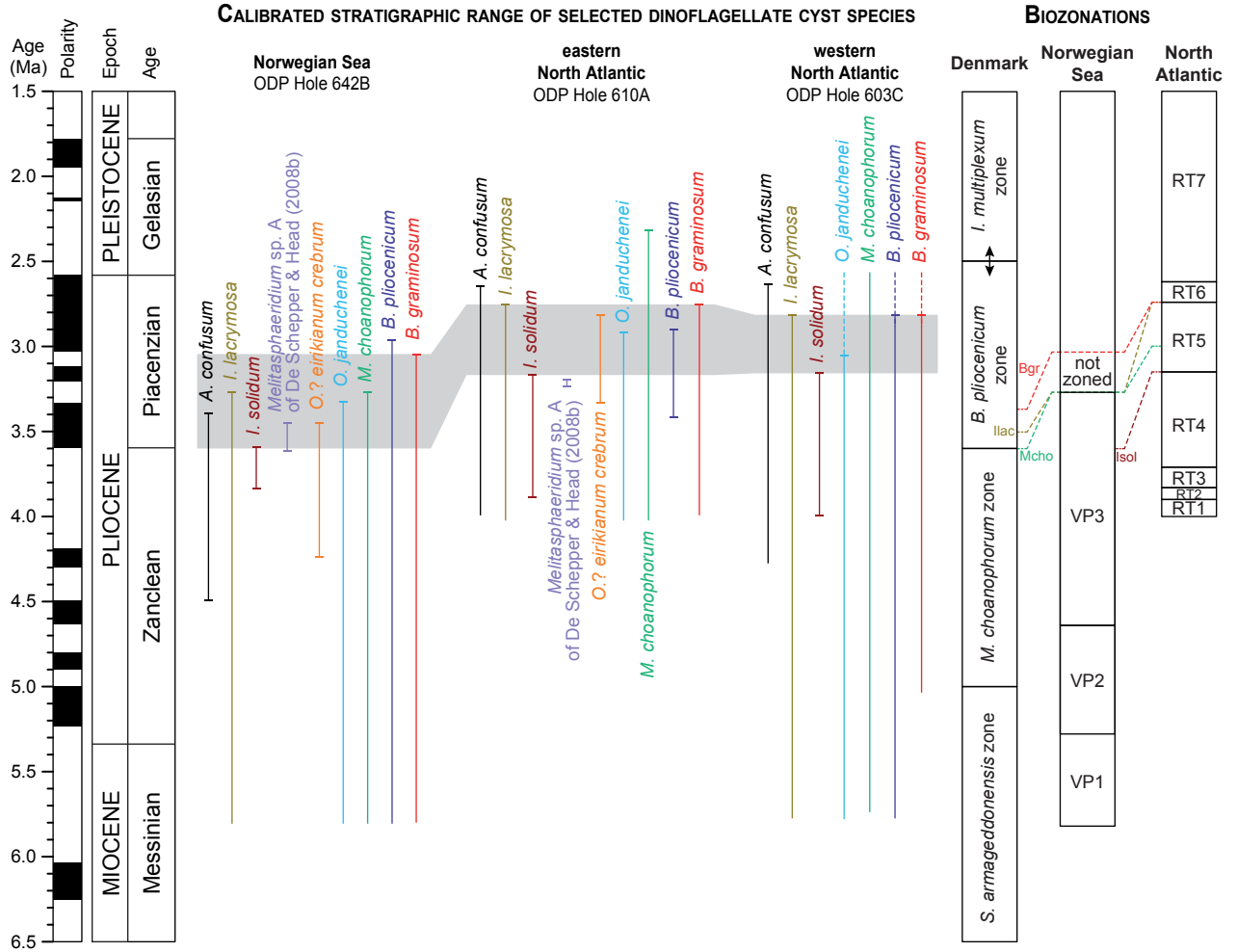
DINOFLAGELLATE CYSTS

ACRITARCHS

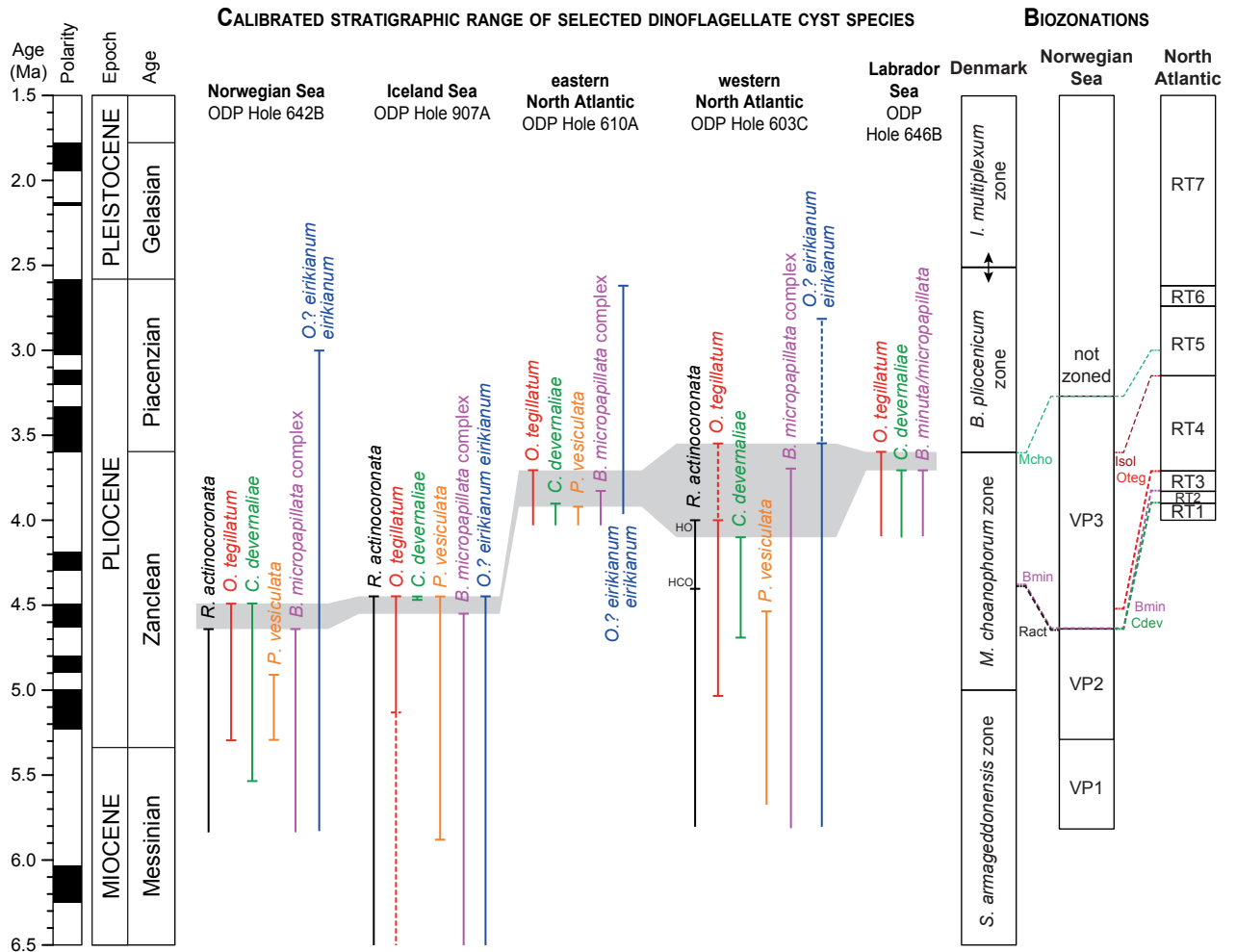




# LATE PLIOCENE



# EARLY PLIOCENE

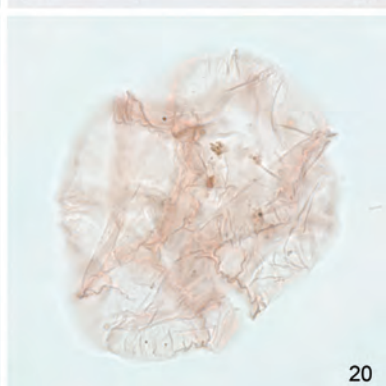
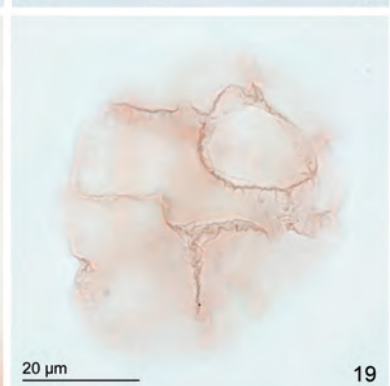
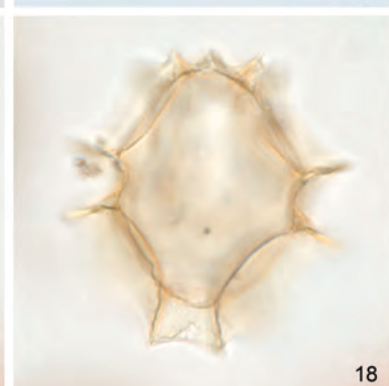
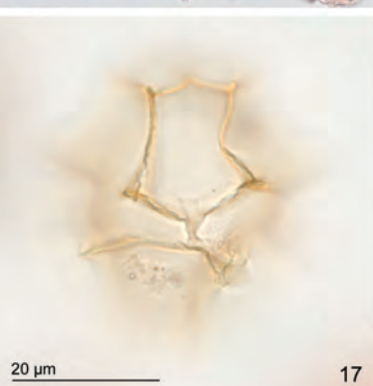
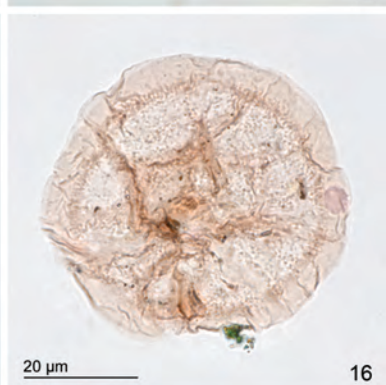
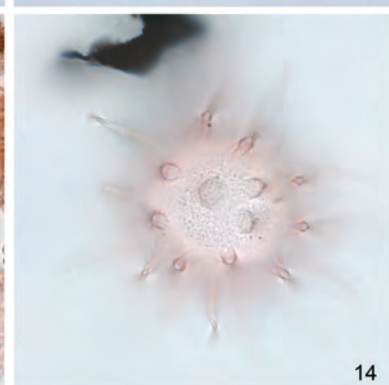
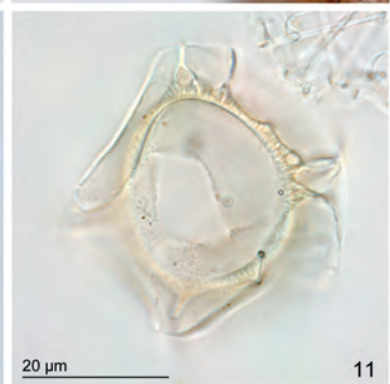
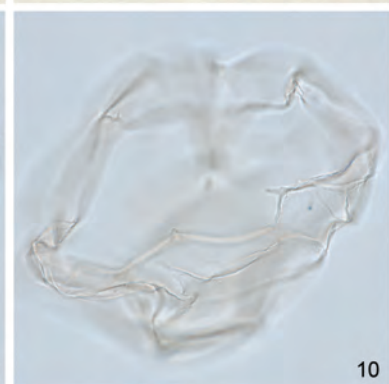
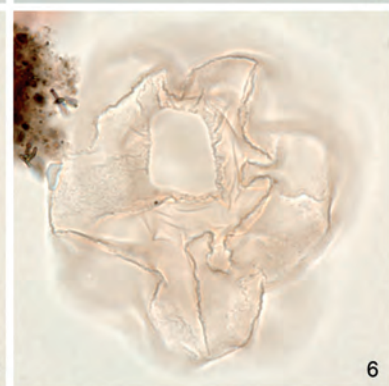
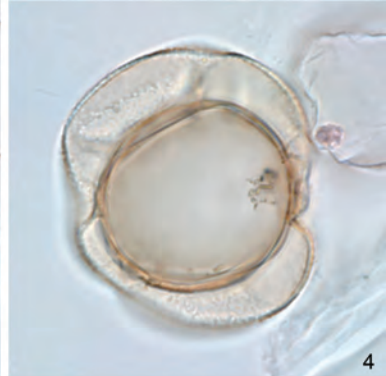
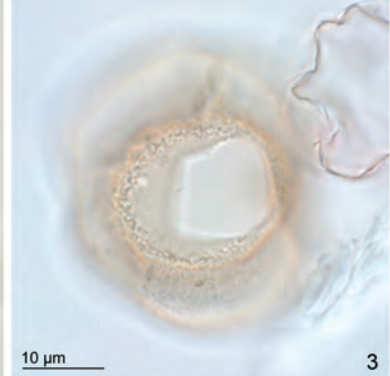
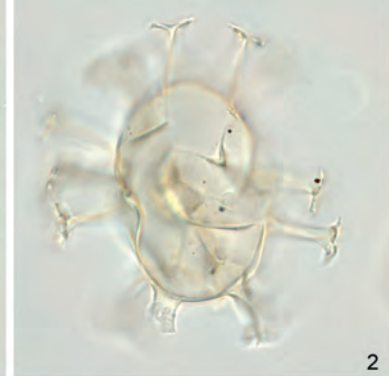
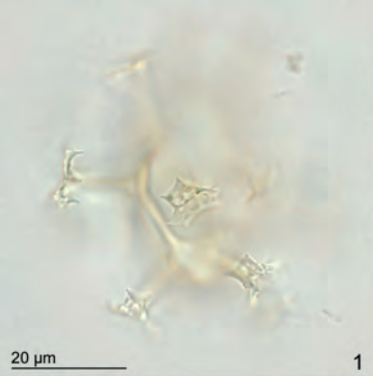


Site	Hole	Core Section	Interval (cm)	Depth (mbsf)	Calibrated ages (Ma)	Dry weight (g)	<i>Lycopodium clavatum</i> tablets added	<i>Lycopodium clavatum</i> tablets batch number	No. of <i>Lycopodium clavatum</i> grains per tablet	Standard deviation	<i>Lycopodium clavatum</i> counted	Dinoflagellate cysts counted (n)	Dinoflagellate cyst concentration (cyst/g)	Dinoflagellate cyst concentration error (cyst/g)	Shannon-Wiener diversity index	Evenness	Reworked dinoflagellate cysts counted (n)	Acritarchs counted (n)	Acritarch concentration (acritarchs/g)	Acritarch concentration error (acritarchs/g)	Terrestrial palynomorphs counted (n)	Terrestrial palynomorphs concentration (grains/g)	Error bar (grains/g)
ODP 642	B	8	14-15	64.54	2.967	9.25	1	483216	18583	1708	248	376	3046	375	2.38	0.71	21	129	1045	149	102	826	123
ODP 642	B	8	94-95	65.34	3.047	14.02	1	483216	18583	1708	824	234	376	44	2.37	0.74	134	90	145	21	624	1004	107
ODP 642	B	8	129-130	65.69	3.20-3.04	14.20	1	483216	18583	1708	454	349	1006	117	2.07	0.70	30	94	271	40	177	510	65
ODP 642	B	9	50-51	66.90	3.20	6.47	1	483216	18583	1708	182	486	7670	970	1.66	0.57	2	20	316	80	21	331	82
ODP 642	B	9	120-121	67.60	3.27	9.53	1	483216	18583	1708	441	332	1468	172	2.34	0.68	9	158	699	91	27	119	26
ODP 642	B	9	20-21	68.10	3.32	6.69	1	483216	18583	1708	208	312	4167	535	2.31	0.66		104	1389	210	132	1763	254
ODP 642	B	9	40-41	68.30	3.35	10.03	1	483216	18583	1708	169	313	3431	455	2.14	0.56	1	137	1502	221	34	373	78
ODP 642	B	9	65-66	68.55	3.39	10.39	1	483216	18583	1708	587	342	1042	119	2.23	0.64	1	42	128	24	52	158	27
ODP 642	B	9	110-111	69.00	3.45	13.93	1	483216	18583	1708	101	535	7066	1005	2.23	0.61	+	668	8823	1243	41	542	112
ODP 642	B	9	120-121	69.10	3.47	15.31	1	483216	18583	1708	40	390	11834	2246	2.32	0.68	1	312	9467	1813	51	1548	356
ODP 642	B	9	55-56	69.95	3.59	10.26	1	483216	18583	1708	110	501	8249	1153	2.12	0.57	+	516	8496	1186	101	1663	275
ODP 642	B	9	60-61	70.00	3.60	9.63	1	483216	18583	1708	88	313	6864	1041	2.12	0.55	1	905	19845	2870	76	1667	303
ODP 642	B	9	70-71	70.10	3.61	10.17	1	483216	18583	1708	156	320	3748	503	2.51	0.68		659	7719	988	58	679	122
ODP 642	B	9	100-101	70.40	3.64	9.87	1	483216	18583	1708	161	435	5087	662	2.05	0.61	1	127	1485	223	20	234	59
ODP 642	B	9	105-106	70.45	3.64	8.54	1	483216	18583	1708	233	470	4389	535	2.22	0.65	3	113	1055	155	26	243	55
ODP 642	B	9	145-146	72.35	3.83	18.76	1	483216	18583	1708	151	446	2926	385	2.69	0.68	3	1596	10470	1312	176	1155	166
ODP 642	B	9	100-101	73.40	3.93	7.40	1	483216	18583	1708	82	425	13015	1974	2.01	0.57	+	1249	38250	5601	112	3430	590
ODP 642	B	9	100-101	74.90	4.08	16.32	1	483216	18583	1708	76	279	4180	663	2.34	0.69	2	2432	36437	5407	104	1558	275
ODP 642	B	9	11-12	75.51	4.14	5.98	1	483216	18583	1708	220	408	5763	716	2.24	0.65	4	1881	26569	3090	200	2825	379
ODP 642	B	10	40-41	76.30	4.23	10.89	1	483216	18583	1708	159	306	3284	441	2.12	0.57	1	1184	12707	1586	80	859	142
ODP 642	B	10	145-146	77.35	4.49	6.74	1	483216	18583	1708	52	324	17179	3013	2.38	0.71	2	1	53	54	10	530	189
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ODP 642	B	10	100-101	78.40	4.69	11.92	1	483216	18583	1708	9	389	67382	23548	2.27	0.60		733	126970	44153	47	8141	3055
ODP 642	B	10	102-103	79.92	4.81	11.84	1	483216	18583	1708	25	432	27121	6111	2.53	0.71	1	425	26682	6014	53	3327	863
ODP 642	B	10	15-16	80.55	4.84	10.15	1	483216	18583	1708	111	560	9237	1281	2.33	0.65	+	709	11694	1606	66	1089	197
ODP 642	B	10	11-12	82.01	4.91	7.83	1	483216	18583	1708	144	324	5340	726	2.11	0.62		1441	23750	3012	140	2307	346
ODP 642	B	10	115-116	83.05	4.98	12.52	1	483216	18583	1708	16	452	41930	11341	2.03	0.56	1	262	24305	6646	67	6215	1821
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ODP 642	B	11	65-66	89.05	5.53	7.65	1	483216	18583	1708	174	484	6757	862	2.55	0.67	4	378	5277	685	103	1438	222
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ODP 642	B	11	15-16	94.55	5.87	6.86	1	483216	18583	1708	94	513	14784	2144	2.44	0.66	+	448	12910	1885	92	2651	459
ODP 642	B	11	55-56	94.95	5.89	10.83	1	483216	18583	1708	64	435	11663	1894	2.34	0.63	+	297	7963	1319	59	1582	320

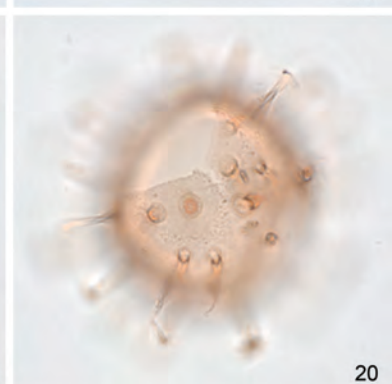
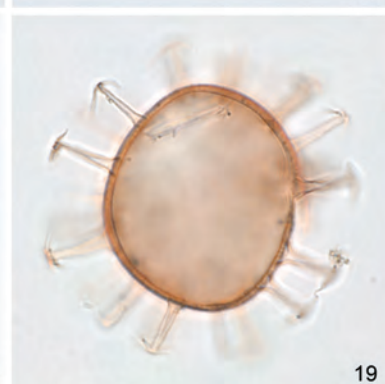
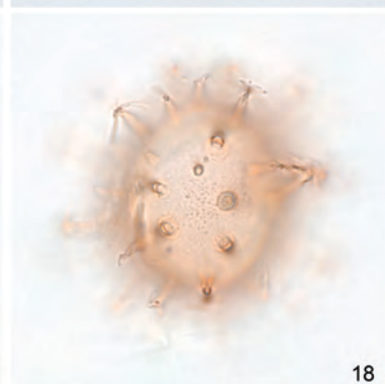
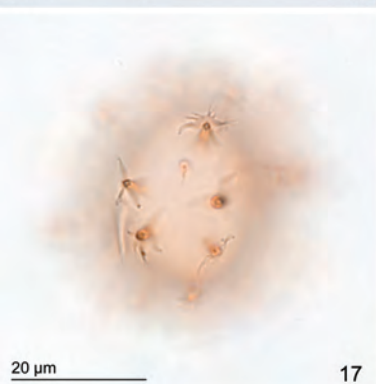
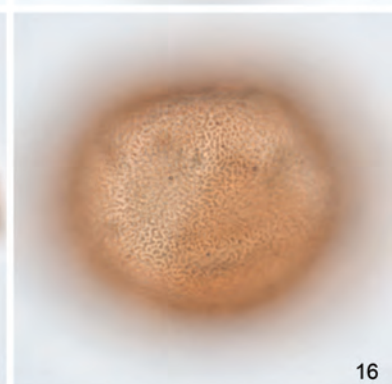
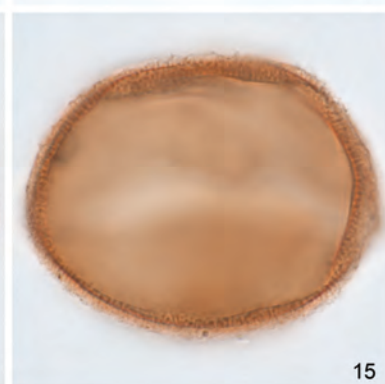
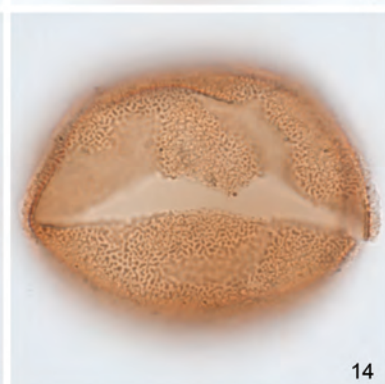
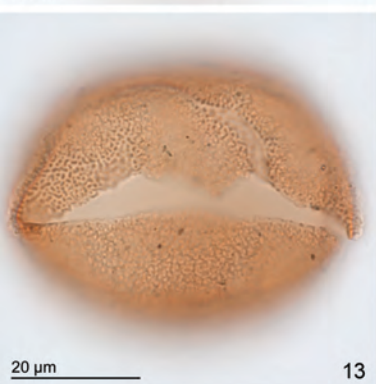
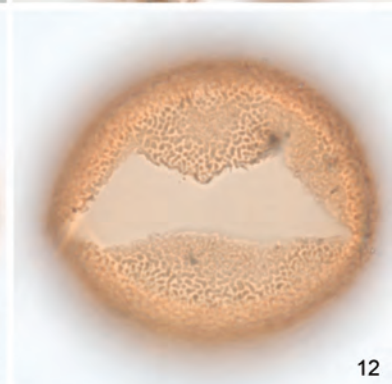
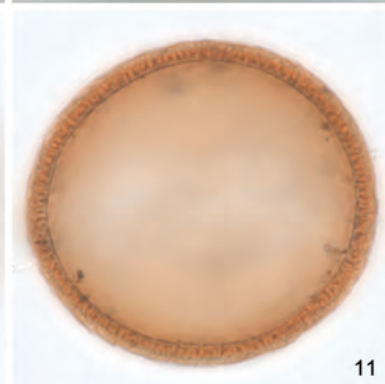
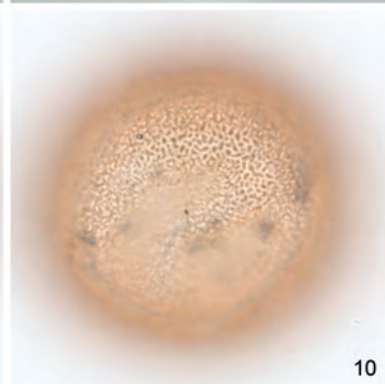
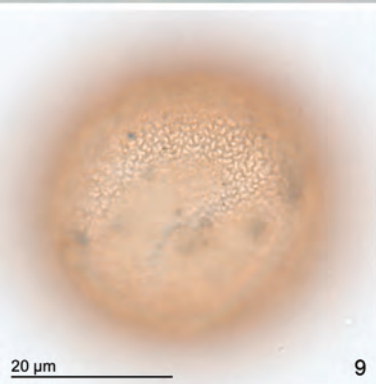
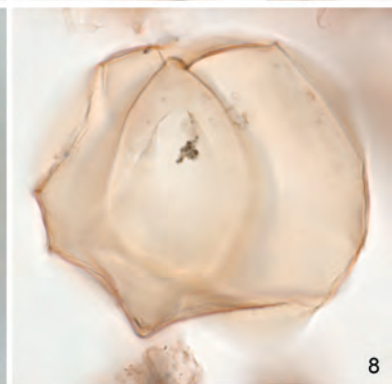
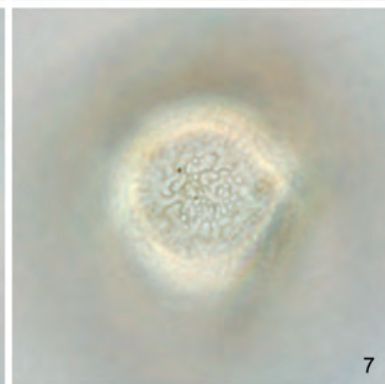
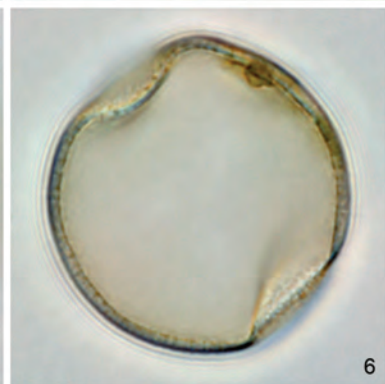
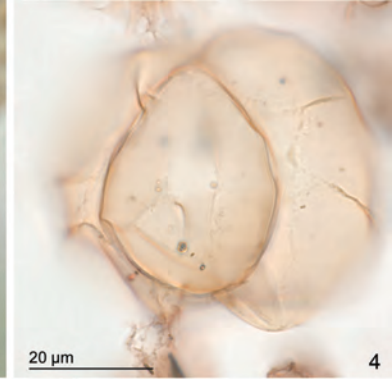
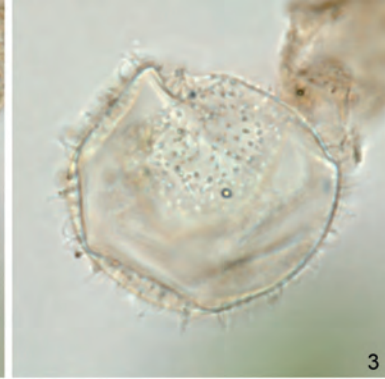
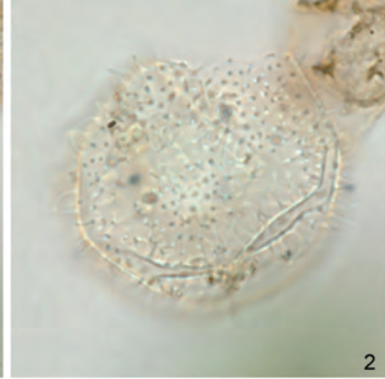
Chron/Subchron (Beil, 1989)	Chron/Subchron (GTS 2012)	Chron boundary name	Upper boundary		Lower boundary		Mid-depth (mbsf)	Age (Ma) in Beil (1989)	Age (Ma) updated to GTS 2012
			Sample	Depth (mbsf)	Sample	Depth (mbsf)			
C2AN-2/C2AR-2	C2An.2n/ C2An.2r	Top Mammoth	9H-1, 41	66.81	9H-1, 71	67.11	66.96	3.08	3.207
C2AR-2/C2AN-3	C2An.2r / C2An.3n	Base Mammoth	9H-2, 11	68.01	9H-2, 41	68.31	68.16	3.18	3.330
C2AN-3/C2AR-3	C2An.3n/C2Ar	Top Gilbert	9H-3, 47	69.87	9H-3,71	70.11	69.99	3.40	3.596
C2AR-3/C3N-1	C2Ar/C3n.1n	Top Cochiti	9H-7, 56	75.96	10H-1, 9	75.99	75.98	3.88	4.187
C3N-1/C3R-1	C3n.1n/C3n.1r	Base Cochiti	10H-1, 70	76.60	10H-1, 101	76.91	76.76	3.97	4.300
C3R-1/C3N-2	C3n.1r/C3n.2n	Top Nunivak	10H-1, 130	77.20	10H-2, 11	77.51	77.36	4.10	4.493
C3N-2/C3R-2	C3n.2n/C3n.2r	Base Nunivak	10H-2, 11	77.51	10H-2, 41	77.81	77.66	4.24	4.631
C3R-2/C3N-3	C3n.2r/C3n.3n	Top Sidufjall	10H-3, 71	79.61	10H-3, 100	79.90	79.76	4.40	4.799
C3N-3/C3R-3	C3n.3n/C3N.3r	Base Sidufjall	10H-4, 126	81.66	10H-5, 11	82.01	81.84	4.47	4.896
C3R-3/C3N-4	C3N.3r/C3n.4n	Top Thvera	10H-5, 130	83.2	10H-6, 10	83.5	83.35	4.57	4.997
C3N-4/C3R-4(?)	C3n.4n/C3r	Base Thvera	10H-6, 71	84.11	10H-6, 100	84.4	84.255	4.77	5.235
C3R-4 (?)(C3A) C3AN-1	C3R/C3An.1n	Top Gilbert	12H-2, 71	97.11	12H-2, 101	97.41	97.26	5.35	6.033

Event	Species	Sample	Depth (mbsf)	Age (Ma)	error (Ma)
HO	<i>Corrudinium harlandii</i>	642B-9H1, 120–121 cm	67.60	3.27	0.07
HO	<i>Corrudinium? labradori</i>	642B-9H1, 120–121 cm	67.60	3.27	0.07
HO	<i>Heteraulacacysta</i> sp. A of Costa and Downie (1979)	642B-9H1, 120–121 cm	67.60	3.27	0.07
HO	<i>Melitasphaeridium choanophorum</i>	642B-9H1, 120–121 cm	67.60	3.27	0.07
HO	<i>Operculodinium janduchenei</i>	642B-9H2, 20–21 cm	68.10	3.32	0.05
HO	<i>Ataxiodinium confusum</i>	642B-9H2, 65–66 cm	68.55	3.39	0.04
HO	<i>Impagidinium solidum</i>	642B-9H3, 55–56 cm	68.55	3.59	0.12
HO	<i>Melitasphaeridium</i> sp. A of Head and De Schepper (2008)	642B-9H2, 110–11 cm	69.00	3.45	0.07
HO	<i>Operculodinium? eirikianum</i> var. <i>crebrum</i>	642B-9H2, 110–11 cm	69.00	3.45	0.07
HO	<i>Corrudinium devernaliae</i>	642B-10H1, 145–146 cm	77.35	4.49	0.26
HO	<i>Operculodinium tegillatum</i>	642B-10H1, 145–146 cm	77.35	4.49	0.26
HO	<i>Batiacasphaera micropapillata</i> complex	642B-10H2, 40–41 cm	77.80	4.64	0.15
HO	Cyst type I of de Vernal and Mudie (1989)	642B-10H2, 40–41 cm	77.80	4.64	0.15
HO	<i>Reticulatosphaera actinocoronata</i>	642B-10H2, 40–41 cm	77.80	4.64	0.15
HO	<i>Selenopemphix brevispinosa</i>	642B-10H2, 40–41 cm	77.80	4.64	0.15
HO	<i>Lavradosphaera lucifer*</i>	642B-10H2, 100–101 cm	78.40	4.69	0.05
HO	<i>Cristadinium diminutivum</i>	642B-10H5, 11–12 cm	82.01	4.91	0.07
HO	<i>Batiacasphaera hirsuta</i>	642B-10H5, 115–116 cm	83.05	4.98	0.07
HO	" <i>Impagidinium densiverrucosum</i> " of Zevenboom and Santarelli in Zevenboom (1995)	642B-10H7, 30–31 cm	85.20	5.29	0.11
HO	<i>Pyxidinospis vesiculata</i>	642B-10H7, 30–31 cm	85.20	5.29	0.11
HO	" <i>Veriplicidium franklinii</i> " of Anstey (1992)*	642B-10H7, 30–31 cm	85.20	5.29	0.11
HO	<i>Cerebrocysta poulsenii</i>	642B-11H1, 80–81 cm	86.20	5.35	0.06
HO	<i>Selenopemphix armageddonensis</i>	642B-11H2, 15–16 cm	87.05	5.41	0.06
HPO	<i>Trinovantedinium glorianum</i>	642B-10H7, 30–31 cm	85.20	5.29	0.11
HPO/HCO	<i>Batiacasphaera hirsuta</i>	642B-11H3, 65–66 cm	89.05	5.53	0.09
HCO	<i>Lavradosphaera crista*</i>	642B-9H1, 120–121 cm	67.60	3.27	0.07
LO	<i>Melitasphaeridium</i> sp. A of Head and De Schepper (2008)	642B-9H3, 70–71 cm	70.10	3.61	0.03
LO	<i>Impagidinium solidum</i>	642B-9H4, 145–146 cm	72.35	3.83	0.10
LO	<i>Operculodinium? eirikianum</i> var. <i>crebrum</i>	642B-10H1, 40–41 cm	76.30	4.23	0.26
LO	<i>Ataxiodinium confusum</i>	642B-10H1, 145–146 cm	77.35	4.49	0.15
LO	Cyst type I of de Vernal and Mudie (1989)	642B-10H5, 11–12 cm	82.01	4.91	0.07
LO	<i>Filisphaera filifera</i> subsp. <i>filifera</i>	642B-10H6, 65–66 cm	84.05	5.18	0.11
LO	<i>Bitectatodinium tepikiense</i>	642B-10H6, 65–66 cm	84.05	5.18	0.11
LO	<i>Heteraulacacysta</i> sp. A of Costa and Downie (1979)	642B-10H7, 30–31 cm	85.20	5.29	0.06
LO	<i>Operculodinium tegillatum</i>	642B-10H7, 30–31 cm	85.20	5.29	0.06
LO	<i>Pyxidinospis vesiculata</i>	642B-10H7, 30–31 cm	85.20	5.29	0.06
LO	<i>Corrudinium devernaliae</i>	642B-11H3, 65–66 cm	89.05	5.53	0.10

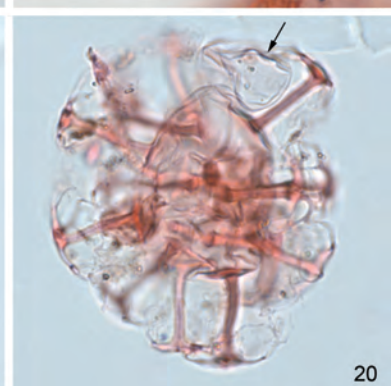
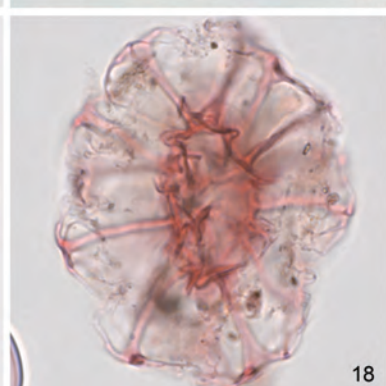
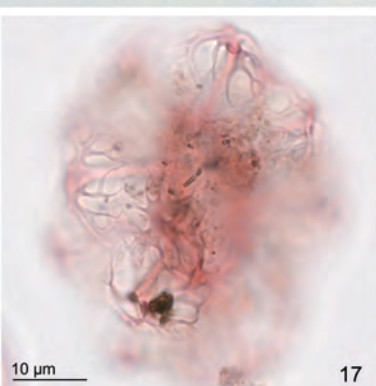
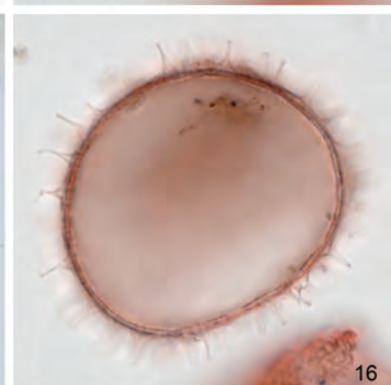
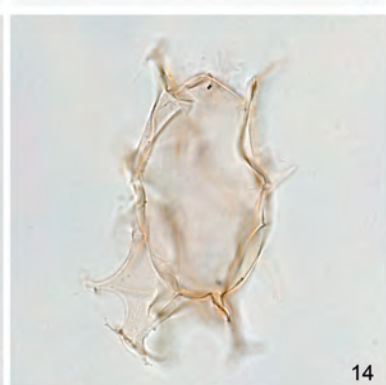
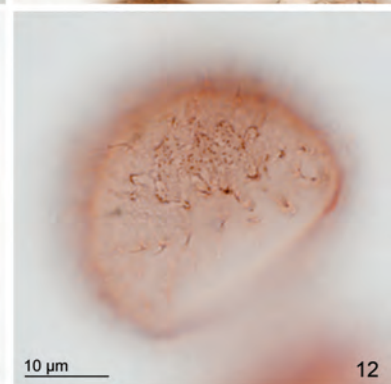
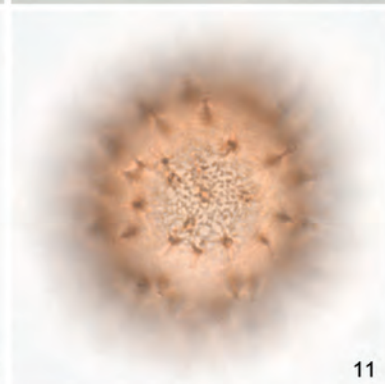
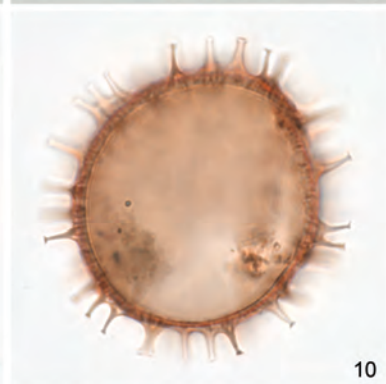
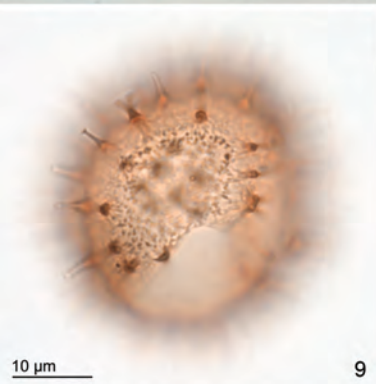
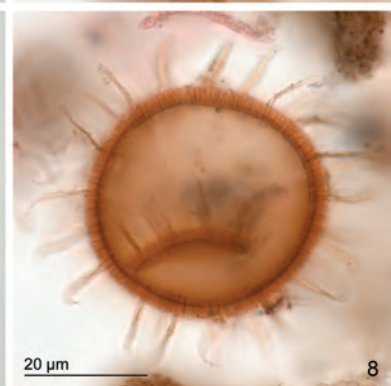
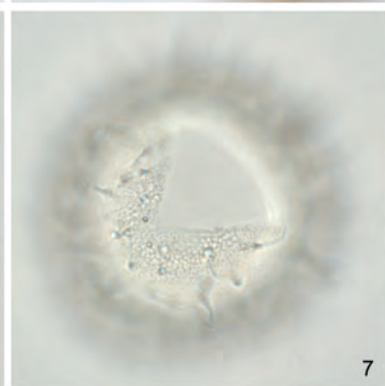
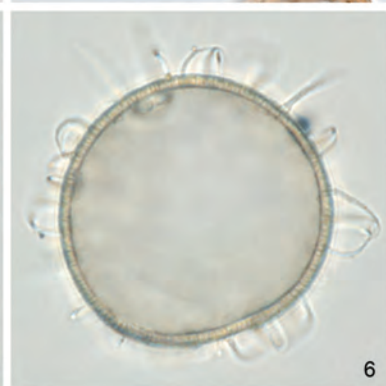
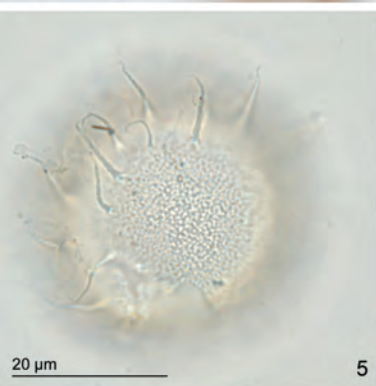
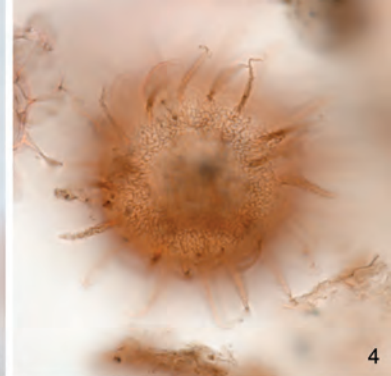
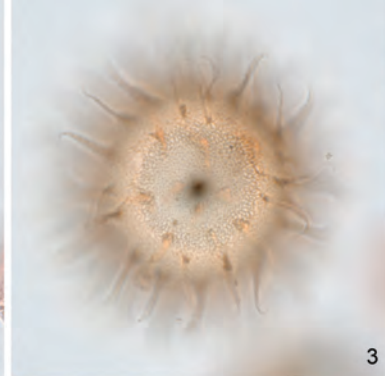
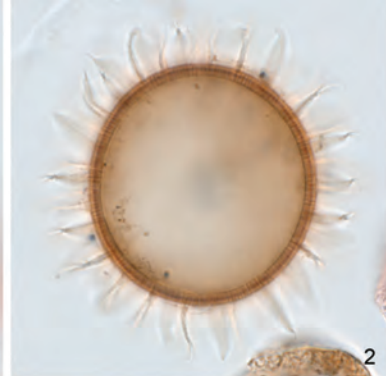
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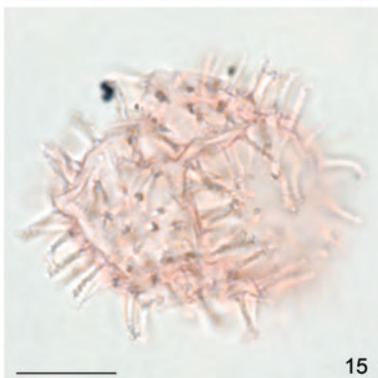
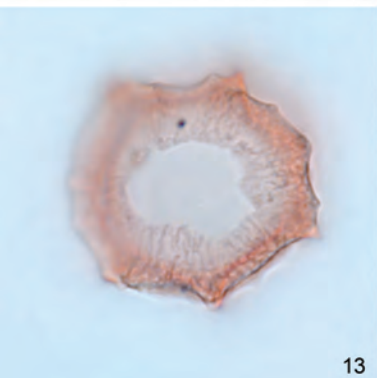
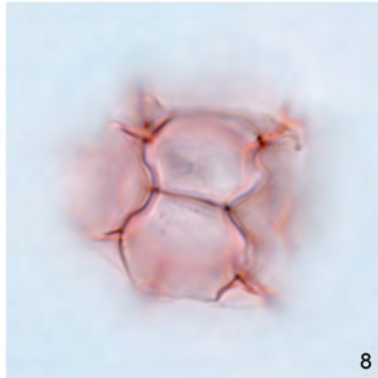
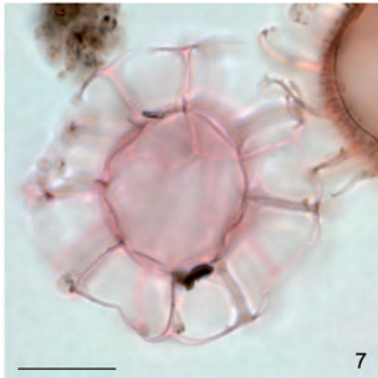
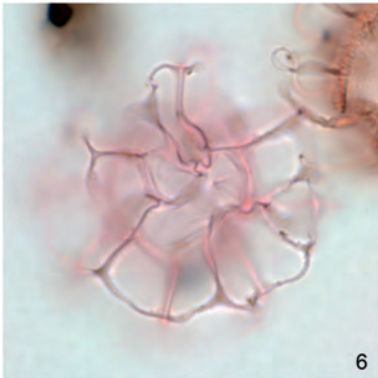
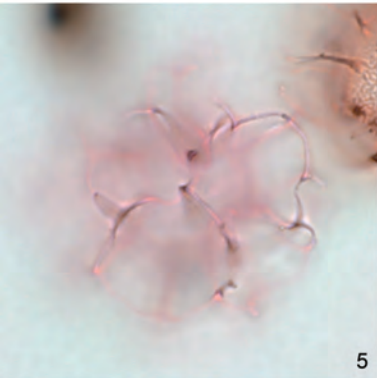
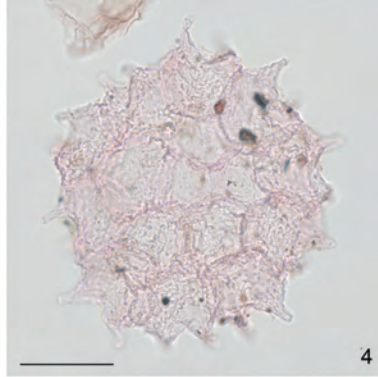




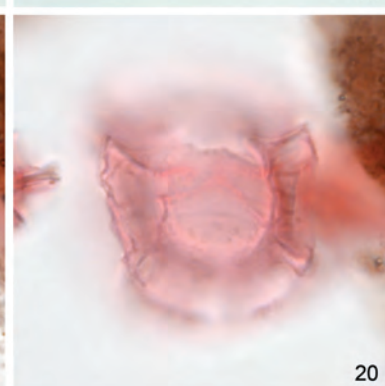
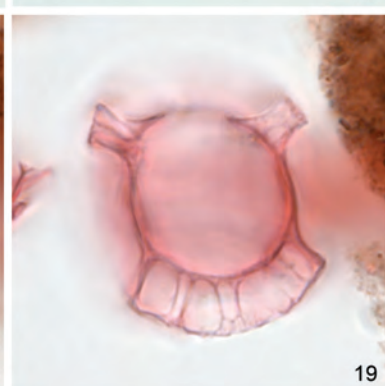
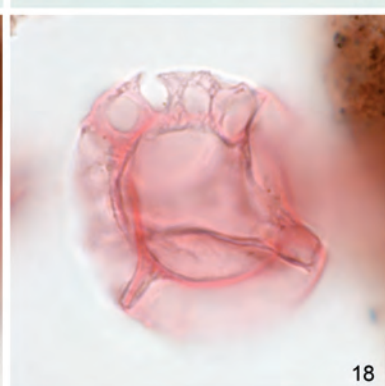
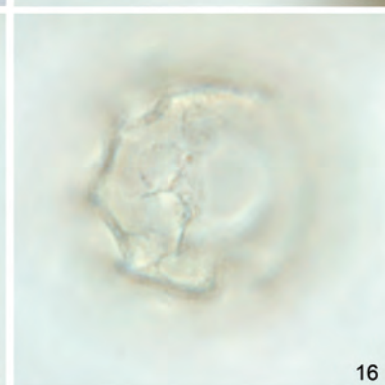
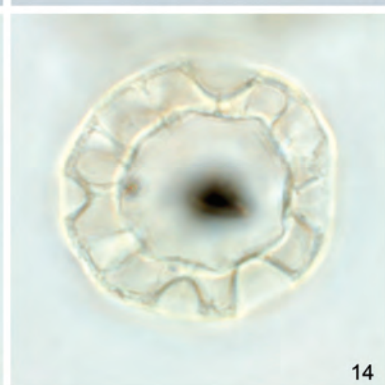
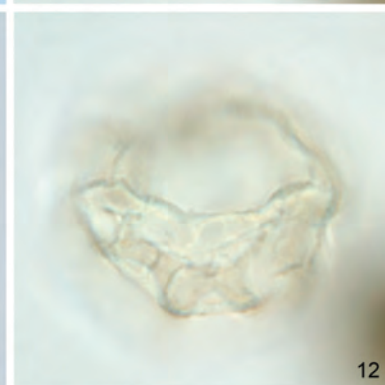
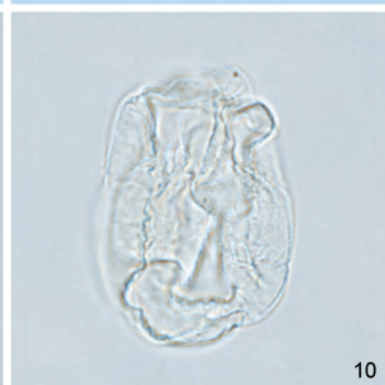
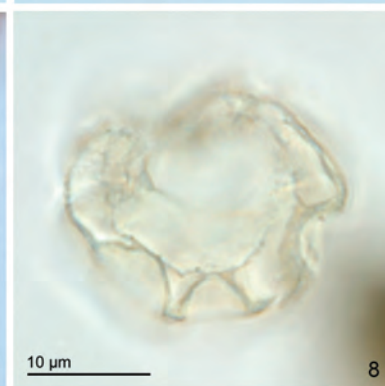
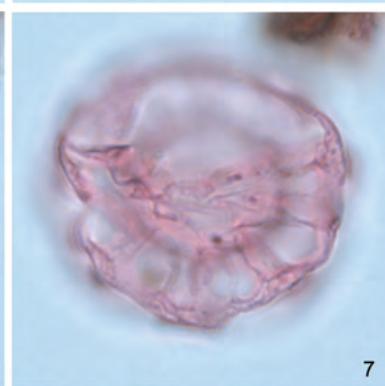
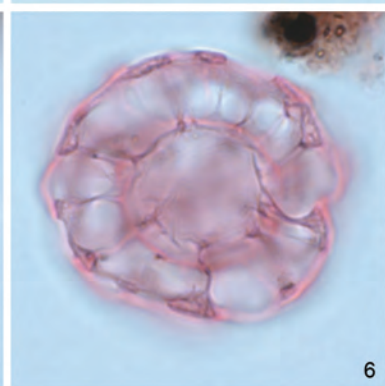
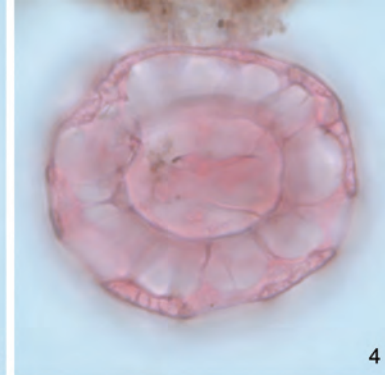
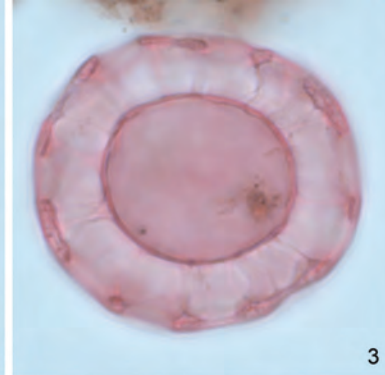
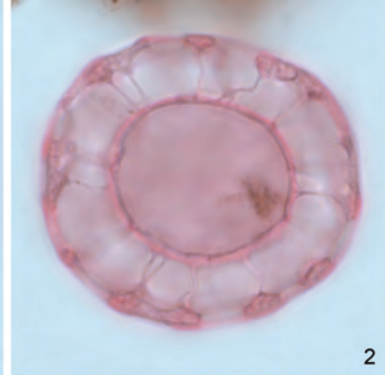
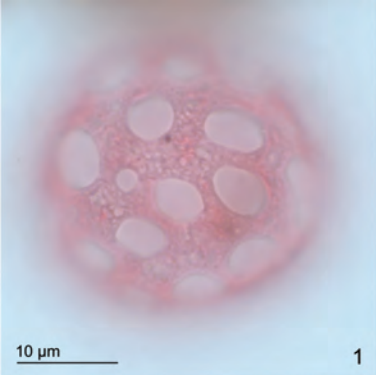














Site	Hole	Core	Section	Interval (cm)	Depth (mbsf)	Calibrated ages (Ma)	Cymatiosphaera? invaginata	Small spiny acritarchs	Nannobarrhophora walldalei	Cymatiosphaera? aegirii	Cymatiosphaera? icenorum	Acritarchs spp. indet.	Cymatiosphaera spp. indet.	Lavradosphaera crista	Cymatiosphaera? fensomei	Lavradosphaera sp. cf. canalis	Lavradosphaera spp. indet.	Lavradosphaera lucifer	Acritarch sp. 1 of Head, Norris & Mudie (1989)	"Veriplicidium franklinii" in Anstey (1992)	Leiosphaeridia rockhallengensis?	Cyclopsiella granosa	Lavradosphaera canalis	Lavradosphaera sp. 2 of Schreck et al. (2013)	Biozones	
ODP 642	B	8	6	14–15	64.54	2.96	30	61	19	6	12			1												Not zoned
ODP 642	B	8	6	94–95	65.34	3.04	7	79	4																	
ODP 642	B	8	6	129–130	65.69	3.20–3.04	10	17	53	3	1	6	4													
ODP 642	B	9	1	50–51	66.90	3.20	1	17	2																	
ODP 642	B	9	1	120–121	67.60	3.27	16	87	16	2	17		12	8												VP3
ODP 642	B	9	2	20–21	68.10	3.32	23	58	1		4	1		16	1											
ODP 642	B	9	2	40–41	68.30	3.35	11	68	17		23	6		12												
ODP 642	B	9	2	65–66	68.55	3.39	+	35	1	2	1			3	+											
ODP 642	B	9	2	110–111	69.00	3.45	101	169	87	14	244			53												
ODP 642	B	9	2	120–121	69.10	3.47	60	50	57	11	93			41												
ODP 642	B	9	3	55–56	69.95	3.59	160	92	11	8	126			116		3										
ODP 642	B	9	3	60–61	70.00	3.60	277	87	15	28	256	13	1	228												
ODP 642	B	9	3	70–71	70.10	3.61	220	94	10	25	188			119	3											
ODP 642	B	9	3	100–101	70.40	3.64	4	81	38		3			1												
ODP 642	B	9	3	105–106	70.45	3.64		65	48	+	+															
ODP 642	B	9	4	145–146	72.35	3.83	128	492	20	19	768	8	2	158		+	1									
ODP 642	B	9	5	100–101	73.40	3.93	152	352	19	3	605			118												
ODP 642	B	9	6	100–101	74.90	4.08	590	995	27	18	459			341			2									
ODP 642	B	9	7	11–12	75.51	4.14	232	541	48	18	811			231												
ODP 642	B	10	1	40–41	76.30	4.23	184	494	31	7	245			223												
ODP 642	B	10	1	145–146	77.35	4.49	1																			
ODP 642	B	10	2	40–41	77.80	4.64	453	180	3		8			7												
ODP 642	B	10	2	100–101	78.40	4.69	469	118	20		34	10		4			3	3								
ODP 642	B	10	3	102–103	79.92	4.81	216	144	12		25	3		13			3	6		+						
ODP 642	B	10	4	15–16	80.55	4.84	338	305	16		16		2	21				11								
ODP 642	B	10	5	11–12	82.01	4.91	1234	89	1		21	3	6	3	1			81	1							
ODP 642	B	10	5	115–116	83.05	4.98	144	37	8	1	1	4		26				41								
ODP 642	B	10	6	65–66	84.05	5.18	242	21	2			2	4	2				47								
ODP 642	B	10	7	30–31	85.20	5.29	49	14	1		2			12	1		1	6		7	1					
ODP 642	B	11	1	80–81	86.20	5.35	100	95				+		8			3			59						
ODP 642	B	11	2	15–16	87.05	5.41	139	145	1	1	1			10						17						
ODP 642	B	11	2	65–66	87.55	5.44	19	132		2		7					3			23						
ODP 642	B	11	3	65–66	89.05	5.53	128	196	15	1	+			12					5	21						
ODP 642	B	11	4	70–71	90.60	5.62	48	123	1	1		+		6			+			13						
ODP 642	B	11	5	60–61	92.00	5.71	68	121				+	+	2			1			68		1				
ODP 642	B	11	6	20–21	93.10	5.78	205	166				3		4			6			13		+	10			
ODP 642	B	11	6	70–71	93.60	5.81	81	74						4			1			7			9			
ODP 642	B	11	6	110–111	94.00	5.83	104	169	19								3	7	4	4			1			
ODP 642	B	11	7	15–16	94.55	5.87	68	320				9		2			4	8	3	32			2			
ODP 642	B	11	7	55–56	94.95	5.89	30	235				7		7			1			16				1		