

DINOFLAGELLATE CYST DISTRIBUTION DURING THE MIDDLE EOCENE IN THE DRAKE PASSAGE AREA: PALEOCEANOGRAPHIC IMPLICATIONS



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Abstract. The middle–late Eocene (45–36 Ma) dinoflagellate cyst distribution in high latitudes of the Southern Hemisphere has been explained by a surface ocean circulation pattern characterized by extensive subpolar gyres around Antarctica. Based on global paleoclimate models it has been proposed that these surface ocean gyres might favored the development of an Antarctic–endemic dinoflagellate cyst assemblage, which was consistently abundant until the end of the Eocene. During the late Eocene the deepening of the Tasman Gateway and the Drake Passage would have generated a circumpolar water flow responsible for the disruption of the local gyre system and the subsequent extinction of the endemic assemblage. Some authors have recently suggested that during the middle Eocene shallow water flows had already developed through incipient openings of the Tasman Gateway and the Drake Passage. In this review we have compared the middle Eocene dinoflagellate cysts assemblages dominated by the Antarctic–endemic species from localities of the Drake Passage area and performed a multivariate analysis to evaluate this hypothesis. Our results point out a clear differentiation between localities to the north and south of what now is the Drake Passage. While localities from the Antarctic Peninsula and Scotia Sea would have been affected by Antarctic surface waters, the Austral Basin would have been flooded by Antarctic waters together with Pacific waterflows developed through an incipient Drake Passage during the middle Eocene.

Key words. Dinoflagellate cysts. Middle Eocene. Drake Passage. Early opening.

Resumen. DISTRIBUCIÓN DE QUIESTES DE DINOFLAGELADOS DURANTE EL EOCENO MEDIO EN EL ÁREA DEL PASAJE DE DRAKE: IMPLICANCIAS PALEOCEANOGRÁFICAS. La distribución de quistes de dinoflagelados del Eoceno medio–tardío (45–36 Ma) en altas latitudes del Hemisferio Sur ha sido explicada por un patrón de circulación oceánica superficial caracterizado por amplios giros subpolares alrededor de Antártida. Sobre la base de modelos paleoclimáticos globales se ha propuesto que dichos giros podrían haber favorecido el desarrollo de una asociación de quistes de dinoflagelados endémica–Antártica, la que habría prevalecido hasta el final del Eocene. Durante el Oligoceno la profundización del Conducto de Tasmania y el Pasaje de Drake habrían generado un flujo de agua circumpolar responsable de la disruptión del sistema de giros y de la subseciente extinción de la asociación endémica. Recientemente algunos autores han sugerido que durante el Eocene medio ya se habrían desarrollado flujos de aguas poco profundas a través de aperturas incipientes del Conducto de Tasmania y el Pasaje de Drake. En este trabajo se compararon las asociaciones de quistes de dinoflagelados dominadas por la asociación endémica Antártica de localidades del área del Pasaje de Drake y se realizó un análisis multivariado para evaluar esta hipótesis. Los resultados obtenidos señalan una clara separación entre las localidades al norte y al sur de lo que hoy es el Pasaje de Drake. Mientras que la Península Antártica y el Mar de Scotia habrían sido afectadas por aguas superficiales originadas en Antártida, la Cuenca Austral habría sido inundada por aguas antárticas junto con flujos de agua del Pacífico desarrollados a través de un incipiente Pasaje de Drake durante el Eocene medio.

Palabras clave. Quistes de dinoflagelados. Eoceno medio. Pasaje de Drake. Apertura temprana.

FOSSIL organic-walled dinoflagellate cysts are abundant in continental shelf sediments and their spatial distribution depends on the physico-chemical characteristics of the water-masses in which the dinoflagellates developed, and on past ocean circulation patterns (Sluijs *et al.*, 2005). During the Paleogene, dinoflagellate cyst assemblages in the Southern Ocean evidenced major modifications as a consequence of the climatic and tectonic changes that occurred during this period (Stickley *et al.*, 2004; Guerstein *et al.*, 2010; Bijl *et al.*, 2013a).

Paleocene and early Eocene circum-Antarctic assemblages were widely dominated by cosmopolitan early Paleogene taxa and endemic species were scarce (Bijl *et al.*, 2011, 2013a, b). At the beginning of the middle Eocene (~50 Ma) dinoflagellate cyst assemblages began to experience a significant change, resulting in the dominance of recorded endemic taxa in the Southern south of 45°S (Bijl *et al.*, 2011). This assemblage, originally called “Transantarctic Flora” by Wrenn and Beckmann (1982), was dominated by endemic and bipolar

taxa differing from species with cosmopolitan and tropical distributions (Lentin and Williams, 1980; Wrenn and Hart, 1988; Bijl *et al.*, 2011, 2013a). As proposed by Huber *et al.* (2004), the middle–late Eocene dinoflagellate cyst distribution reflected an ocean-circulation scheme with broad clockwise gyres surrounding Antarctica. The Antarctic-endemic assemblage prevailed until the Eocene/ Oligocene transition, when the endemic species were replaced by Oligocene cosmopolitan taxa (Houben *et al.*, 2013). The extinction of the endemic assemblage may have occurred as a result of the deepening of the Tasmanian Gateway and the Drake Passage (Sluijs *et al.*, 2003; Stickley *et al.*, 2004; Guerstein *et al.*, 2008a; Houben *et al.*, 2011, 2013). Such tectonic changes and the subsequent development of an unrestricted circum-polar waterflow during the early Oligocene would have disrupted the subpolar gyres and altered the environmental conditions that favored dinoflagellate endemism (Huber *et al.*, 2004).

Bijl *et al.* (2013a) have recently calibrated dinoflagellate cyst assemblages from sediment cores drilled on the Tasman shelves with the Geomagnetic Polarity TimeScale (GPTS) of Vandenberghe *et al.* (2012). These authors also proposed a high-resolution Southern Pacific Ocean dinoflagellate cyst zonation for the late Palaeocene to the late Eocene (58–36 Ma) and confirmed that the Antarctic-endemic dinoflagellate cyst assemblage was dominant during the mid to late Eocene (~ 45–39 Ma). Thus, the mid-Paleogene dinoflagellate cyst

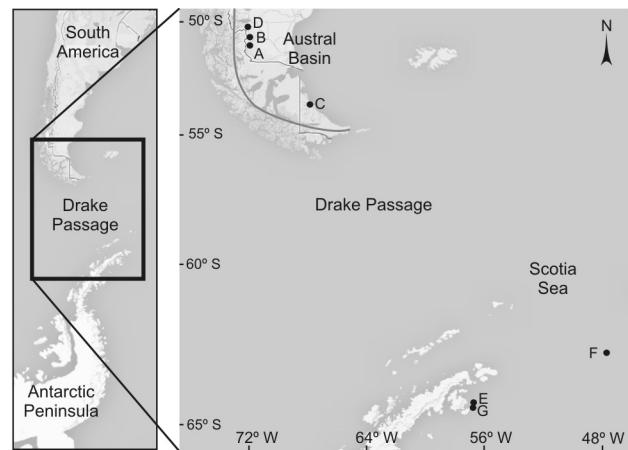


Figure 1. South America and Antarctic Peninsula map, indicating present-day location of the studied sections: **A**, Upper member Río Turbio Fm. (González Estebenet, 2014); **B**, Upper member Río Turbio Fm. (Archangelsky, 1969; Guerstein *et al.*, 2014); **C**, Leticia Fm. in Río de la Turba section (Guerstein *et al.*, 2008b); **D**, Man Aike Fm. (Guerstein *et al.*, 2014); **E**, La Meseta Fm. (Cocozza and Clarke, 1992); **F**, Bruce Bank (Mao and Mohr, 1995); **G**, La Meseta Fm. (Wrenn and Hart, 1988). From Google Maps (<https://maps.google.com.ar>).

stratigraphic distribution provides a correlation tool that can be applied to other sites of the Southern Ocean.

In recent years, interest has grown in the progressive opening of the Southern Ocean Gateways. Bijl *et al.* (2013b) suggested an incipient opening of the Tasman Gateway with a consequent onset of surface water throughflow starting at about 50–49 Ma during the Ypresian (early Eocene). Regarding the Drake Passage, the time of its opening and

TABLE 1. Published works considered in this study.

Lithostratigraphy	Location	Latitude and Longitude	Sources
A Upper member of the Río Turbio Fm.	Santa Cruz, Austral Basin	51°31'13"S, 72°15'11"W	González Estebenet <i>et al.</i> , 2014
B Upper member of the Río Turbio Fm.	Santa Cruz, Austral Basin	three integrated cores 50°45'14"S, 72°01'39"W; 50°56'21"S, 72°02'08"W; 50°35'59"S, 72°13'40"W	Archangelsky, 1969 Guerstein <i>et al.</i> , 2014
C Leticia Fm.	Río de la Turba section. Tierra del Fuego, Austral Basin	53°58'S, 68°28'W	Guerstein <i>et al.</i> , 2008b
D Man Aike Fm.	Santa Cruz, Austral Basin	50°21'45"S, 72°14'30"W	Guerstein <i>et al.</i> , 2014
E La Meseta Fm. Associations 2 and 3 (between 43 and 165 m)	Cape Wiman, Seymour Island, Antarctic Peninsula	64°13'5"S, 56°38'W	Cocozza and Clarke, 1992
F Core IO1578-59 in Bruce Bank	Scotia Sea, Antarctica	60°33.6'S, 40°13.2'W	Mao and Mohr, 1995
G La Meseta Fm. Section 3 (between 47 and 188 m)	Seymour Island, Antarctic Peninsula	64°14'S, 56°49'W*	Wrenn and Hart, 1988 Douglas <i>et al.</i> , 2014

*Geographical coordinates from Google Earth.

TABLE 2. Taxonomic list of species of dinoflagellate cysts present in the localities A-G (see Table 1 for abbreviations).

Species	Bioevents (Ma)	A	B	C	D	E	F	G
Deflandrea antarctica Wilson, 1967a *	FO: 58.20 (± 0.30)	x	x	x	x	x	x	x
Enneadocysta dictyostila (Menéndez) Fensome et al., 2006 *	FCO: 45.20 (± 0.20); LO: 33.40 (± 0.30)	x	x	x	x	x	x	x
Spinidinium spp. *		x	x	x	x	x	x	x
Enneadocysta brevistila Fensome et al., 2006 *		x	x	x	x	x		
Arachnodynium antarcticum Wilson and Clowes, 1982 *	FO: 48.70 (± 0.20); LO: 36 (± 0.3)	x	x	x		x	x	x
Vozzhennikovia apertura Wilson, 1967a *	FO: 65.10 (± 0.40)	x	x	x		x	x	x
Deflandrea cygniformis Pôthe de Baldis, 1966 *					x		x	x
Octodinium askiniae Wrenn and Hart, 1988 *	FO: 53.90 (± 0.10)					x	x	x
Thalassiphora pelagica (Eisenack) Eisenack and Gocht, 1960	FO: 55.30 (± 0.20)	x	x	x	x	x		
Selenopemphix spp.	FO: 49.50 (± 0.30)	x	x	x	x			x
Hystrichosphaeridium truswelliae Wrenn and Hart, 1988	LO: 36 (± 0.3)	x	x	x	x			x
Lejeunecysta spp.		x	x	x	x			x
Operculodinium centrocarpum (Deflandre and Cookson, 1955) Wall, 1967		x	x	x	x			x
Turbiosphaera filosa Wilson, 1967a		x	x	x				
Achomosphaera sp. A		x			x			
Hystrichosphaeridium tubiferum (Ehrenberg, 1838) Davey and Williams, 1966				x			x	
Pyxidinopsis delicata Wilson, 1988				x			x	
Palaeocystodinium sp.				x				x
Alterbidinium sp.		x			x	x	x	
Impletosphaeridium clavus Wrenn and Hart, 1988				x	x	x		
Phthanoperidinium stockmansii (de Coninck) Lentini and Williams, 1977				x	x	x		
Impagidinium spp.		x			x	x	x	
Cordosphaeridium minimum (Morgenroth, 1966a) Benedek, 1972				x	x			
Paucisphaeridium inversibuccinum (Davey and Williams, 1966) Bujak et al., 1980				x	x			
Deflandrea spp.		x			x	x	x	
Operculodinium bergmannii (Archangelsky, 1969) Stover and Evitt, 1978				x	x	x		

Species classified as endemic or bipolar according to Bijl et al. (2011, 2013a) and Houben et al. (2013) are indicated with asterisk (*). Magnetostratigraphically calibrated ages of key species refer to: FO, first occurrences; FCO, first common occurrences and LO, last occurrences.

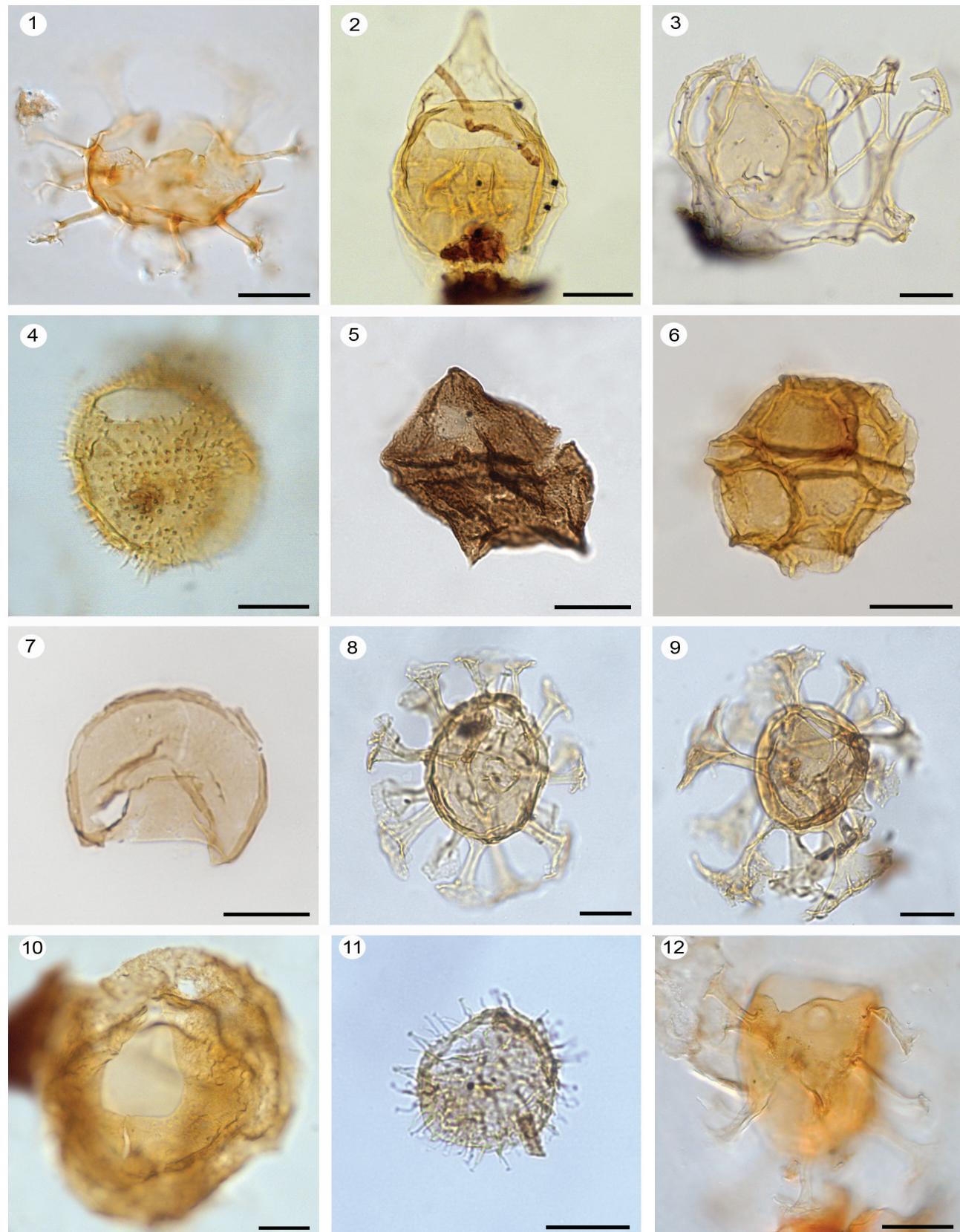
deepening is still under discussion, with ages ranging from the early–middle Eocene (Livermore et al., 2005, 2007; Scher and Martin, 2006) to the early Miocene (Barker, 2001). Scher and Martin (2006), Livermore et al. (2007) and Lagabrielle et al. (2009) suggested an early opening for this passage, with shallow waterflows from the Pacific to the Atlantic Oceans. The aim of this review is to compare the middle Eocene dinoflagellate cyst assemblages from different localities of the Drake Passage area using multivariate data analysis and relate

the results with the tectonic and paleoceanographic settings for the Drake Passage during this critical interval.

MATERIALS AND METHODS

We reviewed the published middle Eocene dinoflagellate cyst data drawn from assemblages dominated by Antarctic–endemic species of sections from the Drake Passage area. We considered the following units for this analysis: (A) upper member of the Río Turbio Fm. studied in outcrop sections

Figure 2. Species from the Leticia Formation and the upper member of the Río Turbio Formation, Austral Basin. Specimens are identified by sample number/England Finder references. 1, **Enneadocysta dictyostila**, LE-RT-6/Q32. 2, **Deflandrea antarctica**, RT 12/1-9/P29/3. 3, **Arachnodynium antarcticum**, RT 12/1-4/N31/2; 4, **Vozzhennikovia apertura**, RT 14/1-1/P45/1. 5, **Lejeunecysta sp.**, RT 13/1-1/H17/4; 6, **Impagidinium parvireticulatum**, RT 15/1-6/M45/2. 7, **Selenopemphix nephroides**, RT 15/1-6/L28. 8, **Hystrichosphaeridium truswelliae**, RT 13/1-3/H33. 9, **Achomosphaera sp. A**, RT13/1-3/J30/2. 10, **Thalassiphora pelagica**, RT 12/1-6/Z37. 11, **Operculodinium centrocarpum**, RT 12/1-9/X45. 12, **Hystrichosphaeridium tubiferum**, LE-RT-5/R49. Scale bar= 20µm.



(González Estebenet *et al.*, 2014); (B) upper member of the Río Turbio Formation in YCP coreholes (Archangelsky, 1969; Guerstein *et al.*, 2014); (C) Leticia Fm. in Río de la Turba section (Guerstein *et al.*, 2008b); (D) Man Aike Fm. (Guerstein *et al.*, 2014); (E) La Meseta Fm. (Cocozza and Clarke, 1992); (F) Bruce Bank in core IO1578-59 (Mao and Mohr, 1995); (G) La Meseta Fm. (Wrenn and Hart, 1988; Douglas *et al.*, 2014) (Fig. 1; Tab. 1).

The taxonomic updating of dinoflagellate cyst species recorded from each locality was based on Williams *et al.* (2004), Fensome *et al.* (2006, 2008) and Sluijs *et al.* (2009). Table 2 shows a presence-absence data list defined with the species that appeared in at least two localities, including their biogeographic distribution according to Bijl *et al.* (2011, 2013a) and Houben *et al.* (2013). The magneto-stratigraphically calibrated ages of the diagnostic species in the Southern Ocean are provided by First Occurrence datum (FO), First Common Occurrence data (FCO), Last Occurrence datum (LO), including the South Pacific Dinocyst Zones (SPDZ) of Bijl *et al.* (2013a). In this work we selected the stratigraphic interval above the FCO of *Enneadocysta dictyostila* (Menéndez) Fensome *et al.*, 2006 in each locality. This dinoflagellate event represents the basis of Zone SPDZ11 of Bijl *et al.* (2013b) dated at 45.2 Ma (middle Lutetian). The youngest age of the analyzed sections is determined by the LO data of *Arachnодиниум antarcticum* and/or *Hystrichosphaeridium truswelliae* at 36 Ma in Zone SPDZ13 (Bartonian–Priabonian, 40.0 Ma–35.95 Ma). A similar biostratigraphic age model was applied by Douglas *et al.* (2014) for La Meseta Formation in Seymour Island. These authors considered that this formation was deposited between 45 and 34 Ma based on the FO of *Enneadocysta dictyostila* and the abundance of other Antarctic-endemic dinoflagellate cyst species, in contrast with previous early to late Eocene age assignments (Wrenn and Hart, 1988).

The palynological samples from the Leticia Formation and the upper member of the Río Turbio Formation were processed at the Museo Argentino de Ciencias Naturales, Bernardino Rivadavia, Buenos Aires, Argentina using hydrochloric and hydrofluoric acids. The residues were sieved through screens of 10, 25 and 180 µm and stained with Bismarck C. Strew mounts were prepared with gelatin glycerin as a mounting medium. The slides are housed at the Laboratorio de Palinología, Instituto Geológico del Sur, Bahía Blanca, Argentina (LPUNS). Light microscopy observation

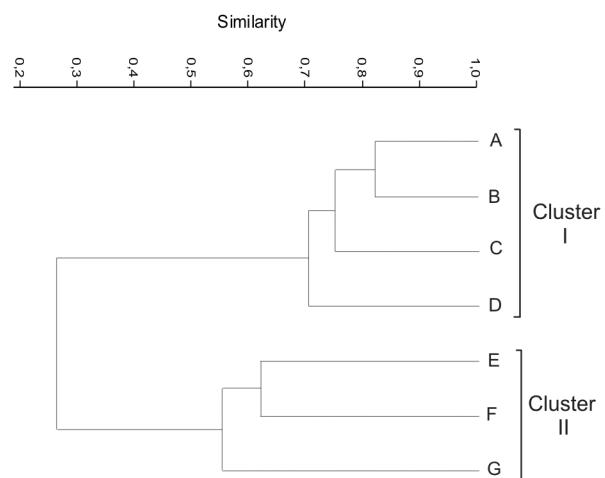


Figure 3. Dendrogram of localities based on non-endemic dinoflagellate cysts showing two clusters separated at 0.27 units of similarity. Cluster I comprised localities from the Austral Basin (A–D). Cluster II involves the localities from Antarctic Peninsula and Scotia Sea (E–G). For localities abbreviations see Table 1. Dendrogram obtained using the Unweighted Pair Group Method (UPGMA) and Dice index to compare similarities.

was carried out using a Nikon Eclipse 600 microscope and an attached Micrometrics high-resolution digital camera. Most of the species recorded in the Austral Basin are illustrated in Figure 2. Sample and slide number with England Finder (EF) references are provided for each illustrated specimen.

In order to visualize the differences among the dinoflagellate cyst assemblages from the localities compared we analyzed only the distribution of non-endemic taxa. The endemic species were recorded in all the tested sites since their consistent presence was the requirement to select the units for the comparison. Cluster analyses were carried out on both the localities and the species present in these sites using the Unweighted Pair Group Method (UPGMA) and Dice similarity index. Statistical analyses and charts were performed using PAST, version 2.17 (Hammer *et al.*, 2001).

RESULTS

The dendrogram based on the localities (Fig. 3) allows distinguishing two clusters separated at 0.27 units of similarity. Cluster I (0.71 units of similarity) is defined by the localities from the Austral Basin (localities A, B, C and D), and cluster II (0.56 units of similarity) comprises the localities from Antarctic Peninsula and Scotia Sea (localities E, F and G). Fig. 4 illustrates the results of the cluster analysis based on non-endemic species. The dendrogram defines two clusters

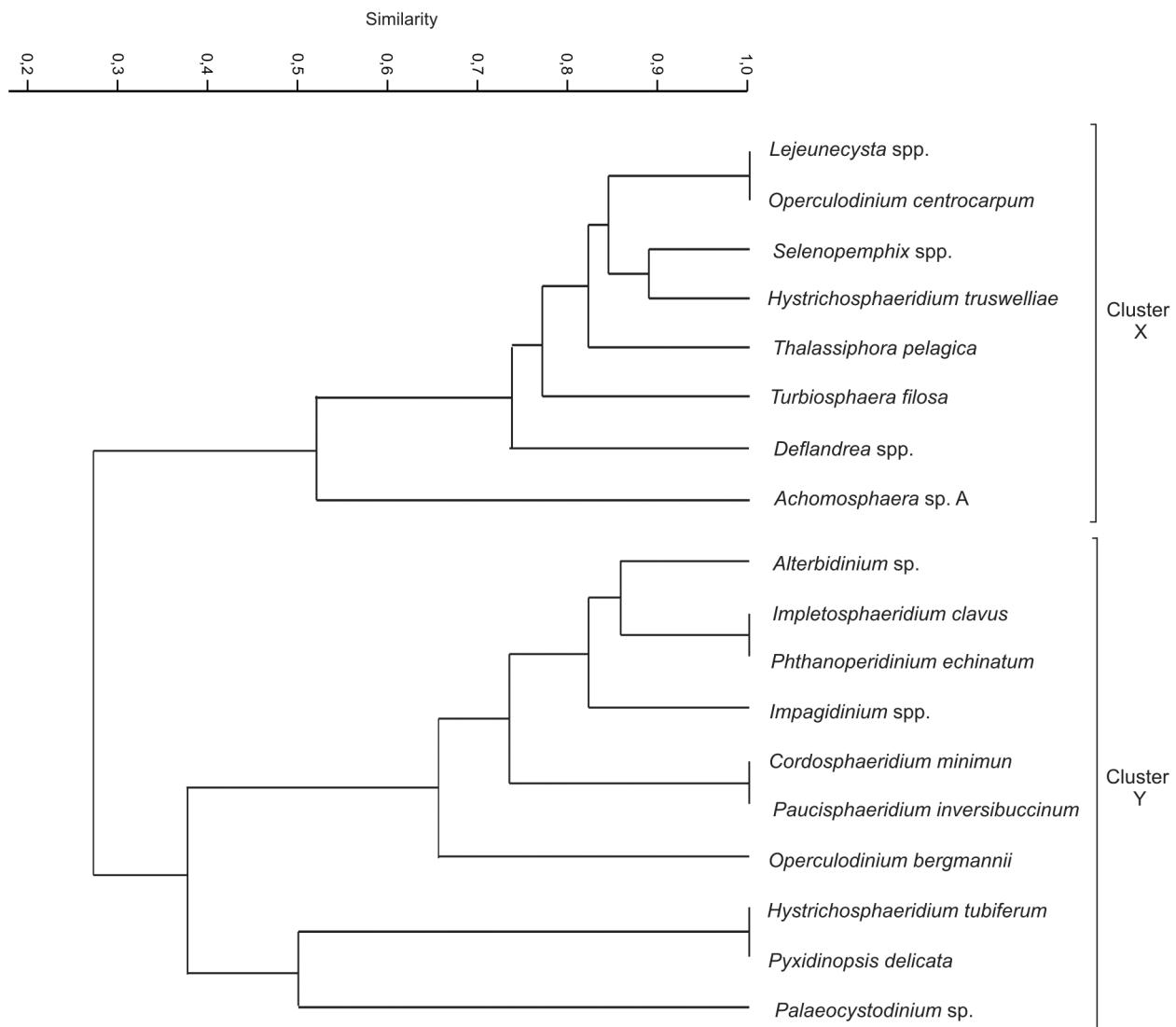


Figure 4. Dendrogram of the non-endemic dinoflagellate cysts showing two clusters (X and Y) separated by 0.27 units of similarity. Dendrogram obtained using the Unweighted Pair Group Method (UPGMA) and Dice similarity index.

separated at 0.27 units of similarity. Cluster X includes *Deflandrea* spp., *Lejeuneocysta* spp., *Operculodinium centrocarpum*, *Selenopemphix* spp., *Thalassiphora pelagica*, *Turbiosaera filosa*, *Achomosphaera* sp. A, and *Hystrichosphaeridium truswelliae*. Cluster Y is defined by *Alterbidinium* sp., *Cordosphaeridium minimun*, *Hystrichosphaeridium tubiferum*, *Impagidinium* spp., *Impletosphaeridium clavus*, *Operculodinium bergmannii*, *Palaeocystodinium* sp., *Paucisphaeridium inversibuccinum*, *Phthanoperidinium echinatum* and *Pyxidinopsis delicata*. Figure 5 depicts the results of a two-way cluster analysis indicating the relationship between taxa and sites based on the dendrograms represented in Figs. 3 and 4. The

result emphasizes that cluster I (including the localities from the Austral Basin; Fig. 3) comprises the species from cluster X (Fig. 4), and cluster II (comprising the localities from the Antarctic Peninsula and Scotia Sea; Fig. 3) is represented by the species of cluster Y (Fig. 4).

DISCUSSION

The palaeogeographic distribution of dinoflagellate cyst assemblages in the areas surrounding the Drake Passage shows the conspicuous presence of the Antarctic-endemic assemblage components. Although, the statistical analysis based on non-endemic taxa exhibited a clear difference between the locali-

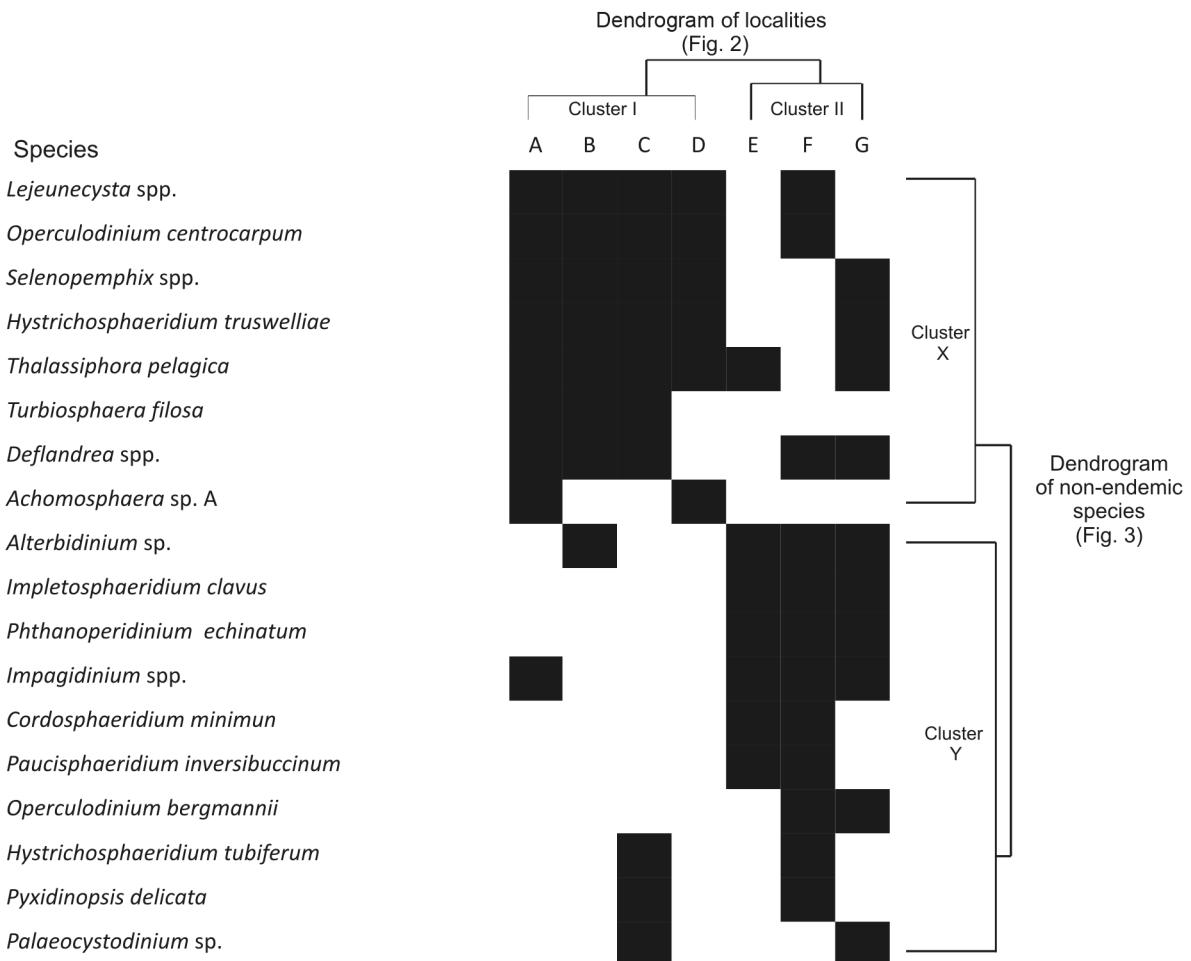


Figure 5. Two-way cluster analyses of dinoflagellate cyst species and localities. Localities and species are arranged following the dendograms in Figures 2 and 3. Species present in each locality are indicated with black squares.

ties to the north (Austral Basin) and south (Antarctic Peninsula and the Scotia Sea) of what today is the Drake Passage (Fig. 5). What palaeogeographic and palaeoceanographic configuration of the Drake Passage area could explain a broad distribution of the Antarctic-endemic assemblage, and at the same time a latitudinal differentiation based on non-endemic taxa?

Huber *et al.* (2004) proposed for the middle–late Eocene a surface ocean circulation model with extensive polar clockwise gyres (Fig. 6.1) and suggested that such gyres had generated a selective intrinsic mechanism called “cold trap”, which led to the endemism of species tolerant to relatively low surface-water temperatures. Guerstein and Daners (2010) considered that the mechanism proposed for the South Pacific could be similarly applied to the South Atlantic and thus con-

jectured the existence of the clockwise proto-Weddell Gyre (p-WG) equivalent to the proto-Ross Gyre (p-RG) (Fig. 6.1). Subsequently, this ocean circulation model was confirmed by circum-Antarctic dinoflagellate cyst biogeographic patterns (Bijl *et al.*, 2011) that would explain the presence of the Antarctic-endemic assemblage in all the analysed localities.

Throughout the past decades, general studies have described the progressive openings of the Tasmanian Gateway and the Drake Passage. Based on data from the IODP cores from the Wilkes Land Margin (eastern Antarctica), Bijl *et al.* (2013b) analysed the organic geochemical biomarker data along with the dinoflagellate cysts records suggested a connection developed between the Pacific Ocean and the Australo-Antarctic Gulf about 49–50 Ma ago. This connection would have resulted in a shallow waterflow through the Tas-

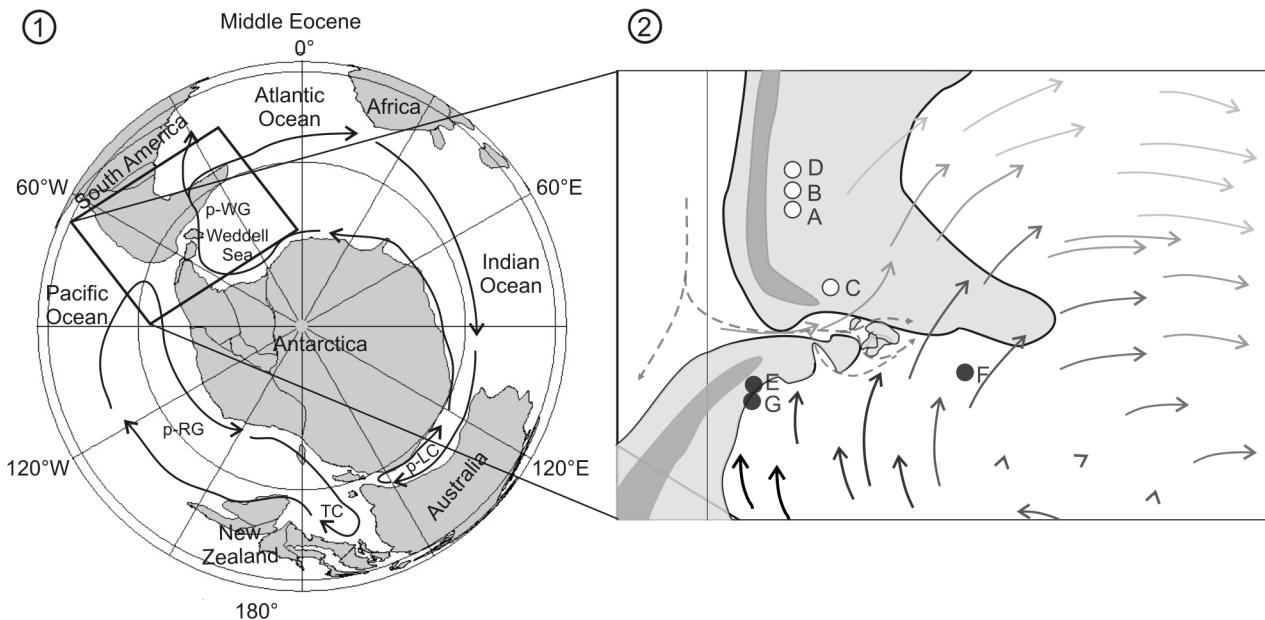


Figure 6.1, Middle Eocene paleogeographic map derived from Plate Tectonic Reconstructions On-line Paleogeographic Mapper (Schettino and Scotese, 2005). Gray areas indicate continental crust. Black lines indicate schematic oceanic circulation (modified after Huber *et al.*, 2004 and Guerstein and Daners, 2010). **2**, Middle Eocene Paleogeographic configuration of the Drake Passage with location of the studied units (A–G, see Tab. 1). Light gray areas indicate continental crust; dark gray areas indicate the maximum of the middle Eocene marine transgression (after Livermore *et al.*, 2007; Rodríguez Raisin, 2010). Dash-line arrows indicate the shallow water flow from the Pacific Ocean to the Atlantic Ocean through an incipient opening of the Drake Passage (modified from Livermore *et al.*, 2007). Full-line arrows represent the western part of the p-WG (after Huber *et al.*, 2004; Warnaar 2006 and Thomas, 2006). The darkest arrows represent the lowermost surface water temperatures. Abbreviations: **p-WG**, proto-Weddell Gyre; **p-RG**, proto-Ross Gyre; **TC**, Tasman Current; **p-LC**, proto-Leeuwin Current.

man Gateway 17 Ma earlier than the deepening of the Drake Passage. These authors interpreted that the current westward flow would have induced surface-water cooling along the Antarctic coast during the middle Eocene.

Livermore *et al.* (2005, 2007) and Eagles *et al.* (2006), based on magnetic anomaly profiles adjacent to the South American margin and in the southwest Scotia Sea, proposed the subsidence on the continental shelves at the beginning of the middle Eocene (~50 Ma). This subsidence could have developed an ocean passage characterized by shallow ocean connections between the Pacific and Atlantic oceans (Livermore *et al.*, 2007 figs. 2b–c). The shallow flows could have established a barrier for the continental fauna and might be probably responsible for the endemism of Eocene marsupials from La Meseta Formation on Seymour Island (Reguero *et al.*, 2002). Eagles *et al.* (2006) and Livermore *et al.* (2007) proposed at ~41 Ma, the development of two relatively deep-water gaps together with restricted current flows of shallow waters. This hypothesis is enhanced by the study of Scher and

Martin (2006), who suggested that the influx of shallow Pacific waters into the South Atlantic Ocean started at approximately 41 Ma, based on shifts of neodymium isotope ratios from fossil fish teeth in drill cores.

Considering the previous hypothesis on the tectonic evolution of the Drake Passage, the differences between the non-endemic dinoflagellate cysts in the localities of the Austral Basin and western of Antarctica might be influenced by surface waters from different sources. The sites located to the south of the Drake Passage (Antarctic Peninsula and Scotia Sea) would reflect the influx of surface waters originated in Antarctic coasts and directed to the North through the p-WG. Otherwise, the shallow flows through the incipient opening of the Drake Passage –as proposed by Scher and Martin (2006) and Livermore *et al.* (2007)– would have allowed the input of waters with cosmopolitan taxa from the Pacific to the Atlantic Ocean. These taxa would have been subsequently transported together with the Antarctic-endemic components into the Austral Basin by the northwestern sector of the p-WG (Fig. 6.2).

CONCLUSIONS

The dinoflagellate cyst distribution in the Southern Ocean has proven to be a powerful tool in reconstructing surface circulation patterns during the Paleogene and their modifications as a consequence of tectonic changes (Huber *et al.*, 2004; Stickley *et al.*, 2004; Bijl *et al.*, 2013b). In this work, we apply the analysis of middle Eocene dinoflagellate cyst assemblages from different localities close to the Drake Passage to test the hypothesis proposed by Scher and Martin (2006) and Livermore *et al.* (2007) about the onset of shallow connections between the Pacific and Atlantic oceans through an incipient Drake Passage during this interval.

The comparison of dinoflagellate cyst assemblages in the Drake Passage area shows a broad distribution of the Antarctic-endemic assemblage typical of the middle Eocene. However, a cluster analysis points out a pronounced difference between the non-endemic species from localities to the north (Austral Basin) and south (Antarctic Peninsula and the Scotia Sea) of what is nowadays the Drake Passage. We propose that the assemblages in the Antarctic Peninsula and the Scotia Sea reflect the influence of Antarctic surface waters conducted by the p-WG, while the dinoflagellate cyst assemblages from the Austral Basin may be the result of the confluence of Pacific water flows through an incipient Drake Passage and Antarctic waters transported northwards by the northwestern part of the p-WG. Thus, the palaeogeographic distribution of dinoflagellate cysts reinforces the hypothesis of an early opening of the Drake Passage during the middle Eocene.

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