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Benthic foraminifera associated to cold-water coral ecosystems

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There are three kinds of lies: lies, damned lies and statistics.

Mark Twain

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ABSTRACT

Cold-water coral reef ecosystems occur worldwide and are especially developed along the European margin, from northern Norway to the Gulf of Cadiz and into the Western Mediterranean Sea. The dominant reef builder in these areas is the scleractinian coral *Lophelia pertusa*, often associated with the scleractinian coral *Madrepora oculata*. These species settle on suitable hard substrates, in environments characterized by elevated currents and high food availability. Along the European margin cold-water coral reefs developed during different times and with different morphologies. In particular, on the Norwegian shelf and upper slope, extended active/living reefs have developed on elevated hard substrata. Along the Irish margin on the Rockall Bank, on the Porcupine Bank, and in the Porcupine Seabight, *L. pertusa* has built large fossil and/or active carbonate mounds. In the Gulf of Cadiz and in the Alboran Sea buried reefs and patch reefs with often strongly fragmented coral rubble are generally found in association with mud volcanoes below a (hemi-)pelagic sediment cover.

Cold-water corals have been known since the eighteenth century but the development of new technologies has resulted in the discovery of huge cold-water coral ecosystems and carbonate mounds during the last few decades. Their widespread occurrence presents a challenge to understand their development, preservation and possible importance in the geologic record.

In modern oceans, they provide important ecological niches for the marine benthic fauna in the deep-sea. In comparison to the macrofauna the microfauna, particularly the foraminifera associated with these systems, are poorly known. The present study focuses on the foraminiferal assemblages associated with cold-water coral ecosystems from the European continental margin. Samples were collected in three key regions:

surface sediments along the Norwegian margin and in the Porcupine-Rockall region, gravity cores were taken on the top of two mud volcanoes in the Alboran Sea. Planktonic and benthic foraminiferal assemblages were quantitatively analyzed. Since patterns of community structures are often not readily apparent, hierarchical cluster analysis, multidimensional scaling (nMDS), and diversity analysis were applied to emphasize differences in the foraminiferal assemblages.

Planktonic foraminiferal assemblages provide information about the conditions in surface waters. The assemblage of planktonic foraminifera in the coral-rich layer of two sediment cores in the Alboran Sea and in surface sediments from the Norwegian margin indicate similar conditions in the uppermost waters of both regions: nutrient-rich with enhanced phytoplankton blooms. At the transition from the coral-rich layer to the overlying (hemi-)pelagic sediments in the Alboran Sea a main shift in the planktonic foraminiferal assemblage indicate a reorganization of water masses towards more oligotrophic conditions. Low food availability could be the main reason for the decline of the cold-water coral reefs in this area.

The benthic foraminiferal assemblages in the Porcupine/Rockall region on the Irish margin are strictly related to the distribution of different sedimentary facies. On the Norwegian margin, benthic foraminiferal assemblages show a clear separation of cold-water coral reef associated fauna and off-reef associated fauna. However, different assemblages for each facies are only weakly defined and grade one into the other preventing a strict facies attribution as that observed along the Irish margin. The benthic assemblages from coral-rich layers in the Alboran Sea and those from cold-water coral reefs associated to carbonate mounds in the Porcupine/Rockall and cold-water

coral reefs along the Norwegian margin also show remarkable similarities. In particular, the assemblages are dominated by epifaunal-attached species such as *Discanomalina coronata*, *Cibicides refulgens*, and *Lobatula lobatula* but also infaunal foraminifera such as *Globocassidulina* spp., *Epistominella* spp., *Cassidulina* spp. are highly abundant. The benthic fauna provide information on currents, oxygenation and organic matter content. In particular, the benthic fauna associated with cold-water coral ecosystems indicates an environment characterized by high energy, well oxygenated waters and high organic matter supply derived from phytoplankton blooms and reaching the sea floor. In the investigated areas *D. coronata* is restricted to living cold-water coral reefs facies only and/or in co-occurrence with coral fragments. Based on these observations *D. coronata* is interpreted to require similar ecological conditions to cold-water corals and, therefore, proposed as an indicator species for healthy cold-water coral ecosystems on the European continental margin.

In conclusion, our data suggest that although cold-water coral ecosystems occur at different latitudes, the associated foraminiferal assemblages are consistent from Norway to the Western Mediterranean. Thus they can be used to identify these ecosystems even in the geologic record, when the corals are often strongly dissolved like in the Alboran Sea.

ZUSAMMENFASSUNG

Kaltwasserkorallenriffe treten weltweit auf, sind aber am stärksten verbreite entlang des europäischen Kontinentalabhangs zwischen Nordnorwegen und dem Golf von Cadiz. Sie kommen aber auch im Mittelmeer vor. Die Steinkoralle *Lophelia pertusa* ist die dominierende riffbildende Art und wird oft von einer zweiten Steinkorallenart, *Madrepora oculata*, begleitet. Die Larven dieser Arten besiedeln ausschliesslich Hartsubstrat. Starke bodennahe Strömungen und ein hohes Nahrungsangebot sind weitere Voraussetzungen für Kaltwasserkorallenwachstum. Entlang des europäischen Kontinentalabhangs etablierten sich Kaltwasserkorallenriffe während verschiedenen geologischen Zeiträumen und mit unterschiedlichen Erscheinungsbildern. Seit Beginn des Holozäns entwickelten sich grossflächige Riffe auf dem norwegischen Schelf und dem oberen Kontinentalabhang. Am irischen Kontinentalabhang, auf der Rockall Bank, Porcupine Bank und in der Porcupine Seabight, hat hauptsächlich *L. pertusa* seit dem späten Pliozän riesige fossile und teilweise noch aktive wachsende Karbonat-Hügel aufgebaut. Im Golf von Cadiz und in der Alboransee (westliches Mittelmeer) sind lebende Kaltwasserkorallenriffe sehr selten, begrabene fossile Riffe und „Patch“-Riffe aber häufig. Sie bestehen aus mehr oder weniger stark verwitterten Korallenbruchstücken und sind häufig mit Schlammvulkanen assoziiert.

Die Existenz von Kaltwasserkorallen ist schon seit dem achtzehnten Jahrhundert bekannt. Aber erst die Entwicklung von neuen Technologien in der Ozeanforschung in den letzten zwanzig Jahren hat zur Entdeckung der weitläufigen Kaltwasserkorallen-Ökosysteme und der riesigen Karbonat-Hügel geführt. Die grosse Verbreitung der Kaltwasserkorallen wirft viele Fragen auf, z.B. was sind die ökologischen Bedingungen für Kaltwasserkorallenwachstum, wie gut bleibt das Korallenskelett in der Tiefsee erhalten

oder wie verbreitet waren die Kaltwasserkorallen in der Vergangenheit.

Heutzutage sind Kaltwasserkorallen-Ökosysteme wichtige Nischen in der Tiefsee mit einer hohen Diversität an mariner benthischer Fauna. Im Vergleich zur Makrofauna ist über die Mikrofauna und speziell über die Foraminiferenvergesellschaftungen assoziiert mit Kaltwasserkorallen fast nichts bekannt. Die vorgelegte Studie befasst sich mit der Foraminiferenvergesellschaftung von Kaltwasserkorallenökosystemen entlang des europäischen Kontinentalabhangs. Planktonische und benthische Foraminiferen wurden in Sedimentoberflächenproben entlang des irischen und norwegischen Kontinentalabhangs und in Schwerelotkernen von Schlammvulkanen quantitative analysiert. Da einzelne Vergesellschaftungen in einem grossen Datensatz oft nicht gleich sichtbar sind wurden statistische Methoden wie Hierarchical Cluster Analysis, Multidimensional Scaling (nMDS) und Diversitätsanalysen angewandt.

Die planktonische Foraminiferenvergesellschaftung in den korallebruchreichen Schichten von zwei Sedimentkerne aus der Alboransee und in den Oberflächensedimenten von norwegischen Korallenriffen weisen auf hohe Nährstoffkonzentrationen in den oberflächennahen Wasserschichten und damit verbundenen hohe Primärproduktion hin. Beim Übergang von den korallenreichen Schichten zu den darüberliegenden (hemi-)pelagischen Schichten in den Alboransee-Bohrkernen verändert sich die planktonischen Foraminiferenvergesellschaftung dramatisch und weist auf eine Reorganisation der Wassermassen zu nährstoffärmeren Bedingungen mit geringer Primärproduktion hin. Die dadurch geringere

Nährstoffverfügbarkeit für die Korallen mag der Hauptgrund für das Absterben der Riffe sein.

Die benthische Foraminiferenvergesellschaftung in der Porcupine/Rockall Region am irischen Kontinentalabhang ist stark abhängig von der Fazies in der sie leben. Auf dem norwegischen Schelf und Kontinentalabhang unterscheidet sich die benthische Foraminiferenfauna zwischen der Vergesellschaftung, die im Riff lebt von derjenigen die ausserhalb lebt. Die starke Faziesabhängigkeit der einzelnen Vergesellschaftungen wie in der Porcupine/Rockall Region, ist in Norwegen nur schwach definiert. Der Grund dafür ist in Faziesunterschieden liegen, dies sich auf kleinstem Raum ändern und sich stark überlappen. Die benthische Foraminiferenvergesellschaftung von den korallereichen Schichten der Alboransee-Sedimentbohrkerne, den Oberflächensedimenten von den Karbonat-Hügeln in der Porcupine/Rockall Region sowie den norwegischen Korallenriffen zeigen grosse Ähnlichkeiten. Speziell epifaunale Arten wie *Discanomalina cornata*, *Cibicides refulgens* und *Lobatula lobatula* dominieren die Korallenriff-Vergesellschaftung. Aber auch infaunal lebende Foraminiferen wie *Globocassidulina* spp., *Epistominella* spp. und *Cassidulina* spp. sind häufig. Einzelne Arten benthischer Foraminiferen geben Auskunft über Strömungsgeschwindigkeiten, Sauerstoff- und Nährstoffgehalt des Wassers. Diese Arten zeigen auf, dass starke bodennahe Strömungen, sauerstoffreiche Wassermassen und ein hoher Eintrag an Nährstoffen, hauptsächlich von Phytoplanktonblüten, charakteristisch für dieses Ökosystem sind. Ebenfalls kennzeichnend für das Kaltwasserkorallenökosystem ist die Art *Discanomalina coronata*. Sie kommt nur in den lebenden Korallenriffen vor und lebt hauptsächlich auf abgestorbenen Korallenästen und anderem vorhandenen Hardsubstrat. Ausserhalb der Korallenriffe kommt *D. coronata* hingegen nicht vor. Es scheint, dass *D. coronata* die gleichen ökologischen Ansprüche wie die Kaltwasserkorallen und im speziellen *L. pertusa* hat. Aus diesem Grund kann *D. coronata* als Indikator-Art für gesunde Kaltwasserkorallenökosysteme des europäischen Kontinentalabhangs definiert werden.

Zusammenfassend kann gesagt werden, dass Kaltwasserkorallenökosysteme entlang des europäischen Kontinentalabhangs in verschiedenen geografischen Breiten vorkommen. Die Foraminiferenvergesellschaftung für diese Ökosysteme stimmen von Nordnorwegen bis ins westliche Mittelmeer überein. Diese kaltwasserkorallenassoziierten Foraminiferen können in Zukunft zur Identifikation

dieser Ökosysteme in der geologischen Abfolge genutzt werden auch wenn die Korallenbruchstücke, wie in der Alboransee, stark aufgelöst.

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1 - INTRODUCTION

1.1 COLD-WATER CORALS

For most people corals are marine animals living in the tropics, thriving in well-illuminated surface waters and may build up enormous coral reefs which host a very high biodiversity like the Great Barrier Reef on the north east coast of Australia. These ecosystems are known since hundreds of years and are very well investigated. Only a few people are presently aware that coral reefs from warm latitudes have a cold-water counterpart.

Cold-water corals were first discovered by Norwegian fisherman in the eighteen century and first described by Pontoppidan (1755), and Gunnerus (1768). However their potential to build up huge carbonate mounds is a very recent discovery and they are under investigation since a couple of decades only. Thanks to the development of new technologies and tools in deep-sea exploration such as manned and robotic submersibles, and sophisticated camera systems we are now able to investigate these fascinating ecosystems.

Comprehensive reviews about cold-water corals are given in the books “Cold-water Corals and Ecosystems” edited by Freiwald and Roberts (2005), “Deep-water Coral Reefs: Unique Biodiversity Hot-spots” (Hovland, 2008) and “Cold-water Corals: The Biology and Geology of Deep-Sea Coral Habitats” (Roberts et al., 2009). Additionally, the United Nations Environmental Program (UNEP) – World Conservation Monitoring Centre (WCMC) has published an overview about cold-water corals: “Cold-water Coral Reefs” (Freiwald et al., 2004) and short summary about the state-of-the-art is given in an article published by Roberts et al., 2006. However, the functioning of these ecosystems is far away to be fully understood and their global distribution is not mapped as yet as it is for warm water reefs.

However, we can assume that almost all oceans and seas worldwide may host cold-water corals when their ecological requirements are met. To the present they have been documented in Fjords and on the continental shelf of Norway (Fosså et al., 2002; Freiwald et al., 2002), along the upper continental slope from the Faroe-Shetland Channel south to central Africa, in the northwest Atlantic from Canada, Gulf of Mexico down to Brasil but also in the Indian and Pacific Oceans (Stetson et al., 1962; Freiwald et al., 1999; Paull et al., 2000; Heifetz, 2002; Reed, 2002; Freiwald et al., 2004; Gass and Willison, 2005; Mortensen and Buhl-Mortensen, 2005; Schroeder et al., 2005; Reyes et al., 2005). On deep-sea banks cold-water corals have been documented on the Rockall Bank in the northeast Atlantic, Galicia Bank close to Spain and the Chatman Rise and Campbell Plateau bear New Zealand and on several seamounts in the Atlantic and Pacific Oceans (Jaques, 1972; Grigg, 1984; Wilson and Kaufman, 1987; Richer de Forges, 1990; Grigg, 1993; Richer de Forges, 1993; Probert et al., 1997; Koslow et al., 2001; Andrews et al., 2002; Gubbay, 2003; Baco and Shank, 2005), and on mud volcanoes in the Gulf of Cadiz and in the Mediterranean (Van Rensbergen et al., 2005; Foubert et al., 2008; Freiwald et al., 2009; Wienberg et al., 2009). Peculiar and spectacular cold-water coral settings are the scleractinian coral reef build-ups, allowing the European margin from northern Norway down to the Mediterranean.

Lophelia pertusa is the main reef forming scleractinian cold-water coral (Freiwald et al., 2004). Minor components are *Madrepora oculata* and *Desmophyllum* spp. *Lophelia pertusa* may form white, orange or red bush-like colonies often of several meter heights. Its distribution is cosmopolitan but it occurs very frequently in the NE Atlantic Ocean (Zibrowius, 1980; Freiwald, 2004). Nevertheless, the full extend of its present geographical distribution is still unknown

(Freiwald et al., 2004). The shallowest occurrence has been recorded at 39 m depth in the Trondheimsfjord, the deepest from the New England Seamount chain in the North Atlantic, at 3383 m, and off Morocco, at 2775 m (Zibrowius, 1980).

The most significant environmental factors for cold-water coral distribution and growth are temperature, salinity, and nutrient supply. *Lophelia pertusa* tolerates a temperature range between 4 and 14 °C (Freiwald et al., 1997; Freiwald et al., 2002), and a salinity range between 32 psu and 38.8 psu (Strømgren, 1971; Taviani et al., 2005a). A combination of these two parameters is expressed in the density gradient sigma-theta (σ_θ). Recent studies show that *L. pertusa* coral reefs within a density gradient of sigma-theta (σ_θ) = 27.35 to 27.65 kg m⁻³ in the NE Atlantic Ocean (Dullo et al., 2008) and sigma-theta (σ_θ) = 29.08 and 29.13 kg m⁻³ in the Mediterranean (Freiwald et al., 2009). Hard substratum is required for initial attachment of the larvae. These factors are more important than specific depth range (Freiwald et al., 2004).

Lophelia pertusa is a suspension feeder (Messing et al., 1990; Jensen and Frederiksen, 1992) and needs to be supplied by a diverse range of food from live zooplankton to particle aggregates of marine snow and resuspended material (Mortensen et al., 2001; Freiwald, 2002; Kiriakoulakis et al., 2004; 2005; Duineveld et al., 2007). In regions where *L. pertusa* is abundant, high primary productivity by surface phytoplankton is observed (Duineveld et al., 2004). This is important to trigger the zooplankton blooms. Bottom current patterns also provide the cold-water corals with food, remove the waste products and limits sediment smothering (Klitgaard et al., 1997; Duineveld et al., 2004; Freiwald et al., 2004; White et al., 2007; Thiem et al., 2006).

The exploration of cold-water coral ecosystems with sophisticated camera systems and manned submersible dives has showed that these ecosystems can undergo serious damage e.g., habitat losses. The main threats are caused by commercial bottom trawling and other bottom fishing techniques, hydrocarbon exploration and production, cable and pipeline placement, bioprospecting and destructive scientific sampling, waste disposal and dumping, which are documented in the UNEP-WCMC report “Cold-water Coral Reefs” (Freiwald et al., 2004). It will be important for the future of these unique and diverse ecosystems, that scientist and policy makers put some effort to protect them.

1.2 COLD-WATER CORAL REEFS

Scleractinian cold-water corals can form biogenic reef frameworks, very similar in morphology to their warm-water counterpart, through complex interaction between biological and geological processes under suitable hydrodynamic conditions. A cold-water coral reef starts with the settlement of coral larvae on suitable hard substratum such as pre-existing height, moraine ridges, iceberg plough mark levees (Freiwald et al., 1999; Mortensen et al., 2001), mud breccia extruded from mud volcanoes (Chapter 4), and skeletal debris (Roberts et al., 2005). Under favourable environmental conditions like permanently or episodically strong currents and food supply, small coral colonies are able to grow (Dons, 1944; Frederiksen et al., 1992). Under stable physical oceanographic conditions in terms of temperature, salinity, food supply and strong currents, the colonies may continue their growth, colonize larger areas to form coral thicket (Dons, 1944; Freiwald, 2002). These thickets provide support and protection for other organisms, which form together a complex reef biocoenosis (Dons, 1944; Burdon-Jones and Tambs-Lyche, 1960; Jensen and Frederiksen, 1992, Fosså and Mortensen, 1998; Rogers, 1999; Fosså et al., 2000; Freiwald et al., 2004). The continuous growth of the reef results in a separation between the live reef and the dead framework providing different habitats resulting in distinct faunal zonation. Bioeroders, dominantly sponges and fungi, attack the dead corals (Beuck and Freiwald, 2005). This process produce the formation of extended fields of coral rubbles, which provide additional different habitat for distinct fauna but also the substrata for renewed coral settlement supporting horizontal reef growth. The bottom circulation pattern may produce a facies zonation, which can be identified by the presence of abundant exposed glacial dropstones in the northern regions. This facies harbour distinct communities, different from the more sheltered areas (Mullins et al., 1981; Messing et al., 1990).

Several reef complexes have been described along the Norwegian margin. The largest reefs are the Sula Reef at 64°N and the Røst Reef at 67° N. The Sula Reef is located on the continental shelf at around 290 m water depth measuring 14 km by 350 m with a hight of around 25 m (Freiwald et al., 2002). The Røst Reef is located on the continental shelf break between 300 and 400 m water depth, is 35 – 40 km long, and up to 3 km wide (Fosså et al., 2005). Cold-water coral reefs also often settle on the sill at fjord entrances like in the Stjernsundet at a water depth around 260 m (Freiwald et al., 1997). The northernmost coral reef “Korallen” is

located at almost 71° N on the Norwegian continental shelf at around 210 m water depth (Chapter 3).

1.3 COLD-WATER CORAL CARBONATE MOUNDS

Cold-water coral can also colonize structural high and form carbonate mounds. The known occurrence of cold-water coral carbonate mounds is generally confined to the upper and mid-slope of continental margins like the Porcupine Seabight, the Gulf of Cadiz, the Moroccan and Mauritanian margins, the Florida-Hatteras Straight, the Blake Plateau (Florida), the eastern USA and the Gulf of Mexico (e.g., Newton et al., 1987; Colman et al., 2005; De Mol et al., 2005; Grasmueck et al., 2006, Foubert and Henriet, 2009). The growth rate of coral carbonate mounds is high (up 0.05 mm yr⁻¹ under favourable conditions for cold-water corals) in comparison to off-mound sedimentation rates (Freiwald et al., 1999; Lindberg et al., 2007). For this reason cold-water coral carbonate mounds can be also called “carbonate factories” (Tucker and Wright 1990; James and Bourque 1992) although most of them occur in mixed carbonatic – siliciclastic domains.

These mounds are interpreted to be formed by cyclic development of cold-water coral, which includes a number of processes acting in different ranges of temporal and spatial scales as described in several models (De Mol et al., 2002; Kenyon et al., 2003; De Mol et al., 2005; Dorschel et al., 2005; Huvenne et al., 2005; Kozachenko et al., 2005; Roberts et al., 2006; Rüggeberg et al., 2007; Huvenne et al., 2009). All these models have a common point, which is the widely accepted mechanism of cold-water coral mound initiation from a cold-water coral reef (Roberts et al., 2009). According to this theory, mounds develop from extended cold-water coral reefs (Williams et al., 2006; Kano et al., 2007) by vertical coral growth on accumulated coral rubble, sediment accumulation of biogenic and authigenic carbonate and sediment baffled in the coral framework. The majority of these models show also that climatic changes from interglacial to glacial causes fundamental changes in the environmental conditions and in the sedimentation rates, thus in the corals development.

Rüggeberg et al. (2007) showed that the decrease in temperature, nutrient supply, current speed and increase in sediment input during glacial times produces unfavourable conditions for cold-water coral growth. They show that the return to interglacial/interstadial conditions is marked by the return to relatively warmer temperatures and by the re-establishment of high speeds

in the circulation patterns with consequent removal of the glacio-marine deposits, thus producing again the favourable conditions for cold water coral growth. In the northern hemisphere glacial/interglacial cycles occurred many times over the last 2.7 Ma (Bartoli et al., 2005). Consequences of this cyclicity are the typical mound sequences with fine grained sediments accumulated during glacial times and coarser deposits accumulated during interglacial/interstadial (Dorschel et al., 2005; Rüggeberg et al., 2007; Huvenne et al., 2009).

Cyclic sedimentation is also responsible for the accumulations of thick mound deposits. When mounds reach a relevant size, their top may become isolated from bedload transport, thus they cannot longer expand and they may result embedded within sediment drifts whose accumulation rate is higher than the mound growth rate (Van Rooij et al., 2003, 2007a, b). In some case large mounds can shape their own hydrodynamics with the establishment of a circulation pattern that erodes the mound itself (Wheeler et al., 2005; White et al., 2005; Dorschel et al., 2007a; Wheeler et al., 2007) until it is completely buried (Huvenne et al., 2003, 2007; Van Rooij et al., 2008).

1.4 FORAMINIFERA

Foraminifera are unicellular amoeboid protists distributed worldwide and mainly restricted to marine environments. They constitute one of the most diverse groups of shelled organisms in modern oceans and are a major component of marine communities. Around 10,000 living foraminiferal species are presently known (Vickerman, 1992). The majority of them are benthic, e.g., they live on the sea floor or in the sediment from the seashore to the deep-sea (e.g., Murray, 2006). Only about 40-50 species are planktonic (floating in the water mass; Kennet and Srinivasan, 1983). Most foraminifera produce a mineralized shell, generally calcitic, which can have either one or multiple chambers, and can be fossilized.

Shells of foraminifera have been noticed in shore sands since the 17th century but the first modern description of foraminifera were made by Beccarius in 1731. In the 18th and 19th century foraminiferal research was only based on taxonomical work and the discovery of new species (e.g. Fichtel and Moll, 1798, d'Orbigny, 1826). In the 20th century foraminiferal studies intensified with the requirements of oil companies for dating and correlating sediments (e.g. Cushman, 1928) and first paleoceanographic and ecological applications were

made in the middle of the 20th century. For example, Emiliani (1955) introduced the study of stable oxygen and carbon isotopes from foraminiferal shells to reconstruct paleoclimates and Boltovskoy (1956) and Phleger (1960) did pioneer works on foraminiferal ecology. Since then, foraminifera have been one of the most important tools for geologic, biostratigraphic and paleoceanographic reconstructions because as pointed out in Spezzaferri and Spiegler (2005):

- They are highly diversified, numerous, easily recognizable, and often their shell is well preserved.
- They occur in high numbers in all marine environments from the Polar region in both hemispheres to the Equator, in marginal to deep basins, floating in surface waters down to living within the sediments.
- Benthic foraminifera first occur in the Cambrian, whereas planktonic foraminifera exist since the Jurassic.
- Some genera and species appear, evolve and become extinct in a very short time and, therefore, represent excellent marker fossils.
- The composition of assemblages reflects ecological conditions and is highly sensitive to environmental influences.
- Several species of planktonic and benthic foraminifera build their shells in equilibrium with seawater and can be used to trace variations occurring in the chemistry of the water masses.

Benthic foraminiferal taxonomy is primarily based on the morphology of the tests, whereas the taxonomy of planktonic foraminifera is based on the characteristic of the wall textures and the function and significance of structures. Spines, pustules, keels have been unraveled as evolutionary-driven life strategies and adaptation to changing paleoecological settings (e.g., Hemleben et al. 1989). A relatively new field in systematics started with genetic studies on foraminifera (e.g., Pawłowski et al., 1994; Darling et al., 1996), which will have a major impact on foraminiferal taxonomy in future.

The criteria and the rules on which the taxonomy of foraminifera is based are formulated by the “International Commission on Zoological Nomenclature” and reported in the “International Code of Zoological Nomenclature (ICZN)” (Ride et al., 2000). The code fixes the criteria of selection and naming of holotypes of each known taxon. The holotype is a single physical example or illustration of an organism, known to have been used when the species was formally described. Holotypes are listed and illustrated in the Ellis and Messina catalogues

(Ellis and Messina (1940 and later). At genus level the recent foraminiferal taxonomy is based on Loeblich and Tappan (1987) “Foraminiferal Genera and their Classification”. The taxonomic concepts from both the ICZN, Ellis and Messina (1940 and later) as well as Loeblich and Tappan (1987) are applied to foraminiferal species reported in the photographic appendix (Plate 1 to 42).

1.4.1 Deep-sea benthic foraminiferal ecology

Benthic foraminifera are an important component of the deep-sea biomass in the present oceans. They are adapted to cold, dark and often extremely oligotrophic environments. The benthic foraminiferal fauna is highly diverse and many species have a cosmopolitan distribution. These organisms convey a remarkable amount of information about the present and past ecological conditions at the ocean seafloor and have played an important role to understand the functioning of the marine systems.

Many investigations have so far proved and documented the relationship between foraminiferal species and different ecological parameters (including physico-chemical) and how these parameters control spatial and temporal dynamics of foraminiferal communities (e.g. Van der Zwaan, 1999; Gooday, 2003; Murray, 2006; Jorissen et al., 2007).

Foraminifera can be considered as reliable indicators for the origin, quality, quantity, and periodicity of organic matter reaching the seafloor (Lutze and Coulbourn, 1984; Altenbach and Sarnthein, 1989; Herguera and Berger, 1991; Gooday, 1994; Altenbach et al., 1999; Loubere and Fariduddin, 1999). One of the major environmental parameter structuring deep-sea benthic foraminiferal faunas is the organic matter flux mainly derived from primary production at the sea surface. In eutrophic environmental settings, the organic particles cannot be consumed directly by benthic organisms, the organic matter can accumulate and thus, sediment pore waters become anoxic. In these cases the limiting variable for benthic foraminiferal distribution become the oxygen (Koutsoukos et al., 1990; Hermelin, 1992; Alve, 1995; Jorissen et al., 1995; Bernhard and Sen Gupta, 1999).

Additional limiting environmental parameters for benthic foraminiferal are sediment grain size and current velocity (e.g. Miller and Lohmann, 1982; Lutze and Coulbourn, 1984; Mackensen et al., 1990, 1995; Schmiedel et al., 1997; Schönfeld, 2002b). Water

mass properties such as salinity and temperature and bathymetry play only a minor role in their distribution (Jorissen et al., 2007).

1.5 MOTIVATION AND MAIN OBJECTIVES

Only recently scientists have started to understand the complex interaction of ecological variables controlling cold-water coral ecosystems. Further investigations are still needed to obtain a complete picture of cold-water coral reefs and their ecology. It is known that these ecosystems are “hot-spots” for marine life and host thousands of species of sponges, hydrozoans, mollusks, bryozoans, echinoderms, polychaetes, crustaceans, and fishes with a comparable biodiversity as observed for their warm-water analogues (Dons, 1944; Burdon-Jones and Tambs-Lyche, 1960; Jensen and Frederiksen, 1992; Fosså and Mortensen, 1998; Rogers, 1999; Fosså et al., 2000; Freiwald et al., 2004). Until now, studies on cold-water coral associated faunas mainly focused on the mega- and macrofauna (e.g., Jensen and Frederiksen, 1992; Costello et al., 2005; Henry and Roberts, 2007) or microfauna (e.g., Penn et al., 2006; Neulinger et al., 2008; Schrötter et al., 2009).

Studies on foraminifera associated to cold-water corals are very few. Freiwald and Schönfeld (1996) focused on the single parasitic foraminifera *Hyrrokkin sarcophaga*, (Hawkes and Scott, 2005) investigated the benthic foraminifera associated to an “octocoral garden” on the east coast of Canada and (Rüggeberg et al., 2007) focused on benthic foraminiferal assemblages in sediment cores on a carbonate mound in the Porcupine Seabight.

Since 2005 the micropaleontology group of Fribourg and the marine geologists of the GEOMAR-Kiel and Renard Center of Marine Geology of Ghent started a collaboration to study these organisms in detail. The goal of this thesis, founded by the Swiss National Foundation Project Ref. 200020-117928, is to present an extensive picture about recent and sub-recent benthic foraminifera associated to scleractinian cold-water coral ecosystems along the European margin from northern Norway down to the Mediterranean.

Three key regions for cold-water coral carbonate mounds and reefs were investigated:

Chapter 2: The Porcupine Seabight-Rockall region. Surface sediment samples from different facies and different coral carbonate mounds in the Porcupine

Seabight, on the Porcupine Bank, and on the Rockall Bank were investigated using multivariate statistical methods on quantitative benthic foraminiferal census. The results indicate that different benthic foraminiferal assemblages characterize different facies along cold-water coral carbonate mounds and are related to the different environmental conditions and available substrate. Therefore benthic foraminiferal assemblage analysis provides an independent tool to identify the different facies, which characterize cold-water coral mounds in this region. The *Off-mound Facies* is dominated by infaunal species indicating high organic matter flux to the seafloor. The *Dropstone Facies* is dominated by *Cibicidoides* sp., living attached to dropstones and thriving in well-oxygenated high-energy environment. The *Dead Coral Facies* is characterized by epifaunal and infaunal species indicating strong bottom currents. The *Living Coral Facies* is dominated by opportunistic infaunal species indicating low food supply on the seafloor. Cold-water corals may consume the major part of the food and only a small amount passes through the coral thicket to the seafloor. The *Sandwave Facies* is characterized by high abundances of epifaunal species. *Discanomalina coronata* is only highly abundant in the *Living Coral Facies* and *Sandwave Facies*, where living cold-water corals thrive and is completely absent in adjacent sediments/facies. Therefore, *D. coronata* is proposed to be a indicator species for living/healthy cold-water coral mounds in this area.

Chapter 3: Norwegian Margin. On the Norwegian shelf and on the upper continental slope, extensive reefs settle on post-glacial structures. Surface sediment samples were taken from different reefs from the Skagerrak to northern Norway. Planktonic and benthic foraminifera were quantitatively analyzed, and the different assemblages characterized using multivariate statistics. Two different planktonic foraminiferal assemblages reflect the two major surface current systems on the Norwegian margin: *Neogloboquadrina incompta* dominates the North Atlantic Current whereas very small specimens of *Globigerinita glutinata*, *Turborotalita quinqueloba*, and *Globigerinita uvula* dominate the Norwegian Coastal Current. The benthic foraminifera show different assemblages for on-reef and off-reef samples but also regional differences related to the different environmental conditions. In particular, the foraminiferal assemblage associated to the healthy reefs on the mid Norwegian slope and northern Norwegian shelf are typical of environments with high energy, well-oxygenated waters and high amount of labile organic matter, derived from seasonal phytoplankton blooms. The benthic foraminifera

associated to the coral reefs from the Skagerrak indicate that the amount of organic matter input to the sea floor is high but the labile components usable for corals are low. The low amount of labile organic matter may explain the extensive occurrence of dead corals in this area.

In the Norwegian reefs benthic foraminiferal habitats are only weakly defined and grade one into the other preventing sharp facies separation as observed on the cold-water coral carbonate mounds in the Rockall/Porcupine region (Chapter 2). The gradual changes in the assemblages observed on the Norwegian margin can be attributed to the changes in the reef facies, which are restricted within tens of meters, whereas the facies on the carbonate mounds in the Porcupine/Rockall region occur over long distance. *Discanomalina coronata* is only abundant in healthy cold-water coral reefs indicating that this species require similar ecological conditions as cold-water corals. Therefore *D. coronata* is proposed as an indicator species for living cold-water coral ecosystems on the Norwegian margin.

Chapter 4: The Alboran Sea – Western Mediterranean. Samples from sediment cores of two Mud Volcanoes (Maya and Dakha) in the Alboran Sea - Western Mediterranean were investigated using facies description and multivariate statistical methods on quantitative planktonic and benthic census. These investigations were carried out to test if there is a causal link between *D. coronata* and cold-water coral ecosystems during the Holocene. *Discanomalina coronata* was observed only in those layers containing partly dissolved cold-water coral fragments. The co-occurrence of these elements shows a striking similarity to their common distribution pattern observed in the Porcupine/Rockall Region (Chapter 2) and on the Norwegian margin (Chapter 3), thus suggesting that the ecosystems in the Alboran Sea are in-situ. Coral growth on the two mud volcanoes are observed in two phases, on the Dakha MV between 2230 ± 59 years BP and slightly older than 4175 ± 62 years BP. On the Maya MV cold-water corals occur between 7616 ± 38 years BP and slightly older than 15583 ± 185 years BP. On both mud volcanoes cold-water corals occur on the top of the extruded mud breccia, which provides the nucleation point for colonization and development of the corals. The period of cold-water coral development on both mud volcanoes is characterized by planktonic foraminifera like *Neogloboquadrina incompta*, indicating high nutrient availability, triggered by upwelling and/or strong currents that mobilized horizontal nutrient fluxes. The decline of the cold-water corals on both

mud volcanoes co-occurs with a shift of *N. incompta* dominated period to a *Globorotalia inflata* dominated period, which reflects more oligotrophic conditions observed in the modern Western Mediterranean.

2 - BENTHIC FORAMINIFERA AS BIOINDICATOR FOR COLD-WATER CORAL REEF ECOSYSTEMS ALONG THE IRISH MARGIN

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ABSTRACT

Cold-water coral ecosystems building cold-water carbonate mounds occur worldwide and are especially developed along the European margin, from northern Norway to the Gulf of Cadiz. A remarkable mound province is documented southwest of Ireland along the Porcupine and Rockall Banks. In this area carbonate mounds are formed in water depths between 500 and 1200 m and are often densely settled by cold-water coral ecosystems offering many ecological niches for benthic foraminifera. We investigated total (unstained) benthic foraminiferal assemblages from surface sediments (0–1 cm, >63 µm size fraction) of this region with the aim to trace their distribution patterns and to test if they can be used as bioindicators for facies characterization in the different parts of carbonate mound systems. Our quantitative data were further statistically treated with non-metric multidimensional scaling (nMDS) based on Bray-Curtis similarity matrix to highlight community patterns that were not readily apparent. Our results indicate that different benthic foraminiferal assemblages characterize different facies along cold-water carbonate mounds and are related to the environmental conditions and available substrates. The following facies can be described: (1) the *Off-Mound Facies* is dominated by uvigerinids and other infaunal species; (2) the *Dropstone Facies* is characterized by infaunal *Globocassidulina subglobosa* and attached-epifaunal *Cibicidoides* sp.; (3) the *Dead Coral Facies* is characterised by epifaunal species (e.g., *Planulina ariminensis*, *Hanzawaia boueana*) and infaunal species (*Spiroplectinella wrightii*, *Angulogerina angulosa*, *Epistominella vitrea*); (4) the *Living Coral Facies* includes both infaunal and epifaunal species, but is dominated by the epifaunal *Discanomalina coronata*; (5) the *Sandwave Facies* contains high abundances of epifaunal species including *D. coronata*. Based on this distribution, we propose *D. coronata*, as an indicator species to identify active mounds and/or living cold-water coral ecosystems. Our results also emphasise the importance of studying the small size fractions that yield many infaunal species. A causal link exists between distribution patterns of benthic foraminifera and cold-water coral facies, thus providing an independent tool to identify and describe the different facies in this setting.

2.1 INTRODUCTION

Cold-water coral ecosystems occur worldwide and are developed along the European margin, from northern Norway (Fosså et al. 2002; Freiwald et al. 1997, 1999; Hovland et al. 1998; Lindberg and Mienert 2005; Mortensen et al. 1995) along the Irish margin (De Mol et al. 2002; Hovland et al. 1994; Kenyon et al. 2003; Van Rooij et al. 2003) down to the Gulf of Cadiz (Pinheiro et al. 2003; Somoza et al. 2003), and in the Mediterranean Sea (Taviani et al. 2005; Zibrowius, 1980). These ecosystems have been known since the last century. Cold-water corals were collected and successively studied by Duncan (1879, 1873, 1878), Gravier (1915, 1920), and mapped by Joubin (1922a, b, 1923). The investigation of these ecosystems

proceeded more quickly in Norway where cold-water corals were more easily accessible. However, research accelerated only in the last two decades after Hovland et al. (1994) described a group of seabed mounds, suspected to be modern bioherms off western Ireland. Since then, research has progressed quickly within the framework of several European and international projects (ECOMOUND, GEOMOUND, ACES, MOUNDFORCE, MICROSYSYTEM, HERMES).

On the Rockall and Porcupine Banks, cold-water corals, particularly the scleractinians *Lophelia pertusa* and *Madrepora oculata*, build up carbonate mounds. These mounds considerably differ in size and some are partially or fully buried (De Mol et al. 2002; Huvenne et al. 2003). Recent investigations in this area revealed

that successive phases of coral growth might be superimposed on fossil coral debris building mounds

up to a few hundred meters in height (Wheeler et al. 2007). The amount of living corals is presently used

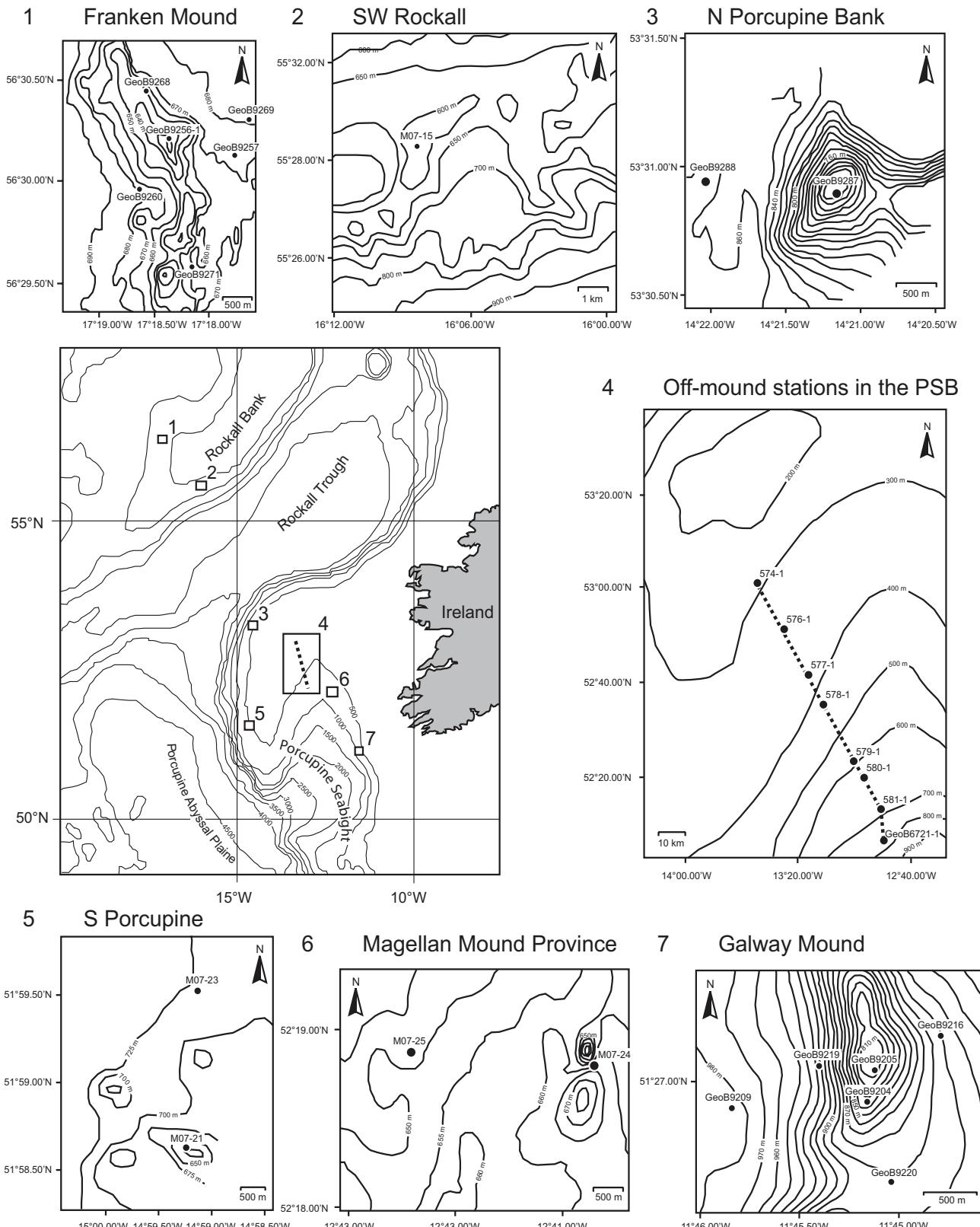


Figure 2.1 Location map of the surface sediment samples investigated in this study. The different carbonate mound provinces along the Rockall Bank (1, 2), the Porcupine Bank (3, 5), and the Porcupine Seabight (4, 6, 7) are highlighted.

as the main parameter for “activity” of cold-water carbonate mounds. If living corals are abundant, the mound is considered to be in a growing state - a so-called “active” or “active growing” mound (Henriet et al. 2002; Rüggeberg et al. 2007). If living corals are absent and pelagic sediment covers the mound, it is then defined as a “buried” mound and is only visible on seismic sections (De Mol et al. 2002; Freiwald 2002).

Presently, cold-water coral mounds are described and classified based on their sedimentary facies and type of coral coverage. Foubert et al. (2005) introduced 12 different facies for seabed classification and used them for interpreting ROV video surveys in the Belgica Mound Province. The study of Dorschel et al. (2007a), focused on Galway Mound, reduced the number of the different facies to seven from mound top down to the basin: (1) dense coral coverage (mostly alive); (2) dense coral coverage (mostly dead); (3) sediment clogged dead coral framework and/or coral rubble; (4) patchy distribution of mostly live (or dead) coral on un-rippled seabed; (5) patchy distribution of mostly dead coral on rippled seabed; (6) sandwaves covered with corals (overgrown sandwaves); (7) patchy distribution of dropstones. Dorschel et al. (2007a) also demonstrated that the spatial distribution of the different facies correlates with different small-scale environmental settings and in particular, with local hydrodynamic regimes.

However, until now only a few studies targeted foraminiferal assemblages and their relation to cold-water coral ecosystems along the European continental margin (e.g., Freiwald and Schönfeld 1996; Hawkes and Scott 2005; Rüggeberg et al. 2007). The present study focuses on the distribution patterns of total (unstained) benthic foraminiferal assemblages that are associated with different cold-water coral mounds on the Rockall and Porcupine Bank area (Fig. 2.1). The aims of this study are to: (a) identify the distribution patterns of benthic foraminifera on cold-water coral mounds in the investigated area; (b) relate these to the different facies; and (c) propose a model for assemblage versus facies distribution that can be applied to other carbonate mounds in similar settings.

2.2 STUDY AREA

The Porcupine and the Rockall Banks are situated in the Northeast Atlantic west of Ireland (Fig. 2.1). Steep flanks characterize the Rockall Trough, a deep-sea channel between the Porcupine Bank to the east and

the Rockall Bank to the west. The Porcupine Bank and the Irish Mainland Shelf delimit a small basin, the Porcupine Seabight, which opens to the west onto the Porcupine Abyssal Plain. Along the flanks of the Porcupine Bank and Porcupine Seabight numerous carbonate mounds of different sizes are documented (e.g., De Mol et al. 2002; Foubert et al. 2007; Huvenne et al. 2002, 2003; Mienis et al. 2006). In particular, three carbonate mound provinces are located along the margin of the Porcupine Seabight: (a) the Belgica Mound Province, which is located along the steep eastern slope, (b) the Hovland Mound Province along the margin of the central Porcupine Seabight and (c) the Magellan Mound Province north of the Hovland Mound Province (De Mol et al. 2002; Henriet et al. 1998; Huvenne et al. 2003; Van Rooij et al. 2003).

Several mounds from the Porcupine and Rockall Bank region have been investigated for the present study (Fig. 2.1). (1) The Franken Mound is situated on the steep western slope of the Rockall Bank. Its summit rises up to 600-675 m water depth. It is 2.5 km long and 600-700 m wide. It has a more irregular shape compared to the ovate shape of the Galway and the Propeller mounds in the Porcupine Seabight (Wienberg et al. 2007). (2) Single mounds on the southeastern Rockall margin, up to 1-2 kilometres in diameter and 50-100 m of elevation. They are all located in a water depth between 650 and 900 m (Akhmetzhanov et al. 2003; Mienis et al. 2006; Rogers, 1999; Van Weering et al. 2003). (3) The Connaught Mound located on the steep slope of the Northern Porcupine Bank. It is orientated north-south, about 3 km long and 1.7 km wide with an elevation above the seafloor of 180 m and a water depth at the summit of 680 m. (4) The Propeller Mound is situated in the Hovland Mound Province. Its summit is at a water depth of 680 m, its elevation above the surrounding seafloor is 140 m (Dorschel et al. 2005, 2007b; Rüggeberg et al. 2005, 2007). (5) Unnamed mound from the western Porcupine Bank discovered in 2008 during the ‘Pelagia’ cruise M07II. (6) Unnamed half buried mounds from the Magellan Mound Province (De Mol, 2002; Huvenne et al. 2003). (7) The Galway Mound, which is part of the Belgica Mound Province. Its summit is at 782 m water depth and its elevation above the seafloor is 160 m (Dorschel et al. 2007a). It belongs to an “active” and “growing” mound chain with a dense coverage of living corals (De Mol et al. 2002; Foubert et al. 2005; Huvenne et al. 2005; Wheeler et al. 2005).

2.3 MATERIAL AND METHODS

During four different cruises (RV ‘Meteor’ cruise M61-3, RV ‘Poseidon’ cruises P265, and P292, and RV ‘Pelagia’ cruise M07II) carbonate mounds and pelagic sediments were sampled at 27 stations in the Rockall and Porcupine Bank region (Tab. 2.1). A giant box corer with a sampling area of 50x50 cm allowed a maximum 50 cm penetration into the sediments. All retrieved samples were taken from cold-water coral mounds and/or the adjacent pelagic sediments. Samples P292/564-1, P292/576-1, P292/577-1, P292/578-1, P292/579-1, P292/580-1, P292/581-1, and GeoB6721-1 were retrieved along a bathymetric transect from the Porcupine Bank at ~200 m down to the Porcupine Seabight at ~750 m (Fig. 2.1). A sample from the uppermost two centimetres of each box core was taken with a teflon spatula and stored in plastic containers. Additionally, video records obtained by an underwater video camera or a Remotely Operated Vehicle (ROV ‘Quest’ and ‘Cherokee’ of MARUM, University Bremen) were used to characterize the seafloor, together with descriptions of cores from unpublished cruise reports (De Haas, 2007; Freiwald et al. 2000; Freiwald et al. 2002; Ratmeyer et al. 2004).

Sediment samples for micropaleontological analyses were processed at the University of Fribourg following Spezzaferri and Coric (2001). Samples were washed through a set of 250 µm, 125 µm and 63 µm mesh

sieves to obtain three size fractions. The obtained residues were dried at room temperature and weighted. If the residue contained more than 400 foraminiferal specimens in one fraction, the volume was reduced by splitting. A maximum of 200 benthic specimens per fraction (600 specimens per sample) were picked, collected in Plummer cell-slides, sorted at species level, fixed with glue, and counted. In the case of samples containing less than 100 specimens per fraction, all specimens were counted. The 124 benthic foraminiferal species identified are listed in Appendix A.

Multivariate statistical treatment on the compositional faunal data was performed with the software PRIMER 5 (Clarke, 1993; Clarke and Warwick, 2001). Data were double-square root transformed in order to limit the contribution of most abundant, ubiquitous species (Field et al. 1982). Bray-Curtis (dis-)similarities were calculated following Clifford and Stephenson (1975). The resulting similarity matrix was used to obtain the nMDS plot (non-metric MultiDimensional Scaling; Kruskal 1964; Kruskal and Wish 1978). The nMDS plot has no dimensions and no axes and can be arbitrarily scaled, rotated, located or inverted as it gives simply the relationship of samples relative to each other (Clarke and Warwick 2001; Warwick and Clark 1991). Based on the clusters given in the nMDS plot, the Similarity Percentage Analysis (SIMPER) was obtained to highlight the contribution of each species to the total average (dis-) similarity between different groups and

Table 2.1 Sample number, geographical position, water depth, region, mound region and facies of the investigated samples.

Station	Latitude	Longitude	Depth [m]	Region	Mound Region	Facies
GeoB 9220	51°26.69'N	11°45.04'W	892	Porcupine Seabight	Belgica	sandwave
GeoB 9209-2	51°26.89'N	11°45.81'W	982	Porcupine Seabight	Belgica	sandwave
GeoB 9204-1	51°26.94'N	11°45.16'W	838	Porcupine Seabight	Belgica	living coral
GeoB 9205-1	51°27.04'N	11°45.12'W	810	Porcupine Seabight	Belgica	living coral
GeoB 9219-1	51°27.05'N	11°45.40'W	920	Porcupine Seabight	Belgica	living coral
GeoB 9216-1	51°27.09'N	11°44.81'W	890	Porcupine Seabight	Belgica	living coral
M07-21	51°58.65'N	14°59.18'W	627	Porcupine Seabight	SW Porcupine Bank	dead coral
M07-23	51°59.54'N	14°59.05'W	721	Porcupine Seabight	SW Porcupine Bank	off-mound
GeoB 6721-1	52°09.22'N	12°46.31'W	696	Porcupine Seabight	Hovland	off-mound
P292/581-1	52°13.40'N	12°50.24'W	736	Porcupine Seabight		off-mound
M07-24	52°18.86'N	12°40.78'W	663	Porcupine Seabight		off-mound
M07-25	52°18.87'N	12°42.42'W	647	Porcupine Seabight	Magellan	dead coral
P292/580-1	52°20.46'N	12°56.72'W	630	Porcupine Seabight	Magellan	off-mound
P292/579-1	52°23.62'N	13°01.56'W	554	Porcupine Seabight		off-mound
P292/578-1	52°35.67'N	13°10.74'W	450	Porcupine Seabight		off-mound
P292/577-1	52°41.78'N	13°16.43'W	356	Porcupine Seabight		off-mound
P292/576-1	52°51.40'N	13°25.39'W	249	Porcupine Seabight		off-mound
P292/574-1	53°00.80'N	13°34.04'W	202	Porcupine Seabight		off-mound
GeoB 9287	53°30.91'N	14°21.16'W	696	Porcupine Bank	N Porcupine Bank	living coral
GeoB 9288	53°31.06'N	14°21.77'W	870	Porcupine Bank	N Porcupine Bank	dropstones
M07-15	55°29.18'N	16°08.24'W	552	Rockall Bank	SE Rockall Trough	living coral
GeoB 9271	56°29.58'N	17°18.16'W	664	Rockall Bank	SW Rockall Bank	dropstones
GeoB 9260	56°29.98'N	17°18.63'W	683	Rockall Bank	SW Rockall Bank	dropstones
GeoB 9257	56°30.13'N	17°17.77'W	678	Rockall Bank	SW Rockall Bank	dropstones
GeoB 9256-1	56°30.20'N	17°18.37'W	629	Rockall Bank	SW Rockall Bank	dead coral
GeoB 9269	56°30.28'N	17°17.63'W	686	Rockall Bank	SW Rockall Bank	dropstones
GeoB 9268	56°30.44'N	17°18.62'W	656	Rockall Bank	SW Rockall Bank	dead coral

within one group (e.g., Basso and Spezzaferri 2000; Clarke and Warwick 2001; Kruskal 1964; Kruskal and Wish 1978;). Taxonomic notes, census data, and compositional data of the benthic foraminiferal taxa used for multivariate statistic are given in Appendix A.

2.4 RESULTS

2.4.1 Facies description

Field observations (video images taken on board with underwater video camera, ROV camera, and core description) and sediments recovered in core samples allowed to identify five different facies types (Tab. 2.1):

- 1) *Off-Mound Facies*: the distal parts of the cold-water carbonate mounds, the fine-grained pelagic sediments contain sand-sized benthic and planktonic foraminifera, echinoids, molluscs, and terrigenous components.
- 2) *Dropstone Facies*: large mounds (e.g., Galway Mound) are flanked by erosional moats formed by bottom currents (De Mol et al. 2002). These moats consist of sandy and silty sediments, sometimes rippled, containing dropstones up to 10 cm in diameter. Fragments of bryozoans, molluscs, and corals, generally characterize this facies.
- 3) *Dead Coral Facies*: this facies consists mainly of dead coral debris accumulating at the lower flanks and/or base of the mounds. Fine-grained sediments, mud and silt, trapped by the coral debris, are deposited to form a soft-sediment substratum containing pieces of molluscs and echinoids. The amount of coral debris decreases towards the distal parts of this facies.
- 4) *Living Coral Facies*: mainly developed on the upper flanks of the mounds and partially on their summits, this facies consists of a dense cover of living cold-water corals, mainly *Lophelia pertusa* and *Madrepora oculata*. Corals may colonize a hard substratum composed of dead coral debris or coarse sand. Also in this facies, muddy and silty sediments are trapped by the coral build-ups. However, the trapped sediments can be finer than in the *Dead Coral Facies*.
- 5) *Sandwave Facies*: the sandwave facies can co-exist with all the other facies. However, this facies is generally developed on the mound flanks

(e.g., Galway and Franken Mounds) or close to the summit (Propeller Mound) within or close to the *Living Coral Facies* (Rüggeberg et al. 2007). Single branches or small patches of living cold-water corals colonize the sandy sediments and/or fragments of dead corals.

2.4.2 Benthic foraminiferal assemblages in surface sediments

A total of 124 total (unstained) benthic foraminiferal species belonging to 75 genera were identified in the studied region (Appendix A). The samples collected along the bathymetric transect (water-depths between 202 and 736 m) from the Porcupine Bank into the Seabight document the variation of benthic foraminiferal assemblages with depth in this area (Fig. 2.1, 2.2; Tab. 2.1).

In the shallowest part of the Porcupine Bank (202 m, 249 m, and 356 m) *Cassidulina carinata* dominates the assemblage (50 % at 202 m, 24.9 % at 249 m, and 10.7 % at 356 m, respectively). The accompanying species at 202 m include abundant *Globocassidulina subglobosa* (up to 14.9 %), and at 249 m abundant *Cibicidoides pachyderma* (20.5 %). The abundances of uvigerinids increase with increasing depth reaching a maximum of 40.6 % at 554 m, but they show abundances still around ~30 % down to 736 m. *Hyalinea balthica* and *Bulimina marginata* are abundant in the whole transect with a maximum between 356 and 630 m water depth. Below 450 m water depth *Melonis barleeanum*, *Epistominella vitrea* and *Epistominella exigua* are important representatives of the community. Other deep-water taxa are present in lower amounts. For example, the highest abundance of *Angulogerina angulosa* (9.3 %) is recorded at 356 m. Sample GeoB6721-1 is located within the Hovland mound region on the summit of a cold-water coral mound. In comparison to other mounds in this region the sediment does not contain coral rubble, dropstones or sandwaves, indicating no recent settling of *Lophelia pertusa*. Therefore we classify it as off-mound sediment but relatively close to cold-water corals. Transport could be the reason for the abundance of epifaunal attached foraminiferal species like *Planulina ariminensis*, *Discanomalina coronata*, *Cibicidoides ungerianus*, and *Fontbotia wuellersdorfi*.

2.4.3 Statistical treatment

Since patterns of community structures are often not readily apparent (Clark and Warwick, 2001), we have applied the Bray-Curtis Similarity clustering to our data, which is an excellent method to treat community data in order to highlight patterns in benthic

foraminiferal distribution (Kenkel and Orloci, 1986). At the 100% of Bray-Curtis similarity, 5 clusters (1 to 5) can be distinguished. On the basis of the same similarity matrix samples are ordinated by non-metric MultiDimensional - nMDS - (Kruskal, 1964; Kruskal and Wish, 1978), with a stress value of 0.19 (Fig. 2.3, Tab. 2.3).

Cluster 1 groups samples 574-1, 576-1, 577-1, 578-1, 579-1, 580-1, 581-1, GeoB6721-1, M07-23, and M07-25 (Bray-Curtis Similarity 49 %). Eleven species and/or groups account for 90.4 % of the average similarity of this cluster (Tab. 2.3). Cluster 2 combines samples 9257, 9269, 9260, 9271, and 9288 (Bray-Curtis Similarity 58 %). Ten species and/or groups account for 91.1 % of the average similarity of this cluster (Tab. 2.3). Cluster 3 groups samples M07-21, M07-24, 9256-1, and 9268 (Bray-Curtis Similarity 51.1 %). Eleven species and/or groups account for 90.2 % of the average similarity of this cluster (Tab. 2.3). Cluster 4 assembles samples 9204-1, 9205-1, 9216-1, 9219-1, 9287, and M07-15 (Bray-Curtis Similarity 51.1 %). Fourteen species and/or groups account for 90.9 % of the average similarity of this cluster (Tab. 2.3). Cluster 5 combines only two samples, 9209-2 and 9220 (Bray-

Curtis Similarity 49.5%). Eleven species and/or groups account for 90.4 % of the average similarity of this cluster (Tab. 2.3).

2.5 DISCUSSION

2.5.1 Potential indicator species

We investigated total (unstained) assemblages from surface sediments recovered in the Porcupine Seabight and Rockall Bank carbonate mound systems. A total assemblage may result from complex interaction of environmental parameters, including the original microhabitat of the living forms, reworking, winnowing, dissolution, and bacterial decomposition of organic test components, which create differences between the live and dead components (Licari and Mackensen, 2005; Loubere, 1989; Mackensen and Douglas, 1989). For this reason we probably miss some of the monothalamous and soft-shelled foraminiferal species (Gooday and Hughes, 2002). Nevertheless, the aim of this study is to define benthic foraminiferal assemblages that can help to identify different mound facies in the paleo-record. We therefore exclude soft-shelled species since they are rarely preserved in the

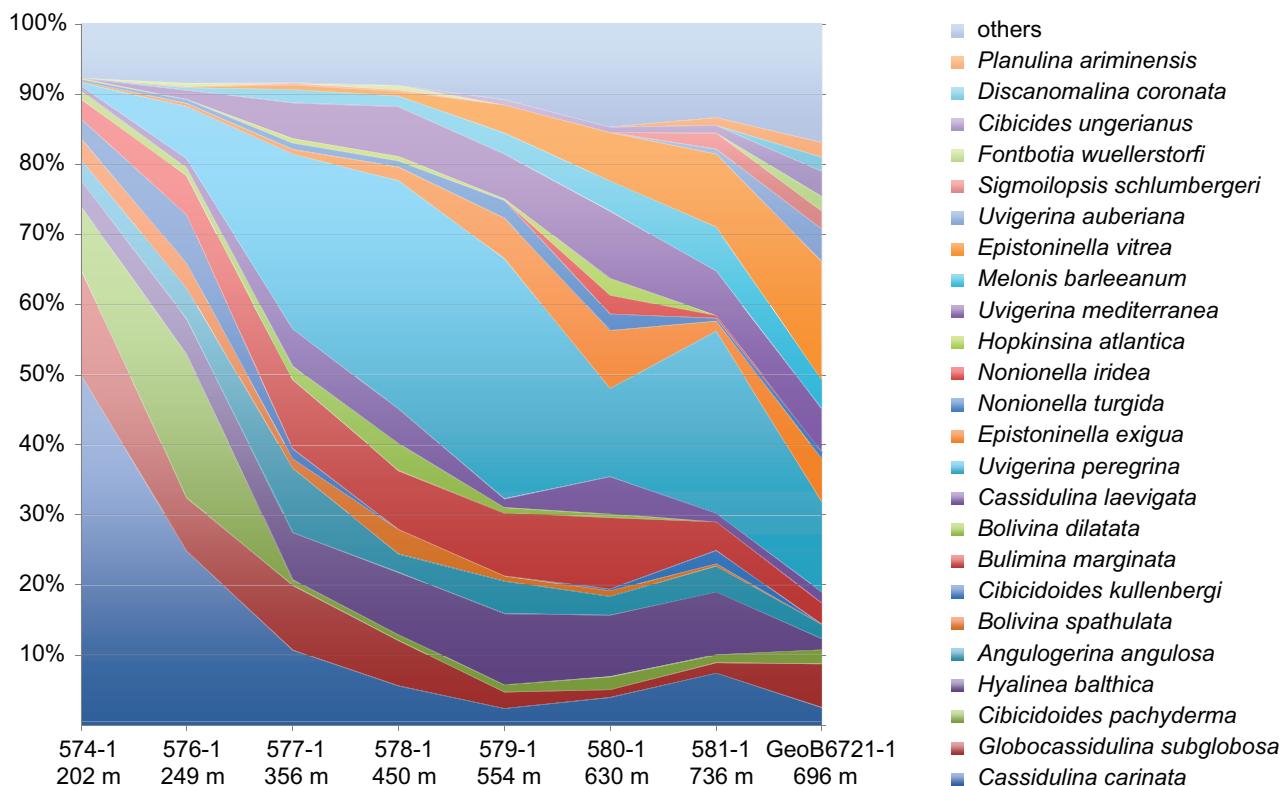


Figure 2.2 Percent distribution of the most abundant species (>2%) along a bathymetric transect comprising 8 off-mound-station in the Porcupine Seabight.

fossil record. Therefore, although, monothalamous and soft-shelled species are very important components of cold-water coral ecosystems, they are rarely preserved in the fossil record and are not treated in the present study.

2.5.2 Environmental interpretation

The spatial distribution of benthic foraminifera is controlled by a combination of environmental parameters (e.g., Gooday 2003; Grimsdale and Van Morkhoven 1955; Gupta 1999; Jorissen et al. 2007; Lutze and Coulbourn 1984; Murray 1991, 2006; Natland 1933; Pflium and Frerichs 1976; Schönfeld 2002a, b; Van der Zwaan et al. 1999; Tab. 2.2). Among the most important parameters controlling the distribution of these organisms in the deep sea are the organic flux to the seafloor and the bottom water oxygenation (e.g. Jorissen 1988; Lutze and Coulbourn 1984; Mackensen et al. 1990, 1995; Miller and Lohmann 1982; Schmiedl and Mackensen 1997). The quality of the organic matter, and the degree of seasonality in its delivery to

the seafloor, are also important (Fontanier et al. 2002, 2005). Where current velocities are high, bottom flow, together with substrate characteristics, play a central role in the distribution of benthic foraminiferal assemblages (Schönfeld 1997, 2002a, b).

Weston (1985) compared living and dead foraminiferal assemblages ($>125 \mu\text{m}$) from surface samples south of $51^{\circ} 50' \text{N}$ in the Porcupine Seabight, the area where big cold-water coral mounds on elevated substrates were originally discovered (Hovland et al. 1994; Rüggeberg et al. 2007). The assemblages of Weston (1985) show zonation with respect to water-depth. In particular, Weston found high numbers of *Bulimina marginata* and *Cassidulina carinata* above 700 m water depth and attached living species, e.g. *Lobatula lobatula*, *Cibicides refulgens*, below this depth. In the area that we investigated, benthic foraminiferal assemblages are dominated by infaunal species. In particular, *Globocassidulina subglobosa*, *Uvigerina peregrina*, *Uvigerina mediterranea*, *Epistominella*

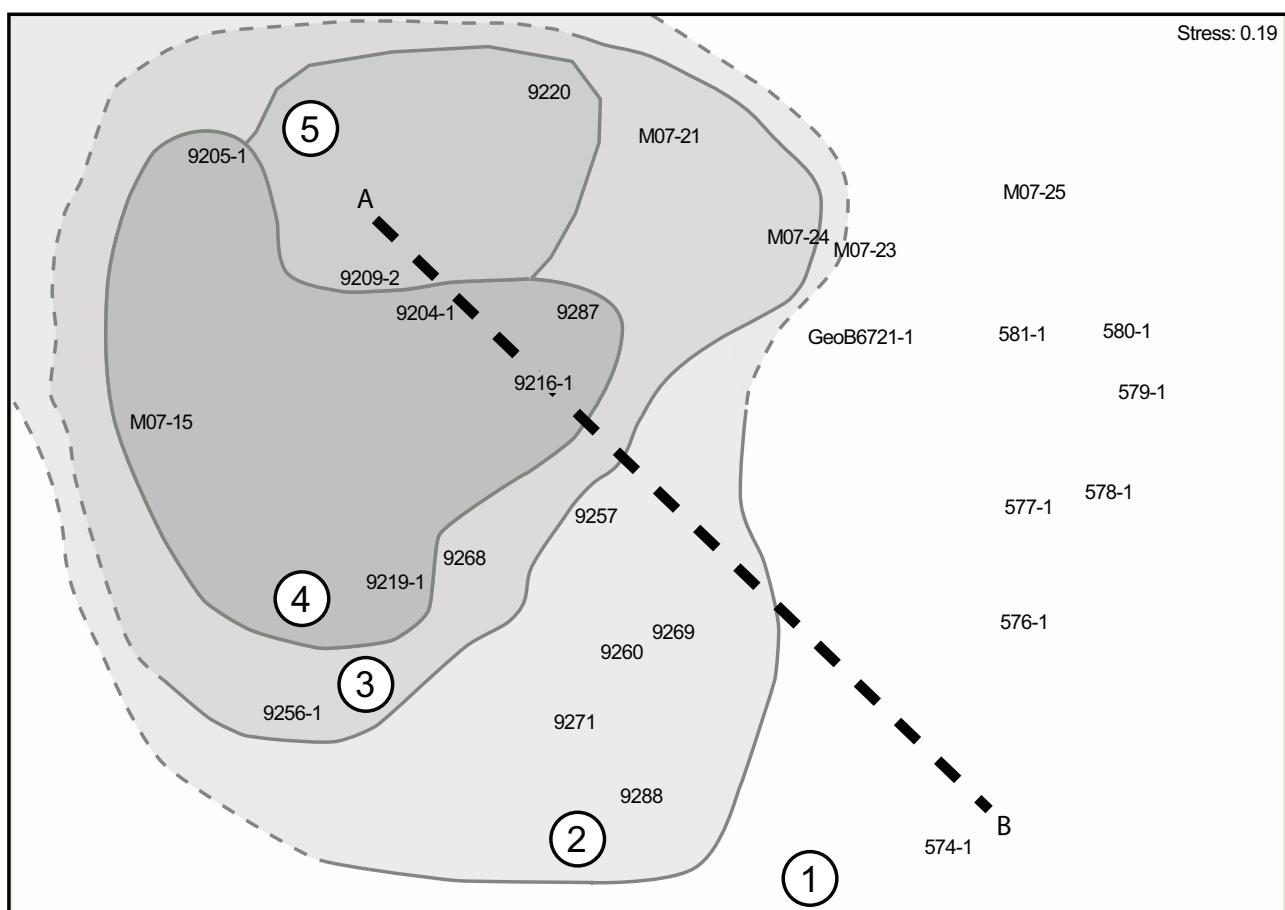


Figure 2.3 Non-metric MultiDimensional Scaling (nMDS) plot obtained from Bray-Curties similarity matrix of benthic foraminifera compositional data. The different clusters (1 to 5) are illustrated in different greyscales. The cluster arrangement corresponds to the facies distribution of an active cold-water coral mound. Transect A–B is illustrated in figure 2.4. The stress represents the distortion involved in compressing the data from a multidimensional space into a smaller number of dimensions (Field et al. 1982).

Table 2.2 Living strategy, preferred substratum and other ecological attributes of selected benthic foraminifera according to the literature. The five facies identified in this study are also plotted.

Species	facies	Living strategy	Preferred substratum	Comments	Reference
<i>Hyalinea bathica</i>	off-mound	epifaunal	mud to silt	opportunistic, phytodetritus feeder, seasonal food fluxes, large tolerance to varying organic flux	Gooday, 1988; Gooday et al. 1993; Loubere and Fariduddin, 1999; Murray, 2006; Smart, 2008; Smart et al. 1994; Thomas et al. 1995; Thomas and Gooday, 1996
<i>Epistominella exigua</i>	off-mound	epifaunal, shallow infaunal	mud	passive suspension feeder, deep sea habitat, oligotrophic environment, oxic, stable physico-chemical conditions	Fariduddin and Loubere, 1997; Jorissen et al. 1998; Morigi et al., 2001; Murray, 2006; Schmiedl et al. 2000; Woodruff et al. 1980
<i>Cibicidoides kullenbergi</i>	off-mound	shallow infaunal	mud	rich supply on labile organic matter, superior competitor in food- and oxygen-rich sediments, less tolerant to suboxic conditions than <i>U. peregrina</i>	Altenbach et al. 1999; Altenbach and Santhein, 1989; Fontanier et al. 2002
<i>Uvigerina mediterranea</i>	off-mound	shallow infaunal	mud	rich supply on labile organic matter and high concentration of bacteria	Altenbach et al. 1999; De Stigter et al. 1998; Fontanier et al. 2002
<i>Uvigerina peregrina</i>	off-mound	infaunal	mud	opportunist, suboxic, reacts fast to phytodetritus blooms	Duchemin et al. 2007; Murray, 2006
<i>Nanionella iridea</i>	off-mound	infaunal	mud	suboxic, disoxic	Kaiho, 1994; Murray, 2006
<i>Nanionella turrida</i>	off-mound	infaunal	mud to fine sand	high carbon flux rates; tolerates low oxygen	Brüchert et al. 2000; Mackensen et al. 1993; Mackensen et al. 1990; Mullins et al. 1985; Murray, 2006; Seidenkrantz et al. 2000
<i>Buliminina</i> spp.	dead coral facies	epifaunal attached	hard substrates	oxic suspension feeder	Murray, 2006; Spezzaferri and Goris, 2001
<i>Hanzawaia boueana</i>	dead coral facies	epifaunal attached	hard substrates	high energy areas, can withstand permanent winnowing and redeposition	Lütze and Thiel, 1987; Schönenfeld, 2002a
<i>Planulina ariminensis</i>	dead coral facies	epifaunal, attached	sand	opportunist, high food availability, low water energy, hypoxic	Gross, 2000
<i>Spiroreticinella Wrightii</i>	dead coral facies	epifaunal, attached	sand	high and Voren, 1984; Jarke, 1980; Mackensen et al. 1985; Murray 1971; Qvale and Van Weering, 1985; Schönenfeld, 2002a; Seijrup et al. 1981	Hald and Voren, 1984; Jarke, 1980; Mackensen et al. 1985; Murray 1971; Qvale and Van Weering, 1985; Schönenfeld, 2002a; Seijrup et al. 1981
<i>Angulogerina angulosa</i>	dead coral facies	infaunal	mud	opportunist, high food availability, low water energy, hypoxic	Jorissen et al. 1992; Murray, 2006; Osterman et al. 2005
<i>Epistominella vitrea</i>	dead coral facies	infaunal	hard substrates	Murray, 2006	
<i>Gavelinopsis praegeri</i>	living coral facies	epifaunal attached, mobile	hard substrates, coarse sediments	suspension feeder, high energy areas	Lütze and Thiel, 1989; Murray, 1971; Murray, 2006; Schönenfeld, 2002a
<i>Lobatula lobatula</i>	living coral facies	epifaunal, attached	hard substrates	opportunist, moderate to high carbon flux rates, preference for high quality food	Altenbach et al. 1999; Gupta and Thomas 1999; Hess et al. 2005; Jorissen et al. 2007; Nomura 1983a, b
<i>Cassidulina carinata</i>	living coral facies, off-mound	shallow infaunal, epifaunal	sand	high carbon flux rates, tolerates low oxygen	Mackensen and Hald, 1988; Murray, 2003; Murray, 2006
<i>Cassidulina laevigata</i>	living coral facies, off-mound	infaunal	mud	high carbon flux rates; tolerates low oxygen	Loubere, 1996; Mackensen et al. 1995; Mullins et al. 1985; Murray, 2006; Seidenkrantz et al. 2000
<i>Bulinina</i> spp.	living coral facies	infaunal	mud	high carbon flux rates, low oxygen, variable food flux	Murray, 2006
<i>Pulnella</i> spp.	living coral facies	infaunal	mud	facultative anaerobic phytodetritus feeder, preferentially ingest fresh diatoms, oligotrophic	Conilis and Chen, 1988; Gupta and Thomas, 1999; Loubere, 1998; Conilis, 1979; Fariduddin and Loubere, 1997; Gooday, 1994; Mackensen et al. 1995; Murray, 2006; Suri et al. 2003
<i>Trifarina brady</i>	living coral facies	infaunal	mud	strong bottom currents	Hawkes and Scott, 2005; Schönenfeld, 1997, 2002a, b
<i>Globocassidulina subglobosa</i>	living coral facies, dropstone facies	infaunal	hard substrates	passive suspension feeder, high energy, oxic, stable physico-chemical conditions	Zwaan, 1982; Murray, 2006
<i>Discanomalina coronata</i>	sandwave facies	epifaunal attached	hard substrates	passive suspension feeder, high energy, oxic, stable physico-chemical conditions	Kaiho, 1994, 1999; Kouwenhoven, 2000; Murray, 2006; Van der Zwaan, 1982
<i>Cibicides refugens</i>	living coral facies	epifaunal attached	hard substrates	passive suspension feeder, high energy, oxic, stable physico-chemical conditions	Jorissen et al. 2007; Kaiho, 1994, 1999; Kouwenhoven, 2000 Murray, 2006; Van der Zwaan, 1982
<i>Astronion</i> spp.	sandwave facies	epifaunal attached, infaunal	mud	passive suspension feeder, preferentially ingest fresh diatoms, oligotrophic environment, high energy, oxic, stable physico-chemical conditions	Almogi-Labin et al. 2000; Miao & Thunell, 1993; Murray, 2006; Schmiedl et al. 2000
<i>Cibicides ungernius</i>	sandwave facies	epifaunal, shallow infaunal	mud	eutrophic environments, does not depend on labile organic matter	De Nooijer et al. 2008; Duijnstee et al. 2003; Ernst 2002; Ernst et al. 2005
<i>Cibicides pachyderma</i>	sandwave facies	epifaunal, shallow infaunal	fine sand	<10°C, high POM, lives in high productivity waters, lives on the redox front	Murray, 2006
<i>Eggerella scabra</i>	sandwave facies	infaunal	mud	suboxic, disoxic	Carap, 1989; Corliss, 1985; Gooday, 1986; Koho et al. 2008; Loubere, 1991; Murray, 2006
<i>Kamerella brady</i>	sandwave facies	epifaunal	mud	Mullins et al. 1985; Murray, 2006	
<i>Melonis barleeanum</i>	off-mound	infaunal	mud		
<i>Gyroidina soldanii</i>	sandwave facies	epifaunal	mud		

vitrea, *Cassidulina carinata*, *Cassidulina laevigata*, *Angulogerina angulosa*, and *Buliminia marginata* are generally very abundant (Appendix A). Clear depth-related patterns in these assemblages, of the kind

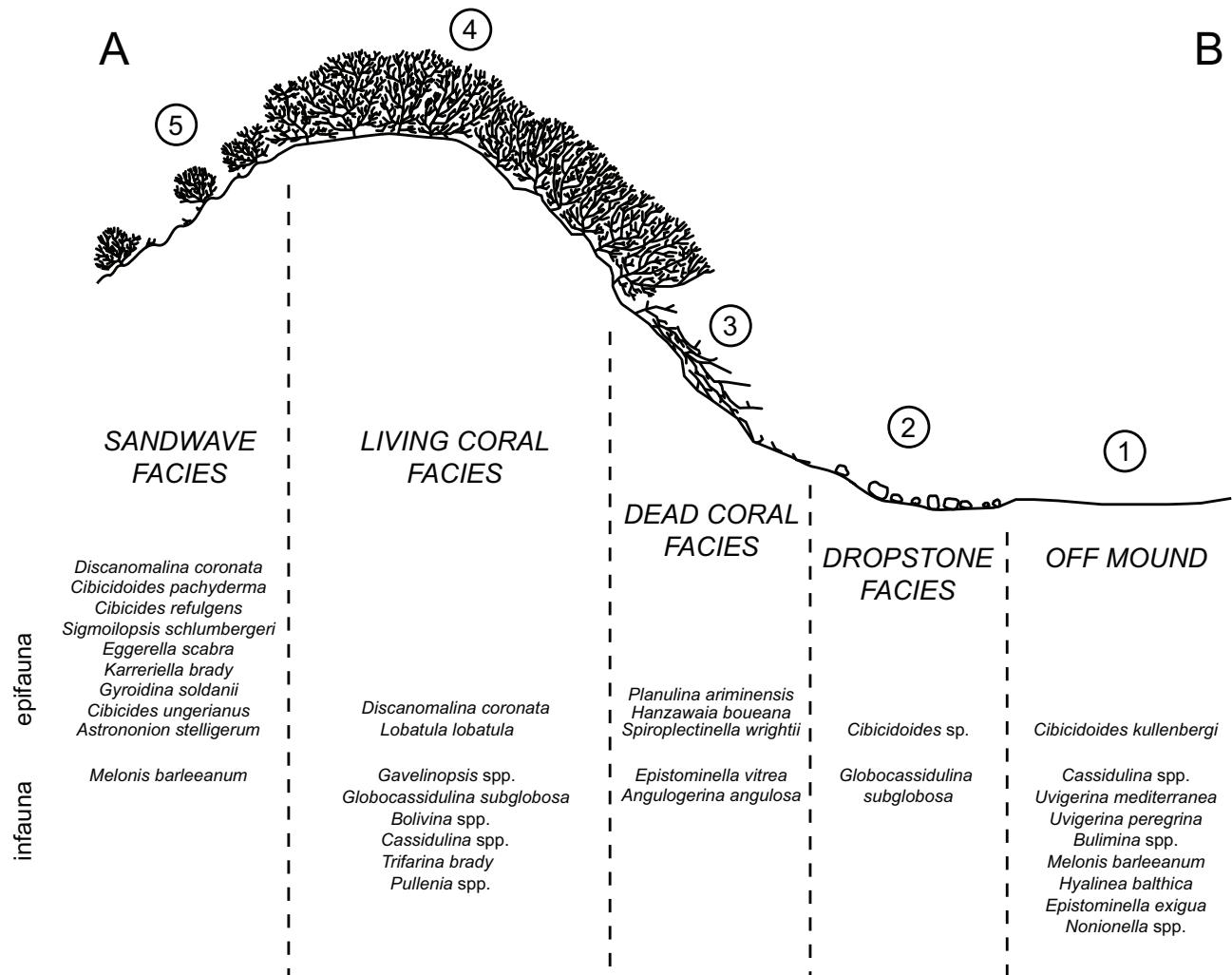


Figure 2.4 Model representing the distribution of benthic foraminiferal assemblages, based on the nMDS ordination, along the A-B transect. Numbers 1 to 5 refer to the five clusters estimated in the nMDS and representing the different facies on an active cold-water coral mound. For each cluster the most important benthic foraminifers are listed according to their living strategy and preference (Tab. 2.3).

display the highest abundance in off-mound surface sediments, whereas *G. subglobosa* and *E. vitrea* dominate on-mound. The most remarkable feature is the high occurrence of the epibenthic species *Discanomalina coronata* in on-mound sediments and its absence in the off-mound sediments.

The nMDS plot provides an overview of the similarities between assemblages (Clarke and Warwick, 2001; Everitt, 1978). This method is particularly suitable for the analysis of coral mound faunas, where the boundaries between different facies are often gradual (Clarke and Warwick, 2001). Our samples have been

observed by Weston (1985), are not obvious (Fig. 2.1, Appendix A). Instead, benthic foraminiferal distribution patterns reveal a link between assemblages and facies. In particular, the infaunal uvigerinids and cassidulinids

collected from different mounds at different water depth and from different facies in the Porcupine-Rockall Bank region (Tab. 2.1). The clustering of samples in the nMDS plot based on species corresponds to the distribution of facies on a typical mound in this area (e.g., Galway Mound, Fig. 2.4). The nMDS ordination (Fig. 2.3) and SIMPER analysis (Tab. 2.3; 2.4), together with information about the ecology of individual species and species groups (e.g., Tab. 2.2), leads to the following interpretations of the five clusters.

Cluster 1 groups 10 samples from the *Off-Mound Facies* (Fig. 2.3; Tab. 2.2; 2.3; 2.4). The sediment

Table 2.3 List of species and statistical parameters associated with the similarity in cluster 1 to 5. Average similarity within the group of station, average abundance, average similarity, contribution (%), and cumulative contribution (%) are given for each species with respect to the total similarity for each cluster.

Group 1				
Average similarity: 49.49%				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>U. peregrina</i>	17.69	11.04	22.31	22.31
<i>Cassidulina</i> spp.	14.51	7.55	15.26	37.56
<i>Bulimina</i> spp.	7.18	4.97	10.04	47.6
<i>H. balthica</i>	6.49	4.6	9.29	56.89
<i>U. mediterranea</i>	10.09	4.52	9.12	66.01
<i>A. angulosa</i>	4.05	2.9	5.87	71.88
<i>G. subglobosa</i>	5.39	2.8	5.67	77.55
<i>E. vitrea</i>	5.92	2.26	4.56	82.11
<i>Bolivina</i> spp.	3.45	1.94	3.92	86.03
<i>M. barleeanum</i>	2.53	1.29	2.6	88.63
<i>C. pachyderma</i>	3.73	0.87	1.75	90.38
Group 2				
Average similarity: 57.80%				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>G. subglobosa</i>	29.74	25.51	44.14	44.14
<i>E. vitrea</i>	10.25	6.34	10.97	55.1
<i>Cibicidoides</i> sp.	7.47	4.98	8.62	63.72
<i>Cassidulina</i> spp.	8.13	3.99	6.9	70.62
<i>Bulimina</i> spp.	4.55	3.37	5.83	76.45
<i>A. angulosa</i>	4.92	2.97	5.13	81.59
<i>C. pachyderma</i>	2.46	1.67	2.89	84.48
<i>U. peregrina</i>	4.44	1.45	2.51	86.99
<i>L. lobatula</i>	1.56	1.26	2.18	89.16
<i>P. ariminensis</i>	2.24	1.1	1.91	91.07
Group 3				
Average similarity: 51.10%				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>E. vitrea</i>	16.88	13.16	25.75	25.75
<i>G. subglobosa</i>	20.52	12.88	25.21	50.97
<i>A. angulosa</i>	9.16	4.79	9.38	60.35
<i>Cassidulina</i> spp.	6.37	4.08	7.99	68.34
<i>P. ariminensis</i>	4.62	2.46	4.81	73.15
<i>H. boueana</i>	2.54	1.93	3.78	76.93
<i>Discanomalina</i> spp.	2.64	1.62	3.16	80.09
<i>U. peregrina</i>	4.65	1.59	3.12	83.21
<i>U. mediterranea</i>	3.14	1.51	2.96	86.17
<i>Bulimina</i> spp.	2.89	1.06	2.08	88.25
<i>L. lobatula</i>	1.44	0.99	1.94	90.19
Group 4				
Average similarity: 51.10%				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>G. subglobosa</i>	19.67	15.51	30.36	30.36
<i>Bolivina</i> spp.	10.15	7.47	14.62	44.98
<i>Cassidulina</i> spp.	11.09	7.33	14.35	59.33
<i>Discanomalina</i> spp.	9.03	4.28	8.38	67.71
<i>P. ariminensis</i>	3.02	1.89	3.71	71.41
<i>L. lobatula</i>	2.14	1.3	2.55	73.96
<i>Cibicidoides</i> sp.	2.13	1.29	2.53	76.49
<i>E. vitrea</i>	4.27	1.29	2.52	79.01
<i>A. stelligerum</i>	2.11	1.18	2.31	81.32
<i>Bulimina</i> spp.	2.48	1.17	2.28	83.61
<i>A. angulosa</i>	3.32	1.09	2.14	85.75
<i>U. peregrina</i>	2.13	0.91	1.79	87.53
<i>T. bradyi</i>	1.47	0.87	1.69	89.23
<i>Gavelinopsis</i> spp.	1.88	0.86	1.67	90.9
Group 5				
Average similarity: 49.19%				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>G. subglobosa</i>	12.1	11.29	22.95	22.95
<i>Discanomalina</i> spp.	14.11	10.48	21.31	44.26
<i>Cassidulina</i> spp.	10.08	5.65	11.48	55.74
<i>E. vitrea</i>	6.05	4.03	8.2	63.93
<i>A. stelligerum</i>	4.44	2.42	4.92	68.85
<i>Bolivina</i> spp.	3.63	2.42	4.92	73.77
<i>U. mediterranea</i>	4.84	1.61	3.28	77.05
<i>K. brady</i>	2.42	1.61	3.28	80.33
<i>L. lobatula</i>	2.42	1.61	3.28	83.61
<i>M. barleeanum</i>	2.02	1.61	3.28	86.89
<i>H. boueana</i>	1.61	1.61	3.28	90.16

Table 2.4 (continued)

Species	Average dissimilarity = 68.5					Species	Average dissimilarity = 49.5					
	Group 1		Group 5				Group 3		Group 2			
	Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%		Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%	
<i>U. peregrina</i>	17.7	0.4	8.7	12.7	12.7	<i>G. subglobosa</i>	20.5	29.7	6.5	13.1	13.1	
<i>Discanomalina</i> spp.	0.2	14.1	7.0	10.2	22.8	<i>E. vitrea</i>	16.9	10.3	4.2	8.4	21.5	
<i>Cassidulina</i> spp.	14.5	10.1	4.9	7.1	29.9	<i>Cibicidoides</i> sp.	0.2	7.5	3.7	7.4	28.8	
<i>C. pachyderma</i>	3.7	8.9	4.6	6.7	36.6	<i>A. angulosa</i>	9.2	4.9	3.3	6.6	35.5	
<i>U. mediterranea</i>	10.1	4.8	3.9	5.7	42.3	<i>U. mediterranea</i>	3.1	5.7	3.2	6.5	42.0	
<i>G. subglobosa</i>	5.4	12.1	3.6	5.3	47.6	<i>Cassidulina</i> spp.	6.4	8.1	2.9	5.9	47.9	
<i>H. balthica</i>	6.5	0.4	3.0	4.4	52.0	<i>U. peregrina</i>	4.7	4.4	2.4	4.9	52.8	
<i>Bulimina</i> spp.	7.2	1.2	3.0	4.4	56.4	<i>P. ariminensis</i>	4.6	2.2	1.6	3.3	56.1	
<i>E. vitrea</i>	5.9	6.1	2.7	3.9	60.3	<i>Bulimina</i> spp.	2.9	4.6	1.5	3.1	59.2	
<i>C. refulgens</i>	0.5	4.4	2.1	3.1	63.4	<i>S. wrightii</i>	2.6	0.0	1.3	2.7	61.8	
<i>A. angulosa</i>	4.1	0.0	2.0	3.0	66.4	<i>C. pachyderma</i>	0.0	2.5	1.2	2.5	64.3	
<i>A. stelligerum</i>	0.9	4.4	1.9	2.8	69.2	<i>Discanomalina</i> spp.	2.6	0.2	1.2	2.5	66.8	
<i>S. schlumbergeri</i>	0.7	3.2	1.6	2.4	71.5	<i>C. kullenbergi</i>	1.6	1.7	1.1	2.2	69.0	
<i>E. exigua</i>	2.5	0.0	1.3	1.9	73.4	<i>H. boueana</i>	2.5	0.5	1.0	2.1	71.1	
<i>Nonionella</i> spp.	2.5	0.0	1.2	1.8	75.2	<i>Bolivina</i> spp.	2.0	1.6	1.0	1.9	73.0	
<i>P. ariminensis</i>	0.3	2.4	1.2	1.8	76.9	<i>C. refulgens</i>	1.5	0.6	0.8	1.6	74.6	
<i>Bolivina</i> spp.	3.5	3.6	1.1	1.6	78.6	<i>M. barleeanum</i>	1.8	1.6	0.7	1.5	76.0	
<i>K. brady</i>	0.3	2.4	1.1	1.6	80.2	<i>E. scabra</i>	1.4	0.0	0.7	1.5	77.5	
						<i>A. stelligerum</i>	0.6	1.2	0.6	1.2	78.6	
						<i>Fissurina</i> spp.	0.8	0.8	0.6	1.2	79.8	
						<i>S. schlumbergeri</i>	0.0	1.0	0.5	1.1	80.8	
Species	Average dissimilarity = 52.4					Species	Average dissimilarity = 61.6					
	Group 4		Group 5				Group 5		Group 2			
	Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%		Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%	
<i>Discanomalina</i> spp.	9.0	14.1	4.8	9.1	9.1	<i>G. subglobosa</i>	12.1	29.7	8.8	14.3	14.3	
<i>C. pachyderma</i>	0.0	8.9	4.5	8.5	17.6	<i>Discanomalina</i> spp.	14.1	0.2	7.0	11.3	25.6	
<i>G. subglobosa</i>	19.7	12.1	4.4	8.4	26.0	<i>C. pachyderma</i>	8.9	2.5	4.4	7.2	32.8	
<i>Bolivina</i> spp.	10.2	3.6	3.5	6.6	32.6	<i>U. mediterranea</i>	4.8	5.7	3.7	6.0	38.8	
<i>Cassidulina</i> spp.	11.1	10.1	3.0	5.8	38.4	<i>Cassidulina</i> spp.	10.1	8.1	3.4	5.6	44.4	
<i>E. vitrea</i>	4.3	6.1	2.5	4.7	43.1	<i>Cibicidoides</i> sp.	0.8	7.5	3.3	5.4	49.8	
<i>U. mediterranea</i>	2.0	4.8	2.0	3.9	46.9	<i>E. vitrea</i>	6.1	10.3	3.2	5.2	55.0	
<i>C. refulgens</i>	1.2	4.4	2.0	3.9	50.8	<i>A. angulosa</i>	0.0	4.9	2.5	4.0	59.0	
<i>A. angulosa</i>	3.3	0.0	1.7	3.2	54.0	<i>C. refulgens</i>	4.4	0.6	2.1	3.5	62.4	
<i>S. schlumbergeri</i>	0.9	3.2	1.6	3.1	57.1	<i>U. peregrina</i>	0.4	4.4	2.1	3.4	65.8	
<i>A. stelligerum</i>	2.1	4.4	1.4	2.7	59.8	<i>Bulimina</i> spp.	1.2	4.6	1.7	2.7	68.6	
<i>P. ariminensis</i>	3.0	2.4	1.3	2.4	62.2	<i>S. schlumbergeri</i>	3.2	1.0	1.6	2.6	71.2	
<i>E. scabra</i>	0.0	2.0	1.0	1.9	64.1	<i>A. stelligerum</i>	4.4	1.2	1.6	2.6	73.8	
<i>U. peregrina</i>	2.1	0.4	1.0	1.9	66.0	<i>Bolivina</i> spp.	3.6	1.6	1.2	2.0	75.8	
<i>K. brady</i>	0.5	2.4	1.0	1.9	67.9	<i>P. ariminensis</i>	2.4	2.2	1.2	2.0	77.8	
<i>Cibicidoides</i> sp.	2.1	0.8	0.9	1.7	69.6	<i>E. scabra</i>	2.0	0.0	1.0	1.6	79.4	
<i>Bulimina</i> spp.	2.5	1.2	0.8	1.6	71.2	<i>K. brady</i>	2.4	0.5	1.0	1.5	80.9	
<i>G. soldanii</i>	0.0	1.6	0.8	1.5	72.7							
<i>Gavelinopsis</i> spp.	1.9	0.8	0.8	1.5	74.2							
<i>M. barleeanum</i>	0.7	2.0	0.8	1.4	75.6							
<i>M. pompilioides</i>	0.9	0.8	0.7	1.3	77.0							
<i>L. lobatula</i>	2.1	2.4	0.7	1.2	78.2							
<i>C. kullenbergi</i>	1.3	0.0	0.6	1.2	79.4							

varies from sandy silt to coarse sand. The water depths at which samples were taken range between 202 and 736 m (Tab. 2.1). Cluster 1 is represented by the infaunal species *U. peregrina*, *U. mediterranea*, *H. balthica*, *M. barleeanum*, *E. exigua* and *Cassidulina* spp. (*C. carinata* and *C. laevigata*), *Bulimina* spp. (*B. aculeata*, *B. marginata*, and *B. striata*), and *Nonionella* spp. (*N. iridea*, *N. labradorica*, and *N. turgida*). These species are known to be typical for muddy/silty to sandy substrata, to prefer high organic matter input, and to thrive under suboxic-dysoxic conditions (e.g., Fontanier et al. 2002; Jorissen et al. 2007; Murray, 2006; Schönfeld and Altenbach, 2005). In particular, uvigerinids require high quantity and quality of nutrients supplied to the seafloor (Jorissen et al. 1988, Schönfeld and Altenbach, 2005).

Cluster 2 combines 5 samples from the *Dropstone Facies*. Sediment consists of a sandy fraction with dropstones indicating strong bottom currents (De Mol et al. 2002). *Globocassidulina subglobosa* is highly abundant in these samples. This infaunal species (e.g., Fariduddin and Loubere, 1997; Murray, 2006) is characteristic for oligotrophic areas with high bottom current velocities (Jorissen, 1988; Mackensen et al. 1995). *Cibicidoides* sp., the dominant epibenthic species, appears to be specialised to this high-energy environment.

Clusters 3, 4, and 5 represent the facies types typical of cold-water carbonate mounds. The boundaries between these facies types are gradual. Some species are represented in all of them, although their relative

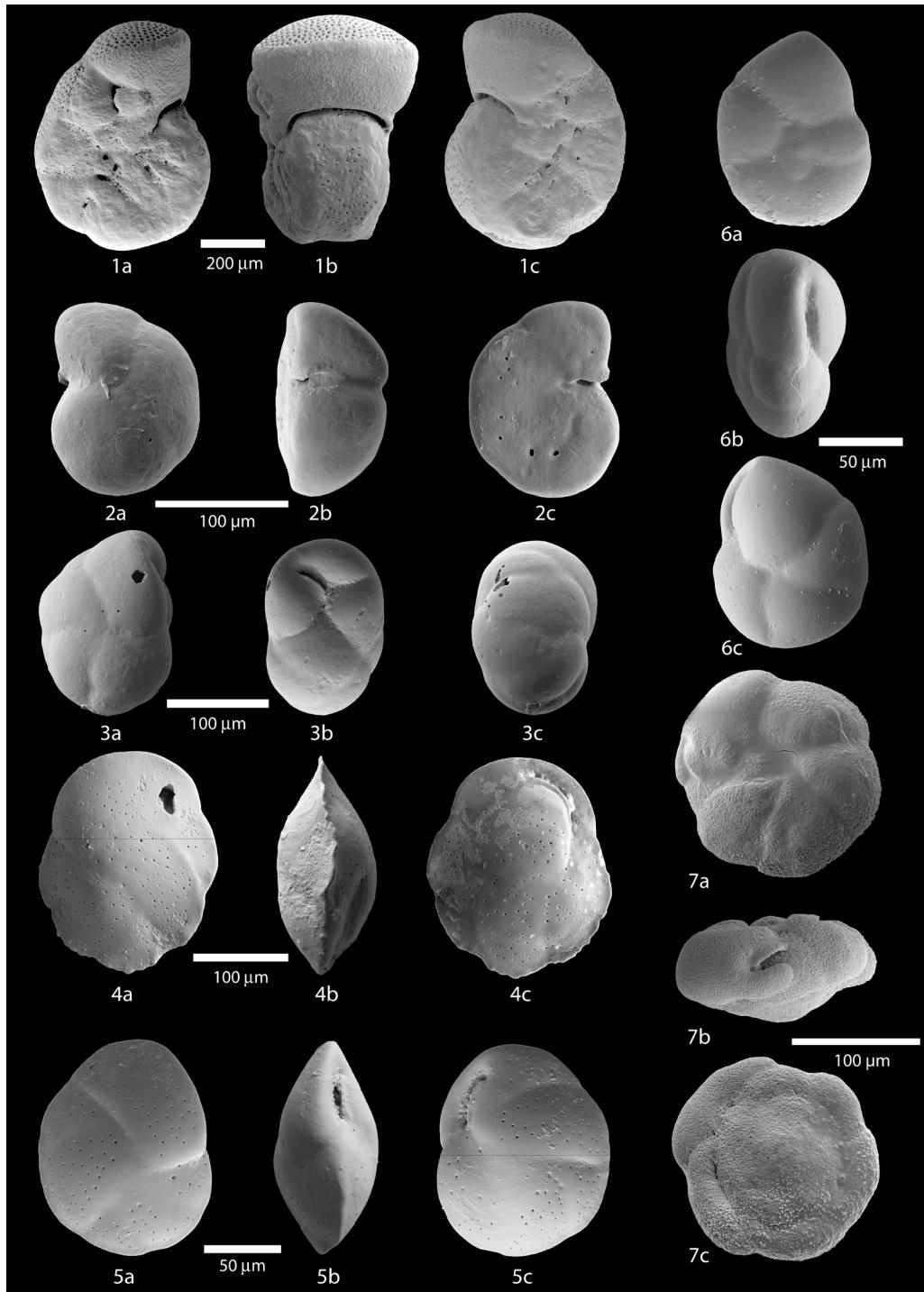


Figure 2.5 Important species collected on cold-water coral mounds at the Rockall and Porcupine Bank: (1a-c) *Discanomalina coronata* (Parker and Jones, 1857), sample M07-15. (2a-c) *Cibicidoides* sp., sample 9260. (3a-c) *Globocassidulina subglobosa* (Brady, 1881), sample 9260. (4a-c) *Cassidulina carinata* (Silvestri, 1896), sample 574-1. (5a-c) *Cassidulina laevigata* (d'Orbigny, 1826), sample 577-1. (6a-c) *Epistominella exigua* (Brady, 1884), sample 580-1. (7a-c) *Epistominella vitrea* (Parker, 1953), sample 9256-1. a = spiral views, b = side views, c = umbilical views.

abundance varies. Attached epifaunal foraminiferal species are considerably more abundant on- than off-mound.

Cluster 3 groups 4 samples from the *Dead Coral*

Facies. A variety of different sediment types are associated with this facies. The framework created by the coarse coral debris traps fine sediment. Coral debris offers substrates favourable for the proliferation of epifaunal-attached species such as *Planulina*

ariminensis, *Hanzawaia boueana*, and *Spiroplectinella wrightii* (Gross 2000; Lutze and Thiel 1989; Murray 2006). *Angulogerina angulosa* and *Epistominella vitrea* are very abundant in this facies. *Angulogerina angulosa*, a small infaunal species often associated with strong bottom currents, is typical for coarse grained sediments (Mackensen et al. 1985; Schönfeld 2002a). *Epistominella vitrea* is an opportunistic species that responds to high food availability and occurs in muddy sediments under hypoxic conditions (e.g., Jorissen et al. 1992; Murray 2006; Osterman et al. 2005). It dominates in microhabitats that are more protected from high current velocities (Mackensen et al. 1995).

Cluster 4 groups 6 samples from the *Living Coral Facies*. Living corals grow on debris and/or pebbly mud that is clogged by fine-grained sediments (mud and silt). Foraminiferal assemblages from this facies mainly comprise *G. subglobosa*, *Bolivina* spp., *Cassidulina* spp., *Trifarina brady*, and *Pullenia* spp., species that are generally considered to be infaunal (e.g., Gooday, 2003; Murray 2006; Jorissen et al. 2007). According to Jorissen (1988), *G. subglobosa* prefers more oligotrophic environments than the uvigerinids. Since *G. subglobosa* dominates the *Living Coral Facies* on-mound and uvigerinids dominate the *Off-Mound Facies* we can interpret this distribution pattern as related to the current hydrography. In particular, the higher current energy probably keeps the organic particles in suspension in the water column, and therefore unavailable to the infaunal species *G. subglobosa*. On the other hand corals feed on high amounts of fresh organic matter (Duineveld et al. 2007), which may influence the quantity and/or quality of the flux material reaching the sea floor.

The epifaunal-attached species in the *Living Coral Facies* consist of *Discanomalina coronata*, *Lobatula lobatula*, and *Gavelinopsis praegeri*, which are characteristic for active mounds (Rüggeberg et al. 2007). *Discanomalina coronata* lives attached to dead biogenic fragments that make up the substratum in the *Living Coral Facies*. The high density of *L. pertusa* in this facies offers wide habitats for *D. coronata*.

Cluster 5 groups 2 samples from the *Sandwave Facies*. In this facies sandwaves are the result of bottom currents. The sandwaves are colonized by patches of living corals and a highly diverse fauna, including hydroids, sponges, soft corals, and bivalves (Wienberg et al. 2007). Typical for this facies are high abundances of epifaunal species such as *Cibicidoides pachyderma*, *Karreriella brady*, *Gyroidina soldanii*,

Cibicidoides ungerianus, *Astrononion stelligerum* and epifaunal-attached forms like *Cibicides refulgens* (e.g., Miao and Thunell 1993; Murray 2006; Schmiedl et al. 2000). *Discanomalina coronata* is a very abundant component of this cluster and dominates the epifaunal group (Rüggeberg et al. 2007; Schönfeld 1997, 2002a). Infaunal species are dominated by *Globocassidulina subglobosa*, *Cassidulina* spp., *Melonis barleanum* and *Eggerella scabra*.

In summary, cold-water carbonate mounds generally develop in environments with high current velocities (e.g., De Mol et al. 2002, White 2005) and high nutrient flux (Kiriakoulakis et al. 2004). Various types of elevated substrata provide different ecological niches for epifaunal foraminifera. The epifauna in the *Living Coral Facies* has a distinctive composition compared to other facies. In particular, *D. coronata* is the dominant epifaunal species in the *Living Coral Facies* and in the *Sandwave Facies* and seems to occupy the ecological niches made available by dead fragments of cold-water corals (Jensen and Frederiksen 1992).

2.6 CONCLUSION

- (1) Benthic foraminiferal assemblages provide an independent tool to identify different facies of cold-water coral mounds in the Porcupine-Rockall Bank region. The *Off-Mound Facies* is dominated by uvigerinids and other infaunal species. The *Dropstone Facies* is characterized by both infaunal (*G. subglobosa*) and epifaunal-attached (*Cibicidoides* sp.) species. The *Dead Coral Facies* is characterized by a combination of infaunal (*A. angulosa*, *E. vitrea*) and epifaunal (*P. ariminensis*, *H. boueana*, *S. wrightii*) species. The *Living Coral Facies* is characterized by epifauna and infauna. The species typical of this facies is *D. coronata*. The *Sandwave Facies* contains high abundances of epifauna and *D. coronata*.
- (2) A potential problem in environmental interpretation is the fact that most of the indicator species (e.g., phytodetritus feeders) are small-sized (63-125 µm) and therefore have not been studied very often (Jorissen et al. 2007). Our results underline the importance of studying the small fraction to highlight the contribution of small-sized species.
- (3) *Discanomalina coronata* is present and abundant only in sediments associated with cold-water coral mounds and is not present off-mound. The

fact that it is mainly found in the *Living Coral* and *Sandwave Facies* makes this species useful as an indicator for active cold-water coral mounds.

- (4) Foraminiferal species associated with cold-water coral carbonate mounds provide potentially useful indicators for these environments in the fossil record.

3 - BENTHIC FORAMINIFERA ASSOCIATED TO NORWEGIAN COLD-WATER CORAL REEFS: TOWARDS THE ASSESSMENT OF REGIONAL BIOINDICATORS

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submitted to Marine Micropaleontology

ABSTRACT

Cold-water coral ecosystems have a worldwide occurrence. Along the European margin the species *Lophelia pertusa* in particular, forms widespread reefs. On the Norwegian shelf and upper slope from the Skagerrak (Oslofjord), mid Norwegian slope (Sula Reef, Røst Reef, Trænadjudpet) to the northern coral-reef areas (Korallen, Lophavet, Stjernsundet, Sveinsgrunnen), extensive reefs have settled on post-glacial structures. These ecosystems offer many ecological niches for benthic foraminifera. In this study we quantitatively investigated the total (unstained) planktonic and benthic foraminiferal assemblage from surface sediments (0–2 cm, > 63 µm size fraction) from the Norwegian cold-water coral reefs. Aim of the study is to highlight their distribution pattern and to identify and trace possible indicator species for these ecosystems in different regions. Based on diversity analysis, hierarchical cluster analysis, and multidimensional scaling (nMDS), discrete patterns in the planktonic and benthic foraminiferal distribution could be identified. Planktonic foraminifera can be grouped into two assemblages. One occurring on the shelf and slope, influenced by the North Atlantic Current and by seasonal phytoplankton blooms, the other one is influenced by the low salinity Norwegian Coastal Current and possibly also by seasonal phytoplankton blooms. The total (unstained) benthic species can be grouped into six different assemblages: I) deep-sea below 1800 m water depth; II) deep-sea between 1800 and 800 m water depth; III) off-reef Skagerrak; IV) on-reef Skagerrak; V) mid-Norwegian coral reefs, and VI) northern Norwegian coral reefs. This study reveals that benthic foraminifera associated with cold-water coral reefs indicate an environment characterized by high energy, well oxygenated waters and high organic matter supply, derived from seasonal phytoplankton blooms and reaching the sea floor. The benthic fauna associated with the coral reefs in the Skagerrak (Oslo Fjord) is depleted of phytodetritus-feeding species. This datum suggests that probably a low amount of labile organic matter and/or nutrients reached the sea floor making the environmental setting unfavourable for coral growth, thus explaining the high amounts of dead corals in this area. High diversity of benthic foraminifera characterizes “on-reef”, especially in the central reef parts, whereas the “off-reef” diversity is lower.

This study reveals that the relation of benthic foraminiferal assemblages with different sedimentary facies is not well expressed for Norwegian coral reefs as it is for cold-water coral reefs on the carbonate mounds along the Irish margin. This difference possibly results from small-scale facies changes within tens of metres in the Norwegian reefs. The abundance of the benthic species *Discanomalina coronata* is restricted to cold-water coral reefs and absent off-reef. A similar distribution has been observed for cold-water coral reefs associated to carbonate mounds along the Irish margin, thus *D. coronata* is here interpreted to have ecological requirement similar to cold-water corals.

3.1 INTRODUCTION

Cold-water corals (CWCs) were first described from Norway in the eighteenth century (Pontoppidan, 1755; Gunnerus, 1768). Since these pioneering investigations many CWC ecosystems have been reported from around the world and especially from along the European margins. Cold-water coral reefs

occur in the Mediterranean Sea (Zibrowius, 1980; Taviani et al., 2005; Margreth et al. in review.), the Gulf of Cadiz (e.g., Pinheiro et al., 2001; Somoza et al., 2001; Fouquet et al., 2008; Wienberg et al. 2009), along the Irish margin (e.g., Hovland et al., 1994; De Mol et al., 2002; Kenyon et al., 2003; Van Rooij et al., 2003) up to northern Norway (e.g., Mortensen et al., 1995; Freiwald et al., 1997; Hovland et al., 1998;

Freiwald et al., 1999; Fosså et al., 2002; Lindberg and Mienert, 2005).

Lophelia pertusa is the dominant species in these ecosystems. It is a colonial, reef forming scleractinian coral (Roberts et al., 2009). Along the Irish margin *L. pertusa* forms large coral carbonate mounds, which originated in the late Pliocene/early Pleistocene (De Mol et al., 2002; Kano et al., 2007, Raddatz et al., in review.). On the Norwegian continental shelf the *L. pertusa* colonies are of Holocene age and have been developing since the end of the last glacial stage (e.g., Freiwald et al., 1997, 1999; Lindberg et al., 2007). In this region the corals build elongated reef-like structures on elevated hard substrates and can reach heights of 40 m and lengths of several kilometres (Freiwald et al., 1999; Freiwald et al., 2002). Their distribution is controlled by pre-existing topographical heights on the sea floor, such as moraine ridges and iceberg plough mark levees (Freiwald et al., 1999; Mortensen et al., 2001; Freiwald et al., 2002; Fosså et al., 2005). Living corals colonize the top and the upper slopes of these structures and preferentially occur at water depths between 150 and 400 m with the exception of a few shallower occurrences in fjord settings. The steep flanks of the reefs are generally characterized by *in situ* dead corals, while bio-eroded coral rubble accumulates around the base. These reefs provide abundant and diverse microhabitats for benthic organisms (Mortensen et al., 1995).

The biodiversity in CWC-reefs is high compared to that of the surrounding pelagic environments (Henry and Roberts, 2007). This is especially the case for suspension and filter feeders such as sponges, molluscs, brachiopods, cnidarians, bryozoans, ophiuroids, and crinoids, which are typical for these ecosystems (Mortensen et al., 2001). A high diversity foraminiferal population occurs in these ecosystems. In particular, recent studies on CWC-mounds in the Rockall Bank and the Porcupine Seabight have shown that a characteristic benthic foraminiferal community with a high number of epifaunal-attached species colonizes these habitats (Rüggeberg et al., 2007; Margreth et al., 2009). Furthermore, the benthic species *Discanomalina coronata* is generally associated with living CWC-reefs in this region (Margreth et al., 2009).

The present study focuses on the foraminiferal assemblages associated with CWC-reefs from northern Norway down to the Oslofjord (Fig. 3.1). The aims of this study are (1) to compare on- and off-reef total (unstained) planktonic and benthic foraminiferal assemblage from different CWC-reefs; (2) to compare

the foraminiferal assemblages from the Rockall Bank and the Porcupine Seabight with those from the Norwegian margin; and (3) to identify foraminiferal indicator species for CWC-reefs along the Norwegian margin.

3.2 OCEANOGRAPHIC SETTINGS AND CORAL REEF OCCURRENCE

Along the Norwegian coast, surface water flow is controlled by two northward-trending current systems, the Norwegian Coastal Current (NCC) and further off the shelf the North Atlantic Current (NAC; Mork, 1981). The primary origins of the NCC are freshwater discharge from the Baltic Sea and freshwater runoff from Norway (Mork, 1981). The waters are low saline with a high temperature gradient between summer and winter (Sætre, 1999; Mitchelson-Jacob and Sundby, 2001). The core of the NAC flows along the shelf break and is characterized by the northward transport of warm water (6–14°C) to high latitudes. During winter the mixing of these two current systems, combined with the effects of wind and topography are responsible for seasonal upwelling and eddies. In summer the two current systems flow parallel and the differences in their respective salinities prevent mixing (Sætre, 1999; Mitchelson-Jacob and Sundby, 2001). Blindheim (1990) groups the NCC and the NAC current systems into the Norwegian Atlantic Current (NwAC), which flows above the Arctic Intermediate Water (AIW). The AIW is characterized by a salinity minimum at depths of ~ 600 m. The Norwegian Sea Deep Water (NSDW) and the Norwegian Sea Bottom Water (NSBW) currents with temperatures below 0°C govern the water masses on the deep continental slope (Peterson and Rooth, 1976).

Along the Norwegian margin CWC-reefs occur at different water depths. In the Oslofjord CWCs develop between 90 and 140 m. They thrive in the saline and well-oxygenated oceanic Atlantic Water Inflow (AWI), only a few tens of meters beneath the permanently brackish surface water layer of the Baltic Waters (Svansson, 1975; Wissak et al., 2005). At Sula Reef they occur at depths of 240–315 m, at Røst Reef at 300–400 m, and at the northern-most reefs near the Nordkapp they thrive in the NwAC at 140–320 m (Fosså et al., 2002; Freiwald et al., 2004; Rüggeberg et al., in review.; Tab. 3.1; Fig. 3.1).

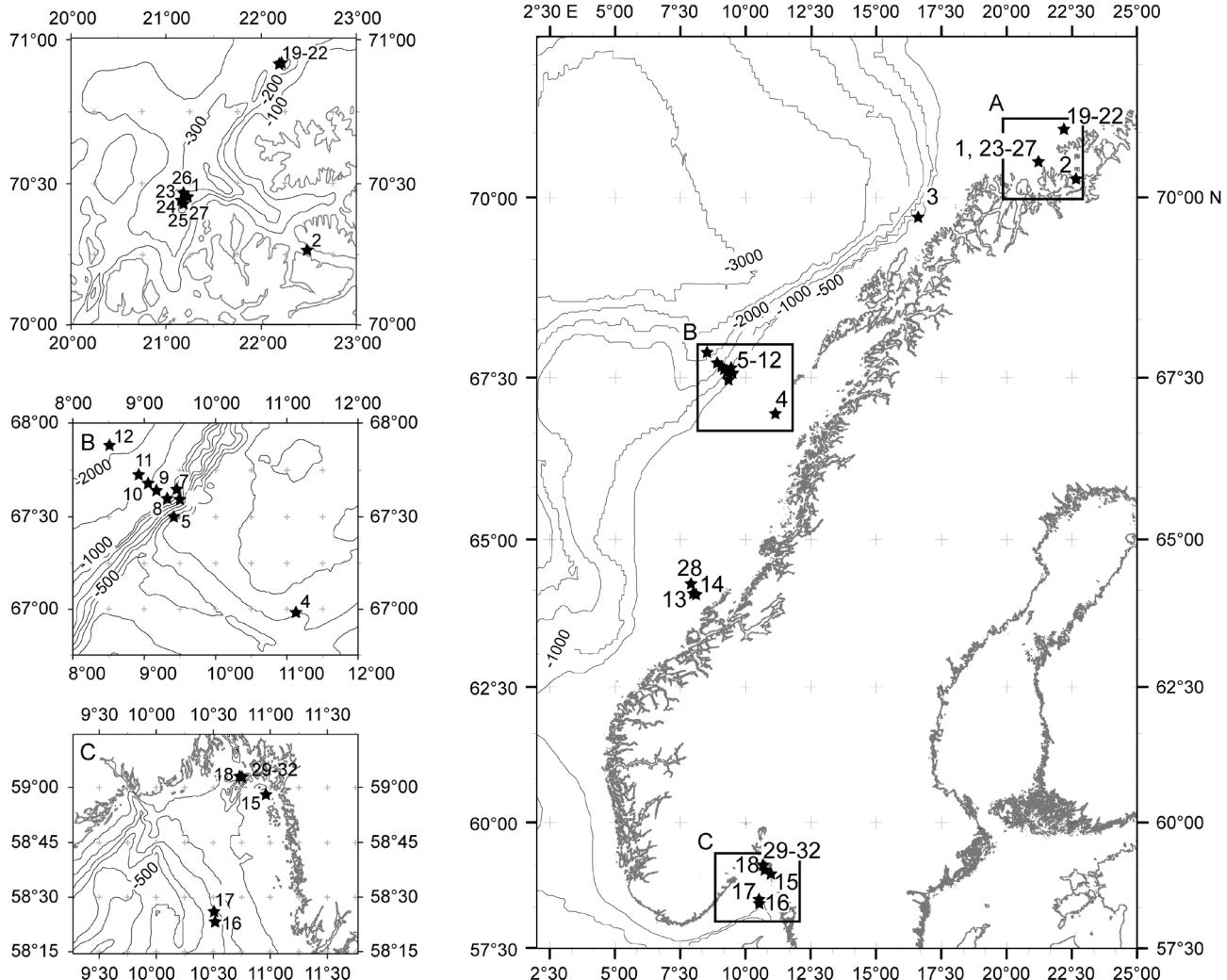


Figure 3.1 Bathymetry and location map of the surface sediment samples investigated in this study.

3.3 MATERIAL AND METHODS

Surface sediments were taken along the Norwegian margin (Fig. 3.1) using a grab sampler operated by the submarine JAGO, a Giant Box corer and a Van-Veen grabber during six different cruises (Tab. 3.1). Dead coral frameworks from steep reef flanks (inclinations $>40^\circ$) were not sampled. From the living coral reefs samples were taken only by submarine operated grab samplers to avoid damaging the reef. A few samples between the coral branches could be taken with a special submarine operated small sampler especially developed on board for this purpose. Surface sub-samples from the uppermost 2 cm were transferred to plastic bottles. Sediment samples for micropaleontological analyses were processed at the University of Fribourg following the method described in Spezzaferri and Coric (2001). Thirty-two samples were dried at room temperature, weighed, washed through a set of sieves ($63\mu\text{m}$, $125\mu\text{m}$, $250\mu\text{m}$), dried again at room temperature, and

weighed. The samples were split to obtain at least 300 dead (unstained) benthic and 900 planktonic specimens for each sample. In the case of samples containing less than these amounts, all specimens were counted. The 19 planktonic and the 213 benthic foraminiferal species identified are listed in Appendix B and C. Video records obtained by Remotely Operated Vehicle (ROV) or images taken with the submersible JAGO were used to characterize the seafloor, together with the descriptions of the cores, which are published in the cruise reports.

To identify significant differences in foraminiferal assemblages, univariate and multivariate statistical analysis were performed on the foraminiferal assemblage data (Appendix A) with the PRIMER software (version 6.1.9; Clarke, 1993, Clarke and Warwick, 2001). The diversity of the benthic foraminiferal faunas is expressed by the species richness S and by the Fisher's α index (Fisher et al.,

1943). To highlight similarities among assemblages, the Bray-Curties similarity matrix was calculated using square-root transformation (Clifford and Stephenson, 1975). The resulting similarity matrix was used for the hierarchical clustering analysis including the similar profile analysis (SIMPROF test, with $\rho = 0.05$) and the non-metric multidimensional scaling (nMDS) ordination plot (Kruskal, 1964; Kruskal and Wish, 1978). The SIMPROF was used to test the significance of the clusters. The null hypothesis of no internal group structures in the full set of samples was rejected when the significance level ρ was < 0.05 . For the foraminiferal assemblages the percentages of similarity (SIMPER) were calculated to highlight the contribution of each species to the total average (dis)similarity between different groups and within one group (e.g., Kruskal, 1964; Kruskal and Wish, 1978; Basso and Spezzaferri, 2000; Clarke and Warwick, 2001). Census data of the benthic foraminiferal taxa used for multivariate statistic is provided in Appendix B.

Abundance curves for each functional group of benthic foraminifera (indicators for phytodetritus flux, organic carbon flux, oxygen level and current strength) were constructed by calculating the abundances of species displaying similar ecological preferences following Murray (2006) and Margreth et al. (2009). These are

listed in Appendix D.

3.4 RESULTS

3.4.1 Surface sediments

The surface sediment samples collected along the Norwegian continental margin were obtained from a relatively wide range of water depths. Between 170 and 330 m at the northern coral reefs (Korallen, Stjernsundet, Lophavet, and Sveinsgrunnen), between 300 m and 2100 m on- and off-reef at the mid-Norwegian margin (Sula Reef, Røst Reef, Trænadjudpet), and between 90 m and 300 m in the northern Skagerrak close to the Norwegian-Swedish border (Tab. 3.1; Fig. 3.1). Field observations made during manned submersible JAGO missions and from underwater imaging material obtained by ROV as well as from the sediments recovered in the core samples, contributed to differentiate five sedimentary facies as also identified and described in Mortensen et al. (1995; 2001), Freiwald et al. (2002) and Rüggeberg et al. (in review). Facies characteristics are subsequently summarized and represented in Fig. 3.2:

(I) *Living coral framework facies*: This facies occurs at the top of the CWC-reefs. It consists

Table 3.1 Sample number, geographical position, water depth, region, mound region and facies of the investigated samples.

Nr.	Campaign label	Event label	Device	Latitude	Longitude	Depth (m)	Region	Facies
1	G.O.Sars Hermi-1	Hermi-1_1	Grab	70°27.71	21°12.77	300	Lophavet	<i>coral rubble facies</i>
2	POS 325	POS325 455	Giant box corer	70°16.13	22°29.46	270	Stjernsundet	<i>coral rubble facies</i>
3	ARK-XXII/1a	PS70/011-1	Giant box corer	69°44.21	16°33.27	327	Sveinsgrunnen	<i>coral rubble facies</i>
4	ARK-XXII/1a	PS70/023-3	van Veen Grab	66°58.12	11°07.79	324	Trænadjudpet	<i>coral rubble facies</i>
5	ARK-XXII/1a	PS70/002-2	Giant box corer	67°30.40	9°25.55	304	Røst Reef	<i>sediment clogged coral framework</i>
6	ARK-XXII/1a	PS70/029-3	Giant box corer	67°35.23	9°28.92	604	Røst Reef	<i>pebbly sand facies</i>
7	ARK-XXII/1a	PS70/028-2	Giant box corer	67°38.05	9°26.98	761	Røst Reef	<i>pebbly sand facies</i>
8	ARK-XXII/1a	PS70/037-2	Giant box corer	67°35.15	9°19.22	889	continental slope	<i>mud facies (deep)</i>
9	ARK-XXII/1a	PS70/038-2	Giant box corer	67°37.77	9°10.30	1214	continental slope	<i>mud facies (deep)</i>
10	ARK-XXII/1a	PS70/039-2	Giant box corer	67°40.15	9°03.00	1514	continental slope	<i>mud facies (deep)</i>
11	ARK-XXII/1a	PS70/033-2	Giant box corer	67°43.00	8°55.00	1824	continental slope	<i>mud facies (deep)</i>
12	ARK-XXII/1a	PS70/032-2	Giant box corer	67°52.22	8°30.72	2098	continental slope	<i>mud facies (deep)</i>
13	AL 316	AL316-320	van Veen Grab	64°06.30	8°04.80	296	Sula Reef	<i>pebbly sand facies</i>
14	AL 316	AL316 321	van Veen Grab	64°05.88	8°05.35	278	Sula Reef	<i>coral rubble facies</i>
15	AL 232	AL 232 1022	Giant box corer	58°59.88	10°57.80	91	Oslo Fjord	<i>coral rubble facies</i>
16	AL 232	AL 232 1025	Giant box corer	58°25.88	10°31.05	326	Oslo Fjord	<i>mud facies (shallow)</i>
17	AL 232	AL 232 1026	Giant box corer	58°27.75	10°30.31	287	Oslo Fjord	<i>mud facies (shallow)</i>
18	AL 232	AL 232 1155	Giant box corer	59°04.71	10°43.90	106	Oslo Fjord	<i>sediment clogged coral framework</i>
19	POS 391	POS 391 534-1	van Veen Grab	70°55.26	22°10.71	214	Korallen	<i>pebbly sand facies</i>
20	POS 391	POS 391 535-1	van Veen Grab	70°55.14	22°11.26	201	Korallen	<i>coral rubble facies</i>
21	POS 391	POS 391 539-1	van Veen Grab	70°56.09	22°11.00	247	Korallen	<i>pebbly sand facies</i>
22	POS 391	POS 391 544-2	van Veen Grab	70°56.03	22°12.35	172	Korallen	<i>coral rubble facies</i>
23	POS 391	POS 391 550-1	JAGO Grab	70°26.72	21°10.36	233	Lophavet	<i>living coral framework facies</i>
24	POS 391	POS 391 555-1	van Veen Grab	70°26.58	21°10.01	232	Lophavet	<i>coral rubble facies</i>
25	POS 391	POS 391 556-2	van Veen Grab	70°26.64	21°11.61	320	Lophavet	<i>coral rubble facies</i>
26	POS 391	POS 391 558-1	van Veen Grab	70°28.29	21°11.48	330	Lophavet	<i>pebbly sand facies</i>
27	POS 391	POS 391 559-1	van Veen Grab	70°26.93	21°11.10	230	Lophavet	<i>pebbly sand facies</i>
28	POS 391	POS 391 562-1	van Veen Grab	64°04.40	08°01.20	287	Sula Reef	<i>coral rubble facies</i>
29	POS 391	POS 391 567-1	JAGO Grab	59°06.78	10°47.46	140	Oslo Fjord	<i>sediment clogged coral framework</i>
30	POS 391	POS 391 570-2	van Veen Grab	59°05.62	10°47.95	110	Oslo Fjord	<i>coral rubble facies</i>
31	POS 391	POS 391 571-1	JAGO Grab	59°05.96	10°47.67	117	Oslo Fjord	<i>living coral framework facies</i>
32	POS 391	POS 391 584-1	Giant box corer	59°03.96	10°48.37	290	Oslo Fjord	<i>mud facies (shallow)</i>

- of large deposits of dead *Lophelia pertusa* (up to 2 m thick) with the living parts at their periphery. The living corals appear pristine white or orange and are covered by the translucent mucus produced by *L. pertusa*. This permanent mucus film actively prevents most organisms from attaching to the living polyps. Samples 23 and 31 represent this facies.
- (2) *Sediment-clogged coral framework*: this facies is found on the slopes. The construction of both *in situ* and dislocated coral framework is filled with silty clay or sandy deposits enriched by skeletal elements of bivalves and bio-eroded sponge chips. The facies could only be sampled where the dip of the slope was gentle (Tab. 3.1). Samples 5, 18, and 29 represent this facies.
- (3) *Coral rubble facies*: this external reef facies consists mainly of dead coral debris, which accumulates on the lower flanks and/or at the base of the living reefs. Fine-grained sediments, mud and silt, trapped by the coral debris form a soft-sediment substratum containing skeletal remains of molluscs and echinoids. The amount of coral debris decreases away from the reef. The inclination of the slope where this facies occurs is generally between 5° and 25°. Samples 1 to 4, 14 to 15, 20, 22, 24 to 25, 28, and 30 represent this facies.
- (4) *Pebby sand facies*: the transitional zone from the external reef facies to the “off-reef” habitats, is characterized by sandy and silty sediments, sometimes rippled and containing dropstones. Fragments of bryozoans, molluscs, and corals, may also be present. Samples 6 to 7, 13, 19, 21, 26, and 27 represent this facies. Sample 26 is in the transition zone between *pebbly sand facies* and *mud facies*, indicated by fine-grained sediment including coarser clasts.
- (5) a) *Mud facies (deep)*: this facies occurs at water depths deeper than 800 m under the influence of the NSDW in the very distal and deep parts off the mid-Norwegian shelves. Sediments consist of fine-grained (hemi-) pelagic sediments with benthic and planktonic foraminifera, echinoids, molluscs, and various terrigenous components. Samples 8 to 12 represent this facies.
- b) *Mud facies (shallow)*: This facies is represented by fine-grained sediments containing skeletal elements of foraminifera, echinoids and molluscs as well as various terrigenous components. The hemipelagic sediments from the northern Skagerrak are very fine-grained and a colour change marks the boundary from a thin oxygenated surface layer to the dysoxic sediments below. Planktonic foraminifera are almost absent. Samples 16 to 17, and 32 represent this facies.

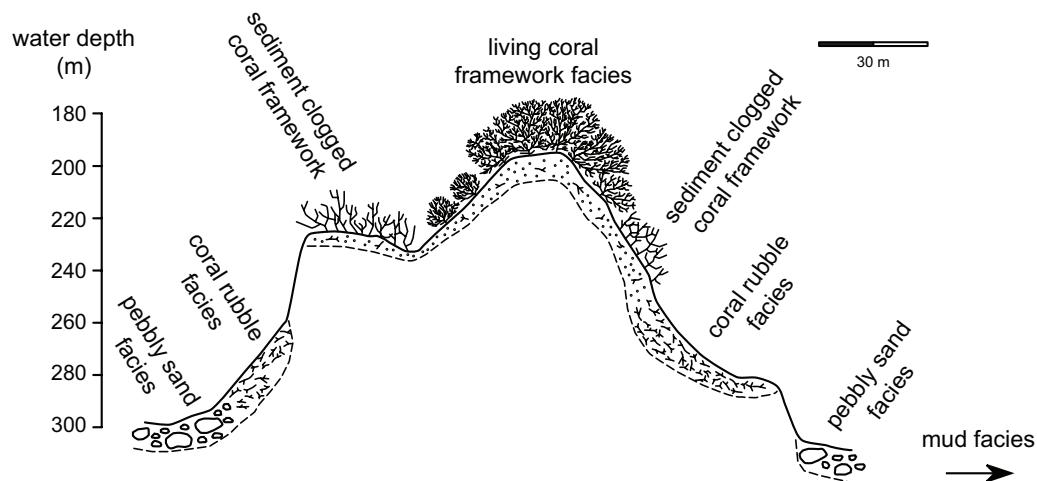


Figure 3.2 Sketch of *Lophelia*-reef facies pattern on a typical cold-water coral reef at Lophavet. The data presented are derived from multibeam maps (Institute of Marine Research, Bergen, Norway) and direct observation with the manned submersible JAGO, IFM-GEOMAR, Kiel, Germany. Facies after Freiwald et al. (2002)

3.4.2 Planktonic foraminifera

Nineteen planktonic foraminiferal species were identified in twenty samples (Appendix C). Samples containing less than 30 planktonic specimens were not considered for cluster analysis and are not listed in Appendix C.

At the 62% level of the Bray-Curties Similarity (Fig. 3.3A) two clusters can be distinguished. Cluster I comprises samples from the outer shelf and continental slope environments, including: Sveinsgrunnen (327 m), Trænadjudupet (324 m), Røst Reef (304 m, 604 m, 761 m), Mid Norwegian continental slope (889 m, 1214 m, 1514 m, 1824 m, 2098 m), and Sula Reef (296 m, 278 m). Six species and/or groups account for 95.1%

of the average similarity (Tab. 3.2; Fig. 3.3B). Cluster II groups samples from Korallen (172 m, 201 m), and Lophphavet (230 m, 232 m, 233 m, 320 m, 330 m) on the inner shelf close to the shore. Four species and/or groups of species account for 93.44% of the average similarity (Tab. 3.2; Fig. 3.3B). The dissimilarities between the two groups are given in Table 3.3.

3.4.3 Benthic foraminifera

In total, 213 (unstained) benthic foraminiferal species were identified in the studied samples (Appendix B). Assemblages are generally well preserved and show a high diversity. They are characterized by epibenthic species such as *Cibicidoides pachyderma*, *Fontbotia wuellerstorfi*, *Cribrostomoides subglobosum*,

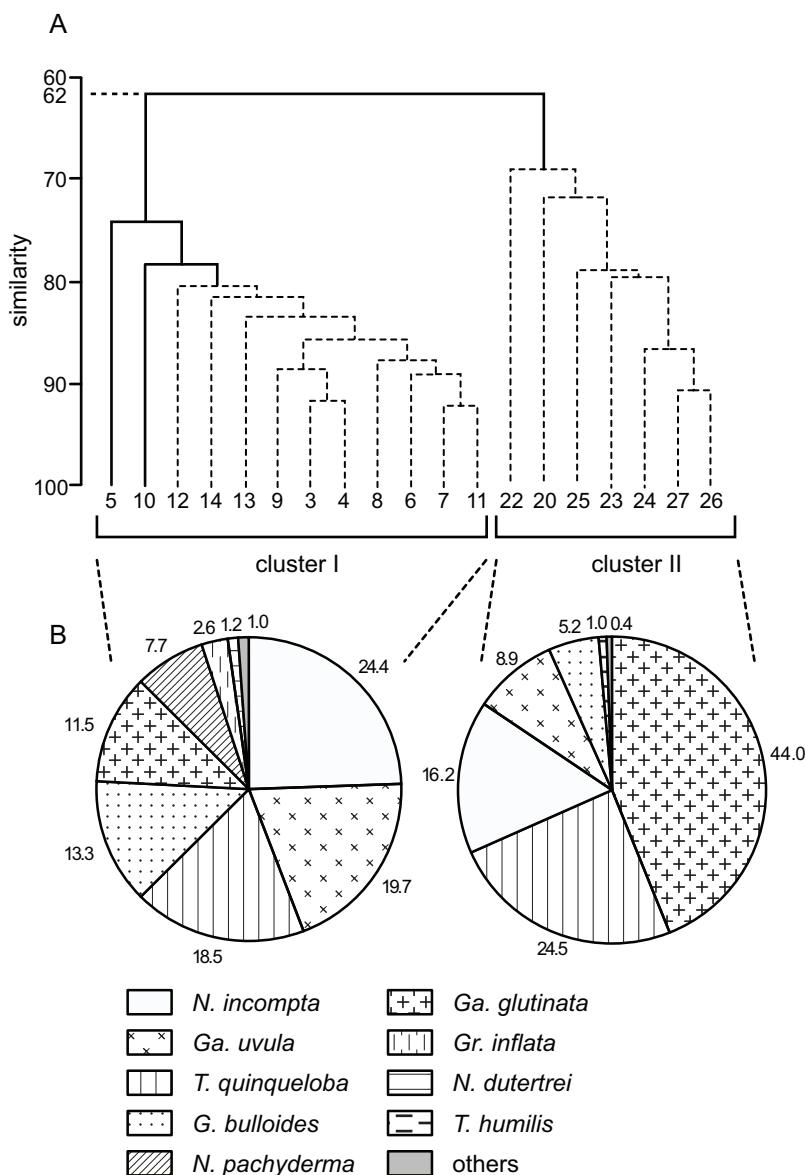


Figure 3.3 A: Hierarchical Cluster Analysis based on the Bray-Curties similarity matrix of planktonic foraminifera compositional data. Dashed lines indicate non-significantly differing samples (SIMPROF $p > 0.05$). B: Contribution (%) of planktonic foraminiferal species to the two separated groups I and II.

Lobatula lobatula, and *Discanomalina coronata*, and infaunal species such as *Angulogerina angulosa*, *Melonis barleeanum*, *Uvigerina peregrina*, and *Globocassidulina* spp., *Cassidulina* spp., *Epistominella* spp.

3.4.3.1. Depth zonation of benthic foraminifera

During the R/V POLARSTERN Cruise ARK-XXII/1a samples were retrieved along a bathymetric transect. The investigations along this transect revealed a zonation in the distribution pattern of the benthic foraminifera from the shelf area of Trænadjudpet and Røst Reef down the slope to the abyssal area at 2098 m water depth (Fig. 3.4). In the shallowest part *A. angulosa* and *C. pachyderma* dominate the assemblages (Fig. 3.4). *Uvigerina peregrina* and *Hyalinea balthica* show their highest abundance at 327 m water depth at Trænadjudpet, while *D. coronata* is more abundant at the Røst Reef at 304 m. Further downslope the numbers of *C. pachyderma* decrease. *Angulogerina angulosa* displays a similar decreasing trend with increasing water depth. *Melonis barleeanum* and the *Cassidulina laevigata* group is very abundant on the midslope. At a water depth of 1214 m the agglutinated species *Reophax scorpiurus* and *Reophax diffugiformis* are highly abundant. In the deepest part

of the transect *Epistominella exigua*, *C. subglobosum*, and *F. wuellerstorfi* dominate the assemblage. *Pyrgo sarsi* is only abundant at 2098 m (Fig. 3.4).

3.4.3.2. Diversity

To compare the diversity patterns of the thanatocoenoses (unstained) of benthic foraminifera from different sediment facies, the species richness S and the Fisher's α were calculated (Tab. 3.4). These univariate statistical parameters were calculated for the species and groups of species listed in the Appendix B. The benthic species richness (S) has values between 19 (sample 21) and 59 (sample 31). Fisher's α has a minimum value of 4.83 in sample 21, and a maximum value of 21.05 in sample 5 (Tab. 3.4).

3.4.3.3. Assemblages

The hierarchical clustering based on the Bray-Curties Similarity matrix (Fig. 3.5A) and nMDS (Fig. 3.5B) is used to highlight potential patterns in the benthic foraminiferal assemblages, since they are not often readily apparent (Clark and Warwick, 1994). The hierarchical cluster analysis shows two high-level clusters A and B separated at 28% of the Bray-Curties Similarity (Fig. 3.5). The two clusters are bathymetrically distinct. Cluster A includes bathyal

Table 3.2 List of planktonic species and statistical parameters associated with the similarity in Clusters I and II. Average similarity within the group of stations, average abundance, average similarity, contribution (%), and cumulative contribution (%) are given for each species with respect to the total similarity for each group.

Cluster I		Average similarity: 81.64		
Species	Av.Aband	Av.Sim	Contrib%	Cum.%
<i>N. incompta</i>	5.51	19.92	24.4	24.4
<i>Ga. uvula</i>	4.55	16.12	19.74	44.14
<i>T. quinqueloba</i>	4.3	15.1	18.49	62.63
<i>G. bulloides</i>	3.17	10.85	13.29	75.92
<i>Ga. glutinata</i>	2.88	9.4	11.52	87.44
<i>N. pachyderma</i>	2.2	6.25	7.66	95.1
Cluster II		Average similarity: 75.82		
Species	Av.Aband	Av.Sim	Contrib%	Cum.%
<i>Ga. glutinata</i>	7.18	33.34	43.97	43.97
<i>T. quinqueloba</i>	4.29	18.55	24.46	68.44
<i>N. incompta</i>	3.74	12.25	16.15	84.59
<i>Ga. uvula</i>	2.17	6.71	8.85	93.44

and abyssal samples from depths between 889 and 2098 m. Cluster B includes shelf and upper bathyal samples from depths between 91 to 761 m. Six sub-clusters separate at the lower level of 49% of the Bray-Curties Similarity (Fig. 3.5). Cluster I includes the deepest bathyal and abyssal samples from the mid Norwegian continental slope (1824 m, 2098 m; Bray-Curties Similarity 66.6%). Nine species and/or groups account for 80.7% of the average similarity (Tab. 3.5).

Cluster II includes shallower bathyal samples from the mid Norwegian slope (889–1514 m; Bray-Curties Similarity 61.6%). Thirteen species and/or groups account for 81.6% of the average similarity (Tab. 3.5). Cluster III includes “off-reef” samples from the Skagerrak (287–326 m; Bray-Curties Similarity 58.1%). Sixteen species and/or groups account for 82.0% of the average similarity (Tab. 3.5). Cluster IV includes “on-reef” samples from the Skagerrak

Table 3.3 List of planktonic species and statistical parameter associated with the dissimilarity between each pair of Clusters I and II. For each pair of groups the average dissimilarity is shown. For each species the average abundance, average dissimilarity, contribution (%), and cumulative contribution (%) to the total similarity are also given.

Species	Average dissimilarity = 38.30				
	Cluster I	Cluster II	Av.Abund	Av.Abund	Av.Diss
<i>Ga. glutinata</i>	2.88	7.18	9.79	25.55	25.55
<i>Ga. uvula</i>	4.55	2.17	5.53	14.44	39.99
<i>N. incompta</i>	5.51	3.74	5.46	14.27	54.26
<i>G. bulloides</i>	3.17	1.07	4.73	12.35	66.61
<i>N. pachyderma</i>	2.2	0.48	4.13	10.79	77.39
<i>T. quinqueloba</i>	4.3	4.29	2.58	6.73	84.12
<i>Gr. inflata</i>	0.86	0	1.93	5.04	89.16
<i>N. dutertrei</i>	0.55	0	1.21	3.16	92.31
<i>T. humilis</i>	0	0.53	1.17	3.05	95.36

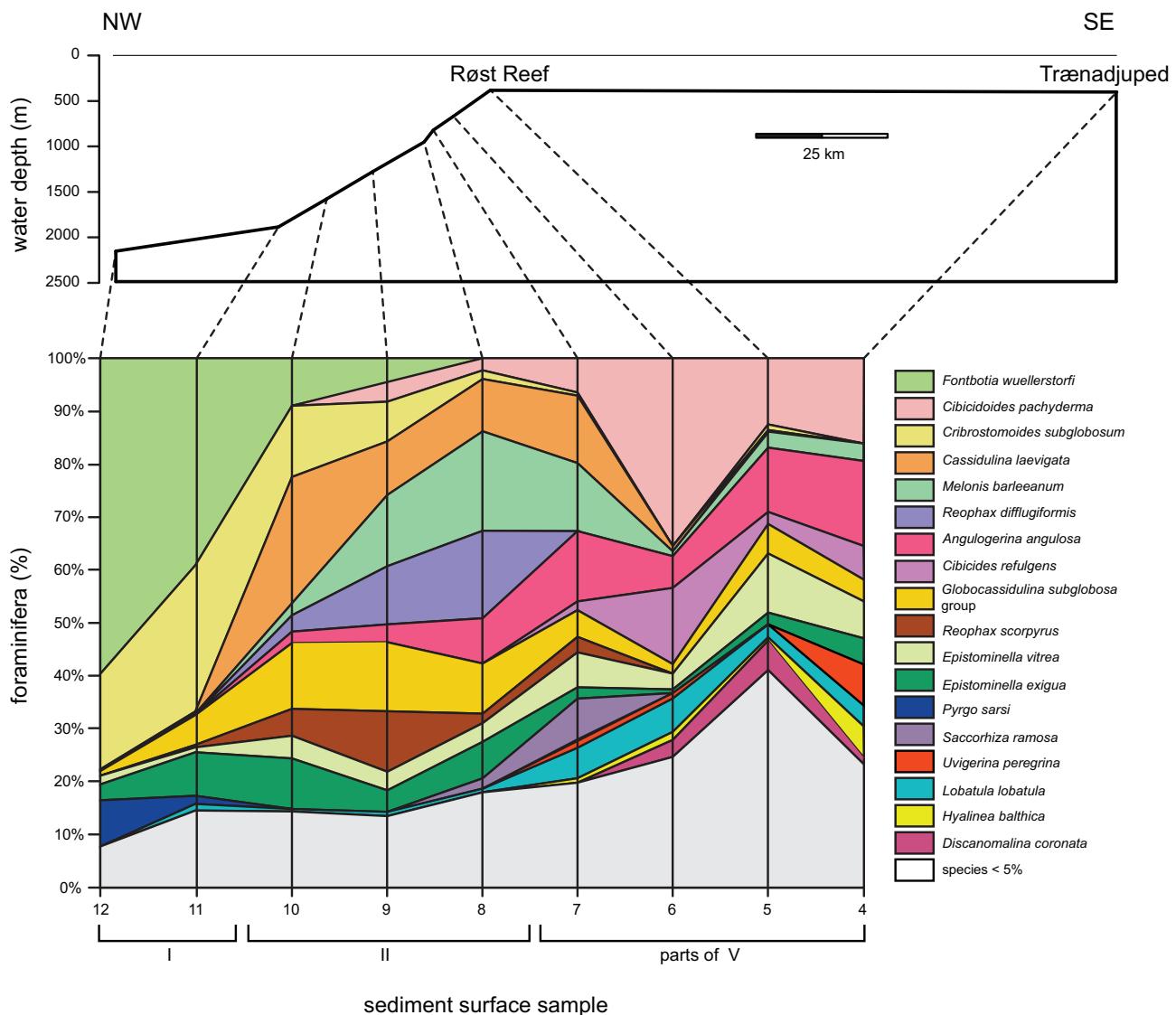


Figure 3.4 Percent distribution of the most abundant benthic foraminiferal species (> 5%) along a bathymetric transect across the Mid-Norwegian shelf at 67°N comprising 5 off-reef stations (11 and 12, group I; 8, 9, and 10, group II) and 4 reef associated stations (4, 5, 6, and 7, parts of group V).

Table 3.4 Species numbers (S) and Fisher's α of the dead (unstained) benthic foraminiferal fauna of surface sediments (0-2 cm sediment depth) associated to Clusters I to VI.

Sample	Facies	Species richness (S)	Fisher's α Index
23	<i>living coral framework facies</i>	50	15.04
31	<i>living coral framework facies</i>	55	17.14
5	<i>sediment clogged coral framework</i>	53	19.88
18	<i>sediment clogged coral framework</i>	45	14.03
29	<i>sediment clogged coral framework</i>	44	13.91
1	<i>coral rubble facies</i>	47	12.37
2	<i>coral rubble facies</i>	46	12.83
3	<i>coral rubble facies</i>	54	15.17
4	<i>coral rubble facies</i>	46	11.93
14	<i>coral rubble facies</i>	35	10.39
15	<i>coral rubble facies</i>	50	15.58
20	<i>coral rubble facies</i>	36	10.64
22	<i>coral rubble facies</i>	30	8.61
24	<i>coral rubble facies</i>	40	10.49
25	<i>coral rubble facies</i>	33	9.34
28	<i>coral rubble facies</i>	50	16.28
30	<i>coral rubble facies</i>	51	17.62
6	<i>pebbly sand facies</i>	52	14.22
7	<i>pebbly sand facies</i>	38	12.05
13	<i>pebbly sand facies</i>	31	8.01
19	<i>pebbly sand facies</i>	32	8.16
21	<i>pebbly sand facies</i>	19	4.83
27	<i>pebbly sand facies</i>	38	9.99
16	<i>mud facies (shallow)</i>	44	9.94
17	<i>mud facies (shallow)</i>	34	8.69
26	<i>mud facies (shallow)</i>	48	13.21
32	<i>mud facies (shallow)</i>	35	9.77
8	<i>mud facies (deep)</i>	37	10.86
9	<i>mud facies (deep)</i>	34	8.11
10	<i>mud facies (deep)</i>	34	9.10
11	<i>mud facies (deep)</i>	32	7.13
12	<i>mud facies (deep)</i>	23	4.69

(91–140 m; Bray-Curties Similarity 61.9%). Twenty-two species and/or groups account for 81.2% of the average similarity (Tab. 3.5). Cluster V includes samples from the mid Norwegian reefs (278–761 m; Bray-Curties Similarity 55.1%). Fifteen species and/or groups account for 81.5% of the average similarity (Tab. 3.5). Cluster VI includes samples from the northern Norwegian reefs (172–327 m; Bray-Curties Similarity 58.9%). Seventeen species and/or groups account for 80.3% of the average similarity (Tab. 3.5). The dissimilarities between each group are given in Table 3.6.

3.5 DISCUSSION

3.5.1 Planktonic foraminifera

Seasonal plankton blooms have been described for coastal zones and shelf areas (Drebes, 1974; Le Fèvre, 1986). On the Norwegian continental margin and shelf plankton blooms are related to high nutrient transport caused by seasonal upwelling and eddies (Berner and Wefer, 1994; Sætre, 1999; Mitchelson-Jacob and Sundby, 2001). As demonstrated by Schiebel et al. (2001) and Schiebel and Hemleben (2005) for the NAC, pulses of primary productivity result in a predictable seasonally

mixed assemblage of planktonic foraminiferal species in the sediment record. Areas influenced by upwelling are characterized by small opportunistic species of *Globigerina bulloides*, *Turborotalita quinqueloba*, and *Globigerinita glutinata* (e.g., Bé and Tolderlund, 1971; Thiede, 1975; Hemleben et al., 1989; Marchant et al., 1998; Rutherford et al., 1999; Schmidt et al., 2004) whereas the presence of *Neogloboquadrina incompta* indicates warmer and low-productive waters (Sautter and Thunnel, 1991; Schiebel et al., 2001).

Cluster I groups samples from the Norwegian continental slope and outer shelf area. The high proportion of *N. incompta* in the assemblages is interpreted to reflect the dominance of relatively warm and nutrient-poor waters of the NAC during summer whereas the high abundance of *T. quinqueloba* and *G. bulloides* indicates the prevalence of cool and nutrient-rich conditions during upwelling in wintertime. Cluster II groups the samples from the inner shelf areas, which are dominated by the NCC. Very small specimens of *G. glutinata*, *T. quinqueloba* and *G. uvula*, ranging in size from 63 µm to 125 µm, are the dominant species in this cluster. These species indicate cool and nutrient-rich conditions and must, therefore, be related to upwelling and eddies in winter.

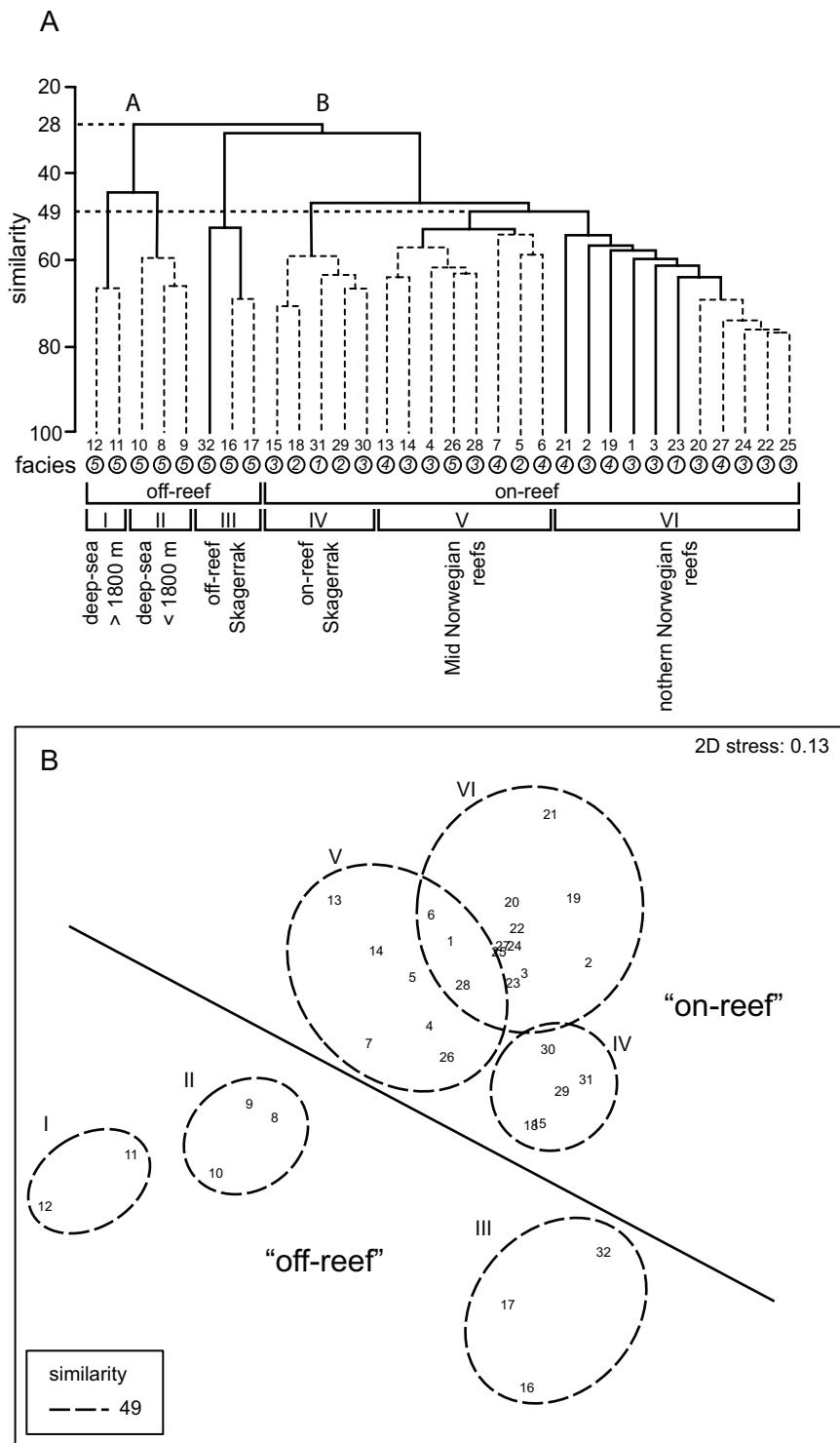


Figure 3.5 A) Hierarchical Cluster Analysis based on the Bray-Curties similarity matrix of benthic foraminifera compositional data. A and B indicate high-level cluster separating at 28% of Bray-Curties Similarity. I, II, III, IV, V, and VI display clusters at a lower level separating at 49% of Bray-Curties Similarity. Dashed lines indicate non-significantly differing samples (SIMPROF $p > 0.05$). The different facies are also specified: 1) living coral framework facies, 2) sediment-clogged coral facies, 3) coral rubble facies, 4) pebbly sand facies, 5) mud facies. B) Non-metric MultiDimensional Scaling (nMDS) plot obtained from Bray-Curties similarity matrix of benthic foraminifera compositional data.

Planktonic foraminifera do not tolerate hyposaline conditions (e.g., Kucera, 2007). During summer the NCC dominantly originate from the discharge of

freshwaters from the Baltic Sea and freshwater runoff from Norway (Mork, 1981), therefore, their influence may explain the low contribution of warm and nutrient-

Table 3.5 List of benthic species and statistical parameters associated with the similarity in Clusters I to VI. Average similarity within the group of station, average abundance, average similarity, contribution (%), and cumulative contribution (%) are given for each species with respect to the total similarity for each group.

Cluster I		Average similarity: 66.56			
Species		Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Fontbotia wuelllerstorfi</i>		6.99	18.23	27.39	27.39
<i>Cribrastomoides subglobosum</i>		4.74	12.32	18.5	45.89
<i>Epistominella exigua</i>		2.35	5.23	7.85	53.75
<i>Oridorsalis umbonatus</i>		1.47	4.04	6.07	59.82
<i>Pyrgo sarsi</i>		2.04	3.5	5.26	65.08
<i>Fissurina</i> spp.		1.02	2.86	4.3	69.38
<i>Lobatula lobatula</i>		1.01	2.61	3.93	73.31
<i>Epistominella vitrea</i>		1.05	2.61	3.92	77.23
<i>Globocassidulina subglobosa</i> group		1.49	2.34	3.51	80.74
Cluster II		Average similarity: 61.64			
Species		Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Globocassidulina subglobosa</i> group		3.41	7.01	11.38	11.38
<i>Cassidulina laevigata</i> group		3.73	6.81	11.04	22.42
<i>Reophax diffugiformis</i>		3.08	5.01	8.12	30.54
<i>Epistominella exigua</i>		2.61	4.93	8	38.54
<i>Melonis barleeanum</i>		3.17	4.77	7.74	46.29
<i>Epistominella vitrea</i>		1.94	4.08	6.61	52.9
<i>Cribrastomoides subglobosum</i>		2.6	4.04	6.55	59.45
<i>Reophax scorpiurus</i>		2.34	3.65	5.91	65.36
<i>Angulogerina angulosa</i>		2.04	3.38	5.49	70.85
<i>Cibicidoides pachyderma</i>		1.25	1.74	2.82	73.67
<i>Paratrochammina challengerii</i>		1.01	1.64	2.66	76.33
<i>Rhabdammina abyssorum</i>		0.92	1.64	2.66	78.99
<i>Lobatula lobatula</i>		0.82	1.61	2.61	81.6
Cluster III		Average similarity: 58.08			
Species		Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Globobulimina affinis</i>		4.53	7.99	13.76	13.76
<i>Eggerelloides scaber</i>		2.52	4.32	7.44	21.2
<i>Hyalinea balthica</i>		2.73	3.8	6.55	27.75
<i>Bigenerina cylindrica</i>		1.99	3.69	6.35	34.1
<i>Textularia tenuissima</i>		2.12	3.28	5.64	39.74
<i>Melonis barleeanum</i>		1.7	2.98	5.12	44.86
<i>Bulimina marginata</i>		1.78	2.65	4.56	49.43
<i>Bolivina</i> spp.		1.98	2.6	4.47	53.9
<i>Cassidulina carinata</i>		2	2.48	4.26	58.16
<i>Melonis pompilioides</i>		1.9	2.35	4.05	62.21
<i>Cassidulina laevigata</i> group		1.69	2.3	3.97	66.18
<i>Stainforthia fusiformis</i>		1.25	2.12	3.65	69.83
<i>Uvigerina peregrina</i>		1.44	2.01	3.46	73.28
<i>Trochammina robertsoni</i>		1.14	1.88	3.23	76.52
<i>Pullenia subcarinata</i>		2.1	1.8	3.1	79.62
<i>Nonionella turgida</i>		0.78	1.36	2.34	81.96
Cluster IV		Average similarity: 61.92			
Species		Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Lobatula lobatula</i>		3.65	5.51	8.9	8.9
<i>Cibicidoides pachyderma</i>		3.05	4.57	7.39	16.28
<i>Melonis pompilioides</i>		2.91	4.56	7.36	23.64
<i>Melonis barleeanum</i>		2.29	3.6	5.81	29.45
<i>Bulimina marginata</i>		2.44	3.3	5.32	34.77
<i>Cassidulina laevigata</i> group		2.22	3.01	4.85	39.63
<i>Cibicides refulgens</i>		1.87	2.34	3.79	43.41
<i>Hyalinea balthica</i>		1.81	2.32	3.74	47.15
<i>Epistominella exigua</i>		1.54	2.17	3.5	50.65
<i>Cibicidoides</i> sp.		1.46	2.11	3.42	54.07
<i>Astrononion gallowayi</i>		1.4	1.91	3.08	57.15
<i>Angulogerina angulosa</i>		1.4	1.88	3.03	60.18
<i>Nonionella iridea</i>		1.33	1.86	3.01	63.19
<i>Gavelinopsis</i> spp.		1.37	1.85	2.99	66.18
<i>Spiroplectinella wrightii</i>		1.11	1.7	2.74	68.92
<i>Globocassidulina subglobosa</i> group		1.09	1.43	2.31	71.23
<i>Cassidulina carinata</i>		1.39	1.43	2.3	73.53
<i>Bolivina</i> spp.		1.06	1.08	1.74	75.27
<i>Quinqueloculina seminula</i>		0.66	0.97	1.56	76.83

Table 3.5 (continued)

<i>Adercotryma wrightii</i>	0.97	0.96	1.55	78.38
<i>Eggerelloides scaber</i>	0.93	0.87	1.41	79.79
<i>Fissurina</i> spp.	0.86	0.85	1.37	81.16
Cluster V				
	Average similarity: 55.11			
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Angulogerina angulosa</i>	3.67	6.24	11.32	11.32
<i>Cibicidoides pachyderma</i>	3.41	5.05	9.17	20.48
<i>Epistominella vitrea</i>	2.66	4.36	7.91	28.39
<i>Globocassidulina subglobosa</i> group	2.34	3.79	6.87	35.26
<i>Cibicides refulgens</i>	2.25	3.36	6.09	41.35
<i>Melonis barleeanum</i>	1.96	2.91	5.27	46.63
<i>Cibicidoides kullenbergi</i>	1.53	2.32	4.22	50.84
<i>Hyalinea balthica</i>	1.56	2.16	3.93	54.77
<i>Cassidulina carinata</i>	1.49	2.13	3.86	58.63
<i>Lobatula lobatula</i>	1.62	2.01	3.65	62.28
<i>Uvigerina peregrina</i>	1.94	1.88	3.41	65.68
<i>Epistominella exigua</i>	1.29	1.76	3.2	68.88
<i>Discanomalina coronata</i>	1.49	1.53	2.77	71.66
<i>Fissurina</i> spp.	0.91	1.4	2.55	74.2
<i>Cassidulina laevigata</i> group	1.37	1.4	2.54	76.75
<i>Hanzawaia boueana</i>	1.17	1.31	2.37	79.12
<i>Cibicidoides</i> sp.	1.17	1.29	2.34	81.46
Cluster VI				
	Average similarity: 58.98			
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Cibicidoides pachyderma</i>	4.57	8.51	14.43	14.43
<i>Lobatula lobatula</i>	2.81	4.69	7.94	22.38
<i>Globocassidulina subglobosa</i> group	2.6	4.68	7.94	30.31
<i>Discanomalina coronata</i>	2.93	4.54	7.7	38.02
<i>Cibicides refulgens</i>	2.56	3.73	6.33	44.35
<i>Angulogerina angulosa</i>	2.15	3.39	5.74	50.09
<i>Cassidulina laevigata</i> group	1.81	3.02	5.12	55.21
<i>Melonis barleeanum</i>	1.66	2.82	4.78	59.99
<i>Cibicidoides</i> sp.	1.7	2.56	4.34	64.33
<i>Astronion gallowayi</i>	1.38	1.91	3.25	67.58
<i>Gavelinopsis</i> spp.	1.13	1.81	3.06	70.64
<i>Fissurina</i> spp.	1.16	1.79	3.04	73.68
<i>Epistominella vitrea</i>	1.14	1.58	2.68	76.37
<i>Nonionella iridea</i>	0.72	1.25	2.12	78.49
<i>Epistominella exigua</i>	0.89	1.08	1.83	80.32

poor indicator species. Samples associated to Cluster II are dominated by small opportunistic species, which are interpreted to be mainly deposited during high productivity in winter.

In summary, planktonic foraminiferal assemblages in both clusters reflect a seasonal mixed signal. Cluster I represents the seasonal mixed assemblage on the continental margin and shelf associated with the NAC with a contribution of both cold and warm water species. Cluster II is dominated by the fauna associated with high winter productivity in the NCC. In this cluster warm water species although present contribute very little to the assemblage.

Cool and deep waters are undersaturated in calcium carbonate and result in the dissolution of foraminiferal tests (e.g., Hemleben et al., 1989). Continental shelves in cold regions (Alexandersson, 1978), slopes and basins deeper than the carbonate compensation depth

(e.g., Murray, 2006) may have corrosive bottom waters with respect to the CaCO_3 . Cluster I includes planktonic foraminifera from sample obtained in water depths ranging between 278 m (sample 14) and 2098 m (sample 12) from the Norwegian slope (Tab.3.1; Fig. 3.3). Since no remarkable changes in composition or abundance have been observed in the planktonic assemblage, it is reasonable to assume that dissolution does not affect the fragile planktonic foraminiferal tests. Consequently, the benthic foraminiferal assemblages can also be considered as being unaffected by carbonate dissolution confirming the observations on pristine benthic assemblages by Mackensen et al. (1985) from the Southwest Norwegian slope.

3.5.2 Benthic foraminiferal assemblage

3.5.2.1 High-level cluster A and B: the bathymetric imprint
Benthic foraminiferal communities show conspicuous

Table 3.6 List of benthic species and statistical parameter associated with the dissimilarity between each pair of Clusters I to VI. For each pair of cluster the average dissimilarity is shown. For each species the average abundance, average dissimilarity, contribution (%), and cumulative contribution (%) to the total similarity are also given.

Species	Average dissimilarity = 51.33				
	Cluster VI	Cluster V	Av.Diss	Contrib%	Cum.%
	Av.Abund	Av.Abund			
<i>Discanomalina coronata</i>	2.93	1.49	1.91	3.71	3.71
<i>Uvigerina peregrina</i>	0.25	1.94	1.89	3.68	7.39
<i>Cibicidoides pachyderma</i>	4.57	3.41	1.77	3.45	10.84
<i>Angulogerina angulosa</i>	2.15	3.67	1.62	3.15	14
<i>Epistominella vitrea</i>	1.14	2.66	1.54	3.01	17
<i>Cibicidoides kullenbergi</i>	0.2	1.53	1.4	2.74	19.74
<i>Lobatula lobatula</i>	2.81	1.62	1.37	2.66	22.4
<i>Cibicides refulgens</i>	2.56	2.25	1.33	2.58	24.98
<i>Hyalinea balthica</i>	0.54	1.56	1.3	2.53	27.52
<i>Cassidulina carinata</i>	0.63	1.49	1.13	2.19	29.71
<i>Cassidulina laevigata</i> group	1.81	1.37	1.08	2.1	31.81
<i>Cibicidoides</i> sp.	1.7	1.17	0.97	1.9	33.71
<i>Cibicides/Cibicidoides</i> juv	1.01	0.28	0.95	1.86	35.56
<i>Hanzawaia boueana</i>	0.83	1.17	0.88	1.71	37.27
<i>Gavelinopsis</i> spp.	1.13	0.39	0.87	1.69	38.97
<i>Astrononion gallowayi</i>	1.38	0.63	0.83	1.61	40.58
<i>Melonis barleeanum</i>	1.66	1.96	0.71	1.39	41.97
<i>Epistominella exigua</i>	0.89	1.29	0.71	1.39	43.36
<i>Discorbiniella bertheloti</i>	0.72	0.21	0.71	1.39	44.74
<i>Miliolinella subrotunda</i>	0.74	0.29	0.71	1.38	46.13
<i>Globocassidulina subglobosa</i> group	2.6	2.34	0.69	1.35	47.48
<i>Pullenia osloensis</i>	0.82	0.69	0.67	1.3	48.78
<i>Melonis pomphiloides</i>	0.53	0.39	0.65	1.26	50.04
<i>Cibicides ungerianus</i>	0.58	0.56	0.64	1.25	51.29
<i>Hanzawaia concentrica</i>	0.65	0.16	0.64	1.25	52.54
<i>Pullenia subcarinata</i>	0.26	0.7	0.63	1.23	53.77
<i>Planulina ariminensis</i>	0	0.62	0.62	1.22	54.98
<i>Patellina corrugata</i>	0.67	0.16	0.58	1.13	56.12
<i>Bolivina</i> spp.	0.73	0.27	0.55	1.08	57.19
<i>Sigmoilopsis schlumbergeri</i>	0	0.52	0.53	1.03	58.22
<i>Bulimina marginata</i>	0.05	0.54	0.52	1.01	59.24
<i>Quinqueloculina viennensis</i>	0.47	0.29	0.52	1.01	60.24
<i>Fissurina</i> spp.	1.16	0.91	0.51	1	61.25
<i>Pullenia bulloides</i>	0.05	0.51	0.5	0.98	62.23
<i>Sphaeroidina bulloides</i>	0.11	0.46	0.5	0.98	63.21
<i>Sacchorhiza ramosa</i>	0.07	0.46	0.5	0.96	64.17
<i>Nonionella iridea</i>	0.72	0.58	0.49	0.96	65.14
<i>Nonionella turgida</i>	0.11	0.47	0.47	0.92	66.06
<i>Spiroplectinella wrightii</i>	0.4	0.45	0.46	0.89	66.95
<i>Pullenia quinqueloba</i>	0.22	0.34	0.43	0.84	67.79
<i>Trochammina inflata</i>	0.04	0.44	0.43	0.83	68.62
<i>Uvigerina mediterranea</i>	0	0.39	0.42	0.82	69.44
<i>Pullenia quadriloba</i>	0.17	0.39	0.41	0.8	70.24
<i>Hyrrokin sarcophaga</i>	0.04	0.43	0.41	0.79	71.04
<i>Verneulina propinqua</i>	0.24	0.29	0.39	0.76	71.8
<i>Rosalina globularis</i>	0.21	0.29	0.36	0.7	72.5
<i>Haplofragmoides membranaceum</i>	0.05	0.36	0.36	0.7	73.2
<i>Quinqueloculina lamarciana</i>	0.18	0.31	0.34	0.67	73.87
<i>Elphidium frigidum</i>	0.08	0.32	0.34	0.66	74.53
<i>Cribrostomoides jeffreysii</i>	0.33	0.06	0.34	0.66	75.19
<i>Nonionella labradorica</i>	0.37	0.14	0.34	0.65	75.85
<i>Reophax scorpiurus</i>	0.12	0.25	0.33	0.64	76.49
<i>Favulinina</i> spp.	0.27	0.33	0.32	0.62	77.1
<i>Homalohedra borealis</i>	0.15	0.28	0.32	0.61	77.72
<i>Gyroidinoides laevigatus</i>	0.04	0.31	0.31	0.61	78.33
<i>Eggerella bradyi</i>	0.05	0.27	0.3	0.58	78.9
<i>Paratrocchamina challengerii</i>	0.04	0.3	0.29	0.56	79.46
<i>Elphidium clavatum</i>	0.26	0	0.28	0.54	80
<i>Gyroidina</i> spp.	0.08	0.25	0.27	0.52	80.53

Table 3.6 (continued)

Species	Average dissimilarity = 79.30				
	Cluster VI Av.Abund	Cluster I Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Fontbotia wuellerstorfi</i>	0.05	6.99	8.58	10.82	10.82
<i>Cibrostomoides subglobosum</i>	0	4.74	5.78	7.29	18.11
<i>Cibicidoides pachyderma</i>	4.57	0.2	5.46	6.89	25
<i>Discanomalina coronata</i>	2.93	0	3.72	4.68	29.68
<i>Cibicides refulgens</i>	2.56	0.2	3.04	3.83	33.51
<i>Pyrgo sarsi</i>	0	2.04	2.56	3.23	36.74
<i>Lobatula lobatula</i>	2.81	1.01	2.19	2.76	39.51
<i>Angulogerina angulosa</i>	2.15	0.4	2.17	2.73	42.24
<i>Cibicidoides sp.</i>	1.7	0	2.07	2.61	44.84
<i>Oridorsalis umbonatus</i>	0	1.47	1.81	2.28	47.12
<i>Epistominella exigua</i>	0.89	2.35	1.79	2.26	49.38
<i>Melonis barleeanum</i>	1.66	0.28	1.69	2.13	51.51
<i>Globocassidulina subglobosa</i> group	2.6	1.49	1.5	1.89	53.39
<i>Cassidulina laevigata</i> group	1.81	0.63	1.43	1.81	55.2
<i>Astrononion gallowayi</i>	1.38	0.2	1.43	1.8	57
<i>Gavelinopsis</i> spp.	1.13	0	1.37	1.73	58.73
<i>Cibicides/Cibicidoides</i> juv	1.01	0	1.21	1.52	60.25
<i>Rhabdammina abyssorum</i>	0.09	0.94	1.08	1.36	61.62
<i>Pullenia osloensis</i>	0.82	0	0.99	1.25	62.87
<i>Cycloforina angularis</i>	0	0.8	0.98	1.24	64.1
<i>Hanzawaia boueana</i>	0.83	0	0.97	1.22	65.33
<i>Miliolinella subrotunda</i>	0.74	0	0.91	1.15	66.48
<i>Bolivina</i> spp.	0.73	0	0.87	1.1	67.57
<i>Discorbinella bertheloti</i>	0.72	0	0.86	1.09	68.66
<i>Hormonisella guttifera</i>	0.04	0.73	0.84	1.06	69.72
<i>Cassidulina carinata</i>	0.63	0.8	0.84	1.06	70.78
<i>Gyroidinoides laevigatus</i>	0.04	0.69	0.8	1.01	71.8
<i>Patellina corrugata</i>	0.67	0	0.79	1	72.8
<i>Hanzawaia concentrica</i>	0.65	0	0.78	0.98	73.78
<i>Pullenia subcarinata</i>	0.26	0.73	0.73	0.92	74.7
<i>Cibicides ungerianus</i>	0.58	0	0.7	0.89	75.58
<i>Epistominella vitrea</i>	1.14	1.05	0.66	0.83	76.41
<i>Botellina labyrinthica</i>	0	0.56	0.66	0.83	77.24
<i>Melonis pomphiloides</i>	0.53	0	0.61	0.77	78.01
<i>Hyalinea balthica</i>	0.54	0	0.61	0.76	78.78
<i>Pullenia quinqueloba</i>	0.22	0.4	0.59	0.74	79.52
<i>Quinqueloculina viennensis</i>	0.47	0	0.58	0.73	80.26

Species	Average dissimilarity = 76.44				
	Cluster V Av.Abund	Cluster I Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Fontbotia wuellerstorfi</i>	0	6.99	8.16	10.67	10.67
<i>Cibrostomoides subglobosum</i>	0.25	4.74	5.19	6.79	17.46
<i>Angulogerina angulosa</i>	3.67	0.4	3.84	5.02	22.48
<i>Cibicidoides pachyderma</i>	3.41	0.2	3.71	4.85	27.33
<i>Pyrgo sarsi</i>	0	2.04	2.42	3.16	30.49
<i>Cibicides refulgens</i>	2.25	0.2	2.39	3.12	33.62
<i>Uvigerina peregrina</i>	1.94	0	2.33	3.05	36.67
<i>Melonis barleeanum</i>	1.96	0.28	1.97	2.58	39.25
<i>Epistominella vitrea</i>	2.66	1.05	1.86	2.43	41.68
<i>Hyalinea balthica</i>	1.56	0	1.84	2.4	44.08
<i>Cibicidoides kullenbergi</i>	1.53	0	1.82	2.38	46.46
<i>Discanomalina coronata</i>	1.49	0	1.66	2.18	48.63
<i>Oridorsalis umbonatus</i>	0.14	1.47	1.55	2.03	50.66
<i>Hanzawaia boueana</i>	1.17	0	1.36	1.78	52.44
<i>Cibicidoides</i> sp.	1.17	0	1.35	1.76	54.2
<i>Epistominella exigua</i>	1.29	2.35	1.29	1.69	55.89
<i>Globocassidulina subglobosa</i> group	2.34	1.49	1.2	1.56	57.45
<i>Lobatula lobatula</i>	1.62	1.01	1.07	1.4	58.85
<i>Cassidulina carinata</i>	1.49	0.8	0.98	1.29	60.14
<i>Cassidulina laevigata</i> group	1.37	0.63	0.97	1.27	61.41
<i>Cycloforina angularis</i>	0	0.8	0.93	1.21	62.62
<i>Rhabdammina abyssorum</i>	0.15	0.94	0.92	1.2	63.82
<i>Hormonisella guttifera</i>	0	0.73	0.84	1.09	64.92
<i>Gyroidinoides laevigatus</i>	0.31	0.69	0.78	1.02	65.93

Table 3.6 (continued)

<i>Pullenia osloensis</i>	0.69	0	0.77	1.01	66.95
<i>Planulina ariminensis</i>	0.62	0	0.72	0.95	67.9
<i>Cibicides ungerianus</i>	0.56	0	0.66	0.87	68.76
<i>Reophax scorpiurus</i>	0.25	0.45	0.64	0.83	69.6
<i>Botellina labyrinthica</i>	0	0.56	0.62	0.81	70.41
<i>Sigmoilopsis schlumbergeri</i>	0.52	0	0.61	0.8	71.21
<i>Nonionella iridea</i>	0.58	0.48	0.59	0.77	71.98
<i>Astrononion gallowayi</i>	0.63	0.2	0.57	0.74	72.72
<i>Nonionella turgida</i>	0.47	0	0.55	0.72	73.44
<i>Pullenia subcarinata</i>	0.7	0.73	0.55	0.72	74.17
<i>Sphaeroidina bulloides</i>	0.46	0	0.55	0.72	74.88
<i>Sacchorhiza ramosa</i>	0.46	0	0.54	0.7	75.59
<i>Bulimina marginata</i>	0.54	0.2	0.53	0.69	76.27
<i>Elphidium frigidum</i>	0.32	0.28	0.51	0.67	76.95
<i>Pullenia bulloides</i>	0.51	0.2	0.51	0.66	77.61
<i>Trochammina inflata</i>	0.44	0	0.5	0.65	78.26
<i>Uvigerina mediterranea</i>	0.39	0	0.5	0.65	78.9
<i>Spiroplectinella wrightii</i>	0.45	0	0.49	0.64	79.54
<i>Hyrrokkin sarcophaga</i>	0.43	0	0.47	0.62	80.16

Average dissimilarity = 66.01

Species	Cluster VI Av.Abund	Cluster II Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Cibicidoides pachyderma</i>	4.57	1.25	3.62	5.48	5.48
<i>Discanomalina coronata</i>	2.93	0	3.22	4.88	10.36
<i>Reophax diffugiformis</i>	0.2	3.08	3.07	4.65	15.01
<i>Cibicides refulgens</i>	2.56	0	2.8	4.25	19.25
<i>Cribrostomoides subglobosum</i>	0	2.6	2.79	4.23	23.48
<i>Reophax scorpiurus</i>	0.12	2.34	2.4	3.63	27.11
<i>Cassidulina laevigata</i> group	1.81	3.73	2.12	3.21	30.31
<i>Lobatula lobatula</i>	2.81	0.82	2.11	3.2	33.51
<i>Epistominella exigua</i>	0.89	2.61	1.87	2.83	36.34
<i>Melonis barleeanum</i>	1.66	3.17	1.84	2.78	39.12
<i>Fontbotia wuellerstorfi</i>	0.05	1.7	1.82	2.76	41.89
<i>Cibicidoides</i> sp.	1.7	0	1.8	2.73	44.62
<i>Paratrochammina challengerii</i>	0.04	1.01	1.04	1.57	46.19
<i>Astrononion gallowayi</i>	1.38	0.44	1.02	1.54	47.73
<i>Pullenia quinqueloba</i>	0.22	1.01	1.01	1.53	49.26
<i>Cibicides/Cibicidoides</i> juv	1.01	0.15	1.01	1.52	50.79
<i>Cassidulina carinata</i>	0.63	0.88	0.96	1.45	52.23
<i>Gavelinopsis</i> spp.	1.13	0.25	0.95	1.44	53.67
<i>Globocassidulina subglobosa</i> group	2.6	3.41	0.93	1.41	55.08
<i>Rhabdammina abyssorum</i>	0.09	0.92	0.93	1.41	56.49
<i>Angulogerina angulosa</i>	2.15	2.04	0.92	1.39	57.87
<i>Epistominella vitrea</i>	1.14	1.94	0.91	1.37	59.25
<i>Hormonisella guttifera</i>	0.04	0.85	0.88	1.33	60.58
<i>Pullenia quadriloba</i>	0.17	0.87	0.87	1.31	61.89
<i>Pullenia osloensis</i>	0.82	0.19	0.81	1.22	63.11
<i>Miliolinella subrotunda</i>	0.74	0	0.79	1.2	64.32
<i>Hanzawaia boueana</i>	0.83	0.26	0.78	1.18	65.5
<i>Bolivina</i> spp.	0.73	0	0.76	1.15	66.65
<i>Discorbinella bertheloti</i>	0.72	0	0.75	1.14	67.79
<i>Patellina corrugata</i>	0.67	0	0.69	1.05	68.84
<i>Hanzawaia concentrica</i>	0.65	0	0.68	1.03	69.87
<i>Adercotryma wrightii</i>	0.12	0.62	0.63	0.95	70.82
<i>Fissurina</i> spp.	1.16	0.71	0.62	0.94	71.77
<i>Cibicides ungerianus</i>	0.58	0.19	0.6	0.91	72.67
<i>Stainforthia fusiformis</i>	0.1	0.55	0.56	0.84	73.51
<i>Melonis pompilioides</i>	0.53	0	0.54	0.82	74.33
<i>Hyalinea balthica</i>	0.54	0	0.53	0.81	75.14
<i>Pelosina cylindrica</i>	0	0.5	0.52	0.78	75.92
<i>Pullenia subcarinata</i>	0.26	0.48	0.51	0.77	76.69
<i>Quinqueloculina viennensis</i>	0.47	0	0.51	0.77	77.46
<i>Sacchorhiza ramosa</i>	0.07	0.46	0.5	0.76	78.22
<i>Pullenia bulloides</i>	0.05	0.42	0.44	0.67	78.9
<i>Cycloforina angularis</i>	0	0.41	0.43	0.66	79.55
<i>Spiroplectinella wrightii</i>	0.4	0	0.42	0.63	80.19

Table 3.6 (continued)

Species	Average dissimilarity = 62.41				
	Cluster V Av.Abund	Cluster II Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Reophax difflugiformis</i>	0	3.08	3.12	4.99	4.99
<i>Cassidulina laevigata</i> group	1.37	3.73	2.49	4	8.99
<i>Cribrastomoides subglobosum</i>	0.25	2.6	2.41	3.87	12.86
<i>Cibicides fulgens</i>	2.25	0	2.29	3.66	16.52
<i>Cibicidoides pachyderma</i>	3.41	1.25	2.18	3.49	20.02
<i>Reophax scorpiurus</i>	0.25	2.34	2.17	3.48	23.5
<i>Uvigerina peregrina</i>	1.94	0.15	1.93	3.09	26.59
<i>Fontbotia wuellerstorfi</i>	0	1.7	1.75	2.81	29.4
<i>Angulogerina angulosa</i>	3.67	2.04	1.73	2.77	32.16
<i>Melonis barleeanum</i>	1.96	3.17	1.61	2.58	34.75
<i>Hyalinea balthica</i>	1.56	0	1.61	2.58	37.32
<i>Cibicidoides kullenbergi</i>	1.53	0	1.59	2.55	39.87
<i>Discanomalina coronata</i>	1.49	0	1.47	2.35	42.22
<i>Epistominella exigua</i>	1.29	2.61	1.37	2.2	44.42
<i>Cibicidoides</i> sp.	1.17	0	1.18	1.9	46.32
<i>Globocassidulina subglobosa</i> group	2.34	3.41	1.11	1.78	48.1
<i>Cassidulina carinata</i>	1.49	0.88	1.09	1.74	49.84
<i>Hanzawaia boueana</i>	1.17	0.26	1.08	1.73	51.57
<i>Lobatula lobatula</i>	1.62	0.82	1.05	1.69	53.26
<i>Hormonisella guttifera</i>	0	0.85	0.87	1.39	54.65
<i>Pullenia quinqueloba</i>	0.34	1.01	0.82	1.31	55.96
<i>Rhabdammina abyssorum</i>	0.15	0.92	0.8	1.28	57.24
<i>Epistominella vitrea</i>	2.66	1.94	0.78	1.26	58.49
<i>Pullenia quadriloba</i>	0.39	0.87	0.76	1.22	59.71
<i>Sacchorhiza ramosa</i>	0.46	0.46	0.76	1.21	60.93
<i>Paratrochammina challengerii</i>	0.3	1.01	0.73	1.17	62.09
<i>Planulina ariminensis</i>	0.62	0	0.63	1.02	63.11
<i>Adercotryma wrightii</i>	0.08	0.62	0.6	0.95	64.06
<i>Pullenia osloensis</i>	0.69	0.19	0.57	0.91	64.97
<i>Stainforthia fusiformis</i>	0	0.55	0.55	0.88	65.85
<i>Pullenia subcarinata</i>	0.7	0.48	0.55	0.88	66.73
<i>Nonionella iridea</i>	0.58	0.82	0.54	0.86	67.6
<i>Sigmoilopsis schlumbergeri</i>	0.52	0	0.54	0.86	68.46
<i>Pelosina cylindrica</i>	0.12	0.5	0.54	0.86	69.31
<i>Cibicides ungerianus</i>	0.56	0.19	0.51	0.82	70.13
<i>Trochammina inflata</i>	0.44	0.32	0.48	0.77	70.9
<i>Sphaeroidina bulloides</i>	0.46	0	0.48	0.77	71.67
<i>Bulimina marginata</i>	0.54	0.17	0.48	0.76	72.44
<i>Gavelinopsis</i> spp.	0.39	0.25	0.45	0.72	73.15
<i>Nonionella turgida</i>	0.47	0.39	0.44	0.71	73.87
<i>Spiroplectinella wrightii</i>	0.45	0	0.43	0.69	74.56
<i>Uvigerina mediterranea</i>	0.39	0	0.43	0.69	75.24
<i>Pullenia bulloides</i>	0.51	0.42	0.43	0.68	75.93
<i>Hyrokkin sarcophaga</i>	0.43	0	0.42	0.67	76.59
<i>Cycloforina angularis</i>	0	0.41	0.41	0.66	77.26
<i>Astrononion gallowayi</i>	0.63	0.44	0.41	0.66	77.91
<i>Melonis pomphiloides</i>	0.39	0	0.4	0.64	78.56
<i>Oridorsalis umbonatus</i>	0.14	0.33	0.38	0.6	79.16
<i>Haplofragmoides membranaceum</i>	0.36	0	0.36	0.57	79.73
<i>Cibicides/Cibicidoides</i> juv	0.28	0.15	0.35	0.56	80.29
<i>Quinqueloculina lamarckiana</i>	0.31	0.19	0.34	0.54	80.83

Species	Average dissimilarity = 55.74				
	Cluster I Av.Abund	Cluster II Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Fontbotia wuellerstorfi</i>	6.99	1.7	6.64	11.91	11.91
<i>Cassidulina laevigata</i> group	0.63	3.73	3.89	6.97	18.88
<i>Melonis barleeanum</i>	0.28	3.17	3.61	6.47	25.36
<i>Reophax difflugiformis</i>	0.2	3.08	3.56	6.39	31.75
<i>Cribrastomoides subglobosum</i>	4.74	2.6	2.62	4.7	36.45
<i>Pyrgo sarsi</i>	2.04	0	2.61	4.68	41.14
<i>Globocassidulina subglobosa</i> group	1.49	3.41	2.45	4.4	45.54
<i>Reophax scorpiurus</i>	0.45	2.34	2.42	4.34	49.88
<i>Angulogerina angulosa</i>	0.4	2.04	2.07	3.71	53.59
<i>Oridorsalis umbonatus</i>	1.47	0.33	1.43	2.56	56.15

Table 3.6 (continued)

<i>Cibicidoides pachyderma</i>	0.2	1.25	1.32	2.37	58.52
<i>Epistominella vitrea</i>	1.05	1.94	1.11	2	60.52
<i>Pullenia quadriloba</i>	0	0.87	1.08	1.93	62.45
<i>Cassidulina carinata</i>	0.8	0.88	1.04	1.86	64.31
<i>Paratrochammina challengerii</i>	0.4	1.01	0.86	1.54	65.85
<i>Gyroidinoidea laevigatus</i>	0.69	0	0.82	1.47	67.31
<i>Epistominella exigua</i>	2.35	2.61	0.8	1.43	68.75
<i>Pullenia quinqueloba</i>	0.4	1.01	0.77	1.39	70.14
<i>Adercotryma wrightii</i>	0	0.62	0.77	1.38	71.51
<i>Botellina labyrinthica</i>	0.56	0	0.67	1.2	72.71
<i>Pelosina cylindrica</i>	0	0.5	0.6	1.08	73.79
<i>Stainforthia fusiformis</i>	0.28	0.55	0.59	1.05	74.84
<i>Sacchorhiza ramosa</i>	0	0.46	0.56	1	75.84
<i>Hormonisella guttifera</i>	0.73	0.85	0.54	0.97	76.81
<i>Nonionella turgida</i>	0	0.39	0.49	0.89	77.69
<i>Cycloforina angularis</i>	0.8	0.41	0.49	0.89	78.58
<i>Hormosina globulifera</i>	0.4	0.17	0.48	0.87	79.45
<i>Astrononion gallowayi</i>	0.2	0.44	0.47	0.83	80.28

Average dissimilarity = 52.29

Species	Cluster VI Av.Abund	Cluster IV Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Discanomalina coronata</i>	2.93	0.22	2.67	5.1	5.1
<i>Bulimina marginata</i>	0.05	2.44	2.28	4.36	9.46
<i>Melonis pomphiloides</i>	0.53	2.91	2.28	4.35	13.81
<i>Cibicidoides pachyderma</i>	4.57	3.05	1.67	3.19	17
<i>Hyalinea balthica</i>	0.54	1.81	1.43	2.74	19.74
<i>Globocassidulina subglobosa</i> group	2.6	1.09	1.42	2.72	22.46
<i>Cibicides refulgens</i>	2.56	1.87	1.33	2.54	25
<i>Lobatula lobatula</i>	2.81	3.65	1.14	2.18	27.18
<i>Cassidulina carinata</i>	0.63	1.39	1.02	1.96	29.14
<i>Angulogerina angulosa</i>	2.15	1.4	0.95	1.82	30.96
<i>Epistominella vitrea</i>	1.14	0.26	0.9	1.73	32.69
<i>Cassidulina laevigata</i> group	1.81	2.22	0.88	1.68	34.37
<i>Eggerelloides scaber</i>	0.04	0.93	0.86	1.64	36
<i>Adercotryma wrightii</i>	0.12	0.97	0.85	1.63	37.64
<i>Cibicides/Cibicidoides</i> juv	1.01	0.37	0.83	1.59	39.22
<i>Epistominella exigua</i>	0.89	1.54	0.76	1.45	40.67
<i>Pullenia osloensis</i>	0.82	0.84	0.76	1.44	42.11
<i>Textularia truncata</i>	0	0.76	0.72	1.37	43.48
<i>Textularia tenuissima</i>	0.04	0.77	0.71	1.37	44.85
<i>Spiroplectinella wrightii</i>	0.4	1.11	0.7	1.34	46.19
<i>Melonis barleeanum</i>	1.66	2.29	0.69	1.31	47.5
<i>Hanzawaia boueana</i>	0.83	0.76	0.67	1.29	48.79
<i>Hanzawaia concentrica</i>	0.65	0.71	0.67	1.28	50.07
<i>Cibicidoides</i> sp.	1.7	1.46	0.66	1.27	51.34
<i>Hopkinsina atlantica</i>	0	0.7	0.66	1.27	52.61
<i>Bolivina</i> spp.	0.73	1.06	0.66	1.27	53.88
<i>Astrononion gallowayi</i>	1.38	1.4	0.65	1.25	55.13
<i>Cribrostomoides jeffreysii</i>	0.33	0.69	0.64	1.23	56.36
<i>Elphidium frigidum</i>	0.08	0.69	0.64	1.22	57.58
<i>Fissurina</i> spp.	1.16	0.86	0.63	1.2	58.78
<i>Uvigerina peregrina</i>	0.25	0.66	0.63	1.2	59.98
<i>Stainforthia fusiformis</i>	0.1	0.71	0.62	1.18	61.16
<i>Miliolinella subrotunda</i>	0.74	0.2	0.61	1.17	62.33
<i>Nonionella iridea</i>	0.72	1.33	0.59	1.14	63.47
<i>Discorbina bertheloti</i>	0.72	0.68	0.59	1.12	64.59
<i>Bigenerina cylindrica</i>	0	0.61	0.58	1.12	65.71
<i>Quinqueloculina seminula</i>	0.09	0.66	0.55	1.05	66.75
<i>Cibicides ungerianus</i>	0.58	0.1	0.53	1.02	67.77
<i>Patellina corrugata</i>	0.67	0.33	0.53	1.01	68.79
<i>Gavelinopsis</i> spp.	1.13	1.37	0.53	1	69.79
<i>Pullenia subcarinata</i>	0.26	0.54	0.51	0.97	70.76
<i>Globobulimina affinis</i>	0.17	0.63	0.51	0.97	71.73
<i>Quinqueloculina viennensis</i>	0.47	0.27	0.44	0.84	72.57
<i>Trochammina robertsoni</i>	0.05	0.46	0.44	0.83	73.4
<i>Elphidium clavatum</i>	0.26	0.3	0.41	0.78	74.19
<i>Trochammina inflata</i>	0.04	0.43	0.4	0.76	74.94

Table 3.6 (continued)

<i>Favulina</i> spp.	0.27	0.4	0.34	0.65	75.6
<i>Robertinoides</i> spp.	0.13	0.37	0.33	0.64	76.23
<i>Nonionella labradorica</i>	0.37	0.25	0.33	0.64	76.87
<i>Paratrochammina challengerii</i>	0.04	0.33	0.3	0.58	77.45
<i>Rosalina globularis</i>	0.21	0.2	0.3	0.58	78.03
<i>Sigmoilopsis schlumbergeri</i>	0	0.32	0.3	0.58	78.61
<i>Lenticulina inornata</i>	0.07	0.31	0.28	0.54	79.14
<i>Reophax diffugiformis</i>	0.2	0.26	0.28	0.53	79.68
<i>Nonionella turgida</i>	0.11	0.26	0.28	0.53	80.21
<i>Homalohedra apiopleura</i>	0.17	0.26	0.28	0.53	80.74
<i>Recurvoides trochanminiformis</i>	0.14	0.21	0.27	0.52	81.26
<i>Quinqueloculina lamarckiana</i>	0.18	0.21	0.27	0.51	81.78
<i>Verneulina propinquia</i>	0.24	0.12	0.26	0.49	82.27
<i>Reophax scorpiurus</i>	0.12	0.21	0.25	0.48	82.75
<i>Parafissurina</i> spp.	0.25	0	0.24	0.46	83.21
<i>Cibicidoides kullenbergi</i>	0.2	0.12	0.24	0.45	83.66
<i>Pyrgo elongata</i>	0.23	0.1	0.23	0.43	84.09
<i>Labrospira crassimargo</i>	0.04	0.22	0.23	0.43	84.53
<i>Textularia pseudogrammen</i>	0	0.23	0.22	0.42	84.95
<i>Stainforthia schreibersiana</i>	0.08	0.21	0.21	0.4	85.35
<i>Hyrrokkin sarcophaga</i>	0.04	0.21	0.21	0.4	85.75
<i>Spirillina vivipara</i>	0.22	0	0.21	0.39	86.14
<i>Cycloforina stalkeri</i>	0	0.21	0.2	0.39	86.53
<i>Pullenia quinqueloba</i>	0.22	0	0.2	0.38	86.91
<i>Elphidium incertum</i>	0.12	0.12	0.2	0.38	87.29
<i>Biloculinella globula</i>	0.21	0	0.2	0.38	87.67
<i>Protelphidium anglicum</i>	0	0.21	0.2	0.37	88.04
<i>Bigenerina nodosaria</i>	0	0.2	0.19	0.36	88.4
<i>Homalohedra borealis</i>	0.15	0.1	0.19	0.36	88.76
<i>Bulimina aculeata</i>	0.04	0.16	0.18	0.34	89.09
<i>Gyroidinoides laevigatus</i>	0.04	0.16	0.18	0.34	89.43
<i>Lenticulina gibba</i>	0.04	0.15	0.16	0.31	89.73
<i>Quinqueloculina cuvieriana</i>	0.09	0.12	0.16	0.3	90.04

Average dissimilarity = 55.79

Species	Cluster V	Cluster IV	Av.Diss	Contrib%	Cum. %
	Av.Abund	Av.Abund			
<i>Melonis pomphiloides</i>	0.39	2.91	2.28	4.09	4.09
<i>Epistominella vitrea</i>	2.66	0.26	2.17	3.89	7.98
<i>Angulogerina angulosa</i>	3.67	1.4	2.06	3.7	11.68
<i>Lobatula lobatula</i>	1.62	3.65	1.85	3.31	15
<i>Bulimina marginata</i>	0.54	2.44	1.74	3.11	18.11
<i>Uvigerina peregrina</i>	1.94	0.66	1.4	2.51	20.62
<i>Cibicidoides kullenbergi</i>	1.53	0.12	1.31	2.35	22.97
<i>Cassidulina laevigata</i> group	1.37	2.22	1.21	2.17	25.14
<i>Discanomalina coronata</i>	1.49	0.22	1.17	2.11	27.24
<i>Globocassidulina subglobosa</i> group	2.34	1.09	1.17	2.1	29.34
<i>Cibicidoides pachyderma</i>	3.41	3.05	0.97	1.74	31.08
<i>Gavelinopsis</i> spp.	0.39	1.37	0.96	1.72	32.8
<i>Eggerelloides scaber</i>	0	0.93	0.84	1.5	34.3
<i>Bolivina</i> spp.	0.27	1.06	0.83	1.48	35.78
<i>Cibicides refulgens</i>	2.25	1.87	0.82	1.47	37.26
<i>Adercotryma wrightii</i>	0.08	0.97	0.82	1.47	38.73
<i>Hanzawaia boueana</i>	1.17	0.76	0.78	1.41	40.14
<i>Cassidulina carinata</i>	1.49	1.39	0.78	1.39	41.53
<i>Hyalinea balthica</i>	1.56	1.81	0.76	1.37	42.9
<i>Nonionella iridea</i>	0.58	1.33	0.73	1.31	44.21
<i>Astrononion gallowayi</i>	0.63	1.4	0.72	1.29	45.49
<i>Textularia tenuissima</i>	0	0.77	0.7	1.26	46.75
<i>Textularia truncata</i>	0	0.76	0.69	1.23	47.99
<i>Cibicidoides</i> sp.	1.17	1.46	0.68	1.22	49.2
<i>Melonis barleeanum</i>	1.96	2.29	0.66	1.19	50.4
<i>Pullenia osloensis</i>	0.69	0.84	0.65	1.17	51.57
<i>Stainforthia fusiformis</i>	0	0.71	0.64	1.15	52.72
<i>Hopkinsina atlantica</i>	0	0.7	0.64	1.14	53.86
<i>Discorbinella bertheloti</i>	0.21	0.68	0.63	1.13	54.99
<i>Spiroplectinella wrightii</i>	0.45	1.11	0.63	1.13	56.11
<i>Hanzawaia concentrica</i>	0.16	0.71	0.62	1.12	57.23

Table 3.6 (continued)

<i>Cribrastomoides jeffreysii</i>	0.06	0.69	0.61	1.1	58.33
<i>Elphidium frigidum</i>	0.32	0.69	0.61	1.09	59.42
<i>Quinqueloculina seminula</i>	0	0.66	0.6	1.08	60.49
<i>Epistominella exigua</i>	1.29	1.54	0.56	1.01	61.5
<i>Bigenerina cylindrica</i>	0	0.61	0.56	1	62.5
<i>Pullenia subcarinata</i>	0.7	0.54	0.54	0.97	63.47
<i>Globobulimina affinis</i>	0.06	0.63	0.53	0.95	64.42
<i>Planulina ariminensis</i>	0.62	0.11	0.52	0.94	65.36
<i>Cibicides ungerianus</i>	0.56	0.1	0.48	0.86	66.22
<i>Pullenia bulloides</i>	0.51	0	0.47	0.84	67.06
<i>Sigmoilopsis schlumbergeri</i>	0.52	0.32	0.44	0.79	67.85
<i>Fissurina</i> spp.	0.91	0.86	0.43	0.77	68.62
<i>Sphaeroidina bulloides</i>	0.46	0	0.43	0.76	69.38
<i>Cibicides/Cibicidooides</i> juv	0.28	0.37	0.43	0.76	70.15
<i>Sacchorhiza ramosa</i>	0.46	0	0.42	0.75	70.9
<i>Trochammina robertsoni</i>	0.06	0.46	0.42	0.74	71.64
<i>Nonionella turgida</i>	0.47	0.26	0.41	0.74	72.38
<i>Trochammina inflata</i>	0.44	0.43	0.41	0.73	73.11
<i>Uvigerina mediterranea</i>	0.39	0	0.38	0.68	73.79
<i>Reophax scorpiurus</i>	0.25	0.21	0.36	0.65	74.44
<i>Quinqueloculina viennensis</i>	0.29	0.27	0.36	0.65	75.09
<i>Hyrrokkin sarcophaga</i>	0.43	0.21	0.36	0.65	75.74
<i>Pullenia quadriloba</i>	0.39	0	0.35	0.63	76.37
<i>Rosalina globularis</i>	0.29	0.2	0.34	0.61	76.98
<i>Gyroidinoides laevigatus</i>	0.31	0.16	0.34	0.61	77.59
<i>Favulinina</i> spp.	0.33	0.4	0.33	0.59	78.18
<i>Robertinooides</i> spp.	0.17	0.37	0.33	0.59	78.77
<i>Miliolinella subrotunda</i>	0.29	0.2	0.33	0.59	79.37
<i>Patellina corrugata</i>	0.16	0.33	0.33	0.59	79.96
<i>Haplofragmoides membranaceum</i>	0.36	0	0.32	0.57	80.53

Average dissimilarity = 80.66

Species	Cluster I		Cluster IV		
	Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Fontbotia wuellerstorfi</i>	6.99	0	7.65	9.48	9.48
<i>Cribrastomoides subglobosum</i>	4.74	0	5.13	6.36	15.84
<i>Melonis pomphiloides</i>	0	2.91	3.16	3.92	19.76
<i>Cibicidooides pachyderma</i>	0.2	3.05	3.13	3.88	23.64
<i>Lobatula lobatula</i>	1.01	3.65	2.86	3.55	27.18
<i>Bulimina marginata</i>	0.2	2.44	2.45	3.03	30.22
<i>Pyrgo sarsi</i>	2.04	0	2.26	2.81	33.02
<i>Melonis barleeanum</i>	0.28	2.29	2.21	2.73	35.76
<i>Hyalinea balthica</i>	0	1.81	1.99	2.47	38.22
<i>Cibicides refulgens</i>	0.2	1.87	1.83	2.27	40.49
<i>Cassidulina laevigata</i> group	0.63	2.22	1.74	2.15	42.65
<i>Oridorsalis umbonatus</i>	1.47	0	1.6	1.98	44.63
<i>Cibicidooides</i> sp.	0	1.46	1.59	1.97	46.6
<i>Gavelinopsis</i> spp.	0	1.37	1.48	1.84	48.43
<i>Astrononion gallowayi</i>	0.2	1.4	1.3	1.61	50.05
<i>Spiroplectinella wrightii</i>	0	1.11	1.2	1.49	51.53
<i>Bolivina</i> spp.	0	1.06	1.16	1.44	52.98
<i>Angulogerina angulosa</i>	0.4	1.4	1.12	1.39	54.36
<i>Adercotryma wrightii</i>	0	0.97	1.04	1.29	55.65
<i>Rhabdammina abyssorum</i>	0.94	0	1.01	1.26	56.91
<i>Eggerelloides scaber</i>	0	0.93	1	1.24	58.15
<i>Cassidulina carinata</i>	0.8	1.39	0.93	1.15	59.3
<i>Epistominella exigua</i>	2.35	1.54	0.92	1.14	60.44
<i>Nonionella iridea</i>	0.48	1.33	0.91	1.13	61.57
<i>Pullenia osloensis</i>	0	0.84	0.91	1.12	62.69
<i>Cycloforina angularis</i>	0.8	0	0.87	1.08	63.77
<i>Epistominella vitrea</i>	1.05	0.26	0.86	1.07	64.84
<i>Textularia tenuissima</i>	0	0.77	0.84	1.05	65.89
<i>Hanzawaia boueana</i>	0	0.76	0.82	1.02	66.91
<i>Textularia truncata</i>	0	0.76	0.82	1.02	67.94
<i>Globocassidulina subglobosa</i> group	1.49	1.09	0.79	0.98	68.91
<i>Hormonisella guttifera</i>	0.73	0	0.78	0.97	69.88
<i>Hanzawaia concentrica</i>	0	0.71	0.77	0.95	70.84
<i>Hopkinsina atlantica</i>	0	0.7	0.76	0.95	71.78

Table 3.6 (continued)

<i>Cibrostomoides jeffreysii</i>	0	0.69	0.75	0.93	72.71
<i>Discorbinella bertheloti</i>	0	0.68	0.74	0.92	73.63
<i>Gyroldinoides laevigatus</i>	0.69	0.16	0.72	0.9	74.53
<i>Uvigerina peregrina</i>	0	0.66	0.72	0.9	75.43
<i>Quinqueloculina seminula</i>	0	0.66	0.72	0.89	76.32
<i>Elphidium frigidum</i>	0.28	0.69	0.68	0.85	77.16
<i>Globobulimina affinis</i>	0	0.63	0.68	0.84	78
<i>Bigenerina cylindrica</i>	0	0.61	0.67	0.83	78.84
<i>Stainforthia fusiformis</i>	0.28	0.71	0.59	0.73	79.57
<i>Botellina labyrinthica</i>	0.56	0	0.58	0.72	80.29
<i>Trochammina robertsoni</i>	0	0.46	0.51	0.64	80.93

Species	Average dissimilarity = 68.04				
	Cluster II Av.Abund	Cluster IV Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Melonis pomphiloides</i>	0	2.91	2.8	4.11	4.11
<i>Lobatula lobatula</i>	0.82	3.65	2.71	3.99	8.1
<i>Reophax diffugiformis</i>	3.08	0.26	2.7	3.97	12.07
<i>Cibrostomoides subglobosum</i>	2.6	0	2.51	3.69	15.76
<i>Globocassidulina subglobosa</i> group	3.41	1.09	2.25	3.3	19.06
<i>Bulimina marginata</i>	0.17	2.44	2.19	3.23	22.28
<i>Reophax scorpiurus</i>	2.34	0.21	2.06	3.03	25.31
<i>Cibicides refulgens</i>	0	1.87	1.8	2.65	27.96
<i>Cibicidoides pachyderma</i>	1.25	3.05	1.76	2.58	30.54
<i>Hyalinea balthica</i>	0	1.81	1.76	2.58	33.13
<i>Fontbotia wuellerstorfi</i>	1.7	0	1.66	2.43	35.56
<i>Epistominella vitrea</i>	1.94	0.26	1.62	2.38	37.94
<i>Cassidulina laevigata</i> group	3.73	2.22	1.45	2.13	40.08
<i>Cibicidoides</i> sp.	0	1.46	1.4	2.06	42.14
<i>Melonis barleeanum</i>	3.17	2.29	1.37	2.01	44.15
<i>Gavelinopsis</i> spp.	0.25	1.37	1.09	1.6	45.75
<i>Spiroplectinella wrightii</i>	0	1.11	1.06	1.56	47.31
<i>Epistominella exigua</i>	2.61	1.54	1.03	1.52	48.83
<i>Bolivina</i> spp.	0	1.06	1.03	1.51	50.34
<i>Cassidulina carinata</i>	0.88	1.39	1.03	1.51	51.85
<i>Pullenia quinqueloba</i>	1.01	0	0.98	1.44	53.29
<i>Astronionion gallowayi</i>	0.44	1.4	0.92	1.35	54.64
<i>Eggerelloides scaber</i>	0	0.93	0.89	1.31	55.94
<i>Rhabdammina abyssorum</i>	0.92	0	0.89	1.31	57.25
<i>Pullenia quadriloba</i>	0.87	0	0.83	1.22	58.47
<i>Hormonisella guttifera</i>	0.85	0	0.82	1.21	59.68
<i>Angulogerina angulosa</i>	2.04	1.4	0.79	1.16	60.83
<i>Pullenia osloensis</i>	0.19	0.84	0.77	1.13	61.96
<i>Textularia tenuissima</i>	0	0.77	0.75	1.1	63.06
<i>Textularia truncata</i>	0	0.76	0.73	1.07	64.13
<i>Hanzawaia concentrica</i>	0	0.71	0.68	1	65.13
<i>Hanzawaia boueana</i>	0.26	0.76	0.68	1	66.12
<i>Adercotryma wrightii</i>	0.62	0.97	0.66	0.97	67.1
<i>Cibrostomoides jeffreysii</i>	0	0.69	0.66	0.97	68.07
<i>Elphidium frigidum</i>	0	0.69	0.66	0.97	69.04
<i>Discorbinella bertheloti</i>	0	0.68	0.66	0.97	70.01
<i>Paratrochammina challengerii</i>	1.01	0.33	0.65	0.96	70.96
<i>Quinqueloculina seminula</i>	0	0.66	0.64	0.94	71.9
<i>Bigenerina cylindrica</i>	0	0.61	0.59	0.87	72.77
<i>Hopkinsina atlantica</i>	0.19	0.7	0.57	0.83	73.61
<i>Uvigerina peregrina</i>	0.15	0.66	0.55	0.81	74.42
<i>Nonionella iridea</i>	0.82	1.33	0.53	0.78	75.2
<i>Stainforthia fusiformis</i>	0.55	0.71	0.52	0.76	75.96
<i>Globobulimina affinis</i>	0.17	0.63	0.5	0.74	76.7
<i>Pullenia subcarinata</i>	0.48	0.54	0.47	0.69	77.39
<i>Pelosina cylindrica</i>	0.5	0	0.47	0.69	78.07
<i>Trochammina robertsoni</i>	0	0.46	0.45	0.67	78.74
<i>Trochammina inflata</i>	0.32	0.43	0.45	0.66	79.4
<i>Sacchorhiza ramosa</i>	0.46	0	0.43	0.64	80.04

Table 3.6 (continued)

<i>Cribrastomoides jeffreysii</i>	0.06	0.69	0.61	1.1	58.33
<i>Elphidium frigidum</i>	0.32	0.69	0.61	1.09	59.42
<i>Quinqueloculina seminula</i>	0	0.66	0.6	1.08	60.49
<i>Epistominella exigua</i>	1.29	1.54	0.56	1.01	61.5
<i>Bigenerina cylindrica</i>	0	0.61	0.56	1	62.5
<i>Pullenia subcarinata</i>	0.7	0.54	0.54	0.97	63.47
<i>Globobulimina affinis</i>	0.06	0.63	0.53	0.95	64.42
<i>Planulina ariminensis</i>	0.62	0.11	0.52	0.94	65.36
<i>Cibicides ungerianus</i>	0.56	0.1	0.48	0.86	66.22
<i>Pullenia bulloides</i>	0.51	0	0.47	0.84	67.06
<i>Sigmoilopsis schlumbergeri</i>	0.52	0.32	0.44	0.79	67.85
<i>Fissurina</i> spp.	0.91	0.86	0.43	0.77	68.62
<i>Sphaeroidina bulloides</i>	0.46	0	0.43	0.76	69.38
<i>Cibicides/Cibicidoides</i> juv	0.28	0.37	0.43	0.76	70.15
<i>Sacchorhiza ramosa</i>	0.46	0	0.42	0.75	70.9
<i>Trochammina robertsoni</i>	0.06	0.46	0.42	0.74	71.64
<i>Nonionella turgida</i>	0.47	0.26	0.41	0.74	72.38
<i>Trochammina inflata</i>	0.44	0.43	0.41	0.73	73.11
<i>Uvigerina mediterranea</i>	0.39	0	0.38	0.68	73.79
<i>Reophax scorpiurus</i>	0.25	0.21	0.36	0.65	74.44
<i>Quinqueloculina viennensis</i>	0.29	0.27	0.36	0.65	75.09
<i>Hyrrokkin sarcophaga</i>	0.43	0.21	0.36	0.65	75.74
<i>Pullenia quadriloba</i>	0.39	0	0.35	0.63	76.37
<i>Rosalina globularis</i>	0.29	0.2	0.34	0.61	76.98
<i>Gyroidinoides laevigatus</i>	0.31	0.16	0.34	0.61	77.59
<i>Favulina</i> spp.	0.33	0.4	0.33	0.59	78.18
<i>Robertinoidea</i> spp.	0.17	0.37	0.33	0.59	78.77
<i>Miliolinella subrotunda</i>	0.29	0.2	0.33	0.59	79.37
<i>Patellina corrugata</i>	0.16	0.33	0.33	0.59	79.96
<i>Haplofragmoides membranaceum</i>	0.36	0	0.32	0.57	80.53

Average dissimilarity = 80.66

Species	Cluster I		Cluster IV		Cum.%
	Av.Abund	Av.Abund	Av.Diss	Contrib%	
<i>Fontbotia wuellerstorfi</i>	6.99	0	7.65	9.48	9.48
<i>Cribrastomoides subglobosum</i>	4.74	0	5.13	6.36	15.84
<i>Melonis pompilioides</i>	0	2.91	3.16	3.92	19.76
<i>Cibicidoides pachyderma</i>	0.2	3.05	3.13	3.88	23.64
<i>Lobatula lobatula</i>	1.01	3.65	2.86	3.55	27.18
<i>Bulimina marginata</i>	0.2	2.44	2.45	3.03	30.22
<i>Pyrgo sarsi</i>	2.04	0	2.26	2.81	33.02
<i>Melonis barleeanum</i>	0.28	2.29	2.21	2.73	35.76
<i>Hyalinea balthica</i>	0	1.81	1.99	2.47	38.22
<i>Cibicides refulgens</i>	0.2	1.87	1.83	2.27	40.49
<i>Cassidulina laevigata</i> group	0.63	2.22	1.74	2.15	42.65
<i>Oridorsalis umbonatus</i>	1.47	0	1.6	1.98	44.63
<i>Cibicidoides</i> sp.	0	1.46	1.59	1.97	46.6
<i>Gavelinopsis</i> spp.	0	1.37	1.48	1.84	48.43
<i>Astrononion gallowayi</i>	0.2	1.4	1.3	1.61	50.05
<i>Spiroplectinella wrightii</i>	0	1.11	1.2	1.49	51.53
<i>Bolivina</i> spp.	0	1.06	1.16	1.44	52.98
<i>Angulogerina angulosa</i>	0.4	1.4	1.12	1.39	54.36
<i>Adercotryma wrightii</i>	0	0.97	1.04	1.29	55.65
<i>Rhabdammina abyssorum</i>	0.94	0	1.01	1.26	56.91
<i>Eggerelloides scaber</i>	0	0.93	1	1.24	58.15
<i>Cassidulina carinata</i>	0.8	1.39	0.93	1.15	59.3
<i>Epistominella exigua</i>	2.35	1.54	0.92	1.14	60.44
<i>Nonionella iridea</i>	0.48	1.33	0.91	1.13	61.57
<i>Pullenia osloensis</i>	0	0.84	0.91	1.12	62.69
<i>Cycloforina angularis</i>	0.8	0	0.87	1.08	63.77
<i>Epistominella vitrea</i>	1.05	0.26	0.86	1.07	64.84
<i>Textularia tenuissima</i>	0	0.77	0.84	1.05	65.89
<i>Hanzawaia boueana</i>	0	0.76	0.82	1.02	66.91
<i>Textularia truncata</i>	0	0.76	0.82	1.02	67.94
<i>Globocassidulina subglobosa</i> group	1.49	1.09	0.79	0.98	68.91
<i>Hormonisella guttifera</i>	0.73	0	0.78	0.97	69.88
<i>Hanzawaia concentrica</i>	0	0.71	0.77	0.95	70.84
<i>Hopkinsina atlantica</i>	0	0.7	0.76	0.95	71.78

Table 3.6 (continued)

<i>Cribrostomoides jeffreysii</i>	0.06	0.69	0.61	1.1	58.33
<i>Elphidium frigidum</i>	0.32	0.69	0.61	1.09	59.42
<i>Quinqueloculina seminula</i>	0	0.66	0.6	1.08	60.49
<i>Epistominella exigua</i>	1.29	1.54	0.56	1.01	61.5
<i>Bigenerina cylindrica</i>	0	0.61	0.56	1	62.5
<i>Pullenia subcarinata</i>	0.7	0.54	0.54	0.97	63.47
<i>Globobulimina affinis</i>	0.06	0.63	0.53	0.95	64.42
<i>Planulina ariminensis</i>	0.62	0.11	0.52	0.94	65.36
<i>Cibicides ungerianus</i>	0.56	0.1	0.48	0.86	66.22
<i>Pullenia bulloides</i>	0.51	0	0.47	0.84	67.06
<i>Sigmoilopsis schlumbergeri</i>	0.52	0.32	0.44	0.79	67.85
<i>Fissurina</i> spp.	0.91	0.86	0.43	0.77	68.62
<i>Sphaeroidina bulloides</i>	0.46	0	0.43	0.76	69.38
<i>Cibicides/Cibicidoides</i> juv	0.28	0.37	0.43	0.76	70.15
<i>Sacchorhiza ramosa</i>	0.46	0	0.42	0.75	70.9
<i>Trochammina robertsoni</i>	0.06	0.46	0.42	0.74	71.64
<i>Nonionella turgida</i>	0.47	0.26	0.41	0.74	72.38
<i>Trochammina inflata</i>	0.44	0.43	0.41	0.73	73.11
<i>Uvigerina mediterranea</i>	0.39	0	0.38	0.68	73.79
<i>Reophax scorpiurus</i>	0.25	0.21	0.36	0.65	74.44
<i>Quinqueloculina viennensis</i>	0.29	0.27	0.36	0.65	75.09
<i>Hyrrokkin sarcophaga</i>	0.43	0.21	0.36	0.65	75.74
<i>Pullenia quadriloba</i>	0.39	0	0.35	0.63	76.37
<i>Rosalina globularis</i>	0.29	0.2	0.34	0.61	76.98
<i>Gyroidinoides laevigatus</i>	0.31	0.16	0.34	0.61	77.59
<i>Favulina</i> spp.	0.33	0.4	0.33	0.59	78.18
<i>Robertinoides</i> spp.	0.17	0.37	0.33	0.59	78.77
<i>Miliolinella subrotunda</i>	0.29	0.2	0.33	0.59	79.37
<i>Patellina corrugata</i>	0.16	0.33	0.33	0.59	79.96
<i>Haplofragmoides membranaceum</i>	0.36	0	0.32	0.57	80.53

Average dissimilarity = 80.66

Species	Cluster I Av.Abund	Cluster IV Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Fontbotia wuellerstorfi</i>	6.99	0	7.65	9.48	9.48
<i>Cribrostomoides subglobosum</i>	4.74	0	5.13	6.36	15.84
<i>Melonis pomphiloides</i>	0	2.91	3.16	3.92	19.76
<i>Cibicidoides pachyderma</i>	0.2	3.05	3.13	3.88	23.64
<i>Lobatula lobatula</i>	1.01	3.65	2.86	3.55	27.18
<i>Bulimina marginata</i>	0.2	2.44	2.45	3.03	30.22
<i>Pyrgo sarsi</i>	2.04	0	2.26	2.81	33.02
<i>Melonis barleeanum</i>	0.28	2.29	2.21	2.73	35.76
<i>Hyalinea balthica</i>	0	1.81	1.99	2.47	38.22
<i>Cibicides refulgens</i>	0.2	1.87	1.83	2.27	40.49
<i>Cassidulina laevigata</i> group	0.63	2.22	1.74	2.15	42.65
<i>Oridorsalis umbonatus</i>	1.47	0	1.6	1.98	44.63
<i>Cibicidoides</i> sp.	0	1.46	1.59	1.97	46.6
<i>Gavelinopsis</i> spp.	0	1.37	1.48	1.84	48.43
<i>Astrononion gallowayi</i>	0.2	1.4	1.3	1.61	50.05
<i>Spiroplectinella wrightii</i>	0	1.11	1.2	1.49	51.53
<i>Bolivina</i> spp.	0	1.06	1.16	1.44	52.98
<i>Angulogerina angulosa</i>	0.4	1.4	1.12	1.39	54.36
<i>Adercotryma wrightii</i>	0	0.97	1.04	1.29	55.65
<i>Rhabdammina abyssorum</i>	0.94	0	1.01	1.26	56.91
<i>Eggerelloides scaber</i>	0	0.93	1	1.24	58.15
<i>Cassidulina carinata</i>	0.8	1.39	0.93	1.15	59.3
<i>Epistominella exigua</i>	2.35	1.54	0.92	1.14	60.44
<i>Nonionella iridea</i>	0.48	1.33	0.91	1.13	61.57
<i>Pullenia osloensis</i>	0	0.84	0.91	1.12	62.69
<i>Cycloforina angularis</i>	0.8	0	0.87	1.08	63.77
<i>Epistominella vitrea</i>	1.05	0.26	0.86	1.07	64.84
<i>Textularia tenuissima</i>	0	0.77	0.84	1.05	65.89
<i>Hanzawaia boueana</i>	0	0.76	0.82	1.02	66.91
<i>Textularia truncata</i>	0	0.76	0.82	1.02	67.94
<i>Globocassidulina subglobosa</i> group	1.49	1.09	0.79	0.98	68.91
<i>Hormonisella guttifera</i>	0.73	0	0.78	0.97	69.88
<i>Hanzawaia concentrica</i>	0	0.71	0.77	0.95	70.84
<i>Hopkinsina atlantica</i>	0	0.7	0.76	0.95	71.78

Table 3.6 (continued)

Species	Average dissimilarity = 73.75				
	Cluster II Av.Abund	Cluster III Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Globobulimina affinis</i>	0.17	4.53	4.63	6.27	6.27
<i>Globocassidulina subglobosa</i> group	3.41	0.28	3.33	4.51	10.79
<i>Reophax difflugiformis</i>	3.08	0	3.24	4.4	15.19
<i>Hyalinea balthica</i>	0	2.73	2.92	3.95	19.14
<i>Eggerelloides scaber</i>	0	2.52	2.67	3.62	22.76
<i>Textularia tenuissima</i>	0	2.12	2.24	3.03	25.79
<i>Cassidulina laevigata</i> group	3.73	1.69	2.16	2.93	28.72
<i>Bolivina</i> spp.	0	1.98	2.12	2.87	31.59
<i>Bigenerina cylindrica</i>	0	1.99	2.1	2.85	34.44
<i>Cribrostomoides subglobosum</i>	2.6	0.66	2.07	2.81	37.24
<i>Reophax scorpiurus</i>	2.34	0.44	2.04	2.76	40.01
<i>Pullenia subcarinata</i>	0.48	2.1	2.04	2.76	42.77
<i>Melonis pompilioides</i>	0	1.9	1.99	2.7	45.47
<i>Epistominella exigua</i>	2.61	0.84	1.87	2.54	48.01
<i>Angulogerina angulosa</i>	2.04	0.28	1.86	2.53	50.54
<i>Fontbotia wuellerstorfi</i>	1.7	0	1.83	2.48	53.01
<i>Epistominella vitrea</i>	1.94	0.28	1.77	2.4	55.41
<i>Melonis barleeanum</i>	3.17	1.7	1.73	2.34	57.76
<i>Bulimina marginata</i>	0.17	1.78	1.71	2.31	60.07
<i>Hopkinsina atlantica</i>	0.19	1.55	1.56	2.12	62.19
<i>Cassidulina carinata</i>	0.88	2	1.55	2.1	64.29
<i>Uvigerina peregrina</i>	0.15	1.44	1.35	1.84	66.13
<i>Trochammina robertsoni</i>	0	1.14	1.2	1.63	67.76
<i>Pullenia quinqueloba</i>	1.01	0	1.08	1.46	69.22
<i>Pullenia quadriloba</i>	0.87	0	0.91	1.24	70.46
<i>Hormonisella guttifera</i>	0.85	0	0.91	1.23	71.69
<i>Paratrochammina challengerii</i>	1.01	0.2	0.85	1.15	72.84
<i>Pullenia osloensis</i>	0.19	0.79	0.84	1.14	73.98
<i>Nonionella labradorica</i>	0	0.78	0.82	1.12	75.1
<i>Cibicidooides pachyderma</i>	1.25	0.72	0.82	1.12	76.21
<i>Stainforthia fusiformis</i>	0.55	1.25	0.79	1.07	77.28
<i>Nonionella iridea</i>	0.82	0.31	0.67	0.9	78.19
<i>Stainforthia schreibersiana</i>	0	0.62	0.65	0.88	79.07
<i>Elphidium frigidum</i>	0	0.58	0.61	0.82	79.89
<i>Adercotryma wrightii</i>	0.62	0.53	0.55	0.75	80.64

Species	Average dissimilarity = 56.37				
	Cluster IV Av.Abund	Cluster III Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Globobulimina affinis</i>	0.63	4.53	3.68	6.53	6.53
<i>Lobatula lobatula</i>	3.65	0.74	2.72	4.83	11.36
<i>Cibicidooides pachyderma</i>	3.05	0.72	2.2	3.91	15.27
<i>Pullenia subcarinata</i>	0.54	2.1	1.79	3.18	18.45
<i>Cibicides refulgens</i>	1.87	0.25	1.52	2.69	21.14
<i>Eggerelloides scaber</i>	0.93	2.52	1.5	2.66	23.8
<i>Textularia tenuissima</i>	0.77	2.12	1.35	2.4	26.2
<i>Bigenerina cylindrica</i>	0.61	1.99	1.29	2.28	28.48
<i>Melonis pompilioides</i>	2.91	1.9	1.24	2.2	30.68
<i>Hopkinsina atlantica</i>	0.7	1.55	1.23	2.19	32.87
<i>Hyalinea balthica</i>	1.81	2.73	1.23	2.18	35.04
<i>Cibicidooides</i> sp.	1.46	0.18	1.2	2.13	37.17
<i>Gavelinopsis</i> spp.	1.37	0.12	1.17	2.08	39.25
<i>Cassidulina carinata</i>	1.39	2	1.13	2.01	41.26
<i>Bolivina</i> spp.	1.06	1.98	1.07	1.9	43.16
<i>Angulogerina angulosa</i>	1.4	0.28	1.07	1.89	45.05
<i>Nonionella iridea</i>	1.33	0.31	0.96	1.7	46.75
<i>Bulimina marginata</i>	2.44	1.78	0.95	1.69	48.44
<i>Spiroplectinella wrightii</i>	1.11	0.12	0.93	1.65	50.09
<i>Cassidulina laevigata</i> group	2.22	1.69	0.91	1.62	51.71
<i>Pullenia osloensis</i>	0.84	0.79	0.88	1.55	53.26
<i>Globocassidulina subglobosa</i> group	1.09	0.28	0.79	1.41	54.67
<i>Uvigerina peregrina</i>	0.66	1.44	0.79	1.4	56.07
<i>Epistominella exigua</i>	1.54	0.84	0.73	1.29	57.36
<i>Textularia truncata</i>	0.76	0	0.71	1.27	58.62
<i>Hanzawaia boueana</i>	0.76	0	0.71	1.26	59.88

Table 3.6 (continued)

<i>Astrononion gallowayi</i>	1.4	0.67	0.7	1.25	61.13
<i>Adercotryma wrightii</i>	0.97	0.53	0.67	1.19	62.32
<i>Hanzawaia concentrica</i>	0.71	0	0.66	1.18	63.49
<i>Trochammina robertsoni</i>	0.46	1.14	0.65	1.16	64.65
<i>Fissurina</i> spp.	0.86	0.3	0.63	1.12	65.78
<i>Quinqueloculina seminula</i>	0.66	0	0.62	1.1	66.88
<i>Cibrostomoides jeffreysii</i>	0.69	0.31	0.62	1.1	67.98
<i>Cibrostomoides subglobosum</i>	0	0.66	0.62	1.1	69.08
<i>Melonis barleeanum</i>	2.29	1.7	0.62	1.1	70.18
<i>Discorbarella bertheloti</i>	0.68	0.18	0.61	1.08	71.25
<i>Stainforthia fusiformis</i>	0.71	1.25	0.58	1.03	72.29
<i>Elphidium frigidum</i>	0.69	0.58	0.54	0.95	73.24
<i>Nonionella labradorica</i>	0.25	0.78	0.53	0.95	74.19
<i>Rhabdammina abyssorum</i>	0	0.56	0.52	0.91	75.1
<i>Stainforthia schreibersiana</i>	0.21	0.62	0.51	0.91	76.01
<i>Nonionella turgida</i>	0.26	0.78	0.51	0.91	76.92
<i>Cibicides/Cibicidoides</i> juv	0.37	0.36	0.46	0.81	77.73
<i>Stainforthia skagerakensis</i>	0	0.47	0.44	0.78	78.51
<i>Cibicides ungerianus</i>	0.1	0.48	0.43	0.77	79.27
<i>Trochammina inflata</i>	0.43	0	0.4	0.71	79.98
<i>Reophax scorpiurus</i>	0.21	0.44	0.39	0.69	80.67

patterns of depth segregation in virtually all marine basins (e.g., Mackensen et al., 1985). The reason for this partitioning is a combination of many ecological parameters such as temperature, salinity, hydrodynamics, oxygen concentration of the bottom waters, and organic matter flux (e.g., Sen Gupta, 1999; Van der Zwaan et al. 1999; Gooday, 2003; Murray, 2006) that vary with depth. Two bathymetry-controlled clusters separate at a high level of the cluster analysis (Fig. 3.5). Cluster A includes all samples from 889 to 2098 m and cluster B integrates all samples above 889 m (Fig. 3.4).

The depth distribution of the dead (unstained) benthic foraminifera in the studied samples generally corresponds to the known depth of occurrence of these species on the Norwegian slope (Mackensen et al. 1985; Fig. 3.4). *Cibrostomoides subglobosum*, *E. exigua*, *F. wuellerstorfi*, the *Globocassidulina subglobosa* group, the *C. laevigata* group, and *Epistominella vitrea* dominate the deep-water fauna (Fig. 3.4). They thrive in the cold waters of the NSDW and NSBW. Contrarily, *Cibicidoides pachyderma*, *L. lobatula*, *M. barleeanum*, *A. angulosa*, the *G. subglobosa* group, *Cibicides refulgens*, and the *C. laevigata* group dominate the shallow-water fauna. They thrive in the high bottom currents (Mackensen et al., 1985) mainly produced by internal tides (Holbrook et al., 2009). The close similarity between living (stained) and dead foraminiferal assemblages described by Mackensen et al. (1985) suggests that our assemblages also strongly represent the actual living fauna.

3.5.2.2 Characterization of “off-reef” sites

A separation between “on-reef” and “off-reef” can only

be distinguished in the lower level clusters (Fig. 3.5). Six clusters separate at the 49% of the Bray-Curties Similarity. Cluster I combines all the deep-sea samples from below 1800 m water depth. The foraminiferal assemblage is dominated by *F. wuellerstorfi*, *C. subglobosum*, *Oridorsalis umbonatus*, *P. sarsi*, and *E. exigua* (Tab. 3.5; Fig. 3.4). *Fontbotia wuellerstorfi* and *C. subglobosum* live in well-ventilated waters with low organic carbon flux (Lutze and Thiel, 1989; Linke and Lutze, 1993; Mackensen et al., 1995; Gooday, 2003; Murray, 2006). *Epistominella exigua* indicates seasonal deposits of phytodetritus (Gooday, 1993; Loubere, 1998; Smart et al., 1994; Thomas et al., 1995; Thomas and Gooday, 1996; Suhr et al., 2003) caused by seasonal phytoplankton. *Oridorsalis umbonatus* is a typical species of nutrient-poor, cold, deep Atlantic waters (Belanger and Streeter, 1980; Mackensen et al., 1985; Murray et al., 1986). Therefore, this cluster is interpreted to represent nutrient-poor deep-sea environments influenced by seasonal pulses of nutrient availability.

Cluster II combines samples from the continental slope between 1800 m and 800 m water depth. The fauna is dominated by the *G. subglobosa* group, the *C. laevigata* group, *Reophax* spp., *Epistominella* spp., *M. barleeanum*, and *C. subglobosum* (Tab. 3.5; Fig. 3.4). The *G. subglobosa* group and *Epistominella* spp. are indicator species for seasonally enhanced phytodetritus fluxes. The abundance of the *C. laevigata* group, and *M. barleeanum* indicate a generally high organic carbon flux (Corliss, 1985; Gooday, 1986; Mackensen and Hald, 1988; Caralp, 1989; Murray, 2006). Therefore, this cluster is interpreted to represent environments affected by a higher background nutrient flux than

Cluster I but also showing seasonal pulses of nutrient availability.

Cluster III combines all “off-reef” samples from the Skagerrak. The fauna is dominated by *Globobulimina affinis*, *Eggerelloides scaber*, *H. balthica*, *Bigenerina cylindrica*, *Textularia tenuissima*, *Bulimina marginata*, *Melonis* spp., and *Bolivina* spp. These species/groups are generally found in poorly oxygenated environments (Bernhard, 1986; Sen Gupta and Machain-Castillo, 1993; Jorissen et al., 1995; Bernhard and Sen Gupta, 1999) in fine-grained sediments rich in refractory organic matter (Fontanier et al., 2002, 2005), e.g., in the Northern Skagerrak region (Corliss and Van Weering, 1993; Bergsten et al., 1996; Alve and Murray, 1997). Therefore, this cluster is interpreted

to represent an environment with high organic matter (possibly refractory) flux and low oxygen levels.

3.5.2.3 Characterization of “on-reef” sites

Cluster IV combines all samples from the coral reefs in the Oslofjord. The benthic foraminiferal assemblage is dominated by *L. lobatula*, *C. pachyderma*, *Melonis* spp., *B. marginata*, the *C. laevigata* group, and *C. refulgens*. *Lobatula lobatula* and *C. refulgens* live attached to elevated substratum (Murray, 1971; Van der Zwaan, 1982; Lutze and Thiel, 1989; Kaiho, 1994, 1999; Kouwenhoven, 2000; Schönfeld, 2002; Murray, 2006) and together with *C. pachyderma*, they are interpreted to indicate high bottom currents and well-oxygenated environments. The *C. laevigata* group, *B. marginata* and *Melonis* spp. are interpreted

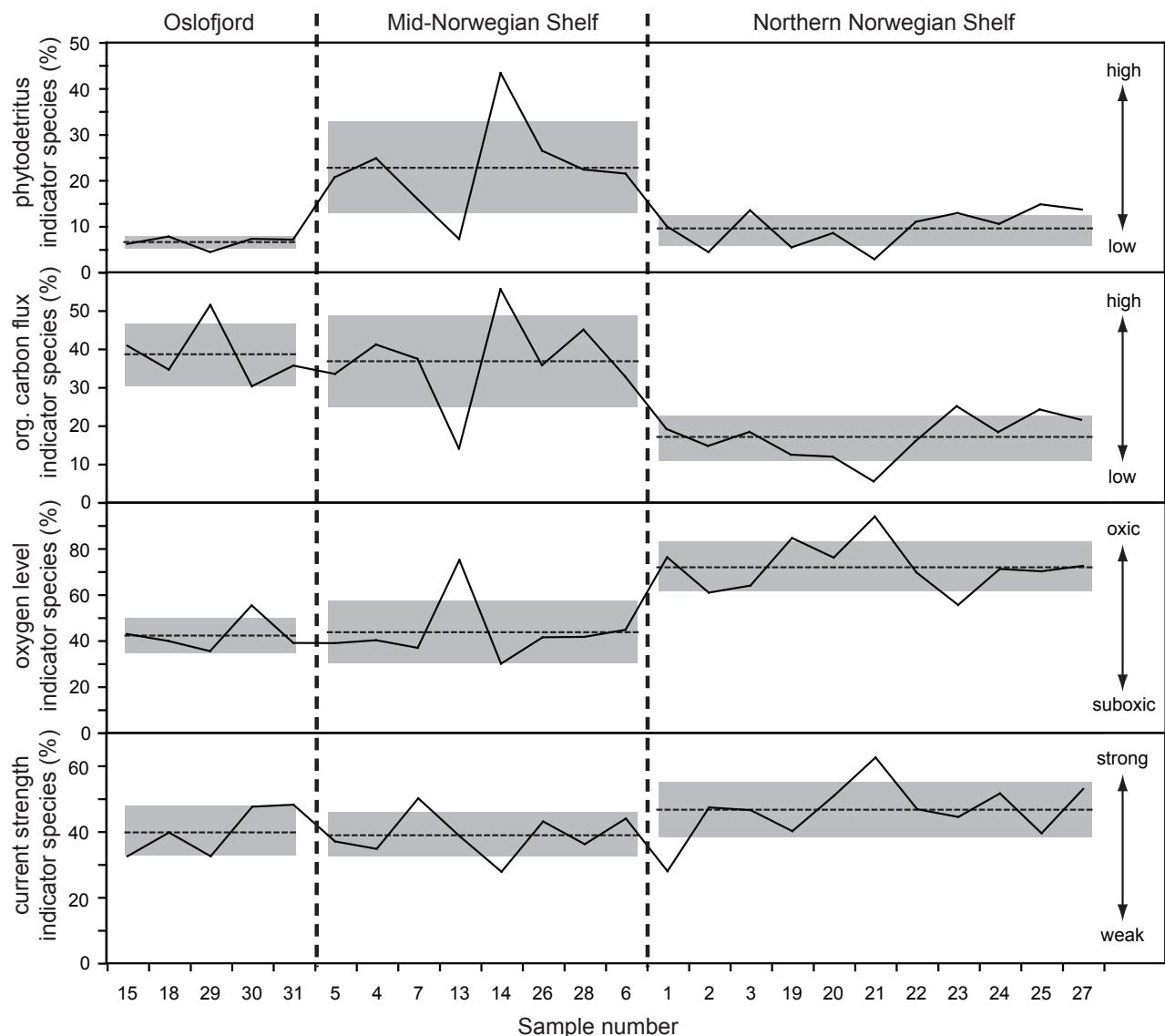


Figure 3.6 Indicator species for phytodetritus flux, organic carbon flux, oxygen level and current strength (according to Murray, 2006 and Margreth et al., 2009) for Clusters IV to VI: Oslofjord, Mid- and Northern Norwegian Shelf. Dotted line indicates the mean values with standard deviation (grey area).

to indicate an elevated organic carbon flux, with a high proportion of refractory components (e.g., Corliss, 1985; Mullins et al. 1985, Gooday, 1986; Mackensen and Hald, 1988; Mackensen et al., 1990). Therefore, this cluster should represent environments with strong bottom currents, high oxygen levels and an elevated organic carbon flux with high proportions of probably refractory components. This is also supported by the absence of phytodetritus indicator species (Fig. 3.6).

Cluster V combines the samples from the Mid Norwegian CWC-reefs. The benthic fauna is dominated by *A. angulosa*, *C. pachyderma*, *E. vitrea*, and the *G. subglobosa* group, *C. refulgens*, *M. barleeanum*. Passive suspension feeders like *C. refulgens* living epifaunal-attached on elevated substrate together with *A. angulosa* are interpreted to indicate high bottom currents (Murray 1971; Van der Zwaan 1982; Mackensen et al. 1985; Lutze and Thiel 1989; Kaiho 1994; Schönfeld, 1997, Kaiho, 1999; Kouwenhoven, 2000; Schönfeld, 2002; Murray, 2006; Margreth et al., 2009). *Epistominella vitrea* is an opportunistic species capable of responding to elevated food availability (Jorissem et al., 1992) whereas the *G. subglobosa* group is known to feed on phytodetritus (Suhr et al., 2003). The infaunal species *M. barleeanum* lives in muddy to silty sediment and is related to high levels of organic matter flux (e.g., Corliss, 1985; Gooday, 1986). Therefore, this cluster is interpreted to represent an environment characterized by strong bottom currents, elevated oxygen levels and a high flux of organic matter including phytodetritus (Fig. 3.6).

Cluster VI combines the northern Norwegian reef samples. The fauna is dominated by *C. pachyderma*, *L. lobatula*, the *G. subglobosa* group, *D. coronata*, *C. refulgens*, and *A. angulosa*. *Lobatula lobatula*, *D. coronata*, *C. refulgens* are passive suspension feeders living epifaunal-attached on elevated substratum and together with *A. angulosa* indicate strong bottom currents. The abundance of *G. subglobosa* may indicate an elevated phytodetritus flux but generally lower than on the Mid-Norwegian shelf. Therefore, this cluster is interpreted to represent strong bottom currents, high oxygen levels and a moderate organic matter flux mainly composed of seasonal pulses of phytodetritus input (Fig. 3.6).

In summary, the benthic foraminiferal assemblages indicate changes in specific environmental parameters such as bathymetry, substrate availability, current strength, oxygenation level, and quantity/quality of the organic carbon flux. The deep-sea environments below 1800 m water depth, are oligotrophic with only

seasonally controlled organic matter fluxes, whereas the “off-reef” environments on the upper continental slope above 1800 m show evidence of an elevated nutrient flux to the sea floor.

“On-reef”, the ecological demands of the recovered planktonic and benthic foraminiferal assemblages point to conditions of strong bottom currents, a high surface productivity and, therefore, an elevated flux of organic matter to the sediment. The ecological requirements of the benthic foraminifera from this cluster are identical with the ecological requirements of CWCs. They occur exclusively in environments characterised by strong currents (Frederiksen et al., 1992; Freiwald et al., 2002; Rüggeberg et al., 2007) and large amounts of organic matter in the form of phytoplankton detritus (Duineveld et al., 2004). The strong currents transport the food to the coral polyps and prevent them from getting buried by fine-grained sediments. The organic matter feeds the zooplankton, which is the main food source for CWCs (Mortensen, 2001; Freiwald et al., 2002).

In the Oslofjord, observations made from the submersible JAGO showed extended reef complexes, which were however, mainly composed of dead corals and coral rubble. Octocorals are generally highly abundant on CWC-reefs (Freiwald et al., 2004) but completely absent in the Skagerrak. Only relatively small patches of living *L. pertusa* were observed indicating that conditions for the growth of cold-water corals were not optimal. We suggest, that the reason for this large amount of dead corals may be the nature of the organic matter. According to Carney (1989) two types of organic matter reach the sea floor, the labile and the refractory organic matter. The refractory component mainly consists of terrestrial organic matter and the labile component mainly consist of phytodetritus. This latter is rapidly remineralised at the sediment-water interface using aerobic pathways and can be used immediately by living organisms. Although, benthic foraminiferal assemblages in this region indicates an elevated flux of organic matter to the sea floor, the very low abundance of phytodetritus-feeding species (Fig. 3.6) suggest that the labile component of the organic matter is very rare.

3.5.2.4 “On- and off-reef” diversity

Warm-water coral reefs commonly show high faunal diversity compared to “off-reef” environments (e.g., Connell, 1978, Carpenter et al., 1998). This is also in agreement with diversity studies comparing the macrofauna of cold-water coral reefs and the surrounding “off-reef” environments. Henry and

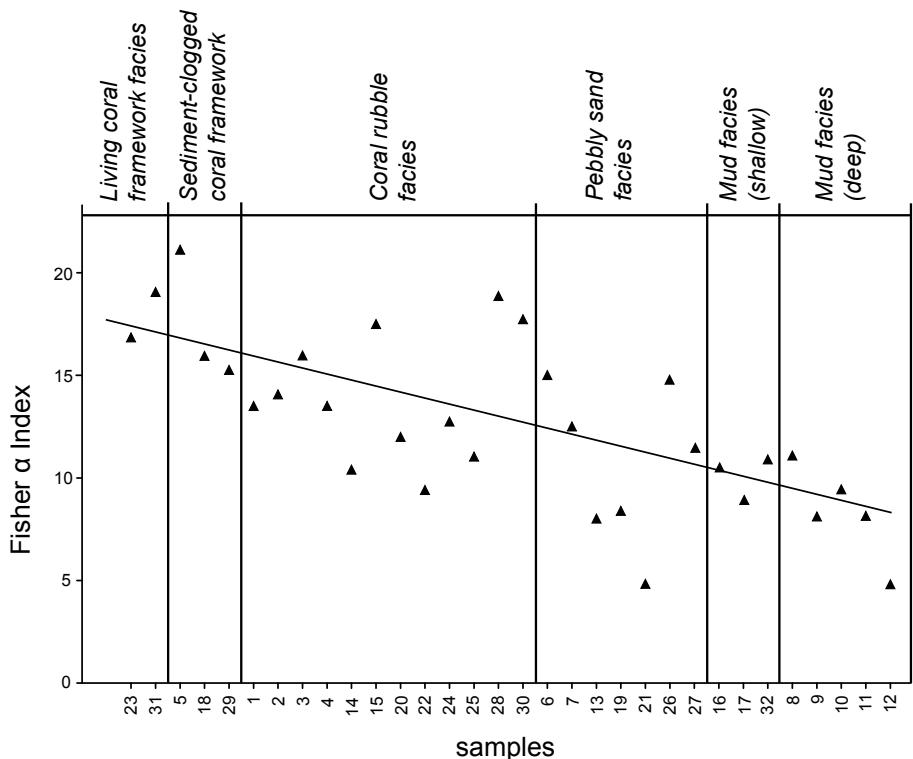


Figure 3.7 Fisher's α Diversity Index of the benthic foraminifera for the different sedimentary facies.

Roberts (2007) concluded that the high vertical habitat heterogeneity on a cold-water coral reef provides a diverse range of niches over small spatial scales that support a characteristic “reef-fauna”. Langer and Lipps (2003) showed that heterogeneity of microhabitats within warm-water coral reefs are an important factor for high foraminiferal species diversity. This seems to be valid also for cold-water coral reefs where the highest foraminiferal diversity is observed “on-reef” represented by Clusters IV, V, and VI. A possible reason for the observed “on-reef” high diversity is substrate availability for epifaunal and epifaunal-attached species, which preferentially live on the skeletal framework and hard substrates whereas, infaunal species prefer baffled fine-grained sediment (Fig. 3.7).

3.5.2.5 Norwegian cold-water coral reefs versus reefs associated to carbonate mounds along the Irish margin CWC-reefs on carbonate mounds on the Irish margin are wide spread and typically different facies are well distinct (Margreth et al., 2009). Benthic foraminifera are highly abundant and the assemblages reflect the different facies of the CWC-reefs. Uvigerinids are characteristic for “off-reef” sediments. The dropstone facies is dominated by *G. subglobosa* and *Cibicidoides* spp., whereas the *Dead Coral Facies* is characterized by *Planulina ariminensis*, *Hanzawaia boueana*, *E. vitrea*, and *A. angulosa*. For the *Living Coral Facies*

the epifaunal *D. coronata*, *L. lobatula* but also highly abundant infaunal species like *Gavelinopsis* spp., *G. subglobosa*, *Bolivina* spp. are typical. The *Sandwave Facies* is dominated by epifaunal-attached species like *D. coronata*, *C. refulgens* and the epifaunal species *Sigmoilopsis schlumbergeri*, *C. pachyderma* (Margreth et al., 2009). A distinct relation of the benthic foraminiferal assemblages to a specific sedimentary facies, such as observed in the CWC-reefs associated to carbonate mounds along the Irish margin, is not well expressed in the Norwegian coral reefs.

However, the assemblage on the Norwegian reefs show a strong latitudinal gradient and are not significantly different for the “on-reef” facies 1-4 (Fig. 3.5) indicating that benthic foraminiferal habitats on Norwegian coral reefs are weakly defined and grade one into the other. The gradual changes in assemblages observed on the Norwegian margin can be attributed to the changes in reef facies that are restricted within tens of metres whereas different facies are extensive on the Irish carbonate mounds (Freiwald et al., 2002, Hühnerbach et al., 2008, Margreth et al., 2009).

However, the species and/or groups of species between the two areas show a remarkable resemblance, indicating that the ecological conditions on the CWC-reefs along the Norwegian margin are similar to the reefs associated to the carbonate mounds along the

Irish margin. A high abundance of indicator species for elevated organic matter flux conditions and strong currents can be observed in both areas. On the Irish margin these indicator species are mainly associated with living coral colonies in the central parts of the carbonate mounds (Margreth et al., 2009) along the Norwegian margin they are concentrated "on reef".

3.5.2.6 Benthic foraminiferal species characteristic for cold-water coral reefs

The distribution of benthic foraminiferal species depends on a variety of ecological parameters (Murray, 2006). Species for which the threshold for certain ecological parameters is constrained can be used for indicating particular conditions in paleoenvironmental reconstructions.

Rüggeberg et al. (2007) studied the evolution through time of benthic foraminifera from a CWC-reef associated with a carbonate mound in the Porcupine Seabight along the Irish margin. The assemblage was composed of reef-related forms with a high abundance of epibenthic-attached species including *D. coronata* as a dominant species. *Discanomalina coronata* has also been described from sediment samples on Pleistocene and early Holocene CWC-reefs in the Mediterranean (Remia and Taviani, 2005; Margreth et al. in review.). A more recent and focused study on surface samples from different CWC-bearing carbonate mounds found specific coral-associated assemblages along the Irish margin (Margreth et al., 2009) and *Discanomalina coronata* was identified as an indicator species for cold-water coral reefs in this region.

Similarly to the Irish margin, the occurrence of *D. coronata* along the mid and northern Norwegian margin is restricted to the living CWC-habitats. In the Oslofjord (Skagerrak), where large areas of the CWC-reefs are dead, *D. coronata* is also almost absent (Fig. 8). The co-occurrence of *D. coronata* and CWC along the Irish margin and on the Norwegian reefs indicates that their ecological and hydrographic requirements are related. Therefore, *D. coronata* is here proposed as an indicator species for living and healthy CWC also on Norwegian cold-water coral reefs.

3.6 CONCLUSION

Cluster analysis, nMDS and diversity studies on dead (unstained) planktonic and benthic foraminiferal assemblages along the Norwegian margin have been used to characterize the CWC ecosystems along the Norwegian margin. They revealed that the distribution of planktonic foraminiferal assemblages is related to different current systems. *Neogloboquadrina incompta*, *G. uvula*, *T. quinqueloba* and *G. bulloides* occur on the Norwegian shelf and slope, influenced by the NAC, and suggest possible seasonal phytoplankton blooms. *Globigerinata glutinata*, *T. quinqueloba*, and *G. uvula* indicated upwelling on the inner continental shelf. High similarity of the fragile planktonic foraminiferal assemblage over a depth range of more than 2000 m indicates that carbonate dissolution in the area does not affect tests. The distribution of the benthic fauna is statistically separated into six different assemblages: I) deep-sea below 1800 m water depth; II) deep-sea between 1800 and 800 m water depth; III) "off-

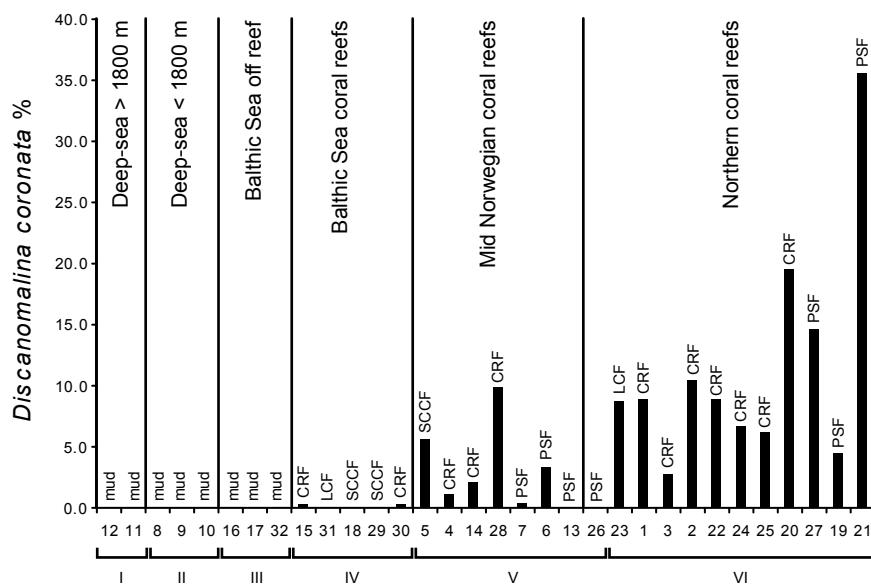


Figure 3.8 Abundance of *Discanomalina coronata* for Clusters I to VI and different sedimentary facies.

reef” Skagerrak; IV) “on-reef” Skagerrak; V) mid-Norwegian coral reefs, and VI) northern Norwegian coral reefs. The benthic fauna provides information on currents, oxygenation and organic matter content. In particular, benthic foraminifera associated to CWC-reefs indicate an environment characterized by high energy, well-oxygenated waters and high amounts of organic matter (derived from seasonal phytoplankton blooms) reaching the sea floor. The benthic foraminiferal fauna associated with coral reefs in the Skagerrak indicates that, although the flux of organic matter is high, the labile component usable for CWC growth is low and explains the extensive occurrence of dead corals in this area. High diversity of benthic foraminifera is observed “on-reef”. Diversity is highest in the central parts of the reefs, in the *living coral framework facies* and *sediment clogged coral framework facies*, with a dense coral framework. However, the benthic foraminiferal habitats are only weakly defined and grade one into the other preventing a sharp facies separation as observed along the Irish margin where the assemblages are strictly related to facies. The gradual changes in the assemblages observed on the Norwegian margin can be attributed to the changes in the reef facies, which are restricted within tens of metres, whereas on the Irish carbonate mounds different facies occur over long distances. *Discanomalina coronata* is only abundant on healthy CWC-reefs along the Mid Norwegian shelf and northern Norway. It is almost absent in the Oslofjord and completely absent “off-reef”. Co-occurrence of *D. coronata* and healthy CWC has been observed along the Irish margin. Based on these observations *D. coronata* is interpreted to require similar ecological conditions as CWCs and, therefore, proposed as an indicator species for healthy CWC-reefs also on the Norwegian margin.

4 - DEVELOPMENT OF COLD-WATER CORAL ECOSYSTEMS ON MUD VOLCANOES IN THE WEST ALBORAN BASIN: PALEOCEANOGRAPHIC IMPLICATIONS

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submitted to Marine Geology

The Dhaka and Maya mud volcanoes (MV), located in the Mud Diapir Province in the Western Alboran Basin along the Moroccan Coasts, were cored during the TTR-17, Leg 1 cruise. Cores were taken on the top of the volcanoes at a water depth of 370 m on the Dhaka MV (core TTR17-MS411G) and of 410 m on the Maya MV (core TTR17-MS419G). On both mud volcanoes the extruded mud breccia provides the nucleation point for the colonization and development of cold-water corals and associated ecosystems. Two phases of cold-water coral growth are observed: (1) between 2230 +/- 59 years BP and slightly older than 4175 +/- 62 years BP at Dhaka, and (2) between 7613 +/- 38 years BP and slightly older than 15583 +/- 185 years BP at Maya. On the top of the Maya MV only a small patch reef and/or isolated corals proliferated, whereas a more extended patch reef colonized the top of the Dhaka MV. At both sites the cold-water corals development was triggered by high nutrient availability related to upwelling and/or strong currents that mobilized nutrient horizontal fluxes. During the intervals of coral growth planktonic foraminiferal assemblages were dominated by *Neogloboquadrina incompta*. The decline of coral ecosystems on the mud volcanoes is accompanied at surface by a shift from the *N. incompta* dominated assemblage to a *Globorotalia inflata* dominated assemblage, possibly reflecting more modern paleoceanographic conditions and a deep pycno-nutricline. This shift seems to occur at the passage from wet to arid conditions at the end of the African Humid Period (Maya MV) and as a combined effect of an early human impact on a fragile environment, which was already stressed by desiccation at the time of the development of complex human society along the Mediterranean coasts (Dhaka MV). No relation with seepage is observed at both volcanoes.

4.1 INTRODUCTION

Mud volcanoes and mud diaps are common features in the Alboran Sea in the Western Mediterranean (e.g., Sautkin et al., 2003) and along the Atlantic margin of the associated Gulf of Cadiz (e.g., Fernandez-Puga et al., 2007; Pinheiro et al., 2003; Somoza et al., 2003). Mud volcanism and associated phenomena such as cold seepage, hydrocarbon venting and gas hydrates in the Gulf of Cadiz have been investigated since 1996 (Baraza et al., 1999; Baraza and Ercilla, 1996; Gardner, 2001; Ivanov et al., 2000, 2001; Kenyon et al., 2001; Masurenko et al., 2002; Pinheiro et al., 2003; Somoza et al., 2000). An exploratory cruise of R/V Belgica in 2002 on an accretionary setting in the Gulf of Cadiz (Gutscher et al., 2002) off Larache (Morocco), led to the discovery of a cluster of nine mud volcanoes: the El Arrache mud volcanoes field (Van Rensbergen et al., 2005). These structures are generally associated to cold-water corals carbonate mounds, which are up to 60 m high and located in water depths of 500-600 m on the Pen Duik Escarpment (Foubert et al., 2008; Van Rensbergen et al., 2005; Wienberg et al., 2009).

The existence of mud volcanoes in the Alboran Sea was first documented in 1999 during the UNESCO-IOC Training Through Research Program (TTR-9) Leg 2 along the Moroccan margin. A further survey carried out in 2002 by TTR-12 Leg 3 revealed the existence of mud volcanism along both Spanish (Northern Mud-Volcano Field) and Moroccan (Southern Mud-Volcano Field) margins of the West Alboran Basin. Additional mud volcanoes were discovered during the TTR-14 between the northern and southern mud-volcano fields (Comas and Ivanov, 2006; Comas et al., 2000, 2003a, 2003b; Sautkin et al., 2003; Talukder et al., 2003). In 2007, a cruise of the R/V Hesperides focusing on mud volcanoes in the Alboran Sea (Comas and Pinheiro, 2007) unveiled an outcropping mound province off Melilla, which displays striking affinities with the Cadiz cold-water coral carbonate mounds and with those discovered in the North Atlantic (e.g., Henriet et al., 1998). The Melilla field was cored for the first time in June 2008 during the TTR-17 cruise (Comas and the SAGAS08 - TTR17, Leg 1 - Scientific Party, unpublished). Other targets of the same cruise were

Table 4.1 Geographic position and water depths of the two investigated cores.

Gravity Core	Latitude	Longitude	Site	Length [m]	Depth [m]
TTR17-MS411G	35°25,43'N	04°31,89'W	Dhaka Mud Volcano	162	370
TTR17-MS419G	35°27,11'N	04°37,14'W	Maya Mud Volcano	167	410

also the mud volcanoes from the southern part of the Alboran Sea to test if cold-water coral carbonate mounds in the Alboran Sea were also associated to mud volcanoes as observed in the Gulf of Cadiz.

Aim of this research is the study of the sedimentary pelagic sequence deposited on the top of the Dhaka and Maya mud volcanoes in the Alboran Sea (Fig. 4.1) to reconstruct the paleoceanographic setting and to identify a possible causal link between these geological structures.

4.2 Regional setting

The Alboran Sea is a 400 km long and 200 km wide basin with a water depth not exceeding 2 km, located in the westernmost part of the Mediterranean Sea (e.g., Comas et al., 1999). Its complex seafloor morphology shows ridges, seamounts and troughs and it is divided in three sub-basins: the West Alboran, the East Alboran and the South Alboran Basins (Fig. 4.1). The West Alboran and the South Alboran Basins are separated by the Alboran Ridge, a prominent NE-SW linear relief, locally forming the small Alboran Island.

The formation of the Alboran Basin as part of the Gibraltar Arc started in the late Cretaceous as a consequence of crustal extension in a setting of overall convergence of the African and Eurasian plates that have had variable directions of relative motion since the late Cretaceous (e.g., Dewey et al., 1989). Seismic data show that present-day plate tectonics contribute to the actual deformation of the Alboran Basin (e.g., Fernandez-Ibañez et al., 2007, and references therein; Frizon de Lamotte et al., 2006). Early Miocene under-compacted shales and olistostromic sediments mobilized by fluid flows in a back-arc basin setting characterized by coeval extensional tectonics have been suggested as the source layer for the Mud Diapir Province in the Western Alboran Basin, which during the post-Messinian compressive tectonics developed pierced diapirs and subsequent mud volcanoes (Chalouan et al., 1997; Comas et al., 1992, 1999; Jurado and Comas, 1992; Pérez-Beluz et al., 1997; Sautkin et al., 2003; Talukder et al., 2003).

4.2.1 Dhaka and Maya Mud Volcanoes

The Dhaka and Maya mud volcanoes are located in the Mud Diapir Province in the Western Alboran Basin (35°25.43'N; 04°31.89'W and 35°27.11'N; 04°37.14'W respectively) along the Moroccan Coasts. The Dhaka mud volcano is a semi-circular structure with a diameter of about 1 km. The Maya mud volcano is elongated in an E-W direction and about 120 m long. Cores were taken on the top of the volcanoes at a water depth of 370 m on the Dhaka mud volcano (TTR17-MS411G) and of 410 m on the Maya mud volcano (TTR17-MS419G) (Fig. 4.1; Tab. 4.1).

According to the on-board description modified after shore-based investigations, the cored material at the Dhaka mud volcano includes from top to bottom an hemi-pelagic drape (from the top of the core to 30 cm), a cold-water coral fragments-rich unit (from 35 cm to 60 cm), a typical mud-breccia layer (from 65 cm to 145 cm), and possibly a lowermost level of pelagic sediments mixed with the mud breccia (from 145 cm to the bottom of the core).

The cored material at the Maya mud volcano includes, from top to bottom, a hemi-pelagic drape (from the top of the core to 94 cm), a cold-water coral fragments-rich unit (from 100 cm to 126 cm), and a typical mud-breccia layer (from 127 cm to the bottom of the core). Both volcanoes are presently not active as testified by the presence of a pelagic drape on their top.

4.3 MATERIAL AND METHODS

The sediments representing the hemi-pelagic drape have been sampled at five cm resolution (Dhaka) and two to ten cm resolution (Maya) for the determination of planktonic and benthic foraminifera assemblages. Sediment samples for micropaleontological analyses were processed following Spezzaferri and Coric (2001). Samples were dried at room temperature, weighted and washed through 250 µm, 125 µm and 63 µm mesh sieves. The obtained three size fractions of residue were dried at room temperature and weighted. When a residue contained more than 400 specimens, the volume was reduced by splitting. About 200 specimens per fraction (600 specimens per sample) were picked,

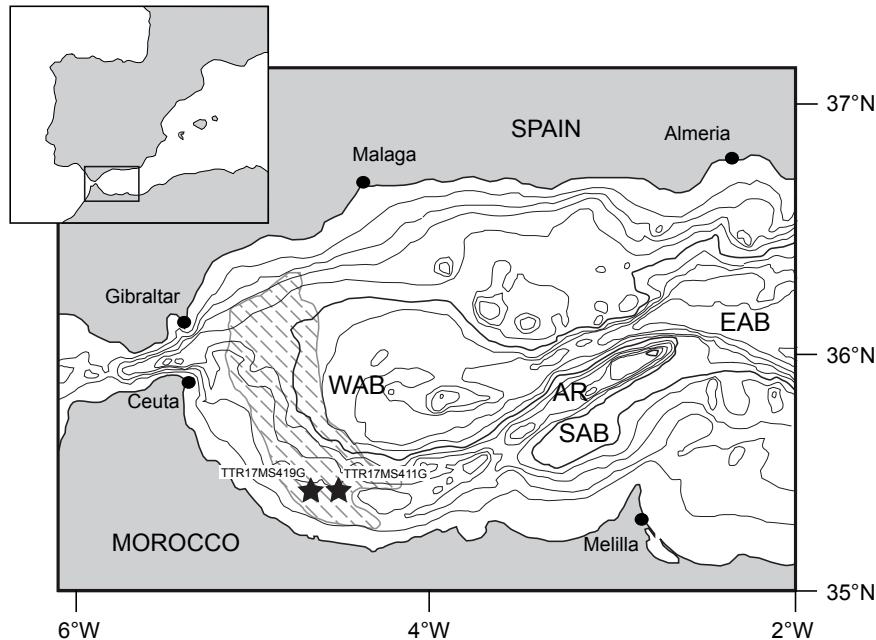


Figure 4.1 Location map of the studied cores. Shaded area: Mud Diapir Province. WAB: West Alboran Basin; EAB: East Alboran Basin; SAB: South Alboran Basin; AR: Alboran Ridge. Modified from Arnone et al. (1990).

collected in slides, sorted at species level and counted. In case of samples containing less than 100 specimens per fraction, all specimens were counted. The identified different benthic and planktonic foraminiferal species are listed in Appendix E and F.

A multivariate statistical treatment on faunal data was performed with the software PRIMER 6 (Clarke, 1993; Clarke and Gorley, 2006; Clarke and Warwick, 2001). Data have been double-square root transformed to limit the contribution of the most abundant and ubiquitous species (Field et al., 1982). Bray-Curtis (dis-) similarities were calculated following Clifford and Stephenson (1975). The same similarity matrix was used to obtain the non-metric MultiDimensional Scaling (nMDS plot) (Kruskal, 1964; Kruskal and Wish, 1978). The significance and potential of the nMDS plot has been discussed in detail in Spezzaferri and Coric (2001) and Margreth et al. (2009). Based on the clusters given in the nMDS plot, the Similarity Percentage Analysis (SIMPER) was obtained to highlight the contribution of each species to the total average (dis-) similarity between different groups and within one group (e.g., Basso and Spezzaferri, 2000; Clarke and Warwick, 2001; Kruskal, 1964; Kruskal and Wish, 1978). Quantitative data of benthic and planktonic foraminifera used for the multivariate statistic are given in Appendix E and F.

The chronology of the two cores retrieved on Dhaka and Maya mud volcanoes (TTR17-MS411G and

TTR17-MS419G, respectively) is based on 8 AMS ^{14}C age determination (Tab. 4.2) obtained at the Leibniz Laboratory for Age Determinations and Isotope Research at the University of Kiel. For each sample, at least 5 mg of *Globorotalia inflata* were handpicked to provide 0.2 to 1.2 mg of carbonate. The organic material was removed in an ultrasonic bath with 15% H_2O_2 . The hydrolysis of the residual pure calcite was obtained at 90°C with 100% orthophosphoric acid. The carbon dioxide was converted to graphite with hydrogen on an iron catalyst. The iron-graphite mixture was measured on the AMS mass spectrometer. A reservoir correction of 400 years (Siani et al., 2000) was subtracted to the obtained radiocarbon conventional ages, which were calibrated using the program Calib 5.1 (Stuiver and Reimer, 1993).

Stable oxygen and carbon isotope analyses ($\delta^{18}\text{O}$ and $\delta^{13}\text{C}$) on about 5-10 specimens of *Discanomalina coronata* were performed on 3 samples from the coral-rich level from the Dhaka mud volcano (Tab. 4.3). These isotope ratios were measured on a Finnigan MAT 252 mass spectrometer with a Kiel II preparation device at the IFM-GEOMAR, Leibniz-Institute of Marine Sciences (Kiel University). The reproducibility was $\pm 0.035\text{\textperthousand}$ for $\delta^{18}\text{O}$ and $\pm 0.029\text{\textperthousand}$ for $\delta^{13}\text{C}$. The isotopes ratios are presented relative to Vienna Pee Dee Belemnite standard (VPDB) based on calibrations of National Bureau of Standards (NBS).

Table 4.2. Result of AMS ^{14}C dating on planktonic foraminifera *G. inflata*. Calibration using the software Calib 5.1 (Stuiver and Reimer, 1993). A reservoir effect of 400 years was subtracted (Siani et al., 2000).

Gravity Core	Depth [cm]	Species	Lab Code	Conventional [yr BP]	Calibrated Age [yr BP]
TTR17-MS411G	0	<i>Gr. inflata</i>	KIA 39074	1145 ±30	689 ±27
TTR17-MS411G	35	<i>Gr. inflata</i>	KIA 39075	2550 ±40	2230 ±59
TTR17-MS411G	55	<i>Gr. inflata</i>	KIA 39076	4120 ±35	4175 ±62
TTR17-MS411G	150	<i>Gr. inflata</i>	KIA 39078	4240 ±40	4345 ±60
TTR17-MS419G	0	<i>Gr. inflata</i>	KIA 39197	1475 ±25	1013 ±40
TTR17-MS419G	25	<i>Gr. inflata</i>	KIA 39198	2425 ±35	2059 ±53
TTR17-MS419G	98	<i>Gr. inflata</i>	KIA 39199	7145 ±40	7613 ±38
TTR17-MS419G	115	<i>Gr. inflata</i>	KIA 39200	13580 ±70	15583 ±185

4.4 RESULTS

4.4.1 Micropaleontology

A hemi-pelagic drape and a cold-water coral fragments-rich unit, even if at different depth in the cores, characterize both Dhaka and Maya mud volcanoes. The hemi-pelagic drape, from the top of the core to 30 cm in Dhaka MV and from the top of the core to 94 cm in Maya MV, yields very similar foraminiferal assemblages. Planktonic foraminifera include very abundant *Globorotalia inflata*, *Globigerinoides* spp., *Turborotalita quinqueloba*, *Globigerinita glutinata* and benthic foraminifera such as *Globocassidulina subglobosa*, *Uvigerina mediterranea*, *Cassidulina carinata*, *Bolivina difformis* (Appendix E). In the cold-water coral fragments-rich unit (from 35 cm to 60 cm in Dhaka MV and from 100 cm to 126 cm in Maya MV) planktonic fauna is represented by dominant *Neogloboquadrina incompta*, *Turborotalita quinqueloba* and associated less abundant *Globigerinita glutinata* and *Globigerina bulloides*. In this unit, epifaunal-attached benthic foraminifera are more abundant than in the above hemi-pelagic sediments and are characterized by *Discanomalina coronata*, a species that is interpreted as bioindicator of living cold water coral ecosystems along the Irish coasts in the North Atlantic (Margreth et al., 2009), *Cibicidoides* sp., *Cibicidoides pachyderma*, *Angulogerina angulosa*, and *Lobatula lobatula* (Appendix E). The residues used for foraminiferal investigations contain also fragments of cold-water corals such as *Madrepora oculata*, *Lophelia pertusa* and solitary species such as *Desmophyllum* sp. and/or *Dendrophyllia* sp. (Fig. 4.2).

4.4.2 Quantitative analyses and statistical treatment of data

Quantitative analyses of planktonic and benthic foraminifera were performed on samples from the hemi-pelagic drape of both mud volcanoes in order to evaluate species abundance and diversities and to plot abundance curves of selected species (Fig. 4.3).

Planktonic foraminifera (Dhaka MV) – A clear positive peak of abundance (almost 50%) is observed for *N. incompta* at about 50 cm (Fig. 4.3A) within the interval containing coral fragments. Abundance of *G. inflata* increases from 45 cm up to over 30% in the upper part of the core with a negative peak (10%) at 35 cm. In the hemi-pelagic sediments above the coral-rich interval the abundances of *G. incompta* and *Globigerinoides* spp. remain below 10%.

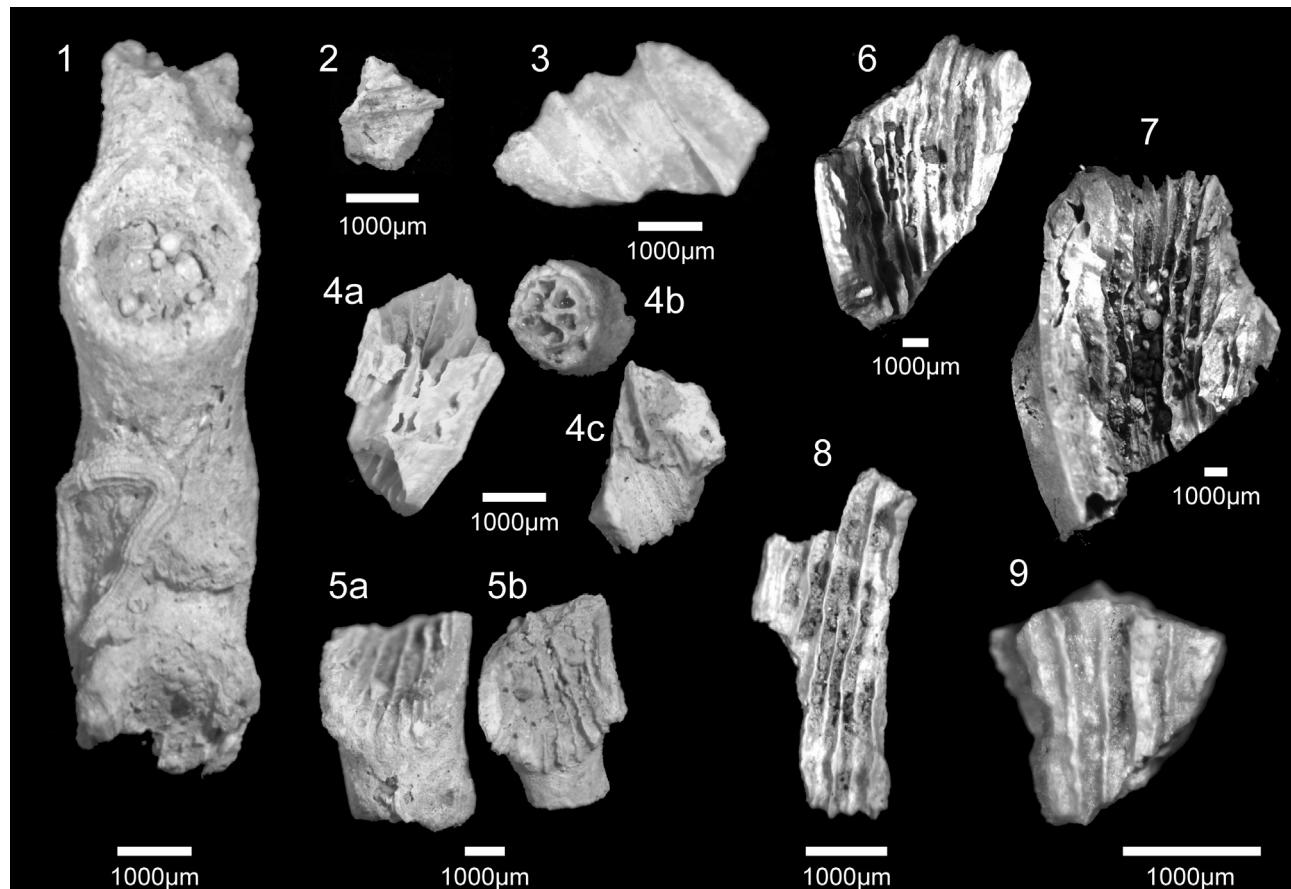
Benthic foraminifera (Dhaka MV) – In the interval containing also coral fragments *D. coronata* is more abundant (around 15%; Fig. 4.3B). From the base of this interval to the top of the core, the abundance of *G. subglobosa* increases from a minimum of around 10 to 25%, respectively. Uvigerinids (*U. mediterranea* and *U. peregrina*) also increase in abundance but never exceed 10%.

Planktonic foraminifera (Maya MV) - The lower abundances of *G. inflata* and *Globigerinoides* spp. are observed in the level containing also coral fragments (Fig. 4.3C), whereas *N. incompta* is very abundant (maximum value 40%). At about 85 cm, the curves of the three species reverse their trend: *G. inflata* becomes abundant (almost 30%) while *N. incompta* and *Globigerinoides* spp. never exceed 10%, showing a strict resemblance to the trend of the same species at Dhaka mud volcano.

Benthic foraminifera (Maya MV) – *D. coronata* is low abundant (1%) only in the coral-rich level in this core (Fig. 4.3D). *G. subglobosa* increases in abundance from the base of the coral-rich horizon (7%) to the top of the core (22%). Uvigerinids are not abundant, with values fluctuating from 3 to 8%. However, *U. peregrina* highest abundance of around 15% is observed in the coral-rich level. Very often patterns of community structures are not clear and/or readily apparent (Clarke and Warwick, 1994). Therefore, to highlight them, we have statistically treated our complete benthic and planktonic foraminifera data to better identify

Table 4.3 Benthic $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ measured on *Discanomalina coronata* from three samples of core TTR17-MS411G (Dhaka MV).

Gravity Core	Depth [cm]	Species	$\delta^{13}\text{C}$ [%‰ VPDB]	$\delta^{18}\text{O}$ [%‰ VPDB]
TTR17-MS411G	35	<i>D. coronata</i>	0.013	2.933
TTR17-MS411G	55	<i>D. coronata</i>	-0.218	3.542
TTR17-MS411G	70	<i>D. coronata</i>	-0.661	3.118

Figure 4.2 Scleractinian cold-water coral fragments, most certainly *Madrepora oculata* (1), *Lophelia pertusa* and solitary species like *Desmophyllum* sp. or *Dendrophyllia* sp. from the two investigated cores: (1) D25; TTR17-MS411G; 25 cm, (2) D45; TTR17-MS411G; 45 cm, (3) D40; TTR17-MS411G; 40 cm, (4a-c) D30; TTR17-MS411G; 30 cm, (5a-b) D25; TTR17-MS411G; 25 cm, (6) M115; TTR17-MS419G; 115 cm, (7) M107; TTR17-MS419G; 107 cm, (8) M100; TTR17-MS419G; 100 cm, (9) M102; TTR17-MS419G; 102 cm

and characterize changes in the assemblage structures and possibly relate them to changing environmental conditions. The multivariate statistical analyses, including agglomerative clustering based on the Bray-Curtis Similarity, is shown in Fig. 4.4. Two clusters separate for planktonic foraminifera and three clusters separate for benthic as shown in the nMDS plots in Fig. 4.3. Species and groups accounting for the average similarity and dissimilarity in all clusters are listed in order of decreasing contribution in Tab. 4.4, 4.5, 4.6 and 4.7.

4.4.3 AMS ^{14}C dating and sedimentation rate

The results of the AMS ^{14}C dating are reported in Tab. 4.2. The sediments from Core TTR17-MS411G (Dhaka MV) yield ages ranging from 689 ± 27 years at the top to 4346 ± 60 years at the bottom. The coral-rich interval is comprised within about 2230 ± 59 years BP and slightly older than 4175 ± 62 years BP. The sediments from Core TTR17-MS419 G (Maya MV) yield ages ranging from 1013 ± 40 years at the top to 15583 ± 185 years at the bottom. The coral-rich interval is comprised within 7613 ± 38 years BP and the breccia layers starting 10 cm below the sample dated as old as 15583 ± 185 years BP. The AMS ^{14}C

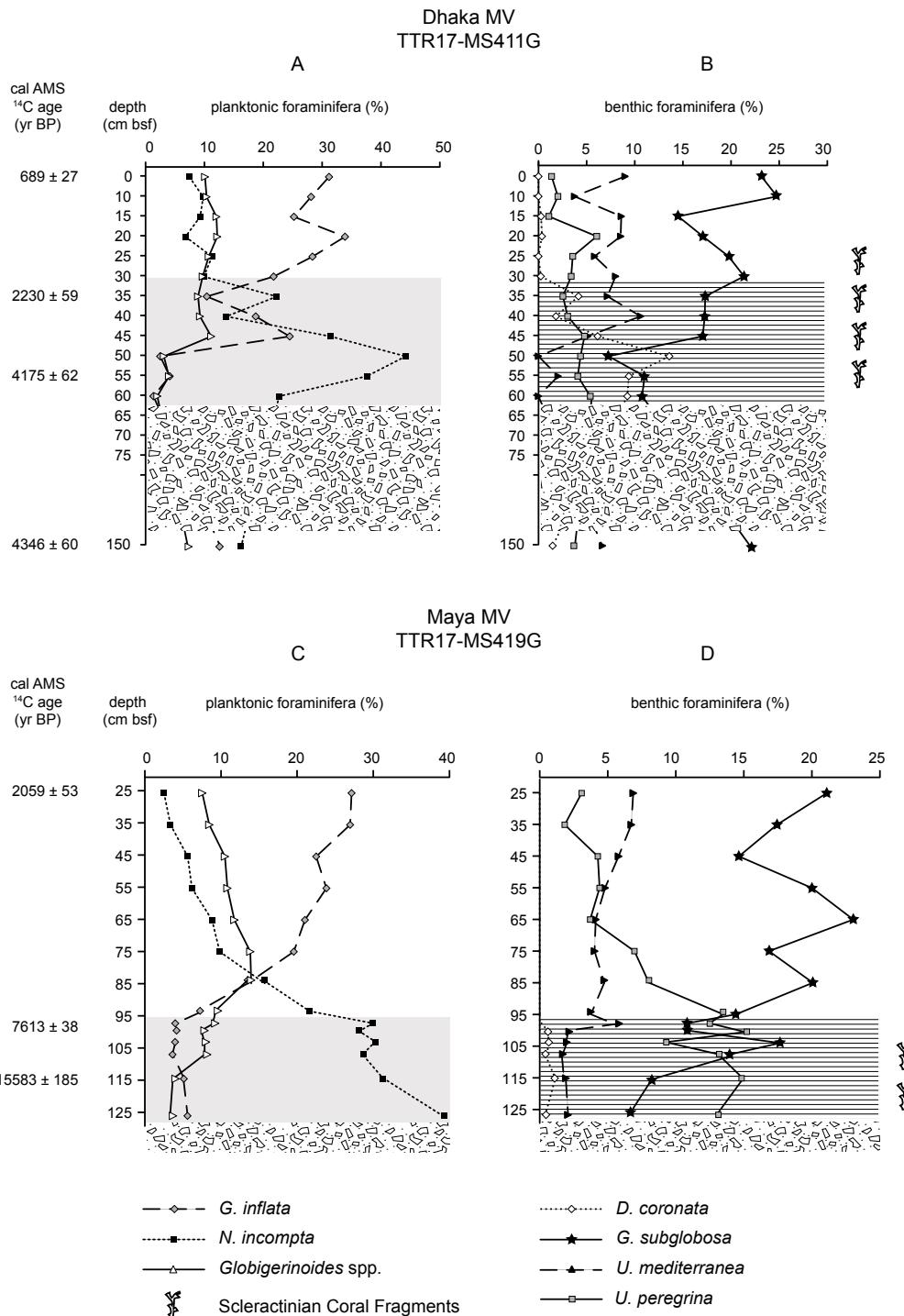


Figure 4.3 Relative abundance of the most significant planktonic (A/C) and benthic (B/D) foraminiferal species for the two investigated cores. Shaded areas: *N. incompta* period in the planktonic foraminiferal assemblage; striped areas: *D. coronata* period in the benthic foraminiferal assemblage; patterned areas: mud breccia level.

dating allowed plotting the sedimentation rate curves for the hemi-pelagic drape at the two sites (Fig. 4.5). The higher values of sedimentation rate are observed between 0 and 35 cm at Dhaka MV (22.7 cm kyr^{-1}) and from 0 to 25 cm at Maya MV (23.9 cm kyr^{-1}). The lower values correspond to the cold-water coral-rich layers and are 10.3 cm kyr^{-1} and 2.1 cm kyr^{-1} , respectively.

4.5 DISCUSSION

It has been demonstrated that the basic requirements of cold-water corals are hard substratum for their settlement and colonization, nutrient supply and low sedimentation rates (e.g., Alvarez-Pérez et al., 2005; De Mol et al., 2002; Frederiksen et al., 1992;

Table 4.4 List of planktonic foraminiferal species and statistical parameters associated with the similarity in cluster I to II. Average similarity within the group of station, average abundance, average similarity, contribution (%), and cumulative contribution (%) are given for each species with respect to the total similarity for each cluster.

Cluster I				
Species	Average similarity: 84.72			
	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Gr. inflata</i>	4.72	12.61	14.89	14.89
<i>Tr. quinqueloba</i>	4.61	12.58	14.85	29.74
<i>Ga. glutinata</i>	4.05	10.58	12.49	42.23
<i>N. incompta</i>	3.17	7.77	9.17	51.4
<i>Gs. ruber</i>	2.78	7.57	8.93	60.33
<i>G. bulloides</i>	2.74	6.97	8.23	68.56
<i>Gs. elongatus</i>	1.23	3.04	3.59	72.15
<i>Ga. uvula</i>	1.27	2.99	3.53	75.68
<i>Tr. humilis</i>	1.09	2.68	3.16	78.84

Cluster II				
Species	Average similarity: 84.15			
	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Tr. quinqueloba</i>	5.62	17.65	20.97	20.97
<i>N. incompta</i>	5.43	16.61	19.73	40.71
<i>Ga. glutinata</i>	3.92	11.95	14.2	54.9
<i>G. bulloides</i>	2.19	6.51	7.73	62.64
<i>Gs. ruber</i>	2.22	6.2	7.36	70
<i>Gr. inflata</i>	2.12	6.01	7.15	77.14

Freiwald, 2002; Roberts et al., 2006; Rogers, 1999). Also important are long-term stable cool conditions of temperature and current-strength (Freiwald et al., 2004). Recently Dullo et al. (2008) have demonstrated that cold-water corals in the Atlantic can adapt to a wide range of environmental conditions but they only thrive in a density gradient between 27.35 to 27.65 kg m⁻³. Freiwald et al. (2009) identified the limiting density in the Mediterranean between 29.08 and 29.13 kg m⁻³.

More than 20 mud volcanoes are presently known to be colonized by fossil scleractinian corals in the Gulf of Cadiz (Pinheiro et al., 2003; Somoza et al., 2003). Some of them are located along the Spanish margin in water depths between 530 and 1100 m and others are

located along the Moroccan margin in a much wider water depth range of 420 – 1300 m (Wienberg et al., 2009). Cold-water corals were previously documented in the Gibraltar Strait (Alvarez-Pérez et al., 2005). However, no data concerning the sedimentary sequences containing fossil cold-water coral remains on top of the Alboran mud volcanoes were available until the TTR-17-Leg 1 research cruise.

The first step in our study was to demonstrate that the cold-water coral intervals observed in the Dhaka and Maya mud volcanoes at the top of the extruded mud breccia were in situ and not coral debris reworked and/or displaced from older layers during mud volcanic activity. Since the sedimentological analysis of the cores did not provide any conclusive evidence,

Table 4.5 List of planktonic species and statistical parameter associated with the dissimilarity between each pair of clusters I to II. The average dissimilarity is shown. For each species the average abundance, average dissimilarity, contribution (%), and cumulative contribution (%) to the total similarity are also given.

Species	Average dissimilarity = 23.66				
	Cluster I	Cluster II	Av.Diss	Contrib%	Cum.%
	Av.Abund	Av.Abund			
<i>Gr. inflata</i>	4.72	2.12	4.02	16.97	16.97
<i>N. incompta</i>	3.17	5.43	3.55	15	31.98
<i>Tr. quinqueloba</i>	4.61	5.62	1.6	6.77	38.75
<i>Ga. glutinata</i>	4.05	3.92	1.11	4.7	43.45
<i>G. bulloides</i>	2.74	2.19	1.06	4.47	47.92
<i>Gs. ruber</i>	2.78	2.22	1	4.25	52.17
<i>Ta. rubescens</i>	1.24	0.91	0.93	3.93	56.1
<i>Gs. immaturus</i>	0.85	0.3	0.88	3.72	59.82
<i>Gs. sacculifer</i>	0.71	0.19	0.85	3.59	63.4
<i>N. pachyderma</i>	0.73	0.6	0.79	3.32	66.72
<i>Gs. elongatus</i>	1.23	0.9	0.7	2.94	69.66
<i>Gr. scitula</i>	0.61	0.49	0.66	2.8	72.46
<i>N. dutertrei</i>	0.29	0.5	0.66	2.77	75.23
<i>Tr. humilis</i>	1.09	0.8	0.64	2.71	77.94

Table 4.6 List of benthic foraminiferal species and statistical parameters associated with the similarity in cluster 1 to 3. Average similarity within the group of station, average abundance, average similarity, contribution (%), and cumulative contribution (%) are given for each species with respect to the total similarity for each cluster.

Cluster 1				
Average similarity: 63.31				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Globocassidulina subglobosa</i> group	19.6	17.73	28	28
<i>Uvigerina mediterranea</i>	5.78	4.31	6.81	34.81
<i>Bolivina difformis</i>	4.77	3.79	5.98	40.79
<i>Bulimina marginata</i>	4.8	3.71	5.86	46.65
<i>Uvigerina peregrina</i>	3.88	2.73	4.32	50.97
<i>Cassidulina carinata</i>	3.38	2.39	3.78	54.75
<i>Gavelinopsis praegeri</i>	3.39	2.27	3.58	58.33
<i>Cibicidoides</i> sp.	2.92	1.95	3.08	61.41
<i>Melonis barleeanum</i>	2.71	1.94	3.07	64.48
<i>Hyalinea balthica</i>	2.64	1.73	2.74	67.22
<i>Amphicoryna scalaris</i>	2.41	1.52	2.4	69.63
<i>Bigenerina nodosaria</i>	1.86	1.3	2.05	71.67
<i>Hoeglundina elegans</i>	1.64	1.08	1.7	73.37
<i>Cibicidoides pseudoungerianus</i>	1.63	0.9	1.42	74.79
<i>Lobatula lobatula</i>	1.38	0.87	1.38	76.17
<i>Sigmoilopsis schlumbergeri</i>	1.27	0.76	1.2	77.37
<i>Sphaeroidina bulloides</i>	1.38	0.73	1.16	78.53
<i>Cassidulina laevigata</i>	1.35	0.65	1.03	79.56
<i>Quinqueloculina viennensis</i>	1.25	0.65	1.03	80.59

Cluster 2				
Average similarity: 71.06				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Uvigerina peregrina</i>	13.09	12.01	16.9	16.9
<i>Globocassidulina subglobosa</i> group	11.86	9.51	13.38	30.28
<i>Bulimina marginata</i>	7.97	6.87	9.66	39.94
<i>Hyalinea balthica</i>	7.14	5.71	8.03	47.98
<i>Melonis barleeanum</i>	3.99	3.21	4.52	52.5
<i>Amphicoryna scalaris</i>	3.03	2.31	3.25	55.75
<i>Bulimina striata</i>	3.03	2.28	3.21	58.97
<i>Cassidulina carinata</i>	2.96	2.25	3.16	62.13
<i>Uvigerina mediterranea</i>	2.78	2.01	2.82	64.95
<i>Sigmoilopsis schlumbergeri</i>	2.16	1.82	2.56	67.51
<i>Cibicidoides pseudoungerianus</i>	2.02	1.72	2.42	69.94
<i>Sphaeroidina bulloides</i>	1.94	1.41	1.98	71.92
<i>Bolivina difformis</i>	1.87	1.35	1.9	73.82
<i>Bulimina aculeata</i>	1.58	1.09	1.53	75.35
<i>Hoeglundina elegans</i>	1.4	1.04	1.46	76.81
<i>Spiroplectinella wrighti</i>	1.31	1	1.41	78.22
<i>Bolivina spathulata</i>	1.39	0.97	1.36	79.58

Cluster 3				
Average similarity: 63.23				
Species	Av.Abund	Av.Sim	Contrib%	Cum.%
<i>Discanomalia coronata</i>	10.74	9.28	14.68	14.68
<i>Globocassidulina subglobosa</i> group	9.69	8.44	13.36	28.04
<i>Bolivina difformis</i>	5.75	4.57	7.23	35.27
<i>Uvigerina peregrina</i>	4.61	4.18	6.61	41.88
<i>Cibicidoides</i> sp.	6.85	3.44	5.44	47.32
<i>Cibicidoides pachyderma</i>	3.68	2.93	4.63	51.96
<i>Hyalinea balthica</i>	2.91	2.59	4.1	56.05
<i>Melonis barleeanum</i>	2.76	2.39	3.77	59.83
<i>Gavelinopsis praegeri</i>	2.57	2.27	3.59	63.42
<i>Angulogerina angulosa</i>	2.71	2	3.16	66.58
<i>Bolivina subaenariensis</i>	2.37	1.86	2.94	69.52
<i>Bulimina marginata</i>	2.25	1.63	2.57	72.09
<i>Cassidulina carinata</i>	2.43	1.57	2.48	74.57
<i>Cassidulina laevigata</i>	2.92	1.57	2.48	77.05
<i>Lobatula lobatula</i>	2.21	1.24	1.96	79.02
<i>Cibicidoides pseudoungerianus</i>	1.65	1.16	1.83	80.85

planktonic and benthic foraminiferal analyses were performed and the results compared with the study of Margreth et al. (2009) on modern foraminiferal assemblages from cold-water coral mounds in the

Porcupine Basin and Rockall Bank off-shore Ireland. These authors showed that benthic foraminiferal assemblages provide an independent tool to identify and describe the different facies related to cold-

Table 4.7 List of benthic species and statistical parameter associated with the dissimilarity between each pair of clusters 1 to 3. For each pair of clusters the average dissimilarity is shown. For each species the average abundance, average dissimilarity, contribution (%), and cumulative contribution (%) to the total similarity are also given.

Species	Average dissimilarity = 45.19				
	Cluster 1		Cluster 2		
	Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Uvigerina peregrina</i>	3.88	13.09	4.61	10.19	10.19
<i>Globocassidulina subglobosa</i> group	19.6	11.86	3.95	8.75	18.94
<i>Hyalinea balthica</i>	2.64	7.14	2.31	5.12	24.05
<i>Uvigerina mediterranea</i>	5.78	2.78	1.73	3.84	27.89
<i>Bulimina marginata</i>	4.8	7.97	1.71	3.78	31.68
<i>Bolivina difformis</i>	4.77	1.87	1.48	3.27	34.95
<i>Gavelinopsis praegeri</i>	3.39	1.08	1.22	2.69	37.64
<i>Bulimina striata</i>	0.76	3.03	1.13	2.51	40.15
<i>Cibicidoides</i> sp.	2.92	1.12	0.96	2.12	42.27
<i>Melonis barleeanum</i>	2.71	3.99	0.91	2.02	44.29
<i>Cassidulina carinata</i>	3.38	2.96	0.83	1.83	46.11
<i>Amphicoryna scalaris</i>	2.41	3.03	0.79	1.76	47.87
<i>Cassidulina laevigata</i>	1.35	1.42	0.64	1.42	49.29
<i>Sphaeroidina bulloides</i>	1.38	1.94	0.63	1.4	50.69
<i>Discanomalina coronata</i>	1.18	0.38	0.62	1.38	52.07
<i>Cibicidoides pseudoungerianus</i>	1.63	2.02	0.59	1.3	53.38
<i>Bulimina aculeata</i>	0.57	1.58	0.56	1.23	54.61
<i>Sigmaiolopsis schlumbergeri</i>	1.27	2.16	0.55	1.22	55.83
<i>Bigenerina nodosaria</i>	1.86	1.19	0.55	1.21	57.03
<i>Gyroidina altiformis</i>	0.81	1.08	0.52	1.14	58.18
<i>Quinqueloculina viennensis</i>	1.25	0.44	0.51	1.12	59.3
<i>Lobatula lobatula</i>	1.38	1.35	0.5	1.11	60.4
<i>Cibicidoides pachyderma</i>	1.06	0.16	0.49	1.09	61.49
<i>Hoeglundina elegans</i>	1.64	1.4	0.47	1.03	62.53
<i>Gyroldina neosoldanii</i>	1.24	0.63	0.46	1.02	63.55
<i>Bolivina spathulata</i>	1.18	1.39	0.46	1.02	64.57
<i>Spiroplectinella wrighti</i>	1.09	1.31	0.44	0.97	65.54
<i>Melonis pomphiloides</i>	0.45	1.21	0.42	0.94	66.48
<i>Bolivina dilatata</i>	0.84	1.23	0.42	0.93	67.41
<i>Bolivina suspinescens</i>	0.41	1.05	0.4	0.89	68.3
<i>Cibicides refulgens</i>	0.71	0.44	0.4	0.89	69.19
<i>Bolivina striatula</i>	0.34	1.05	0.4	0.88	70.07
<i>Cibicidoides kullenbergi</i>	1.06	0.57	0.38	0.84	70.91
<i>Planulina ariminensis</i>	0.56	0.96	0.37	0.82	71.73
<i>Nuttalides umboniferus</i>	0.72	0	0.36	0.79	72.52
<i>Globobulimina affinis</i>	0.81	0.57	0.35	0.77	73.29
<i>Rectuvigerina elongatastriata</i>	0.72	0.05	0.35	0.77	74.06
<i>Angulogerina angulosa</i>	0.66	0.57	0.34	0.75	74.81
<i>Textularia agglutinans</i>	0.69	0.03	0.33	0.74	75.55
<i>Lenticulina inornata</i>	0.74	0.22	0.33	0.73	76.28
<i>Bolivina subaenariensis</i>	0.12	0.67	0.31	0.68	76.96
<i>Discorbina bertheloti</i>	0.42	0.74	0.3	0.67	77.63
<i>Hanzawaia boueana</i>	0.91	0.45	0.3	0.67	78.31
<i>Cycloforina laevigata</i>	0.31	0.6	0.3	0.67	78.98
<i>Rosalina globularis</i>	0.67	0.51	0.29	0.64	79.62

Species	Average dissimilarity = 50.06				
	Cluster 1		Cluster 3		
	Av.Abund	Av.Abund	Av.Diss	Contrib%	Cum.%
<i>Globocassidulina subglobosa</i> group	19.6	9.69	4.96	9.9	9.9
<i>Discanomalina coronata</i>	1.18	10.74	4.78	9.55	19.45
<i>Uvigerina mediterranea</i>	5.78	0.68	2.57	5.14	24.59
<i>Cibicidoides</i> sp.	2.92	6.85	2.27	4.54	29.12
<i>Cibicidoides pachyderma</i>	1.06	3.68	1.33	2.65	31.77
<i>Bulimina marginata</i>	4.8	2.25	1.31	2.63	34.4
<i>Bolivina subaenariensis</i>	0.12	2.37	1.12	2.24	36.64
<i>Cassidulina laevigata</i>	1.35	2.92	1.07	2.14	38.78
<i>Bolivina difformis</i>	4.77	5.75	1.03	2.06	40.84
<i>Angulogerina angulosa</i>	0.66	2.71	1.03	2.06	42.9
<i>Uvigerina peregrina</i>	3.88	4.61	0.92	1.83	44.73
<i>Cassidulina carinata</i>	3.38	2.43	0.89	1.79	46.52
<i>Bigenerina nodosaria</i>	1.86	0.14	0.87	1.74	48.26
<i>Gavelinopsis praegeri</i>	3.39	2.57	0.84	1.68	49.93

Table 4.7 (continued)

<i>Amphicoryna scalaris</i>	2.41	1.04	0.81	1.61	51.55
<i>Cibicides refulgens</i>	0.71	1.49	0.74	1.49	53.03
<i>Marginulina subalata</i>	0.06	1.46	0.74	1.48	54.51
<i>Hyalinea balthica</i>	2.64	2.91	0.69	1.38	55.89
<i>Lobatula lobatula</i>	1.38	2.21	0.68	1.37	57.25
<i>Bolivina spathulata</i>	1.18	0.95	0.68	1.37	58.62
<i>Pyrgo anomala</i>	0.08	1.3	0.64	1.27	59.89
<i>Bulimina aculeata</i>	0.57	1.66	0.63	1.25	61.14
<i>Quinqueloculina viennensis</i>	1.25	0.73	0.62	1.24	62.39
<i>Cibicidoides pseudoungerianus</i>	1.63	1.65	0.59	1.17	63.56
<i>Hoeglundina elegans</i>	1.64	1.55	0.59	1.17	64.73
<i>Sigmoilopsis schlumbergeri</i>	1.27	0.16	0.58	1.16	65.89
<i>Sphaeroidina bulloides</i>	1.38	0.65	0.56	1.12	67.01
<i>Melonis barleeanum</i>	2.71	2.76	0.55	1.11	68.11
<i>Miliolinella subrotunda</i>	0.46	1.36	0.55	1.1	69.21
<i>Spiroplectinella wrighti</i>	1.09	0.26	0.49	0.98	70.19
<i>Gyroidina neosoldanii</i>	1.24	0.68	0.49	0.97	71.16
<i>Cibicidoides kullenbergi</i>	1.06	1.35	0.47	0.93	72.1
<i>Hanzawaia boueana</i>	0.91	1.57	0.44	0.88	72.98
<i>Bolivina dilatata</i>	0.84	1.3	0.41	0.82	73.8
<i>Gyroidina altiformis</i>	0.81	0	0.4	0.81	74.61
<i>Globobulimina affinis</i>	0.81	0.14	0.37	0.74	75.34
<i>Gyroidina soldanii</i>	0.5	0.84	0.37	0.73	76.08
<i>Rectuvigerina elongatastriata</i>	0.72	0	0.36	0.72	76.8
<i>Textularia agglutinans</i>	0.69	0	0.34	0.68	77.48
<i>Nuttalides umboniferus</i>	0.72	0.08	0.34	0.67	78.15
<i>Fissurina marginata</i>	0.09	0.73	0.33	0.66	78.82
<i>Lenticulina inornata</i>	0.74	0.26	0.33	0.66	79.48

Average dissimilarity = 54.15

Species	Cluster 2		Cluster 3		Cum.%
	Av.Abund	Av.Abund	Av.Diss	Contrib%	
<i>Discanomalina coronata</i>	0.38	10.74	5.18	9.56	9.56
<i>Uvigerina peregrina</i>	13.09	4.61	4.24	7.83	17.39
<i>Cibicidoides</i> sp.	1.12	6.85	2.86	5.28	22.68
<i>Bulimina marginata</i>	7.97	2.25	2.86	5.28	27.96
<i>Hyalinea balthica</i>	7.14	2.91	2.12	3.91	31.87
<i>Bolivina difformis</i>	1.87	5.75	1.94	3.58	35.45
<i>Globocassidulina subglobosa</i> group	11.86	9.69	1.78	3.29	38.74
<i>Cibicidoides pachyderma</i>	0.16	3.68	1.76	3.25	41.99
<i>Bulimina striata</i>	3.03	0.73	1.15	2.12	44.11
<i>Angulogerina angulosa</i>	0.57	2.71	1.1	2.03	46.14
<i>Uvigerina mediterranea</i>	2.78	0.68	1.07	1.98	48.12
<i>Amphicoryna scalaris</i>	3.03	1.04	1	1.85	49.97
<i>Sigmoilopsis schlumbergeri</i>	2.16	0.16	1	1.85	51.82
<i>Cassidulina laevigata</i>	1.42	2.92	1	1.84	53.66
<i>Bolivina subaenariensis</i>	0.67	2.37	0.85	1.57	55.23
<i>Melonis barleeanum</i>	3.99	2.76	0.78	1.44	56.67
<i>Cassidulina carinata</i>	2.96	2.43	0.75	1.38	58.05
<i>Gavelinopsis praegeri</i>	1.08	2.57	0.75	1.38	59.43
<i>Marginulina subalata</i>	0	1.46	0.73	1.34	60.77
<i>Sphaeroidina bulloides</i>	1.94	0.65	0.71	1.31	62.08
<i>Bolivina spathulata</i>	1.39	0.95	0.71	1.31	63.39
<i>Lobatula lobatula</i>	1.35	2.21	0.69	1.27	64.66
<i>Cibicides refulgens</i>	0.44	1.49	0.67	1.24	65.9
<i>Pyrgo anomala</i>	0.06	1.3	0.64	1.18	67.08
<i>Bulimina aculeata</i>	1.58	1.66	0.58	1.07	68.14
<i>Miliolinella subrotunda</i>	0.23	1.36	0.57	1.05	69.2
<i>Hanzawaia boueana</i>	0.45	1.57	0.57	1.05	70.25
<i>Spiroplectinella wrightii</i>	1.31	0.26	0.56	1.04	71.29
<i>Gyroidina altiformis</i>	1.08	0	0.54	1	72.28
<i>Bigenerina nodosaria</i>	1.19	0.14	0.53	0.98	73.27
<i>Hoeglundina elegans</i>	1.4	1.55	0.53	0.97	74.24
<i>Bolivina suspinescens</i>	1.05	0	0.52	0.97	75.21
<i>Melonis pompilioides</i>	1.21	0.24	0.49	0.91	76.11
<i>Cibicidoides kullenbergi</i>	0.57	1.35	0.46	0.84	76.95
<i>Planulina ariminensis</i>	0.96	0.08	0.44	0.81	77.76
<i>Quinqueloculina viennensis</i>	0.44	0.73	0.44	0.81	78.57
<i>Spiroloculina excavata</i>	0	0.87	0.43	0.8	79.37

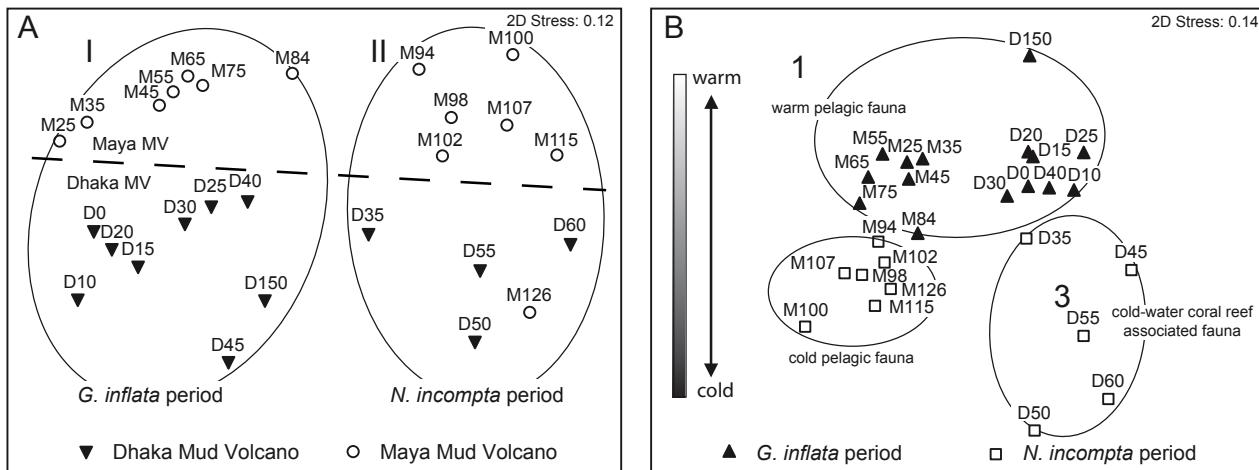


Figure 4.4 Non-metric Multi Dimensional Scaling (nMDS) plot obtained from the Bray-Curties similarity matrix of planktonic (A) and benthic (B) foraminifera compositional data. (A) Planktonic foraminifera clusters in the nMDS in a *G. inflata* dominated group (I) and in an *N. incompta* dominated group (II; Tables 4 and 5). (B) Benthic foraminiferal assemblages plot in three clusters: (1) pelagic benthic foraminifera; (2) pelagic benthic foraminifera under more eutrophic conditions and lower oxygen; (3) cold-water coral associated foraminifera (Tab. 4.6 and 4.7).

water coral mounds. In particular, they identified the species *D. coronata*, associated with other epi- and infaunal species, as a bioindicator for cold-water coral ecosystems and uvigerinids as bioindicators for off-mound sediments. The distribution pattern of the benthic assemblages in the samples from the two mud volcanoes is consistent with their distribution pattern observed in the cold-water coral mounds off-shore Ireland. In particular, *D. coronata* and the associated benthic assemblage seem to be consistently present in

association with coral fragments (Fig. 4.6; Appendix E). The above and below sediments contain the off-mound typical association described for the Porcupine Basin and Rockall Bank, which is not present in the coral-rich levels, thus suggesting that the coral-rich layers are in situ. The fragmentation of the corals can be explained as a later effect of a weak and/or partial reactivated mud volcanic activity (e.g., weak seepage) that may produce dissolution of aragonites as observed by e.g., Foubert et al. (2008) and Kopf et al. (2004).

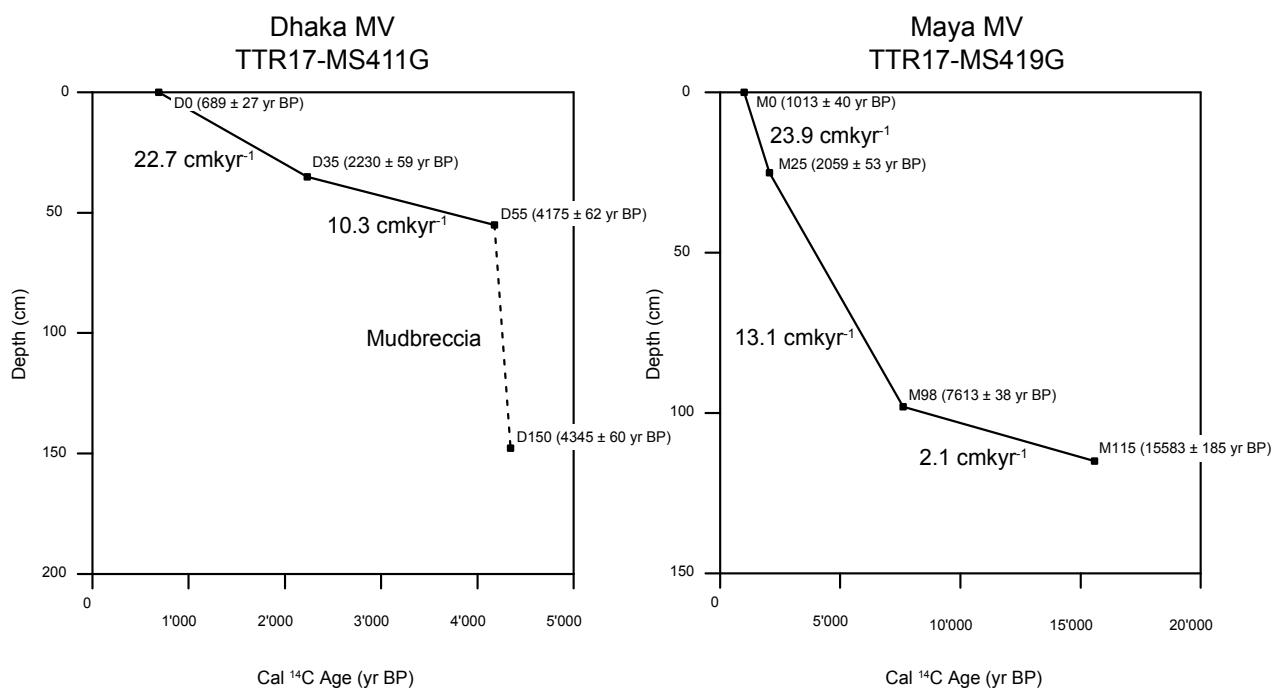


Figure 4.5 Calibrated ^{14}C ages (reservoir corrected) versus depth and sedimentation rates in cores TTR17-MS411G (Dhaka MV) and TTR17-MS419G (Maya MV).

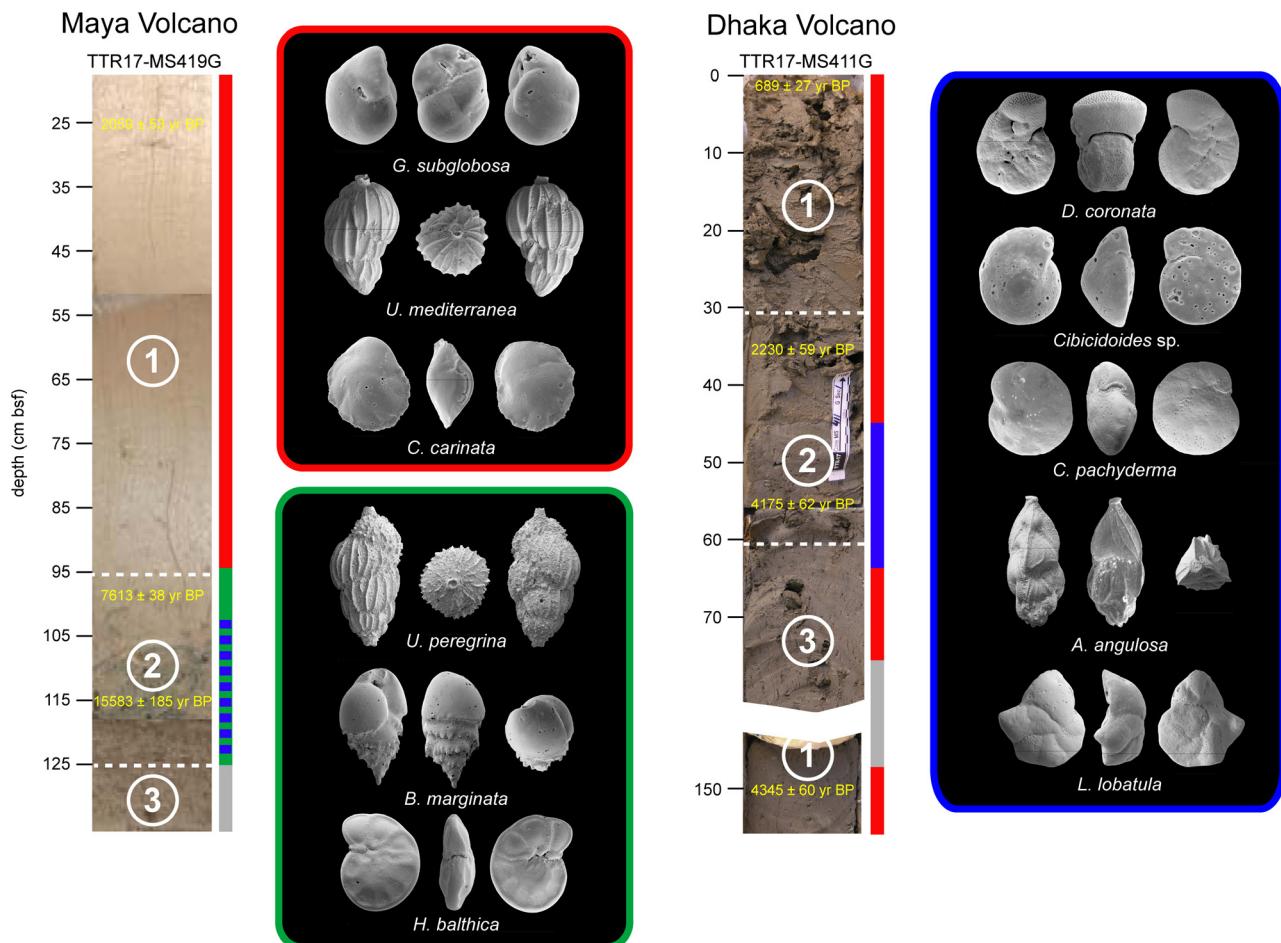


Figure 4.6 Benthic foraminiferal assemblages derived from the three nMDS clusters in cores TTR17-MS411G (Dhaka MV) and TTR17-MS419G (Maya MV). Red: pelagic benthic foraminiferal community; green: pelagic sediments with a higher nutrient supply and higher abundance of low oxygen tolerable foraminifera; blue: attached-epifaunal species typically associated to cold-water coral reefs. 1) *G. inflata* period, 2) *N. incompta* period, 3) mud breccia.

4.5.1 Similarities and differences between Dhaka and Maya mud volcanoes

Suitable substratum, low sedimentation rate, relatively long duration of stable conditions and similar density values seem to be a common feature for both volcanoes. León et al. (2007) suggested that colonization of mud volcanoes by non-chemosynthetic organisms such as corals might be related to periods of slow seepage activity inducing the formation of carbonate crusts. Sediment cores from various mud volcanoes in the Gulf of Cadiz show sequences of mud breccia with intercalation of intervals rich in cold-water coral debris (Akhmetzhanov et al., 2007; Foubert et al., 2008). However, in the cores from the Dhaka and Maya mud volcanoes, no carbonate crusts are observed. Therefore, the clasts contained in the mud breccia extruded by both mud volcanoes very reasonably acted as nucleation points and as a suitable hard substratum for cold-water coral colonization (Fig.

4.3), which was also facilitated by the elevated position of these structures (Masson et al., 2003). The AMS ^{14}C dating indicate that, according to the distribution of coral fragments and of the typical cold-water coral-associated benthic foraminiferal fauna in the cores, the duration of cold-water coral colonization was at least 2000 years at Dhaka and 7000 years at Maya, thus long enough to provide stable paleoceanographic conditions for cold-water coral development. Low sedimentation rate is one of the favourable factors for cold-water coral ecosystem development (Huvenne et al., 2002), because corals must sufficiently keep up with hemi-pelagic sedimentation (Dorschel et al., 2005; Rüggeberg et al., 2005; 2007). In both volcanoes the coral-rich layers correspond to the lower values of sedimentation rates for their respective site e.g., 10.3 cm kyr^{-1} for Dhaka and 2.1 cm kyr^{-1} for Maya mud volcano, respectively, thus confirming this essential thriving requirement and supporting our hypothesis of in-situ coral development. Dullo et al. (2008) have

demonstrated that in the North Atlantic cold-water corals only thrive in density gradients between 27.35 to 27.65 kg m⁻³, whereas Freiwald et al. (2009) show that in the Mediterranean Sea they grow between 29.08 and 29.13 kg m⁻³. These Mediterranean water density values correspond to the boundary between light and dense Mediterranean Water at 29.08 kg m⁻³ (Millot, 2009). Today the boundary between the two water masses is approximately at 430 m (Millot, 2009), thus too deep for coral growth on the two investigated mud volcanoes that are at 370 m water depth (Dhaka) and 410 m water depth (Maya). During the deglaciation times, Rogerson et al. (2008) show a shoaling of the interface between these intermediate and deep water, that we suggest shallowed up to the coral site on the mud volcanoes.

4.5.2 Paleoceanographic setting

A preliminary indication of similarity between the paleoceanographic conditions during the deposition of the hemi-pelagic drapes on top of the mud breccia of both mud volcanoes is given by the nMDS plots obtained with multivariate statistical analysis of the complete data set. In particular we observe that a *G. inflata* dominated period and a *N. incompta* dominated period are clearly discriminated and distinct in both volcanoes (Fig. 4.4A; Tab. 4.4 and 4.5). Additionally, also benthic foraminifera show clear separation in the two mud volcanoes according to a warm pelagic fauna (cluster 1) dominated by *G. subglobosa*, *U. mediterranea*, *B. marginata* and *B. difformis*, a cold pelagic fauna (cluster 2) dominated by *U. peregrina*, *G. subglobosa*, *B. marginata* and *H. balthica*, and a cold-water corals associated fauna (cluster 3) dominated by *B. difformis*, *U. peregrina*, *Cibicidoides* spp. and *D. coronata* (Figs. 4.4B and 4.6; Tab. 4.6 and 4.7). However, since the cold-water corals development on the two mud volcanoes occurred at different times, we discuss them separately to investigate if these ecosystems responded to similar paleoceanographic settings and/or to identify a causal link to possible more extended and/or global paleoceanographic events.

4.5.3 Maya Mud Volcano

An indication of the paleoceanographic conditions of surface waters can be obtained by the distribution patterns of planktonic foraminiferal assemblages. The sediments corresponding to the coral-rich layer spanning a time interval from 7613 +/- 38 years BP to slightly older than 15583 +/- 185 years BP, were deposited on the top of the mud breccia and are dominated by *N. incompta* with very low contribution of warm water species (e.g., *Globigerinoides* spp.) (Figs. 4.3C and 4.4A; Tab. 4.4 and 4.5). This interval

corresponds to a deglaciation time including the Bølling-Allerød period, the Younger Dryas, and the Early Holocene. It also contains a few specimens of *D. coronata*, *U. peregrina*, *B. marginata*, *H. balthica* and *A. angulosa* (Fig. 4.4B; Tables 4.6 and 4.7). According to Margreth et al. (2009), these lines of evidence support the interpretation of an environment characterized by enhanced nutrient supply to the surface of the Alboran Sea (e.g., Rogerson et al., 2008) and by cooler and nutrient richer waters at the sea floor, as also corroborated by high Total Organic Carbon values in the Alboran deep basin (e.g., Jimenez-Espejo et al., 2008; Rogerson et al., 2008). Presently the Alboran Sea is considered as one of the highest productivity areas associated with upwelling activity (Jimenez-Espejo et al., 2008; Morel, 1991) in the otherwise oligotrophic Mediterranean Sea (Cruzado, 1985). The occurrence of a few specimens of *D. coronata* indeed, underlines the presence of cold-water corals, but its low contribution points to a reef further away, to a small patch reef, or to the presence of solitary corals e.g., *Desmophyllum* spp. This period is also interpreted, based on other proxies, to be characterized by fresh, enhanced vertical mixing, and wet conditions on continent (Frisia et al., 2006; Zanchetta et al., 2007). According to Jimenez-Espejo et al. (2008) and Pérez-Folgado et al. (2003) these conditions ended between 7.7 and 7.2 ka BP, when a major oceanographic change occurred, giving rise to the modern setting of eddies and frontal circulation patterns of the Western Mediterranean Sea. The abundance curve of *N. incompta* reverses its trend with respect to the *Globigerinoides* spp. and *G. inflata* at 85 cm, 13 cm above the sample dated at 7613 +/- 38 years BP (Fig. 4.3C). A similar reversal was also observed at about 8 ka by Rohling et al. (1995) and at 7/8 ka by Rohling and De Rijk (1999), attributed to a shallowing of the pycnocline/nutricline at this time due to the on-set of modern conditions (Fairbanks and Wiebe, 1980; Fairbanks et al., 1982; Reynolds and Thunell, 1986; Rohling et al., 1995; Sautter and Thunell, 1989; Weaver and Pujol, 1988). Our data confirm the age of the circulation shift as reported by these previous authors and allow to link it to the end of the African Humid Period which started as early as 7 – 8 ka in many North African areas (Liu et al., 2007). The slight discrepancy in the age between the present research and the cited literature may be due to two factors: 1) an additional reversal younger than that observed in Rohling et al. (1995) cannot be ruled out; 2) Rohling et al. (1995) derive their age model on core KS310 from three non calibrated data points obtained by Pujol and Vergnaud-Grazzini (1989) on Core KS8232 adjacent to core KS310. The hemi-pelagic drape above the interval, where the reverse of

N. incompta occurs, contains dominant *G. inflata* and reflects paleoceanographic conditions more similar to the modern ones with a relatively deep pycnocline (~200 m) and a surface layer containing actively meandering jets of Modified Atlantic Waters (Rohling et al., 1995).

4.5.4 Dhaka Mud Volcano

The coral-rich interval at Dhaka mud volcano developed between 4175 ± 62 and 2230 ± 59 years BP contains planktonic and benthic foraminiferal assemblages strictly similar to those observed at Maya mud volcano. Planktonic assemblages are dominated by *N. incompta*, suggesting a shallow pycnocline/nutricline and high nutrient availability (Figs. 4.3A and 4.4A). The benthic foraminiferal assemblages contain abundantly the cold-water coral associated *D. coronata* (Margreth et al., 2009), together with other epifaunal species and *A. angulosa*. The presence of this species points toward an environment characterized by high energy and strong currents (Mackensen et al., 1985), that are important for the lateral transport of sufficient fresh food particle to the suspension feeding cold-water corals (Frederiksen et al., 1992; White et al., 2005, 2007) and are conducive of their development and growth (e.g., Roberts et al., 2006). Therefore, we suggest that the top of the mud breccia at Dhaka MV was colonized by a more patch reef-like ecosystem than at Maya MV.

The shift between the *N. incompta* and the *G. inflata* dominated assemblages is less marked than at Maya MV and is observed at about 30 cm (~2230 ± 59 years BP). The interpretation of this shift in the circulation from high productivity and shallow pycno/nutricline to a more modern system dominated by less nutrient supply and a deeper pycno/nutricline is more difficult to explain with respect to the oldest similar event documented at Maya MV. Pollen records identify a progressive climatic aridity along the Mediterranean coasts between 6000 and 4000 years BP, roughly corresponding to the end of the African Humid Period at 5500 years BP, with dry conditions observed on the continental records (Barriendos and Martin-Vide, 1998; De Menocal et al., 2000; Liu et al., 2007; Swezey, 2001). At about 4000 years BP the progressive development of complex human society occurred along the Mediterranean coasts (Roberts, 1989) resulting in a drastic anthropogenic-related change in the natural vegetation in the Eastern Mediterranean clearly detectable in pollen diagrams (Van Zeist et al., 1975) and culminating at about 3400 to 2400 years BP (Baruch, 1994; Roberts, 1989). Riehl (2009) observed consistent changes in vegetation at

about 2000 years BP in correlation to increased aridity along the Eastern Mediterranean coasts. Although this early human impact is mainly recorded in the Eastern Mediterranean, we suggest that it may have had also affected the western basin as a combined effect of human impact on a fragile environment which was already stressed by desiccation.

4.5.5 Relation to mud volcano-related seepage activity

When cold-water coral ecosystems were discovered in the North Atlantic, Hovland and Thomsen (1997) proposed a relation between the development of these peculiar ecosystems and seepage of light hydrocarbons as trigger of local primary production. In this context, the mud volcanism in the Alboran Sea has the potential to support cold-water coral ecosystems as many mud volcanoes in the region are still active, e.g., the Carmen mud volcano, where gas bubbling was documented during the TTR-17 Leg 1 (Comas et al., in press). However, Becker et al. (2009) showed that the presence of cold-water corals is not primarily due to nutrients linked to seepage activity but rather to the formation of seep-related carbonate crusts on elevated topography characterized by intense bottom currents and mixing resulting in increased food supply (Roberts et al., 2006). The Dhaka and Maya mud volcanoes are presently not active, and is therefore difficult to prove a possible relationship between seepage, nutrient supply and cold-water corals. However, oxygen and carbon isotopic values from shells of *Discanomalina coronata* from the Dhaka mud volcano (Tab. 4.3) do not show the strong isotopic anomaly found for foraminifera associated to methane seeps as e.g., in Hill et al. (2003), Rathburn et al. (2000) and Sen Gupta et al. (1997). Therefore, seepage activity during cold-water coral development on the Dhaka mud volcano can be ruled out.

4.6 CONCLUSION

Cold-water coral-rich layers were found at the top of the Dhaka and Maya mud volcanoes in the Alboran Sea. On both mud volcanoes the extruded mud breccia provides the nucleation point for the colonization and development of the corals. The co-occurrence of cold-water corals and the benthic foraminifera *D. coronata* together with the associated in- and epi-fauna as observed in the Porcupine Basin and on the Rockall Bank (Margreth et al., 2009; Fig. 4.6) suggests that these ecosystems are in situ. Two phases of cold-water corals growth are observed: at Dhaka MV they occur between 2230 ± 59 years BP and slightly older than

4175 ± 62 years BP; at Maya they occur between 7616 ± 38 years BP and slightly older than 15583 ± 185 years BP. The development of these ecosystems at different times on two mud volcanoes that are geographically close is probably related to the nature of the substratum, which plays a very important role for cold-water corals colonization. Our data indicate the mud breccia as the potential substratum, and that the two volcanoes were active before 4175 ± 62 years BP and 15583 ± 185 years BP, respectively, thus providing suitable substratum in different time slices. Our data suggest that on the top of the Maya mud volcano only a small patch reef and/or isolated corals proliferated, whereas a more extended patch reef colonized the top of the Dhaka mud volcano. At both mud volcanoes the cold-water corals development is related to times characterized by high nutrient availability, triggered by upwelling and/or strong currents that mobilized the nutrient horizontal fluxes. These intervals are characterized by high abundances of the planktonic foraminifera *N. incompta*. The shift from the *N. incompta* dominated period to the *G. inflata* dominated period, which reflects more modern paleoceanographic conditions and a deep pycno-nutricline, seems to occur at the passage from wet to arid conditions, at the end of the African Humid Period (Maya MV) and at the initiation of the complex human society along the Mediterranean coasts with consequent impact on the natural vegetation culminating at about 3400 to 2400 years BP (Dhaka MV). At both volcanoes no relation with seepage is observed.

5 - CONCLUSIONS

5. CONCLUSIONS

The presented work provides the first detailed documentation of foraminiferal fauna associated to cold-water coral ecosystems. Foraminiferal assemblages have been quantitatively investigated at three key sites for cold-water coral growth along the European continental margin:

- Giant cold-water carbonate mounds supporting cold-water coral ecosystems in the Porcupine/Rockall area (Chapter 2).
- Widespread cold-water coral reefs on the Norwegian margin and in its upper slope area (Chapter 3).
- Buried reefs in the Western Mediterranean (Chapter 4).

Investigations on these sites allow to describe in details all species of benthic foraminifera associated to these ecosystems, to identify indicator species to demonstrate the link between foraminiferal assemblages and sedimentary facies, and to highlight the potential of this group of organisms to detect cold-water coral ecosystems in the past. In parallel, the study of planktonic foraminifera has provided the scenario for the environmental conditions of the surface water masses.

Based on the results presented and discussed in previous chapters, the following conclusions can be highlighted:

- Benthic foraminiferal assemblages associated to cold-water coral ecosystems show remarkable similarities from northern Norway to the Western Mediterranean. Such similarity may be related to environmental requirements of foraminifera similar to those of cold-water corals. In particular, the cold-water coral associated assemblages are dominated by epifaunal-attached species such as *Discanomalina coronata*, *Cibicides refulgens*, and

Lobatula lobatula but also infaunal foraminifera such as *Globocassidulina* spp., *Epistominella* spp., and *Cassidulina* spp.

- The benthic fauna can efficiently provide relevant information on currents, oxygenation and organic matter content in the sediments within and adjacent the cold-water coral reefs and/or mounds. In particular, these organisms indicate the preference of these ecosystems for strong bottom currents, well-oxygenated waters and high organic matter supply derived from seasonal phytoplankton and reaching the sea floor.
- Several species of benthic foraminifera that give important environmental indications are small-sized (63-125 µm). Our results underline the importance of analysing the small fractions to highlight the contribution of small-sized species to the interpretation of complex environmental settings.
- The planktonic fauna provides information on nutrient availability and primary productivity in surface waters. In particular, active/living reefs are accompanied by planktonic foraminiferal assemblages typical of surface waters particularly rich in nutrients. The decline of the cold-water coral ecosystems in the Alboran Sea co-occurred together with a shift in the planktonic foraminiferal assemblage from eutrophic (nutrient-rich) to more oligotrophic (nutrient-poor) conditions at the surface.
- *Discanomalina coronata* is only abundant in healthy cold-water coral ecosystems and completely absent outside the reefs and mounds. For this reason *D. coronata* is interpreted to respond to similar conditions as cold-water corals and, therefore, it is here proposed as an indicator species for healthy cold-water coral ecosystems along the European margin.
- Benthic foraminiferal assemblages provide an independent tool to identify different sedimentary facies on cold-water coral mounds on the Irish

margin, where these facies are strongly defined. Each facies is clearly identified with respect to the adjacent ones over a distance of several hundreds of meters (over 2000 m).

- In the cold-water coral reefs of the Norwegian margin, benthic foraminiferal habitats are only weakly defined and grade one into the other. The gradual changes in foraminiferal assemblages can be attributed to changes in the reef facies, which are restricted within tens of meter, whereas on the carbonate mounds the different facies occur over long distances.
- Foraminiferal diversity on the Norwegian and in along the Irish margins reefs is higher in the living parts of the cold-water coral reefs and gradually decreases towards the distal parts.
- The peculiar cold-water coral associated foraminiferal assemblage can be used to identify these ecosystems in the geologic record, e.g., even when the aragonitic corals are strongly dissolved like in the Alboran Sea.

will clarify these issues and give us an answer.

5.1. Follow-up

The study of foraminifera associated to cold-water coral ecosystem is still very far to be completed as our understanding of the dynamics of foraminiferal assemblages living in these ecosystems is still in a preliminary phase. Although presently we have a better understanding of ecological and environmental control on foraminifera and corals, we still do not completely know how these ecosystems worked in different time periods.

Several questions are still open such as:

- Can we use the same bioindicators also for living cold-water coral ecosystems in regions far from the Atlantic margin e.g., New Zealand?
- Are benthic foraminiferal assemblages associated to these ecosystems similar worldwide?
- What foraminifera do tell us about paleoceanographic changes in terms of sea level, climate, productivity and ocean chemistry in the geological past of these ecosystems?
- Can we find the same bioindicators also for past ecosystems?
- If not, which species respond to similar ecological condition as cold-water corals?
- When did foraminifera acquire these similar requirements?
- Further studies on the geochemistry of the shells, in different locations and on the geological record

REFERENCES

- Akhmetzhanov, A.M., Ivanov, M.K., Kenyon, N.H., Mazzini, A., 2007. Deep-water cold seeps, sedimentary environments and ecosystems of the Black and Tyrrhenian Seas and Gulf of Cadiz. Preliminary results of investigations during the TTR-15 cruise of Professor Logachev, June-August 2005. IOC Technical Series 72, UNESCO.
- Akhmetzhanov, A.M., Kenyon, N.H., Ivanov, M., Wheeler, A.J., Shashkin, P.V., Van Weering, T.C.E., 2003. Giant carbonate mounds and current-swept seafloors on the slopes of the southern Rockall Trough. In: Mienert, J., Weaver, P. (Eds.), European Margin Sediment Dynamics. Springer, Berlin, 203–209.
- Alexandersson, E.T., 1978. Destructive diagenesis of carbonate sediments in Eastern Skagerrak, North Sea. *Geology*, 6(6), 324-327.
- Almogi-Labin, A., Schmiedl, G., Hemleben, C., Siman-Tov, R., Segl, M., Meischner, D., 2000. The influence of the NE winter monsoon on productivity changes in the Gulf of Aden, NW Arabian Sea, during the last 530 ka as recorded by foraminifera. *Mar. Micropaleontol.* 40, 295-319.
- Altenbach, A.V., Pflaumann, U., Schiebel, R., Thies, A., Timm, S., Trauth, M., 1999. Scaling percentages and distributional patterns of benthic foraminifera with flux rates of organic carbon. *J. Foraminifer. Res.* 29 (3), 173-185.
- Altenbach, A.V., Sarnthein, M., 1989. Productivity Record in Benthic Foraminifera. In: Berger, W.H., Smetacek, V.S., Wefer, G. (Eds.), Productivity of the Oceans: Present and Past. Dahlem Conferences, 255-269.
- Alvarez-Pérez, G., Busquets, P., De Mol, B., Sandoval, N.G., Canals, M., Casamor, J.L., 2005. Deep-water coral occurrences in the Straits of Gibraltar. In: Freiwald, A., Murray, J.M. (Eds.). Cold-water corals and ecosystems. Springer-Verlag Berlin, 207-221.
- Alve, E., 1995. Benthic foraminiferal distribution and recolonization of formerly anoxic environments in Drammensfjord, southern Norway. *Mar. Micropaleontol.* 25, 169-186.
- Alve, E., Murray, J.W., 1997. High benthic fertility and taphonomy of foraminifera: a case study of the Skagerrak, North Sea. *Mar. Micropaleontol.* 31, 157-175.
- Andrews, A.H., Cordes E.E., Mahoney M.M., Munk K., Coale K.H., Cailliet G.M., Heifetz J., 2002. Age, growth and radiometric age validation of a deep-sea, habitat-forming gorgonian (*Primnoa resedaeformis*) from the Gulf of Alaska. *Hydrobiologia* 471, 101-110.
- Arnone, R.A., Wiesenburg, D.A., Saunders, K., 1990. The origin and characteristics of the Algerian current. *J. Geophys. Res.* 95, 1587-1598.
- Baco, A.R., Shank, T.M., 2005. Population-genetic structure of the Hawaiian precious coral *Corallium lauuense* (Octocorallia: Corallidae) using macrosatellites. In: A. Freiwald and J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, 663-678.
- Baraza, J., Ercilla, G., 1996. Gas-charged sediments and large pockmark like features on the Gulf of Cadiz slope (SW Spain). *Mar. Petr. Geol.* 13, 253-261.
- Baraza, J., Ercilla, G., Nelson, C.H., 1999. Potential geologic hazards on the eastern Gulf of Cadiz slope (SW Spain). *Mar. Geol.* 155 (1– 2), 191–215.
- Bartoli, G., Sarnthein, M., Weinelt, M., Erlenkeuser, H., Garbe-Schönberg, D., Lea, D.W., 2005. Final closure of Panama and the onset of northern hemisphere glaciation. *Ear. Planet. Sci. Letters* 237, 33-44.
- Barriendos, M., Martin-Vide, J., 1998. Secular climatic oscillations as indicated by catastrophic floods in the Spanish Mediterranean coastal area (14th – 19th centuries). *Clim. Change* 38, 473–491.
- Baruch, U., 1994. The Late Quaternary pollen record of the Near East. In: O., Bar-Yosef, R.S., Kra (Eds.), Late Quaternary chronology and paleoclimate of

- the Eastern Mediterranean. Radiocarbon, 103-119.
- Basso, D., Spezzaferri, S., 2000. The distribution of living (stained) benthic foraminifera in Iskenderun Bay: a statistical approach. Boll. Soc. Paleont. It. 29, 356-370.
- Bé, A.W.H., Tolderlund, D.S., 1971. Distribution and ecology of living planktonic foraminifera in surface waters of the Atlantic and Indian Oceans. In: B.M. Funnel and W.R. Riedel (Eds.), Micropaleontology of the Oceans. Cambridge University Press, London, 105-149.
- Becker, E.L., Cordes, E.E., Macko, S.A., Fisher, C.R., 2009. Importance of seep primary production to *Lophelia pertusa* and associated fauna in the Gulf of Mexico. Deep-Sea Res. I 56, 786-800.
- Belanger, B.E., Streeter, S.S., 1980. Distribution and ecology of benthic foraminifera in the Norwegian-Greenland Sea. Mar. Micropaleontol. 5, 401-428.
- Bergsten, H., Nordberg, K., Malmgren, B., 1996. Recent benthic Foraminifera traces of water masses along a transect in the Skagerrak, north-eastern North Sea. J. Sea Res. 35(1-3), 111-121.
- Berner, H., Wefer, G., 1994. Clay-mineral flux in the Fram Strait and Norwegian Sea. Mar. Geol., 116, 327-345.
- Brückert, V., Perez, M.E., Lange, C.B., 2000. Coupled primary production, benthic foraminiferal assemblage, and sulfur diagenesis in organic-rich sediments of the Benguela upwelling system. Mar. Geol. 163, 27-40.
- Bernhard, J.M., 1986. Characteristic assemblages and morphologies of benthic foraminifera from anoxic, organic-rich deposits - Jurassic through Holocene. J. Foraminiferal Res., 16(3), 207-215.
- Bernhard, J.M., Sen Gupta, B.K., 1999. Foraminifera of oxygen-depleted environments. In: B.K. Sen Gupta (Ed.), Modern Foraminifera. Kluwer Academic Press, New York, 201-216.
- Beuck, L., Freiwald, A., 2005. Bioerosion patterns in deep-water *Lophelia pertusa* (Scleractinia) thicket (Propeller Mound), northern Porcupine Seabight). In: Freiwald, A., Roberts, J.M. (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 915-936.
- Blindheim, J., 1990. Arctic intermediate water in the Norwegian Sea. Deep Sea Res. Part I, 35, 1475-1489.
- Boltovskoy, E., 1956. Applications of chemical ecology in the study of foraminifera. Micropal., 2, 321-325.
- Burdon-Jones, C., Tamps-Lyche, H., 1960. Observations on the fauna of the North Bratholmen stone-coral reef near Bergen. Arb. Univ. Bergen, Mat-Naturv. Ser. 4, 1-24.
- Caralp, M.H., 1989. Size and morphology of the benthic foraminifer *Melonis barleeanum*: Relationships with marine organic matter. J. Foraminifer. Res. 19 (3), 235-245.
- Carney, R.S., 1989. Examining relationship between organic carbon flux and deep-sea depositfeeding. In: G. Lopez, G. Taghon and J. Levinton (Eds.), Ecology of marine deposit feeders. Lecture Notes on Coastal and Estuarine Studies. Springer, New York, 31, 24-58.
- Carpenter, K.E., Niem, V.H., 1998. FAO Species Identification Guide for Fishery Purposes, The living resources of the Western Central Pacific. Volume 1. Seaweeds, corals, bivalves and gastropods. Food and Agriculture Organisation of the United Nations, Rome.
- Chalouan, A., Saji, R., Michard, A., Bally, A.W., 1997. Neogene tectonic evolution of the Southwestern Alboran Basin as inferred from seismic data off Morocco. A.A.P.G. Bull. 81, 1161-1184.
- Clarke, K., 1993. Non-parametric multivariate analyses of changes in community structure. Aust. J. Ecol. 18, 117-143.
- Clarke, K.R., Gorley, R.N., 2006. PRIMER v6: User Manual/Tutorial, PRIMER-E, Plymouth.
- Clarke, K.R., Warwick, R.M., 1994. Similarity-based testing for community pattern: the 2-way layout with no replication. Mar. Biol. 118, 167-176.
- Clarke, K.R., Warwick, R.M., 2001a. A further biodiversity index applicable to species lists: variation in taxonomic distinctness. Mar. Ecol. Prog. Ser. 216, 265-278.
- Clarke, K., Warwick, R., 2001b. Change in marine communities: an approach to statistical analysis and interpretation. PRIMER-E, Plymouth.
- Clifford, D.H.T., Stephenson, W., 1975. An introduction to numerical classification. Academic Press, New York.
- Colman, J.G., Gordon, D.M., Lane, A.P., Forde, M.J., Fitzpatrick, J.J., 2005. Carbonate mounds off Mauritania, Northwest Africa: status of deep-water corals and implications for management of fishing and oil exploration activities. In: A. Freiwald and J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, 417-441.
- Comas, M.C., Ivanov, M., 2006. Eastern Alboran margin: the transition between the Alboran and the Balearic-Algerian Basin. Introduction. In: N.H. Kenyon, M.K. Ivanov, A.M. Akhmetzhanov, and E.V. Kozlova (Eds.), Interdisciplinary geoscience studies of the Gulf of Cadiz and Western

- Mediterranean basin. IOC Tech. Ser. 70, UNESCO, 48-49.
- Comas, M.C., Pinheiro, L.M., 2007. Mud volcanism, gas hydrates and hydrocarbon-rich fluid escape structures in the Gibraltar Arc System. Exploring Escarpment Mud Mound Systems and Mud Volcanoes with new European Strategies for sustainable mid-depth coring, Magellan Workshop. Murten, Switzerland.
- Comas, M.C., Garcia-Dueñas, V., Jurado, M.J., 1992. Neogene tectonic evolution of the Alboran Sea. In: A. Maldonado (Ed.). The Alboran Sea. Geo-Mar. Lett. Spec. Issue 12, 157-164.
- Comas, M.C., Platt, J., Soto, J.I., Watts, T., 1999. The origin and tectonic history of the Alboran Basin: insights from Leg 161 results. In: R. Zahn, M.C., Comas, A. Klaus (Eds.), ODP Proc., Sci. Results 161. College Station, Texas A & M University, 555-580.
- Comas, M.C., Soto, J.I., BASALCALB cruise (TTR-9 Leg 3) Scientific Party, 2000. A tectonic overview on the mud diapirs and related mud volcanoes in the Alboran Basin. Geological Processes on European Continental Margins (TTR-9 Post-Cruise Conference), Granada, Spain. IOC Workshop Report 168, 29-30.
- Comas, M.C., Soto, J.I., Talukder, A.R., TTR-12 Leg 3 (MARSIBAL-1) Scientific Party, 2003a. Discovering active mud volcanoes in the Alboran Sea (Western Mediterranean). IOC Workshop Report 187, UNESCO, 14-16.
- Comas M.C., Talukder, A.R., Soto, J.I., 2003b. Mud volcanoes in the Alboran Sea: learning from occurrences in the western Mediterranean. Geophys. Res. Abstr. 5, 12617.
- Connell, J.H., 1978. Diversity in Tropical Rain Forests and Coral Reefs. Science 199, 1302-1310.
- Corliss, B.H., 1979. Quaternary Antarctic bottom-water history: Deep-sea benthonic foraminiferal evidence from the Southeast Indian Ocean — Inferred bottom-water routes and ecological implications. Mar. Geol 31, 115–138.
- Corliss, B.H., 1985. Microhabitats of benthic foraminifera within deep-sea sediments. Nature 314, 435-438.
- Corliss, B.H., Chen, C., 1988. Morphotype Patterns of Norwegian Sea Deep-Sea Benthic Foraminifera and Ecological Implications. Geology 16 (8), 716-719.
- Corliss, B.H., Van Weering, T.C.E., 1993. Living (stained) benthic foraminifera within surficial sediments of the Skagerrak. Mar. Geol. 111 (3-4), 323-335.
- Costello, M.J., McCrea, M., Freiwald, A., Lundälv, T., Jonsson, L., Bett, B.J., Van Weering, C.E., De Haas, H., Roberts, J.M., Allen, D., 2005. Role of cold-water *Lophelia pertusa* coral reefs as fish habitat in the NE Atlantic. In: A. Freiwald and J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, 771-805.
- Cruzado, A., 1985. Chemistry of Mediterranean waters. In: R. Margalef (Ed.), Western Mediterranean, Pergamon, Oxford, U.K., 126–147.
- Cushman, J.A., 1928. Foraminifera: Their classification and economic use. Cushman Laboratory for Foraminiferal Research Special Publication, 1.
- Darling, K.F., Kroon, D., Wade, C.M., Leigh Brown, A.J., 1996. Molecular phylogeny of the planktonic foraminifera. J. Foram. Res. 26, 324-330.
- De Haas, H., 2007. RV Pelagia cruise M07II (64PE276), Galway – Galway. Unpublished report.
- De Menocal, P. D., Ortiz, J., Guilderson, T., Sarnthein, M., 2000. Coherent high-and low-latitude climate variability during the Holocene warm period. Science 288, 2198– 2202.
- De Mol, B., Henriet, J.-P., Canals, M., 2005. Development of coral banks in Porcupine Seabight: do they have Mediterranean ancestors? In: A. Freiwald and J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, 515-533.
- De Mol, B., Van Rensbergen, P., Pillen, S., Van Herreweghe, K., Van Rooij, D., McDonnell, A., Huvenne, V., Ivanov, M., Swennen, R., Henriet, J.P., 2002. Large deep-water coral banks in the Porcupine Basin, southwest of Ireland. Mar. Geol. 188, 193-231.
- De Nooijer, L.J., Duijnstee, I.A.P., Bergman, M.J.N., Van der Zwaan, G.J., 2008. The ecology of benthic foraminifera across the Frisian Front, southern North Sea. Estuar. Coast. Shelf. Sci. 78 (4), 715-726.
- De Stigter, H.C., Jorissen, F.J., Van der Zwaan, G.J., 1998. Bathymetric distribution and microhabitat partitioning of live (Rose Bengal stained) benthic foraminifera along a shelf to bathyal transect in the southern Adriatic Sea. J. Foraminifer. Res. 28 (1), 40-65.
- Dewey, J.F., Helma, M.L., Turco, E., Hutton, D.W.H., Knott, S.D., 1989. Kinematics of the western Mediterranean. In: M.P. Coward, D. Dietrich, R.G. Park (Eds.), Alpine Tectonics. Geol. Soc. Lond. Spec. Publ., 265-283.
- Dons, C., 1944. Norges korallrev. Det Kongelige Norske Videnskabers Selskab. Forhandlinger 16, 37-82.
- D'Orbigny, A., Tableau Méthodique de la Class des Céphalopodes. Annales des Sciences Naturelles,

- Paris, Séries 1(7), 245-314.
- Dorschel, B., Hebbeln, D., Foubert, A., White, M., Wheeler, A.J., 2007a. Hydrodynamics and cold-water coral facies distribution related to recent sedimentary processes at Galway Mound west of Ireland. *Mar. Geol.* 244, 184-195.
- Dorschel, B., Hebbeln, D., Rüggeberg, A., Dullo, C., 2007b. Carbonate budget of a deep water coral mound: Propeller Mound, Porcupine Seabight. *Int. J. Earth Sci.* 96, 73-83.
- Dorschel, B., Hebbeln, D., Rüggeberg, A., Dullo, C., Freiwald, A., 2005. Growth and erosion of a cold-water coral covered carbonate mound in the Northeast Atlantic during the Late Pleistocene and Holocene. *Earth Planet. Sci. Lett.* 233, 33-44.
- Drebes, G., 1974. *Marines Phytoplankton. Eine Auswahl der Helgoländer Planktonalgen (Diatomeen. Peridineen).* Georg Thieme, Stuttgart.
- Duchemin, G., Fontanier, C., Jorissen, F.J., Barras, C., Griveaud, C., 2007. Living small-sized (63-150 µm) foraminifera from mid-shelf to mid-slope environments in the Bay of Biscay. *J. Foraminifer. Res.* 37 (1), 12-32.
- Duijnste, I.A.P., Ernst, S.R., Van der Zwaan, G.J., 2003. Effect of anoxia on the vertical migration of benthic foraminifera. *Mar. Ecol. Prog. Ser.* 246, 85-94.
- Duineveld, G.C.A., Lavaleye, M.S.S., Berghuis, E.M., 2004. Particle flux and food supply to a seamount cold-water coral community (Galicia Bank, NW Spain). *Mar. Ecol. Prog. Ser.* 277, 13-23.
- Duineveld, G.C.A., Lavaleye, M.S.S., Bergman, M.J.N., Stigter, H.C.D., Mienis, F., 2007. Trophic structure of a cold-water coral Mound community (Rockall Bank, NE Atlantik) in relation to the near-bottom particle supply and current regime. *Bull. Mar. Sci.* 81 (3), 449-467.
- Dullo, W.C., Flögel, S., Rüggeberg, A., 2008. Cold-water coral growth in relation to the hydrography of the Celtic and Nordic European continental margin. *Mar. Ecol. Prog. Ser.* 371, 165-176.
- Duncan, P.M., 1870. A description of the *Madreporaria* dredged up during the expeditions of HMS Porcupine. *Proc. Roy. Soc. Lond.* 18, 289-301.
- Duncan, P.M., 1873. A description of the *Madreporaria* dredged up during the expeditions of HMS Porcupine in 1869 and 1870. *Trans. Zool. Soc. Lond.* Part 1 (8), 303-344.
- Duncan, P.M., 1878. A description of the *Madreporaria* dredged up during the expeditions of HMS Porcupine in 1869 and 1870. *Trans. Zool. Soc. Lond.* Part 2 (10), 235-249.
- Emiliani, C., 1955. Pleistocene temperatures. *J. Geol.*, 63, 538-578.
- Ellis, B.F., Messina, A.R., 1940 and later. Catalogue of Foraminifera. American Museum of Natural History, New York.
- Ernst, S.R., 2002. An experimental study on the proxy value of benthic foraminifera - The impact of physical disturbance, oxygen depletion and organic flux. Unpublished Ph.D. Thesis, University of Utrecht, Netherlands.
- Ernst, S.R., Bours, R., Duijnste, I., Van der Zwaan, B., 2005. Experimental effects of an organic matter pulse and oxygen depletion on a benthic foraminiferal shelf community. *J. Foraminifer. Res.* 35 (3), 177-197.
- Everitt, B., 1978. *Graphical techniques for multivariate data.* Heinemann, London.
- Fairbanks, R.G., Sverdrup, M., Free, R., Wiebe, P.H., Bé, A.W.H., 1982. Vertical distribution of living planktonic foraminifera from the Panama Basin. *Nature* 298, 841-844.
- Fairbanks, R.G., Wiebe, P.H., 1980. Foraminifera and Chlorophyll Maximum: vertical distribution, seasonal succession, and paleoceanographic significance. *Science* 209, 1524-1526.
- Fariduddin, M., Loubere, P., 1997. The surface ocean productivity response of deeper water benthic foraminifera in the Atlantic Ocean. *Mar. Micropaleontol.* 32 (3-4), 289-310.
- Fernández-Ibáñez, F., Soto, J.I., Zoback, M.D., Morales, J., 2007. Present-day stress field in the Gibraltar Arc (western Mediterranean). *J. Geophys. Res.* 112, B08404, doi: 10.1029/2006JB004683.
- Fernandez-Puga, M.C., Vasquez, J.T., Somoza, L., Diaz del Rio, V., Medialdea, T., Mata, M.P., Léon, R., 2007. Gas-related morphologies and diapirism in the Gulf of Cadiz. *Geo-Mar. Lett.* 27 (2-4), 213-221.
- Fichtel, L., Moll, J.P.C., 1798. *Testacea Microscopica Aliaque Minuta ex Generibus Argonauta et Nautilus ad Naturam Delineata et Descripta*, Anton Pichler, Vienna.
- Field, J.G., Clarke, K.R., Warwick, R.M., 1982. A practical strategy for analysing multispecies distribution patterns. *Mar. Ecol. Prog. Ser.* 8, 37-52.
- Fisher, R.A., Corbet, A.S., Williams, C.B., 1943. The relationship between the number of species and the number of individuals in a random sample of an animal sample. *J. Anim. Ecol.* 12, 42-58.
- Fontanier, C., Jorissen, F.J., Chaillou, G., Anschutz, P., Gremare, A., Griveaud, C., 2005. Live foraminiferal faunas from a 2800 m deep lower canyon station from the Bay of Biscay: Faunal response to focusing of refractory organic matter. *Deep Sea Res. I* 52 (7), 1189-1227.

- Fontanier, C., Jorissen, F.J., Licari, L., Alexandre, A., Anschutz, P., Carbonel, P., 2002. Live benthic foraminiferal faunas from the Bay of Biscay: faunal density, composition, and microhabitats. Deep Sea Res. I 49, 751–785.
- Fosså, J.H., Mortensen, P.B., 1998. Artsmangfoldet på *Lophelia*-korallrev og metoder for kartlegging og overvåkning. Fisken og Havet, 17, 1-95.
- Fosså, J.H., Lindberg, B., Christensen, O., Lundälv, T., Svellingen, I., Mortensen, P.B., Alvsvåg, J., 2005. Mapping of *Lophelia* reefs in Norway: experiences and survey methods. In: A. Freiwald and J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, 359-391.
- Fosså, J.H., Mortensen, P.B., Furevik, D.M., 2000. Lophelia-korallrev langs norskekysten forekomst og tilstand. Fisken og Havet, 2, 1-94.
- Fosså, J.H., Mortensen, P.B., Furevik, D.M., 2002. The deep-water coral *Lophelia pertusa* in Norwegian waters: distribution and fishery impacts. Hydrobiologia 471, 1–12.
- Foubert, A., Henriet, J.-P., 2009. Nature and Significance of the Recent Carbonate Mound Record, the Challenger Code. Springer Verlag, Dordrecht, Heidelberg, London, New York.
- Foubert, A., Beck, T., Wheeler, A.J., Opderbecke, J., Grehan, A., Klages, M., Thiede, J., Henriet, J.-P., Party, P.A.-X.a.S., 2005. New view of the Belgica Mounds, Porcupine Seabight, NE Atlantic: preliminary results from the Polarstern ARK-XIX/3a ROV cruise. In: A. Freiwald, J.M., Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 403-415.
- Foubert, A., Depreiter, D., Beck, T., Maignien, L., Pannemans, B., Frank, N., Blamart, D., Henriet, J.-P., 2008. Carbonate mounds in a mud volcano province off north-west Morocco: key to processes and controls. Mar. Geol. 248, 74-96.
- Foubert, A., Van Rooij, D., Blamart, D., Henriet, J.-P., 2007. X-ray imagery and physical core logging as a proxy of the content of sediment cores in cold-water coral mound provinces: a case study from Porcupine Seabight, SW of Ireland. Int. J. Earth Sci. 96, 141-158.
- Frederiksen, R., Jensen, A., Westerberg, H., 1992. The distribution of the scleractinian coral *Lophelia pertusa* around the Faroe islands and the relation to internal tidal mixing. Sarsia, 77: 157-171.
- Freiwald, A., 2002. Reef-Forming Cold-Water Corals. In: G. Wefer, D. Billett, D. Hebbeln, B.B. Jørgensen, M. Schlüter, T.C.E. Van Weering, (Eds.), Ocean Margin Systems. Springer Verlag, Berlin, Heidelberg, New York, 365-385.
- Freiwald, A., Dullo, C., shipboard party, 2000. RV Poseidon Cruise 265, Thórshavn – Galway – Kiel. Unpublished report.
- Freiwald, A., Roberts, J.M., 2005. Cold-water coral ecosystems. Springer-Verlag, Berlin, Heidelberg.
- Freiwald, A., Schönfeld, J., 1996. Substrate pitting and boring pattern of *Hyrrokkin sarcophaga* Cedhagen, 1994 (Foraminifera) in a modern deep-water coral reef mound. Mar. Micropaleontol. 28, 199-207.
- Freiwald, A., shipboard party, 2002. RV Poseidon Cruise 292, Reykjavík – Galway. Unpublished report.
- Freiwald, A., Beuck, L., Rüggeberg, A., Taviani, M., Hebbeln, D., 2009. The white coral community in the central Mediterranean Sea revealed by ROV surveys. Oceanography 22, 58-74.
- Freiwald, A., Fosså, J.H., Grehan, A., Koslow, T., Roberts, J.M., 2004. Cold-water Coral Reefs, UNEP-WCMC, Cambridge, UK.
- Freiwald, A., Heinrich, R., Pätzold, J., 1997. Anatomy of a deep-water coral reef mound from Stjernsund, West-Finnmark, northern Norway. In: N. James, J. Clarke (Eds.), Cool-water carbonates. SEPM Special Publication, 141-161.
- Freiwald, A., Hühnerbach, V., Lindberg, B., Wilson, J.B., Campbell, J., 2002. The Sula Reef Complex, Norwegian shelf. Facies 47, 179-200.
- Freiwald, A., Wilson, J.B., Henrich, R., 1999. Grounding Pleistocene icebergs shape recent deep-water coral reefs. Sed. Geol. 125, 1-8.
- Frisia, S., Borsato, A., Mangini, A., Spötl, C., Madonia, G., Sauro, U., 2006. Holocene climate variability from a discontinuous stalagmite record and the Mesolithic to Neolithic Transition. Quat. Res. 66, 388–400.
- Frizon de Lamotte, D., Michard, A., Saddiqi, O., 2006. Quelques développements récents sur la géodinamique du Maghreb. C. R. Géosci. 338 (1-2), 1-10.
- Gardner, J.M., 2001. Mud volcanoes revealed and sampled on the western Moroccan continental margin. Geophys. Res. Lett. 28, 334-342.
- Gass, S.E., Willison, J.H., 2005. An assessment of the distribution of deep-sea corals in Atlantic Canada by using both scientific and local forms of knowledge. In: A. Freiwald, J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 223-245.
- Gooday, A.J., 1986. Meiofaunal foraminiferans from the bathyal Porcupine Seabight (northeast Atlantic): size structure, standing stock, taxonomic composition, species diversity and vertical distribution in the sediments. Deep Sea Res. I 33 (10), 1345-1373.

- Gooday, A.J., 1988. A response by benthic foraminifera to the deposition of phytodetritus in the deep sea. *Nature* 322, 70-73.
- Gooday, A.J., 1994. The biology of deep-sea foraminifera: A review of some advances and their applications in paleoceanography. *Palaios* 9, 14–31.
- Gooday, A.J., 2003. Benthic foraminifera (Protista) as tools in deep-water palaeoceanography: Environmental influences on faunal characteristics. *Adv. Mar. Biol.* 46, 1–90.
- Gooday, A.J., Hughes, J.A., 2002. Foraminifera associated with phytodetritus deposits at a bathyal site in the northern Rockall Trough (NE Atlantic): seasonal contrasts and a comparison of stained and dead assemblages. *Mar. Micropaleontol.* 46, 83–110.
- Gooday, A.J., Bett, B.J., Pratt, D.N., 1993. Direct observation of episodic growth in an abyssal xenophyophore (Protista). *Deep Sea Res. I* 40, 2131–2143.
- Grasmueck, M., Eberli, G.P., Viggiano, D.A., Correa, T., Rathwell, G., Luo, J., 2006. Autonomous underwater vehicle (AUV) mapping reveals coral mound distribution, morphology, and oceanography in deep water of the Straits of Florida. *Geoph. Res. Letters* 33, 1-6.
- Gravier, C., 1915. Note préliminaire sur les *Madréporaires* recueillis au cours des croisières de la Princesse-Alice et de l'Hirondelle II, de 1893 à 1913 inclusivement. *Bull. Inst. Océanogr. Monaco* 12 (304), 1–22.
- Gravier, C., 1920. *Madréporaires* provenant des campagnes des yachts Princesse Alice et Hirondelle II (1893–1913). Résultats des Campagnes Scientifiques du Prince Albert Ier de Monaco 55 (1–123).
- Grigg, R.W., 1984. Resource management of precious corals: a review and application to shallow water reef building corals. *Mar. Ecol.* 5, 57–74.
- Grimsdale, T.F., Van Morkhoven, F.P.C.M., 1955. The ratio between pelagic and benthonic foraminifera as a means of estimating depth of deposition of sedimentary rocks. 4th World Petroleum Congress, 473–491.
- Gross, O., 2000. Influence of temperature, oxygen and food availability on the migrational activity of bathyal benthic foraminifera: evidence by microcosm experiments. *Hydrobiologia* 426, 123–137.
- Gubbay, S., 2003. Seamounts of the North-East Atlantic. WWF-OASIS Reports.
- Gunnerus, J.E., 1768. Om nogle Norske coraller. Kongl norske videnskelsks 4, 38–73.
- Gupta, A.K., 1997. Paleoceanographic and paleoclimatic history of the Somali Basin during the Pliocene-Pleistocene: Multivariate analyses of benthic foraminifera from DSDP Site 241 (Leg 25). *J. Foraminifer. Res.* 27 (3), 196–208.
- Gupta, A.K., Thomas, E., 1999. Latest Miocene through Pleistocene paleoceanographic evolution of the northwestern Indian Ocean (DSDP Site 219): global and regional factors. *Paleoceanography* 14, 62–73.
- Gutscher, M.A., Malod, J., Rehault, J.P., Contrucci, I., Klingelhoefer, F., Mendes-Victor, L., Spakman, W., 2002. Evidence for active subduction beneath Gibraltar. *Geology* 30, 1071–1074.
- Hald, M., Vorren, T., 1984. Modern and Holocene foraminifera and sediments on the continental shelf off Troms, North Norway. *Boreas* 13, 133–154.
- Hawkes, A.D., Scott, D.B., 2005. Attached benthic Foraminifera as indicators of past and present distribution of the coral *Primnoa resedaeformis* on the Scotian Margin. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 881–894.
- Heifetz, J., 2002. Coral in Alaska: distribution, abundance, and species associations. *Hydrobiologia*, 471, 19–28.
- Hemleben, C., Spindler, M., Anderson, O.R., 1989. Modern planktonic foraminifera. Springer Verlag, New York.
- Henriet, J.P., De Mol, B., Pillen, S., Vanneste, M., Van Rooij, D., Versteeg, W., Croker, P.F., Shannon, P.M., Unnithan, V., Bouriak, S., Chachkine, P., 1998. Gas hydrate crystals may help build reefs. *Nature* 391, 648–649.
- Henriet, J.P., Guidard, S., Team, O.P., 2002. Carbonate Mounds as a Possible Example for Microbial Activity in Geological Processes. In: G. Wefer, D. Billett, D. Hebbeln, B.B. Jørgensen, M. Schlüter, T.C.E. Van Weering (Eds.), *Ocean Margin Systems*. Springer Verlag, Berlin, Heidelberg, New York, 437–455.
- Henry, L.A., Roberts, J.M., 2007. Biodiversity and ecological composition of macrobenthos on cold-water coral mounds and adjacent off-mound habitat in the bathyal Porcupine Seabight, NE Atlantic. *Deep Sea Res. Part I*, 54, 654–672.
- Herguera, J.C., Berger, W.H., 1991. Paleoproductivity from benthic foraminifera abundance: glacial to postglacial change in the west-equatorial Pacific. *Geology* 19, 1173–1176.
- Hermelin, J.O.R., 1992. Variations in the benthic foraminiferal fauna of the Arabian Sea: response to changes in upwelling intensity? In: C.P.

- Summerhayes, W.J., Prell, K.C., Emeis (Eds.), Upwelling Systems: Evolution since the Miocene. Geol. Soc. Spec. Publ. 64, 51-166.
- Hess, S., Jorissen, F.J., Venet, V., Abu-Zied, R., 2005. Benthic foraminiferal recovery after recent turbidite deposition in Cap Breton Canyon (Bay of Biscay). *J. Foraminifer. Res.* 35, 114-129.
- Hill, T.M., Kennett, J.P., Spero, H.J., 2003. Foraminifera as indicators of methane-rich environments, a study of modern methane seeps in Santa Barbara Channel, California. *Mar. Micropaleontol.* 49, 123-138.
- Holbrook, W.S., Fer, I., Schmitt, R.W., 2009. Images of internal tides near the Norwegian continental slope. *Geophys. Res. Lett.*, 36, 5.
- Hovland, M., 2008. Deep-water coral reefs. Unique biodiversity hot-spots. Springer Praxis Books.
- Hovland, M., Thomsen, E., 1997. Cold-water corals— are they hydrocarbon seep related? *Mar. Geol.* 137, 159–164.
- Hovland, M., Croker, P.F., Martin, M., 1994. Fault-associated seabed mounds (carbonate knolls?) off western Ireland and north-west Australia. *Mar. Pet. Geol.* 11 (2), 232-246.
- Hovland, M., Mortensen, P.B., Brattegård, T., Strass, P., Rokoengen, K., 1998. Ahermatypic coral banks of mid-Norway: evidence for a link with seepage of light hydrocarbons. *Palaios* 13, 189-200.
- Hühnerbach, V., Blondel, P., Huvenne, V.A.I. and Freiwald, A., 2008. Habitat mapping of a cold-water coral reef off Norway, with a comparison of visual and computer-assisted methods to interpret sidescan sonar data. In: B.J. Todd and H.G. Greene (Eds.), Mapping the seafloor for habitat characterisation. *Geol. Assoc. Canada, St Johns NL, Canada*, 291-302.
- Huvenne, V.A.I., Bailey, W.R., Shannon, P.M., Neath, J., Di Primio, R., Henriet, J.-P., Horsfield, B., De Haas, H., Wheeler, A.J., Olu-Le Roy, K., 2007. The Magellan mound province in the Porcupine Basin. *Int. J. Earth Sci.*, 96, 85-101.
- Huvenne, V.A.I., Beyer, A., De Haas, H., Dekindt, K., Henriet, J.P., Kozachenko, M., Olu-Le Roy, K., Wheeler, A.J., Participants, T.P.a.C.c., 2005. The seabed appearance of different coral bank provinces in the Porcupine Seabight, NE Atlantic: results from sidescan sonar and ROV sea-bed mapping. In: Freiwald, A., Roberts, J.M. (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 535-569.
- Huvenne, V.A.I., Blondel, P., Henriet, J.P., 2002. Textural analyses of sidescan sonar imagery from two mound provinces in the Porcupine Seabight. *Mar. Geol.* 189 (3-4), 323-341.
- Huvenne, V.A.I., De Mol, B., Henriet, J.-P., 2003. A 3D seismic study of the morphology and spatial distribution of buried coral banks in the Porcupine Basin, SW of Ireland. *Mar. Geol.* 198, 5-25.
- Huvenne, V.A.I., Masson, D.G., Wheeler, A.J., 2009. Sediment dynamics of sandy contourite: the sedimentary context of the Darwin cold-water coral mounds, Northern Rockall Trough. *Int. J. Earth. Sci.*, 98, 865-884.
- Ivanov, M. K., Kenyon, N., Nielsen, T., Wheeler, A., Monteiro, J., Gardner, J., Comas, M., Akhmanov, G., Akhmetzhanov, A., Scientific Party of the TTR-9 cruise, 2000. Goals and principal results of the TTR-9 cruise. *Proc. Int. Conf. Geological Processes on European Continental Margins. IOC/UNESCO Workshop Report* 168, 3-4.
- Ivanov M.K., Pinheiro, L., Stadnitskaia, A., Blinova, V., 2001. Hydrocarbon seeps on Deep Portuguese Margin. Final Proc. 11th Meeting of the E. G. U., Strasbourg, France, CC11, 160.
- James, N.P., Bourque, P.A., 1992. Reefs and Mounds. In: R.G. Walker, N.P. James (Eds.), Facies models. *Geol. Assoc. Can., St. Jones*, 323-345.
- Jaques, T.G., 1972. On the occurrence of deep-sea corals on one of the Corner Seamounts. Graduate School of Oceanography, Rhode Island, Research Project OCG 540.
- Jarke, J., 1960. Beitrag zur Kenntnis der Foraminiferenfauna der mittleren und westlichen Barents-See. *Int. Rev. Ges. Hydrobiol.* 45, 581-654.
- Jensen, A., Frederiksen, R., 1992. The fauna associated with the bank-forming deepwater coral *Lophelia pertusa* (Scleractinia) on the Faroe shelf. *Sarsia* 77, 53-69.
- Jimenez-Espejo, F.J., Martinez-Ruiz, F., Sakamoto, T., Iijima, K., Gallego-Torres, D., Harada, N., 2008. Paleoenvironmental changes in the Western Mediterranean since the last glacial maximum: high resolution multiproxy record from the Algero-Balearic basin. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 246, 292-306.
- Jorissen, F.J., 1988. Benthic foraminifera from the Adriatic Sea; Principles of phenotypic variation. *Utr. Micropaleontol. Bull.* 37, 176.
- Jorissen, F.J., Barmawidjaja, D.M., Puskaric, S., Van der Zwaan, G.J., 1992. Vertical distribution of benthic Foraminifera in the Northern Adriatic Sea. The relation with high organic flux. *Mar. Micropaleontol.* 19, 131–146.
- Jorissen, F.J., De Stigter, H.C., Widmark, J.G.V., 1995. A conceptual model explaining benthic foraminiferal microhabitats. *Mar. Micropaleontol.* 22, 3–15.

- Jorissen, F.J., Fontanier, C., Thomas, E., 2007. Paleoceanographical Proxies Based on Deep-Sea Benthic Foraminiferal Assemblage Characteristics. In: C. Hillaire-Marcel, A. De Vernal (Eds.), *Proxies in late Cenozoic Paleoceanography*. Elsevier, 263-325.
- Jorissen, F.J., Wittling, I., Peypouquet, J.P., Rabouille, C., Relexans, J.C., 1998. Live benthic foraminiferal faunas off Cape Blanc, NW-Africa: Community structure and microhabitats. *Deep Sea Res. I* (45), 2157-2188.
- Joubin, L., 1922a. Les coraux de mer profonde nuisibles aux chalutiers. *Notes et Mémoires. Ov. Sci. Tech. Pêches Maritimes* 18, 5-16.
- Joubin, L., 1922b. Distribution géographique de quelques coraux abyssaux dans les mers occidentales européennes. *Comptes Rendus, Académie des Sciences de Paris* 175, 930-933.
- Joubin, L., 1923. Les coraux de mer profonde du plateau continental atlantique. Rapports et procès-verbaux des réunions., International Council for the Exploration of the Sea, Copenhagen, Denmark, 43-46.
- Kaiho, K., 1994. Benthic foraminiferal dissolved-oxygen index and dissolved-oxygen levels in the modern ocean. *Geology* 22, 719-722.
- Kaiho, K., 1999. Effect of organic carbon flux and dissolved oxygen on the benthic foraminiferal oxygen index (BFOI). *Mar. Micropaleontol.* 37, 67-76.
- Kano, A., Ferdelman, T.G., Williams, T., Henriet J.P., Ishikawa, T., Kawagoe, N., Takashima, C., Kakizaki, Y., Abe, K., Sakai, S., Browning, E.L., Li, X., Integrated Ocean Drilling Program Expedition 307 Scientists, 2007. Age constraints on the origin and growth history of a deep-water coral mound in the northeast Atlantic drilled during Integrated Ocean Drilling Program Expedition 307. *Geology* 35(11), 1051-1054.
- Kenkel, N.C., Orloci, L., 1986. Applying metric and nonmetric multidimensional scaling to some ecological studies: some new results. *Ecology* 67, 919-928.
- Kennet, J.P., Srinivasan, M.S., 1983. *Neogene Planktonic Foraminifera: a Phylogenetic Atlas*, Hutchinson Ross Publishing Co., Stroudsberg, Pennsylvania.
- Kenyon, N.H., Akhmetzhanov, A.M., Wheeler, A.J., Van Weering, T.C.E., de Hass, H., Ivanov, M.K., 2003. Giant carbonate mud mounds in the southern Rockall Trough. *Mar. Geol.* 195, 5-30.
- Kenyon, N., Ivanov, M., Akhmetzhanov, A., Akhmanov, G., 2001. Interdisciplinary Geoscience Research on the North East Atlantic Margin, IOC UNESCO Tech. Ser. 76.
- Kiriakoulakis, K., Bett, B.J., White, M., Wolff, G.A., 2004. Organic biogeochemistry of the Darwin Mounds, a deep-water coral ecosystem, of the NE Atlantic. *Deep Sea Res. I* 51 (12), 1937-1954.
- Kiriakoulakis, K., Fisher, E., Wolff, A.G., Freiwald, A., Grehan, A., Roberts, J.M., 2005. Lipids and nitrogen isotopes of two deep-water corals from the North-East Atlantic: initial results and implications for their nutrition. In: Freiwald, A., Roberts, J.M. (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 715-729.
- Klitgaard, A.B., Tendal, O.S., Westerberg, H., 1997. Mass occurrences of large sponges (Porifera) in Faroe Island (NE Atlantic) shelf and slope areas: characteristics, distribution and possible causes. In: L.E. Hawkins, S. Hutchinson (Eds.), *The responses of marine organisms to their environments. Proceedings of the 30th European Marine Biology Symposium*, Southampton, 129-142.
- Koho, K.A., García, R., De Stigter, H.C., Epping, E., Koning, E., Kouwenhoven, T.J., Van der Zwaan, G.J., 2008. Sedimentary labile organic carbon and pore water redox control on species distribution of benthic foraminifera: A case study from Lisbon-Setúbal Canyon (southern Portugal). *Prog. Oceanogr.* 79, 55-82.
- Kopf, A., Bannert, B., Brückmann, W., Dorschel, B., Foubert, A.T.G., Grevemeyer, I., Gutscher, M.A., Hebbeln, D., Heesemann, B., Hensen, C., Kaul, N.E., Lutz, M., Magalhaes, V.H., Marquardt, M.J., Marti, A.V., Nass, K.S., Neubert, N., Niemann, H., Nuzzo, M., Poort, J.P.D., Rosiak, U.D., Sahling, H., Schneider, J., Somoza, L., Thiebot, E., Wilkop, T.P., 2004. Report and preliminary results of Sonne cruise SO175, Miami-Bremerhaven, 12.11. – 30.12.2003. Berichte Fachbereich Geowissenschaften Universität Bremen 228.
- Koslow J.A., Gowlett-Holmes K., Lowry J.K., O'Hara T., Poore G.C.B., Williams A., 2001. Seamount benthic macrofauna off southern Tasmania: community structure and impacts of trawling. *Mar. Ecol. Prog. Ser.* 213, 111-125.
- Koutsoukos, E.A.M., Leary, P.N., Hart, M.B., 1990. Latest Cenomanian-earliest Turonian low-oxygen tolerant benthonic foraminifera: a case study from the Sergipe basin (N.E. Brazil) and the western Anglo-Paris basin (southern England). *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 77, 145-177.
- Kouwenhoven, T.J., 2000. Survival under stress: benthic foraminiferal patterns and Cenozoic biotic crises. *Geol. Utraiectina* 186, 1-206.

- Kozachenko, M., 2005. Present and past environments of the Belgica mounds (deep-water coral carbonate mounds), Eastern Porcupine Seabight, NE Atlantic. PhD Thesis, Departement of Geology and Environmental Research Institute Cork, Ireland.
- Kruskal, J.B., 1964. Multidimensional scaling by optimizing goodness of fit to a nonmetric hypothesis. *Psychometrika* 29, 1-27.
- Kruskal, J.B., Wish, M., 1978. Multidimensional scaling. Sage Publication, Beverly Hills, California.
- Kucera, M., 2007. Planktonic Foraminifera as Tracers of past Oceanic Environments. In: C. Hillaire-Marcel and A. De Vernal (Eds.), *Developments in Marine Geology*. Elsevier, Amsterdam, 213-262.
- Langer, M.R., Lipps, J.H., 2003. Foraminiferal distribution and diversity, Madang Reef and Lagoon, Papua New Guinea. *Coral Reefs* 22(2), 143-154.
- Le Fèvre, J., 1986. Aspects of the Biology of Frontal Systems. *Adv. Mar. Biol.* 23, 163-299.
- León, R., Somoza, L., Medialdea, T., González, F.J., Díaz-del-Río, V., Fernández-Puga, M.C., Maestro, A., Mata, M.P., 2007. Sea-floor features related to hydrocarbon seeps in deepwater carbonate-mud mounds of the Gulf of Cádiz: from mud flows to carbonate precipitates. *Geo-Mar. Lett.* 27, 237-247.
- Licari, L., Mackensen, A., 2005. Benthic foraminifera off West Africa (1°N to 32°S): Do live assemblages from the topmost sediment reliably record environmental variability? *Mar. Micropaleontol.* 55, 205-233.
- Lindberg, B., Mienert, J., 2005. Sedimentological and geochemical environment of the Fugløy Reef off northern Norway. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 633-650.
- Lindberg, B., Berndt, C., Mienert, J., 2007. The Fugløy Reef at 70°N: acoustic signature, geologic, geomorphologic and oceanographic setting. *J. Int. Earth Sci.*, 96, 201-213.
- Linke, P. and Lutze, G.F., 1993. Microhabitat preferences of benthic foraminifera - a static concept or a dynamic adaption to optimize food acquisition? *Mar. Micropaleontol.* 20, 215-234.
- Liu, Z., Wang, Y., Gallimore, R., Gasse, F., Johnson, T., de Menocal, P., Adkins, J., Notaro, M., Prentice, I.C., Kutzbach, J., Jacob, R., Behling, P., Wang, L., Ong, E., 2007. Simulating the transient evolution and abrupt change of Northern Africa atmosphere-ocean-terrestrial ecosystem in the Holocene. *Quat. Sci. Rev.* 26, 1818– 1837.
- Loeblich, A.R., Tappan, H., 1987. Foraminiferal genera and their classification. Van Nostrand Reinhold Company, New York.
- Loubere, P., 1989. Bioturbation and sedimentation rate control of benthic microfossil taxon abundances in surface sediments: a theoretical approach to the analysis of species microhabitats. *Mar. Micropaleontol.* 14, 317–325.
- Loubere, P., 1991. Deep-sea benthic foraminiferal assemblage response to a surface ocean productivity gradient: a test. *Paleoceanography* 6 (2), 193-204.
- Loubere, P., 1996. The surface ocean productivity and bottom water oxygen signals in deep water benthic foraminiferal assemblages. *Mar. Micropaleontol.* 28, 247-261.
- Loubere, P., 1998. The impact of seasonality on the benthos as reflected in the assemblages of deep-sea foraminifera. *Deep Sea Res. I* 45 (2-3), 409-432.
- Loubere, P., Fariduddin, M., 1999. Quantitative estimation of global patterns of surface ocean biological productivity and its seasonal variation on timescales from centuries to millennia. *Global Biogeochem. Cy.* 13 (1), 115-133.
- Lutze, G.F., Coulbourn, W.T., 1984. Recent benthic foraminifera from the continental margin of northwest Africa: Community structure and distribution. *Mar. Micropaleontol.* 8, 361–401.
- Lutze, G.F., Thiel, H., 1987. *Cibicidoides wuellerstorfi* and *Planulina ariminensis*, elevated epibenthic Foraminifera. *Berichte SFB* 313 6, 17-30.
- Lutze, G.F., Thiel, H., 1989. Epibenthic foraminifera from elevated microhabitats: *Cibicidoides wuellerstorfi* and *Planulina ariminensis*. *J. Foraminifer. Res.* 19 (2), 153-158.
- Mackensen, A., Douglas, R.G., 1989. Down-core distribution of live and dead deep-water benthic foraminifera in box cores from the Weddell Sea and the California continental borderland. *Deep Sea Res. I* 36 (6), 879– 900.
- Mackensen, A., Hald, M., 1988. *Cassidulina teretis* TAPPAN and *C. laevigata* d'ORBIGNY: Their modern and late quaternary distribution in Northern seas. *J. Foraminifer. Res.* 18 (1), 16-24.
- Mackensen, A., Fütterer, D. K., Grobe, H., Schmiedl, G., 1993. Benthic foraminiferal assemblages from the eastern South Atlantic Polar Front region between 35° and 57°S: Distribution, ecology and fossilization potential. *Mar. Micropaleontol.* 22 (1993), 33-69.
- Mackensen, A., Grobe, H., Kuhn, G., Fütterer, D.K., 1990. Benthic foraminiferal assemblages from the eastern Weddell Sea between 68 and 73°S: Distribution, ecology and fossilization potential. *Mar. Micropaleontol.* 16, 241-283.

- Mackensen, A., Schmiedl, G., Harloff, J., Giese, M., 1995. Deep-sea foraminifera in the South Atlantic Ocean: ecology and assemblage generation. *Micropaleontology* 41 (4), 342-358.
- Mackensen, A., Sejrup, H., Jansen, E., 1985. The distribution of living benthic foraminifera on the continental slope and rise off Southwest Norway. *Mar. Micropaleontol.* 9 (7), 275-306.
- Marchant, M., Hebbeln, D., Wefer, G., 1998. Seasonal flux patterns of planktic foraminifera in the Peru-Chile Current. *Deep Sea Res. Part I* 45, 1161-1185.
- Margreth, S., Gennari, G., Rüggeberg, A., Comas, M.C., Pinheiro, L.M., Spezzaferri, S., in review. Development of cold-water coral ecosystems on mud volcanoes in the West Alboran Basin: paleoceanographic implications. *Mar. Geol.*
- Margreth, S., Rüggeberg, A., Spezzaferri, S., 2009. Benthic foraminifera as bioindicator for cold-water coral reef ecosystems along the Irish margin. *Deep Sea Res. Part I* 56, 2216-2234.
- Masson, D.G., Bett, B.J., Billett, D.S.M., Jacobs, C.L., Wheeler, A.J., Wynn, R.B., 2003. The origin of deep-water, coral-topped mounds in the northern Rockall Trough, Northeast Atlantic. *Mar. Geol.* 194, 159-180.
- Mazurenko, L.L., Soloviev, V.A., Belenkaya, I., Ivanov, M.K., Pinheiro, L.M., 2002. Mud volcano gas hydrates in the Gulf of Cadiz. *Terra Nova*, 14, 321-329.
- Messing C.G., Neumann A.C., Lang J.C., 1990. Biozonation of deep-water lithoherms and associated hardgrounds in the Northeastern Straits of Florida. *Palaeos*, 5, 15-33.
- Miao, Q., Thunell, R.C., 1993. Recent deep-sea benthic foraminiferal distributions in the South China and Sulu Seas. *Mar. Micropaleontol.* 22, 1-32.
- Mienis, F., Van Weering, T.C.E., de Haas, H., de Stigter, H., Huvenne, V.A.I., Wheeler, A., 2006. Carbonate mound development at the SW Rockall Trough margin based on high resolution TOBI and seismic recording. *Mar. Geol.* 233 (1-4), 1-19.
- Miller, K.G., Lohmann, G.P., 1982. Environmental distribution of recent benthic foraminifera on the northeast United States continental slope. *Geol. Soc. Am. Bull.* 93, 200-206.
- Millot, C., 2009. Another description of the Mediterranean Sea outflow. *Prog. Oceanogr.* 82 (2), 101-124.
- Mitchelson-Jacob, G., Sundby, S., 2001. Eddies of Vestfjorden, Norway. *Cont. Shelf Res.* 21(16-17), 1901-1918.
- Morel, A., 1991. Light and marine photosynthesis: A spectral model with geochemical and climatological implications. *Prog. Oceanogr.* 26, 263 – 306.
- Morigi, C., Jorissen, F.J., Gervais, A., Guichard, S., Borsetti, A.M., 2001. Benthic foraminiferal faunas in surface sediments off NW Africa: relationship with organic flux to the ocean floor. *J. Foraminifer. Res.* 31 (4), 350-368.
- Mork, M., 1981. Circulation phenomena and frontal dynamics of the Norwegian coastal current. *Philos. T. Roy. Soc. A* 302, 635-647.
- Mortensen, P.B., 2001. Aquarium observations on the deep-water coral *Lophelia pertusa* (L., 1758) (scleractinia) and selected associated invertebrates. *Ophelia*, 54(2): 83-104.
- Mortensen, P.B., Buhl-Mortensen, L., 2005. Morphology and growth of the deep-water gorgonians *Primnoa resedaeformis* and *Paragorgia arborea*. *Mar. Biol.* 147, 755-788.
- Mortensen, P.B., Hovland, M.T., Brattegard, T., Farestveit, R., 1995. Deep water bioherms of the scleractinian coral *Lophelia pertusa* (L.) at 64° N on the Norwegian shelf: structure and associated megafauna. *Sarsia* 80, 145-158.
- Mortensen, P.B., Hovland, M.T., Fosså, J.H., Furevik, D.M., 2001. Distribution, abundance and size of *Lophelia pertusa* coral-reefs in mid-Norway in relation to seabed characteristics. *J. Mar. Biol. Assoc. U.K.* 81, 581-597.
- Mullins, H.T., Newton, C.R., Heath, K., Vanburen, H.M., 1981. Modern deep-water coral mounds north of Little Bahama Bank: criteria for recognition of deep-water coral bioherms in the rock record. *J. Sed. Petr.* 51, 999-1013.
- Mullins, H.T., Thompson, J.B., McDougall, K., Vercoutere, T.L., 1985. Oxygen minimum zone edge effect: evidence from the Central California coastal upwelling system. *Geology* 13, 491-494.
- Murray, J.W., 1971. An Atlas of Recent Foraminiferids. Heinemann Educational Books, London.
- Murray, J.W., 1991. Ecology and Palaeoecology of Benthic Foraminifera. Longman Scientific & Technical, Essex.
- Murray, J.W., 2003. Patterns in the cumulative increase in species from foraminiferal time-series. *Mar. Micropaleontol.* 48, 1-21.
- Murray, J.W., 2006. Ecology and Applications of Benthic Foraminifera. Cambridge University Press, Cambridge.
- Murray, J.W., Weston, J.F., Haddon, C.A. and Powell, A.D.J., 1986. Miocene to recent bottom water masses of the north-east Atlantic: An analysis of benthic foraminifera. In: C.P. Summerhayes and N.J. Shackleton (Eds.), North Atlantic Palaeoceanography. Spec. Pub. Geol. Soc., London, 219-230.

- Natland, M.L., 1933. The temperature- and depth-distribution of some Recent and fossil Foraminifera in the Southern California Region. Bull. Scripps Inst. Oceanogr. 3, 225–230.
- Neulinger, S.C., Järnegren, J., Ludvigsen, M., Lochte, K., Dullo W.C., 2008. Phenotype-specific bacterial communities in the cold-water coral *Lophelia pertusa* (Scleractinia) and their implications for the coral's nutrition, health, and distribution. Appl. Environ. Microbiol. 74, 7272–7285.
- Newton, C.R., Mullins, H.T., Gardulski, A.F., Hine, A.C., Dix, G.R., 1987. Coral mounds on the western Florida Slope: unanswered questions regarding the development of deep-water banks. Palaios 2, 359–367.
- Nomura, R., 1983a. *Cassidulinidae* (foraminiferida) from the uppermost Cenozoic of Japan (part 1). Sci. Rep. Tohoku. Univ. 2 Ser. Geol. 53, 1-101.
- Nomura, R., 1983b. *Cassidulinidae* (foraminiferida) from the uppermost Cenozoic of Japan (part 2). Sci. Rep. Tohoku. Univ. 2 Ser. Geol. 54, 1-93.
- Osterman, L.E., Poore, R.Z., Swarzenski, P.W., Turner, R.E., 2005. Reconstructing a 180 yr record of natural and anthropogenic induced low-oxygen conditions from Louisiana continental shelf sediments. Geology 33 (4), 329-332.
- Paull, C.K., Neumann, A.C., Ende, B.A.A., Ussler, W., Rodriguez, N.M., 2000. Lithiherms on the Florida-Hatteras slope. Mar. Geol. 166, 83-101.
- Pawlowski, J., Bolivar, I., Guard-Maffia, J., Gouy, M., 1994. Phylogenetic position of foraminifera inferred from LSU rRNA gene sequences. Molec. Biol. Evol. 13, 445-450.
- Penn, K., Wu, D.Y., Eisen, J.A., Ward, N., 2006. Characterization of bacterial communities associated with deep-sea corals on Gulf of Alaska Seamounts. Appl. Environm. Microbiol. 72, 1680-1683.
- Pérez-Belzuz, F., Alonso, B., Ercilla, G., 1997. History of mud diapirism and trigger mechanism in the Western Alboran Sea. Tectonophysics 282, 399-422.
- Pérez-Folgado, M., Sierro, F.J., Flores, J.A., Cacho, I., Grimalt, J.O., Zahn, R., Shackleton, N.J., 2003. Western Mediterranean planktonic foraminifera events and millennial climatic variability during the last 70 kyr. Mar. Micropaleontol. 48 (1-2), 49-70.
- Peterson, W.H., Rooth, C.G.H., 1976. Formation and exchange of deep-water in Greenland and Norwegian Seas. Deep Sea Res. Part I 23(4), 273-283.
- Pflum, C.E., Frerichs, W.E., 1976. Gulf of Mexico Deep-Water Foraminifers. Spec. Publ. Cushman Foundation Foram. Res. 14, 125.
- Phleger, F.B., 1960. Ecology and distribution of recent Foraminifera, The Johns Hopkins Press, Baltimore.
- Pinheiro, L.M., Ivanov, M.K., Sautkin, A., Akhmanov, G., Magalhães, V.H., Volkonskaya, A., Monteiro, J.H., Somoza, L., Gardner, J., Hamouni, N., Cunha, M.R., 2003. Mud volcanism in the Gulf of Cadiz: results from the TTR-10 cruise. Mar. Geol. 195, 131–151.
- Pontoppidan, E., 1755. The natural history of Norway, London.
- Probert P.K., McKnight D.G., Grove S.L., 1997. Benthic invertebrate bycatch from a deep-water trawl fishery, Chatham Rise, New Zealand. Aquat. Conserv.: Mar. Freshwat. Ecosyst. 7, 27-40.
- Pujol, C., Vergnaud-Grazzini, C., 1989. Palaeoceanography of the Last Deglaciation in the Alboran Sea (Western Mediterranean): stable isotopes and planktonic foraminiferal records. Mar. Micropaleontol. 15, 153-179.
- Qvale, G., Van Weering, T.C.E., 1985. Relationship of surface sediments and benthic foraminiferal distribution patterns in the Norwegian Channel (northern North Sea). Mar. Micropaleontol. 9, 469-488.
- Raddatz, J., Rüggeberg, A., Margreth, S., Dullo, W.-Chr., in review. Paleoenvironmental reconstruction of deep-water carbonate mound initiation in the Porcupine Seabight, NE Atlantic. Mar. Geol.
- Rathburn, A.E., Levin, L.A., Held, Z.A., Lohmann, K.C., 2000. Benthic foraminifera associated with cold methane seeps on the Northern Californian Margin: ecology and stable isotopic composition. Mar. Micropaleontol. 38, 247-266.
- Ratmeyer, V., shipboard party, 2004. Meteor-Berichte 06-2, Cruise No. 61, Leg 3, Cork – Ponta Delgada. Unpublished report.
- Reed, J.K., 2002. Deep-water *Oculina* coral reefs of Florida: biology, impacts, and management. Hydrobiologica 471, 43-55.
- Remia, A., Taviani, M., 2005. Shallow-buried Pleistocene Madrepora-dominated coral mounds on a muddy continental slope, Tuscan Archipelago, NE Tyrrhenian Sea. Facies 50(3-4), 419-425.
- Reyes, J., Santodomingo, N., Gracia, A., Borrero-Pérez, G., Navas, G., Mejía-Ladino, L.M., Bermudez, A., Benavides, M., 2005. Southern Caribbean azooxanthellate coral communities off Colombia. In: A. Freiwald, J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 309-330.
- Reynolds, L.A., Thunell, R.C., 1986. Seasonal

- production and morphologic variation of *Neogloboquadrina pachyderma* (Ehrenberg) in the Northeast Pacific. *Micropaleontology* 32, 1-18.
- Richer de Forges B., 1993. Deep-sea crabs of the Tasman Seamounts (Crustacea: Decapoda: Brachyura). *Rec. Australian Museum* 45, 11-24.
- Ride, W.D.L., Cogger, H.G., Dupuis, C., Kraus, O., Minelli, A., Thompson, F.C., Tubbs, P.K., 1999. International code of zoological nomenclature. The International Trust for Zoological Nomenclature, London.
- Riehl, S., 2009. Archeaeobotanical evidence for the interrelationship of agricultural decision-making and climate changes in the ancient Near East. *Quat. Int.* 197, 93-114.
- Roberts, J.M., 2005. Reef-aggregating behaviour by symbiotic eunicid polychaetes from cold-water corals: do worms assemble reefs?. *J. Mar. Biol. Assoc. UK* 85, 813-819.
- Roberts, J.M., Wheeler, A.J., Freiwald, A., 2006. Reefs of the deep: the biology and geology of cold-water coral ecosystems. *Science* 312, 543-547.
- Roberts, J.M., Wheeler, A.J., Freiwald, A., Cairns, S., 2009. Cold-water corals. Cambridge University Press, Cambridge.
- Roberts, N., 1989. The Holocene: an environmental history. Blackwell, Oxford.
- Rogers, A.D., 1999. The biology of *Lophelia pertusa* (Linnaeus 1758) and other deep-water reef forming corals and impact from human activities. *Int. Rev. Hydrobiol.* 84, 315-406.
- Rogerson, M., Cacho, I., Jimenez-Espejo, F.J., Reguera, M.I., Sierro, F.J., Martinez-Ruiz, F., Frigola, J., Canals, M., 2008. A dynamic explanation for the origin of the Western Mediterranean organic rich layers. *Geochem. Geophys. Geosyst.* 9, Q07U01, doi:10.1029/2007GC001936.
- Rohling, E.J., De Rijk, S., 1999. The Holocene climate optimum and last glacial maximum in the Mediterranean: the marine oxygen isotopes record. *Mar. Geol.* 153, 57-75.
- Rohling, E.J., Den Dulk, M., Pujol, C., Vergnaud-Grazzini, C., 1995. Abrupt hydrological change in the Alboran sea (western Mediterranean) around 8000 yrs BP. *Deep-Sea Res. I.* 42 (9), 1609-1619.
- Rüggeberg, A., Dorschel, B., Dullo, W.-Chr., Hebbeln, D., 2005. Sedimentary patterns in the vicinity of a carbonate mound in the Hovland Mound Province, northern Porcupine Seabight. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 87-112.
- Rüggeberg, A., Dullo, C., Dorschel, B., Hebbeln, D., 2007. Environmental changes and growth history of Propeller Mound, Porcupine Seabight: Evidence from benthic foraminiferal assemblages. *Int. J. Earth Sci.* 96, 57-72.
- Rüggeberg, A., Flögel, S., Dullo, W.-Chr., Hissmann, K., Freiwald, A., in review. Water mass characteristics and sill dynamics in a polar cold-water coral reef setting at Stjernsund, northern Norway. *Mar. Geol.*
- Rutherford, S., D'Hondt, S., Prell, W., 1999. Environmental controls on the geographic distribution of zooplankton diversity. *Nature* 400(6746), 749-753.
- Sætre, R., 1999. Features of the central Norwegian shelf circulation. *Cont. Shelf Res.* 19(14), 1809-1831.
- Sautter, L.R., Thunell, R.C., 1991. Planktonic foraminiferal response to upwelling and seasonal hydrographic conditions: sediment trap results from San Pedro Basin, southern California Bight. *J. Foraminiferal Res.* 21, 347-363.
- Sautkin, A., Talukder, A.R., Comas, M.C., Soto, J.I., Alekseev, A., 2003. Mud volcanoes in the Alboran Sea: evidence from micropaleontological and geophysical data. *Mar. Geol.* 195, 237-261.
- Sautter, L.R., Thunell, R.C., 1989. Seasonal succession of planktonic foraminifera: results from a four-year series sediment traps experiment in the Northeast Pacific. *J. Foraminifer. Res.* 19 (4), 253-267.
- Schiebel, R., Hemleben, C., 2005. Extant planktic foraminifera: A brief review. *Paläontolog. Zeitsch.* 79, 135-148.
- Schiebel, R., Waniek, J., Bork, M., Hemleben, C., 2001. Planktic foraminiferal production stimulated by chlorophyll redistribution and entrainment of nutrients. *Deep Sea Res. Part I* 48(3), 721-740.
- Schmiedl, G., Mackenesen, A., 1997. Late Quaternary paleoproductivity and deep water circulation in the Eastern South Atlantic Ocean: Evidence from benthic foraminifera. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 130, 43-80.
- Schmiedl, G., Bovée, F.D., Buscail, R., Charrière, B., Hemleben, C., Medernach, L., Picon, P., 2000. Trophic control of benthic foraminiferal abundance and microhabitat in the bathyal Gulf of Lions, western Mediterranean Sea. *Mar. Micropaleontol.* 40, 167-188.
- Schmidt, D.N., Renaud, S., Bollmann, J., Schiebel, R. and Thierstein, H.R., 2004. Size distribution of Holocene planktic foraminifer assemblages: biogeography, ecology and adaptation. *Mar. Micropaleontol.* 50(3-4), 319-338.
- Schönfeld, J., 1997. The impact of the Mediterranean Outflow Water (MOW) on benthic foraminiferal assemblages and surface sediments at the

- southern Portuguese continental margin. Mar. Micropaleontol. 29 (1997), 211-236.
- Schönfeld, J., 2002a. Recent benthic foraminiferal assemblages in deep high-energy environments from the Gulf of Cadiz (Spain). Mar. Micropaleontol. 44, 141-162.
- Schönfeld, J., 2002b. A new benthic foraminiferal proxy for near-bottom current velocities in the Gulf of Cadiz, northeastern Atlantic Ocean. Deep Sea Res. Part I 49, 1853-1875.
- Schönfeld, J., Altenbach, A.V., 2005. Late Glacial to Recent distribution pattern of deep-water Uvigerina species in the north-eastern Atlantic. Mar. Micropaleontol. 57, 1-24.
- Schroeder, W.W., Brooke, S.D., Olson, J.B., Phaneuf, B., McDonough, J.J., Etnoyer, P., 2005. Occurrence of deep-water *Lophelia pertusa* and *Madrepora oculata* in the Gulf of Mexico. In: A. Freiwald, J.M. Roberts (Eds.), Cold-water Corals and Ecosystems. Springer-Verlag, Berlin, Heidelberg, New York, 297-307.
- Schöttner, S., Hoffmann, F., Wild, C., Rapp, H.T., Boetius, A., Ramatte, A., 2009. Inter- and intra-habitat bacterial diversity associated with cold-water corals. ISME J. 3, 756-769.
- Seidenkrantz, M.-S., Kouwenhoven, T.J., Jorissen, F.J., Shackleton, N.J., Van der Zwaan, G.J., 2000. Benthic foraminifera as indicators of changing Mediterranean-Atlantic water exchange in the late Miocene. Mar. Geol. 163, 387-407.
- Sejrup, H.P., Fjærøen, T., Hald, M., Beck, L., Hagen, J., Miljeteig, I., Morvik, I., Norvik, O., 1981. Benthic foraminifera in surface samples from the Norwegian continental margin between 62°N and 65°N. J. Foraminifer. Res. 11, 277-295.
- Sen Gupta, B.K., 1999. Modern foraminifera. Kluwer Academic Publishers, Dordrecht.
- Sen Gupta, B.K., Machain-Castillo, M.L., 1993. Benthic foraminifera in oxygen-poor habitats. Mar. Micropaleontol. 20, 183-201.
- Sen Gupta, B.K., Platon, E., Bernhard, J.M., Aharon, P., 1997. Foraminiferal colonization of hydrocarbon-seep bacterial mats and underlying sediment, Gulf of Mexico slope. J. Foraminif. Res. 27, 292-300.
- Siani, G., Paterne, M., Arnold, M., Bard, E., Métivier, B., Tisnerat, N., Bassinot, F., 2000. Radiocarbon reservoir ages in the Mediterranean Sea and Black Sea. Radiocarbon 42 (2), 271-280.
- Smart, C.W., 2008. Abyssal NE Atlantic benthic foraminifera during the last 15 kyr: Relation to variations in seasonality of productivity. Mar. Micropaleontol. 69 (2), 193-209.
- Smart, C.W., King, S.C., Gooday, A.J., Murray, J.W., Thomas, E., 1994. A benthic foraminiferal proxy of pulsed organic matter paleofluxes. Mar. Micropaleontol. 23, 89-99.
- Somoza, L., Díaz-del-Rio, V., Leon, R., Ivanov, M., Fernandez-Puga, M.C., Gardner, J.M., Hernandez-Molina, F.J., Pinheiro, L.M., Rodero, J., Lobato, A., 2003. Seabed morphology and hydrocarbon seepage in the Gulf of Cádiz mud volcano area: acoustic imagery, multibeam and ultra-high-resolution seismic data. Mar. Geol. 195, 153-176.
- Somoza, L., Ivanov, M.K., Pinheiro, L.M., Maestro, A., Lowrie, A., Vasquez, J.T., Gardner, J., Leon, R., Fernandez-Puga, M.C., 2000. Structural and tectonic control of fluid seeps and mud volcanoes in the Gulf of Cadiz. Proceedings of the TTR-10 Post-Cruise Meeting, Mozenka, Moscow.
- Spezzaferri, S., Coric, S., 2001. Ecology of Karpatian (early Miocene) foraminifers and calcareous nannoplankton from Laa an der Thaya, Lower Austria: A statistical approach. Geol. Carpathica 52 (6), 361-374.
- Spezzaferri, S., Spiegler, D., 2005. Fossil planktonic foraminifera (an overview). Paläontologische Zeitschrift, Stuttgart, 79, 149-166.
- Stetson, T.R., Squires, D.F., 1962. Coral banks occurring in deep water on the Blake Plateau. Am. Mus. Novit. 2114, 1-39.
- Strømgren, T., 1971. Vertical and horizontal distribution of *Lophelia pertusa* (Linné) in Trondheimsfjorden on the West Coast of Norway. Det Kongelige Norske Videnskabers Selskabs Skrifter 6, 1-9.
- Stuiver, M., Reimer, P.J., 1993. Extended C-14 Database and Revised Calib 3.0 C-14 Age Calibration Program. Radiocarbon 35(1), 215-230.
- Suhr, S.B., Pond, D.W., Gooday, A.J., Smith, C.R., 2003. Selective feeding by benthic foraminifera on phydetritus on the western Antarctic Peninsula shelf: evidence from fatty acid biomarker analysis. Mar. Ecol. Prog. Ser. 262, 153-162.
- Svansson, A., 1975. Physical and chemical oceanography in the Skagerrak and Kattegat, 1. Open sea conditions. Fish. Board Swed. Inst. Mar. Res. Rep. 1, 1-88.
- Swezey, C., 2001. Eolian sediment responses to late Quaternary climate changes: temporal and spatial patterns in the Sahara. Palaeogeogr. Palaeoclimatol. Palaeoecol. 167, 119-155.
- Talukder, A.R., Comas, M.C., Soto, J.I., 2003. Pliocene to Recent mud diapirism and related mud volcanoes in the Alboran Sea (Western Mediterranean). In: P. Van Rensbergen, R.R. Hillis, A.J. Maltman, C.K. Morley (Eds.), Subsurface Sediment Mobilization. Geological Society, London, Spec. Publ. 216, 443-459.
- Taviani, M., Freiwald, A., Zibrowius, H., 2005a. Deep

- coral growth in the Mediterranean Sea: an overview. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 137-156.
- Taviani, M., Remia, A., Corselli, C., Freiwald, A., Malinverno, E., Mastrototaro, F., Savini, A., Tursi, A., 2005b. First geo-marine survey of living cold-water Lophelia reefs in the Ionian Sea (Mediterranean basin). *Facies* 50, 409-417.
- Thiede, J., 1975. Distribution of foraminifera in surface waters of a coastal upwelling area. *Nature*, 253, 712-714.
- Thiem, Ø., Ravagnan, E., Fosså, J.H., Bernsten, J., 2006. Food supply mechanisms for cold-water corals along the continental shelf. *J. Mar. Sys.* 60, 207-219.
- Thomas, E., Gooday, A.J., 1996. Cenozoic deep-sea benthic foraminifers: Tracers for changes in oceanic productivity?. *Geology* 24 (4), 355-358.
- Thomas, E., Booth, L., Maslin, M., Shackleton, N.J., 1995. Northeastern Atlantic benthic foraminifera during the last 45,000 years: Changes in productivity seen from the bottom up. *Paleoceanography* 10 (3), 545-562.
- Tucker, M.E., Wright, V.P., 1990. *Carbonate Sedimentology*. Blackwell Science, Oxford.
- Van der Zwaan, G.J., 1982. Paleoecology of Late Miocene Mediterranean foraminifera. *Utr. Micropaleontol. Bull.* 25, 1-202.
- Van der Zwaan, G.J., Duijnstee, I.A.P., Dulk, M.D., Ernst, S.R., Jannink, N.T., Kouwenhoven, T.J., 1999. Benthic foraminifers: proxies or problems? A review of paleoecological concepts. *Earth Sci. Rev.* 46, 213-236.
- Van Rensbergen, P., Depreiter, D., Pannemans, B., Moerkerke, G., Van Rooij, D., Marsset, B., Akhmanov, G., Blinova, V., Ivanov, M., Rachidi, M., Magalhaes, V., Pinheiro, L., Cunha, M., Henriet, J.P., 2005. The El Arraiche mud volcano field at the Moroccan Atlantic slope, Gulf of Cadiz. *Mar. Geol.* 219 (1), 1-17.
- Van Rooij, D., Blamart, D., Richter, T.O., Wheeler, A.J., Kozachenko, M., Henriet, J.-P., 2007a. Quaternary sediment dynamics in the Belgica mound province, Porcupine Seabight: Ice rafting events and contour current processes. *Int. J. Earth Sci.* 96, 121-140.
- Van Rooij, D., De Mol, B., Huvenne, V.A.I., Ivanov, M., Henriet, J.-P., 2003. Seismic evidence of current-controlled sedimentation in the Belgica mound province, upper Porcupine slope, southwest of Ireland. *Mar. Geol.* 195, 31-53.
- Van Rooji, D., Huvenne, V.A.I., Foubert, A., Blamart, D., Staelens, P., Henriet, J.-P., Wheeler, A.J., De Haas, H., 2007b. The lost Enya mounds: causal relationship with sediment drifts and/or pockmarks?. *Int. J. Earth Sci.* 98, 849-863.
- Van Weering, T.C.E., De Haas, H., De Stigter, H.C., Lykke-Anderson, H., Kouvaev, I., 2003. Structure and development of giant carbonate mounds at the SW and SE Rockall Trough margins, NE Atlantic Ocean. *Mar. Geol.* 198, 67-81.
- Van Zeist, W., Woldring, H., Stapert, D., 1975. Late Quaternary vegetation and climate of southwestern Turkey. *Palaeohistoria* 17, 55-142.
- Vickerman, K., 1992. The diversity and ecological significance of Protozoa. *Biodiversity Conserv.* 1, 334-341.
- Warwick, R.M., Clarke, K.R., 1991. A comparison of methodes for analysing changes in benthic community structure. *J. Mar. Biol. Ass. U.K.* 71, 225-244.
- Weaver, P.P.E., Pujol, C., 1988. History of the last deglaciation in the Alboran Sea (Western Mediterranean) and adjacent North Atlantic as revealed by coccolith floras. *Palaeogeogr. Palaeoclimatol. Palaeoecol.* 64, 35-42.
- Weston, J.F., 1985. Comparison between Recent benthic foraminiferal faunas of the Porcupine Seabight and Western Approaches Continental Slope. *J. Micropalaeontol.* 4 (2), 165-183.
- Wheeler, A.J., Beck, T., Thiede, J., Klages, M., Grehan, A., Monteys, F.X., Party, P.A.X.a.S., 2005. Deep-water coral mounds on the Porcupine Bank, Irish Margin: preliminary results from the Polarstern ARK-XIX/3a ROV cruise. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 393-402.
- Wheeler, A.J., Beyer, A., Freiwald, A., De Haas, H., Huvenne, V.A.I., Kozachenko, M., Olu-Le Roy, K., 2007. Morphology and Environment of Deep-water Coral Mounds on the NW European Margin. *Int. J. Earth Sci.* 96, 37-56.
- White, M., Mohn, C., De Stigter, H., Mottram, G., 2005. Deep-water coral development as a function of hydrodynamics and surface productivity around the submarine banks of the Rockall Trough, NE Atlantic. In: A. Freiwald, J.M. Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, New York, 503-514.
- White, M., Roberts, J.M., Weering, T.C.E., 2007. Do bottom intensified diurnal tidal currents shape the alignment of carbonate mounds in NE Atlantic?. *Geo-Mar. Lett.* 27, 391-397.
- Wienberg, C., Beuck, L., Heidkamp, S., Hebbeln, D., Freiwald, A., Pfannkuche, O., Monteys, X., 2007. Franken Mound: facies and biocoenoses

- on a newly-discovered „carbonate mound“ on the western Rockall Bank, NE Atlantic. *Facies* 54 (1), 1-24.
- Wienberg, C., Hebbeln, D., Fink, H.G., Mienis, F., Dorschel, B., Vertino, A., López Correa, M., Freiwald, A., 2009. Scleractinian cold-water corals in the Gulf of Cadiz - First clues about their spatial and temporal distribution. *Deep Sea Res. Part I* 56(10), 1873-1893.
- Wilson R.R., Kaufman R.S. (1987). Seamount biota and biogeography. In: Seamounts, Islands and Atolls. In: B.H. Keating, P. Fryer, R. Batiza, G.W. Backland (Eds.), *Geophysical Monographs*, 43, Washington, 355-377.
- Williams, T., Kano, A., Ferdelman, T., Henriet, J.-P., Abe, K., Andres, M.S., Bierager, M., Browning, E.L., Cragg, B.A., De Mol, B., Foubert, A., Frannk, T.D., Fuwa, Y., Gaillot, P., Gharib, J.J., Gregg, J.M., Huvenne, V.A.I., Léonide, P., Li, X., Mangelsdorf, K., Tanaka, A., Monteys, X., Novosei, I., Sakai, S., Samarkin, V.A., Sasaki, K., Spivack, A.J., Takashima, C., Titschack, J., 2006. Cold-water coral mounds revealed. *EOS*, 87, 525-526.
- Wissak, M., Freiwald, A., Lundälv, T. and Gektidis, M., 2005. The physical niche of the bathyal *Lophelia pertusa* in a non-bathyal setting: environmental controls and palaeoecological implications. In: A., Freiwald, J.M., Roberts (Eds.), *Cold-water Corals and Ecosystems*. Springer-Verlag, Berlin, Heidelberg, 979–1001.
- Woodruff, F., Savin, S.M., Douglas, R.G., 1980. Biological fractionation of oxygen and carbon isotopes by recent benthic foraminifera. *Mar. Micropaleontol.* 5, 3-11.
- Zanchetta, G., Drysdale, R.N., Hellstrom, J.C., Fallick, A.E., Isola, I., Gagan, M.K., Pareschi M.T., 2007. Enhanced rainfall in the Western Mediterranean during deposition of sapropel S1: stalagmite evidence from Corchia cave (Central Italy). *Quat. Sci. Rev.* 26, 279–286.
- Zibrowius, H., 1980. Les Scléractiniaires de la Méditerranée et de l'Atlantique nord-oriental. Mémoires de l' Institut océanographique, Monaco.

SYSTEMATIC DESCRIPTION

to be submitted to Earth-Science Reviews

- Order Foraminiferida Eichwald, 1830
 Suborder Textulariina Delage and Hérouard, 1896
 Superfamily Astrorhizacea Brady, 1881
 Family Astrorhizidae Brady, 1881
 Subfamily Astrorhizinae Brady, 1881
 Genus *Astrorhiza* Sandahl, 1858
 Type species: *Astrorhiza limnicola* Sandahl, 1858, p. 301
 Astrorhiza cf. *catenata* Norman, 1877
 Pl. 1, fig. 1
 Astrorhiza catenata Norman in Norman, 1877, p. 213
 Astrorhiza catenata Norman in Brady, 1879, p. 42, pl. 4, figs. 12-13
 Aschemonella catenata Norman in Jones, 1994, p. 35, pl. 27, fig. 3, pl. 27A, fig. 3
 Family Bathysiphonidae Avnimelech, 1951
 Genus *Bathysiphon* M. Sars, 1872
 Type species: *Bathysiphon filiformis* M. Sars, 1872, p. 251
 Bathysiphon filiformis M. Sars, 1872
 Pl. 1, fig. 2
 Bathysiphon filiformis M. Sars in G. O. Sars, 1872, p. 251, pl. 6, fig. 4
 Bathysiphon filiformis M. Sars in Brady, 1884, p. 248, pl. 26, figs. 15, 17-20
 Bathysiphon filiformis M. Sars in Cushman, 1921, p. 41, pl. 2, fig. 1
 Bathysiphon filiformis M. Sars in Gooday, 1988, p. 98, textfig. 1
 Bathysiphon filiformis M. Sars in Jones, 1994, p. 34, pl. 26, figs 15, 17-20

- Family Rhabdamminidae Brady, 1884
 Subfamily Rhabdammininae Brady, 1884
 Genus *Rhabdammina* M. Sars, 1869
 Type species: *Rhabdammina abyssorum* M. Sars, 1869, in Carpenter, p. 61
 Rhabdammina abyssorum M. Sars, 1869
 Pl. 1, fig. 3
 Rhabdammina abyssorum M. Sars in Carpenter, p. 61
 Rhabdammina abyssorum M. Sars in Brady, 1884, p. 266, pl. 21, figs. 1-8, 10-13
 Rhabdammina abyssorum M. Sars in Höglund, 1947, p. 25, pl. 1, fig. 2
 Rhabdammina abyssorum M. Sars in Jones, 1994, p. 32, pl. 21, figs. 1-8, 10-13
 Family Hippocrepinellidae Loeblich and Tappan, 1984
 Genus *Hippocrepinella* Heron-Allen and Earland, 1932
 Type species: *Hippocrepinella hirudinea* Heron-Allen and Earland, 1932, p. 71
 Hippocrepinella hirudinea Heron-Allen and Earland, 1932
 Pl. 1, fig. 4
 Hippocrepinella hirudinea Heron-Allen and Earland in Heron-Allen and Earland, p. 258, pl. 1, figs. 7-15
 Family Psammosphaeridae Haeckel, 1894
 Subfamily Psammosphaerinae Haeckel, 1894
 Genus *Psammosphaera* Schulze, 1875
 Type species: *Psammosphaera fusca* Schulze, 1875, p. 113

- Psammosphaera fusca* Schulze, 1875
Pl. 5, fig. 5
- Psammosphaera fusca* Schulze in Schulze, 1875, p. 113, pl. 2, fig. 8
- Psammosphaera fusca* Schulze in Brady, 1884, p. 249, pl. 18, figs. 1-8
- Psammosphaera fusca* Schulze in Höglund, 1947, p. 46, pl. 4, figs. 9-14
- Psammosphaera fusca* Schulze in Hofker, 1972, p. 32, pl. 7, figs. 1-3
- Psammosphaera fusca* Schulze in Schröder, 1986, p. 36, pl. 10, figs. 1
- Psammosphaera fusca* Schulze in Jones, 1994, p. 31, pl. 18, figs. 1-8
- Psammosphaera fusca* Schulze var. *testacea* Flint, 1899
Pl. 1, fig. 6
- Psammosphaera fusca* var. *testacea* Flint in Flint, 1899, p. 268, pl. 8, fig. 2
- Family Saccamminidae Brady, 1884
Subfamily Saccamminiae Brady, 1884
- Genus *Lagenammina* Rhumbler, 1911
Type species: *Lagenammina laguncula* Rhumbler, 1911, p. 92, 111
- Lagenammina fusiformis* (Williamson, 1858)
Pl. 1, fig. 7
- Proteonina fusiformis* Williamson in Williamson, 1858, p. 1, pl. 1, fig. 1
- Reophax fusiformis* Williamson in Brady, 1884, p. 290, pl. 30, figs. 7-10, ?11
- Reophax fusiformis* Williamson in Schröder, 1986, p. 44, pl. 15, fig. 9
- Lagenammina fusiformis* Williamson in Cimerman and Langer, 1991, p. 15, pl. 1, fig. 4-5
- Reophax fusiformis* Williamson in Jones, 1994, p. 37, pl. 30, figs. 7-10, ?11
- Lagenammina arenulata* (Williamson, 1858)
Pl. 1, fig. 8
- Reophax diffugiformis* Brady var. *arenulata* Skinner in Skinner, 1961, p. 1239, pl. 30, fig. 5
- Lagenammina arenulata* Skinner in Jones, 1994, p. 37, pl. 30, fig. 5

Genus *Saccammina* Carpenter, 1869
Type species: *Saccammina sphaerica* Brady, 1871, p. 183

Saccammina sphaerica M. Sars, 1872
Pl. 1, fig. 9

- Saccammina sphaerica* M. Sars in G. O. Sars, 1872, p. 250
- Saccammina sphaerica* M. Sars in Brady, 1884, p. 253, pl. 18, figs. 11-15, 17
- Saccammina sphaerica* M. Sars in Cushman, 1918, p. 44, pl. 16, figs 4-5
- Saccammina sphaerica* M. Sars in Höglund, 1947, p. 50, pl. 4, figs 15-17
- Saccammina sphaerica* M. Sars in Loeblich and Tappan, 1964, p. C196, fig. 112.1
- Saccammina sphaerica* M. Sars in Hofker, 1972, p. 44, pl. 12, figs 1-3
- Saccammina sphaerica* M. Sars in Zheng, 1988, p. 33, pl. 4, fig. 12
- Saccammina sphaerica* M. Sars in Jones, 1994, p. 31, pl. 18, figs 11-15, ?17

Superfamily Hippocrepinacea Rhumbler, 1895

Family Hippocrepinidae Rhumbler, 1895
Subfamily Hyperammininae Eimer and Fickert, 1899

Genus *Hyperammina* Brady, 1878
Type species: *Hyperammina elongata* Brady, 1878, p. 433

Hyperammina elongata Brady, 1878
Pl. 1, fig. 10

- Hyperammina elongata* Brady in Brady, 1878, p. 433, pl. 20, fig. 2
- Hyperammina elongata* Brady in Brady, 1884, p. 257, pl. 23, fig. 8
- Hyperammina elongata* Brady in Hofker, 1972, p. 45, pl. 12, figs 4-7
- Hyperammina elongata* Brady in Schröder, 1986, p. 34, pl. 6, fig. 2
- Hyperammina elongata* Brady in Zheng, 1988, p. 28, pl. 4, figs 4-5
- Hyperammina elongata* Brady in Jones, 1994, p. 33, pl. 23, fig. 8

Genus *Saccorrhiza* Eimer and Fickert, 1899
Type species: *Hyperammina ramosa* Brady, 1879, p. 33

Saccorrhiza ramosa (Brady, 1879)

Pl. 1, fig. 11

- Hyperammina ramosa* Brady in Brady, 1879, p. 33, pl. 3, figs. 14-15
Saccorhiza ramosa Brady in Schröder, 1986, p. 35, pl. 7, fig. 1
Saccorhiza ramosa Brady in Loeblich and Tappan, 1987, p. 43, pl. 32, figs. 10-15
Saccorhiza ramosa Brady in Cimerman and Langer, 1991, p. 16, pl. 2, figs. 4-5
Saccorhiza ramosa Brady in Jones, 1994, p. 33, pl. 23, figs 15-19

Family Ammodiscidae Reuss, 1862
Subfamily Ammodiscinae Reuss, 1862

- Genus *Ammodiscus* Reuss, 1862
Type species: *Ammodiscus infimus* L. G. Bornemann, 1874, p. 725

- Ammodiscus incertus* (d'Orbigny, 1839)
Pl. 2, fig. 4

- Operculina incerta* d'Orbigny in d'Orbigny, 1839, p. 49, pl. 6, figs. 16-17
Ammodiscus incertus d'Orbigny in Brady, 1884, p. 330, pl. 38, figs. 1-3
Ammodiscus incertus d'Orbigny in Cushman, 1910, p. 73, figs. 95-96
Ammodiscus incertus d'Orbigny in Schröder, 1986, p. 39, pl. 10, fig. 10
Ammodiscus anguillae Höglund in Jones, p. 43, pl. 38, figs. 1, 3

Subfamily Tolypammininae Cushman, 1928

- Genus *Ammolagena* Eimer and Fickert, 1899
Type species: *Trochammina irregularis* d'Orbigny var. *clavata* Jones and Parker, 1860, p. 304

- Ammolagena clavata* (Jones and Parker, 1860)
Pl. 2, fig. 7

- Trochammina irregularis* d'Orbigny var. *clavata* Jones and Parker in Jones and Parker, 1860, p. 304
Webbina clavata Jones and Parker in Brady, 1884, p. 349, pl. 41, figs. 12-16
Ammolagena clavata Jones and Parker in Cushman, 1921, p. 61, pl. 6, figs 1-4; pl. 10, figs 3-4
Ammolagena clavata Jones and Parker in Saidova, 1961, p. 24, pl. 7, fig. 32
Ammolagena clavata Jones and Parker in Schröder,

- 1986, p. 40, pl. 11, figs. 5-6
Ammolagena clavata Jones and Parker in Jones, 1994, p. 46, pl. 41, figs. 12-16
Ammolagena clavata Jones and Parker in Abu-Zied et al., 2008, p. 51, pl. 1, fig. 1

Subfamily Ammovertellininae Saidova, 1981

- Genus *Glomospira* Rzehak, 1885
Type species: *Trochammina squamata* Jones and Parker var. *gordialis* Jones and Parker, 1860, p. 304

- Glomospira charoides* (Jones and Parker, 1878)
Pl. 2, fig. 5

- Trochammina squamata* Jones and Parker var. *charoides* Jones and Parker in Jones and Parker, 1860, p. 304
Ammodiscus charoides Jones and Parker in Brady, 1884, p. 334, pl. 36, figs. 10-16
Glomospira charoides Jones and Parker in Cushman, 1918, p. 100, pl. 36, figs. 10-15
Glomospira charoides Jones and Parker in Höglund, 1947, p. 129, pl. 3, fig. 11
Glomospira charoides Jones and Parker in Resig, 1981, pl. 9, fig. 8
Repmanina Jones and Parker in Loeblich and Tappan, 1987, p. 52, pl. 39, figs. 24-26
Repmanina Jones and Parker in Cimerman and Langer, 1991, p. 17, pl. 3, figs. 6-9
Usbekistania charoides Jones and Parker in Jones, 1994, p. 43, pl. 38, figs. 10-16
Glomospira charoides Jones and Parker in Abu-Zied et al., 2008, p. 51, pl. 1, figs. 2-3

- Superfamily Hormosinacea Haeckel, 1894
Family Hormosinidae Haeckel, 1894
Subfamily Reophacinae Cushman, 1910

- Genus *Hormosinella* Shchedrina, 1969
Type species: *Reophax distans* Brady, 1881, p. 50

- Hormosinella guttifera* (Brady, 1881)
Pl. 2, fig. 2

- Reophax guttifera* Brady in Brady, 1881, p. 49, pl. 31, figs. 10-15
Hormosinella guttifera Brady in Jones, 1994, p. 38, pl. 32, figs. 18-22
Hormosinella guttifera Brady in Lobegeier and Gupta, 2008, p. 110, pl. 4, fig. 6

Genus *Reophax* de Montfort, 1808
 Type species: *Reophax scorpiurus* de Montfort, 1808,
 p. 331

Reophax agglutinatus Cushman, 1913
 Pl. 2, fig. 1

Reophax agglutinatus Cushman in Cushman, 1913, p.
 637, pl. 79, fig. 6

Reophax agglutinatus Cushman in Jones, 1994, p. 37,
 pl. 30, fig. 13

Reophax scorpiurus de Montfort, 1808
 Pl. 2, fig. 3

Reophax scorpiurus de Montfort in de Montfort,
 1808, p. 330, fig. 130

Reophax scorpiurus de Montfort in Brady, 1884, p.
 291, pl. 30, fig. 15-17

Reophax scorpiurus de Montfort in Cushman, 1921,
 p. 65, pl. 6, fig. 6

Reophax scorpiurus de Montfort in Cimerman and
 Langer, 1991, p. 17, pl. 4, figs. 1-4

Reophax sp. nov. (2) in Jones, 1994, p. 37, pl. 30,
 figs. 15-17

Reophax scorpiurus de Montfort in Majewski and
 Zajaczkowski, p. 112, pl. 1, fig. 10

Superfamily Lituolacea de Bainville, 1827
 Family Haplophragmidae Maync, 1952

Genus *Cribrostomoides* Cushman, 1910
 Type species: *Cribrostomoides bradyi* Cushman,
 1910, p. 108

Cribrostomoides subglobosum (M. Sars, 1868)
 Pl. 2, fig. 6

Lituola subglobosa M. Sars in Carpenter, 1869, p.
 250

Lituola subglobosa M. Sars in G. O. Sars, 1872, p.
 253

Haplophragmoides subglobosa M. Sars in Cushman,
 1910b, p. 105, figs. 162-164

Cribrostomoides bradyi M. Sars in Cushman, 1910b,
 p. 108, fig. 167

Cribrostomoides bradyi M. Sars in Loeblich and
 Tappan, 1987, p. 65, pl. 49, figs 1-3

Cribrostomoides subglobosum M. Sars in Schröder,
 1986, p. 48, pl. 18, figs 15-16

Cribrostomoides subglobosus Cushman in Jones,
 1994, p. 40, pl. 34, figs 8-10

Genus *Haplophragmoides* Cushman, 1910
 Type species: *Nonionina canariensis* d'Orbigny,
 1839, p. 128

Haplophragmoides robertsoni Brady, 1887
 Pl. 3, figs. 1-2

Haplophragmoides robertsoni Brady in Brady, 1887,
 p. 893

Haplophragmoides bradyi Robertson in Robertson,
 1891, p. 388

Haplophragmoides bradyi Robertson in Höglund,
 1947, p. 134, pl. 10, fig. 1; fig. 111

Haplophragmoides bradyi Robertson in Murray,
 1971, p. 25, pl. 5, figs. 1-2

Haplophragmoides bradyi Robertson in Schröder,
 1986, p. 46, pl. 18, fig. 8

Haplophragmoides membranaceum Höglund, 1947
 Pl. 3, fig. 3

Haplophragmoides membranaceum Höglund in
 Höglund, 1947, p. 136, pl. 10, fig. 5

Haplophragmoides membranaceum Höglund in
 Gabel, 1971, p. 30, pl. 3, figs. 25-25

Haplophragmoides membranaceum Höglund in de
 Stigter et al., 1998, p. 45, pl. 1, fig. 8

Genus *Labrospira* Höglund, 1947
 Type species: *Haplophragmium crassimargo*
 Norman, 1892, p. 17

Labrospira jeffreysii (Williamson, 1858)
 Pl. 2, figs. 8-9

Nonionina jeffreysii Williamson in Williamson, 1858,
 p. 34, pl. 3, figs. 72-73

Haplophragmium canariensis d'Orbigny in Brady,
 1884, p. 310, pl. 35, figs 1-3, 5

Cribrostomoids jeffreysii Williamson in Öki, 1989, p.
 71, pl. 1, fig. 14

Veleroninoides jeffreysii Williamson in Jones, 1994,
 p. 41, pl. 35, figs. 1-3, 5

Cribrostomoids jeffreysii Williamson in Murray,
 2003, p. 11, pl. 2, fig. 5

Labrospira jeffreysii Williamson in Majewski and
 Anderson, 2009, p. 138, fig. 6

Family Lituolidae de Bainville, 1827
 Subfamily Ammomarginulininae Podobina, 1978

Genus *Ammobaculites* Cushman, 1910

Type species: *Spirolina agglutinans* d'Orbigny, 1846,
p. 137

Ammobaculites agglutinans (d'Orbigny, 1846)
Pl. 3, fig. 4

Spirolina agglutinans d'Orbigny in d'Orbigny, 1846,
p. 137, pl. 7, figs. 10-12

Haplophragmium agglutinans d'Orbigny in Brady,
1884, p. 301, pl. 32, figs 19-20, 24-26.

Ammobaculites agglutinans d'Orbigny in Schröder,
1986, p. 50, pl. 21, figs 1-4

Ammobaculites agglutinans d'Orbigny in Jones,
1994, p. 39, pl. 32, figs 19-20, 24-26

Superfamily Haplophragmiacea Eimer and Fickert,
1899

Family Ammosphaeroidinidae Cushman, 1927

Subfamily Ammosphaeroidininae Cushman, 1927

Genus *Adercotryma* Loeblich and Tappan, 1952
Type species: *Lituola glomerata* Brady, 1878, p. 433

Adercotryma wrighti Brönnimann and Whittaker,
1987

Pl. 3, fig. 5

Adercotryma wrighti Brönnimann and Whittaker in
Brönnimann and Whittaker, 1987, p. 27, figs. 3B, 7

Superfamily Spirolectamminacea Cushman, 1927

Family Spirolectamminidae Cushman, 1927

Subfamily Spirolectammininae Cushman, 1927

Genus *Spirolectinella* Kisel'man, 1972

Type species: *Spirolecta wrightii* Silvestri, 1903, p.
63

Spirolectinella wrightii (Silvestri, 1903)

Pl. 3, fig. 6

Spirolecta wrightii Silvestri in Silvestri, 1903, p. 59,
figs. 1-6

Textularia sagittula Defrance in Brady, 1884, p. 361,
pl. 42, figs 17-18

Spirorutilus wrightii Silvestri in Banner and Pereira,
1981, p. 104, pl. 6, figs 7-8, 10; pl. 7, figs 1-2, 5

Spirolectinella wrightii Silvestri in Kisel'man, 1972,
p. 135, fig. 1

Spirolectinella wrightii Silvestri in Loeblich and
Tappan, 1987, p. 112, pl. 120, figs 1-10

Spirolectinella wrightii Silvestri in Jones, 1994, p.

47, pl. 42, figs 17-18

Subfamily Vulvulininae Saidova, 1981

Genus *Vulvulina* d'Orbigny, 1826

Type species: *Vulvulina capreolus* d'Orbigny, 1826 =
Nautilus (Ortoceras) pennatula Batsch, 1791

Vulvulina pennatula Batsch, 1791

Pl. 4, fig. 1

Vulvulina pennatula Batsch in Batsch, 1791, p. 3, 5,
pl. 4, fig. 13

Vulvulina pennatula Batsch in Jones, 1994, p. 49, pl.
45, figs. 1-8

Vulvulina pennatula Batsch in Holbourn and
Henderson, 2002, p. 10, pl. 2, figs. 17-18

Superfamily Trochamminacea Schwager, 1877

Family Trochamminidae Schwager, 1877

Subfamily Trochammininae Schwager, 1877

Genus *Portatrocchammina* R. J. Echols, 1971

Type species: *Portatrocchammina eltaninae* R. J.
Echols, 1971

Portatrocchammina antarctica (Parr, 1950)

Pl. 4, fig. 4

Trocchammina antarctica Parr in Parr, 1950, p. 280,
pl. 5, figs. 2-4

Portatrocchammina antarctica Parr in Lobegeier and
Gupta, 2008, p. 110, pl. 4, figs. 14-15

Portatrocchammina antarctica Parr in Majewski and
Anderson, 2009, p. 138, textfig. 3, fig. 7

Genus *Tritaxis* Schubert, 1921

Type species: *Tritaxis fusca* Williamson, 1858, p. 55

Tritaxis fusca (Williamson, 1858)

Pl. 4, fig. 3

Rotalina fusca Williamson in Williamson, 1858, p.
55, pl. 5, figs. 114-115

Tritaxis fusca Williamson in Hedley, Hurdle and
Burdett, 1964, p. 420, fig. 1.

Tritaxis fusca Williamson in Loeblich and Tappan,
1987, p. 122, pl. 128, fig. 1-4

Tritaxis fusca Williamson in Jones, 1994, p. 54, pl.
49, fig. 13

Genus *Trochammina* Parker and Jones, 1859

Type species: *Nautilus inflatus* Montagu, 1808, p. 81

Trochammina labiosa Höglund, 1947
Pl. 4, fig. 2

Trochammina labiosa Höglund in Höglund, 1947, p. 207, pl. 15, fig. 6

Trochammina labiosa Höglund in Gabel, 1971, p. 40, pl. 8, figs. 3-5

Subfamily Polystomammininae Brönnimann and Beurlen, 1977

Genus *Lepidodeuterammina* Brönnimann and Whittaker, 1983

Type species: *Rotalina ochracea* Williamson, 1858, p. 55

Lepidodeuterammina ochracea (Williamson, 1858)
Pl. 4, fig. 5

Rotalina ochracea Williamson in Williamson, 1858, p. 55, pl. 4, figs. 112, pl. 5, figs. 113

Lepidodeuterammina ochracea Williamson in Vazquez Riveiros and Patterson, 2008, p. 10, pl. 4, fig. 3

Superfamily Verneuilinacea Cushman, 1911

Family Verneuilinidae Suleymanov, 1973

Subfamily Verneuilininae Cushman, 1911

Genus *Gaudryina* d'Orbigny, 1839

Type species: *Gaudryina rugosa* d'Orbigny, 1840, p. 44

Gaudryina rufis Wright, 1900

Pl. 4, fig. 6

Gaudryina rufis Wright in Wright, 1900, p. 53, pl. 2, fig. 1

Gaudryina rufis Wright in Gabel, p. 34, pl. 5, figs. 1-2

Gaudryina rufis Wright in Wagener, 1988, p. 126, pl. 21, fig. 15

Connemarella rufis Wright in Cimerman and Langer, p. 23, pl. 8, figs. 1-4

Gaudryina rufis Wright in Murray, 2003, p. 13, pl. 2, figs. 12-13

Gaudryina pseudotrochus (Cushman, 1922)

Pl. 5, fig. 1

Textularia pseudotrochus Cushman in Cushman, 1922, p. 21, pl. 5, fig. 1-3

Textularia pseudotrochus Cushman in Lacroix, p. 12, pl. 5

Textularia pseudotrochus Cushman in Hofker, 1960, p. 238, pl. A, figs. 21-23

Textularia barkeri Hofker in Hofker, 1978, p. 27, pl. 1, fig. 3

Sahulia patelliformis Hofker in Loeblich and Tappan, 1985, p. 203, pl. 14, figs 1-10

Sahulia barkeri Hofker in Jones, 1994, p. 48, pl. 43, figs 15-16, 18-19

Superfamily Textulariacea Ehrenberg, 1838

Family Eggerellidae Cushman, 1937

Subfamily Eggerellinae Cushman, 1937

Genus *Eggerella* Cushman, 1935

Type species: *Verneuilina bradyi* Cushman 1911, p. 54

Eggerella humboldti Todd and Brönnimann, 1957
Pl. 5, fig. 2

Eggerella humboldti Todd and Brönnimann in Todd and Brönnimann, 1957, p. 26, pl. 2, fig. 26

Genus *Eggerelloides* Haynes, 1973

Type species: *Bulimina scabra* Williamson, 1858, p. 65

Eggerelloides scaber (Williamson, 1858)

Pl. 5, fig. 4

Bulimina scabra Williamson in Williamson, 1858, p. 65, pl. 5, figs. 136-137

Eggerelloides scabrus Williamson in Loeblich and Tappan, 1987, p. 170, pl. 189, figs. 5-7

Eggerelloides scabrus Williamson in Cimerman and Langer, 1991, p. 21, pl. 8, fig. 7

Eggerelloides scaber Williamson in Jones, 1994, p. 52, pl. 47, figs. 15-17

Eggerelloides scaber Williamson in Murray, 2003, p. 13, pl. 2, fig. 11

Genus *Karreriella* Cushman, 1933

Type species: *Gaudryina siphonella* Reuss, 1851, p. 78

Karreriella bradyi (Cushman, 1911)

Pl. 5, fig. 3

Gaudryina bradyi Cushman in Cushman, 1911, p. 67,

pl. 107

Gaudryina bradyi Cushman in Cushman, 1921, p. 149, pl. 29, fig. 3
Karreriella bradyi Cushman in Schröder, 1986, p. 55, pl. 22, fig. 8-9
Karreriella bradyi Cushman in Jones, 1994, p. 50, pl. 46, figs. 1-4
Karreriella bradyi Cushman in Loeblich and Tappan, 1994, p. 25, pl. 30, figs 8-16
Karreriella bradyi Cushman in Holbourn and Henderson, 2002, p. 11, pl. 2, fig. 4-5

Family Textulariidae Ehrenberg, 1838

Subfamily Textulariinae, Ehrenberg, 1838

Genus *Bigenerina* d'Orbigny, 1826

Type species: *Bigenerina nodosaria* d'Orbigny, 1826, p. 27

Bigenerina cylindrica Cushman, 1922

Pl. 5, fig. 6

Bigenerina cylindrica Cushman in Cushman, 1922, p. 26, pl. 3, figs. 7-8

Bigenerina cylindrica Cushman in Jones, p. 49, pl. 44, figs. 19-24

Bigenerina nodosaria d'Orbigny, 1826

Pl. 5, fig. 5

Bigenerina nodosaria d'Orbigny in d'Orbigny, 1826, p. 261, pl. 11, figs. 9-12

Bigenerina ? nodosaria d'Orbigny in Cimerman and Langer, p. 21, pl. 9, figs. 1-6

Bigenerina nodosaria d'Orbigny in Jones, p. 49, pl. 44, figs. 14-18

Bigenerina nodosaria d'Orbigny in Murray, 2003, p. 11, pl. 2, fig. 4

Bigenerina nodosaria d'Orbigny in Duchemin et al., 2005, p. 205, pl. 1, fig. 5

Genus *Textularia* Defrance, 1824

Type species: *Textularia sagittula* Defrance in de Bainville, 1824, p. 177

Textularia lateralis Laliker, 1935

Pl. 5, fig. 8

Textularia lateralis Laliker in Laliker, 1935, p. 1, pl. 1, figs 3-5

Textularia lateralis Laliker in Zheng, 1988, p.111, pl. 26, fig. 4

Textularia lateralis Laliker in Loeblich and Tappan,

1994, p. 28, pl. 33, figs. 13-16

Textularia truncata (Höglund, 1947)

Pl. 6, fig. 1

Textularia truncata Höglund in Höglund, 1947, p.

175, pl. 12, figs. 8-9

Textularia truncata Höglund in Gabel, 1971, p. 32, pl. 4, figs. 20-21

Textularia truncata Höglund in Cimerman and Langer, 1991, p. 22, pl. 12, figs. 1-3

Textularia truncata Höglund in Murray, 2003, p. 15, pl. 3, figs. 17-18

Textularia tenuissima Earland, 1933

Pl. 2, fig. 2

Textularia elegans Lacroix in Lacroix, 1931, p. 14

Textularia elegans Lacroix in Lacroix, 1932, p. 8, pl. 4-6

Textularia tenuissima Earland in Earland, 1933, p. 3, pl. 21-30

Textularia tenuissima Earland in Murray, 2003, p. 15, pl. 3, figs. 15-16

Textularia tenuissima Earland in Blais, 1995, p. 92, pl. 2-2, fig. 1

Subfamily Siphotextulariinae, Loeblich and Tappan, 1985

Genus *Siphotextularia* Finlay, 1939

Type species: *Siphotextularia wairoana* Finlay, 1939

Siphotextularia obesa Parr, 1950

Pl. 5, fig. 7

Siphotextularia obesa Parr in Parr, 1950, p. 276, pl. 5, figs. 1

Family Valvulinidae Berthelin, 1880

Subfamily Valvulininae Berthelin, 1880

Genus *Clavulina* d'Orbigny, 1826

Type species: *Clavulina parisiensis* d'Orbigny, 1826, p. 268

Clavulina parisiensis d'Orbigny, 1826

Pl. 6, fig. 3

Clavulina parisiensis d'Orbigny in d'Orbigny, 1826, p. 268

Suborder Spirillinina Hohenegger and Piller, 1975
Family Spirillinidae Reuss and Fritsch, 1861

Genus *Mychostomina* Berthelin, 1881
Type species: *Spirillina vivipara* Ehrenberg var.
revertens Rhumbler, 1906, p. 32

Mychostomina revertens (Rhumbler, 1906)
Pl. 6, fig. 4

Spirillina vivipara Ehrenberg var. *revertens*
Rhumbler in Rhumbler, 1906, p. 32, pl. 2, figs. 8-10
Mychostomina revertens Rhumbler in Jones, 1994, p.
92, pl. 85, fig. 5

Genus *Spirillina* Ehrenberg, 1843
Type species: *Spirillina vivipara* Ehrenberg, 1843

Spirillina vivipara Ehrenberg, 1843
Pl. 6, fig. 5

Spirillina vivipara Ehrenberg in Heron-Allen and
Earland, 1930, p. 178
Spirillina vivipara Ehrenberg in Y. Le Calvez, 1958,
p. 181
Spirillina vivipara Ehrenberg in Barker, 1960, pl. 85,
figs. 1-5
Spirillina vivipara Ehrenberg in Colom, 1974, p. 139,
figs. 23 c, d
Spirillina vivipara Ehrenberg in Loeblich and
Tappan, 1987, p. 304, pl. 318, figs. 4-7
Spirillina vivipara Ehrenberg in Murray, 2003, p. 15,
pl. 4, fig. 1

Family Patellinidae Rhumbler, 1906
Subfamily Patellininae Rhumbler, 1906

Genus *Patellina* Williamson, 1858
Type species: *Patellina corrugata* Williamson, 1858,
p. 46

Patellina corrugata Williamson, 1858
Pl. 6, fig. 6

Patellina corrugata Williamson in Williamson, 1858,
p. 46, pl. 3, figs. 86-89
Patellina corrugata Williamson in Loeblich and
Tappan, 1987, p. 306, pl. 320, figs. 7-14
Patellina corrugata Williamson in Jones, 1994, p. 93,
pl. 86, figs. 1-7
Patellina corrugata Williamson in Cimerman and
Langer, 1991, p. 24, pl. 15, figs. 7-14

Patellina corrugata Williamson in Murray, 2003, p.
24, pl. 9, figs. 6-7

Suborder Miliolina Delage and Hérouard, 1896
Superfamily Cornuspiracea Schultze, 1854
Family Cornuspiridae Schultze, 1854
Subfamily Cornuspirinae Schultze, 1854

Genus *Cornuspira* Schultze, 1854
Type species: *Orbis foliaceus* Philippi, 1844

Cornuspira foliacea (Philippi, 1844)
Pl. 6, fig. 8

Orbis foliaceus Philippi in Philippi, 1844, p. 147, pl.
24, fig. 26
Cornuspira foliacea Philippi in Loeblich and Tappan,
1987, p. 310, pl. 322, figs. 7-8
Cornuspira foliacea Philippi in Cimerman and
Langer, 1991, p. 24, pl. 15, figs. 1-3

Cornuspira involvens (Reuss, 1850)
Pl. 6, fig. 7

Operculina involvens Reuss in Reuss, 1850, p. 370,
pl. 46, fig. 20
Cornuspira involvens Reuss in Cimerman and
Langer, 1991, p. 25, pl. 15, figs. 4-7
Cornuspira involvens Reuss in Jones, 1994, p. 26, pl.
11, figs. 1-3
Cornuspira involvens Reuss in Murray, 2003, p. 15,
pl. 4, fig. 5

Family Hemigordiopsidae A. Nikitina, 1969
Subfamily Hemigordiopsinae A. Nikitina, 1969

Genus *Gordiospira* Heron-Allen and Earland, 1932
Type species: *Gordiospira fragilis* Heron-Allen and
Earland, 1932

Gordiospira elongata Collins, 1958
Pl. 6, fig. 10

Gordiospira elongata Collins in Collins, 1958, p.
347, pl. 1, figs 6-7
Gordiospira elongata Collins in Loeblich and
Tappan, 1994, p. 37, pl. 56, figs. 17-18; pl. 57, figs.
1-4

Gordiospira sp.
Pl. 6, fig. 9

Remarks. Test discoidal, planispiral, proloculus followed by an undivided second chamber, which is irregularly enroled, surface with transverse growth wrinkle.

Superfamily Miliolacea Ehrenberg, 1839
Family Spiroloculinidae Wiesner, 1920

Genus *Spiroloculina* d'Orbigny, 1826
Type species: *Spiroloculina depressa* d'Orbigny, 1826

Spiroloculina dilatata d'Orbigny, 1846
Pl. 7, fig. 2

Spiroloculina dilatata d'Orbigny in d'Orbigny, 1846, p. 271, pl. 16, figs. 16-18

Spiroloculina dilatata d'Orbigny in Wiesner, 1923, p. 35, pl. 4, fig. 26

Spiroloculina dilatata d'Orbigny in Cimerman and Langer, 1991, p. 30, pl. 22, figs. 5-8

Spiroloculina excavata d'Orbigny, 1846
Pl. 7, fig. 3

Spiroloculina excavata d'Orbigny, 1846, p. 271, pl. 16, figs. 19-21

Spiroloculina excavata d'Orbigny in Cimerman and Langer, 1991, p. 30, pl. 23, figs. 1-3

Spiroloculina excavata d'Orbigny in Sgarrella and Moncharmont-Zei, 1993, p. 169, pl. 5, fig. 6

Spiroloculina excavata d'Orbigny in Murray, 1991, p. 17, pl. 4, figs. 13-14

Spiroloculina tenuiseptata Brady, 1884
Pl. 7, fig. 1

Spiroloculina tenuiseptata Brady in Brady, 1884, p. 153, pl. 10, fig. 5

Spiroloculina tenuiseptata Brady in Le Calvez, 1958, p. 162, pl. 1, fig. 7

Spiroloculina tenuiseptata Brady in Cimerman and Langer, 1991, p. 31, pl. 24, figs. 6-9

Family Hauerinidae Schwager, 1876
Subfamily Siphonapertinae Saidova, 1975

Genus *Ammomassilina* Cushman, 1933
Type species: *Massilina alveoliniformis* Millett, 1898, p. 609

Ammomassilina arenaria (Brady, 1884)

Pl. 7, fig. 4

Spiroloculina arenaria Brady in Brady, 1884, p. 153, pl. 8, fig. 12

Subfamily Hauerininae Schwager, 1876

Genus *Cycloforina* Luczkowska, 1972
Type species: *Quinqueloculina contorta* d'Orbigny, 1846, p. 298

Cycloforina laevigata (d'Orbigny, 1839)
Pl. 7, fig. 5

Quinqueloculina laevigata d'Orbigny in d'Orbigny, 1839, p. 301, pl. 3, figs. 31-33

Cycloforina stalkeri (Loeblich and Tappan, 1953)
Pl. 7, fig. 6

Quinqueloculina stalkeri Loeblich and Tappan in Loeblich and Tappan, 1953, p. 40, pl. 5, figs. 5-9
Quinqueloculina stalkeri Loeblich and Tappan in Sgarrella and Moncharmont-Zei, 1993, p. 174, pl. 5, figs. 13-14.

Genus *Quinqueloculina* d'Orbigny, 1826
Type species: *Serpula seminulum* Linné, 1758, p. 786

Quinqueloculina arctica Cushman, 1933
Pl. 8, fig. 1

Quinqueloculina arctica Cushman in Cushman, 1933, p. 2, pl. 1, fig. 3

Quinqueloculina seminula (Linné, 1758)
Pl. 7, fig. 8

Serpula seminula Linné in Linné, 1758, p. 786, pl. 2, figs. 1

Quinqueloculina seminula Linné in Schlumberger, 1893, p. 208, pl. 4, figs. 80-81

Quinqueloculina seminula Linné in Murray, 1971, p. 65, pl. 24, figs. 1-6

Quinqueloculina seminula Linné in Cimerman and Langer, 1991, p. 38, pl. 34, figs. 9-12

Quinqueloculina viennensis Le Calvez and Le Calvez, 1958
Pl. 7, fig. 7

Miliolina cuvieriana d'Orbigny in d'Orbigny, 1839, p. 190, pl. 11, figs. 19-21

Quinqueloculina viennensis Le Calvez and Le Calvez in Le Calvez and Le Calvez, 1958, p. 187, pl. 5, figs. 42, 44, 45

Quinqueloculina viennensis Le Calvez and Le Calvez in Véneç-Peyré, 1984 pl. 4, fig. 1

Quinqueloculina viennensis Le Calvez and Le Calvez in Sgarrella and Moncharmont - Zei, p. 176, pl. 7, fig. 8

Subfamily Miliolinellinae Vella, 1957

Genus *Biloculinella* Wiesner, 1931

Type species: *Biloculina labiata* Schlumberger, 1891, p. 556

Biloculinella depressa (Wiesner, 1923)

Pl. 8, fig. 3

Biloculina labiata Schlumberger var. *depressa* Wiesner in Wiesner, 1923, pp. 89, 90, pl. 18, fig. 263

Biloculina depressa Wiesner in Cimerman and Langer, 1991, p. 39, pl. 36, fig. 11

Biloculina depressa Wiesner in Murray, 2003, p. 15, pl. 4, figs. 2-3

Biloculinella fragilis Le Calvez and Le Calvez, 1958
Pl. 8, fig. 4

Biloculinella fragilis Le Calvez and Le Calvez in Le Calvez and Le Calvez, 1958, pp. 202-203, pl. 16, figs. 182-183

Biloculinella globula (Bornemann, 1855)

Pl. 8, fig. 2

Biloculina globula Bornemann in Bornemann, 1855, p. 349, pl. 19, fig. 3

Biloculinella globula Bornemann in Cimerman and Langer, 1991, p. 40, pl. 36, figs. 1-2

Genus *Miliolinella* Wiesner, 1931

Type species: *Vermiculum subrotundum* Montagu, 1803, p. 1298

Miliolinella subrotunda (Montagu, 1803)

Pl. 8, fig. 5

Vermiculum subrotundum Montagu in Montagu, 1803, p. 521

Miliolinella subrotunda Montagu in Loeblich and Tappan, 1987, p. 340, pl. 350, figs. 1-12

Miliolinella subrotunda Montagu in Cimerman and Langer, 1991, p. 42, pl. 38, figs. 4-9

Miliolinella subrotunda Montagu in Jones, 1994, p. 20, pl. 4, fig. 3

Miliolinella subrotunda Montagu in Murray, 2003, p. 15, pl. 4, fig. 6

Miliolinella subrotunda Montagu in Duchemin et al., 2007, p. 16, pl. 1, fig. 9

Miliolinella elongata Kruit, 1955

Pl. 9, fig. 1

Miliolinella circularis Borneman var. *elongata* Kruit in Kruit, 1955, p. 468, pl. 1, fig. 15

Miliolinella elongata Kruit in Cimerman and Langer, 1991, p. 41, pl. 37, figs. 8

Miliolinella circularis var. *elongata* Kruit in Kruit, Sgarrella and Moncharmont-Zei, 1993, p. 187, pl. 8, fig. 2

Genus *Pyrgo* Defrance, 1824

Type species: *Pyrgo laevis* Defrance, 1824, p. 273

Pyrgo anomala (Schlumberger, 1891)

Pl. 9, fig. 4

Biloculina anomala Schlumberger in Schlumberger, 1891, p. 569, pl. 11, figs. 84-86, pl. 12, fig. 101

Pyrgo anomala Schlumberger in Cimerman and Langer, 1991, p. 44, pl. 41, figs. 3-5

Pyrgo comata (Brady, 1881)

Pl. 9, fig. 3

Biloculina comata Brady in Brady, 1881, p. 45, pl. 3, fig. 9

Biloculina comata Brady in Brady, 1884, p. 144, pl. 3, figs. 9 a, b

Biloculina comata Brady in Schlumberger, 1891, p. 565, pl. 10, figs. 72, 73,

Biloculina comata Brady in Cushman, 1917, p. 81, pl. 34, fig. 1

Biloculina comata Brady in Cimerman and Langer, 1991, p. 44, pl. 41, fig. 9

Biloculina comata Brady in Jones, 1994, p. 19, pl. 3, fig. 9

Pyrgo elongata (d'Orbigny, 1826)

Pl. 10, fig. 5

Biloculina elongata d'Orbigny in d'Orbigny, 1826, p. 298

Pyrgo elongata d'Orbigny in Cimerman and Langer, 1991, p. 44, pl. 41, figs. 6-8

Pyrgo elongata d'Orbigny in Gabel, 1971, p. 38, pl. 7, figs. 27-28

Pyrgo elongata d'Orbigny in Sgarrella and Moncharmont-Zei, 1993, p. 182, pl. 9, fig. 1.

Pyrgo inornata (d'Orbigny, 1846)
Pl. 9, fig. 2

Biloculina inornata d'Orbigny in d'Orbigny, 1846, p. 266, pl. 16, figs. 7-9

Pyrgo lucernula (Schwager, 1866)
Pl. 10, fig. 4

Biloculina lucernula Schwager in Schwager, 1866, p. 202, pl. 4, fig. 17

Pyrgo lucernula Schwager in Cimerman and Langer, p. 45, pl. 41, figs. 10-11

Pyrgo lucernula Schwager in Jones, p. 18, pl. 2, figs. 5-6

Pyrgo murrhina (Schwager, 1866)
Pl. 10, fig. 3

Biloculina murrhina Schwager in Schwager, 1866, p. 203, pl. 4, fig. 15

Pyrgo murrhina Schwager in Mullineaux and Lohmann, 1981, p. 38, pl. 1, fig. 13

Pyrgo murrhina Schwager in Murgese and De Deckker, 2005, p. 34, text-fig. 4, figs. 9-10

Pyrgo murrhina Schwager in Abu-Zied et al., 2008, p. 51, pl. 1, figs. 16-17

Pyrgo sarsi Schlumberger, 1891
Pl. 10, figs. 1-2

Biloculina ringens Lamarck in Brady, 1884, p. 139

Pyrgo sarsi Schlumberger in Schlumberger, 1891, p. 553, pl. 9, figs. 55-59

Pyrgo fornasinii Chapman and Parr in Chapman and Parr, 1935, p. 5

Pyrgo sarsi Schlumberger in Jones, 1994, p. 18, pl. 2, fig. 7

Pyrgo subsphaerica (d'Orbigny, 1840)
Pl. 9, figs. 5

Biloculina subsphaerica d'Orbigny in d'Orbigny, 1839, p. 162, pl. 8, figs. 25-27

Pyrgo williamsoni (Silvestri, 1923)
Pl. 9, figs. 6

Biloculina ringens Lamarck "typica" Williamson in Williamson, 1858, p. 79, pl. 6, 169-170

Biloculina williamsoni Silvestri in Silvestri, 1923, p.

73

Pyrgo williamsoni Silvestri in Haynes, 1973, p. 61, text-fig. 14, nos. 1-3

Pyrgo williamsoni Silvestri in Murray, 2003, p. 17, pl. 4, figs. 7-8

Genus *Triloculina* d'Orbigny, 1826
Type species: *Miliolites trigonula* Lamarck, 1804, p. 351

Triloculina marioni Schlumberger, 1893
Pl. 11, fig. 1

Triloculina marioni Schlumberger in Schlumberger, 1893, p. 62, pl. 1, figs. 38-41

Triloculina marioni Schlumberger in Le Calvez and Le Calvez, 1958, p. 191, pl. 6, figs. 54-56.

Triloculina marioni Schlumberger in Cimerman and Langer, 1991, p. 46, pl. 43, figs. 1-5.

Triloculina tricarinata d'Orbigny, 1826
Pl. 11, fig. 3

Triloculina tricarinata d'Orbigny in d'Orbigny, 1826, p. 299

Triloculina tricarinata d'Orbigny in Bock, 1971, p. 28, pl. 12, figs. 1-2

Triloculina tricarinata d'Orbigny in Cimerman and Langer, 1991, p. 46, pl. 44, figs. 3-4

Triloculina tricarinata d'Orbigny in Sgarrella and Moncharmont - Zei, 1993, p. 187, pl. 9, figs. 14-15

Triloculina tricarinata d'Orbigny in Hottinger et al, 1993, p. 65, pl. 68, figs. 7-12

Triloculina trigonula (Lamarck, 1804)
Pl. 11, fig. 2

Miliolites trigonula Lamarck in Lamarck, 1804, p. 351, pl. 17, fig. 4

Triloculina trigonula Lamarck in Cushman, 1917, p. 65, pl. 25, fig. 3

Triloculina trigonula Lamarck in Whittaker and Hodgkinson, 1979, p. 34, pl. 3, fig. 8

Triloculina trigonula Lamarck in Zheng, 1988, p. 242, pl. 19, fig. 3; pl. 23, fig. 9; pl. 33, fig. 5; text-fig. 59

Triloculina trigonula Lamarck in Jones, 1994, p. 20, pl. 3, figs. 15-16

Triloculina trigonula Lamarck in Hayward et al., 1999, p. 106, pl. 5, figs 31-32

Subfamily Sigmoilinitinae Luczkowska, 1974

- Genus *Sigmoinella* Saidova, 1975
 Type species: *Sigmoinella borealis* Saidova, 1975, p. 157
- Sigmoinella borealis* Saidova, 1975
 Pl. 11, fig. 4
- Sigmoinella borealis* Saidova in Saidova, 1975, p. 158m pl. 45, fig. 2
- Sigmoinella borealis* Saidova in Loeblich and Tappan, 1987, p. 349, pl. 356, figs. 19-20
- Subfamily Sigmoilopsinae Vella, 1957
- Genus *Sigmoilopsis* Finlay, 1947
 Type species: *Sigmoilina schlumbergeri* Silvestri, 1904, p. 267
- Sigmoilopsis schlumbergeri* (Silvestri, 1904)
 Pl. 11, fig. 5
- Sigmoilina schlumbergeri* Silvestri in Silvestri, 1904, pp. 267, 269
- Sigmoilopsis schlumbergeri* Silvestri in Cimerman and Langer, 1991, p. 48, pl. 46, figs. 10-14
- Sigmoilopsis schlumbergeri* Silvestri in Loeblich and Tappan, 1987, p. 350, pl. 356, figs. 8-13
- Sigmoilina schlumbergeri* Silvestri in Kihle and Løfaldli, 1975
- Sigmoilopsis woodi* Atkinson, 1968
 Pl. 11, fig. 6
- Sigmoilopsis woodi* Atkinson in Atkinson, 1968, p. 161, pl. 18, fig. 4
- Suborder Lagenina Delage and Hérouard, 1896
 Superfamily Nodosariacea Ehrenberg, 1838
 Family Nodosariidae Ehrenberg, 1838
 Subfamily Nodosariinae Ehrenberg, 1838
- Genus *Dentalina* Risso, 1826
 Type species: *Nodosaria cuvieri* d'Orbigny, 1826, p. 255
- Dentalina cuvieri* (d'Orbigny, 1826)
 Pl. 12, fig. 1
- Nodosaria cuvieri* d'Orbigny in d'Orbigny, 1826, p. 255, pl. 9, fig. 57
- Dentalina lamarcki* Neugeboren, 1856

Pl. 12, fig. 2

Dentalina lamarcki Neugeboren in Neugeboren, 1856, p. 91, pl. 4, fig. 16Genus *Grigelis* Mikhalevich, 1981
 Type species: *Nodosaria pyrula* d'Orbigny, 1826, p. 253*Grigelis orectus* (Loeblich and Tappan, 1994)
 Pl. 12, fig. 3*Nodosaria pyrula* d'Orbigny in Schwager, p. 217, pl. 5, fig. 38*Nodosaria pyrula* d'Orbigny in Brady, 1884, p. 497, pl. 62, figs 10-12*Nodosaria pyrula* d'Orbigny in Cushman, 1921, p. 187, pl. 33, figs 3-5*Dentalina guttifera* d'Orbigny in Barker, 1960, p. 130, pl. 62, figs 10-12*Grigelis guttifera* d'Orbigny in Loeblich and Tappan, 1987, p. 396, pl. 441, figs 2-3*Grigelis* sp. nov. Jones, 1994, p. 73, pl. 62, figs 10-12*Grigelis orectus* Loeblich and Tappan in Loeblich and Tappan, 1994, p. 64, pl. 115, fig. 22Genus *Laevidentalina* Loeblich and Tappan, 1986
 Type species: *Laevidentalina aphelis* Loeblich and Tappan, 1986, p. 242*Laevidentalina sidebottomi* (Cushman, 1933)
 Pl. 12, fig. 4*Dentalina sidebottomi* Cushman in Cushman, 1933, p. 12, pl. 3, fig. 4Family Vaginulinidae Reuss, 1860
 Subfamily Lenticulininae Chapman, Parr, and Collins, 1934Genus *Lenticulina* Lamarck, 1804
 Type species: *Lenticulites rotulatus* Lamarck, 1823, p. 153*Lenticulina calcar* (Linné, 1758)
 Pl. 12, fig. 5*Nautilus calcar* Linné in Linné, 1758, p. 709, pl. 19, figs. b-c*Lenticulina calcar* Linné in Cimerman and Langer, 1991, p. 51, pl. 53, figs. 1-4*Lenticulina gibba* (d'Orbigny, 1826)
 Pl. 12, fig. 7

- Cristellaria gibba* d'Orbigny in d'Orbigny, 1826, p. 292, no. 17
Lenticulina gibba d'Orbigny in Kihle and Løfaldli, 1975
Lenticulina gibba d'Orbigny in Cimerman and Langer, 1991, p. 51, pl. 53, figs. 7-11
Lenticulina gibba d'Orbigny in Jones, 1994, p. 81, pl. 69, figs. 8-9
- Lenticulina inornata* (d'Orbigny, 1846)
 Pl. 12, fig. 9
- Robulina inornata* d'Orbigny in d'Orbigny, 1846, p. 102, pl. 4, figs. 25-26
- Lenticulina orbicularis* (d'Orbigny, 1846)
 Pl. 12, fig. 8
- Robulina orbicularis* d'Orbigny in d'Orbigny, 1826, p. 288, pl. 15, figs. 8-9
Lenticulina orbicularis d'Orbigny in Cimerman and Langer, 1991, p. 51, pl. 53, fig. 12
Lenticulina orbicularis d'Orbigny in Jones, 1994, p. 81, pl. 69, fig. 17
Lenticulina orbicularis d'Orbigny in Kihle and Løfaldli, 1975
- Lenticulina vortex* (Fichtel and Moll, 1798)
 Pl. 12, fig. 6
- Nautilus vortex* Fichtel and Moll in Fichtel and Moll, 1798, p. 33, pl. 2, figs. d-i
Lenticulina vortex Fichtel and Moll in Jones, 1994, p. 81, pl. 69, figs. 14-16
- Genus *Neolenticulina* McCulloch, 1977
 Type species: *Neolenticulina chathamensis* McCulloch, 1977, p. 8
- Neolenticulina peregrina* (Schwager, 1866)
 Pl. 12, fig. 10
- Cristellaria peregrina* Schwager, 1866, p. 245, pl. 7, figs. 89
Cristellaria variabilis Reuss in Brady, 1884 (not Reuss, 1850), p. 541, pl. 68, figs 11-16.
Lenticulina peregrina Schwager in Cushman and McCulloch, 1950, p. 302, pl. 39, fig. 5.
Dimorphina peregrina Schwager in Hofker, 1978, p. 37, pl. 3, figs 3-4, 7-8.
Neolenticulina variabilis Reuss in Jones, 1994, p. 80, pl. 68, figs 11-16.
Neolenticulina peregrina Schwager in Loeblich and

- Tappan, 1987, p. 406, pl. 447, figs 9-12, 16.
- Genus *Saracenaria* Defrance, 1824
 Type species: *Saracenaria italicica* Defrance, 1824, in de Blainville p. 176
- Saracenaria caribbeana* Hofker, 1976
 Pl. 13, fig. 1
- Cristellaria italicica* Defrance in Brady, 1884, p. 22, pl. 68, fig. 17
Saracenaria caribbeana Hofker in Hofker, 1976, p. 191, pl. 65
Saracenaria caribbeana Hofker in Jones, 1994, p. 80, pl. 68, fig. 17
- Subfamily Marginulininae Wedekind, 1937
- Genus *Amphicoryna* Schlumberger, 1881
 Type species: *Nautilus scalaris* Batsch, 1791, p. 1, 4
- Amphicoryna scalaris* (Batsch, 1791)
 Pl. 13, figs. 2-3
- Nautilus (Ortoceras) scalaris*, Batsch in Batsch, 1791, p. 91, pl. 2, figs. 4 a-b
Nodosaria scalaris Batsch in Cushman, 1921, p. 199, pl. 35, fig. 6
Amphicoryna scalaris Batsch in Barker, 1960, pl. 63, figs. 28-31
Amphicoryna scalaris Batsch in Cimerman and Langer, 1991, p. 52, pl. 54, figs. 1-9
Amphicoryna scalaris Batsch in Jones, 1994, p. 75, pl. 63, figs 28-31
- Genus *Astacolus* de Montfort, 1908
 Type species: *Nautilus crepidula* Fichtel and Moll, 1798, p. 64
- Astacolus beerae* Brenner and McMillan, 1976
 Pl. 13, fig. 4
- Astacolus beerae* Brenner and McMillan in McLachlan, Brenner, and McMillan, 1976, p. 351, pl. 11, figs. 13-18
- Subfamily Vaginulininae Reuss, 1860
- Genus *Planularia* Defrance, 1826
 Type species: *Peneroplis auris* Defrance in de Blainville, 1824, p. 178

- Planularia perculata* McCulloch, 1977
Pl. 13, fig. 5
- Planularia perculata* McCulloch, 1977, p. 10, pl. 96,
fig. 14
- Planularia costata* (d'Orbigny, 1902)
Pl. 13, fig. 7
- Robulina costata* d'Orbigny in d'Orbigny, 1902, p.
43, pl. 44, fig. 3
- Family Lagenidae Reuss, 1862
- Genus *Lagena* Walker and Jacob, 1798
Type species: *Serpula (Lagena) sulcata* Walker and
Jacob, in Kanmacher, 1798, p. 634
- Lagena substriata* Williamson, 1848
Pl. 13, fig. 6
- Lagena substriata* Williamson in Williamson, 1848,
p. 15, pl. 2, fig. 12
- Lagena substriata* Williamson in Murray, 2003, p. 17,
pl. 5, fig. 7
- Lagena substriata* Williamson in Jones, 1994, p. 64,
pl. 57, fig. 19
- Lagena meridionalis* Wiesner, 1931
Pl. 13, fig. 8
- Lagena gracilis* Williamson var. *meridionalis*
Wiesner in Wiesner, 1931, p. 117, pl. 18, fig. 211
- Lagena meridionalis* Williamson in Jones, 1994, p.
66, pl. 58, fig. 19
- Lagena meridionalis* Williamson in Vazquez Riveiros
and Patterson, 2008, p. 16, pl. 6, fig. 4
- Lagena semilineata* var. *spinigera* Earland, 1934
Pl. 13, fig. 9
- Lagena semilineata* Wright var. *spinigera* Earland in
Earland, 1934, p. 173, pl. 7, fig. 21
- Lagena semilineata* Wright var. *spinigera* Earland in
Jones, 1994, p. 65, pl. 58, figs. 4?, 17
- Lagena squamosoalata* Brady, 1881
Pl. 13, fig. 11
- Lagena squamosoalata* Brady in Brady, 1881, p. 61,
pl. 60, fig. 23
- Lagena squamosoalata* Brady in Jones, 1994, p. 70,
- pl. 60, fig. 23
- Lagena trigonolaevigata* Balkwill and Millett, 1884
Pl. 13, fig. 12
- Lagena trigonolaevigata* Balkwill and Millett in
Balwill and Millett, 1884, p. 86, pl. 3, fig. 4
- Genus *Pygmaeoseistron* Patterson and Richardson,
1987
Type species: *Lagena hispidula* Cushman, 1913, p.
14
- Pygmaeoseistron laevis ovalis* (Williamson, 1848)
Pl. 13, fig. 10
- Serpula laevis ovalis* Walker and Boys in Walker and
Boys, 1784, p. 3, pl. 1, fig. 9
- Vermiculum laeve* Montagu in Montagu, 1803, p.
524, pl. 1, fig. 9
- Genus *Hyalinonetrion* Patterson and Richardson,
1987
Type species: *Hyalinonetrion sahulense* Patterson
and Richardson, 1987
- Hyalinonetrion gracillimum* (Costa, 1856)
Pl. 14, fig. 1
- Amphorina gracilis* Costa in Costa, 1856, p. 121, pl.
11, fig. 11a
- Lagena gracillima* Costa in Feyling-Hanssen, 1971,
p. 206, pl. 4, fig. 1
- Lagena gracillima* Costa in Gabel, 1971, p. 45, pl.
10, figs. 1, 2
- Hyalinonetrion gracillimum* Costa in Cimerman and
Langer, , p. 52, pl. 55, figs. 1-2
- Family Polymorphinidae d'Orbigny, 1839
Subfamily Polymorphininae d'Orbigny, 1839
- Genus *Globulina* d'Orbigny, 1839
Type species: *Polymorpha gibba* d'Orbigny, 1826,
p. 266
- Globulina aequalis* d'Orbigny, 1846
Pl. 14, fig. 2
- Globulina aequalis* d'Orbigny in d'Orbigny, 1846, p.
227, pl. 13, figs. 11, 12
- Genus *Pseudopolymorpha* Cushman and Ozawa,
1928

- Type species: *Pseudopolymorphina hanzawai*
Cushman and Ozawa, 1928, p. 15
- Pseudopolymorphina* sp.
Pl. 14, fig. 3
- Genus *Pyrulina* d'Orbigny, 1839
Type species: *Polymorphina gutta* d'Orbigny, 1926,
p. 267, 310
- Pyrulina cylindroids* (Roemer, 1930)
Pl. 14, fig. 4
- Polymorphina cylindroids* Roemer in Roemer, p.
385, pl. 3, fig 26
- Pyrulina cylindroids* Roemer in Feyling-Hannsen, p.
219, pl. 5, figs. 10, 11
- Family Ellipsolagenidae A. Silvestri, 1923
Subfamily Oolininae Loeblich and Tappan, 1961
- Genus *Favulina* Patterson and Richardson, 1987
Type species: *Entosolenia squamosa* Montagu var.
hexagona Williamson, 1848, p. 20
- Favulina squamosa* (Montagu, 1803)
Pl. 14, fig. 5
- Vermiculum squamosum* Montagu in Montagu, 1803,
p. 526, pl. 14, fig. 2
- Oolina squamosa* Montagu in Jones, 1994, p. 66, pl.
58, fig. 32
- Oolina squamosa* Montagu in Kihle and Løfaldli,
1975
- Favulina melo* (d'Orbigny, 1839)
Pl. 14, fig. 6
- Oolina squamosa* Montagu in Jones, 1994, p. 66, pl.
58, fig. 28-30
- Oolina melo* d'Orbigny in Kihle and Løfaldli, 1975
- Favulina melo* d'Orbigny in Vazquez Riveiros and
Patterson, 2008, p. 17, pl. 7, fig. 2
- Favulina hexagona* (Williamson, 1858)
Pl. 14, fig. 7
- Entosolenia squamosa* Montagu var. *hexagona*
Williamson in Williamson, 1848, p. 20, pl. 2, fig. 23
- Oolina hexagona* Williamson in Jones, 1994, p. 66,
pl. 58, fig. 33
- Favulina hexagona* Williamson in Cimerman and
Langer, 1991, p. 55, pl. 58, figs. 8-9
- Genus *Homalohedra* Patterson and Richardson, 1987
Type species: *Lagena guntheri* Earland, 1987, p. 151
- Homalohedra williamsoni* (Alcock, 1865)
Pl. 14, fig. 8
- Entosolenia williamsoni* Alcock in Alcock, 1865, p.
195
- Oolina williamsoni* Alcock in Feyling-Hannsen, p.
227, pl. 18, gigs. 1,2
- Oolina williamsoni* Alcock in Murray, 2003, p. 17, pl.
5, fig. 10
- Homalohedra apiopleura* (Loeblich and Tappan,
1953)
Pl. 14, fig. 9
- Lagena apiopleura* Loeblich and Tappan in Loeblich
and Tappan, 1953, p. 59, pl. 10, figs. 14, 15
- Oolina apiopleura* Loeblich and Tappan in Jones,
1994, p. 65, pl. 57, fig. 32
- Homalohedra apiopleura* Loeblich and Tappan in
Vazquez Riveiros and Patterson, 2008, p. 17, pl. 7,
fig. 3
- Homalohedra borealis* (Loeblich and Tappan, 1954)
Pl. 15, fig. 1
- Entosolenia costata* Williamson in Williamson, 1858,
p. 9, pl. 1, fig. 18
- Oolina costata* Williamson in Loeblich and Tappan,
1953, p. 68, pl. 13, figs. 4-6
- Oolina borealis* Loeblich and Tappan in Loeblich and
Tappan, 1954, p. 384
- Homalohedra borealis* Loeblich and Tappan in
Vazquez Riveiros and Patterson, 2008, p. 17, pl. 7,
fig. 4
- Homalohedra eucostata* (McCulloch, 1977)
Pl. 15, fig. 2
- Oolina eucostata* McCulloch in McCulloch, 1977,
pp. 78, 79, pl. 55, fig. 7
- Genus *Oolina* d'Orbigny, 1839
Type species: *Oolina laevigata* d'Orbigny, 1839, p.
19
- Oolina lineata* subsp. *communis* McCulloch, 1977
Pl. 15, fig. 3
- Oolina lineata* Williamson subsp. *communis*
McCulloch in McCulloch, 1977, p. 81, pl. 55, fig. 18

- Oolina ampulladistoma* (Jones, 1874)
Pl. 15, fig. 4
- Lagena vulgaris* Williamson var. *ampulla-distoma*
Jones in Jones, 1874, p. 63, pl. 19, fig. 5
- Oolina globosa* (Montagu, 1803)
Pl. 15, fig. 5
- Vermiculum globosum* Montagu in Montagu, 1803, p. 3, pl. 1, fig. 8
- Oolina globosa* Montagu in Jones, 1994, p. 61, pl. 56, figs. 1-3
- Oolina laevigata* d'Orbigny, 1839
Pl. 15, fig. 6
- Oolina laevigata* d'Orbigny in d'Orbigny, 1839, p. 19, pl. 5, fig. 3
- Genus *Fissurina* Reuss, 1850
Type species: *Fissurina laevigata* Reuss, 1850, p. 366
- Fissurina agassizi* Todd and Bronnimann, 1957
Pl. 15, fig. 7
- Fissurina agassizi* Todd and Bronnimann in Todd and Bronnimann, 1957, p. 36, pl. 9, fig. 14
- Fissurina annectens* (Burrows and Holland, 1895)
Pl. 16, fig. 4
- Lagena annectens* Burrows and Holland in Jones, 1895, p. 203, pl. 7, fig. 11
- Fissurina annectens* Kihle and Løfaldli, 1975
- Fissurina circularis* Todd, 1954
Pl. 15, fig. 8
- Fissurina circularis* Todd in Todd, 1954, p. 351, pl. 87, fig. 27
- Fissurina lucida* (Williamson, 1848)
Pl. 16, fig. 1
- Entosolenia marginata* Monatgu var. *lucida*
Williamson in Williamson, 1848, p. 17, pl. 2, fig. 17
- Fissurina lucida* Williamson in Kihle and Løfaldli, 1975
- Fissurina lucida* Williamson in Vazquez Riveiros and Patterson, 2008, p. 19, pl. 8, Fig. 2
- Fissurina eburnea* Buchner, 1940
- Pl. 16, fig. 2
- Fissurina eburnea* Buchner in Buchner, 1940, p. 458, pl. 9, figs. 146, 147
- Fissurina eburnea* Buchner in Vazquez Riveiros and Patterson, p. 19, pl. 8, figs. 1, 8
- Fissurina kerguelensis* Parr, 1950
Pl. 16, fig. 3
- Lagena staphyllearia* Schwager in Schwager, 1884, p. 474, pl. 59, figs. 8-11
- Fissurina kerguelensis* Parr in Parr, 1950, p. 305, pl. 8, fig. 7
- Fissurina crassiporosa* McCulloch, 1977
Pl. 16, fig. 5
- Fissurina crassiporosa* McCulloch in McCulloch, 1977, p. 98, pl. 56, figs. 15, 16, 22
- Fissurina longpointensis* McCulloch, 1977
Pl. 16, fig. 6
- Fissurina longpointensis* McCulloch in McCulloch, 1977, p. 114, pl. 59, fig. 9
- Fissurina pseudoorbigniana* (Buchner, 1940)
Pl. 17, fig. 3
- Lagena pseudoorbigniana* Buchner in Buchner, 1940, p. 460, pl. 10, figs. 157-160
- Fissurina dublini* McCulloch, 1977
Pl. 17, fig. 4
- Fissurina dublini* McCulloch in McCulloch, 1977, p. 102, pl. 62, fig. 8
- Fissurina derogata* McCulloch, 1977
Pl. 17, fig. 5
- Fissurina derogata* McCulloch in McCulloch, 1977, p. 101, pl. 65, fig. 6
- Fissurina lacunata* (Burrows and Holland, 1895)
Pl. 17, fig. 6
- Lagena castrensis* Brady in Brady, 1884, p. 485, pl. 1, figs. 1-3
- Lagena lacunata* Burrows and Holland in T.R. Jones, 1895, p. 205, pl. 7, fig. 12
- Fissurina nucelloides* (Buchner, 1940)

Pl. 17, fig. 7

Lagena nucelloides Buchner in Buchner, 1940, p. 518, pl. 22, figs. 476-477

Fissurina pseudolucida Zheng, 1979

Pl. 17, fig. 8

Fissurina pseudolucida Zheng in Zheng, 1979, p. 215, pl. 13, figs. 8-9

Genus *Palliolatella* Patterson and Richardson, 1987

Type species: *Palliolatella avita* Patterson and Richardson, 1987, p. 219

Palliolatella semimarginata (Reuss, 1870)

Pl. 18, fig. 1

Lagena marginata Montagu var. *semimarginata* Reuss in Reuss, 1870, p. 468, pl. 4, figs. 4-6, 10-12

Fissurina semimarginata Reuss in Jones, 1994, p. 68, pl. 59, figs. 17, 19

Subfamily Parafissurininae R. W. Jones, 1984

Genus *Parafissurina* Parr, 1947

Type species: *Lagena ventricosa* A. Silvestri, 1904, p. 10

Parafissurina basispinata McCulloch, 1977

Pl. 17, fig. 1

Parafissurina basispinata McCulloch in McCulloch, 1977, p. 139, pl. 72, figs. 1-3

Parafissurina marginata (Walker and Boys, 1803)

Pl. 17, fig. 2

Serpula (Lagena) marginata Walker and Boys, 1784, p. 2, pl. 1, fig. 7

Vermiculum marginatum Montagu in Montagu, 1803, p. 524, pl. 1, fig. 7

Fissurina marginata Murray in Murray, 2003, p. 17, pl. 5, figs. 3-4

Parafissurina felsinea (Fornasini, 1894)

Pl. 18, fig. 4

Lagena felsinea Fornasini in Fornasini, 1894, p. 1

Parafissurina felsinea Fornasini in Jones, 1994, p. 61, pl. 56, fig. 4

Parafissurina lateralis (Cushman, 1913)

Pl. 18, fig. 2

Lagena lateralis Cushman in Cushman, 1913, p. 9, pl. 1, fig. 1

Parafissurina lateralis Cushman in Jones, 1994, p. 62, pl. 56, fig. 17, 18

Parafissurina robusta (Zheng, 1979)

Pl. 18, fig. 6

Fissurina robusta Zheng in Zheng, 1979, pp. 215-216, pl. 13, fig. 11

Suborder Robertinina Loeblich and Tappan, 1984

Superfamily Ceratobuliminacea Cushman, 1927

Family Epistominidae Wedekind, 1937

Subfamily Epistomininae Wedekind, 1937

Genus *Hoeglundina* Brotzen, 1948

Type species: *Rotalia elegans* d'Orbigny, 1826, p. 272

Hoeglundina elegans (d'Orbigny, 1826)

Pl. 18, fig. 3

Rotalia (Turbulinina) elegans d'Orbigny in d'Orbigny, 1826, p. 276

Hoeglundina elegans d'Orbigny in Hermelin and Scott, 1985, p. 210, pl. 6, fig. 6-7

Hoeglundina elegans d'Orbigny in Cimerman and Langer, 1991, p. 56, pl. 59, figs. 10-12

Høglundina elegans d'Orbigny in Kihle and Løfaldli, 1975

Superfamily Conorboidacea Thalmann, 1952

Family Robertinidae Reuss, 1850

Subfamily Robertininae Reuss, 1850

Genus *Robertinoides* Höglund, 1947

Type species: *Bulimina normani* Goës, 1894, p. 47

Robertinoides bradyi (Cushman and Parker, 1936)

Pl. 18, fig. 7

Bulimina subteres Brady in Brady, 1881, p. 55

Bulimina subteres Brady in Brady, 1884, p. 403, pl. 50, fig. 18

Robertina bradyi Cushman and Parker in Cushman and Parker, 1936, p. 99, pl. 16, fig. 9

Robertinoides bradyi Cushman and Parker in Jones, 1994, p. 55, pl. 50, fig. 18

Robertinoides pumilum Höglund, 1947

Pl. 18, fig. 8

Robertinoides pumilum Höglund in Höglund, 1947,
p. 227, pl. 18, fig. 5

Suborder Globigerinina Delange and Hérouard, 1896
Superfamily Heterohelicacea Cushman, 1927
Family Chiloguembelinidae Reiss, 1963

Genus *Strepolchilus* Brönnimann and Resig, 1971
Type species: *Bolivina tokelauae* Boersma, 1969, p.
329

Strepolchilus tokelauae (Boersma, 1969)
Pl. 23, fig. 6

Bolivina tokelauae Boersma in Kierstead et al., 1969,
p. 329, pl. 1, fig. 1
Strepolchilus tokelauae Boersma in Saito, Thompson,
and Dee Breger, 1981, p. 161, pl. 55, fig. 3

Superfamily Globorotaliacea Cushman, 1927
Family Globorotaliidae Cushman, 1927

Genus *Berggrenia* F. L., Parker, 1976
Type species: *Globanomalina praepumilio* F. L.,
Parker, 1967, p. 148

Berggrenia clarkei (Rögl and Bolli, 1973)
Pl. 22, fig. 5

Globigerina clarkei Rögl and Bolli in Rögl and Bolli,
1973, p. 563, pl. 14, figs. 13-15
Berggrenia clarkei Rögl and Bolli in Saito,
Thompson, and Dee Breger, 1981, p. 87, pl. 26, fig. 3

Genus *Neogloboquadrina* Bandy, Frerichs, and
Vincent, 1967
Type species: *Globigerina dutertrei* d'Orbigny, 1839,
p. 84

Neogloboquadrina dutertrei (d'Orbigny, 1839)
Pl. 22, fig. 4

Globigerina dutertrei d'Orbigny in d'Orbigny, 1839,
p. 84, pl. 4, figs. 19-21
Neogloboquadrina dutertrei d'Orbigny in Kennet and
Srinivasan, 1983, p. 198, pl. 48, figs. 7-9
Neogloboquadrina dutertrei d'Orbigny in Saito,
Thompson, and Dee Breger, 1981, p. 111, pl. 36, fig.
36

Neogloboquadrina incompta (Cifelli, 1961)
Pl. 22, fig. 2

Globigerina incompta Cifelli in Cifelli, 1961, p. 83,
pl. 4, figs. 1-7
Neogloboquadrina incompta Cifelli in Saito,
Thompson, and Dee Breger, 1981, p. 108, pl. 34, fig.
2

Neogloboquadrina pachyderma (Ehrenberg, 1861)
Pl. 22, fig. 3

Aristospira pachyderma Ehrenberg in Ehrenberg,
1861, pp. 276, 277, 303

Aristospira pachyderma Ehrenberg in Ehrenberg,
1872, pl. 1, fig. 4

Neogloboquadrina pachyderma Ehrenberg in Kennet
and Srinivasan, 1983, p. 192, pl. 47, figs. 2, 6-8
Neogloboquadrina pachyderma Ehrenberg in Saito,
Thompson, and Dee Breger, 1981, p. 106, pl. 34, fig.
1

Genus *Globorotalia* Cushman, 1927
Type species: *Pulvinulina menardii* d'Orbigny var.
tumida Brady, 1877, p. 535

Globorotalia inflata (d'Orbigny, 1839)
Pl. 21, fig. 1

Globigerina inflata d'Orbigny in d'Orbigny, 1839, p.
134, pl. 12, figs. 7-9
Globorotalia inflata d'Orbigny in Kennet and
Srinivasan, 1983, p. 118, pl. 27, figs. 7-9
Globorotalia inflata d'Orbigny in Saito, Thompson,
and Dee Breger, 1981, p. 124, pl. 41, fig. 1

Globorotalia scitula (Brady, 1882)
Pl. 21, fig. 2

Pulvinulina scitula Brady in Brady, 1882, p. 716
Globorotalia scitula Brady in Kennet and Srinivasan,
1983, p. 134, pl. 31, figs. 1, 3-5
Globorotalia scitula Brady in Saito, Thompson, and
Dee Breger, 1981, p. 137, pl. 46, fig. 2

Globorotalia hirsuta (d'Orbigny, 1839)
Pl. 21, fig. 3

Rotalia hirsuta d'Orbigny in d'Orbigny, 1839
Globorotalia hirsuta d'Orbigny in Parker, 1962, pl. 5,
figs. 12, 14

Genus *Truncorotalia* Cushman and Bermudez, 1949

Type species: *Rotalina truncatulinoides* d'Orbigny, 1839, p. 132

Truncorotalia crassaformis (Galloway and Wissler, 1927)
Pl. 21, fig. 4

Globigerina crassaformis Galloway and Wissler in Galloway and Wissler, p. 41, pl. 7, fig. 12

Globorotalia crassaformis Galloway and Wissler in Kennet and Srinivasan, 1983, p. 146, pl. 34, figs. 6-8

Globorotalia crassaformis Galloway and Wissler in Saito, Thompson, and Dee Breger, 1981, p. 129, pl. 43, fig. 2

Truncorotalia truncatulinoides (d'Orbigny, 1839)
Pl. 21, fig. 5

Rotalina truncatulinoides d'Orbigny in d'Orbigny, 1839, p. 132, pl. 1, figs. 25-27

Globorotalia truncatulinoides d'Orbigny in Kennet and Srinivasan, 1983, p. 148, pl. 34, figs. 2, pl. 35, figs. 4-6

Globorotalia truncatulinoides Saito, Thompson, and Dee Breger, 1981, p. 158, pl. 54, fig. 1

Family Pulleniatinidae Cushman, 1927

Genus *Pulleniatina* Cushman, 1927

Type species: *Pullenia obliquiloculata* Parker and Jones, 1865, p. 368

Pulleniatina obliquiloculata (Parker and Jones, 1865)
Pl. 22, fig. 1

Pullenia sphaeroides d'Orbigny var. *obliquiloculata* Parker and Jones in Parker and Jones, pp. 365, 368, pl. 19, fig. 4

Pulleniatina obliquiloculata Parker and Jones in Kennet and Srinivasan, 1983, p. 202, pl. 49, fig. 2, pl. 50, figs. 6-9

Pulleniatina obliquiloculata Parker and Jones in Saito, Thompson, and Dee Breger, 1981, p. 98, pl. 31, fig. 3

Superfamily Globorotaliacea Cushman, 1927

Family Candeinidae Cushman, 1927

Subfamily Tenuitallinae Banner, 1982

Genus *Tenuitella* Fleisher, 1974

Type species: *Globorotalia gemma* Jenkins, 1966, p. 1115

Tenuitella anfracta (Parker, 1967)
Pl. 23, fig. 5

Globorotalia anfracta Parker in Parker, 1967, p. 175, pl. 28, figs. 3-8

Globorotalia anfracta Parker in Kennet and Srinivasan, 1983, p. 164, pl. 39, figs. 2, 4, pl. 40, figs. 7-9

Tenuitella iota (Parker, 1962)
Pl. 23, fig. 4

Globigerinita iota Parker in Parker, 1962, p. 250, pl. 10, figs. 26-30

Globigerinita iota Parker in Saito, Thompson, and Dee Breger, 1981, p. 79, pl. 23, fig. 2

Subfamily Globigerinitinae Bermudez, 1961

Genus *Globigerinita* Brönnimann, 1951
Type species: *Globigerinita naparimaensis* Brönnimann, 1951, p. 18

Globigerinita glutinata (Egger, 1893)
Pl. 19, fig. 4

Globigerina glutinata Egger in Egger, 1839, p. 371, pl. 13, figs. 19-21

Globigerinita glutinata Egger in Kennet and Srinivasan, 1983, p. 224, pl. 56, figs. 1, 3-5

Globigerinita glutinata Egger in Saito, Thompson, and Dee Breger, 1981, p. 79, pl. 23, fig. 1

Globigerinita uvula (Ehrenberg, 1861)
Pl. 19, fig. 6

Pylodexia uvula Ehrenberg in Ehrenberg, 1861, p. 276, 277, 308

Globigerinita uvula Ehrenberg in Kennet and Srinivasan, 1983, p. 224, pl. 56, figs. 6-8

Globigerinita uvula Ehrenberg in Saito, Thompson, and Dee Breger, 1981, p. 81, pl. 24, fig. 3

Superfamily Globigerinacea Carpenter, Parker, and Jones, 1862

Subfamily Globigerininae Carpenter, Parker, and Jones, 1862

Genus *Globigerina* d'Orbigny, 1826
Type species: *Globigerina bulloides* d'Orbigny, 1826, p. 36

- Globigerina bulloides* d'Orbigny, 1826
Pl. 19, fig. 1
- Globigerina bulloides* d'Orbigny in d'Orbigny, 1826, p. 277
- Globigerina bulloides* d'Orbigny in Jones, 1994, p. 88, pl. 77, p. 88, pl. 79, figs. 3-7
- Globigerina bulloides* d'Orbigny in Kennet and Srinivasan, 1983, p. 36, pl. 6, figs. 4-6
- Globigerina bulloides* d'Orbigny in Saito, Thompson, and Dee Breger, 1981, p. 40, pl. 7, fig. 1
- Genus *Globigerinella* Cushman, 1927
Type species: *Globigerina aequilateralis* Brady, 1879, p. 285
- Globigerinella calida* (Parker, 1961)
Pl. 19, fig. 2
- Globigerina calida* Parker in Parker, p. 221, pl. 1, figs. 9-13, 15
- Globigerina calida* Parker in Cimerman and Langer, 1991, p. 57, pl. 60, figs. 2-3
- Globigerinella calida* Parker in Kennet and Srinivasan, 1983, p. 240, pl. 60, figs. 7-9
- Globigerinella calida* Parker in Saito, Thompson, and Dee Breger, 1981, p. 32, pl. 4, fig. 2
- Globigerinella siphonifera* (d'Orbigny, 1839)
Pl. 19, fig. 3
- Globigerina siphonifera* d'Orbigny in d'Orbigny, 1839, p. 83, pl. 4, figs. 15-18
- Genus *Globigerinoides* Cushman, 1927
Type species: *Globigerina rubra* d'Orbigny, 1839, p. 82
- Globigerinoides conglobatus* Brady, 1879
Pl. 20, fig. 1
- Globigerinoides conglobatus* Brady in Brady, 1879, p. 289
- Globigerinoides conglobatus* Brady in Brady, 1884, p. 603, pl. 80, figs. 1-5, pl. 82, fig. 5
- Globigerinoides conglobatus* Brady in Kennet and Srinivasan, 1983, p. 58, pl. 12, figs. 4-6
- Globigerinoides conglobatus* Brady in Saito, Thompson, and Dee Breger, 1981, p. 56, pl. 14, fig. 1
- Globigerinoides elongatus* (d'Orbigny, 1826)
Pl. 20, fig. 4
- Globigerina elongata* d'Orbigny in d'Orbigny, 1826, p. 277
- Globigerinoides elongata* d'Orbigny in Cushman, 1941, p. 40, pl. 10, figs. 20-23, pl. 11, fig. 3
- Globigerinoides elongatus* d'Orbigny in Saito, Thompson, and Dee Breger, 1981, p. 62, pl. 16, fig. 2
- Globigerinoides ruber* (d'Orbigny, 1839)
Pl. 20, fig. 2
- Globigerina rubra* d'Orbigny in d'Orbigny, 1839, p. 82, pl. 4, figs. 12-14
- Globigerinoides ruber* d'Orbigny in Kennet and Srinivasan, 1983, p. 78, pl. 10, fig. 6, pl. 17, figs. 1-3
- Globigerinoides ruber* d'Orbigny in Saito, Thompson, and Dee Breger, 1981, p. 59, pl. 15, fig. 1
- Globigerinoides sacculifer* (Brady, 1877)
Pl. 20, fig. 3
- Globigerina sacculifera* Brady in Brady, 1877, p. 535
- Globigerina sacculifera* Brady in Brady, 1884, p. 604, pl. 80, figs. 11-17, pl. 81, fig. 2, pl. 82, fig. 4
- Globigerinoides sacculifer* Brady in Kennet and Srinivasan, 1983, p. 66, pl. 14, fig. 4-6
- Globigerinoides sacculifer* Brady in Saito, Thompson, and Dee Breger, 1981, p. 65, pl. 17, fig. 1-2
- Globigerinoides trilobus* (Reuss, 1850)
Pl. 20, fig. 6
- Globigerina triloba* Reuss in Reuss, 1850, p. 374, pl. 447, fig. 11
- Globigerinoides triloba* Reuss in Kennet and Srinivasan, 1983, p. 62, pl. 10, fig. 4, pl. 13, fig. 1-3
- Genus *Globoturborotalita* Hofker, 1976
Type species: *Globigerina rubescens* Hofker, 1956, p. 234
- Globoturborotalita rubescens* (Hofker, 1956)
Pl. 19, fig. 5
- Globigerina rubescens* Hofker in Hofker, 1956, p. 234, pl. 32, fig. 26, pl. 35, fig. 18-21
- Globigerina rubescens* Hofker in Kennet and Srinivasan, 1983, p. 50, pl. 9, figs. 7-9
- Globigerina rubescens* Hofker in Saito, Thompson, and Dee Breger, 1981, p. 50, pl. 11, fig. 1
- Genus *Turborotalita* Blow and Banner, 1962
Type species: *Truncatulina humilis* Brady, 1884, p. 665

- Turborotalita quinqueloba* (Natland, 1938)
Pl. 23, fig. 1
- Globigerina quinqueloba* Natland in Natland, 1938, p. 149, pl. 6, fig. 7
- Globigerina quinqueloba* Natland in Kennet and Srinivasan, 1983, p. 32, pl. 5, figs. 4-6
- Globigerina quinqueloba* Natland in Saito, Thompson, and Dee Breger, 1981, p. 48, pl. 10, fig. 1-2
- Turborotalita humilis* (Brady, 1884)
Pl. 23, fig. 2
- Truncatulina humilis* Brady in Brady, 1884, p. 665, pl. 94, fig. 7
- Turborotalita humilis* Brady in Kennet and Srinivasan, 1983, p. 167, pl. 41, figs. 1, 3-5
- Turborotalita humilis* Brady in Saito, Thompson, and Dee Breger, 1981, p. 84, pl. 25, fig. 1-2, 4
- Turborotalita cristata* (Heron-Allen and Earland, 1929)
Pl. 23, fig. 3
- Globigerina cristata* Heron-Allen and Earland in Heron-Allen and Earland, 1929, p. 331, pl. 4, figs. 33-39
- Turborotalita cristata* Heron-Allen and Earland in Kennet and Srinivasan, 1983, p. 168, pl. 41, figs. 2, 6-8
- Subfamily Orbulinininae Schultze, 1854
- Genus *Orbulina* d'Orbigny, 1839
Type species: *Orbulina universa* d'Orbigny, 1839, p. 2
- Orbulina universa* d'Orbigny, 1839
Pl. 20, fig. 5
- Orbulina universa* d'Orbigny in d'Orbigny, 1839, p. 2, pl. 1, fig. 1
- Orbulina universa* d'Orbigny in Jones, 1994, p. 90, pl. 81, figs. 8-27
- Orbulina universa* d'Orbigny in Kennet and Srinivasan, 1983, p. 86, pl. 20, figs. 4-6
- Orbulina universa* d'Orbigny in Saito, Thompson, and Dee Breger, 1981, p. 70, pl. 19, fig. 1-6
- Suborder Rotaliina Delange and Hérouard, 1896
Superfamily Bolivinacea Glaessner, 1937
- Family Bolivinidae Glaessner, 1937
- Genus *Bolivina* d'Orbigny, 1839
Type species: *Bolivina plicata* d'Orbigny, 1839, p. 31
- Bolivina alata* (Seguenza, 1862)
Pl. 24, fig. 1
- Valvulina alata* Seguenza in Seguenza, 1862, p. 115, pl. 2, fig. 5
- Bolivina alata* Seguenza in Hofker, p. 248, figs. 85-88
- Brizalina alata* Seguenza in Cimerman and Langer, 1991, pl. 61, figs. 12-14
- Bolivina dilatata* (Reuss, 1850)
Pl. 24, fig. 2
- Brizalina dilatata* Reuss in Reuss, 1850, p. 381, pl. 48, figs 15 a-c.
- Brizalina dilatata* Reuss in Cimerman and Langer, 1991, p. 59, pl. 62, fig. 2.
- Bolivina pseudoplicata* Heron-Allen and Earland, 1930
Pl. 24, fig. 4
- Bolivina pseudo-plicata* Heron-Allen and Earland in Heron-Allen and Earland, 1930, p. 81, pl. 3, figs. 36-40
- Bolivina pseudoplicata* Heron-Allen and Earland in Kihle and Løfaldli, 1975
- Bolivina pseudoplicata* Heron-Allen and Earland in Cimerman and Langer, 1991, pl. 61, fig. 1-3.
- Bolivina difformis* (Williamson, 1858)
Pl. 24, fig. 6
- Textularia variabilis* var. *difformis* Williamson in Williamson, 1858, p. 77, pl. 6, figs. 166, 167
- Bolivina difformis* Williamson in Cushman, 1937, p. 164, pl. 15, figs. 13, 17
- Bolivina pygmaea* Williamson in Brady, 1960, pl. 53, figs. 5, 6
- Bolivina difformis* Williamson in Cimerman and Langer, 1991, pl. 61, fig. 9-11
- Brizalina difformis* Williamson in Murray, 2003, p. 20, pl. 6, fig. 2
- Bolivina spinescens* (Cushman, 1911)
Pl. 24, fig. 7
- Bolivina textilaroides* Brady in Brady, 1884, p. 419, pl. 52, figs. 24, 25
- Bolivina spinescens* Cushman in Cushman, 1911, p.

47, fig. 76

Bolivina subspinescens Cushman, 1922
Pl. 24, fig. 8

Bolivina subspinescens Cushman in Cushman, 1922,
p. 48, pl. 7, fig. 5
Brizalina subspinescens Cushman in Gabel, 1971, p.
54, pl. 15, fig. 4-5

Genus *Bolivinellina* Saidova, 1975
Type species: *Bolivinellina pescicula* Saidova, 1975,
p. 301

Bolivinellina pseudopunctata (Höglund, 1947)
Pl. 24, fig. 3

Bolivina pseudopunctata Höglund in Höglund, 1947,
p. 273, pl. 24, figs. 23-24
Brizalina pseudopunctata Höglund in Murray, 1971,
p. 109, pl. 44, figs. 3-6.
Bolivinellina pseudopunctata Höglund in Murray,
2003, p. 20, pl. 6, fig. 1

Bolivinellina striatula (Cushman, 1922)
Pl. 24, fig. 5

Bolivina striatula Cushman in Cushman, 1922, p. 27,
pl. 3, fig. 10
Bolivina striatula Cushman in Cushman, 1937, p.
154, pl. 18, figs. 30, 31
Brizalina striatula Cushman in Cimerman and
Langer, 1991, pl. 62, fig. 6-9

Genus *Brizalina* O.G. Costa, 1856
Type species: *Brizalina aenariensis* O.G. Costa,
1856, p. 296

Brizalina subaenariensis (Cushman, 1922)
Pl. 25, fig. 1

Bolivina aenariensis O.G. Costa in Brady, 1884, p.
423, pl. 53, figs. 10-11
Bolivina subaenariensis Cushman in Cushman, 1922,
p. 46, pl. 7, fig. 6

Superfamily Cassidulinacea d'Orbigny, 1839
Family Cassidulinidae d'Orbigny, 1839
Subfamily Cassidulininae d'Orbigny, 1839

Genus *Cassidulina* d'Orbigny, 1826
Type species: *Cassidulina laevigata* d'Orbigny, 1826,
p. 282

Cassidulina reniforme Nörvang, 1945
Pl. 25, fig. 3

Cassidulina crassa d'Orbigny var. *reniforme*
Nörvang in Nörvang, 1945, p. 41, fig. 6
Cassidulina reniforme Nörvang in Wollenburg and
Mackensen, 2009, p. 18, fig. 3, fig. 12

Cassidulina laevigata d'Orbigny, 1826
Pl. 25, fig. 4

Cassidulina laevigata d'Orbigny in d'Orbigny, 1826,
p. 282, pl. 15, figs. 4-5
Cassidulina laevigata d'Orbigny in Kihle and
Løfaldli, 1975
Cassidulina laevigata d'Orbigny in Cimerman and
Langer, 1991, p. 61, pl. 63, fig. 1-3
Cassidulina laevigata d'Orbigny in Murray, 2003, p.
21, pl. 6, fig. 10

Cassidulina carinata Silvestri, 1896
Pl. 25, fig. 5

Cassidulina laevigata d'Orbigny var. *carinata*
Silvestri in Silvestri, 1896, p. 104, pl. 2, fig. 10
Cassidulina laevigata carinata Silvestri in Jorissen,
1987, p. 34, pl. 1, fig. 8
Cassidulina laevigata carinata Silvestri in
Seidenkrantz, 1995, p. 148, pl. 1, figs. 7-9, p. 156, pl.
5, figs. 7-8

Cassidulina teretis Tappan, 1951
Pl. 25, fig. 6

Cassidulina teretis Tappan in Tappan, 1951, p. 7, pl.
1, fig. 30
Cassidulina teretis Tappan in Seidenkrantz, 1995, p.
148, pl. 1, fig. 12-13, p. 151, pl. 2, figs. 15-18

Cassidulina neoteretis Seidenkrantz, 1995
Pl. 26, fig. 1

Cassidulina neoteretis Seidenkrantz in Seidenkrantz,
1995, p. 148, pl. 1, figs. 1-6, p. 151, pl. 2, figs. 1-14
Cassidulina neoteretis Seidenkrantz in Wollenburg
and Mackensen, p. 18, fig. 3, fig. 10

Cassidulina crassa d'Orbigny, 1839
Pl. 26, fig. 3

Cassidulina crassa d'Orbigny in d'Orbigny, 1839, p.
56, pl. 7, figs. 18-20
Cassidulina crassa d'Orbigny in Kihle and Løfaldli,

1975

Cassidulina crassa d'Orbigny in Jones, 1994, p. 60, pl. 54, fig. 4

Cassidulina crassa d'Orbigny in Wollenburg and Mackensen, 2009, p. 18, fig. 3, fig. 9

Genus *Takayanagia* Nomura, 1983

Type species: *Cassidulina delicata* Cushman, 1927, p. 168

Takayanagia delicata (Cushman, 1927)

Pl. 26, fig. 2

Cassidulina delicata Cushman in Cushman, 1927, p. 168, pl. 6, fig. 5

Takayanagia delicata Cushman in Loeblich and Tappan, 1987, p. 507, pl. 560, figs. 5-10

Genus *Globocassidulina* Voloshinova, 1960

Type species: *Cassidulina globosa* Hantken, 1876, p. 64

Globocassidulina subglobosa (Brady, 1881)

Pl. 26, fig. 4

Cassidulina subglobosa Brady in Brady, 1881, p. 60, pl. 54, fig. 17

Globocassidulina subglobosa Brady in Jones, 1994, p. 60, pl. 54, fig. 17

Globocassidulina subglobosa Brady in Murray, 2003, p. 24, pl. 8, fig. 7

Globocassidulina subglobosa Brady in Cimeman and Langer, 1991, p. 61, pl. 63, fig. 4-6

Genus *Cassidulinoides* Cushman, 1927

Type species: *Cassidulina parkeriana* Brady, 1881, p. 59

Cassidulinoides bradyi (Norman, 1881)

Pl. 26, fig. 6

Cassidulina bradyi Norman in Norman, 1881, p. 59, pl. 54, figs. 6-10

Genus *Islandiella* Nørvang, 1959

Type species: *Cassidulina islandica* Nørvang, 1945, p. 41

Islandiella norcrossi (Cushman, 1933)

Pl. 26, fig. 5

Cassidulina norcrossi Cushman in Cushman, 1933, p. 7, pl. 2, fig. 7

Islandiella norcrossi Cushman in Wollenburg and

Mackensen, 2009, p. 18, text-fig. 3, fig. 13

Subfamily Ehrenbergininae Cushman, 1927

Genus *Ehrenbergina* Reuss, 1850

Type species: *Ehrenbergina serrata* Reuss, 1850, p. 377

Ehrenbergina trigona Goës, 1896

Pl. 27, fig. 1

Ehrenbergina serrata Brady var. *trigona* Goës in Goës, 1896, p. 434, pl. 55, figs. 2-7

Ehrenbergina trigona Goës in Jones, 1994, p. 61, pl. 55, figs. 2-3, 5

Ehrenbergina trigona Goës in Hermelin and Scott, 1985, p. 206, pl. 4, figs. 15-16

Superfamily Turrilinacea Cushman, 1927

Family Stainforthiidae Reiss, 1963

Genus *Stainforthia* Hofker, 1956

Type species: *Virgulina concava* Höglund, 1947, p. 257

Stainforthia fusiformis (Williamson, 1858)

Pl. 27, fig. 2

Bulimina pupoides var. *fusiformis* Williamson in Williamson, 1858, p. 63,

Stainforthia fusiformis Williamson in Kihle and Løfaldli, 1975

Stainforthia fusiformis Williamson in Murray, 2002, p. 26, pl. 10, figs. 1-4

Stainforthia loeblichi (Feyling-Hanssen, 1954)

Pl. 27, fig. 3

Virgulina loeblichi Feyling-Hanssen in Feyling-Hanssen, 1954, p. 191, pl. 1, figs. 14-18

Stainforthia skagerakensis (Höglund, 1947)

Pl. 27, fig. 5

Virgulina skagerakensis Höglund in Höglund, 1947, p. 255, pl. 23, figs. 1-2

Stainforthia skagerakensis Höglund in Kihle and Løfaldli, 1975

Superfamily Buliminacea Jones, 1875

Family Siphogenerinoididae Saidova, 1981

- Subfamily Siphogenerinoidinae Saidova, 1981
- Genus *Parabrizalina* Zweig-Strykowski and Reiss, 1976
Type species: *Bolivina porrecta* Brady, 1881, p. 57
- Parabrizalina porrectum* (Brady, 1881)
Pl. 25, fig. 2
- Bulimina (Bolivina) porrecta* Brady in Brady, 1881, p. 57
- Bulimina (Bolivina) porrecta* Brady in Brady, 1881, pl. 52, fig. 22
- Subfamily Tubulogenerininae Saidova, 1981
- Genus *Rectuvigerina* Mathews, 1945
Type species: *Siphogenerina multicostata* Cushman and Jarvis, 1929, p. 14
- Rectuvigerina elongatastriata* (Colom, 1952)
Pl. 27, fig. 6
- Uvigerina cf. tenuistriata* Reuss in Colom, 1941, p. 17, pl. 3, figs. 57-58
- Angulogerina elongatastriata* Colom in Colom, 1952, p. 29, pl. 4, figs. 6-9
- Rectuvigerina elongatastriata* Colom in Cimerman and Langer, 1991, p. 61, pl. 63, figs. 7-9
- Genus *Siphogenerina* Schlumberger, 1882
Type species: *Siphogenerina costata* Schlumberger, 1883, p. 26
- Siphogenerina columellaris* (Brady, 1881)
Pl. 27, fig. 7
- Uvigerina columellaris* Brady in Brady, 1881, p. 64, pl. 75, figs. 15-17
- Siphogenerina columellaris* Brady in Jones, 1994, p. 87, pl. 75, figs. 15-17
- Family Buliminidae Jones, 1875
- Genus *Bulimina* d'Orbigny, 1826
Type species: *Bulimina marginata* d'Orbigny, 1826, p. 269
- Bulimina aculeata* d'Orbigny, 1826
Pl. 27, fig. 8
- Bulimina aculeata* d'Orbigny in
- d'Orbigny, 1926, p. 269, n. 7
- Bulimina aculeata* d'Orbigny in Colom, 1974, p. 115, fig. 16g
- Bulimina aculeata* d'Orbigny in Cimerman and Langer, 1991, p. 61, pl. 63, figs. 10-11
- Bulimina aculeata* d'Orbigny in Sgarrella and Moncharmon-Zei, 1993, p. 211, pl. 15, fig. 1
- Bulimina marginata* d'Orbigny, 1826
Pl. 27, fig. 9
- Bulimina marginata* d'Orbigny in d'Orbigny, 1826, p. 269, pl. 12, figs. 10-12
- Bulimina marginata* var. *marginata*, d'Orbigny in Hottinger, Halicz, and Reiss, 1993, p. 99, pl. 125, figs. 1-6
- Bulimina marginata* d'Orbigny in Cimerman and Langer, 1991, p. 62, pl. 64, figs 9-11
- Bulimina marginata* d'Orbigny in Murray, 2003, p. 20, pl. 6, figs. 4-5
- Bulimina striata* d'Orbigny, 1826
Pl. 27, fig. 10
- Bulimina striata* d'Orbigny in d'Orbigny, 1826, p. 269
- Bulimina inflata* Seguenza in Brady, 1884, p. 406, pl. 51, figs 11, 13
- Bulimina striata* var. *notoensis* Asano in Wang et al., 1988, p. 150, pl. 21, fig. 4
- Bulimina striata* d'Orbigny in Akimoto, 1990, p. 194, pl. 16, fig. 8
- Bulimina striata* d'Orbigny in Van Marle, 1991, p. 88, pl. 5, figs 6-8
- Bulimina striata* d'Orbigny in Loeblich and Tappan, 1994, p. 125, pl. 242, figs 8-14
- Genus *Globobulimina* Cushman, 1927
Type species: *Globobulimina pacifica* Cushman, 1927, p. 67
- Globobulimina affinis* (d'Orbigny, 1839)
Pl. 28, fig. 2
- Bulimina affinis* d'Orbigny in d'Orbigny, 1839, p. 105, pl. 2, figs. 25-26
- Bulimina affinis* d'Orbigny in Brady, 1884, p. 400, pl. 50, fig. 14
- Bulimina affinis* d'Orbigny in Cushman, 1911, p. 79, text-fig. 130
- Bulimina affinis* d'Orbigny in Phleger and Parker,

- 1951, p. 15, pl. 7, figs 21-22
Bulimina affinis d'Orbigny in Loeblich and Tappan, 1994, p. 124, pl. 240, figs 12-13
- Globobulimina doliolum* (Terquem and Terquem, 1886)
 Pl. 28, fig. 5
- Bulimina doliolum* Terquem and Terquem in Terquem and Terquem, 1886, p. 333, pl. 11, figs. 17-18
- Globobulimina turgida* (Bailey, 1851)
 Pl. 28, fig. 1
- Bulimina turgida* Bailey in Bailey, 1851, p. 12, figs. 28-31, 67
Globobulimina turgida Bailey in Gabel, 1971, p. 52, pl. 14, fig. 21
- Genus *Praeglobobulimina* Hofker, 1951
 Type species: *Bulimina pyrula* d'Orbigny var. *spinescens* Brady, 1884, p. 400
- Praeglobobulimina ovata* (d'Orbigny, 1846)
 Pl. 28, fig. 4
- Bulimina ovata* d'Orbigny in d'Orbigny, 1846, p. 185, pl. 11, figs. 13-14
Bulimina ovata d'Orbigny in Brady, 1884, p. 400, pl. 50, fig. 13
Praeglobobulimina ovata d'Orbigny in Jones, 1994, p. 54, pl. 50, fig. 13
- Family Buliminellidae Hofker, 1951
 Genus *Buliminella* Cushman, 1911
 Type species: *Bulimina elegantissima* d'Orbigny, 1839, p. 51
- Buliminella spinigera* Cushman, 1922
 Pl. 28, fig. 6
- Buliminella spinigera* Cushman in Cushman, 1922, p. 113, pl. 23, figs. 1-3
- Subfamily Uvigerininae Haeckel, 1894
 Genus *Uvigerina* d'Orbigny, 1826
 Type species: *Uvigerina pygmaea* d'Orbigny, 1826, p. 268
- Uvigerina mediterranea* Hofker, 1932
- Pl. 29, fig. 1
- Uvigerina mediterranea* Hofker in Hofker, 1932, p. 118, figs. 32
Uvigerina mediterranea Hofker in Colom, 1974, p. 122, figs. 19h-n
Uvigerina mediterranea Hofker in Cimerman and Langer, 1991, p. 63, pl. 65, figs. 7-9
Uvigerina mediterranea Hofker in Abu-Zied et al., 2008, p. 52, pl. 2, figs. 17-18
- Uvigerina peregrina* Cushman, 1923
 Pl. 29, fig. 2
- Uvigerina peregrina* Cushman in Cushman, 1923, p. 166, pl. 42, figs. 7-10
Uvigerina peregrina Cushman in Abu-Zied et al., 2008, p. 52, pl. 2, figs. 19-20
Uvigerina peregrina Cushman in Phleger and Parker, 1951, p. 18, pl. 8 , figs 22, 24-26
Uvigerina peregrina Cushman in Lutze, 1986, p. 32, pl. 1, figs 1-6
Uvigerina peregrina Cushman in Ujiié, 1990, p. 31, pl. 13, figs 1-3
- Uvigerina auberiana* d'Orbigny, 1839
 Pl. 29, fig. 3
- Uvigerina auberiana* d'Orbigny in d'Orbigny, 1839, p. 106, pl. 2 , figs 23-24
Uvigerina asperula Czjzek var. *aueriana* d'Orbigny in Brady, 1884, p. 579, pl. 75, fig. 9
Uvigerina asperula Czjzek in Brady, 1884, p. 578, pl. 75, figs 6-8
Uvigerina asperula Czjzek in Ujiié, 1990, p. 31, pl. 13, figs 7-8
Uvigerina auberiana d'Orbigny in Uchio, 1960, p. 65, pl. 7, fig. 11
Uvigerina auberiana d'Orbigny in Jones, 1994, p. 86, pl. 75, figs 6-9
- Uvigerina peregrina parva* Lutze, 1986
 Pl. 29, fig. 4
- Uvigerina peregrina* Cushman var. *parva* Lutze in Lutze, 1986, p. 36, pl. 3, fig. 1-5
Uvigerina bradyana Fornasini in Austin and Evans, 2000, p. 690, fig. 3.1
Uvigerina peregrina Cushman var. *parva* Lutze in Schönfeld, 2006, p. 359, pl. 1, figs. 12-13
- Uvigerina pigmaea* d'Orbigny, 1826
 Pl. 29, fig. 5

Uvigerina pigmea d'Orbigny in d'Orbigny, 1826, p. 269, pl. 12, figs. 8, 9
Uvigerina pigmea d'Orbigny in Thomas, 1980, pl. 3, fig. 3
Uvigerina pigmea d'Orbigny in Timm, 1992, p. 68, pl. 6, figs. 1a-b
Uvigerina pygmea d'Orbigny in Lutze, 1986, p. 36, pl. 3, figs. 6-8
Uvigerina pigmea d'Orbigny in Schönfeld, 2006, p. 357, pl. 1, figs. 6-11

Subfamily *Angulogerininae* Galloway, 1933

Genus *Angulogerina* Cushman, 1927
Type species: *Uvigerina angulosa* Williamson, 1858, p. 67

Angulogerina angulosa (Williamson, 1858)
Pl. 29, fig. 6

Uvigerina angulosa Williamson in Williamson, 1858, p. 67, pl. 5, fig. 140
Angulogerina angulosa Williamson in Y. Le Calvez, 1958, p. 180
Trifarina angulosa Williamson in Barker, pl. 74, figs. 15-16
Trifarina angulosa Williamson in Daniels, p. 83, pl. 6, fig. 4
Trifarina angulosa Williamson in Abu-Zied et al., 2008, p. 52, pl. 2, fig. 21
Angulogerina angulosa Williamson in Loeblich and Tappan, 1987, p. 525, pl. 574, figs. 5-9

Genus *Trifarina* Cushman, 1923
Type species: *Trifarina bradyi* Cushman, 1923, p. 99

Trifarina bradyi (Cushman, 1923)
Pl. 29, fig. 7

Rhabdogonium tricarinatum d'Orbigny in Brady, 1884, p. 525, pl. 67, figs 1-3
Trifarina bradyi Cushman in Cushman, 1923, p. 99, pl. 22, figs 3-9
Trifarina bradyi Cushman in Loeblich and Tappan, 1987, p. 526, pl. 574, figs. 10-13
Trifarina bradyi Cushman in Van Marle, 1991, p. 110, pl. 7, figs 8-9
Trifarina bradyi Cushman in Jones, 1994, p. 78, pl. 67, figs 1-3
Trifarina bradyi Cushman in Loeblich and Tappan, 1994, p. 128, pl. 251, figs 6-16

Trifarina fornasini (Sell, 1948)

Pl. 29, fig. 8

Angulogerina fornasini Sell, 1948, p. 40, pl. 43, figs. 1-4

Superfamily *Fursenkoinacea* Loeblich and Tappan, 1961

Family *Fursenkoinidae* Loeblich and Tappan, 1961

Genus *Fursenkoina* Loeblich and Tappan, 1961
Type species: *Virgulina squammosa* d'Orbigny, 1826, p. 267

Fursenkoina complanata (Egger, 1893)
Pl. 27, fig. 4

Virgulina schreibersiana Cziczek var. *complanata* Egger in Egger, 1893, p. 292, pl. 8, figs. 91-92
Fursenkoina complanata Egger in Jones, 1994, p. 56, pl. 52, figs. 1-3
Fursenkoina complanata Egger in Austin and Evans, 2000, p. 684, pl. 1, fig. n

Superfamily *Stilostomellacea* Finlay, 1947
Family *Stilostomellidae* Finlay, 1947

Genus *Nodogenerina* Cushman, 1927
Type species: *Nodogenerina bradyi* Cushman, 1927, p. 79

Nodogenerina virgula (Brady, 1884)
Pl. 29, fig. 9

Sagrina virgula Brady in Brady, 1884, pl. 76, fig. 8
Nodogenerina virgula Brady in Jones, 1994, p. 88, pl. 76, fig. 8, suppl. pl. 2, figs. 2-3, 14

Superfamily *Discorbacea* Ehrenberg, 1838
Family *Bagginidae* Cushman, 1927
Subfamily *Baggininae* Cushman, 1927

Genus *Cancris* de Montfort, 1808
Type species: *Nautilus auricula* Fichtel and Moll, 1798, p. 108

Cancris auriculus (Fichtel and Moll, 1798)
Pl. 29, fig. 10

Nautilus auricula Fichtel and Moll in Fichtel and Moll, 1798, p. 102, pl. 18, figs. g-i
Cancris auriculatus Fichtel and Moll in Loeblich and

Tappan, 1988, p. 545, pl. 591, figs. 1-4
Cancris auriculus Fichtel and Moll in Jones, 1994, p. 105, pl. 106, fig. 4
Cancris auricula Fichtel and Moll in Murray, 2003, p. 19, pl. 6, figs. 6-7

Genus *Valvulineria* Cushman, 1926
Type species: *Valvulineria californica* Cushman, 1926, p. 59

Valvulineria bradyana (Fornasini, 1900)
Pl. 30, fig. 1

Discorbina bradyana Fornasini in Fornasini, 1900, p. 393, textfig. 43
Valvulineria bradyana Fornasini in Jorissen, p. 26, pl. 4, figs. 1-2

Family Eponididae Hofker, 1951
Subfamily Eponidinae Hofker, 1951

Genus *Ioanella* Saidova, 1975
Type species: *Truncatulina tumidula* Brady, 1884, p. 666

Ioanella tumidula (Brady, 1884)
Pl. 30, fig. 2

Truncatulina tumidula Brady in Brady, 1884, p. 666, pl. 95, fig. 8
Ioanella tumidula Brady in Duchemin et al., 2007, p. 18, pl. 3, figs. 16-18
Ioanella tumidula Brady in Wollenburg and Mackensen, 2009, p. 18, textfig. 3, fig. 4

Subfamily Stomatorbininae Saidova, 1981

Genus *Stomatorbina* Doreen, 1948
Type species: *Lamarcina torrei* Cushman and Bermudez, 1937, p. 21

Stomatorbina concentrica (Parker and Jones, 1864)
Pl. 30, fig. 3

Pulvinulina concentrica Parker and Jones in Parker and Jones, 1864, p. 470, pl. 48, fig. 14
Stomatorbina concentrica Parker and Jones in Cimerman and Langer, 1991, p. 65, pl. 68, figs. 7-9
Mississippina concentrica Parker and Jones in Jones, 1994, p. 104, pl. 105, fig. 1

Family Rosalinidae Reiss, 1963

Genus *Hyrrokkin* Cedhagen, 1994
Type species: *Hyrrokkin sarcophaga* Cedhagen, 1994, p. 66

Hyrrokkin sarcophaga Cedhagen, 1994
Pl. 30, figs. 4-5

Pulvinulina repanda var. *punctulata* d'Orbigny in Parker and Jones, 1865, p. 394, pl. 14, figs. 12-13
Rosalina carnivora Todd in Todd, 1965, pp. 834-835, pl. 1, figs. 1-3, pl. 2, figs. 1-4, pl. 3, figs. 1
Hyrrokkin sarcophaga Cedhagen in Cedhagen, 1994, p. 66, textfigs. 1-17
Hyrrokkin sarcophaga Cedhagen in Freiwald and Schönfeld, 1995, p. 202, textfig. 2

Genus *Gavelinopsis* Hofker, 1951
Type species: *Discorbina praegeri* Heron-Allen and Earland, 1913, p. 122

Gavelinopsis nitida (Williamson, 1858)
Pl. 31, fig. 1

Rotalina nitida Williamson in Williamson, 1858, p. 54, pl. 4, figs. 106-108

Gavelinopsis praegeri (Heron-Allen and Earland, 1913)
Pl. 31, fig. 2

Discorbina praegeri Heron-Allen and Earland in Heron-Allen and Earland, 1913, p. 122, pl. 10, figs. 8-10.

Gavelinopsis praegeri Heron-Allen and Earland in Hofker, 1951, p. 486, text-figs 332-334
Gavelinopsis praegeri Heron-Allen and Earland in Zheng, 1980, p. 167, pl. 5, fig. 1
Gavelinopsis praegeri Heron-Allen and Earland in Loeblich and Tappan, 1987, p. 560, pl. 608, figs 6-12
Gavelinopsis praegeri Heron-Allen and Earland in Loeblich and Tappan, 1994, p. 138, pl. 281, figs 1-10
Gavelinopsis praegeri Heron-Allen and Earland in Murray, 2003, p. 24, pl. 8, figs. 5-6

Gavelinopsis caledonia Murray and Whittaker, 2001
Pl. 31, fig. 3

Gavelinopsis caledonia Murray and Whittaker in Murray and Whittaker, 2001, p. 179, pl. 1, figs. 1-10, pl. 2, figs. 1-7

Gavelinopsis caledonia Murray and Whittaker in Murray, 2003, p. 24, pl. 8, figs. 2-4

Genus *Rosalina* d'Orbigny, 1826
 Type species: *Rosalina globularis* d'Orbigny, 1826,
 p. 62

Rosalina globularis d'Orbigny, 1826
 Pl. 31, fig. 4

Rosalina globularis d'Orbigny in d'Orbigny, 1826, p.
 271, pl. 13,
 figs 1-4.

Discorbina globularis d'Orbigny in Brady, 1884, p.
 643, pl. 86, fig. 13

Rosalina globularis d'Orbigny in Todd, 1965, p. 11,
 pl. 3, fig. 4

Rosalina globularis d'Orbigny in Loeblich and
 Tappan, 1987, p. 561, pl. 610, figs 1-5; pl. 611, figs
 1-3

Rosalina globularis d'Orbigny in Jones, 1994, p. 93,
 pl. 86, fig. 13

Rosalina bradyi (Cushman, 1915)
 Pl. 31, fig. 5

Discorbina globularis var. *bradyi* Cushman in
 Cushman, 1915, p. 12, pl. 8, fig. 1
Discopulvinulina bradyi Cushman in Hofker, 1951, p.
 452, figs. 310
Rosalina bradyi Cushman in Hornbrook and Vella,
 1954, p. 26

Rosalina semipunctata (Bailey, 1851)
 Pl. 32, fig. 1

Rotalina semipunctata Bailey in Bailey, 1851, p. 11,
 figs. 17-19, 67

Family Sphaeroidinidae Cushman, 1927

Genus *Sphaeroidina* d'Orbigny, 1826
 Type species: *Sphaeroidina bulloides* d'Orbigny,
 1826, p. 267

Sphaeroidina bulloides d'Orbigny, 1826
 Pl. 32, fig. 2

Sphaeroidina bulloides d'Orbigny in d'Orbigny,
 1826, p. 267, pl. 2, fig. 58

Sphaeroidina bulloides d'Orbigny in Kihle and
 Løfaldli, 1975

Sphaeroidina bulloides d'Orbigny in Loeblich and
 Tappan, 1987, p. 564, pl. 617, figs. 1-6

Superfamily Glabratellacea Loeblich and Tappan,
 1964
 Family Glabratellidae Loeblich and Tappan, 1964

Genus *Glabratella* Dorreen, 1948
 Type species: *Glabratella crassa* Dorreen, 1948, p.
 294

Glabratella patelliformis (Brady, 1884)
 Pl. 32, fig. 3

Discorbina patelliformis Brady in Brady, 1884, p.
 647, pl. 88, fig. 3, pl. 89, fig. 1
Glabratella patelliformis Brady in Jones, 1994, p. 94,
 pl. 88, fig. 3, p. 95, pl. 89, fig. 1

Genus *Heronallenita* Seiglie and Bermudez, 1965
 Type species: *Heronallenita striatospinata* Seiglie
 and Bermudez, 1965, p. 61

Heronallenita lingulata (Burrows and Holland, 1895)
 Pl. 32, fig. 5

Discorbina lingulata Burrows and Holland in
 Burrows and Holland, 1895, p. 297, pl. 7, fig. 33
Heronallenita lingulata Burrows and Holland in
 Hermelin, 1989, p. 68, pl. 12, figs. 14, 18
Heronallenita lingulata Burrows and Holland in
 Jones, 1994, pp. 96, pl. 91, fig. 3

Superfamily Siphoninacea Cushman, 1927
 Family Siphoninidae Cushman, 1927

Genus *Siphonina* Reuss, 1850
 Type species: *Rotalina reticulata* Czjzek, 1848, p.
 294

Siphonina reticulata (Czjzek, 1848)
 Pl. 32, fig. 4

Rotalia reticulata Czjzek in Czjzek, 1848, p. 145, pl.
 13, figs. 7-9
Siphonina reticulata Czjzek in Cimerman and
 Langer, 1991, p. 69, pl. 73, figs. 11-13
Siphonina reticulata Czjzek in Loeblich and Tappan,
 1987, p. 571, pl. 624, figs. 4-6 and 13-15

Superfamily Discorbinellacea Sigal, 1952
 Family Parrelloididae Hofker, 1956

Genus *Cibicidoides* Thalmann, 1939

Type species: *Truncatulina mundula* Brady, Parker, and Jones, 1939, p. 228

Cibicidoides mundulus (Parker, 1953)
Pl. 33, fig. 1

Truncatulina mundula Brady, Parker, and Jones in Brady, Parker, and Jones, 1888, p. 228, pl. 45, figs. 25
Cibicides kullenbergi Parker in Phleger et al., 1953, p. 49, pl. 11, figs. 7-8

Cibicides kullenbergi Parker in Boltovskoy, 1978, pl. 3, figs. 9-12

Cibicidoides mundulus Brady, Parker and Jones in Hermelin, 1989, p. 86, pl. 17, figs. 9-11

Cibicidoides pachyderma (Rzehak, 1886)
Pl. 33, fig. 2

Truncatulina pachyderma Rzehak in Rzehak, 1886, p. 87, pl. 1, fig. 5

Truncatulina pseudoungeriana Cushman in Cushman, 1922b, p. 97, pl. 20, fig. 9

Cibicides pseudoungeriana Cushman in Cushman, 1931, p. 123, pl. 22, figs 3-7

Cibicidoides pachyderma Rzehak in van Morkhoven et al., 1986, p. 68, pl. 22, fig. 1

Cibicidoides pachyderma Rzehak in Jones, 1994, p. 98, pl. 94, fig. 9

Cibicidoides pseudoungerianus Cushman in Hess, 1998, p. 78, pl. 16, figs 1-2

Family Pseudoparrellidae Voloshinova, 1952
Subfamily Pseudoparrellinae Voloshinova, 1952

Genus *Epistominella* Husezima an Maruhasi, 1944
Type species: *Epistominella pulchella* Husezima an Maruhasi, 1944, p. 397

Epistominella vitrea Parker, 1953
Pl. 33, fig. 3

Epistominella vitrea Parker in Parker et al., 1953, p. 9, pl. 4, fig. 34-36, 40-41

Epistominella vitrea Parker in Hayward et al., 1999, pl. 13, figs. 14-16

Epistominella vitrea Parker in Duchemin et al., 2007, p. 17, pl. 2, figs. 5-6

Epistominella vitrea Parker in Margreth et al., 2009, p. 2230, pl. 1, fig. 7

Epistominella exigua (Brady, 1884)
Pl. 33, fig. 4

Pulvinulina exigua Brady in Brady, 1884, p. 696, pl. 103, figs. 13-14

Epistominella exigua Brady in Hermelin and Scott, 1985, p. 208, pl. 4, fig. 1

Epistominella exigua Brady in Wollenburg and Mackensen, 2009, p. 18, textfig. 3, fig. 5

Epistominella exigua Brady in Margreth et al., 2009, p. 2230, pl. 1, fig. 6

Family Discorbinellidae Sigal, 1952
Subfamily Discorbinellinae Sigal, 1952

Genus *Discorbinella* Cushman and Martin, 1935
Type species: *Discorbinella montereyensis* Cushman and Martin, 1935, p. 89

Discorbinella bertheloti (d'Orbigny, 1839)
Pl. 33, fig. 5

Rosalina bertheloti d'Orbigny in d'Orbigny, 1839b, p. 135, pl. 1 , figs 28-30

Discorbina bertheloti d'Orbigny in Brady, 1884, p . 650, pl. 89, figs 10-12

Discorbis bertheloti d'Orbigny in Cushman, 1931, p . 16, pl. 3, fig. 2

Discopulvinulina bertheloti d'Orbigny in Hofker, 1951, p. 449

Discorbinella bertheloti d'Orbigny in Loeblich and Tappan, 1987, p. 577, pl. 630, figs 4-6

Discorbinella bertheloti d'Orbigny in Jones, 1994, p. 95, pl. 89, figs 10-12

Superfamily Planorbulinacea Schwager, 1877
Family Planulinidae Bermudez, 1952

Genus *Planulina* d'Orbigny, 1826
Type species: *Planulina ariminensis* d'Orbigny, 1826, p. 66

Planulina ariminensis d'Orbigny, 1826
Pl. 34, fig. 1

Planulina ariminensis d'Orbigny in d'Orbigny, 1826, p. 280 pl. 14, fig. 1-3

Planulina ariminensis d'Orbigny in Hermelin and Scott, 1985, p. 214, pl. 4, fig. 9-11

Planulina ariminensis d'Orbigny in Jones, 1994, p. 98, pl. 93, fig. 10-11

Planulina ariminensis d'Orbigny in Abu-Zied et al., 2008, p. 52, pl. 2, figs. 31-32

Genus *Hyalinea* d'Orbigny, 1826

Type species: *Nautilus balthicus* Schröter, 1783, p. 20

Hyalinea balthica (d'Orbigny, 1826)
Pl. 34, fig. 2

Nautilus balthicus d'Orbigny in Schröter, 1783, p. 20,
pl. 1, fig. 2

Hyalinea balthica d'Orbigny in Jones, 1994, p. 110,
pl. 112, figs. 1-2

Hyalinea balthica d'Orbigny in Cimerman and
Langer, 1991, p. 70, pl. 74, figs. 4, 7

Family Cibicididae Cushman, 1927
Subfamily Cibicidinae Cushman, 1927

Genus *Cibicides* de Montfort, 1808

Type species: *Cibicides refulgens* de Montfort, 1808,
p. 122

Cibicides ungerianus (d'Orbigny, 1826)
Pl. 34, fig. 3

Rotalina ungeriana d'Orbigny in d'Orbigny, 1846, p.
157, pl. 8, figs. 16-18

Cibicides ungerianus d'Orbigny in Schweizer et al.,
2009, p. 301, textfig. 1, fig. k-l

Cibicides refulgens de Montfort, 1808
Pl. 34, fig. 4

Cibicides refulgens de Montfort in de Montfort, 1808,
p. 122

Cibicides refulgens de Montfort in Kihle and
Løfaldli, 1975

Cibicides refulgens de Montfort in Cimerman and
Langer, 1991, p. 70, pl. 75, figs. 5-9

Cibicides refulgens de Montfort in Murray, 2003, p.
21, pl. 7, figs. 1-2

Cibicides aravaensis Perelis and Reiss, 1976
Pl. 34, fig. 5

?*Cibicides* sp. Venec-Peyre in Venec-Peyre, 1973, p.
28, pl. C, fig. 2

Cibicides aravaensis Perelis and Reiss in Perelis and
Reiss, 1976, p. 93, pl. 8, figs. 1-7

Genus *Lobatula* Fleming, 1828

Type species: *Nautilus lobatulus* Walker and Jacob in
Kanmacher, 1798, p. 642

Lobatula lobatula (Walker and Jacob, 1798)
Pl. 35, fig. 1

Nautilus lobatulus Walker and Jacob in Kanmacher,
1798, p. 642, pl. 14, fig. 36

Lobatula lobatula Walker and Jacob in Cimerman
and Langer, 1991, p. 71, pl. 75, figs. 1-4

Cibicides lobatulus Walker and Jacob in Jones, 1994,
p. 97, pl. 93, figs. 1, 4-5

Cibicides lobatulus Walker and Jacob in Murray,
2003, p. 21, pl. 6, figs. 13-15

Genus *Fontbotia* Gonzales-Donoso and Linares, 1970

Type species: *Anomalina wuellerstorfi* Schwager,
1866, p. 258

Fontbotia wuellerstorfi (Schwager, 1866)
Pl. 35, fig. 2

Anomalina wüllerstorfi Schwager in Schwager, 1866,
p. 258, pl. 7, figs. 105, 107

Planulina wuellerstorfi Schwager in Cushman, 1931,
p. 110, pl. 19, fig. 5-6

Fontbotia wuellerstorfi Schwager in Gonzales-
Donoso and Linares, 1970, p. 238, pl. 1, fig. 4

Fontbotia wuellerstorfi Schwager in Loeblich and
Tappan, 1987, p. 538, pl. 634, figs. 1-3

Cibicidoides wuellerstorfi Schwager in Jones, 1994,
p. 98, pl. 93, figs. 8-9

Cibicides wuellerstorfi Schwager in Abu-Zied et al.,
2008, p. 53, pl. 3, figs. 6-7

Superfamily Asterigerinacea d'Orbigny, 1839

Family Epistomariidae Hofker, 1954
Subfamily Epistomariinae Hofker, 1954

Genus *Pseudoeponides* Uchio, 1950

Type species: *Pseudoeponides japonicus* Uchio, in
Kawai et al., 1950

Pseudoeponides falsobeccarii Rouvillois, 1974
Pl. 35, fig. 3

Pseudoeponides falsobeccarii Rouvillois in
Rouvillois, p. 4, pl. 1, figs. 1-12

Ammonia falsobeccarii Rouvillois in Gross, p. 69

Ammonia falsobeccarii Rouvillois in Murray, p. 19,
pl. 5, figs. 14-16

Pseudoeponides falsobeccarii Rouvillois in Mojtabahid
et al., 2009, p. 188, pl. 1, fig. 2

Subfamily Nuttallidinae Saidova, 1981

Genus *Nuttallides* Finlay, 1939

- Type species: *Eponides truempyi* Nuttall, 1930
- Nuttallides umbonifera* (Rouville, 1974)
Pl. 35, fig. 4
- Pulvinulinella umbonifera* Cushman in Cushman, 1933, p. 90, pl. 9, fig. 9
Nuttallides umbonifera Cushman in Hermelin, 1989, p. 69, pl. 12, figs. 15-17
- Nuttallides decorata* (Phleger and Parker, 1951)
Pl. 35, fig. 5
- Pseudoparrella? decorata* Phleger and Parker in Phleger and Parker, 1951, p. 28, pl. 15, figs. 4-5
Nuttallides decorata Phleger and Parker in Lobegeier and Gupta, 2008, p. 106, pl. 2, figs. 10, 13
- Family Astigerinatidae Reiss, 1963
- Genus *Astigerinata* Bermudez, 1949
Type species: *Astigerinata dominicana* Bermudez, 1949
- Astigerinata mamilla* (Williamson, 1858)
Pl. 36, fig. 1
- Rotalina mamilla* Williamson in Williamson, 1858, p. 54, pl. 4, figs. 109-111
Astigerinata mamilla Williamson in Cimerman and Langer, p. 73, pl. 82, figs. 1-4
Astigerinata mamilla Williamson in Abu-Zied et al., p. 53, pl. 3, figs. 8-9
- Superfamily Nonionacea Schultze, 1854
Family Nonionidae Schultze, 1854
Subfamily Nonioninae Schultze, 1854
- Genus *Nonion* de Montfort, 1808
Type species: *Nautilus faba* Fichtel and Moll, 1798, p. 103
- Nonion fabum* (Fichtel and Moll, 1798)
Pl. 36, fig. 2
- Nautilus faba* Fichtel and Moll in Fichtel and Moll, 1798, p. 103, pl. 19, figs. a-c
Nonionina boueana d'Orbigny in Brady, 1884, p. 729, pl. 109, figs. 12-13
Nonion fabum Fichtel and Moll in Loeblich and Tappan, 1987, p. 617, pl. 690, figs. 1-7 and 14-16
Nonion fabum Fichtel and Moll in Jones, p. 108, pl.
- 109, figs. 12-13
- Nonion pauperatus* (Balkwill and Wright, 1885)
Pl. 36, fig. 3
- Nonionina pauperata* Balkwill and Wright in Balkwill and Wright, 1885, p. 353, pl. 13, figs. 25-26
Nonion pauperatus Balkwill and Wright in Murray, 2003, p. 24, pl. 9, fig. 1
- Nonion pauciloculum* Cushman, 1944
Pl. 36, fig. 5
- Nonion pauciloculum* Cushman in Cushman, 1944, p. 24, pl. 3, fig. 25
- Genus *Nonionella* Cushman, 1926
Type species: *Nonionella miocenica* Cushman, 1926, p. 64
- Nonionella turgida* (Williamson, 1858)
Pl. 37, fig. 1
- Rotalina turgida* Williamson in Williamson, 1858, p. 50, pl. 4, figs. 95-97
Nonionella turgida Williamson in Murray, 2003, p. 24, pl. 9, figs. 4-5
Nonionella turgida Williamson in Jones, 1994, p. 108, pl. 109, figs. 17-19
Nonionella turgida Williamson in Cimerman and Langer, 1991, p. 74, pl. 84, figs. 6-8
- Nonionella iridea* Heron-Allen and Earland, 1932
Pl. 37, fig. 2
- Nonionella iridea* Heron-Allen and Earland in Heron-Allen and Earland, 1932, p. 438, pl. 16, figs. 14-16
Nonionella iridea Heron-Allen and Earland in Murray, 2003, p. 24, pl. 9, figs. 2-3
- Genus *Nonionellina* Cushman, 1926
Type species: *Nonionina labradorica* Dawson, 1860, p. 192
- Nonionellina labradorica* (Dawson, 1860)
Pl. 36, fig. 4
- Nonionina labradorica* Dawson in Dawson, 1860, p. 192, pl. 4
Nonion labradoricum Dawson in Feyling-Hannsen, p. 262, pl. 10, figs. 1-2
Nonionellina labradorica Dawson in Vazquez Riveiros and Patterson, 2008, p. 29, pl. 12, fig. 7

- Subfamily Astrononioninae Saidova, 1981
- Genus *Astrononion* Cushman and Edwards, 1937
Type species: *Nonionina stelligera* d'Orbigny, 1839, p. 128
- Astrononion gallowayi* Loeblich and Tappan, 1953
Pl. 37, fig. 3
- Astrononion gallowayi* Loeblich and Tappan in Loeblich and Tappan, 1953, p. 90, pl. 17, figs. 4-7
- Astrononion gallowayi* Loeblich and Tappan in Feyling-Hanssen et al., 1971, p. 266, pl. 10, figs. 10-12
- Astrononion gallowayi* Loeblich and Tappan in Hermelin and Scott, 1985, p. 203, pl. 5, fig. 1
- Astrononion antarcticus* Parr, 1950
Pl. 37, fig. 4
- Nonion stelligerus* d'Orbigny in Dryglaski, 1931, pl. 19, fig. 234
- Nonion stelliger* d'Orbigny in Earland, 1934, p. 189
- Astrononion antarcticus* Parr in Parr, 1950, p. 371, pl. 15, figs. 13-14
- Subfamily Pulleniinae Schwager, 1877
- Genus *Melonis* de Montfort, 1808
Type species: *Nautilus pompilioides* Fichtel and Moll, 1798, p. 31
- Melonis pompilioides* (Fichtel and Moll, 1798)
Pl. 37, fig. 5
- Nautilus pompilioides* Fichtel and Moll in Fichtel and Moll, 1798, p. 31, pl. 2, figs. a-c
- Melonis etruscus* de Montfort in de Montfort, 1808, p. 67
- Melonis pompilioides* Fichtel and Moll in Hermelin and Scott, 1985, p. 212, pl. 6, fig. 5
- Melonis pompilioides* Fichtel and Moll in Cimerman and Langer, 1991, p. 74, pl. 85, figs. 1-4
- Melonis barleeanum* (Williamson, 1858)
Pl. 38, fig. 1
- Nonionina barleeana* Williamson in Williamson, 1858, p. 32, pl. 3, figs. 68-69
- Nonion barleeanum* Williamson in Kihle and Løfaldli, 1975
- Nonion barleeanum* Williamson in Hermelin and Scott, 1985, p. 212, pl. 5, fig. 2
- Melonis barleeanum* Williamson in Murray, 2003, p. 24, pl. 8, figs. 11-14
- Genus *Pullenia* Parker and Jones, 1862
Type species: *Nonionina bulloides* d'Orbigny, 1846, p. 107
- Pullenia subcarinata* (d'Orbigny, 1839)
Pl. 38, fig. 2
- Nonionina subcarinata* d'Orbigny in d'Orbigny, 1839, p. 28, pl. 5, figs. 23-24
- Nonionina quinqueloba* d'Orbigny in Reuss, 1851, p. 71, pl. 5, fig. 31
- Pullenia simplex* Rhumbler in Wiesner, 1931, p. 132, pl. 22, fig. 263
- Pullenia quinqueloba* d'Orbigny in Jones, 1994, p. 92, pl. 84, figs. 14-15
- Pullenia bulloides* (d'Orbigny, 1826)
Pl. 38, figs. 3-4
- Nonionina bulloides* d'Orbigny in d'Orbigny, 1826, p. 293
- Nonionina bulloides* d'Orbigny in d'Orbigny, 1846, p. 107, pl. 5, figs. 9-10
- Pullenia sphaerooides* d'Orbigny in Brady, 1884, p. 615, pl. 84, figs. 12-13
- Pullenia bulloides* d'Orbigny in Ujiié, 1990, p. 42, pl. 23, figs. 1-2
- Pullenia bulloides* d'Orbigny in Jones, 1994, p. 92, pl. 84, figs. 12-13
- Superfamily Chilostomellacea Brady, 1881
Family Chilostomellidae Brady, 1881
Subfamily Chilostomellinae Brady, 1881
- Genus *Chilostomella* Reuss, 1849
Type species: *Chilostomella ovoidea* Reuss, 1850, p. 380
- Chilostomella oolina* Schwager, 1878
Pl. 38, fig. 5
- Chilostomella oolina* Schwager in Schwager, 1878, p. 527, pl. 1, fig. 16
- Chilostomella ovoidea* Reuss in Brady, 1884, p. 436, pl. 55, figs. 12-14, 17-18
- Chilostomella oolina* Schwager in Cushman and Todd, 1949, p. 91, pl. 15, figs 23-24
- Chilostomella oolina* Schwager in Jones, 1994, p. 61, pl. 55, figs. 12-14, 17-18

Chilostomella oolina Schwager in Loeblich and Tappan, 1994, p. 160, pl. 349, figs. 12-13

Family Heterolepidae Gonzales-Donoso, 1969

Genus *Anomalinoides* Brotzen, 1942

Type species: *Anomalina pinguis* Jennings, 1936, p. 195

Anomalinoides globulosa (Chapman and Parr, 1937)
Pl. 39, fig. 1

Anomalina globulosa Chapman and Parr in

Chapmann and Parr, 1937, p. 117, pl. 9, fig. 27

Anomalina globulosa Chapman and Parr in Feyling-Hanssen, 1954, p. 258, pl. 9, figs. 1-3

Cibicidoides globulosus Chapman and Parr in Jones, 1994, p. 98, pl. 94, fig 4-5

Family Gavelinellidae Hofker, 1956
Subfamily Gavelinellinae Hofker, 1956

Genus *Discanomalina* Asano, 1951

Type species: *Discanomalina japonica* Asano, 1951, p. 13

Discanomalina coronata (Parker and Jones, 1857)
Pl. 39, fig. 2

Anomalina coronata Parker and Jones in Parker and Jones, 1857, p. 294, pl. 10, figs. 15-16

Discanomalina coronata Jones in Jones, 1994, p. 100, pl. 97, figs. 1-2

Discanomalina coronata Parker and Jones in Schönfeld, 2003, p. 1858, pl. 1, fig. 14

Discanomalina japonica Asano, 1951
Pl. 39, fig. 3

Discanomalina japonica Asano in Asano, p. 13, figs. 3-5

Genus *Gyroidina*, 1826

Type species: *Gyroidina orbicularis* d'Orbigny, 1826, p. 190

Gyroidina laevigata d'Orbigny, 1826
Pl. 39, fig. 4

Gyroidina laevigata d'Orbigny in d'Orbigny, 1826, p. 278

Gyroidina lamarckiana (d'Orbigny, 1839)
Pl. 39, fig. 5

Rotalia lamarckiana d'Orbigny in d'Orbigny, 1839b, p. 131, pl. 2 , figs. 13-15

Gyroidina lamarckiana d'Orbigny in Phleger et al., 1953, p. 41, pl. 8, figs. 33-34

Gyroidina lamarckiana d'Orbigny in Todd, 1965, p. 19, pl. 6, fig. 3

Gyroidina lamarckiana d'Orbigny in Loeblich and Tappan, 1994, p. 163, pl . 361, figs. 7-12

Gyroidina soldanii d'Orbigny, 1839
Pl. 40, fig. 1

Gyroidina soldanii d'Orbigny in d'Orbigny, 1826, p. 278, no. 5

Rotalia soldanii d'Orbigny in d'Orbigny, 1846, p. 155, pl. 8, figs. 10-12

Gyroidina soldanii d'Orbigny in Papp and Schmid, 1985, p. 60, pl. 50, figs. 4-9

Hansenisca soldanii d'Orbigny in Loeblich and Tappan, 1987, p. 639, pl. 719, figs. 5-9

Gyroidina neosoldanii Brotzen, 1936
Pl. 40, fig. 2

Rotalina soldanii d'Orbigny in Brady, 1884, p. 107, figs. 6-7

Gyroidina neosoldanii Brotzen in Brotzen, 1936, p. 158

Gyroidinoides soldanii Brotzen in Jones, 1994, p. 106, pl. 107, figs. 6-7

Gyroidina neosoldanii Brotzen in Hermelin, 1989, p. 81, pl. 15, figs. 16-18

Gyroidina altiformis (Steward and Steward, 1930)
Pl. 40, fig. 3

Gyroidina soldanii d'Orbigny var. *altiformis* Steward and Steward in Cushman et al., 1930, p. 67, pl. 9, fig. 2

Gyroidina soldanii d'Orbigny var. *altiformis* Steward and Steward in Cushman, 1931, p. 41, pl. 8, fig. 10; pl. 9, fig. 1

Gyroidina soldanii d'Orbigny var. *altiformis* Steward and Steward in Renz, 1948, p. 140, pl. 8, fig. 13

Hansenisca altiformis Steward and Steward in Finger, 1990, p. 124-125, figs 1-8; text-fig. 2

Genus *Hanzawaia* Asano, 1944
Type species: *Hanzawaia nipponica* Asano, 1944, p. 98

- Hanzawaia boueana* (d'Orbigny, 1846)
Pl. 40, figs. 4-5
- Truncatulina boueana* d'Orbigny in d'Orbigny, 1846, p. 169, pl. 9, figs. 24-26
- Cibicides boueanus* d'Orbigny in Graham and Militante, 1959, p. 116, pl. 19, fig. 11
- Hanzawaia boueana* d'Orbigny in Zheng, 1980, p. 171, pl. 5, fig. 10
- Hanzawaia boueana* d'Orbigny in Loeblich and Tappan, 1994, p. 164, pl. 364, figs 1-8
- Family Trichohyalidae Saidova, 1981
- Genus *Buccella* Andersen, 1952
Type species: *Eponides hannai* Phleger and F.L. Parker, 1951, p. 21
- Buccella frigida* (Cushman, 1922)
Pl. 41, fig. 1
- Pulvinulina frigida* Cushman in Cushman, 1922, p. 144
- Buccella frigida* Cushman in Feyling – Hanssen et al., 1971, p. 253, pl. 8, figs. 12-14, pl. 19, fig. 1
- Buccella frigida* Cushman in Kihle and Løfaldli, 1975
- Buccella frigida* Cushman in Vazquez Riveiros and Patterson, 2008, p. 30, pl. 13, fig. 3
- Superfamily Orbitoidacea Schwager, 1876
- Family Elphidiinae Galloway, 1933
- Genus *Elphidium* de Montfort, 1808
Type species: *Nautilus macellus* var. β Fichtel and Moll, 1798, p. 66
- Elphidium albiumbilicatum* (Weiss, 1954)
Pl. 41, fig. 2
- Nonion pauciloculum* Cushman subsp. *albiumbilicatum* Weiss in Weiss, 1954, p. 157, pl. 32, figs. 1-2
- Elphidium albiumbilicatum* Weiss in Feyling – Hanssen et al., p. 268, pl. 10, figs. 15-19, pl. 19, figs. 4-8
- Elphidium albiumbilicatum* Weiss in Murray et al., 2003, p. 687, textfig. 6
- Elphidium groenlandicum* Cushman, 1933
Pl. 41, fig. 3
- Elphidium groenlandicum* Cushman in Cushman, 1933, p. 4, pl. 1, fig. 10
- Elphidium groenlandicum* Cushman in Feyling – Hanssen et al., 1971, p. 275, pl. 12, figs. 1-8, pl. 21, figs. 1-3
- Elphidium groenlandicum* Cushman in Kihle and Løfaldli, 1975
- Elphidium hanzawai* Asano, 1939
Pl. 41, fig. 4
- Elphidium hanzawai* Asano in Asano, 1939, p. 426, figs. 3-4
- Elphidium incertum* (Williamson, 1858)
Pl. 42, fig. 1
- Polystomella umbilicata* Walker var. *incerta* Williamson in Williamson, 1858, p. 44, pl. 3, fig. 82a
- Polystomella striatopunctata* Brady in Brady, 1884, p. 739, pl. 109, fig. 23
- Elphidium incertum* Williamson in Feyling – Hanssen et al., 1971, p. 277, pl. 12, figs. 11-12, pl. 21, figs. 8-9
- Elphidium incertum* Williamson in Kihle and Løfaldli, 1975
- Cribrononion incertum* Williamson in Jones, 1994, p. 108, pl. 109, fig. 23
- Elphidium magellanicum* Heron-Allen and Earland, 1932
Pl. 42, fig. 2
- Elphidium (Polystomella) magellanicum* Heron-Allen and Earland in Heron-Allen and Earland, p. 440, pl. 16, figs. 26-28
- Elphidium magellanicum* Heron-Allen and Earland in Kihle and Løfaldli, 1975
- Elphidium magellanicum* Heron-Allen and Earland in Murray, 2003, p. 21, pl. 7, figs. 9-10
- Criboelphidium magellanicum* Heron-Allen and Earland in Vazquez Riveiros and Patterson, p. 34, pl. 15, fig. 4
- Elphidium subarcticum* Cushman, 1944
Pl. 42, fig. 3
- Elphidium subarcticum* Cushman in Cushman, 1944, p. 27, pl. 3, figs. 34-35
- Elphidium subarcticum* Cushman in Kihle and Løfaldli, 1975
- Elphidium subarcticum* Cushman in Scott et al., 2008, p. 244, pl. 4, fig. 5

PLATES

to be submitted to Earth-Science Reviews

PLATE 1

- 1a-b *Astrorhiza* cf. *catenata* Norman, 1877, Hypotype from PS70/038-2
2a-b *Bathysiphon filiformis* M. Sars, 1872, Hypotype from PS70/038-2
3a-b *Rhabdammina abyssorum* M. Sars, 1869, a) Hypotype from AL232 1026; b) Hypotype from PS70/039-2
4a-b *Hippocrepinella hirudinea* Heron-Allen and Earland, 1932, Hypotype from PS70/033-2
5a-b *Psammosphaera fusca* Schulze, 1875, Hypotype from PS70/039-2
6a-b *Psammosphaera fusca* Schulze var. *testacea* Flint, 1899, Hypotype from PS70/037-2
7a-b *Lagenammina fusiformis* (Williamson, 1858), Hypotype from PS70/037-2
8a-b *Lagenammina arenulata* (Williamson, 1858), Hypotype from PS70/039-2
9a-b *Saccammina sphaerica* M. Sars, 1872, Hypotype from AL232 1025
10a-b *Hyperammina elongata* Brady, 1878, Hypotype from POS391 558-1
11a-b *Saccorrhiza ramosa* (Brady, 1879), Hypotype from PS70/028-2

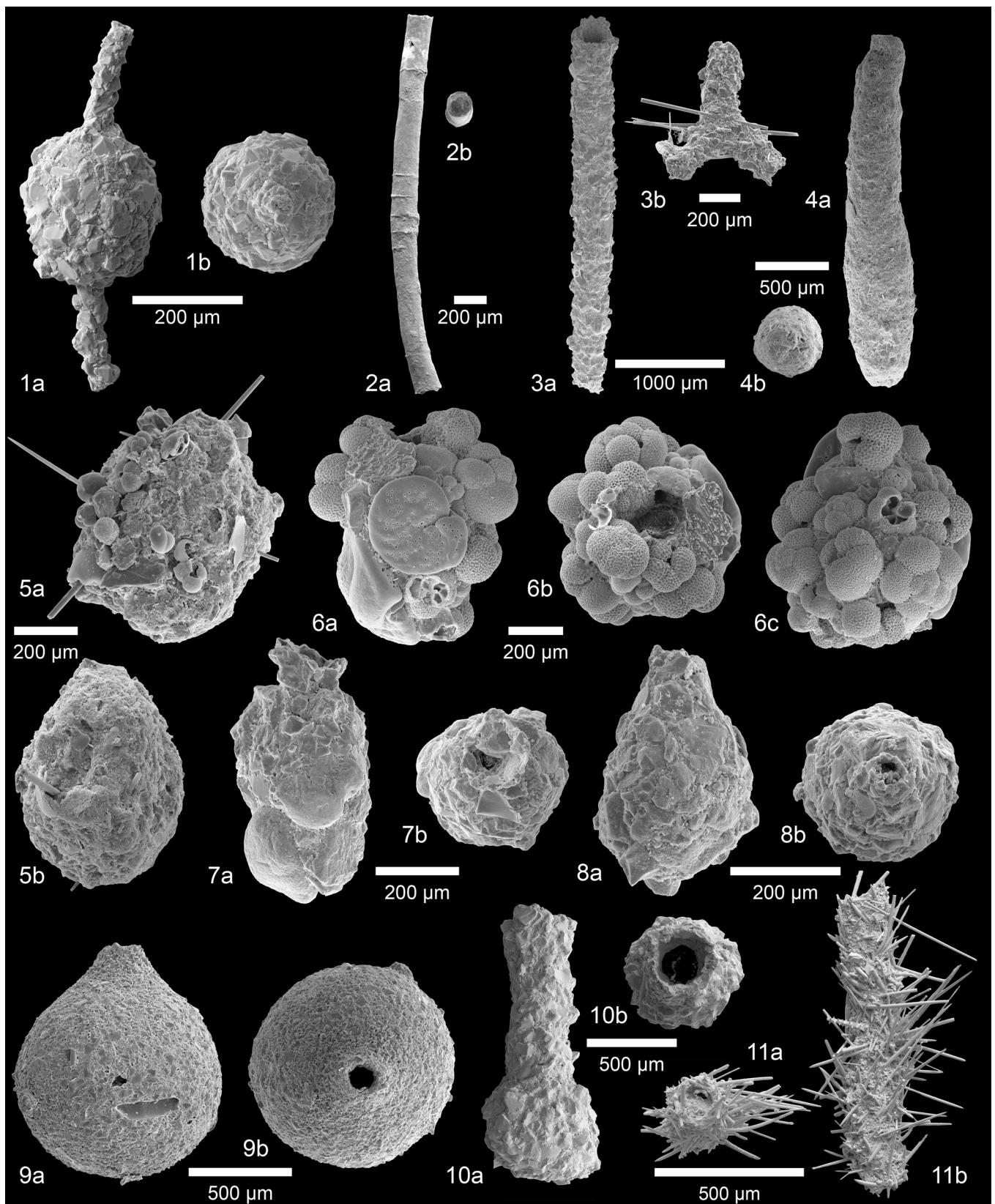


PLATE 2

- 1a-b *Reophax agglutinatus* Cushman, 1913, Hypotype from PS70/037-2
2a-b *Hormosinella guttifera* (Brady, 1881), Hypotype from PS70/039-2
3a-b *Reophax scorpiurus* de Montfort, 1808, Hypotype from P292 577-1
4a-b *Ammodiscus incertus* (d'Orbigny, 1839), Hypotype from PS70/011-1
5a-c *Glomospira charoides* (Jones and Parker, 1878), Hypotype from PS70/011-1
6a-b *Cribrostomoides subglobosum* (M. Sars, 1868), Hypotype from PS70/011-1
7 *Ammolagena clavata* (Jones and Parker, 1860), Hypotype from PS70/002-2
8a-c *Labrospira jeffreysii* (Williamson, 1858), Hypotype from PS70/002-2
9a-c *Labrospira jeffreysii* (Williamson, 1858), Hypotype from AL232 1025

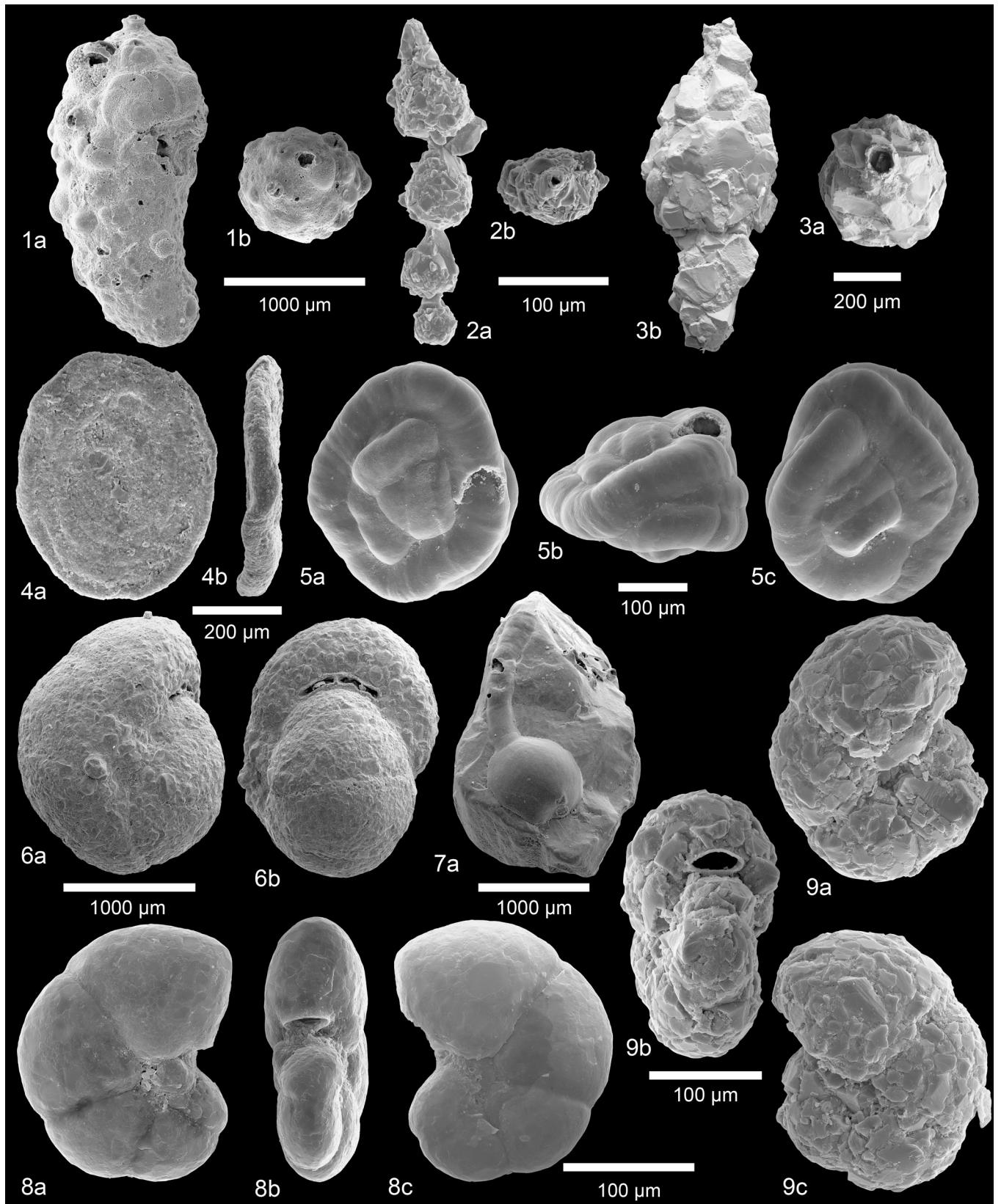


PLATE 3

- 1a-c *Haplophragmoides robertsoni* Brady, 1887, Hypotype from POS391 550-1
2a-c *Haplophragmoides robertsoni* Brady, 1887, Hypotype from PS70/011-1
3a-c *Haplophragmoides membranaceum* Höglund, 1947, Hypotype from PS70/029-3
4a-b *Ammobaculites agglutinans* (d'Orbigny, 1846), Hypotype from PS70/032-2
5a-c *Adercotryma wrighti* Brönnimann and Whittaker, 1987, Hypotype from AL232 1022
6a-c *Spiroplectinella wrightii* (Silvestri, 1903), Hypotype from POS391 550-1

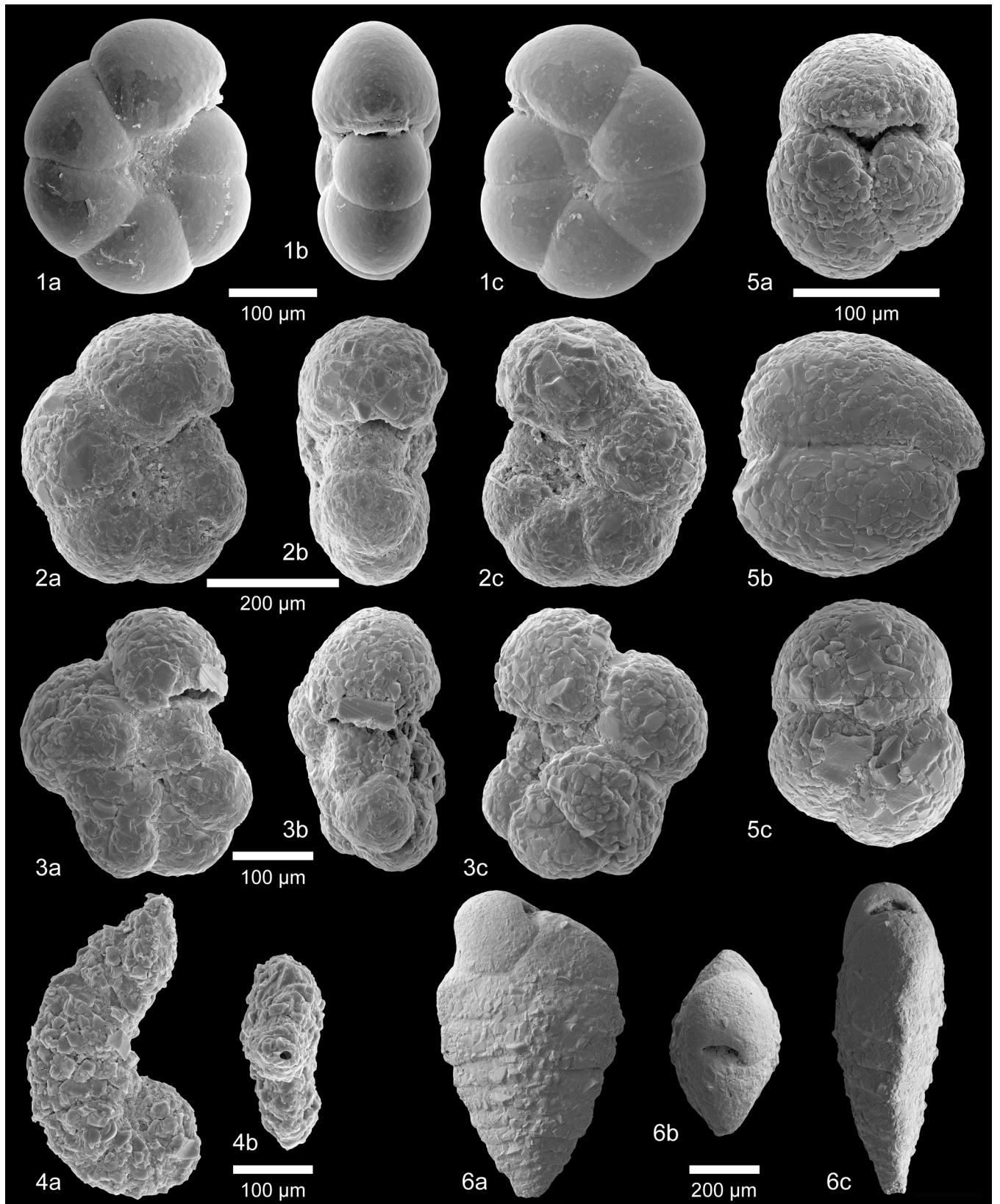


PLATE 4

- 1a-c *Vulvulina pennatula* Batsch, 1791, Hypotype from TTR17 MS411G 15
- 2a-b *Trochammina labiosa* Höglund, 1947, Hypotype from POS391 571-1
- 3a-c *Tritaxis fusca* (Williamson, 1858), P292 580-1
- 4a-c *Portatrocchammina antarctica* (Parr, 1950), Hypotype from PS70/002-2
- 5a-c *Lepidodeuterammina ochracea* (Williamson, 1858), POS325 455
- 6a-c *Gaudryina rufa* Wright, 1900, Hypotype from AL232 1025

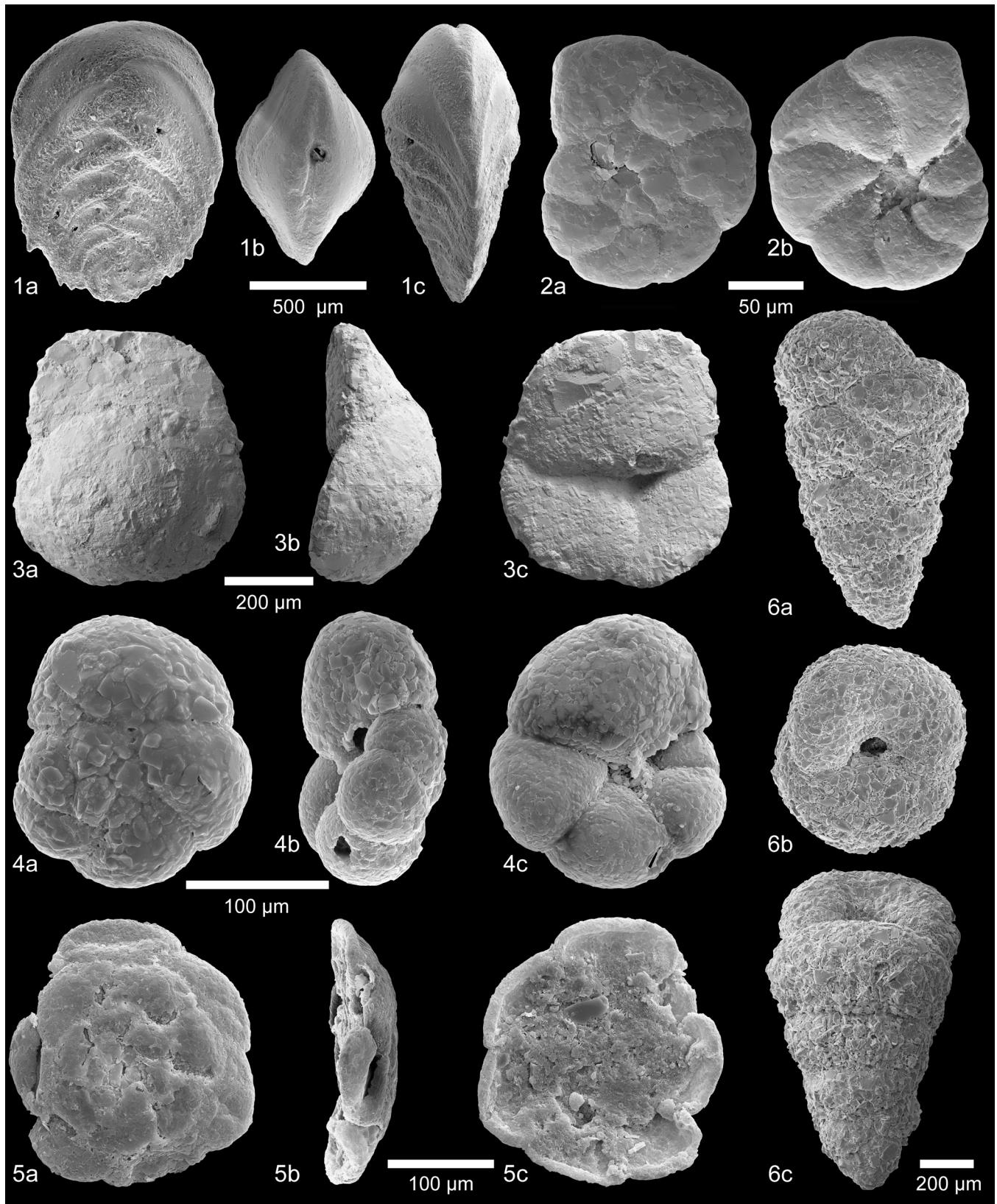


PLATE 5

- 1a-c *Gaudryina pseudotrochus* (Cushman, 1922), Hypotype from GeoB 9204-1
2a-b *Eggerella humboldti* Todd and Brönnimann, 1957, Hypotype from M07-21
3a-c *Karreriella bradyi* (Cushman, 1911), Hypotype from M07-23
4a-b *Eggerelloides scaber* (Williamson, 1858), Hypotype from AL232 1025
5a-c *Bigenerina nodosaria* d'Orbigny, 1826, Hypotype from P292 577-1
6a-b *Bigenerina cylindrica* Cushman, 1922, Hypotype from AL232 1025
7a-c *Siphonostularia obesa* Parr, 1950, Hypotype from P292 578-1
8a-b *Textularia lateralis* Laliker, 1935, TTR-17 MS411G 20

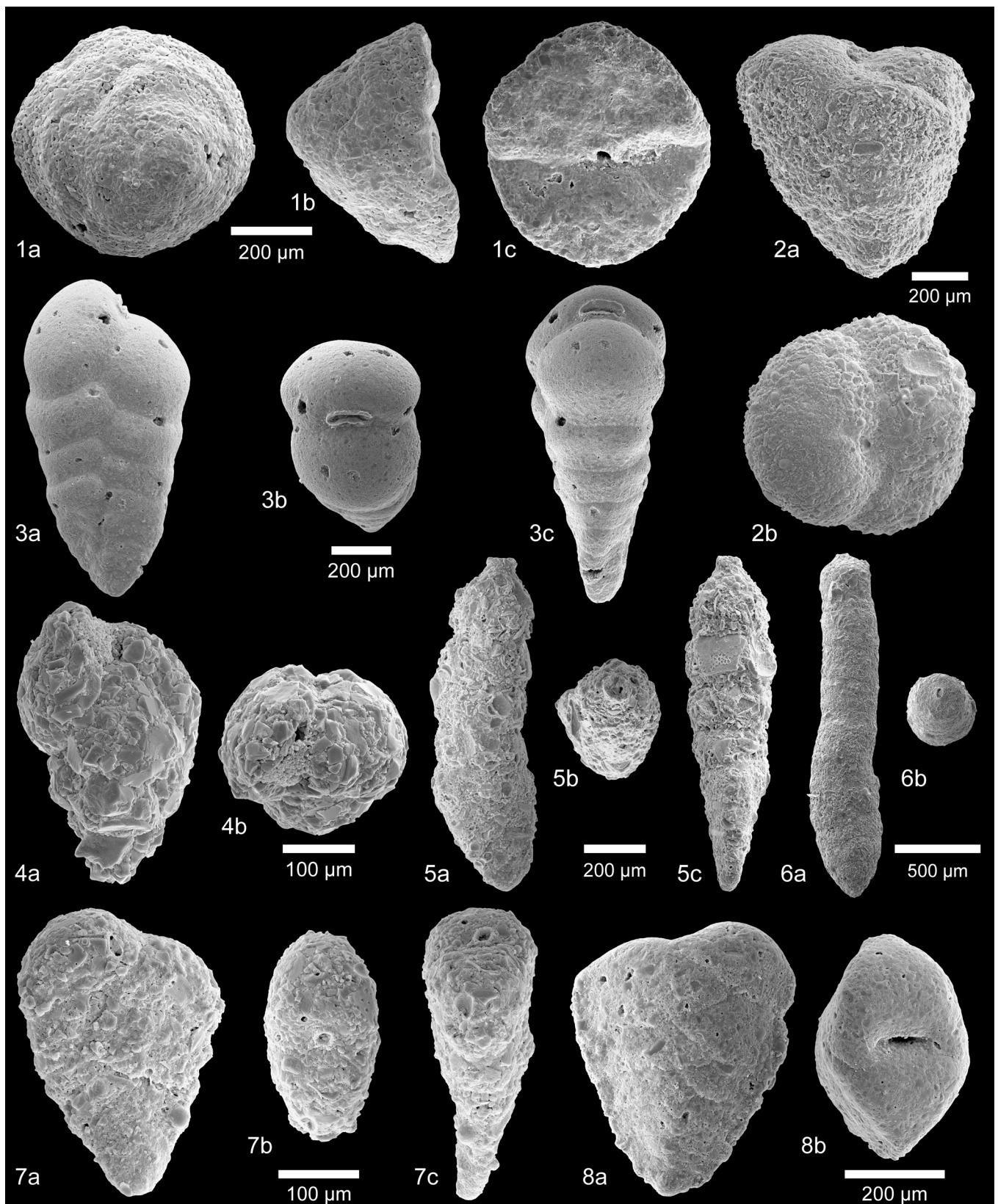


PLATE 6

- 1a-c *Textularia truncata* Höglund, 1947, Hypotype from TTR17 MS411G 25
2a-c *Textularia tenuissima* Earland, 1933, Hypotype from AL232 1025
3a-b *Clavulina parisiensis* d'Orbigny, 1826, Hypotype from TTR17 MS411G 0
4a-c *Mychostomina revertens* (Rhumbler, 1906), Hypotype from PS70/002-2
5a-c *Spirillina vivipara* Ehrenberg, 1843, Hypotype from POS391 550-1
6a-b *Patellina corrugata* Williamson, 1858, Hypotype from POS391 550-1
7a-c *Cornuspira involvens* (Reuss, 1850), Hypotype from PS70/011-1
8a-b *Cornuspira foliacea* (Philippi, 1844), Hypotype from TTR17 MS411G 15
9a-b *Gordiospira* sp., Hypotype from PS70/028-2
10a-b *Gordiospira elongata* Collins, 1958, Hypotype from TTR17 MS411G45

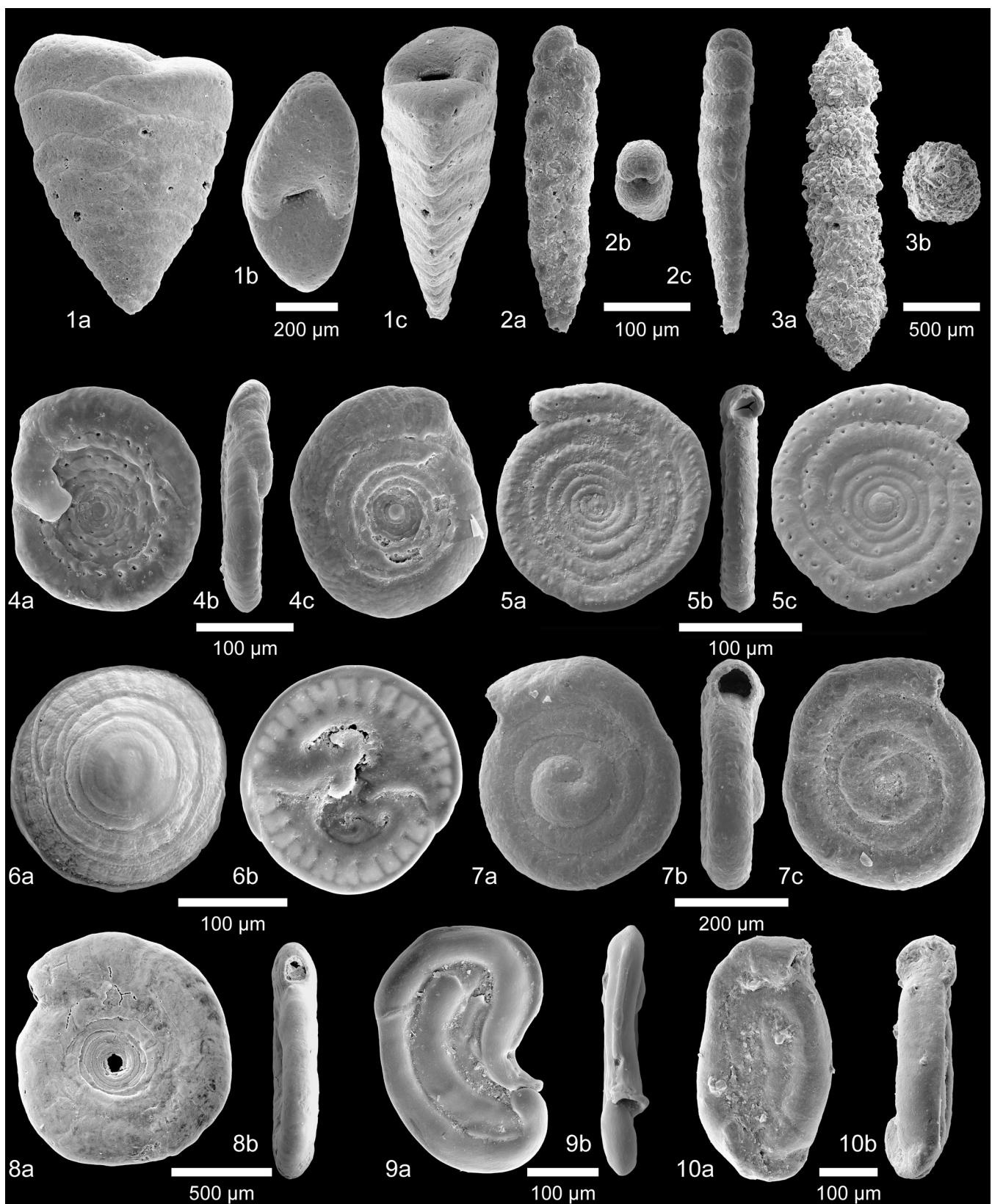


PLATE 7

- 1a-b *Spiroloculina tenuiseptata* Brady, 1884, Hypotype from TTR17 MS411G 30
2a-b *Spiroloculina dilatata* d'Orbigny, 1846, Hypotype from TTR17 MS419G 94
3a-b *Spiroloculina excavata* d'Orbigny, 1846, Hypotype from TTR17 MS411G 10
4a-b *Ammomassilina arenaria* (Brady, 1884), Hypotype from TTR17 MS411G 30
5a-c *Cycloforina laevigata* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 0
6a-c *Cycloforina stalkeri* (Loeblich and Tappan, 1953), Hypotype from TTR17 MS411G 10
7a-c *Quinqueloculina viennensis* Le Calvez and Le Calvez, 1958, Hypotype from POS391 556-2
8a-c *Quinqueloculina seminula* (Linné, 1758), Hypotype from TTR17 MS411G 15

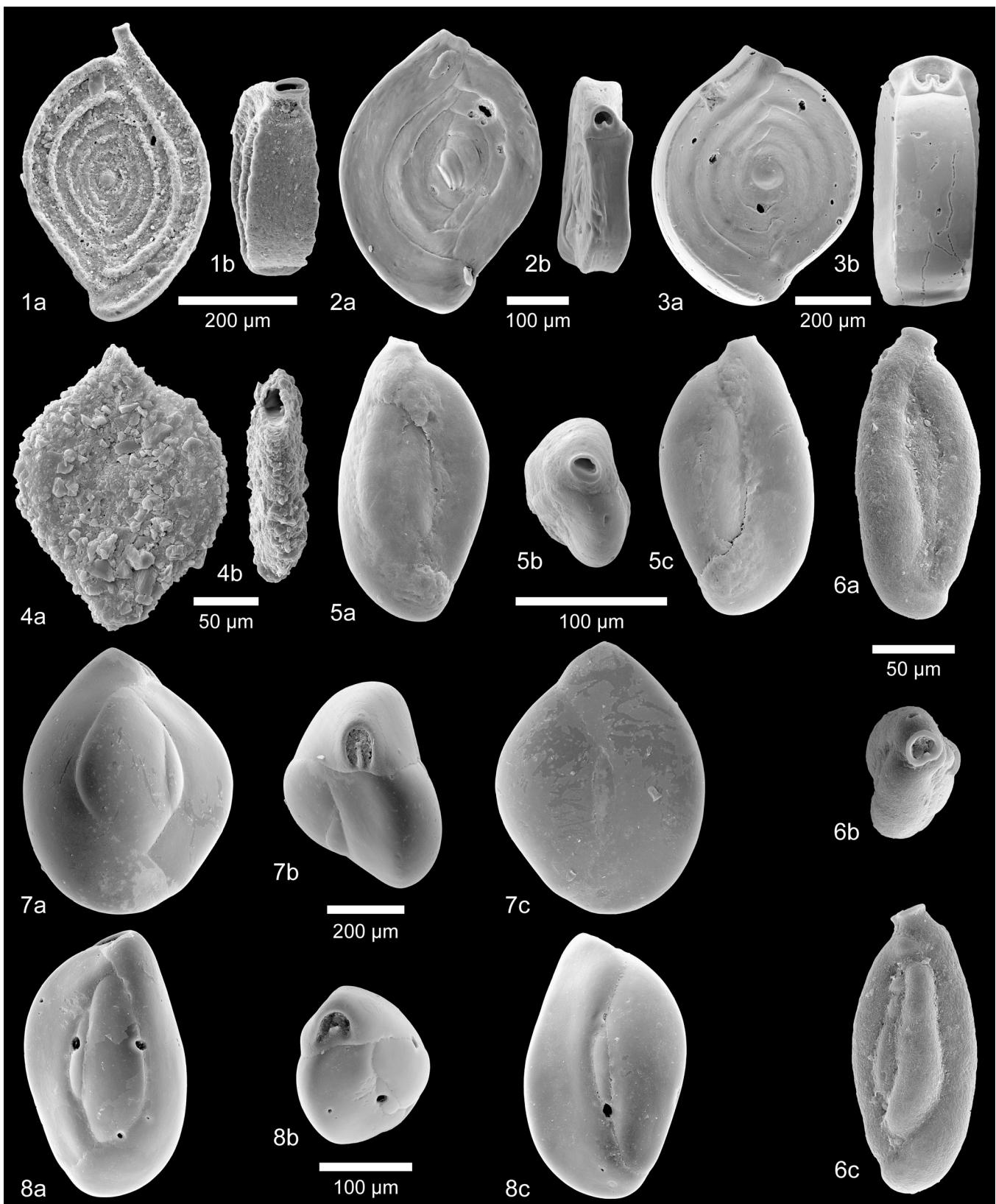


PLATE 8

- 1a-c *Quinqueloculina arctica* Cushman, 1933, Hypotype from GeoB9257
2a-c *Biloculinella globula* (Bornemann, 1855), Hypotype from POS391 534-1
3a-c *Biloculinella depressa* (Wiesner, 1923), Hypotype from POS325 455
4a-c *Biloculinella fragilis* Le Calvez and Le Calvez, 1958, Hypotype from TTR17 MS411G 45
5a-c *Miliolinella subrotunda* (Montagu, 1803), Hypotype from POS325 455

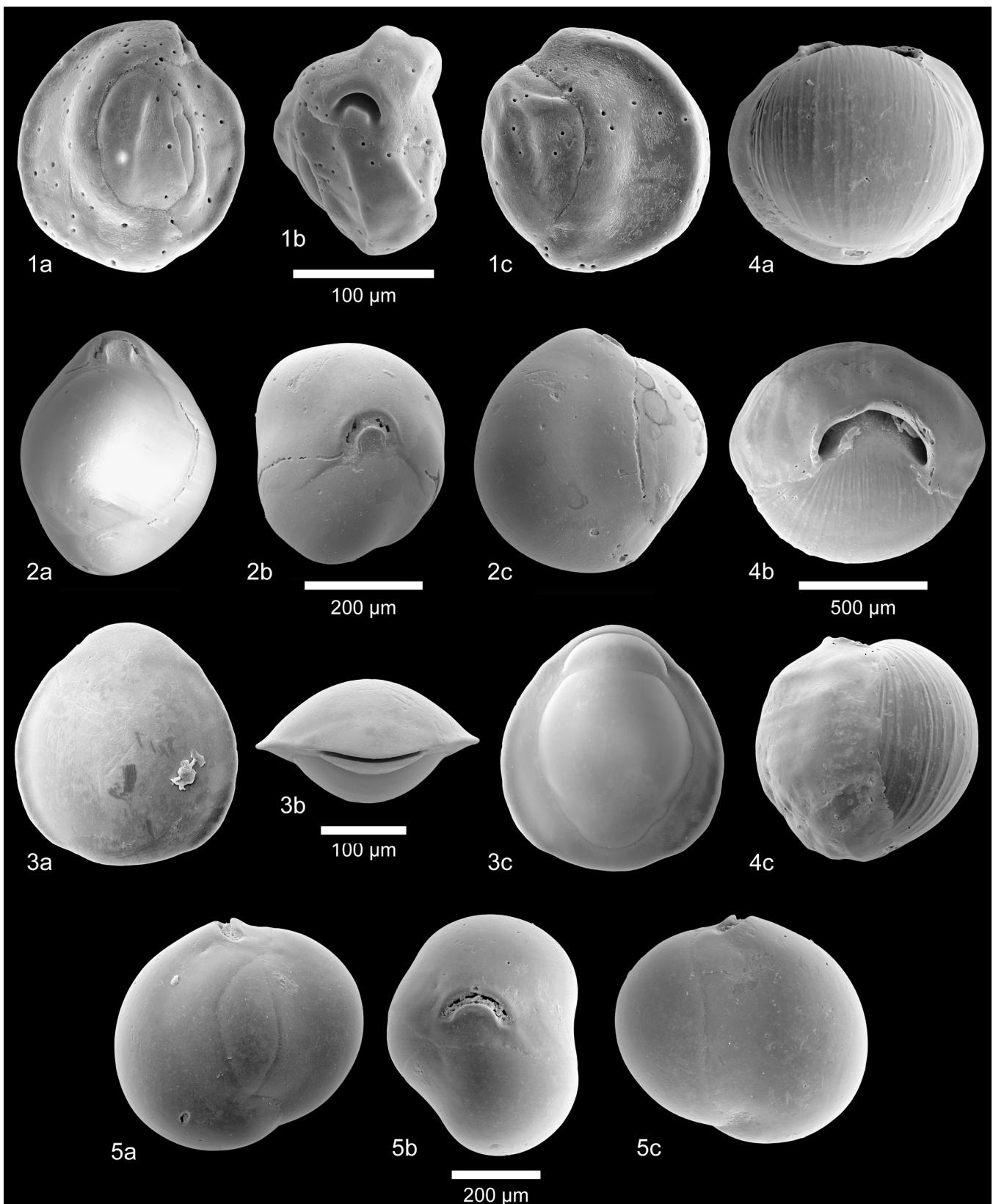


PLATE 9

- 1a-c *Miliolinella elongata* Kruit, 1955, Hypotype from TTR17 MS411G 50
2a-c *Pyrgo inornata* (d'Orbigny, 1846), Hypotype from TTR17 MS411G 55
3a-c *Pyrgo comata* (Brady, 1881), Hypotype from Hermi-1_1
4a-c *Pyrgo anomala* (Schlumberger, 1891), Hypotype from M07-15
5a-c *Pyrgo subsphaerica* (d'Orbigny, 1840), Hypotype from POS391 559-1
6a-c *Pyrgo williamsoni* (Silvestri, 1923), Hypotype from AL232 1022

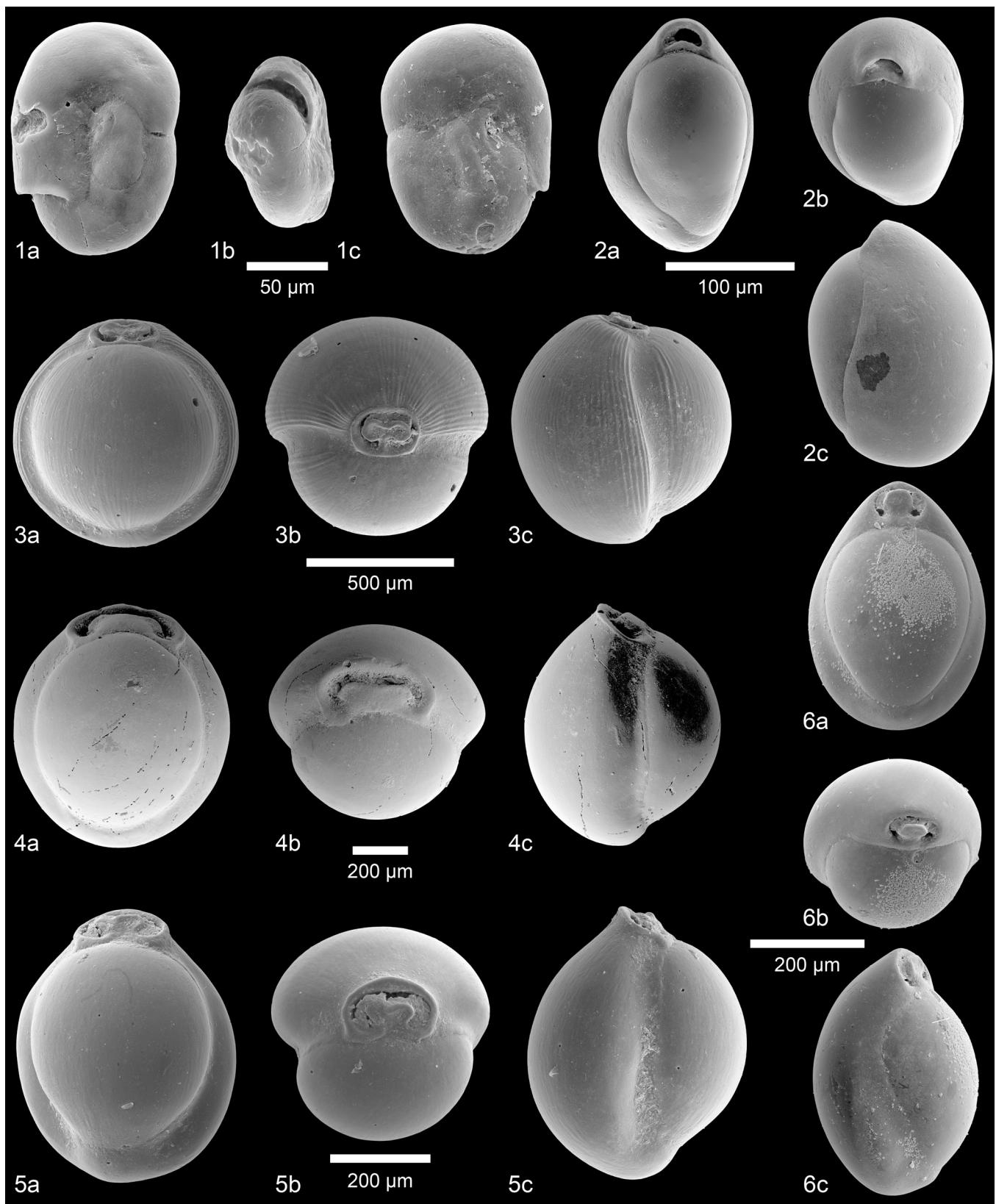


PLATE 10

- 1a-c *Pyrgo sarsi* Schlumberger, 1891, Hypotype from PS70/032-2
2a-c *Pyrgo sarsi* Schlumberger, 1891, Hypotype from PS70/032-2
3a-c *Pyrgo murrhina* (Schwager, 1866), Hypotype from PS70/032-2
4a-c *Pyrgo lucernula* (Schwager, 1866), Hypotype from POS391 562-1
5a-c *Pyrgo elongata* (d'Orbigny, 1826), Hypotype from TTR17 MS411G 15

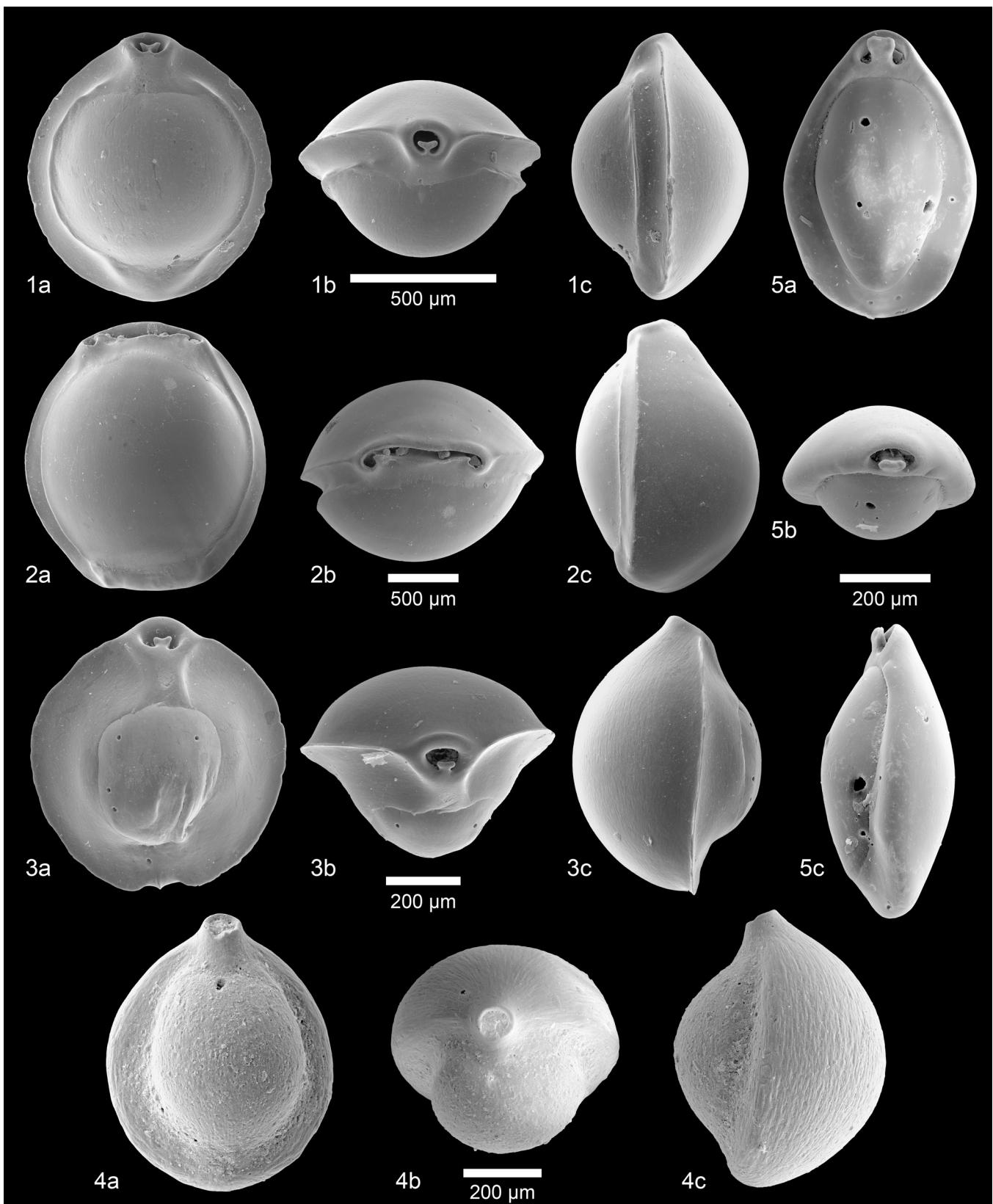


PLATE 11

- 1a-c *Triloculina marioni* Schlumberger, 1893, Hypotype from TTR17 MS411G 15
2a-c *Triloculina trigonula* (Lamarck, 1804), Hypotype from PS70/032-2
3a-c *Triloculina tricarinata* d'Orbigny, 1826, Hypotype from TTR17 MS411G 15
4a-c *Sigmoinella borealis* Saidova, 1975, Hypotype from TTR17 MS411G 25
5a-c *Sigmoilopsis schlumbergeri* (Silvestri, 1904), Hypotype from TTR17 MS411G 0
6a-b *Sigmoilopsis woodi* Atkinson, 1968, Hypotype from POS391 570-2

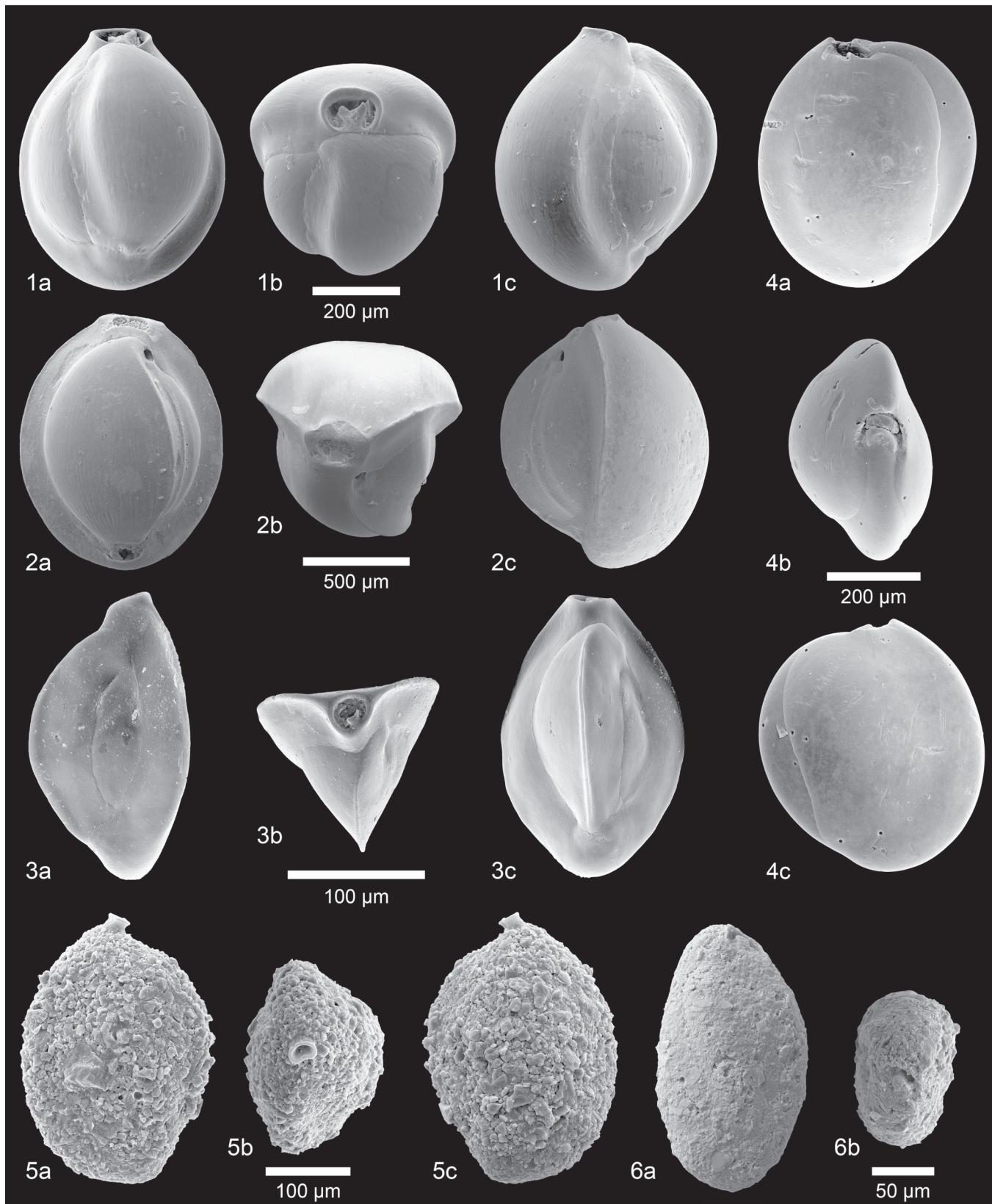


PLATE 12

- 1a-b *Dentalina cuvieri* (d'Orbigny, 1826), Hypotype from P292 574-1
2a-b *Dentalina lamarcki* Neugeboren, 1856, Hypotype from TTR17 MS411G 30
3a-b *Grigelis orectus* (Loeblich and Tappan, 1994), Hypotype from TTR17 MS419G 65
4a-b *Laevidentalina sidebottomi* (Cushman, 1933), Hypotype from TTR17 MS411G 0
5a-b *Lenticulina calcar* (Linné, 1758), Hypotype from TTR17 MS411G 50
6a-b *Lenticulina vortex* (Fichtel and Moll, 1798), Hypotype from TTR17 MS411G 10
7a-b *Lenticulina gibba* (d'Orbigny, 1826), Hypotype from TTR17 MS411G 10
8a-b *Lenticulina orbicularis* (d'Orbigny, 1846), Hypotype from GeoB 6721-1
9a-b *Lenticulina inornata* (d'Orbigny, 1846), Hypotype from M07-23
10a-c *Neolenticulina peregrina* (Schwager, 1866), Hypotype from GeoB 12721-1

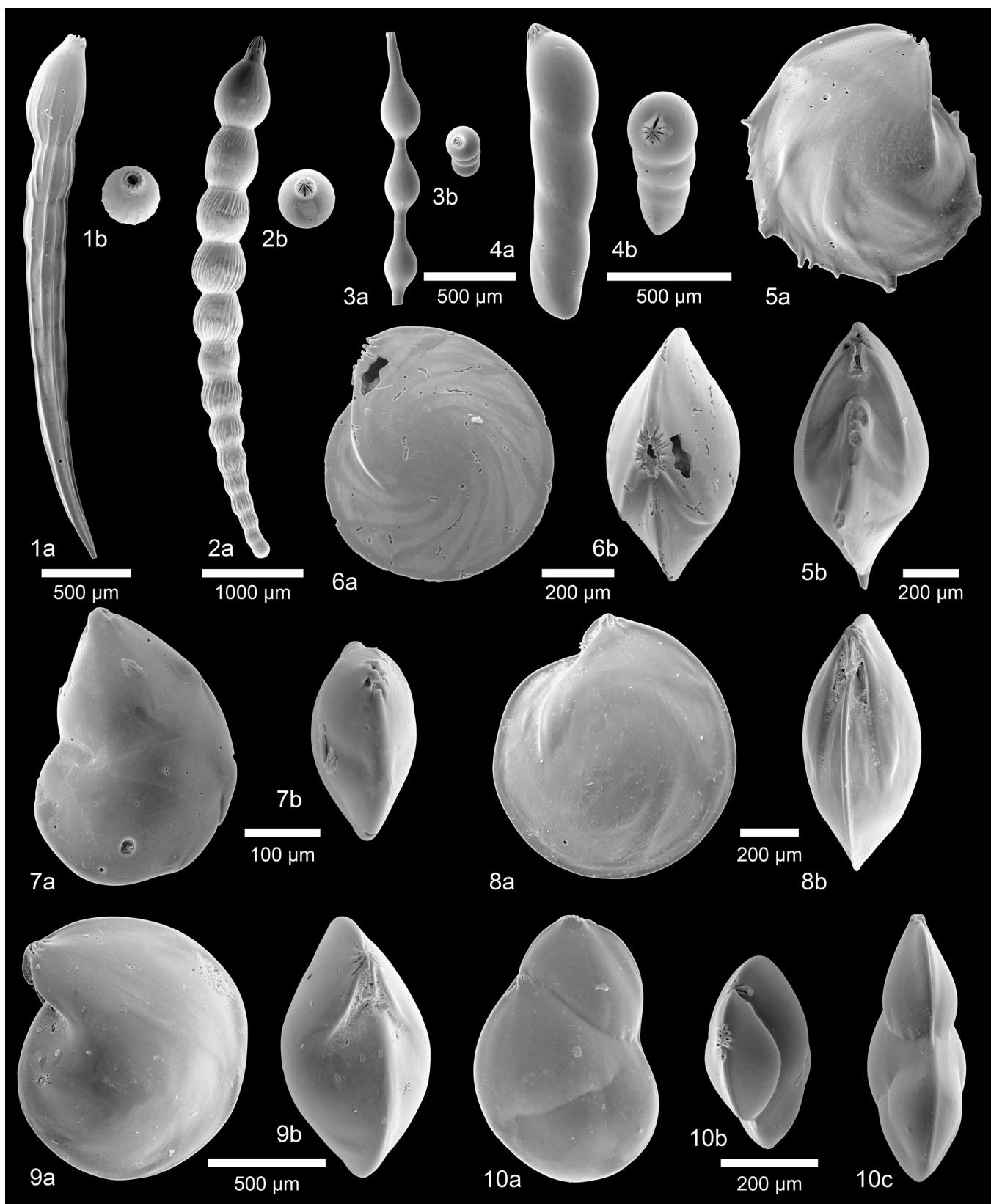


PLATE 13

- 1a-b *Saracenaria caribbeana* Hofker, 1976, Hypotype from GeoB 12729-1
2a-b *Amphicoryna scalaris* (Batsch, 1791), Hypotype from TTR17 MS411G 0
3a-c *Amphicoryna scalaris* (Batsch, 1791), Hypotype from TTR17 MS411G 10
4a-c *Astacolus beerae* Brenner and McMillan, 1976, Hypotype from PS70/028-2
5a-b *Planularia perculata* McCulloch, 1977, Hypotype from TTR17 MS419G 35
6a-c *Lagena substriata* Williamson, 1848, Hypotype from PS70/033-2
7a-b *Planularia costata* (d'Orbigny, 1902), Hypotype from GeoB12722-1
8a-b *Lagena meridionalis* Wiesner, 1931, Hypotype from TTR17 MS411G 55
9a-b *Lagena semilineata* var. *spinigera* Earland, 1934, Hypotype from POS391 556-1
10a-c *Pygmaeoseistron laevis ovalis* (Williamson, 1848), Hypotype from M07-21
11a-b *Lagena squamosoalata* Brady, 1881, Hypotype from GeoB 9271
12a-b *Lagena trigonolaevigata* Balkwill and Millett, 1884, Hypotype from PS70/029-3

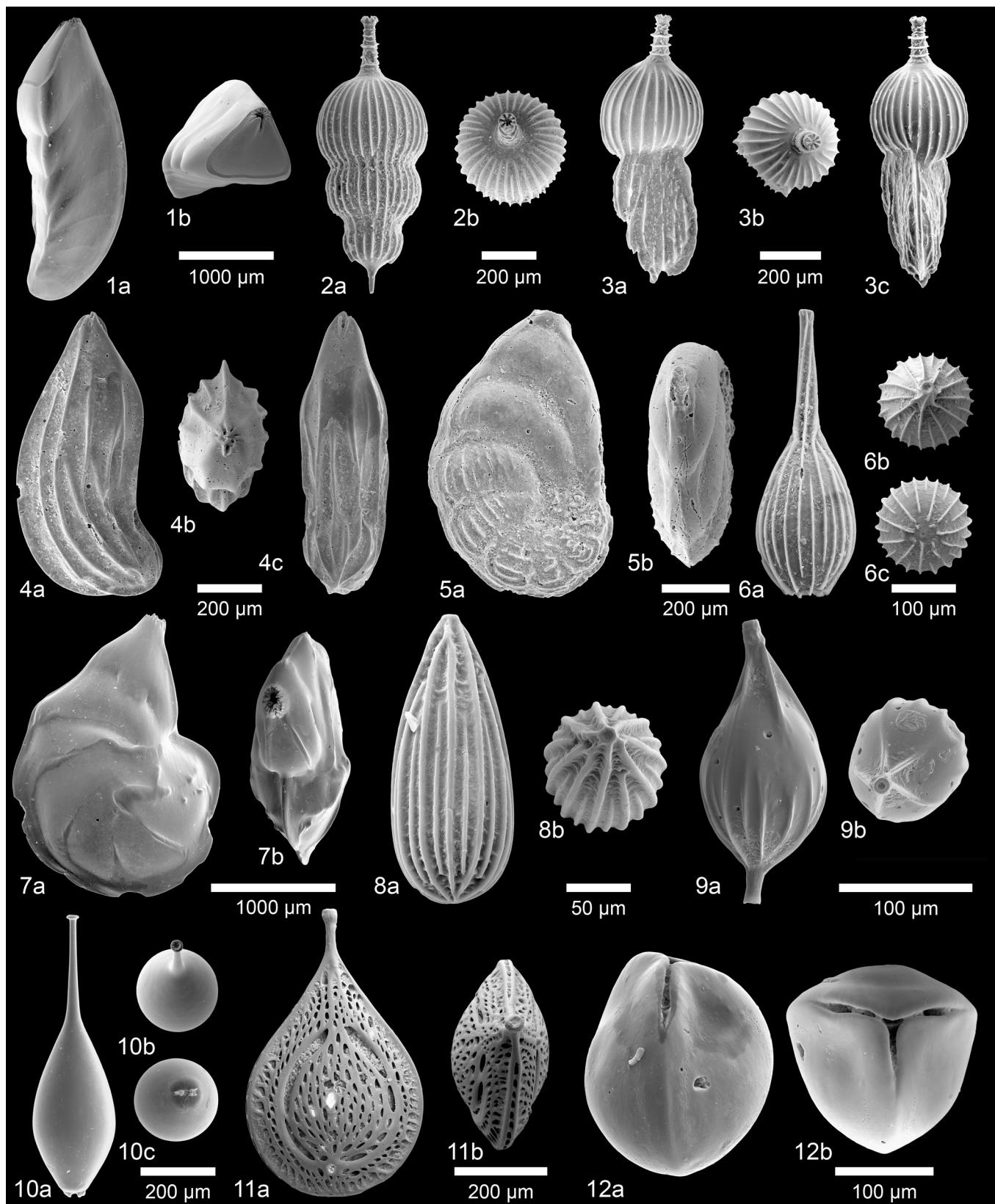


PLATE 14

- 1a-b *Hyalinonetrion gracillimum* (Costa, 1856), Hypotype from PS70/011-1
2a-b *Globulina aequalis* d'Orbigny, 1846, Hypotype from AL232 1022
3a-b *Pseudopolymorphina* sp., Hypotype from GeoB 9219
4a-c *Pyrulina cylindroids* (Roemer, 1930), Hypotype from POS391 550-1
5a-c *Favulina squamosa* (Montagu, 1803), Hypotype from TTR17 MS419G 65
6a-b *Favulina melo* (d'Orbigny, 1839), Hypotype from TTR17 MS419G 35
7a-b *Favulina hexagona* (Williamson, 1858), Hypotype from TTR17 MS411G 0
8a-b *Homalohedra williamsoni* (Alcock, 1865), Hypotype from POS391 550-1
9a-c *Homalohedra apiopleura* (Loeblich and Tappan, 1953), Hypotype from TTR17 MS411G 45

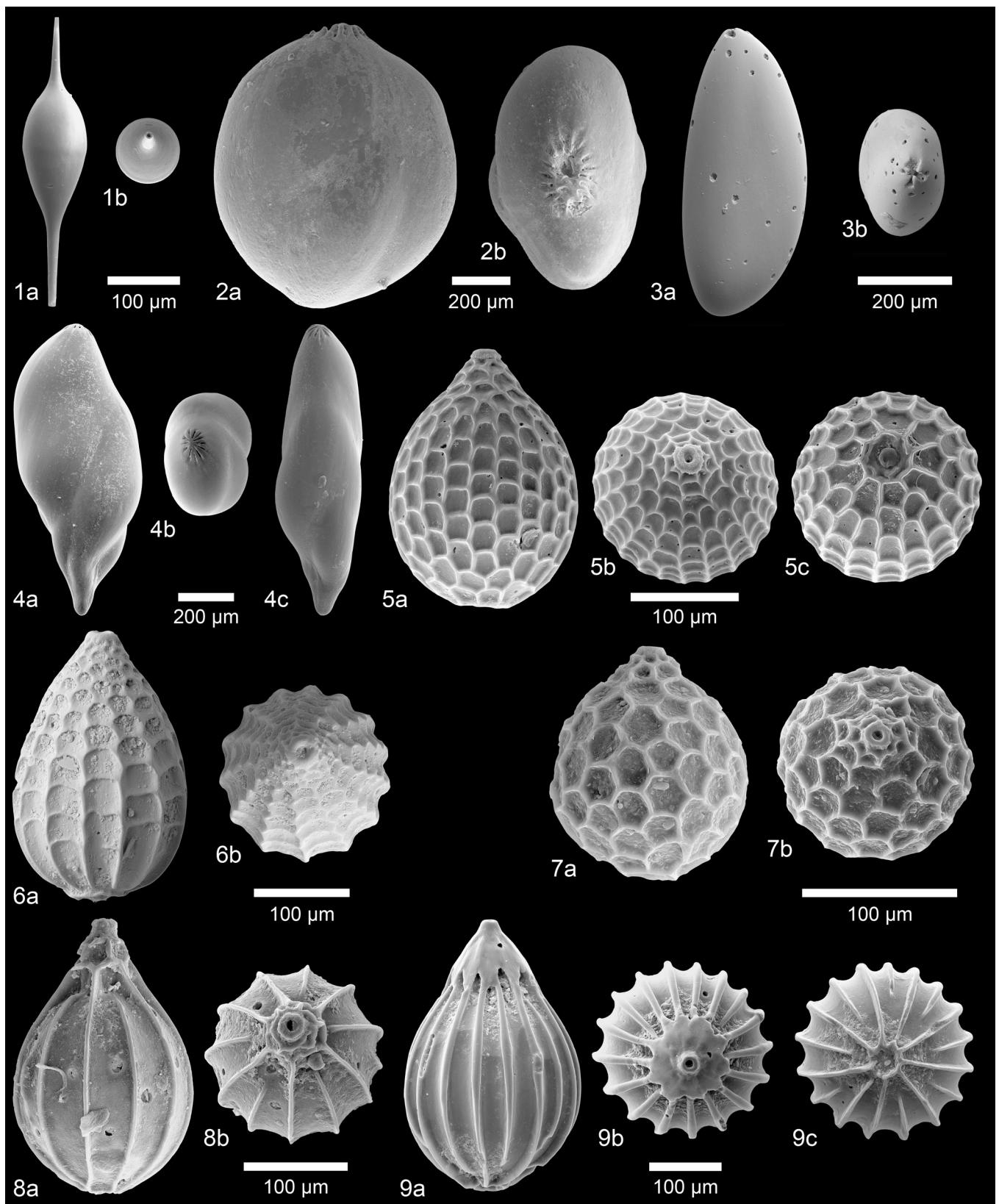


PLATE 15

- 1a-b *Homalohedra borealis* (Loeblich and Tappan, 1954), Hypotype from POS391 550-1
2a-c *Homalohedra eucostata* (McCulloch, 1977), Hypotype from POS391 562-1
3a-c *Oolina lineata* subsp. *communis* McCulloch, 1977, Hypotype from POS391 534-1
4a-c *Oolina ampulladistoma* (Jones, 1874), Hypotype from POS391 558-1
5a-c *Oolina globosa* (Montagu, 1803), Hypotype from POS391 571-1
6a-b *Oolina laevigata* d'Orbigny, 1839, Hypotype from AL232 1022
7a-b *Fissurina agassizi* Todd and Bronnimann, 1957, Hypotype from TTR17 MS419G 45
8a-b *Fissurina circularis* Todd, 1954, Hypotype from GeoB 9271

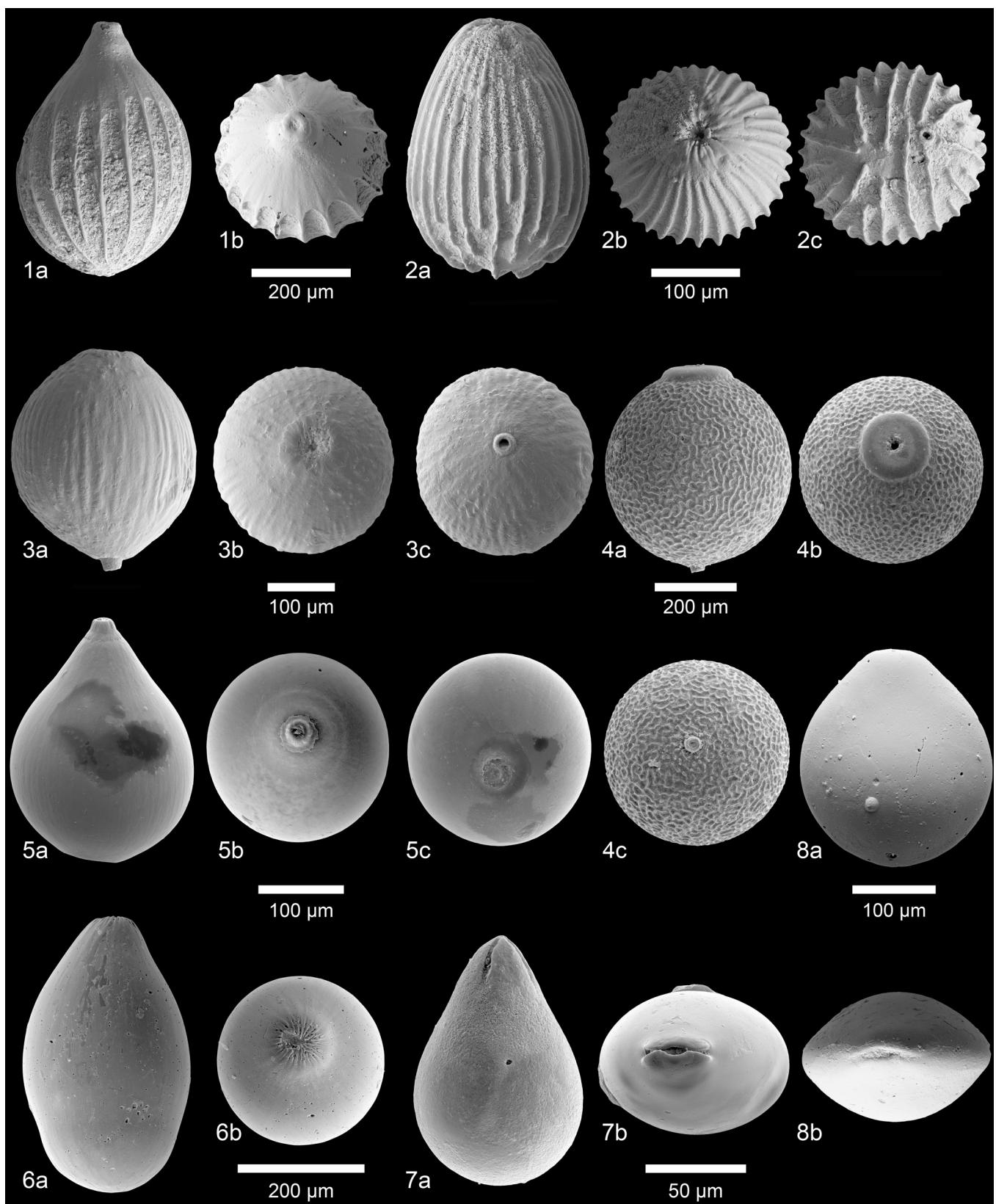


PLATE 16

- 1a-c *Fissurina lucida* (Williamson, 1848), Hypotype from PS70/033-2
2a-c *Fissurina eburnea* Buchner, 1940, Hypotype from POS391 550-1
3a-b *Fissurina kerguelensis* Parr, 1950, Hypotype from TTR17 MS419G 35
4a-c *Fissurina annectens* (Burrows and Holland, 1895), Hypotype from POS391 555-1
5a-b *Fissurina crassiporosa* McCulloch, 1977, Hypotype from TTR17 MS411G 10
6a-b *Fissurina longpointensis* McCulloch, 1977, Hypotype from TTR17 MS411 40

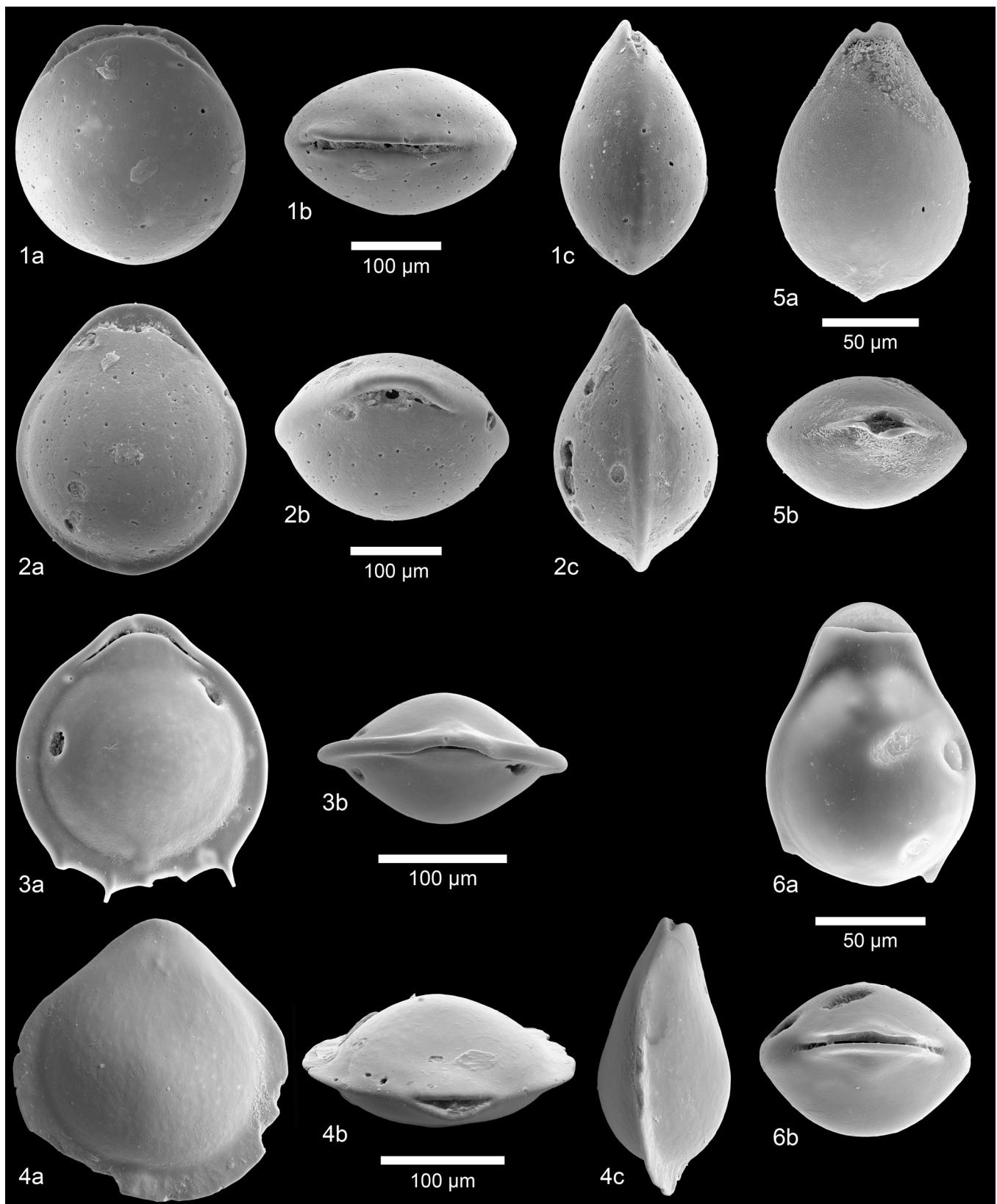


PLATE 17

- 1a-b *Parafissurina basispinata* McCulloch, 1977, Hypotype from TTR17 MS419G 84
2a-b *Parafissurina marginata* (Walker and Boys, 1803), Hypotype from TTR17 MS411G 60
3a-b *Fissurina pseudoorbignyana* (Buchner, 1940), Hypotype from TTR17 MS411G 10
4a-b *Fissurina dublini* McCulloch, 1977, Hypotype from P292 578-1
5a-b *Fissurina derogata* McCulloch, 1977, Hypotype from TTR17 MS411G 35
6a-b *Fissurina lacunata* (Burrows and Holland, 1895), Hypotype from PS70/028-2
7a-b *Fissurina nucelloides* (Buchner, 1940), Hypotype from GeoB9209-2
8a-c *Fissurina pseudolucida* Zheng, 1979, Hypotype from POS391 544-2

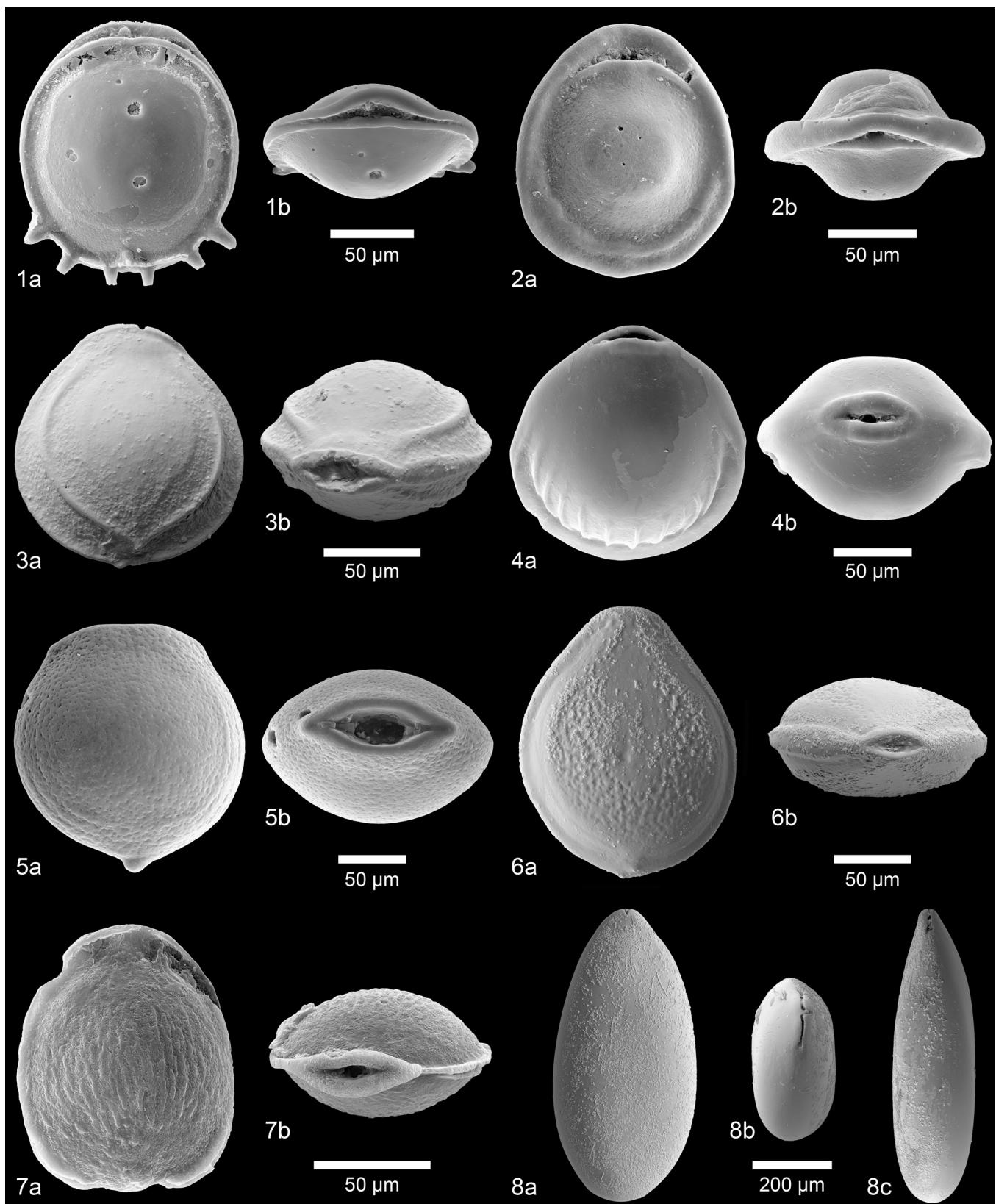


PLATE 18

- 1a-b *Palliolatella semimarginata* (Reuss, 1870), Hypotype from P292 578-1
2a-c *Parafissurina lateralis* (Cushman, 1913), Hypotype from PS70/033-2
3a-c *Hoeglundina elegans* (d'Orbigny, 1826), Hypotype from P292 581-1
4a-b *Parafissurina felsinea* (Fornasini, 1894), Hypotype from PS70/038-2
5a-c *Robertinoides bradyi* (Cushman and Parker, 1936), Hypotype from M07-15
6a-c *Parafissurina robusta* (Zheng, 1979), Hypotype from POS326 455
7a-c *Robertinoides bradyi* d'Orbigny, 1826, Hypotype from PS70/029-3
8a-b *Robertinoides pumilum* Höglund, 1947, Hypotype from TTR17 MS411G 40

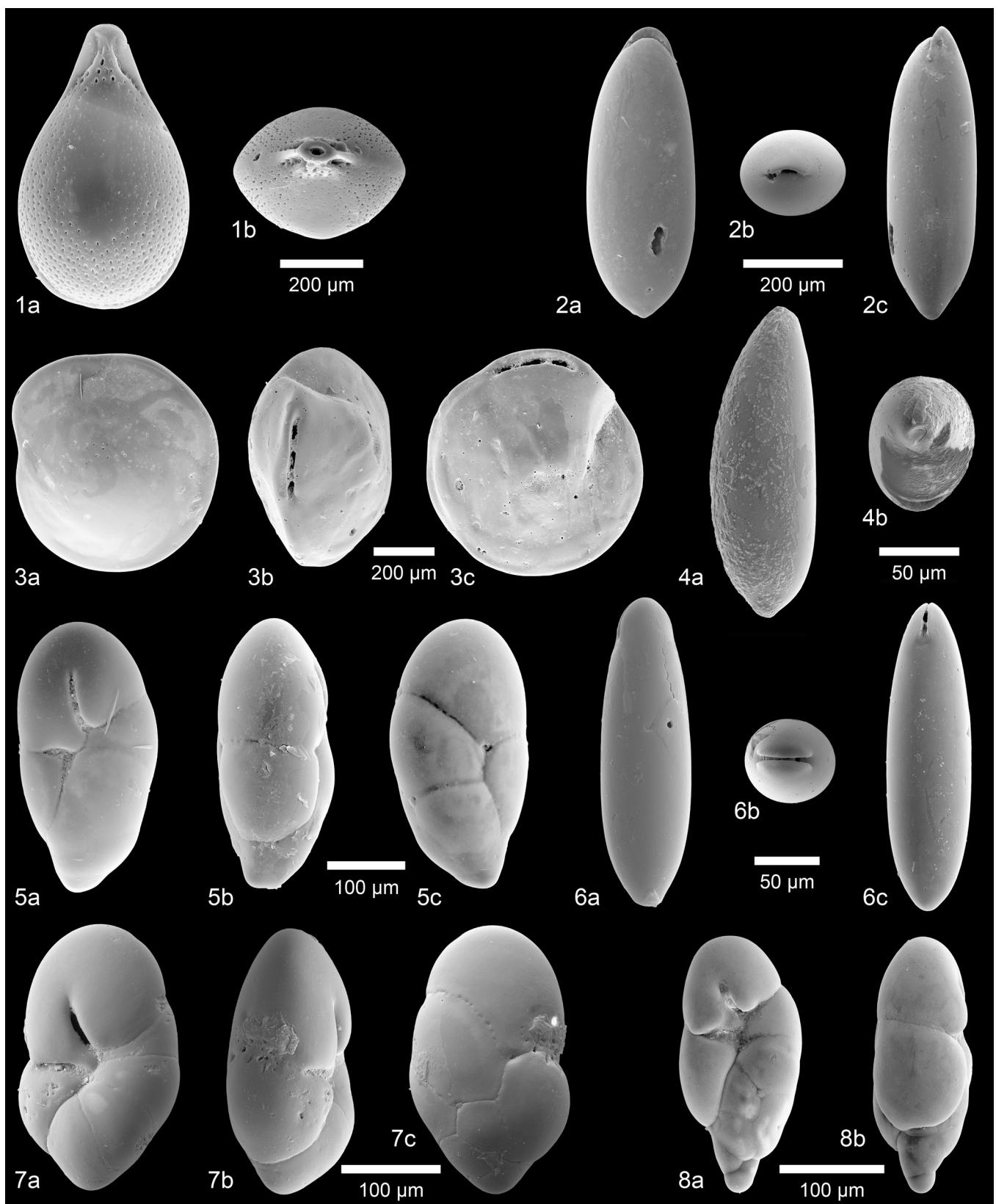


PLATE 19

- 1a-c *Globigerina bulloides* d'Orbigny, 1826, Hypotype from PS70/028-2
2a-c *Globigerinella calida* (Parker, 1961), Hypotype from TTR17 MS411G 0
3a-c *Globigerinella siphonifera* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 15
4a-c *Globigerinella glutinata* (Egger, 1893), Hypotype from PS70/028-2
5a-c *Globoturborotalita rubescens* (Hofker, 1956), Hypotype from TTR17 MS411G 40
6a-c *Globigerinella uvula* (Ehrenberg, 1861), Hypotype from PS70/032-2

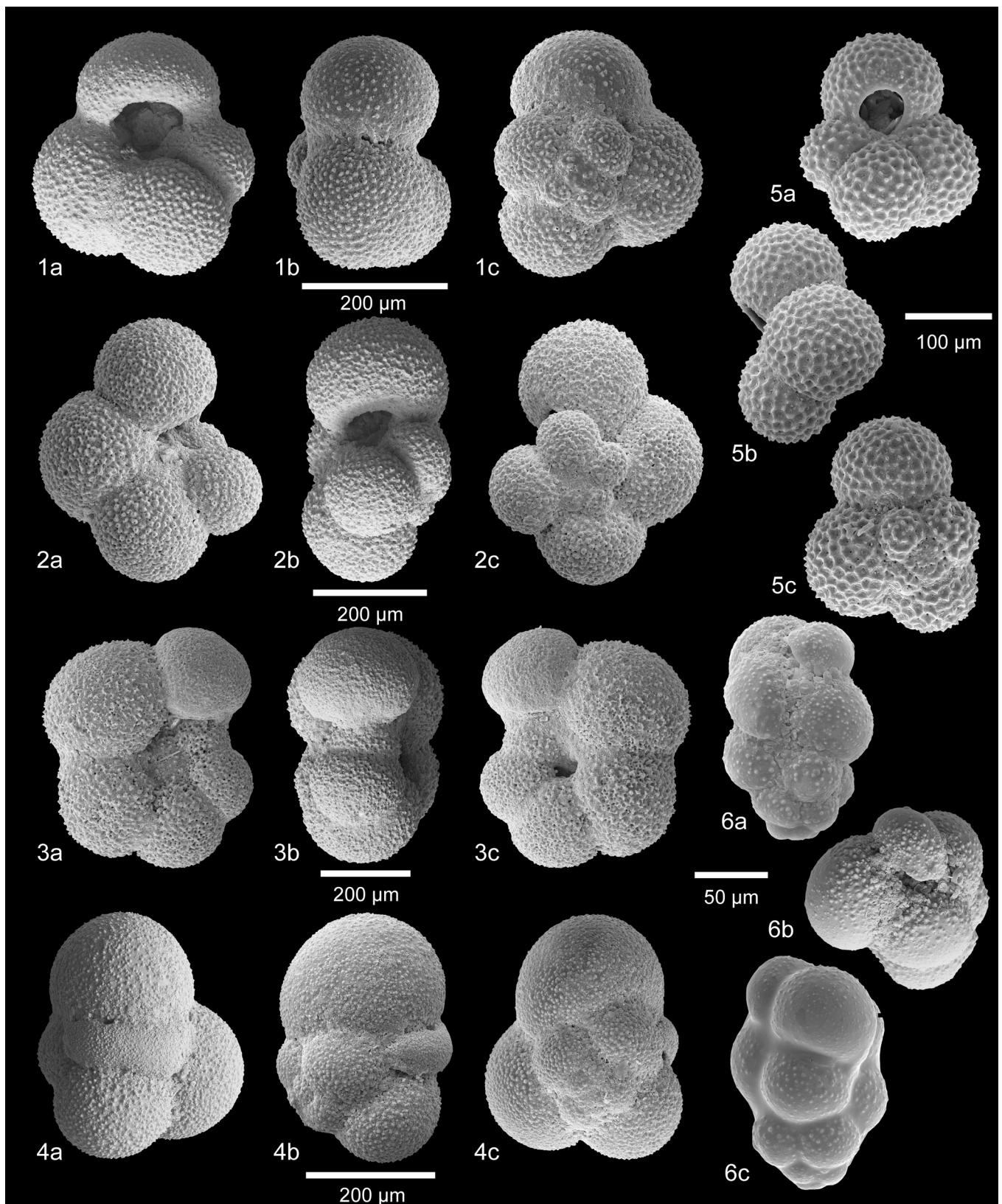


PLATE 20

- 1a-c *Globigerinoides conglobatus* Brady, 1879, Hypotype from TTR17 MS411G 10
2a-c *Globigerinoides ruber* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 0
3a-c *Globigerinoides sacculifer* (Brady, 1877), Hypotype from TTR17 MS411G 0
4a-c *Globigerinoides elongatus* (d'Orbigny, 1826), Hypotype from TTR17 MS411G 0
5 *Orbulina universa* d'Orbigny, 1839, Hypotype from TTR17 MS411G 0
6a-c *Globigerinoides trilobus* (Reuss, 1850), Hypotype from TTR17 MS411G 0

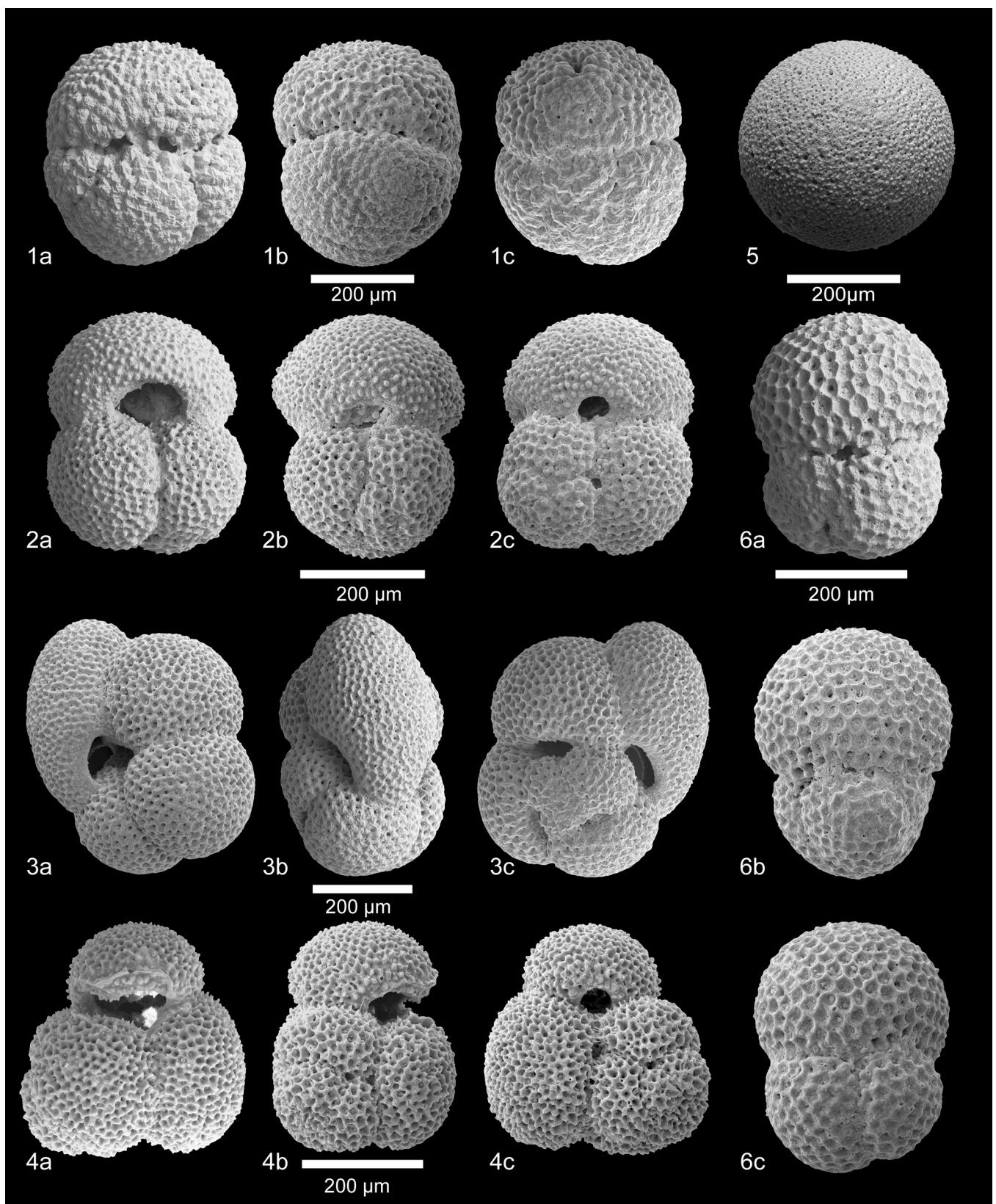


PLATE 21

- 1a-c *Globorotalia inflata* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 0
2a-c *Globorotalia scitula* (Brady, 1882), Hypotype from TTR17 MS411G 0
3a-c *Globorotalia hirsuta* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 0
4a-c *Truncorotalia crassaformis* (Galloway and Wissler, 1927), Hypotype from TTR17 MS411G 25
5a-c *Truncorotalia truncatulinoides* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 0

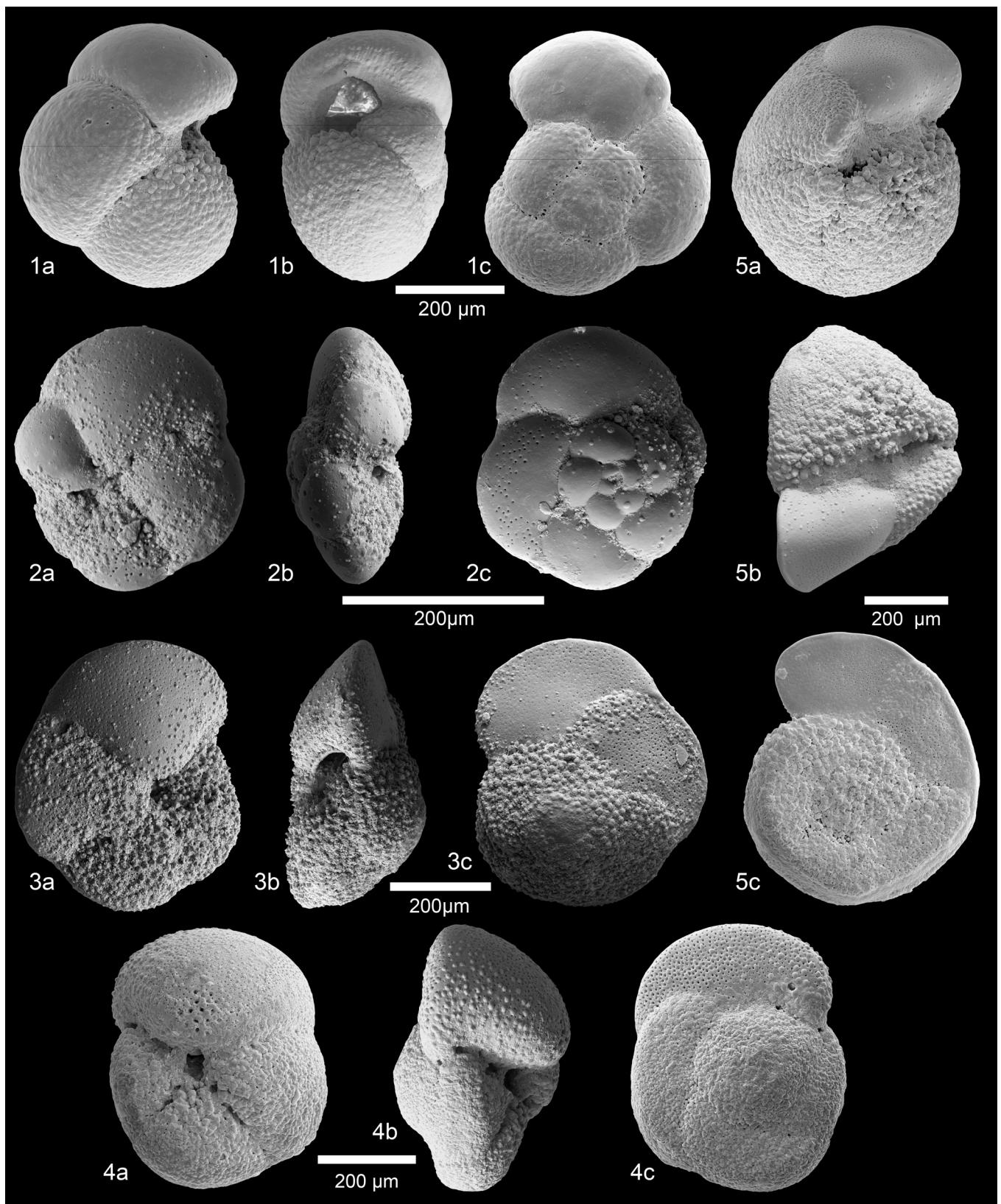


PLATE 22

- 1a-c *Pulleniatina obliquiloculata* (Parker and Jones, 1865), Hypotype from TTR17 MS411G 45
2a-c *Neogloboquadrina incompta* (Cifelli, 1961), Hypotype from PS70/011-1
3a-c *Neogloboquadrina pachyderma* (Ehrenberg, 1861), Hypotype from PS70/028-2
4a-b *Neogloboquadrina dutertrei* (d'Orbigny, 1839), Hypotype from PS70/011-1
5a-c *Berggrenia clarkei* (Rögl and Bolli, 1973), Hypotype from PS70/002-2

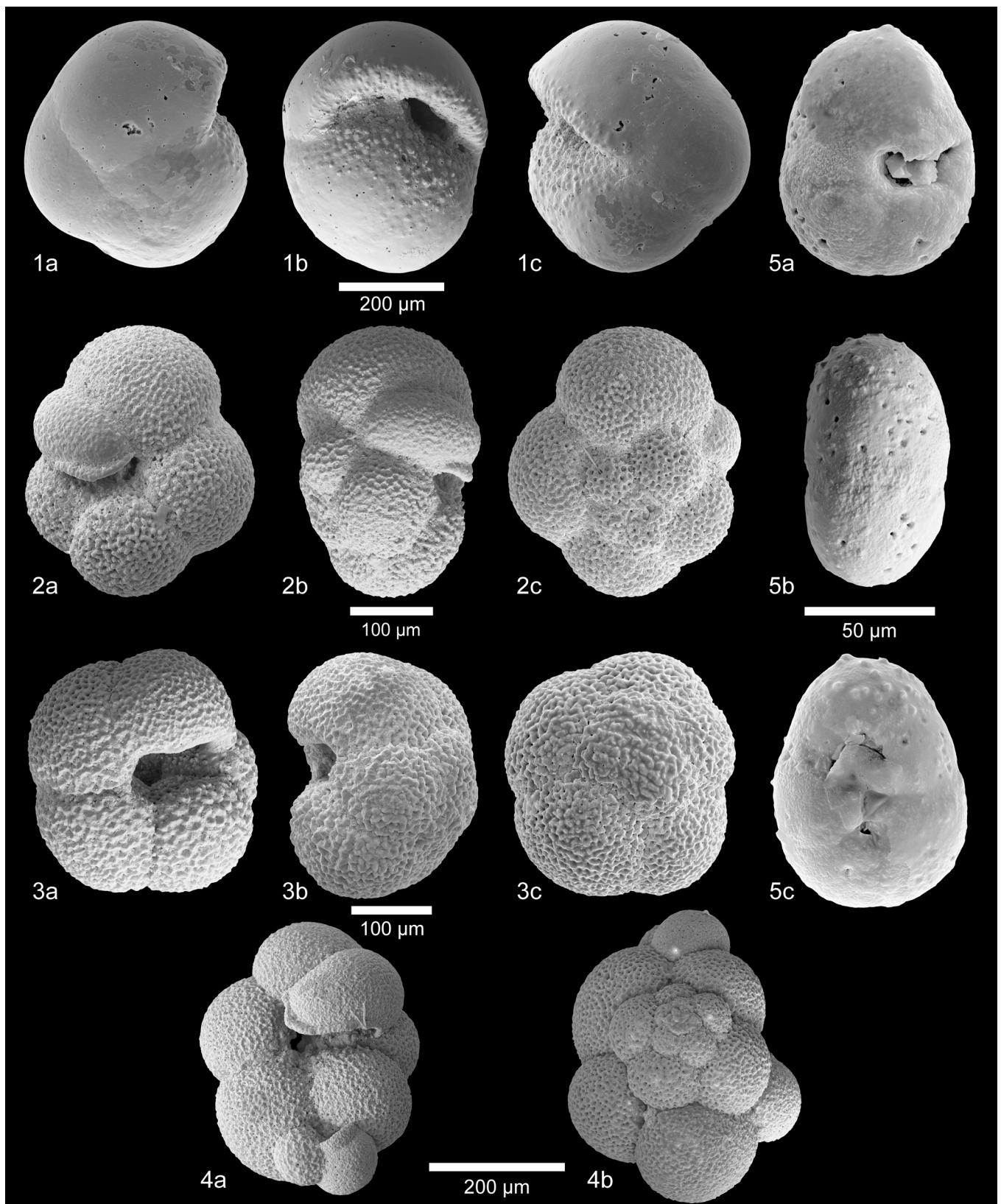


PLATE 23

- 1a-c *Turborotalita quinqueloba* (Natland, 1938), Hypotype from PS70/028-2
2a-c *Turborotalita humilis* (Brady, 1884), Hypotype from TTR17 MS411G 20
3a-c *Turborotalita cristata* (Heron-Allen and Earland, 1929), Hypotype from TTR17 MS411G 20
4a-b *Tenuitella iota* (Parker, 1962), Hypotype from TTR17 MS419G 25
5a-c *Tenuitella anfracta* (Parker, 1967), Hypotype from TTR17 MS419G 55
6a-c *Streporchilus tokelauae* (Boersma, 1969), Hypotype from TTR17 MS419G 25

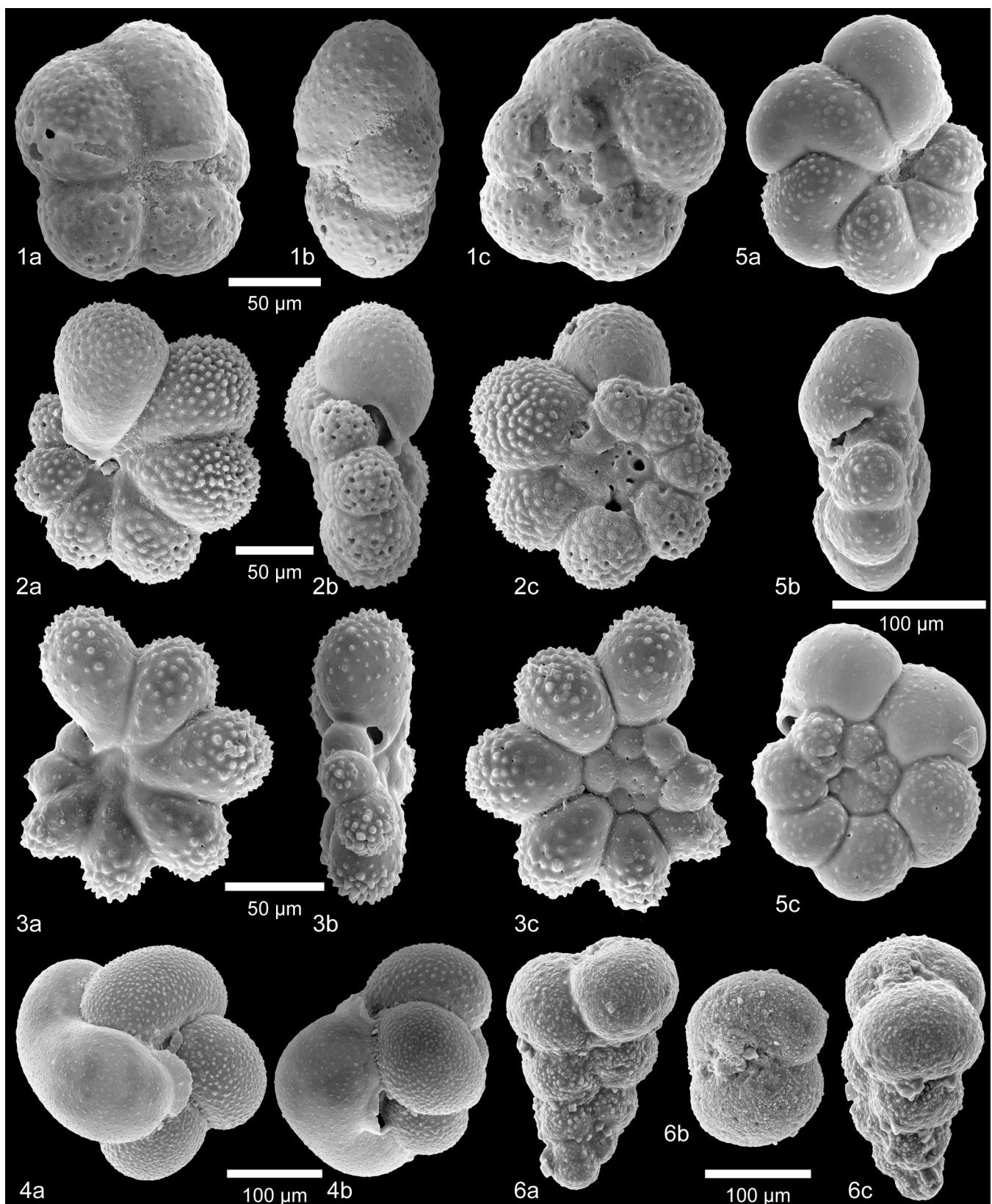


PLATE 24

- 1a-c *Bolivina alata* (Seguenza, 1862), Hypotype from TTR17 MS419G 25
2a-c *Bolivina dilatata* (Reuss, 1850), Hypotype from P292 577-1
3a-c *Bolivinellina pseudopunctata* (Höglund, 1947), Hypotype from GeoB9204-1
4a-c *Bolivina pseudoplicata* Heron-Allen and Earland, 1930, Hypotype from TTR17 MS411G 25
5a-c *Bolivinellina striatula* (Cushman, 1922), Hypotype from TTR17 MS411G 0
6a-c *Bolivina difformis* (Williamson, 1858), Hypotype from P292 576-1
7a-b *Bolivina spinescens* Cushman, 1911, Hypotype from POS391 555-1
8a-c *Bolivina subspinescens* Cushman, 1922, Hypotype from P292 580-1

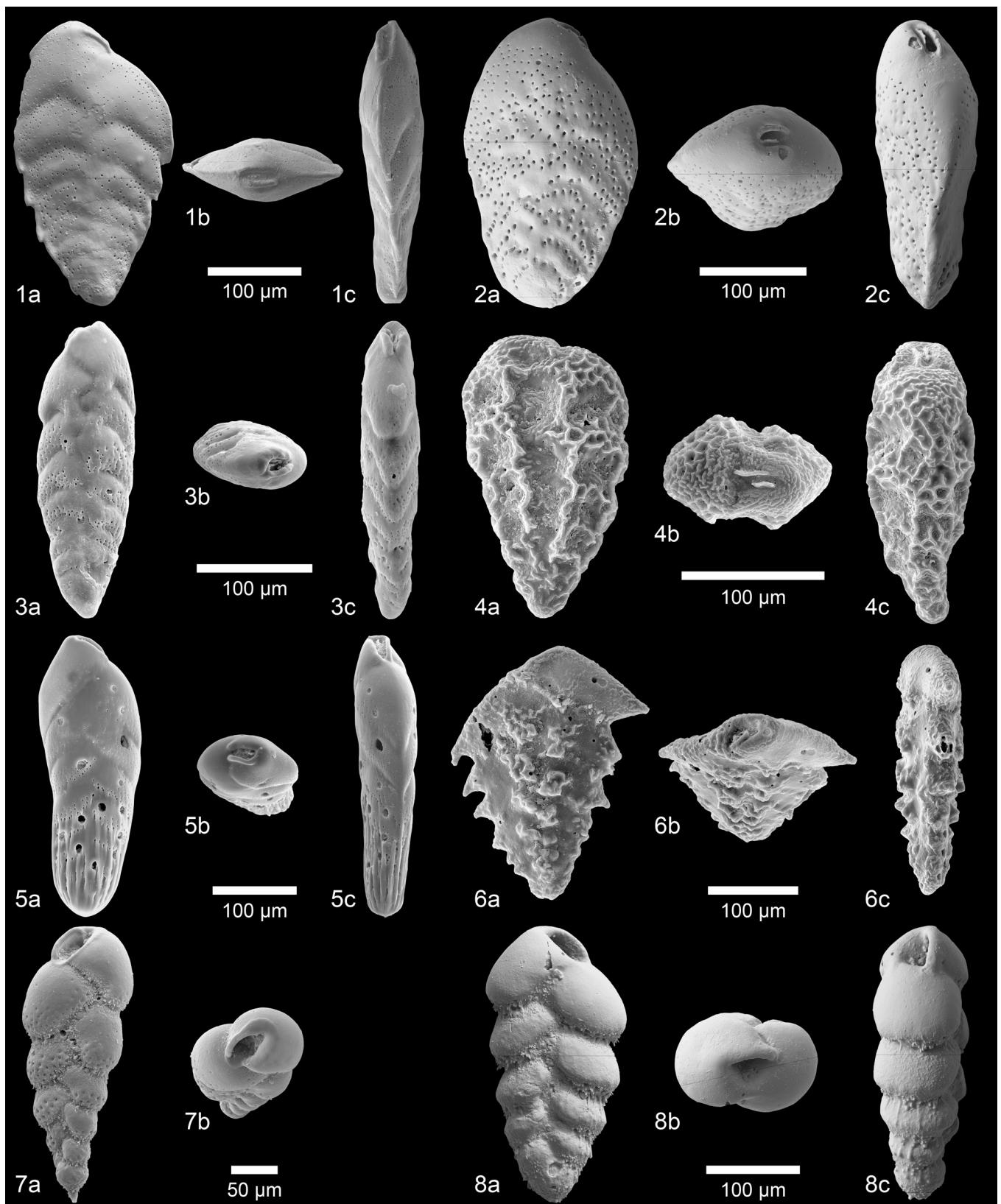


PLATE 25

- 1a-b *Brizalina subaenariensis* (Cushman, 1922), Hypotype from P292 580-1
2a-c *Parabrizalina porrectum* (Brady, 1881), Hypotype from AL232 1025
3a-c *Cassidulina reniforme* Nörvang, 1945, Hypotype from POS391 555-1
4a-c *Cassidulina laevigata* d'Orbigny, 1826, Hypotype from PS70/028-2
5a-c *Cassidulina carinata* Silvestri, 1896, Hypotype from PS70/011-1
6a-c *Cassidulina teretis* Tappan, 1951, Hypotype from PS70/038-2

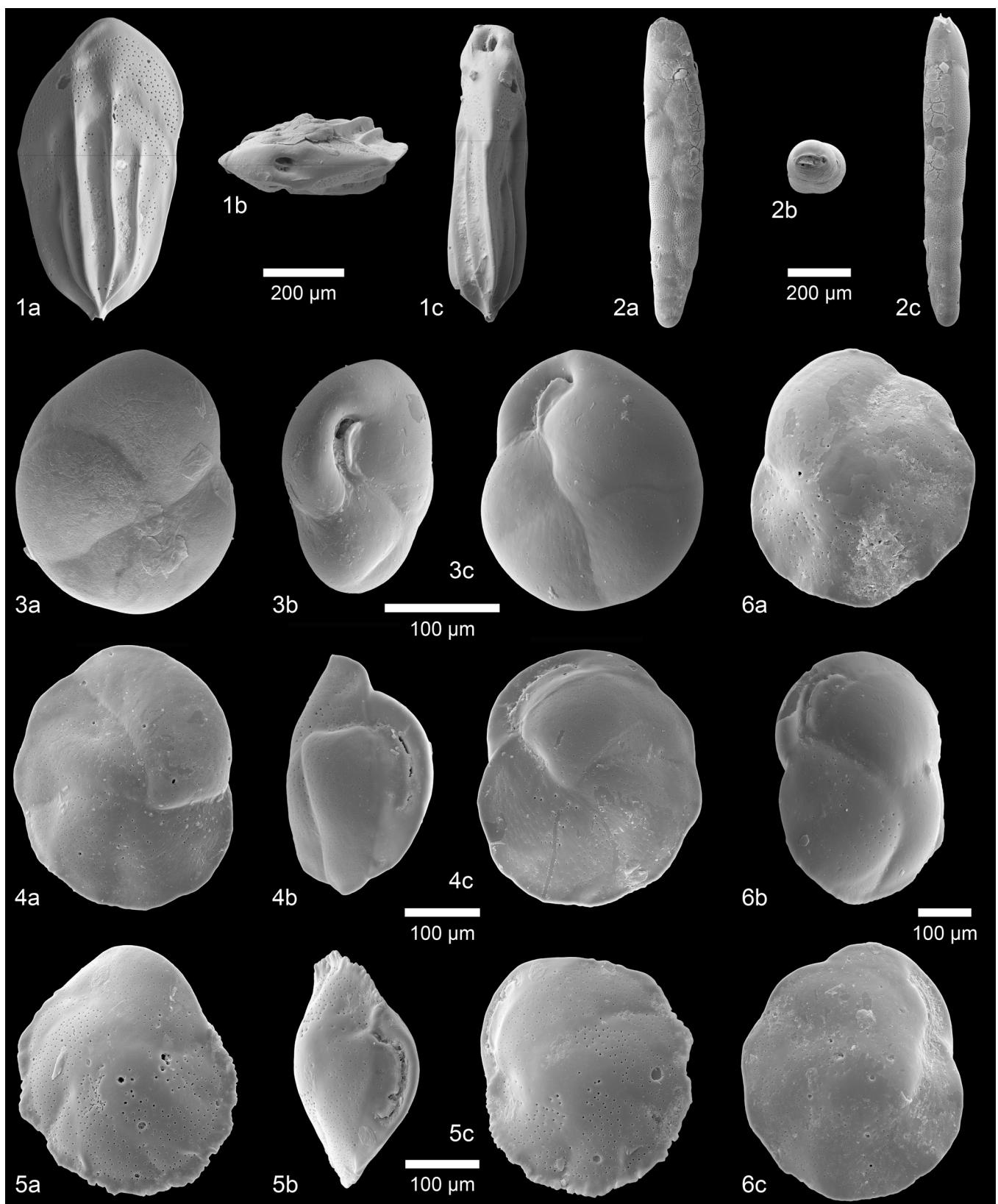


PLATE 26

- 1a-c *Cassidulina neoteretis* Seidenkrantz, 1995, Hypotype from POS391 558-1
2a-c *Takayanagia delicata* (Cushman, 1927), Hypotype from PS292 576-1
3a-c *Cassidulina crassa* d'Orbigny, 1839, Hypotype from PS70/028-2
4a-c *Globocassidulina subglobosa* (Brady, 1881), Hypotype from TTR17 MS411G 0
5a-c *Islandiella norcrossi* (Cushman, 1933), Hypotype from POS391 555-1
6a-b *Cassidulinoides bradyi* (Norman, 1881), Hypotype from TTR17 MS419G 75

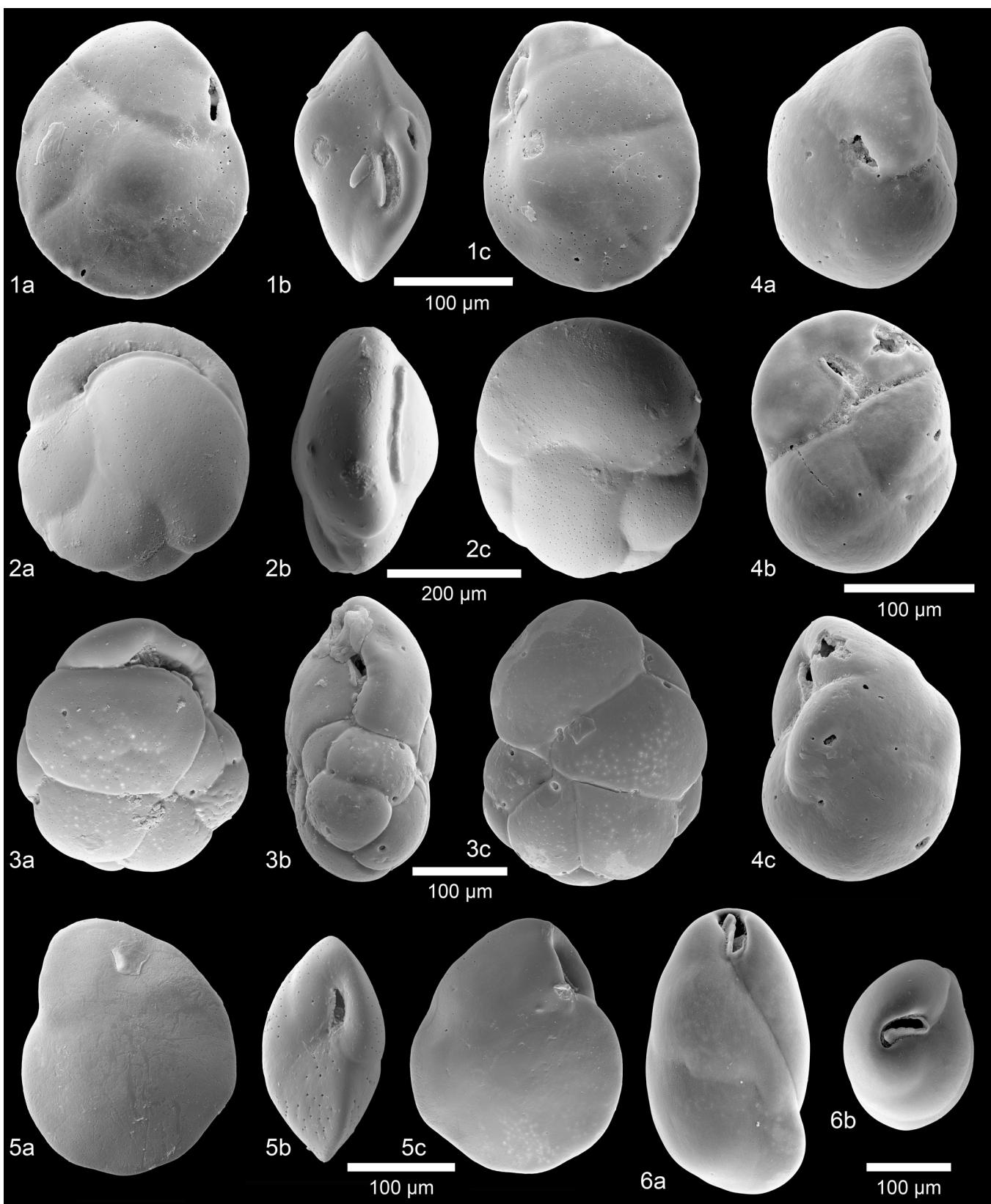


PLATE 27

- 1a-b *Ehrenbergina trigona* Goës, 1896, Hypotype from GeoB 9256-1
2a-c *Stainforthia fusiformis* (Williamson, 1858), Hypotype from AL232 1025
3a-c *Stainforthia loeblichi* (Feyling-Hanssen, 1954), Hypotype from POS391 555-1
4a-c *Furstenkoina complanata* (Egger, 1893), Hypotype from AL232 1025
5a-c *Stainforthia skagerakensis* (Höglund, 1947), Hypotype from AL232 1025
6a-c *Rectuvigerina elongatastriata* (Colom, 1952), Hypotype from TTR17 MS411G 10
7a-b *Siphogenerina columellaris* (Brady, 1881), Hypotype from TTR17 MS411G 15
8a-b *Bulimina aculeata* d'Orbigny, 1826, Hypotype from TTR17 MS419G 94
9a-b *Bulimina marginata* d'Orbigny, 1826, Hypotype from TTR17 MS411G 10
10a-c *Bulimina striata* d'Orbigny, 1826, Hypotype from TTR17 MS411G 15

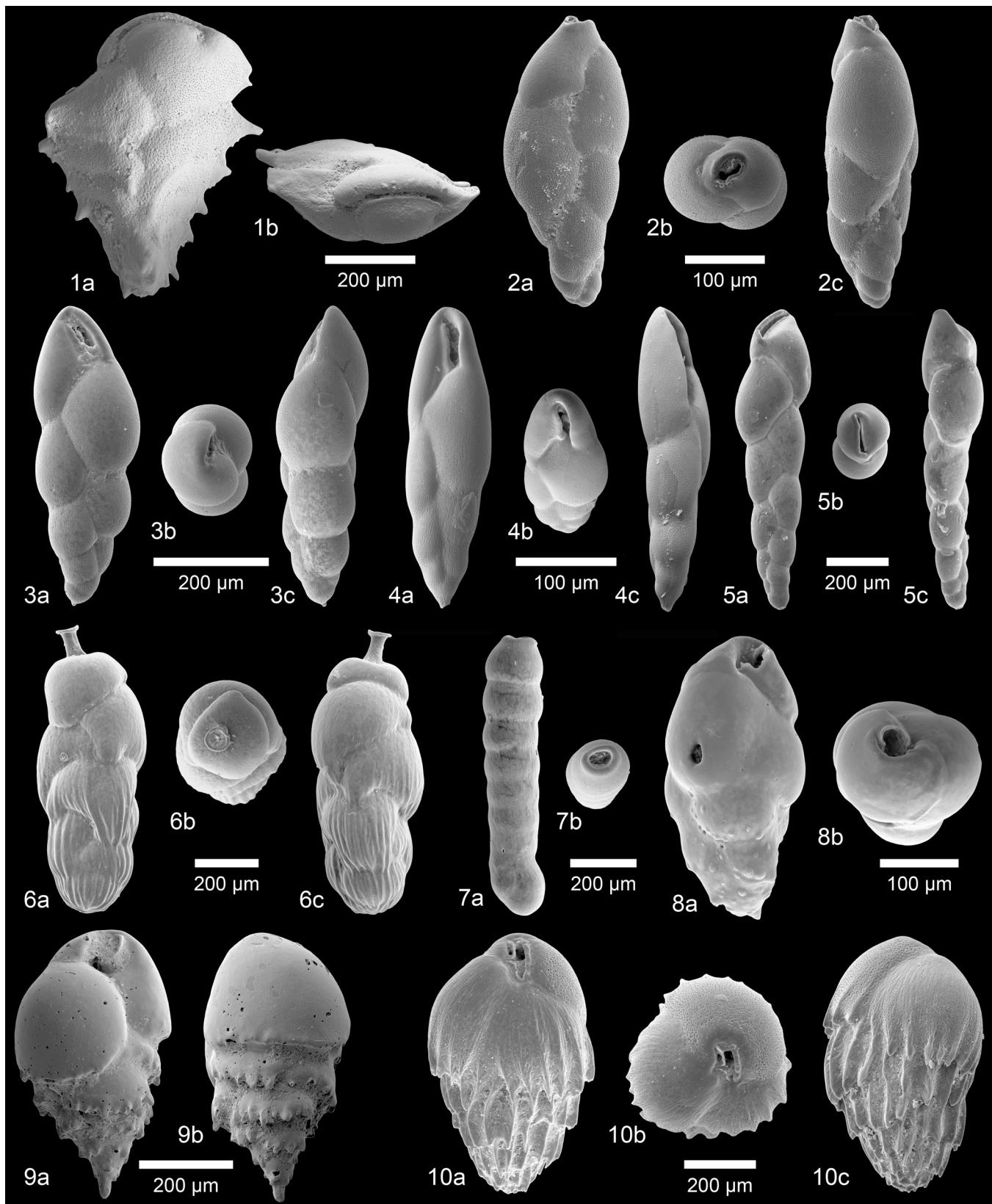


PLATE 28

- 1a-c *Globobulimina turgida* (Bailey, 1851), Hypotype from AL232 1025
2a-c *Globobulimina affinis* (d'Orbigny, 1839), Hypotype from AL232 1025
3a-c *Globobulimina affinis* (d'Orbigny, 1839), Hypotype from AL232 1026
4a-c *Praeglobobulimina ovata* (d'Orbigny, 1846), Hypotype from P292 580-1
5a-c *Globobulimina dololum* (Terquem and Terquem, 1886), Hypotype from POS391 550-1
6a-b *Buliminella spinigera* Cushman, 1922, Hypotype from GeoB12722-1

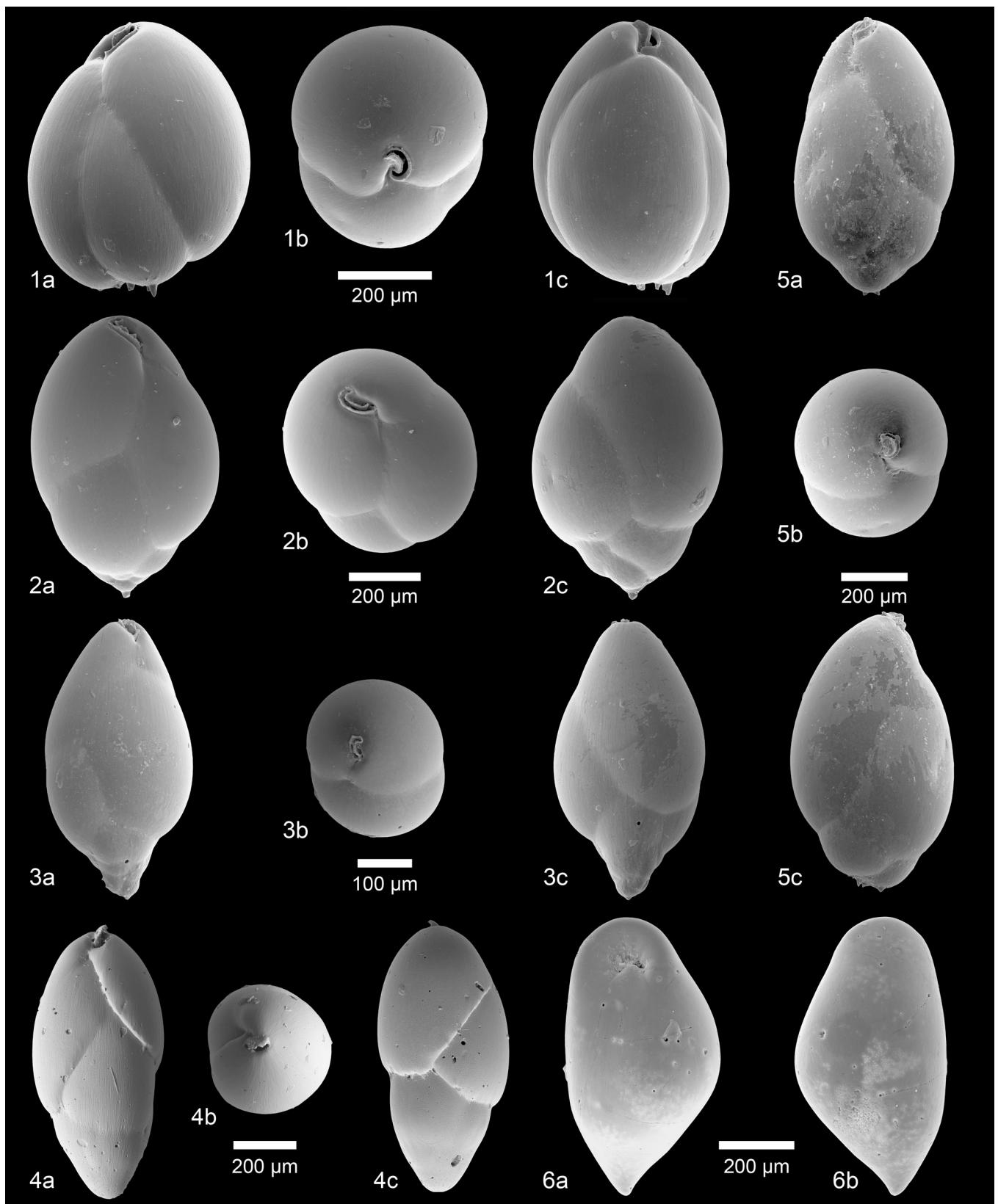


PLATE 29

- 1a-b *Uvigerina mediterranea* Hofker, 1932, Hypotype from P292 580-1
2a-b *Uvigerina peregrina* Cushman, 1923, Hypotype from P292 580-1
3a-b *Uvigerina auberiana* d'Orbigny, 1839, Hypotype from GeoB 6721-1
4a-b *Uvigerina peregrina parva* Lutze, 1986, Hypotype from GeoB 6721-1
5a-b *Uvigerina pigmaea* d'Orbigny, 1826, Hypotype from P292 576-1
6a-b *Angulogerina angulosa* (Williamson, 1858), Hypotype from P292 577-1
7a-b *Trifarina bradyi* (Cushman, 1923), Hypotype from TTR17 MS419G 107
8a-b *Trifarina fornasini* (Sell, 1948), Hypotype from TTR17 MS411G 25
9a-b *Nodogenerina virgula* (Brady, 1884), Hypotype from PS70/033-2
10a-c *Cancris auriculus* (Fichtel and Moll, 1798), Hypotype from TTR17 MS411G 40

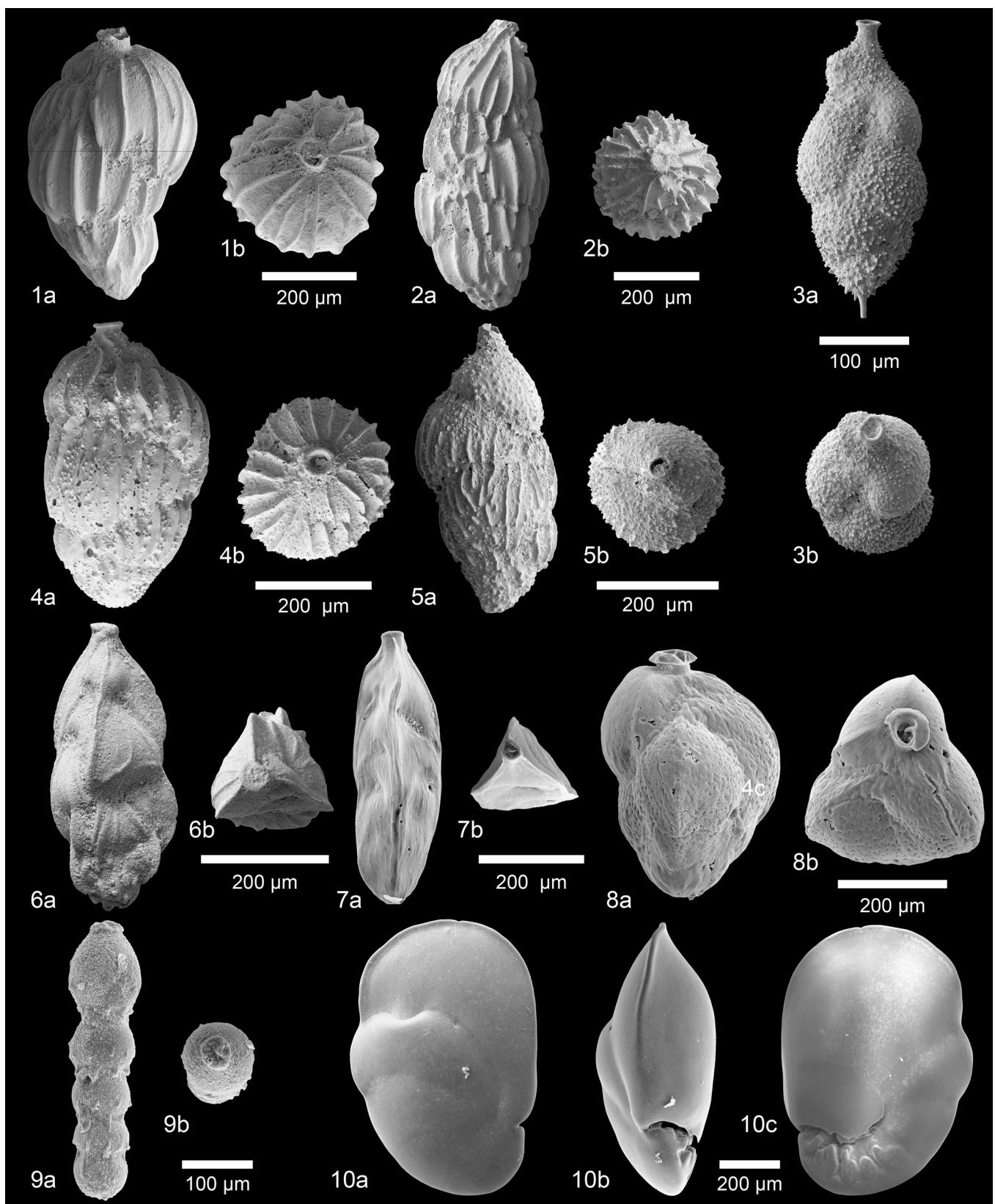


PLATE 30

- 1a-c *Valvularia bradyana* (Fornasini, 1900), Hypotype from TTR17 MS411G 15
- 2a-c *Ioanella tumidula* (Brady, 1884), Hypotype from PS70/033-2
- 3a-c *Stomatorbina concentrica* (Parker and Jones, 1864), Hypotype from GeoB9256-1
- 4a-c *Hyrrokkin sarcophaga* Cedhagen, 1994, Hypotype from POS391 562-1
- 5a-c *Hyrrokkin sarcophaga* Cedhagen, 1994, Hypotype from POS391 550-1

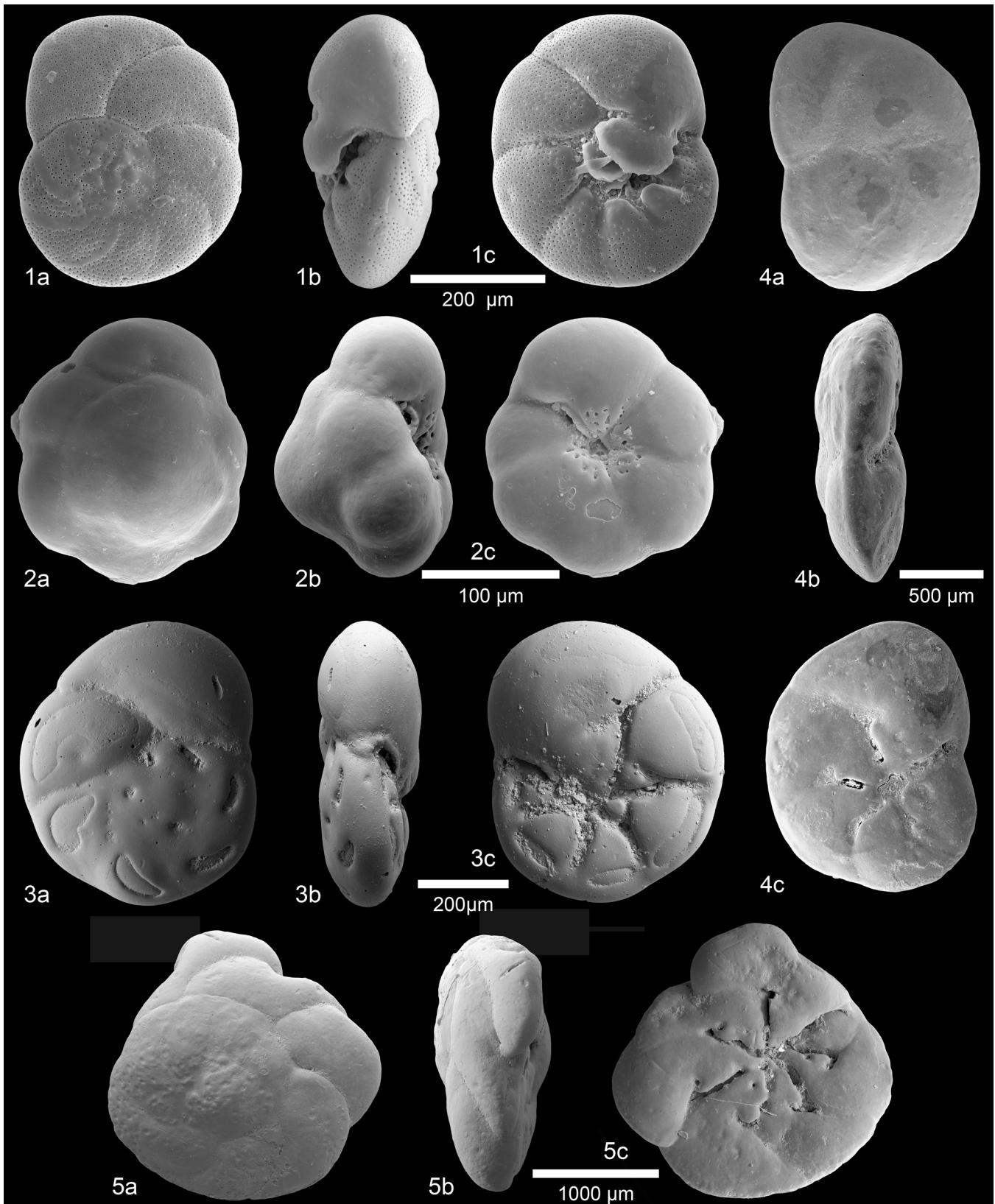


PLATE 31

- 1a-c *Gavelinopsis nitida* (Williamson, 1858), Hypotype from POS391 555-1
2a-c *Gavelinopsis praegeri* (Heron-Allen and Earland, 1913), Hypotype from TTR17 MS411G 0
3a-c *Gavelinopsis caledonia* Murray and Whittaker, 2001, Hypotype from POS325 455
4a-c *Rosalina globularis* d'Orbigny, 1826, Hypotype from PS70/011-1
5a-c *Rosalina brady* (Cushman, 1915), Hypotype from PS70/011-1

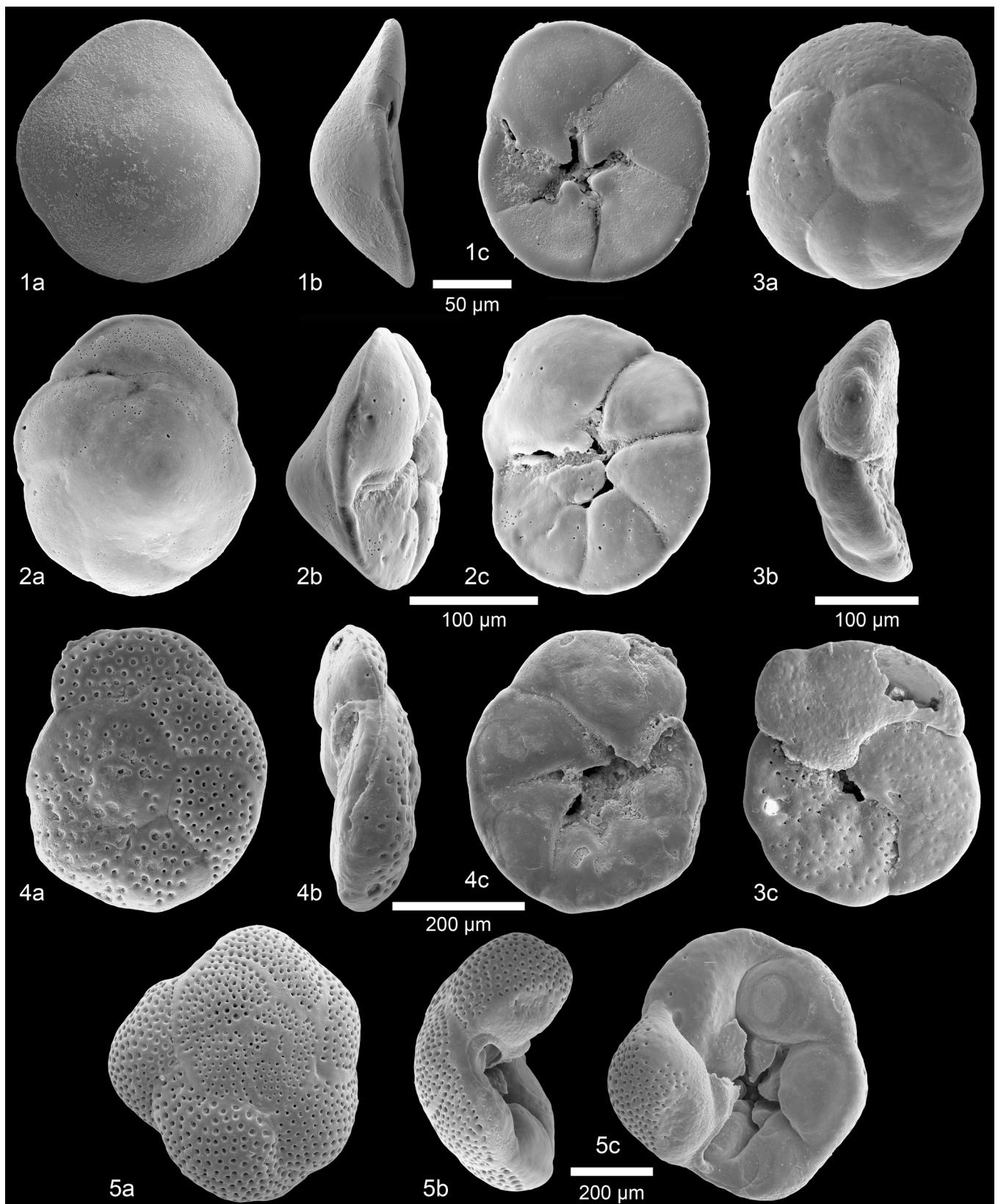


PLATE 32

- 1a-c *Rosalina semipunctata* (Bailey, 1851), Hypotype from GeoB9205-1
2a-c *Sphaeroidina bulloides* d'Orbigny, 1826, Hypotype from TTR17 MS411G 0
3a-c *Glabratella patelliformis* (Brady, 1884), Hypotype from TTR17 MS411G 0
4a-c *Siphonina reticulata* (Czjzek, 1848), Hypotype from TTR17 MS411G 15
5a-c *Heronallenita lingulata* (Burrows and Holland, 1895), Hypotype from TTR17 MS411G 25

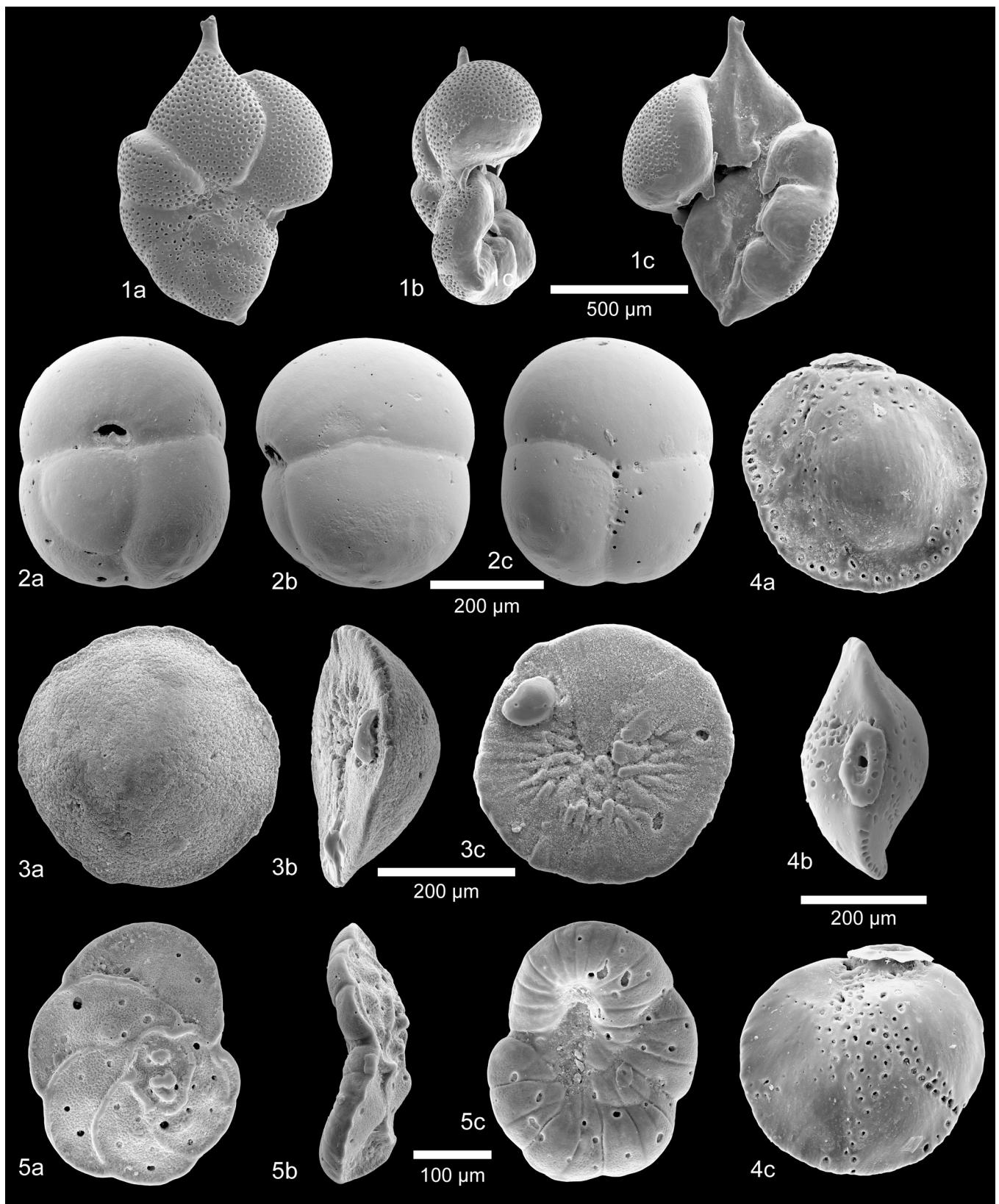


PLATE 33

- 1a-c *Cibicidoides mundulus* (Parker, 1953), Hypotype from TTR17 MS411G 0
2a-c *Cibicidoides pachyderma* (Rzehak, 1886), Hypotype from TTR17 MS411G 0
3a-c *Epistominella vitrea* Parker, 1953, Hypotype from AL316 321
4a-c *Epistominella exigua* (Brady, 1884), Hypotype from TTR17 MS411G 15
5a-c *Discorbinella bertheloti* (d'Orbigny, 1839), Hypotype from POS391 550-1

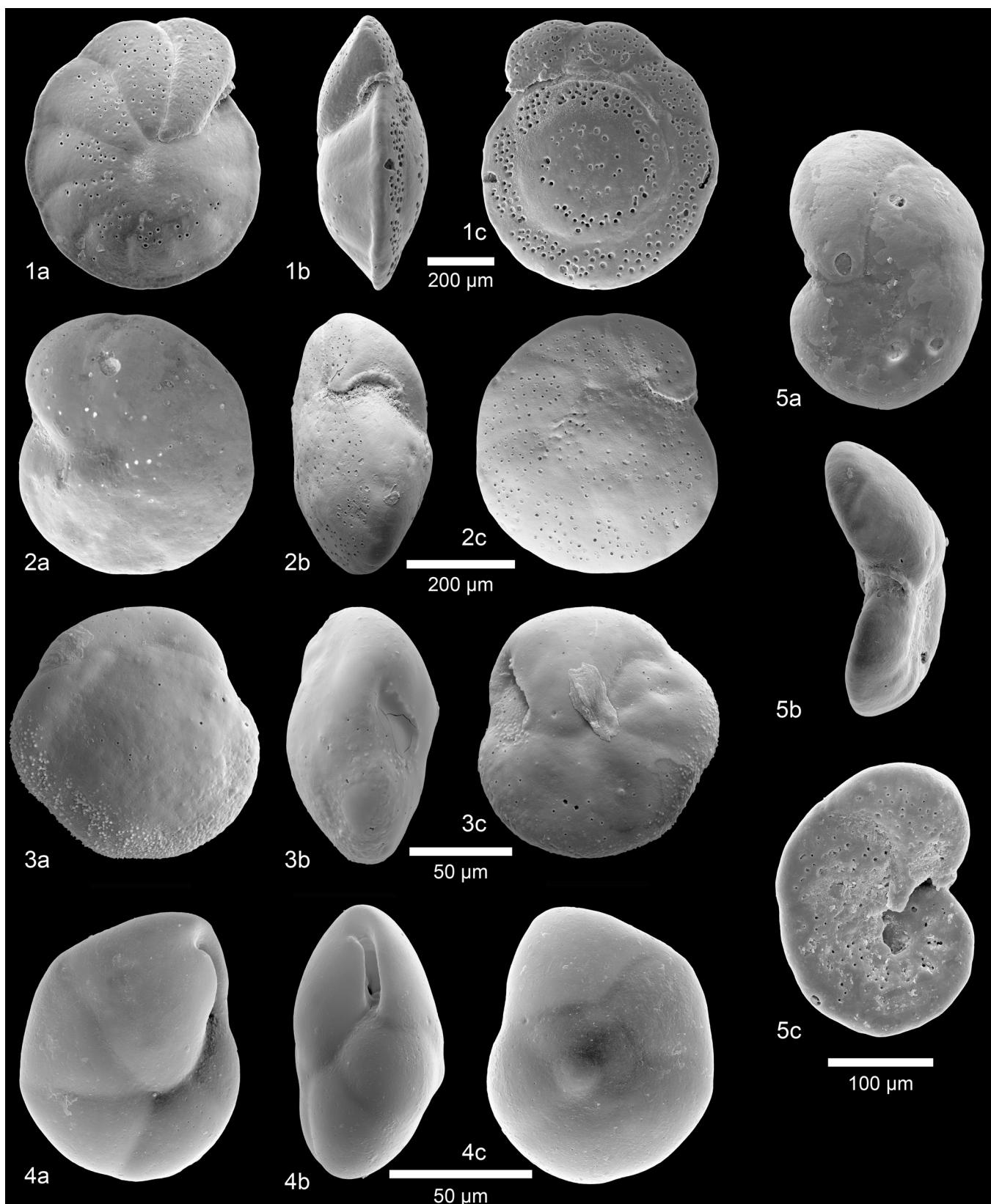


PLATE 34

- 1a-c *Planulina ariminensis* d'Orbigny, 1826, Hypotype from M07-15
2a-c *Hyalinea balthica* (d'Orbigny, 1826), Hypotype from P292 576-1
3a-c *Cibicides ungerianus* (d'Orbigny, 1826), Hypotype from POS391 534-1
4a-c *Cibicides refulgens* de Montfort, 1808, Hypotype from P292 574-1
5a-c *Cibicides aravaensis* Perelis and Reiss, 1976, Hypotype from TTR17 MS411G 50

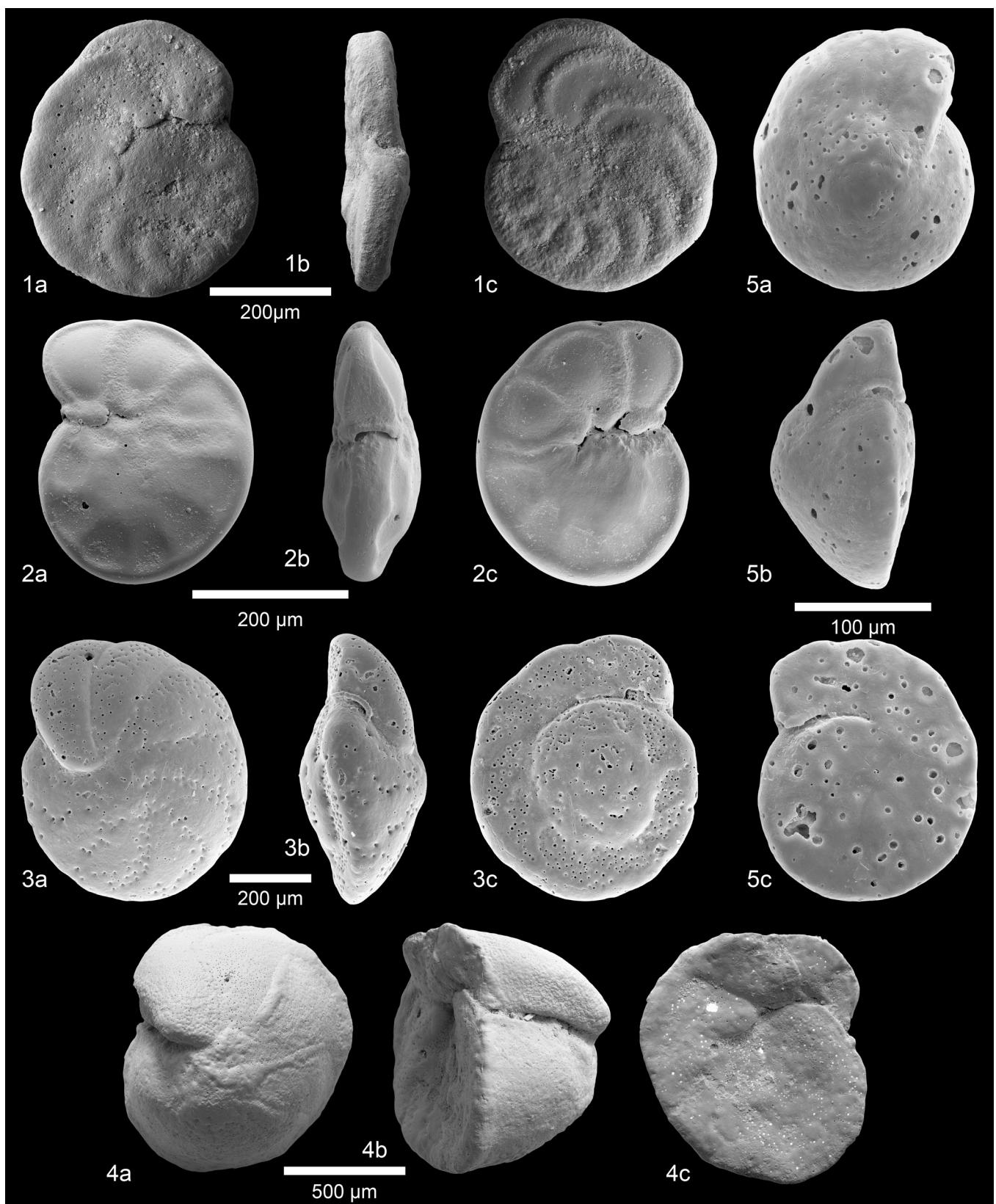


PLATE 35

- 1a-c *Lobatula lobatula* (Walker and Jacob, 1798), Hypotype from TTR17 MS411G 35
2a-c *Fontbotia wuellerstorfi* (Schwager, 1866), Hypotype from PS70/032-2
3a-c *Pseudoeponides falsobeccarii* Rouvillois, 1974, Hypotype from TTR17 MS419G 126
4a-c *Nuttallides umbonifera* (Rouvillois, 1974), Hypotype from TTR17 MS411G 20
5a-c *Nuttallides decorata* (Phleger and Parker, 1951), Hypotype from TTR17 MS411G 10

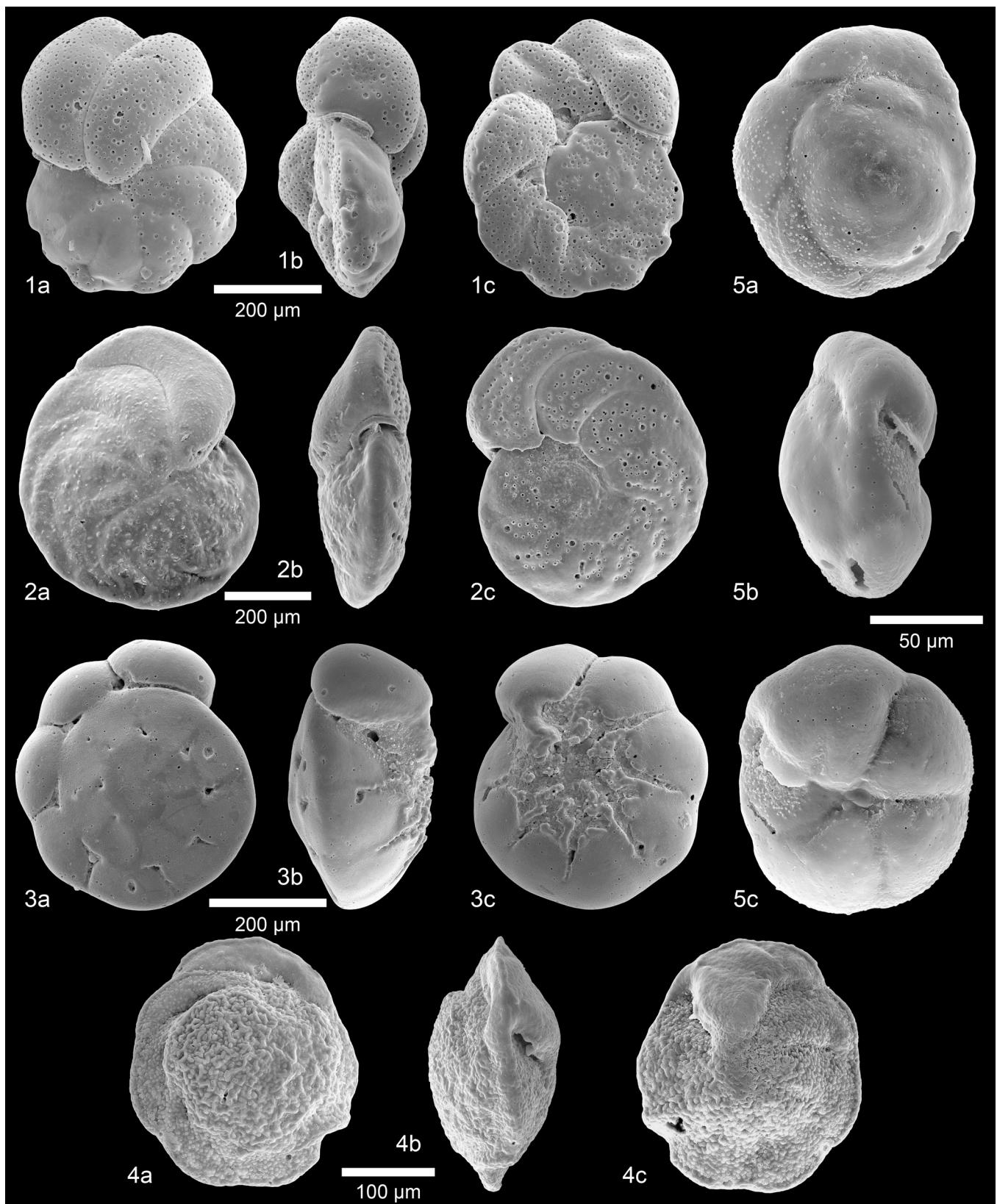


PLATE 36

- 1a-c *Astigerinata mamilla* (Williamson, 1858), Hypotype from TTR17 MS419G 25
2a-c *Nonion fabum* (Fichtel and Moll, 1798), Hypotype from TTR17 MS419G 55
3a-c *Nonion pauperatus* (Balkwill and Wright, 1885), Hypotype from TTR17 MS419G 100
4a-c *Nonionellina labradorica* (Dawson, 1860), Hypotype from AL232 1026
5a-c *Nonion pauciloculum* (Cushman, 1944), Hypotype from PS70/033-2

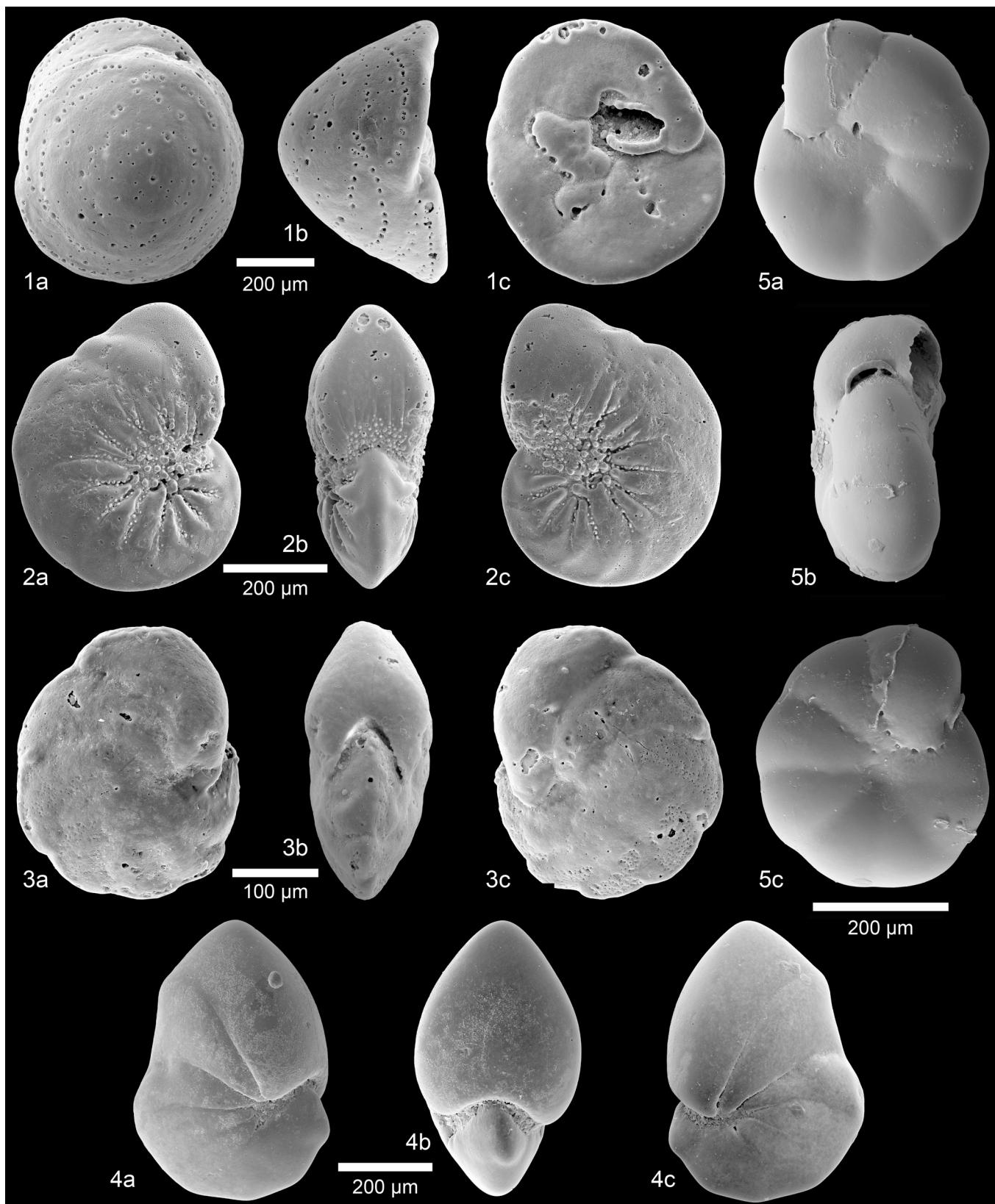


PLATE 37

- 1a-c *Nonionella turgida* (Williamson, 1858), Hypotype from TTR17 MS411G 15
2a-c *Nonionella iridea* Heron-Allen and Earland, 1932, Hypotype from TTR17 MS419G 100
3a-c *Astrononion gallowayi* Loeblich and Tappan, 1953, Hypotype from POS325 455
4a-c *Astrononion antarcticus* Parr, 1950, Hypotype from POS325 455
5a-c *Melonis pompilioides* (Fichtel and Moll, 1798), Hypotype from TTR17 MS419G 98

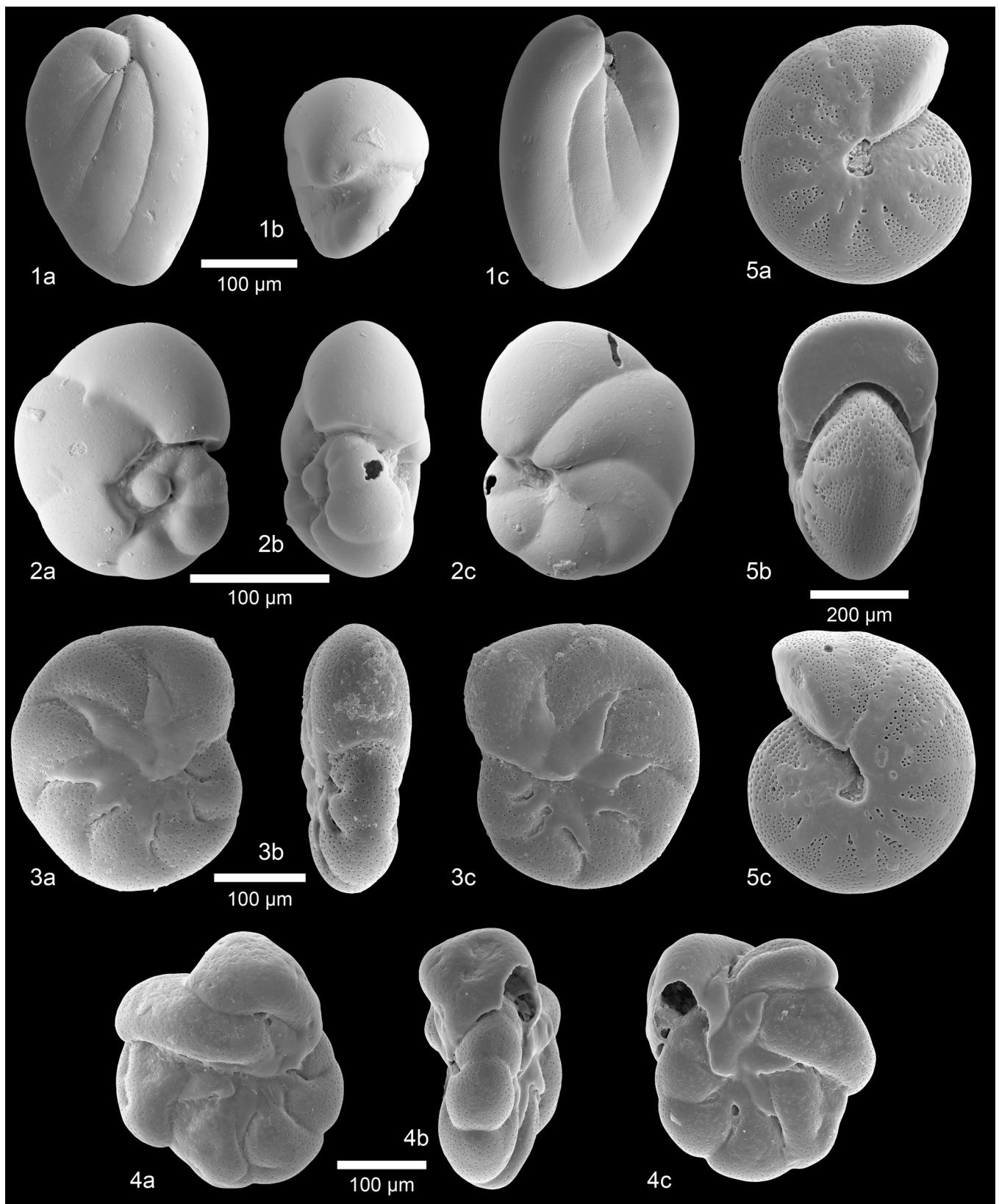


PLATE 38

- 1a-c *Melonis barleeanum* (Williamson, 1858), Hypotype from P292 581-1
2a-c *Pullenia subcarinata* (d'Orbigny, 1839), Hypotype from PS70/023-3
3a-c *Pullenia bulloides* (d'Orbigny, 1826), Hypotype from TTR17 MS419G 25
4a-c *Pullenia bulloides* (d'Orbigny, 1826), Hypotype from Herm1_1
5a-c *Chilostomella oolina* Schwager, 1878, Hypotype from P292 580-1

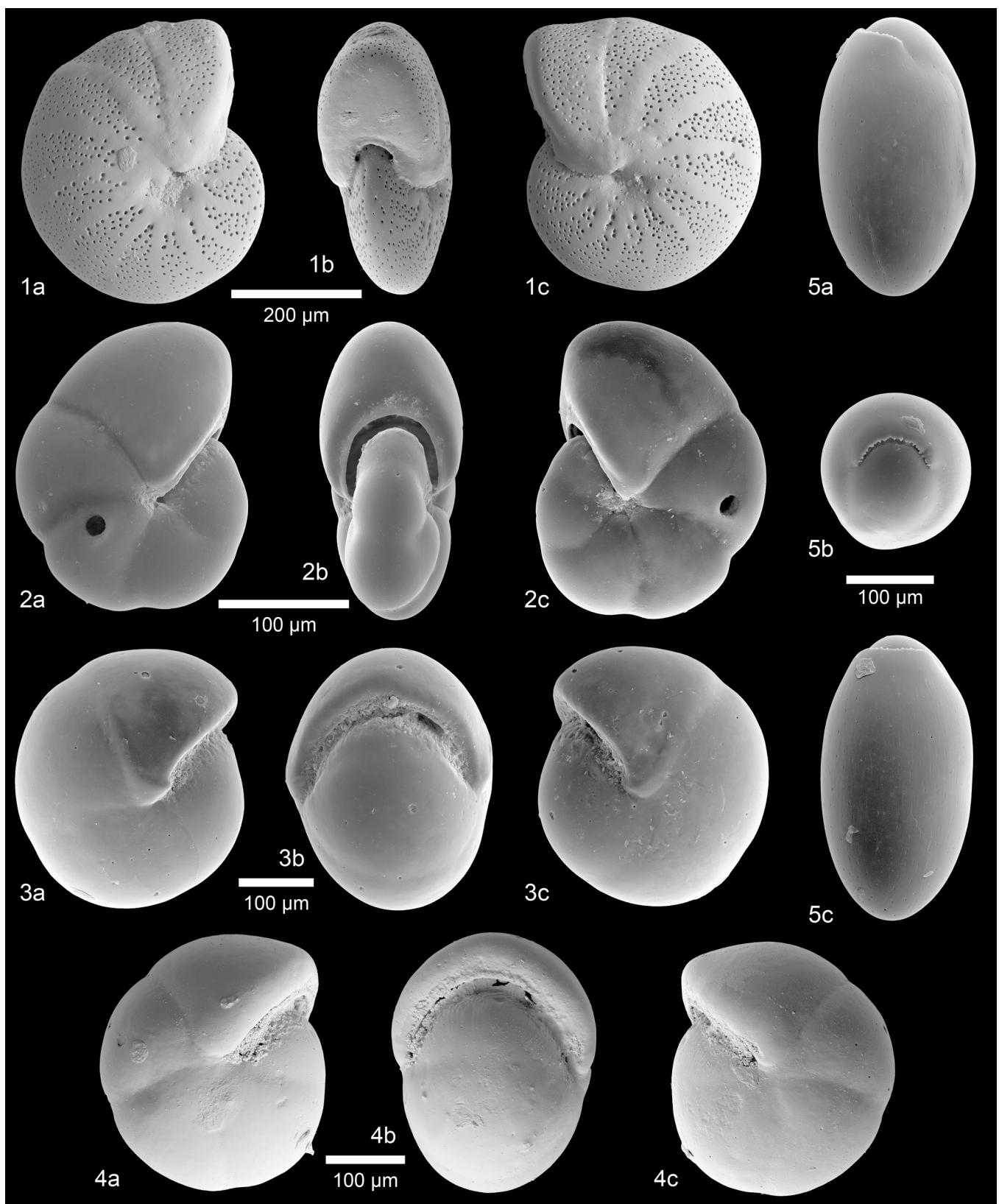


PLATE 39

- 1a-c *Anomalinoides globulosa* (Chapman and Parr, 1937), Hypotype from GeoB 9288
2a-c *Discanomalina coronata* (Parker and Jones, 1857), Hypotype from GeoB9204-1
3a-c *Discanomalina japonica* Asano, 1951, Hypotype from GeoB9205-1
4a-c *Gyroidina laevigata* d'Orbigny, 1826, Hypotype from GeoB 6721-1
5a-c *Gyroidina lamarckiana* (d'Orbigny, 1839), Hypotype from TTR17 MS411G 10

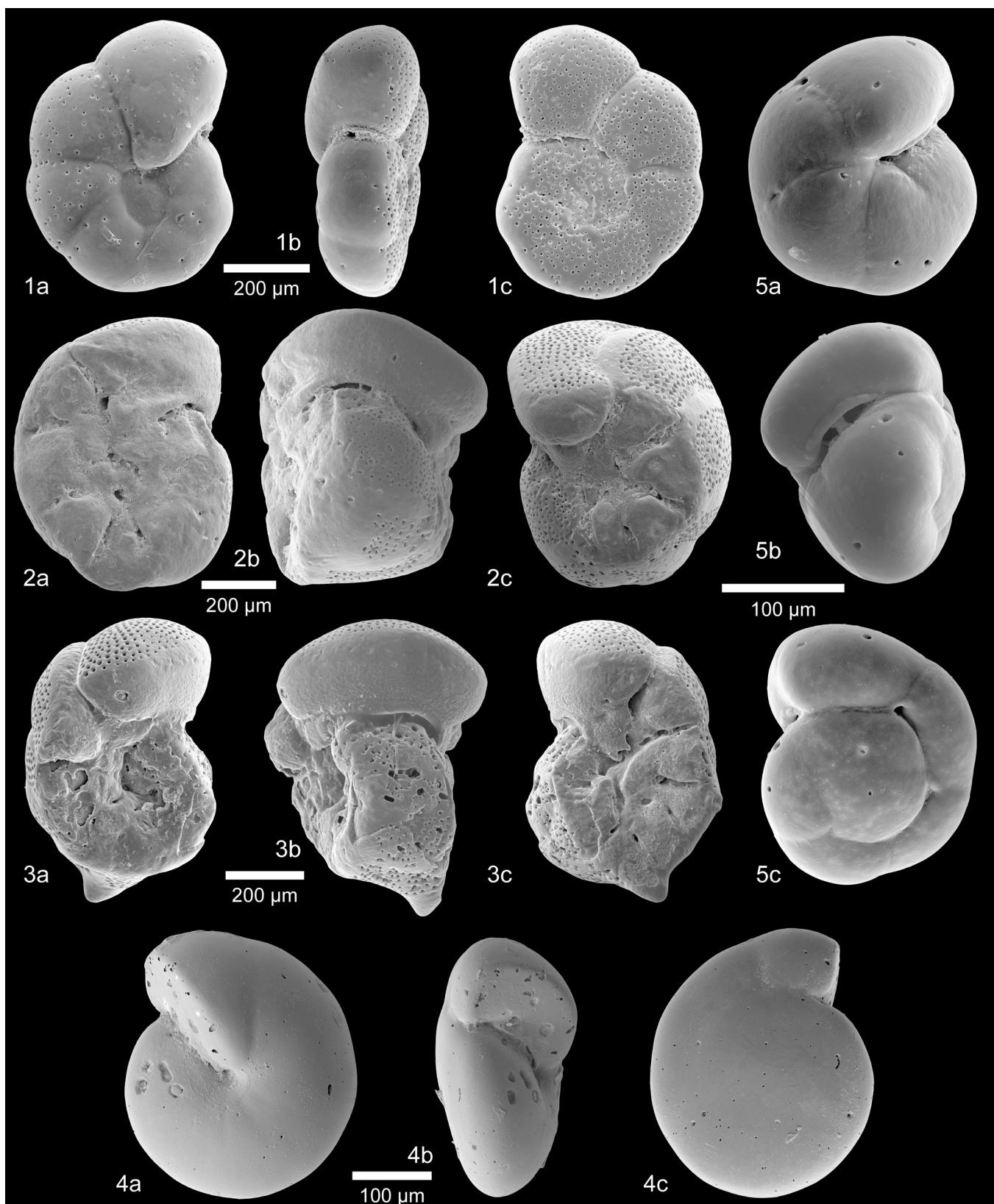


PLATE 40

- 1a-c *Gyroidina soldanii* d'Orbigny, 1839, Hypotype from GeoB 9220
2a-c *Gyroidina neosoldanii* Brotzen, 1936, Hypotype from GeoB9220
3a-c *Gyroidina altiformis* Steward and Steward, 1930, Hypotype from TTR17 MS419G 15
4a-c *Hanzawaia boueana* (d'Orbigny, 1846), Hypotype from GeoB12748-1
5a-c *Hanzawaia boueana* (d'Orbigny, 1846), Hypotype from POS391 558-1

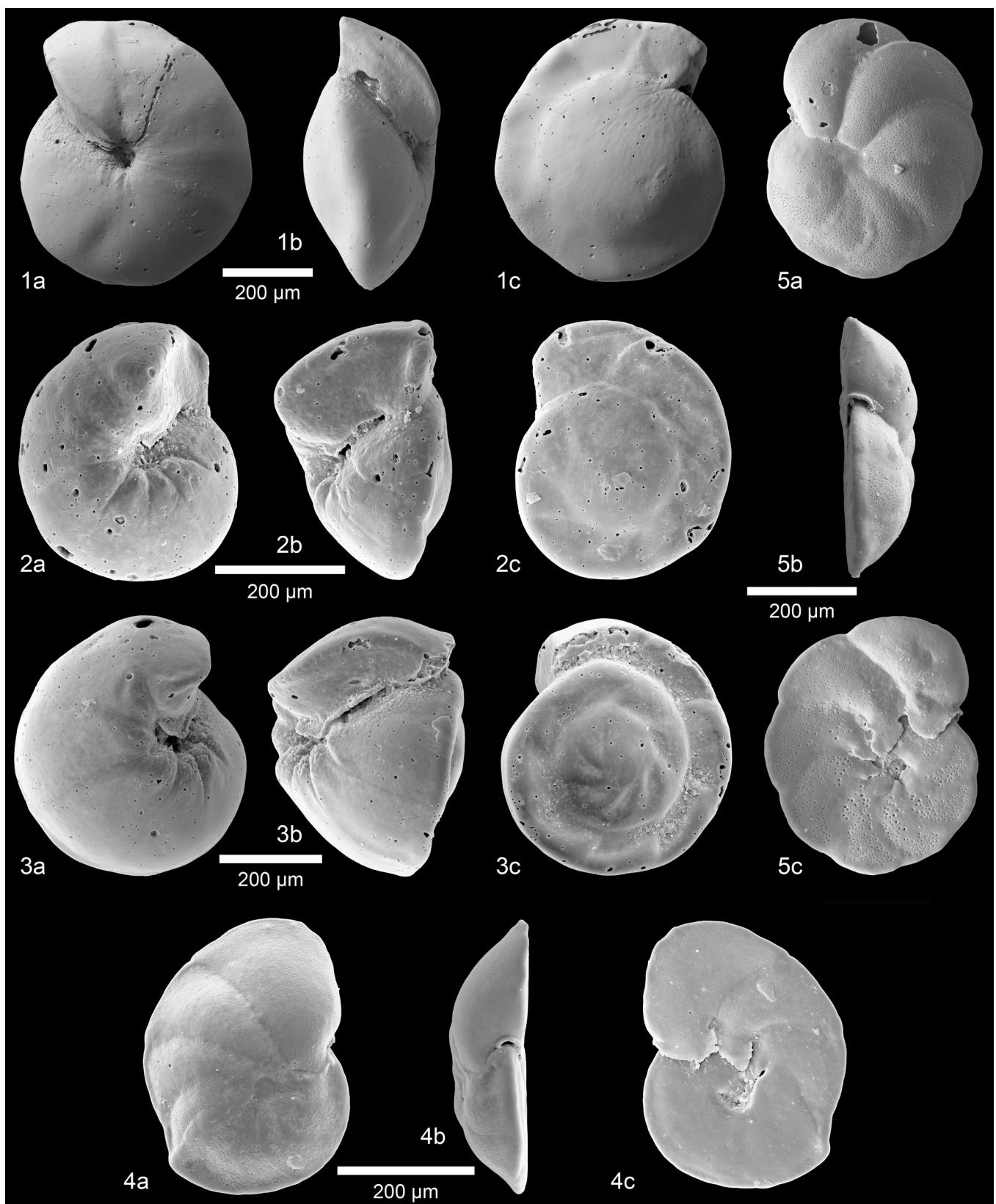


PLATE 41

- 1a-c *Buccella frigida* (Cushman, 1922), Hypotype from POS391 558-1
2a-c *Elphidium albiumbilicatum* (Weiss, 1954), Hypotype from POS391 571-1
3a-c *Elphidium groenlandicum* Cushman, 1933, Hypotype from PS70/039-2
4a-c *Elphidium hanzawai* Asano, 1939, Hypotype from TTR17 MS419G 45

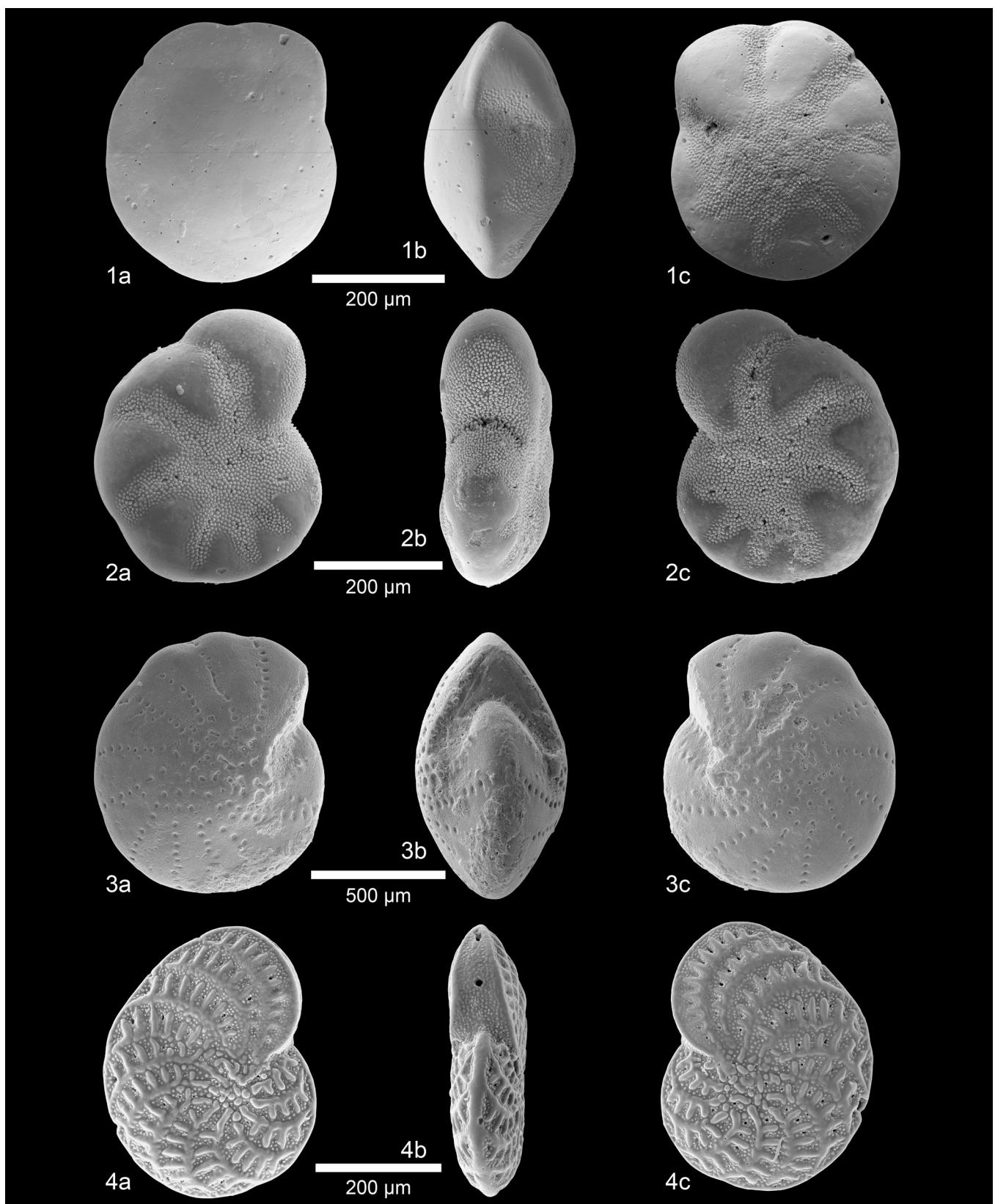
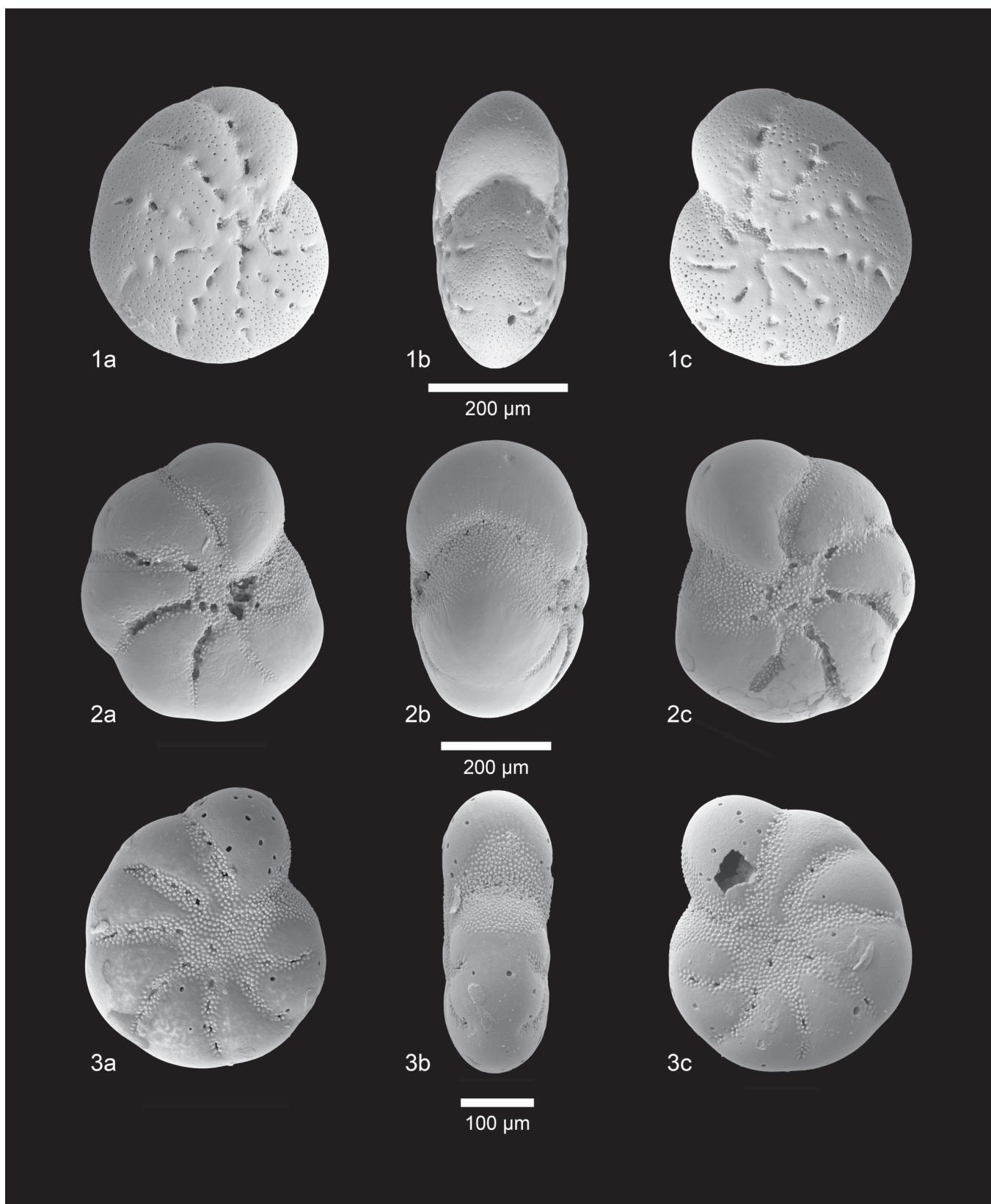


PLATE 42

- 1a-c *Elphidium incertum* (Williamson, 1858), Hypotype from P292 578-1
2a-c *Elphidium magellanicum* Heron-Allen and Earland, 1932, Hypotype from POS391 534-1
3a-c *Elphidium subarcticum* Cushman, 1944, Hypotype from POS391 534-1



ANNEX

Appendix A. Species list and quantitative data of benthic foraminifera from the Porcupine/Rockall region (Chapter 2). Generic names are in accordance with Loeblich and Tappan (1987).

sample number	574-1	576-1	577-1	578-1	579-1	580-1	581-1
sum counts	430	595	627	605	502	375	266
<i>Adercotryma wrigiti</i> Brönnimann and Whittaker, 1987	0	0	0	0	0	0	0
<i>Ammonia beccarii</i> Linne, 1758	0	0	2	3	0	0	0
<i>Pseudoeponides falsobeccarii</i> Rouvillois = <i>Ammonia falsobeccarii</i> Rouvillois, 1974	3	0	0	0	0	0	0
<i>Ammoscalaria pseudospiralis</i> Bätsch = <i>Proteonina pseudospiralis</i> Williamson, 1858	0	0	0	0	0	0	0
<i>Angulogerina scalaris</i> Bätsch = <i>Nautilus scalaris</i> Bätsch, 1791	0	0	0	1	2	1	1
<i>Angulogerina angulosa</i> Williamson = <i>Uvigerina angulosa</i> Williamson, 1858	13	26	58	16	23	10	10
<i>Anomalinoides granosa</i> Hantken, 1875	0	0	0	0	0	0	0
<i>Astacolus crepidulus</i> Fichtel and Moll = <i>Nautilus crepidulus</i> Fichtel and Moll, 1798	0	0	2	0	0	0	0
<i>Astromonion gallorivayi</i> Loeblich and Tappan, 1953	3	0	0	0	0	0	1
<i>Astromonion stelligerum</i> d'Orbigny = <i>Nonionina stelligera</i> d'Orbigny, 1839	0	6	2	1	0	3	0
<i>Bigenenerina nodosa</i> Seguenza = <i>Vulvulinia alata</i> Seguenza, 1862	0	2	5	3	2	1	0
<i>Bolivina dilatata</i> Reuss = <i>Brizalina dilatata</i> Reuss, 1850	5	7	12	23	4	2	0
<i>Bolivina pseudopunctata</i> Heron-Allen and Earland = <i>Bolivina pseudo-plicata</i> Heron-Allen and Earland, 1930	0	0	0	0	0	0	0
<i>Bolivina pseudopunctata</i> Höglund, 1947	0	0	1	1	1	0	0
<i>Bolivina difformis</i> Williamson = <i>Textularia variabilis</i> var. <i>difformis</i> Williamson, 1858	6	1	0	0	0	0	0
<i>Bolivina robusta</i> Brady, 1881	0	0	0	0	0	1	0
<i>Bolivina</i> sp.	0	0	0	0	0	0	0
<i>Bolivina spathulata</i> Williamson = <i>Textularia variabilis</i> var. <i>spathulata</i> Williamson, 1858	13	21	8	21	4	3	1
<i>Bolivina striatula</i> Cushman = <i>Brizalina striatula</i> Cushman, 1922	0	0	0	0	0	0	0
<i>Bolivina subaenaeensis</i> Cushman = <i>Brizalina subaenaeensis</i> Cushman, 1922	0	0	0	0	1	1	0
<i>Bolivina subspinoscens</i> Cushman, 1922	0	0	3	2	0	2	0
<i>Budashhevella multicamerata</i> Voloshinova & Budasheva, 1961	0	0	0	0	0	0	0
<i>Buliminaria aculeata</i> d'Orbigny, 1826	0	10	12	5	8	6	1
<i>Buliminaria marginata</i> d'Orbigny, 1826	12	33	62	51	45	38	11
<i>Buliminaria striata</i> d'Orbigny, 1843	0	0	1	0	0	0	0
<i>Cassidulina carinata</i> Silvestri = <i>Cassidulina laevigata</i> d'Orbigny var. <i>carinata</i> Silvestri, 1896	215	148	67	34	12	15	20
<i>Cassidulina laevigata</i> d'Orbigny, 1826	3	8	33	30	6	20	3
<i>Cibicides refugens</i> de Montfort, 1808	5	0	0	0	0	0	0
<i>Cibicides ungerianus</i> d'Orbigny = <i>Rotalia ungeriana</i> d'Orbigny, 1846	0	0	1	0	4	3	3
<i>Cibicidoides kullenbergi</i> Parker = <i>Cibicides kullenbergi</i> Parker, 1953	12	41	9	0	0	1	5
<i>Cibicidoides pachyderma</i> Rzehak, 1886	39	122	5	5	5	7	3
<i>Cibicides/Cibicidoides</i> juv.	0	0	2	0	15	21	10
<i>Cibicidoides</i> sp.	0	0	0	3	1	0	0
<i>Compspira involvens</i> Reuss = <i>Operculla involvens</i> Reuss, 1850	0	0	0	0	0	0	0
<i>Cribrostomoides jeffreysi</i> Williamson = <i>Nonionina jeffreysi</i> Williamson, 1858	0	0	0	0	0	0	0
<i>Dentalina leguminiformis</i> Bätsch = <i>Nautilus</i> (<i>Orthoceras</i>) leguminiformis Bätsch, 1791	1	0	0	0	0	1	0
<i>Deuterammina rotiformis</i> Heron-Allen and Earland = <i>Trochammina rotiformis</i> Heron-Allen and Earland, 1911	0	0	0	0	0	1	0
<i>Disconammina coronata</i> Parker and Jones = <i>Rotalina semipunctata</i> Bailey, 1857	0	0	0	0	0	0	0
<i>Eggerellida bradyi</i> Cushman, 1911 = <i>Vernuuiina bradyi</i> Cushman, 1911	0	0	0	0	0	0	0
<i>Eggerelloides scabra</i> Williamson = <i>Buliminia scabrum</i> Williamson, 1958	0	0	0	0	0	0	0
<i>Ehrenbergina trigona</i> Goës = <i>Ehrenebergina serrata</i> Brady var. <i>trigona</i> Goës, 1896	0	0	0	0	0	0	0

Appendix A (continued)

sample number	sum counts	574-1	576-1	577-1	578-1	579-1	580-1	581-1
		430	595	627	605	502	375	266
<i>Eiphidium earlandi</i> Cushman, 1936	0	0	0	0	0	0	0	0
<i>Eiphidium gerthii</i> Van Voorthuysen, 1957	0	0	0	0	0	0	0	0
<i>Epistominella exigua</i> Brady = <i>Pulvinulina exigua</i> Brady, 1884	1	3	4	12	29	31	4	
<i>Epistominella vitea</i> Parker, 1953	0	1	4	4	20	26	28	
<i>Fissurina laevigata</i> Reuss, 1850	0	0	0	0	0	0	0	
<i>Fissurina lucida</i> Williamson = <i>Entosolenia marginata</i> Montagu var. <i>lucida</i> Williamson, 1848	0	0	0	0	0	0	0	
<i>Fissurina marginata</i> Montagu = <i>Vermiculum marginatum</i> Montagu, 1803	0	0	0	0	0	0	0	
<i>Fissurina orbigniana</i> Seguenza, 1862	0	0	0	0	0	0	0	
<i>Paliatella semimarginata</i> Reuss = <i>Fissurina semimarginata</i> Reuss, 1870	0	0	0	1	0	0	0	
<i>Fissurina</i> sp.	0	0	0	0	0	0	0	
<i>Fornitella wuerstorffii</i> Schwager = <i>Anomalina wuerstorffii</i> Schwager, 1866	0	3	0	4	0	0	0	
<i>Fusserkoina schreiberiana</i> Carter, 1880 = <i>Virgulina schreiberiana</i> Carter	0	1	0	0	4	0	0	
<i>Gaudryina rufa</i> Wright 1900	0	0	0	0	0	0	0	
<i>Gavelinopsis praegeri</i> Heron-Allen and Earland = <i>Discorbina praegeri</i> Heron-Allen and Earland, 1913	5	2	3	3	3	0	1	
<i>Globobulimina pupoides</i> d'Orbigny = <i>Buliminia pupoides</i> d'Orbigny, 1846	0	0	0	0	0	0	1	
<i>Gavelinopsis</i> sp.	0	0	0	0	0	0	0	
<i>Globocassidulina subglobosa</i> group	64	45	58	39	12	4	4	
<i>Guttulina communis</i> d'Orbigny = <i>Polymorphina (Guttulina) communis</i> d'Orbigny, 1826	0	0	0	0	0	0	0	
<i>Guttulina regina</i> Brady, Parker and Jones = <i>Polymorphina regina</i> Brady, Parker and Jones, 1871	0	0	0	0	0	0	0	
<i>Gyroidea neosoldanii</i> Brocken, 1936	0	0	0	1	2	0	0	2
<i>Gyroidinoides laevigatus</i> d'Orbigny = <i>Gyroidina laevigata</i> d'Orbigny, 1826	0	0	0	0	0	1	0	0
<i>Gyroidina soldanii</i> d'Orbigny, 1826	0	0	0	0	0	0	1	0
<i>Hanzawaia boueana</i> d'Orbigny = <i>Rotalina boueana</i> d'Orbigny, 1846	0	0	0	0	0	0	0	1
<i>Heterolepa haidingeri</i> d'Orbigny = <i>Rotalina haidingeri</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0
<i>Hoeglundina elegans</i> d'Orbigny = <i>Rotalia (Turbulinina) elegans</i> d'Orbigny, 1826	0	1	0	0	0	0	0	1
<i>Hopkinsina atlantica</i> Cushman = <i>Hopkinsina pacifica</i> Cushman, 1933	1	0	4	4	1	9	0	
<i>Hyalinea balthica</i> Schröter = <i>Nautilus balthicus</i> Schröter, 1783	16	30	42	54	51	33	24	
<i>Hyrokkin sarcophaga</i> Cedhagen, 1994	0	0	0	0	0	0	0	
<i>Kameriella bradyi</i> Cushman = <i>Gaudryina bradyi</i> Cushman, 1911	0	0	0	0	0	1	0	
<i>Lagena hispida</i> Reuss, 1863	0	0	0	0	0	0	0	
<i>Lagena spicata</i> Thiele, 1898	0	0	0	0	0	0	0	
<i>Lagena striata</i> d'Orbigny, 1839	1	0	1	0	0	0	0	
<i>Lagena sulcata</i> Walker and Jacob = <i>Serpula sulcata</i> Walker and Jacob, 1798	0	0	0	0	0	1	0	
<i>Lenticulina inornata</i> d'Orbigny = <i>Robulina inornata</i> d'Orbigny, 1846	0	0	0	2	1	0	0	
<i>Lenticulina orbicularis</i> d'Orbigny = <i>Robulina orbicularis</i> d'Orbigny, 1826	0	0	0	0	0	0	1	
<i>Lobatula lobatula</i> Walker and Jacob = <i>Nautilus lobatulus</i> Walker and Jacob, 1798	5	4	1	0	1	0	1	
<i>Melonis bardeeanum</i> Williamson = <i>Nonionina bardeana</i> Williamson, 1858	0	2	12	9	15	16	17	
<i>Melonis pomphiloides</i> Fichtel and Moll, 1798 = <i>Nautilus pomphiloides</i> Fichtel and Moll, 1798	0	0	0	0	0	1	0	
<i>Melonis</i> sp.	0	0	0	0	0	0	0	
<i>Miliolinella subrotunda</i> Montagu, 1803 = <i>Vermiculum subrotundum</i> Montagu, 1803	0	0	0	0	0	0	0	
<i>Neconorbina terquemi</i> Rzehak = <i>Discorbina terquemi</i> Rzehak, 1888	0	0	0	0	0	0	0	
<i>Nomionella iridea</i> Heron-Allen and Earland, 1932	0	0	0	0	0	10	1	
<i>Nomionellina labradorica</i> Dawson = <i>Nonionina labradorica</i> Dawson, 1860	0	1	0	0	0	0	0	

Appendix A (continued)

sample number	574-1	576-1	577-1	578-1	579-1	580-1	581-1
sum counts	430	595	627	605	502	375	266
<i>Nonionella turgida</i> Williamson = <i>Rotalina turgida</i> Williamson, 1858	1	3	4	5	13	9	1
<i>Nuttallides umboniferus</i> Cushman = <i>Pulvinulina umbonifera</i> Cushman, 1933	0	0	0	0	1	0	1
<i>Favulinula hexagona</i> Williamson = <i>Entosolenia squamosa</i> Montagu var. <i>hexagona</i> Williamson, 1858	0	0	0	0	0	0	0
<i>Homalohedra williamsoni</i> Alcock = <i>Entosolenia williamsoni</i> Alcock, 1885	0	0	0	0	1	0	0
<i>Ophthalmaeum filianeris</i> Rhambler, 1936	0	0	0	0	0	0	0
<i>Oridorsalis umbonatus</i> Reuss = <i>Rotalina umbonatus</i> Reuss, 1851	0	0	0	0	1	0	0
<i>Paratrochammina challengerii</i> Brönnimann and Whittaker, 1988	0	0	0	1	0	1	0
<i>Paratrochammina murrayi</i> Brönnimann and Zaninetti, 1984	0	0	1	0	0	0	0
<i>Planulina arminensis</i> d'Orbigny, 1826	0	0	0	0	0	0	3
<i>Porosonion ex gr. Granosum</i> d'Orbigny = <i>Nonionina granosa</i> d'Orbigny, 1826	0	5	9	11	1	2	0
<i>Pullenia bulloides</i> d'Orbigny = <i>Nonionina bulloides</i> d'Orbigny, 1846	0	0	0	0	0	0	0
<i>Pullenia quinqueloba</i> Reuss = <i>Nonionina quinqueloba</i> Reuss, 1851	0	0	0	0	0	2	5
<i>Pyrgo anomala</i> Schlumberger = <i>Biloculina anomala</i> Schlumberger, 1891	0	0	0	0	0	0	0
<i>Pyrgo elongata</i> d'Orbigny = <i>Biloculina elongata</i> d'Orbigny, 1826	0	0	0	0	0	0	0
<i>Pyrgo inornata</i> d'Orbigny = <i>Biloculina elongata</i> d'Orbigny, 1839	0	0	0	0	0	0	0
<i>Pyrgo murrhina</i> Schwager = <i>Biloculina murrhina</i> Schwager, 1866	0	0	0	0	0	0	0
<i>Pyrgo subsphaerica</i> d'Orbigny = <i>Biloculina subsphaerica</i> d'Orbigny, 1840	0	0	0	0	0	0	0
<i>Quinqueloculina arctica</i> Cushman, 1933	0	0	0	0	0	0	0
<i>Cycloforina stalkeri</i> Loeblich and Tappan, 1953	0	0	0	0	0	0	0
<i>Reophax scorpiurus</i> de Montfort, 1808	0	1	0	0	0	0	0
<i>Robertinoides subcylinindrica</i> Brady = <i>Bulimia subcylinindrica</i> Brady, 1881	0	0	0	0	0	0	0
<i>Rosalina</i> sp.	0	0	0	0	0	0	0
<i>Rosalina vilardoebana</i> d'Orbigny, 1839	0	0	0	0	1	0	0
<i>Saccammina sphaerica</i> Sars, 1868	0	0	0	1	0	0	0
<i>Saccorhiza ramosa</i> Brady = <i>Hyperammina ramosa</i> Brady, 1879	0	0	0	0	0	0	0
<i>Sigmollinella tenuis</i> Czizek = <i>Quinqueloculina tenuis</i> Czizek, 1848	0	0	0	0	0	1	1
<i>Sigmollinopsis schlumbergeri</i> Silvestri = <i>Sigmollinella schlumbergeri</i> Silvestri, 1904	0	0	0	1	1	0	6
<i>Sphaeroidina bulloides</i> d'Orbigny, 1826	0	0	0	0	2	0	1
<i>Springilla vivipara</i> Ehrenberg, 1843	0	0	0	0	0	0	0
<i>Spiroplectinella sagittula</i> d'France, 1824	0	0	0	0	1	0	0
<i>Spiroplectinella wrighti</i> Silvestri = <i>Spiroplectla wrighti</i> Silvestri, 1903	2	6	2	1	1	3	3
<i>Stainforthia fusiformis</i> Williamson = <i>Bulimina papoides</i> var. <i>fusiformis</i> Williamson, 1858	1	2	2	0	3	0	2
<i>Stomatorbina concentrica</i> Parker and Jones = <i>Pulvinulina concentrica</i> Parker and Jones, 1864	0	0	0	0	0	0	0
<i>Textularia agglutinans</i> d'Orbigny, 1839	0	0	0	2	2	1	1
<i>Textularia bigenerhoidea</i> Lacroix, 1932	0	0	0	0	1	0	0
<i>Textularia pseudogrammen</i> d'Orbigny, 1839	0	0	0	0	0	0	0
<i>Textularia truncata</i> Höglund, 1947	0	3	0	0	0	0	0
<i>Tosalia hanzawai</i> Takayanagi, 1953	0	0	0	0	0	0	0
<i>Trifaria bradyi</i> Cushman, 1923	0	0	0	0	0	0	0
<i>Uvigerina auberiana</i> d'Orbigny, 1839	0	0	0	0	0	0	2
<i>Uvigerina mediterranea</i> Hofker, 1932	0	8	32	43	32	36	17
<i>Uvigerina peregrina</i> Cushman, 1923	2	44	157	197	172	47	70
<i>Pulvinulina pennatula</i> Bartsch = <i>Nautilus (Orthoceras) pennatula</i> Bartsch, 1791	0	0	0	0	0	0	0

Appendix A (continued)

Appendix A (continued)

sample number	9220	9256-1	9271	9287	9257	9260	9268	9269	9288
sum counts	124	131	79	123	95	63	118	68	81
<i>Adercytoma wrightii</i> Brönnimann and Whittaker, 1987	0	0	0	2	0	0	0	0	0
<i>Ammonia beccarii</i> Linné, 1758	0	0	0	0	0	0	0	0	0
<i>Pseudoponides falsobeccearii</i> Rouvillois = <i>Ammonia falsobeccearii</i> Rouvillois, 1974	0	0	0	0	0	0	0	0	0
<i>Ammoscalaria pseudospiralis</i> Williamson = <i>Proteonina pseudospiralis</i> Williamson, 1858	0	0	0	0	0	0	0	0	0
<i>Amphiconya scalaris</i> Bartsch = <i>Nautilus scalaris</i> Bartsch, 1791	0	0	0	0	0	0	0	0	0
<i>Angulogerina angulosa</i> Williamson = <i>Uvigerina angulosa</i> Williamson, 1858	0	2	3	13	2	3	10	2	9
<i>Anomalinoides granosa</i> Hantken, 1875	0	0	0	0	0	0	0	0	1
<i>Astacolus crepidulus</i> Fichtel and Moll = <i>Nautilus crepidulus</i> Fichtel and Moll, 1798	0	0	0	0	0	0	2	0	0
<i>Astromonion gallawayi</i> Loeblich and Tappan, 1953	0	0	0	0	0	0	0	0	0
<i>Bigennerina nodosaria</i> d'Orbigny = <i>Nanionina stelligera</i> d'Orbigny, 1839	3	1	2	4	2	1	2	0	0
<i>Bolivina alata</i> Sequenza = <i>Vulvulina alata</i> Sequenza, 1862	0	0	0	0	0	0	0	0	0
<i>Bolivina dilatata</i> Reuss = <i>Brizalina dilatata</i> Reuss, 1850	3	0	0	5	0	0	1	0	1
<i>Bolivina pseudoplicata</i> Heron-Allen and Earland = <i>Bolivina pseudo-plicata</i> Heron-Allen and Earland, 1930	0	0	0	0	0	0	0	0	0
<i>Bolivina pseudopunctata</i> Höglund, 1947	0	0	0	1	0	0	1	0	0
<i>Bolivina difformis</i> Williamson = <i>Textularia variabilis</i> var. <i>difformis</i> Williamson, 1858	0	0	1	0	0	0	1	0	0
<i>Bolivina robusta</i> Brady, 1881	0	0	0	0	0	0	0	0	0
<i>Bolivina</i> sp.	0	0	0	0	0	0	0	0	0
<i>Bolivina spathulata</i> Williamson = <i>Textularia variabilis</i> var. <i>spathulata</i> Williamson, 1858	0	1	0	3	1	0	0	0	0
<i>Bolivina striatula</i> Cushman = <i>Brizalina striatula</i> Cushman, 1922	0	0	0	0	0	0	0	0	0
<i>Bolivina subaenariensis</i> Cushman = <i>Brizalina subaenariensis</i> Cushman, 1922	0	0	0	0	0	0	0	0	0
<i>Bolivina subspinifrons</i> Cushman, 1922	0	0	0	0	0	0	0	0	1
<i>Budashnevella multicamerata</i> Voloshinova & Budasheva, 1961	0	0	0	0	0	0	0	0	0
<i>Buliminia aculeata</i> d'Orbigny, 1826	1	1	4	2	3	5	7	2	3
<i>Buliminia marginata</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0
<i>Buliminia striata</i> d'Orbigny, 1843	0	0	0	0	0	0	0	0	0
<i>Cassidulina carinata</i> Silvestri = <i>Cassidulina laevigata</i> d'Orbigny var. <i>carinata</i> Silvestri, 1896	1	0	0	2	1	0	0	4	3
<i>Cassidulina laevigata</i> d'Orbigny, 1826	6	1	3	5	6	1	11	10	3
<i>Cibicides refugens</i> de Montfort, 1808	1	4	0	6	0	2	1	0	0
<i>Cibicides ungerianus</i> d'Orbigny = <i>Rotalia ungeriana</i> d'Orbigny, 1846	2	1	1	0	0	0	0	0	0
<i>Cibicidoides kullenbergi</i> Parker = <i>Cibicides kullenbergi</i> Parker, 1953	0	0	0	0	3	1	1	0	3
<i>Cibicidoides pachyderma</i> Rzehak, 1886	22	0	3	0	0	2	0	2	3
<i>Cibicidoides/Cibicidoides</i> juv.	0	0	0	0	0	0	0	0	0
<i>Cibicidoides</i> sp.	0	0	2	3	10	3	1	5	10
<i>Cornuspira involvens</i> Reuss = <i>Operculina involvens</i> Reuss, 1850	0	1	0	0	0	0	0	0	0
<i>Cribrostomoides jeffreysi</i> Williamson = <i>Nonionina jeffreysi</i> Williamson, 1858	0	0	0	0	0	0	0	0	0
<i>Dentalina leguminiformis</i> Bartsch = <i>Nautilus</i> (<i>Orthoceras</i>) <i>leguminiformis</i> Bartsch, 1791	0	0	0	0	0	0	0	0	0
<i>Deuterammina rotaliformis</i> Heron-Allen and Earland = <i>Trichammina rotaliformis</i> Heron-Allen and Earland, 1911	0	0	0	0	0	0	0	0	0
<i>Discanomolina coronata</i> Parker and Jones = <i>Anomalina coronata</i> Parker and Jones, 1857	13	2	0	4	1	0	6	0	0
<i>Discanomolina semipunctata</i> Bailey, 1851	0	1	0	0	0	0	0	0	0
<i>Eggerelloides bradyi</i> Cushman, 1911 = <i>Vermulima bradyi</i> Cushman, 1911	0	0	0	0	0	0	0	0	0
<i>Eggerelloides scabra</i> Williamson = <i>Bulimina secastrum</i> Williamson, 1958	5	0	0	0	0	3	0	0	0
<i>Ehrenbergina trigona</i> Goës = <i>Ehrenbergina serrata</i> Brady var. <i>trigona</i> Goës, 1896	0	1	0	0	0	0	0	0	0

Appendix B. Species list and quantitative data of benthic foraminifera from the Norwegian Margin (Chapter 3). Generic names are in accordance with Loeblich and Tappan (1987).

sample number	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
sum counts	540	450	518	552	266	537	270	317	528	373	628	626	376	291	370	821
<i>Adercotyma wrightii</i> Brönnimann and Whittaker, 1987	0	4	0	0	0	0	0	4	0	2	0	0	0	1	8	2
<i>Ammobaculites agglutinans</i> d'Orbigny = <i>Spirilolina agglutinans</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0
<i>Ammodiscus incertus</i> d'Orbigny = <i>Opercucina incerta</i> d'Orbigny, 1839	0	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Ammolagena clavata</i> Jones and Parker = <i>Trochammina irregularis</i> d'Orbigny var. <i>clavata</i> Jones and Parker, 1860	0	1	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Amphicorona scalaris</i> Batsch = <i>Nautilus scalaris</i> Batsch, 1791	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Angulogerina angulosa</i> Williamson = <i>Uvigerina angulosa</i> Williamson, 1858	31	13	61	87	32	32	36	27	16	8	4	0	51	71	5	0
<i>Astronion gallicum</i> Loeblich and Tappan, 1953	16	49	9	1	3	0	1	2	0	1	1	0	1	3	7	2
<i>Bathygiphon filiformis</i> M. Sars, 1872	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0
<i>Bigenenerina cylindrica</i> Cushman, 1922	0	0	0	0	0	0	0	0	0	0	0	0	0	0	4	37
<i>Bigenenerina nodosaria</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	0	0	0	0	2	0	1
<i>Biloculinella depreressa</i> Wiesner = <i>Biloculina labiata</i> Schubert var. <i>depreressa</i> Wiesner, 1923	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Biloculinella globula</i> Bonnemain = <i>Biloculina globulus</i> Bonnemain, 1855	0	0	1	0	0	0	0	0	0	1	0	0	0	0	0	0
<i>Bolivina alata</i> Seguenza = <i>Vulvulina alata</i> Seguenza, 1862	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Bolivina difformis</i> Williamson = <i>Textularia variabilis</i> var. <i>difformis</i> Williamson, 1850	1	0	2	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Bolivina dilatata</i> Reuss = <i>Brizalina dilatata</i> Reuss, 1850	0	0	0	0	0	0	0	0	0	0	0	0	0	0	6	0
<i>Bolivina pseudopunctata</i> Heron-Allen and Earland = <i>Bolivina pseudo-plicata</i> Heron-Allen and Earland, 1930	2	1	0	1	0	0	0	0	0	0	0	0	0	0	4	0
<i>Bolivina pseudopunctata</i> Hoglund, 1947	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Bolivina spathulata</i> Williamson = <i>Textularia variabilis</i> var. <i>spathulata</i> Williamson, 1858	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	4
<i>Bolivina spinescens</i> Cushman, 1911	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Bolivina striatula</i> Cushman = <i>Brizalina striatula</i> Cushman, 1922	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Bolivina subspinescens</i> Cushman, 1922	0	1	1	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Botellina labyrinthica</i> Brady, 1881	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Buccella frigida</i> Cushman = <i>Pulvinulina frigida</i> Cushman, 1922	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
<i>Buccella tenerima</i> Bandy = <i>Rotalia tenerima</i> Bandy, 1950	1	0	0	0	0	1	0	0	1	0	0	0	2	0	0	0
<i>Bullimina aculeata</i> d'Orbigny, 1826	0	0	1	0	0	0	0	0	0	0	0	0	0	1	0	0
<i>Bullimina marginata</i> d'Orbigny, 1826	0	0	0	5	1	3	0	0	1	0	1	0	1	2	0	28
<i>Cassidulina carinata</i> Silvestri = <i>Cassidulina laevigata</i> d'Orbigny var. <i>carinata</i> Silvestri, 1896	13	4	29	13	3	11	4	1	23	0	9	1	24	13	11	69
<i>Cassidulina laevigata</i> group	8	7	11	1	1	6	34	31	52	90	2	3	1	5	16	8
<i>Cassidulinoidea oblonga</i> Reuss = <i>Cassidulina oblonga</i> Reuss, 1850	0	0	1	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Cibicides refugens</i> de Montfort, 1808	9	0	17	37	6	78	4	0	0	0	1	0	17	12	3	0
<i>Cibicides ungerianus</i> d'Orbigny = <i>Rotalia ungeriana</i> d'Orbigny, 1846	12	1	0	1	1	7	0	1	0	0	0	2	3	1	1	0
<i>Cibicides/Cibicidoides juv.</i>	42	0	12	13	0	1	0	1	0	1	0	0	0	0	0	0
<i>Cibicidoides kullenbergi</i> Parker = <i>Cibicides kullenbergi</i> Parker, 1953	3	16	31	2	13	7	0	0	0	0	0	0	3	15	10	0
<i>Cibicidoides</i> sp.	192	85	113	88	33	191	17	6	18	1	1	0	32	11	49	0
<i>Cibicidoides pachyderma</i> Rzehak, 1886	0	0	0	0	0	2	0	0	0	0	0	0	0	0	0	0
<i>Cibicidoides pseudoungerianus</i> Cushman = <i>Truncatulina pseudoungeriana</i> Cushman, 1922	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Clavulina primaeva</i> Cushman, 1913	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Comuspira foliacea</i> Philippi = <i>Orbis foliaceus</i> Philippi, 1844	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Comuspira involvens</i> Reuss = <i>Opercucina involvens</i> Reuss, 1850	0	1	4	0	0	0	0	0	0	1	0	0	0	0	0	0
<i>Cribroelphidium excavatum</i> Terquem = <i>Polystomella excavata</i> Terquem, 1875	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	0
<i>Cribrostomoides jeffreysi</i> Williamson = <i>Nonionina jeffreysi</i> Williamson, 1858	0	10	3	0	0	1	0	0	0	0	0	0	0	0	0	0
<i>Cribrostomoides subglobosum</i> Cushman = <i>Haplophragmoides subglobosum</i> , Cushman, 1910	0	0	0	2	0	2	6	41	49	174	111	0	0	14	0	0
<i>Cushmania plumigera</i> Brady = <i>Lagenia plumigera</i> Brady, 1881	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0

Appendix B (continued)

sample number	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
sum counts	540	450	518	552	266	537	270	317	528	373	628	626	376	291	370	821
<i>Cycloforina angularis</i> d'Orbigny = <i>Quinqueloculina angularis</i> d'Orbigny, 1905	0	0	0	0	0	0	0	2	1	0	4	4	0	0	0	0
<i>Cycloforina laevigata</i> d'Orbigny, 1839	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Cycloforina starkeri</i> Loeblich and Tappan, 1953	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Dentalina advena</i> Cushman = <i>Nodosaria advena</i> Cushman, 1923	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Discarmina compressa</i> Goes, 1882	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Discarmonalina coronata</i> Parker and Jones = <i>Anomalina coronata</i> Parker and Jones, 1857	48	47	14	6	15	18	1	0	0	0	0	0	0	6	1	0
<i>Discorbina bertheleti</i> d'Orbigny = <i>Rosalinia bertheloti</i> d'Orbigny, 1839	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Eggerellides scabra</i> Williamson = <i>Buliminina scabrum</i> Williamson, 1958	0	0	1	0	0	0	4	3	0	0	0	0	0	0	0	0
<i>Elphidiella hawaii</i> Cushman and Grant = <i>Elphidium hawaii</i> Cushman and Grant, 1927	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1
<i>Elphidiella albumbilicata</i> Weiss = <i>Nonion pauciloculum</i> Cushman subsp. <i>albumbilicatum</i> Weiss, 1954	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Elphidium clavatum</i> Cushman = <i>Elphidium incertum</i> Williamson var. <i>clavatum</i> Cushman, 1930	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Elphidium frigidum</i> Cushman, 1933	1	1	0	0	2	10	0	0	0	0	0	2	0	0	0	8
<i>Elphidium greenlandicum</i> Cushman, 1933	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Elphidium incertum</i> Williamson = <i>Polystomella umbiliicata</i> Walker var. <i>incerta</i> Williamson, 1858	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Elphidium subarcicum</i> Cushman, 1944	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Epistominella exigua</i> Brady = <i>Psilostomella exigua</i> Brady, 1884	4	2	8	27	6	4	6	22	23	36	53	20	1	2	13	4
<i>Epistominella vitrea</i> Parker, 1953	8	0	11	38	30	16	18	11	19	16	5	9	20	37	1	0
<i>Favolina hexagona</i> Williamson = <i>Entosolenia squamosa</i> Montagu var. <i>hexagona</i> Williamson, 1858	0	1	2	0	0	1	0	0	0	0	0	0	1	0	1	0
<i>Favolina melo</i> d'Orbigny = <i>Oolina melo</i> d'Orbigny, 1839	0	0	0	0	1	0	0	0	0	0	0	0	0	0	2	0
<i>Favolina squamosa</i> Montagu = <i>Vermiculum squamosum</i> Montagu, 1803	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
<i>Fissurina annectens</i> Burrows and Holland = <i>Lagena clathrata</i> Brady, 1884	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina clathrata</i> Brady = <i>Lagena clathrata</i> Brady, 1884	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina curvifibulosa</i> McCulloch, 1977	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina derogata</i> McCulloch, 1977	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina eburnea</i> Buchner, 1940	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina infatiperforata</i> McCulloch, 1977	2	1	1	1	1	1	0	1	0	0	0	0	1	2	0	0
<i>Fissurina kerqueiensis</i> Parr, 1950	0	0	0	1	0	0	0	0	0	0	0	0	1	0	0	0
<i>Fissurina laevigata</i> Reuss, 1850	0	1	1	0	2	0	0	0	0	0	0	0	0	0	2	0
<i>Fissurina lucida</i> Williamson = <i>Entosolenia marginata</i> Montagu var. <i>lucida</i> Williamson, 1848	2	1	1	1	1	1	1	0	0	0	0	0	1	2	0	1
<i>Fissurina marginata</i> Montagu = <i>Vermiculum marginatum</i> Montagu, 1803	0	0	0	1	0	0	0	0	0	0	0	0	1	0	0	0
<i>Fissurina orbigniana</i> Seguenza, 1862	0	0	0	1	0	0	0	0	0	0	0	0	1	0	0	0
<i>Fissurina pseudoorbigniana</i> Buchner = <i>Lagena pseudoorbigniana</i> Buchner, 1940	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina quadricostulata</i> Reuss = <i>Lagena quadricostulata</i> Reuss, 1870	0	0	0	1	0	2	0	1	0	0	0	0	0	0	0	0
<i>Fontotria wuerstorffii</i> Schwager = <i>Anomalina wuerstorffii</i> Schwager, 1866	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Gaudryina rufis</i> Wright 1900	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Gavelinopsis caledonia</i> Murray and Whittaker, 2001	0	11	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Gavelinopsis nitida</i> Williamson = <i>Rotalina nitida</i> , Williamson 1858	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Gavelinopsis praegeri</i> Heron-Allen and Earland = <i>Discorbina praegeri</i> Heron-Allen and Earland, 1913	4	7	12	0	6	2	1	0	3	0	0	0	0	0	0	8
<i>Glandulina laevigata</i> d'Orbigny = <i>Nodosaria laevigata</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	1
<i>Glandulina rotundata</i> Reuss, 1850	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globobulimina affinis</i> d'Orbigny = <i>Bulimina affinis</i> d'Orbigny, 1839	1	1	0	1	0	0	0	0	1	0	0	0	0	0	1	231
<i>Globocassidulina subglobosa</i> group	41	14	46	23	15	9	14	30	70	46	30	4	21	33	3	0

Appendix B (continued)

	sample number	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16
	sum counts	540	450	518	552	268	537	270	317	528	373	628	626	376	291	370	821
<i>Nanionella fabricatorica</i> Dawson = <i>Nanionina labradorica</i> Dawson, 1860		2	0	1	0	0	1	0	0	0	0	0	0	1	0	0	2
<i>Nanionella turgida</i> Williamson = <i>Rotalina turgida</i> Williamson, 1858		4	0	0	10	1	0	0	0	1	2	0	0	2	1	1	3
<i>Nuttallides umboferus</i> Cushman = <i>Pulvinulina umbonifera</i> Cushman, 1933		0	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Oolina globosa</i> Montagu = <i>Vermiculum globosum</i> Montagu, 1803		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Ondorsalis umbonatus</i> Reuss = <i>Rotalina umbonatus</i> Reuss, 1851		0	0	0	5	0	0	0	1	1	0	12	15	0	0	0	0
<i>Ovulina striata</i> Seguenza, 1862		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Palioletella semimarginata</i> Reuss = <i>Fissurina semimarginata</i> Reuss, 1870		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Parafissurina botelliiformis</i> Brady = <i>Lagena botelliiformis</i> Brady, 1881		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Parafissurina felsinea</i> Fornasini = <i>Lagena felsinea</i> Fornasini, 1894		0	0	0	0	0	0	0	0	0	2	0	0	0	0	0	0
<i>Parafissurina marginoradiata</i> McCulloch, 1977		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Parafissurina subquadrata</i> Parr = <i>Fissurina subquadrata</i> Parr, 1945		1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Paratrichammina challengeri</i> Brönnmann and Whittaker, 1988		0	0	0	1	1	1	1	6	2	4	4	0	0	1	3	
<i>Patellina corrugata</i> Williamson, 1858		5	5	0	2	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pelosina cylindrica</i> Brady, 1884		0	0	0	0	0	0	1	0	7	0	0	0	0	0	0	0
<i>Planulina ariminensis</i> d'Orbigny, 1826		0	0	0	0	0	0	1	5	0	0	0	0	5	0	0	0
<i>Procerolagena gracilis</i> Costa = <i>Amphorina gracilis</i> Costa, 1856		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	3
<i>Proteiphidium anglicum</i> Murray, 1965		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pseammoeba fusca</i> Schulze, 1875		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pseudoeponides falsobecarii</i> Rouvillois = <i>Ammonia falsobecarii</i> Rouvillois, 1974		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pullenia bulloides</i> d'Orbigny = <i>Nanionina bulloides</i> d'Orbigny, 1846		2	0	0	1	1	0	0	0	0	0	0	0	0	0	0	0
<i>Pullenia ostensis</i> Feijling-Hanssen = <i>Pullenia quinquefasciata</i> Reuss var. <i>minuta</i> Feijling-Hanssen, 1954		0	3	0	3	2	1	2	1	0	0	0	0	0	1	0	0
<i>Pullenia quadriloba</i> Reuss = <i>Pullenia compressifuscula</i> Reuss var. <i>quadriloba</i> Reuss, 1867		2	1	1	2	0	3	5	6	8	0	0	0	0	0	0	0
<i>Pullenia quinqueloba</i> Reuss = <i>Nanionina quinqueloba</i> Reuss, 1851		10	7	0	1	3	0	2	1	3	11	1	1	0	0	0	0
<i>Pullenia subcarinata</i> d'Orbigny = <i>Nanionina subcarinata</i> d'Orbigny, 1839		0	1	0	4	7	2	1	1	4	0	5	2	0	0	6	59
<i>Pyrgo anomala</i> Schlämberger = <i>Biloculina anomala</i> Schlämberger, 1891		0	1	0	0	1	0	1	0	0	0	0	0	0	0	0	0
<i>Pyrgo constricta</i> Costa = <i>Biloculina constricta</i> Costa, 1866		1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pyrgo elongata</i> d'Orbigny = <i>Biloculina elongata</i> d'Orbigny, 1826		1	2	0	1	0	0	0	0	0	0	0	0	1	0	0	0
<i>Pyrgo lucernula</i> Schwager = <i>Biloculina lucernula</i> Schwager, 1866		0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pyrgo oblonga</i> d'Orbigny = <i>Biloculina oblonga</i> d'Orbigny, 1839		0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pyrgo rotularia</i> Loeblich and Tappan, 1953		1	0	0	0	0	0	0	0	1	0	0	0	0	0	0	2
<i>Pyrgo sarsi</i> Schlämberger = <i>Biloculina sarsi</i> Schlämberger, 1891		0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pyrgo subsphaerica</i> d'Orbigny = <i>Biloculina subsphaerica</i> d'Orbigny, 1840		0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Pyrgo williamsoni</i> Silvestri = <i>Biloculina williamsoni</i> Silvestri, 1923		1	1	0	0	0	0	0	0	0	0	0	0	0	1	0	0
<i>Pyrulina gutta</i> d'Orbigny = <i>Polymorphina</i> (<i>Pyruline</i>) <i>gutta</i> d'Orbigny, 1826		0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Quinqueloculina lamarkiana</i> d'Orbigny, 1859		2	3	0	0	0	2	1	0	0	0	0	0	0	0	0	0
<i>Quinqueloculina arctica</i> Cushman, 1933		0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Quinqueloculina canariensis</i> d'Orbigny, 1839		0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Quinqueloculina viennensis</i> Le Calvez and Le Calvez, 1958		8	0	0	0	0	3	5	0	0	0	0	0	0	0	0	0
<i>Recinvooides trochamminiformis</i> Höglund, 1947		0	0	0	0	0	0	0	0	0	0	0	0	0	4	0	0
<i>Reophax diffugiformis</i> Brady, 1879		1	1	0	0	0	0	0	0	0	0	0	0	0	1	0	0
<i>Reophax nodulosa</i> Brady, 1879		0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0
<i>Reophax scorpiurus</i> de Montfort, 1808		1	0	0	0	0	0	0	0	0	8	6	61	19	5	0	0

Appendix B (continued)

Appendix B (continued)

sample number	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32
sum counts	426	333	404	303	242	272	403	465	310	487	438	335	315	301	407	342
<i>Rhabdammina abyssorum</i> Sars, 1869	4	0	0	0	0	0	0	0	0	6	0	0	0	0	0	0
<i>Robertinoides bradyi</i> Cushman and Parker, 1936	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Robertinoides purillum</i> Höglund, 1947	0	0	0	0	0	0	0	0	1	1	0	0	2	0	2	0
<i>Robertinoides subcylindrica</i> Brady = <i>Bulimia subcylindrica</i> Brady, 1881	0	1	0	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Rosalina anomala</i> Terquem, 1875	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Rosalina bradyi</i> Cushman = <i>Discorbina globularis</i> d'Orbigny var. <i>bradyi</i> Cushman, 1915	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Rosalina globularis</i> d'Orbigny, 1826	0	0	0	1	0	0	0	0	0	0	0	0	0	3	0	0
<i>Rosalina vilardoboana</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Rupertina stabilis</i> Wallich, 1877	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Saccammina sphaerica</i> Sars, 1868	0	0	0	0	0	0	0	0	0	0	0	0	0	2	0	0
<i>Saccorhiza ramosa</i> Brady = <i>Hyperammina ramosa</i> Brady, 1879	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Sigmoidella elegantissima</i> Parker and Jones = <i>Polymorphina elegantissima</i> Parker and Jones, 1865	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0
<i>Sigmoidilites tenuis</i> Czjzek = <i>Quinqueloculina tenuis</i> Czjzek, 1848	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Sigmolopsis schlumbergeri</i> Silvestri = <i>Sigmololina schlumbergeri</i> Silvestri, 1904	2	2	0	0	0	0	0	0	0	3	0	2	0	0	0	0
<i>Sphaeroidina bullata</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	9	0	4	0	0	0	0
<i>Spirillina vivipara</i> Ehrenberg, 1843	0	0	0	0	0	0	0	0	4	3	0	0	0	0	0	0
<i>Spiriloculina excavata</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Spiriloculina tenellaseptata</i> Brady, 1884	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Spiroplectinella wrighti</i> Silvestri = <i>Spiroplecta wrighti</i> Silvestri, 1903	0	6	1	3	0	0	0	0	0	0	0	0	0	0	3	0
<i>Stainforthia fusiformis</i> Williamson = <i>Bulimina pupoides</i> var. <i>fusiformis</i> Williamson, 1858	11	2	0	0	0	0	0	0	0	0	0	0	1	2	0	3
<i>Stainforthia loeblii</i> Feyling-Hanssen = <i>Virgulina loeblii</i> Feyling-Hanssen, 1954	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0
<i>Stainforthia schreibersiana</i> Carter = <i>Virgulina schreibersiana</i> Carter, 1880	5	1	0	0	0	0	0	0	0	0	0	0	0	1	0	0
<i>Stainforthia skagerakensis</i> Höglund = <i>Virgulina skagerakensis</i> Höglund, 1947	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Stomatobina concentrica</i> Parker and Jones = <i>Pulvinilina concentrica</i> Parker and Jones, 1864	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
<i>Textularia agglutinans</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0
<i>Textularia tenuissima</i> Eartland, 1933	30	11	0	0	0	0	0	0	0	0	0	0	3	0	4	0
<i>Textularia truncata</i> Höglund, 1947	0	1	0	0	0	0	0	0	0	0	0	0	2	0	4	0
<i>Thifrina bradyi</i> Cushman, 1923	0	0	0	0	0	0	0	2	0	0	0	0	0	0	0	0
<i>Triloculina tricarinata</i> d'Orbigny, 1826	9	3	0	0	0	0	0	0	0	0	0	0	2	1	0	2
<i>Trisegmentina compressa</i> Wiesner, 1923	0	0	0	0	0	0	0	0	1	0	0	0	0	1	0	0
<i>Tritaxis fusca</i> Williamson = <i>Rotalina fusca</i> Williamson, 1858	0	0	0	0	0	0	0	0	0	0	0	0	0	1	1	0
<i>Trochammina inflata</i> Montagu = <i>Nutilus inflatus</i> Montagu, 1808	0	0	0	0	0	0	0	0	0	0	0	0	1	1	4	0
<i>Trochammina robertsoni</i> Brady, 1887	23	3	0	0	0	0	0	0	0	0	0	0	1	1	1	3
<i>Uzbekistania charoides</i> Jones and Parker = <i>Trochammina squamata</i> Jones and Parker var. <i>charoides</i> Jones and Parker, 1860	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Uvigerina auberiana</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Uvigerina mediterranea</i> Hofker, 1932	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Uvigerina peregrina</i> Cushman, 1923	19	5	1	1	1	1	1	1	1	1	1	1	1	1	1	3
<i>Vavulinaria bradyana</i> Fornasini = <i>Discorbina bradyana</i> Fornasini, 1900	0	0	0	0	0	0	0	1	0	0	0	0	5	1	0	0
<i>Verniculina propinqua</i> Brady, 1884	0	0	0	0	0	0	0	1	0	0	5	1	0	1	0	0

Appendix C. Species list and quantitative data of planktonic foraminifera from the Norwegian Margin (Chapter 3). Generic names are in accordance with Loeblich and Tappan (1987).

sample number	3	4	5	6	7	8	9	10	11	12	13	14	19	20	22	23	24	25	26	27
sum counts	802	664	383	602	875	809	870	831	950	1164	188	832	24	47	43	117	101	122	189	55
<i>Globigerinella bullata</i> d'Orbigny, 1826	91	78	10	62	128	43	86	67	139	218	28	42	3	1	1	0	1	1	3	1
<i>Globigerinella caudata</i> Parker = <i>Globigerinella siphonifera</i> d'Orbigny, 1826	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globigerinella siphonifera</i> d'Orbigny = <i>Globigerinella siphonifera</i> d'Orbigny, 1839	2	1	0	2	2	0	1	0	4	0	1	5	0	0	0	0	0	0	0	0
<i>Globigerinella glutinata</i> Egger = <i>Globigerina glutinata</i> Egger, 1893	98	35	22	19	43	74	73	221	51	43	32	69	1	24	12	79	67	87	74	25
<i>Globigerinella uvula</i> Ehrenberg = <i>Polydexia uvula</i> Ehrenberg, 1861	192	176	192	136	176	213	180	66	157	181	30	113	0	0	0	10	7	17	13	6
<i>Globigerinoides ruber</i> d'Orbigny = <i>Globigerina rubra</i> d'Orbigny, 1839	1	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0
<i>Globigerinoides sacculifer</i> Brady = <i>Globigerina sacculifera</i> Brady, 1877	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globorotalia inflata</i> d'Orbigny = <i>Globigerina inflata</i> d'Orbigny, 1839	3	0	1	9	7	8	20	23	13	19	0	2	0	0	0	0	0	0	0	0
<i>Globorotalia scitula</i> Brady = <i>Pulvinulina scitula</i> Brady, 1882	1	0	0	0	0	0	0	0	0	0	0	0	0	0	2	0	0	0	0	0
<i>Neogloboquadrina duteiri</i> d'Orbigny = <i>Globigerina duteiri</i> d'Orbigny, 1839	5	1	0	0	2	3	8	5	19	15	0	0	0	0	0	0	0	0	0	0
<i>Neogloboquadrina incompta</i> Cifelli = <i>Globigerina incompta</i> Cifelli, 1961	302	262	46	144	275	220	326	200	249	290	63	473	1	2	23	1	13	6	48	14
<i>Neogloboquadrina pachyderma</i> Ehrenberg = <i>Aristospira pachyderma</i> Ehrenberg, 1861	18	17	4	17	39	25	96	69	82	286	6	6	2	0	0	0	2	0	0	0
<i>Orbulina universa</i> d'Orbigny, 1839	0	0	0	0	0	0	0	1	0	0	1	0	0	0	0	0	0	0	0	0
<i>Tenuitella iota</i> Parker = <i>Globigerinella iota</i> Parker, 1962	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Tenuitella parkerae</i> Brönnimann and Resig, 1971	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Truncorotalia truncatulinoides</i> d'Orbigny = <i>Rotolina truncatulinoides</i> d'Orbigny, 1839	0	0	0	0	0	2	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Turborotalita cristata</i> Heron-Allen and Earland, 1929	0	0	0	1	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0
<i>Turborotalita humilis</i> Brady = <i>Truncatulina humilis</i> Brady, 1884	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	1	0	1	0	1
<i>Turborotalita quinqueloba</i> Naitland = <i>Globigerina quinqueloba</i> Naitland, 1938	89	94	108	213	201	221	80	176	236	112	26	121	11	17	7	27	13	8	51	8

Appendix D. Indicator species for phytodetritus flux, organic carbon flux, oxygen level, and current strength on the Norwegian margin according to Murray (2006) and Margreth et al. (2009) and the references cited therein.

phytodetritus indicator species	organic carbon flux indicator species	oxygen level indicator species	current strength indicator species
<i>Epiostomella</i> spp.	<i>Bathyphion filiformis</i> M. Sars, 1872	<i>Bathyphion filiformis</i> M. Sars, 1872	<i>Angulogerina angulosa</i> Williamson, 1858
<i>Globocassidulina</i> spp.	<i>Bolivina</i> spp.	<i>Cassidulina</i> spp.	<i>Cassidulina</i> spp.
<i>Homosinella guttifera</i> Brady, 1881	<i>Bullimina</i> spp.	<i>Cassidulinoidea oblonga</i> Reuss, 1850	<i>Cassidulinoidea oblonga</i> Reuss, 1850
<i>Nonion depressulus</i> Walker and Jacob, 1798	<i>Cribrostomoides</i> spp.	<i>Cibicides</i> spp.	<i>Cibicides refugens</i> de Montfort, 1808
<i>Nonionella iridea</i> Heron-Allen and Earland, 1932	<i>Cibicoides</i> spp.	<i>Cornuspira</i> spp.	<i>Cornuspira</i> spp.
<i>Spiroloculina</i> spp.	<i>Discanomalina coronata</i> Parker and Jones, 1857	<i>Cribrostomoides</i> spp.	<i>Discanomalina coronata</i> Parker and Jones, 1857
<i>Triloculina tricarinata</i> d'Orbigny, 1826	<i>Elphidiella</i> spp.	<i>Discorbinella berthelotii</i> d'Orbigny, 1839	<i>Discorbinella berthelotii</i> d'Orbigny, 1839
<i>Uvigerina mediterranea</i> Hofer, 1932	<i>Epistominella</i> spp.	<i>Elphidiella albulumillicata</i> Weiss, 1954	<i>Elphidiella albulumillicata</i> Weiss, 1954
<i>Uvigerina peregrina</i> Cushman, 1923	<i>Globobassidulina</i> spp.	<i>Elphidium claratum</i> Cushman, 1930	<i>Elphidium claratum</i> Cushman, 1930
<i>Uvigerina</i> spp.	<i>Globocassidulina</i> spp.	<i>Elphidium frigidum</i> Cushman, 1933	<i>Elphidium frigidum</i> Cushman, 1933
<i>Uvigerina affinis</i> d'Orbigny, 1839	<i>Globularia</i> spp.	<i>Elphidium groenlandicum</i> Cushman, 1933	<i>Elphidium groenlandicum</i> Cushman, 1933
<i>Globocassidulina</i> spp.	<i>Globularia</i> spp.	<i>Elphidium incertum</i> Williamson, 1858	<i>Elphidium incertum</i> Williamson, 1858
<i>Hyalinea balthica</i> Schröter, 1783	<i>Globocassidulina</i> spp.	<i>Elphidium subarcticum</i> Cushman, 1944	<i>Elphidium subarcticum</i> Cushman, 1944
<i>Lagenaria</i> spp.	<i>Hyalinea</i> spp.	<i>Fontbotia wuellestorffii</i> Schwager, 1866	<i>Fontbotia wuellestorffii</i> Schwager, 1866
<i>Lenticulina</i> spp.	<i>Lagenaria</i> spp.	<i>Gaudryina nudis</i> Wright 1900	<i>Gaudryina nudis</i> Wright 1900
<i>Melonis</i> spp.	<i>Lenticulina</i> spp.	<i>Gavelinopsis</i> spp.	<i>Gavelinopsis</i> spp.
<i>Nanion</i> spp.	<i>Melonis</i> spp.	<i>Hanzawaia</i> spp.	<i>Hanzawaia</i> spp.
<i>Nanionella</i> spp.	<i>Quinqueloculina</i> spp.	<i>Hyperammina</i> sp.	<i>Hyperammina</i> sp.
<i>Pulenia</i> spp.	<i>Robertinoides</i> spp.	<i>Lobatula lobatula</i> Walker and Jacob, 1798	<i>Lobatula lobatula</i> Walker and Jacob, 1798
<i>Robertinoides</i> spp.	<i>Triloculina tricarinata</i> d'Orbigny, 1826	<i>Miliolinella</i> spp.	<i>Miliolinella</i> spp.
<i>Stainforthia</i> spp.	<i>Pyrgo</i> spp.	<i>Paratrochammima challengeri</i> Brönnimann and Whittaker, 1988	<i>Paratrochammima challengeri</i> Brönnimann and Whittaker, 1988
<i>Uvigerina</i> spp.	<i>Quinqueloculina</i> spp.	<i>Planulina corrugata</i> Williamson, 1858	<i>Planulina corrugata</i> Williamson, 1858
<i>Valvulinaria bradyana</i> Fornasini, 1900	<i>Robertinoides</i> spp.	<i>Planulina ariminensis</i> d'Orbigny, 1826	<i>Planulina ariminensis</i> d'Orbigny, 1826
	<i>Stainforthia</i> spp.	<i>Psammosphaera fusca</i> Schulze, 1875	<i>Psammosphaera fusca</i> Schulze, 1875
	<i>Uvigerina</i> spp.	<i>Reophax</i> spp.	<i>Reophax</i> spp.
	<i>Valvulinaria</i> spp.	<i>Rosalina</i> spp.	<i>Rosalina</i> spp.
		<i>Rupertina stabilis</i> Wallich, 1877	<i>Rupertina stabilis</i> Wallich, 1877
		<i>Saccammina sphaerica</i> Sars, 1868	<i>Saccammina sphaerica</i> Sars, 1868
		<i>Saccorhiza ramosa</i> Brady, 1879	<i>Saccorhiza ramosa</i> Brady, 1879
		<i>Spiroloculina</i> spp.	<i>Spiroloculina</i> spp.
		<i>Spiroplectinella wrighti</i> Silvestri, 1903	<i>Spiroplectinella wrighti</i> Silvestri, 1903
		<i>Triloculina tricarinata</i> d'Orbigny, 1826	<i>Triloculina tricarinata</i> d'Orbigny, 1826
		<i>Tritaxis fusca</i> Williamson, 1858	<i>Tritaxis fusca</i> Williamson, 1858
		<i>Trochammina</i> spp.	<i>Trochammina</i> spp.

Appendix E. Species list and quantitative data of benthic foraminifera from the Alboran Sea (Chapter 4). Generic names are in accordance with Loeblich and Tappan (1987).

Appendix E (continued)

sample number	M25	M35	M45	M55	M65	M75	M84	M94	M98	M100	M102	M107	M115	M126
sum counts	422	327	328	250	243	273	274	319	240	184	301	235	364	434
<i>Eggerelloides scabra</i> Williamson = <i>Bulinima seabrum</i> Williamson, 1958	1	1	2	2	2	3	2	0	0	2	0	0	0	0
<i>Elphidium fichtelianum</i> d'Orbigny = <i>Polystomella fichtelliana</i> d'Orbigny, 1846	0	1	2	0	1	0	0	0	1	1	4	0	3	2
<i>Epistominella exigua</i> Brady = <i>Polyvinula exigua</i> Brady, 1884	0	0	0	0	0	0	0	0	1	0	0	0	0	1
<i>Epistominella vitrea</i> Parker, 1953	0	1	0	0	0	0	0	0	0	0	0	0	0	3
<i>Favulinia hexagona</i> Williamson = <i>Entosolenia squamosa</i> Montagu var. <i>hexagona</i> Williamson, 1855	0	1	2	2	1	0	0	0	0	0	0	0	0	0
<i>Favulinia melo</i> d'Orbigny = <i>Oolina melo</i> d'Orbigny, 1839	0	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Favulinia squamosa</i> Montagu = <i>Vermiculum squamosum</i> Montagu, 1803	0	0	0	0	0	1	0	0	0	0	0	0	0	0
<i>Fissurina kerqueiensis</i> Parr, 1950	0	1	0	0	0	1	0	0	1	0	0	0	0	0
<i>Fissurina laevigata</i> Reuss, 1850	0	0	3	1	0	0	1	1	0	0	0	0	0	0
<i>Fissurina lucida</i> Williamson = <i>Entosolenia marginata</i> Montagu var. <i>lucida</i> Williamson, 1848	0	0	0	1	0	0	2	0	0	0	0	0	0	0
<i>Fissurina marginata</i> Montagu = <i>Vermiculum marginatum</i> Montagu, 1803	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina orbigniana</i> Seguenza, 1862	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina planiformis</i> Buchner = <i>Lagena planiformis</i> Buchner, 1940	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina quadricostulata</i> Reuss = <i>Lagenaria quadricostulata</i> Reuss, 1870	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fontbotia wuellestorfi</i> Schwager = <i>Anomalina wuellestorfi</i> Schwager, 1866	0	1	0	0	0	0	0	1	0	0	0	0	0	2
<i>Gaudryina rufis</i> Wright 1900	0	0	0	0	0	0	0	0	0	0	0	0	1	0
<i>Gavelinopsis praegeri</i> Heron-Allen and Earland = <i>Discorbina praegeri</i> Heron-Allen and Earland, 1913	10	6	9	5	3	5	1	5	4	1	1	3	2	7
<i>Glabratella torrei</i> Bermúdez = <i>Discorbis torrei</i> Bermúdez, 1935	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Glandulinea ovula</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globobulimina affinis</i> d'Orbigny = <i>Bullimina affinis</i> d'Orbigny, 1839	9	5	8	4	1	5	3	1	0	0	2	3	3	4
<i>Globocassidulina subglobosa</i> group	89	57	48	50	56	46	55	46	26	20	54	33	30	29
<i>Gordiospira fragilis</i> Heron-Allen and Earland, 1932	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Guttulina communis</i> d'Orbigny = <i>Polymorphina (Guttulina) communis</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Gyroidina affiformis</i> Steward and Steward = <i>Gyroidina soldanii</i> d'Orbigny var. <i>affiformis</i> Steward and Steward, 1930	2	0	4	1	6	7	7	4	1	1	0	1	3	1
<i>Gyroidina neosoldanii</i> Brozen, 1936	2	4	1	6	1	0	3	2	1	0	0	1	4	3
<i>Gyroidina soldanii</i> d'Orbigny, 1826	6	5	1	0	3	2	1	0	0	0	0	0	5	3
<i>Gyroidinoides laevigatus</i> d'Orbigny = <i>Gyroidina laevigata</i> d'Orbigny, 1826	0	1	0	0	0	2	1	0	0	0	0	0	2	2
<i>Hanzawaia boueana</i> d'Orbigny = <i>Rotalina boueana</i> d'Orbigny, 1846	1	5	2	0	1	0	2	1	1	2	0	0	4	0
<i>Hanzawaia concentrica</i> Cushman = <i>Truncatulina concentrica</i> Cushman, 1918	1	2	1	1	0	1	2	1	0	1	1	1	0	1
<i>Hoeglundina elegans</i> d'Orbigny = <i>Rotalia (Turbulinula) elegans</i> d'Orbigny, 1826	4	3	6	0	1	5	4	3	5	3	7	2	3	5
<i>Homalohedra williamseni</i> Alcock = <i>Entosolenia williamseni</i> Alcock, 1865	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Hyalinaea balthica</i> Schröter = <i>Nautilus balthicus</i> Schröter, 1783	15	18	10	9	9	18	10	19	11	7	26	19	32	44
<i>Hyrokkin sarcophaga</i> Cedhagen, 1994	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Kareniella bradyi</i> Cushman = <i>Gaudryina bradyi</i> Cushman, 1911	0	0	0	0	0	0	0	0	1	0	0	0	0	0
<i>Lagena gibbera</i> Buchner, 1940	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Lagena meridionalis</i> Wiesner = <i>Lagena gracilis</i> Williamson var. <i>meridionalis</i> Wiesner, 1931	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Lagena striata</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	1	0	0	0	0	0
<i>Lagena williamseni</i> Harvey and Bailey, 1854	0	0	0	0	0	0	0	0	1	0	0	0	1	0
<i>Lenticulina calcare Linné</i> = <i>Nauillus calcare</i> Linné, 1758	2	0	1	1	0	1	0	1	0	0	0	0	0	1
<i>Lenticulina gibba</i> d'Orbigny = <i>Cristellaria gibba</i> d'Orbigny, 1826	1	1	0	1	1	0	1	0	0	0	0	0	0	1
<i>Lenticulina ornata</i> d'Orbigny = <i>Robulina ornata</i> d'Orbigny, 1846	0	1	1	1	2	0	0	1	0	0	2	1	0	0
<i>Lenticulina orbicularis</i> d'Orbigny = <i>Robulina orbicularis</i> d'Orbigny, 1826	0	0	0	0	0	1	0	3	1	1	2	1	0	3
<i>Lenticulina peregrina</i> Schwager = <i>Cristellaria peregrina</i> Schwager, 1866	1	1	0	1	0	0	2	1	0	0	0	2	1	0

Appendix E (continued)

sample number	D0	D10	D15	D20	D25	D30	D35	D40	D45	D50	D55	D60	D150
sum counts	367	347	372	281	309	412	433	330	293	412	245	130	36
<i>Eggerelloides scabra</i> Williamson = <i>Bullimina seabrum</i> Williamson, 1958	0	2	4	4	0	0	2	0	0	1	1	0	0
<i>Elphidium fichtelianum</i> d'Orbigny = <i>Polystomella fichtelliana</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Epistominella exigua</i> Brady = <i>Polyvinulina exigua</i> Brady, 1884	0	3	6	2	1	0	0	0	0	0	0	0	1
<i>Epistominella vitrea</i> Parker, 1953	0	2	0	0	1	0	0	0	0	1	1	1	3
<i>Favulinia hexagona</i> Williamson = <i>Entosolenia squamosa</i> Montagu var. <i>hexagona</i> Williamson, 1855	1	0	1	1	0	0	1	0	0	0	1	0	1
<i>Favulinia melo</i> d'Orbigny = <i>Oolina melo</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	0	0	1
<i>Favulinia squamosa</i> Montagu = <i>Vermiculum squamosum</i> Montagu, 1803	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina kerqueensis</i> Parr, 1950	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Fissurina laevigata</i> Reuss, 1850	0	0	0	1	1	0	0	0	0	0	0	0	0
<i>Fissurina lucida</i> Williamson = <i>Entosolenia marginata</i> Montagu var. <i>lucida</i> Williamson, 1848	0	0	0	0	0	0	0	1	0	0	0	2	0
<i>Fissurina marginata</i> Montagu = <i>Vermiculum marginatum</i> Montagu, 1803	1	2	0	0	0	0	0	0	1	1	1	2	0
<i>Fissurina orbigniana</i> Sequenza, 1862	0	1	2	1	0	1	2	1	2	1	2	1	0
<i>Fissurina planiformis</i> Buchner = <i>Lagena planiformis</i> Buchner, 1940	0	1	0	0	0	0	0	0	0	0	1	0	0
<i>Fissurina quadricostulata</i> Reuss = <i>Lagenaria quadricostulata</i> Reuss, 1870	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Fontbotia wuerstorffii</i> Schwager = <i>Anomalina wuerstorffii</i> Schwager, 1866	0	0	0	0	0	0	0	0	0	1	0	0	0
<i>Gaudyina rufa</i> Wright 1900	0	0	0	0	0	1	0	0	0	4	1	1	1
<i>Gavelinopsis praegeri</i> Heron-Allen and Earland = <i>Discorbina praegeri</i> Heron-Allen and Earland, 1913	19	16	14	14	17	20	14	3	9	6	4	2	
<i>Glabratella torrei</i> Bermúdez = <i>Discorbis torrei</i> Bermúdez, 1935	0	0	0	0	1	0	0	0	0	0	0	0	0
<i>Glandulina ovula</i> d'Orbigny, 1846	1	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globobulimina affinis</i> d'Orbigny = <i>Bullimina affinis</i> d'Orbigny, 1839	2	1	1	0	2	1	1	2	0	1	0	0	0
<i>Globocassidulina subglobosa</i> group	85	86	54	48	61	88	75	57	50	30	27	14	30
<i>Gordiospira fragilis</i> Heron-Allen and Earland, 1932	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Guttulina communis</i> d'Orbigny = <i>Polymorphina (Guttulina) communis</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Gyrodina affiformis</i> Steward and Steward = <i>Gyrodina soldanii</i> d'Orbigny var. <i>affiformis</i> Steward and Steward, 1930	0	1	5	4	3	7	4	4	0	0	0	0	1
<i>Gyrodina neosoldanii</i> Brozen, 1936	1	0	2	3	5	3	1	1	2	5	2	0	1
<i>Gyrodina soldanii</i> d'Orbigny, 1826	3	4	0	1	1	1	0	1	0	7	2	0	1
<i>Gyroidinoides laevigatus</i> d'Orbigny = <i>Gyroidina laevigata</i> d'Orbigny, 1826	1	2	0	0	0	0	1	0	3	0	0	0	0
<i>Hanzawaia boueana</i> d'Orbigny = <i>Rotalina boueana</i> d'Orbigny, 1846	5	7	4	3	3	7	4	3	3	6	2	1	
<i>Hanzawaia concentrica</i> Cushman = <i>Truncatulina concentrica</i> Cushman, 1918	0	0	0	0	0	0	0	0	0	3	1	2	0
<i>Hoeeglundina elegans</i> d'Orbigny = <i>Rotalia (Turbulinula) elegans</i> d'Orbigny, 1826	10	11	7	6	0	11	9	4	8	1	7	2	
<i>Homalohedra williamseni</i> Alcock = <i>Entosolenia williamseni</i> Alcock, 1865	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Hyalinea balthica</i> Schröter = <i>Nautilus balthicus</i> Schröter, 1783	6	4	6	5	7	3	5	4	2	13	8	3	4
<i>Hyrtokkin sarcophaga</i> Cedhagen, 1994	0	1	0	0	0	1	0	0	0	0	0	0	0
<i>Kareniella bradyi</i> Cushman = <i>Gaudryina bradyi</i> Cushman, 1911	0	0	0	0	0	0	0	0	1	0	0	0	0
<i>Lagena gibbera</i> Buchner, 1940	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Lagena meridionalis</i> Wiesner = <i>Lagena gracilis</i> Williamson var. <i>meridionalis</i> Wiesner, 1931	1	0	0	0	0	0	0	0	1	0	0	0	0
<i>Lagena striata</i> d'Orbigny, 1854	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Lenticulina calcare</i> Linne = <i>Nauillus calcare</i> Linne, 1758	0	0	0	0	0	0	0	0	0	1	0	0	0
<i>Lenticulina gibba</i> d'Orbigny = <i>Cristellaria gibba</i> d'Orbigny, 1826	0	2	0	0	0	0	0	0	0	0	0	0	0
<i>Lenticulina inornata</i> d'Orbigny = <i>Robulina inornata</i> d'Orbigny, 1846	4	6	2	2	4	0	0	0	0	0	0	0	0
<i>Lenticulina orbicularis</i> d'Orbigny = <i>Robulina orbicularis</i> d'Orbigny, 1826	1	1	4	4	1	8	2	3	4	0	2	0	0
<i>Lenticulina peregrina</i> Schwager = <i>Cristellaria peregrina</i> Schwager, 1856	0	0	0	0	0	2	0	0	0	0	0	0	0

Appendix E (continued)

sample number	D0	D10	D15	D20	D25	D30	D35	D40	D45	D50	D55	D60	D65	D70	D75	D80	D85	D90	D95	D100
sum counts	367	347	372	281	309	412	433	330	293	412	245	130	136							
<i>Lenticulina vortex</i> Fichtel and Moll = <i>Nautilus vortex</i> Fichtel and Moll, 1798	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Lobatula lobatula</i> Walker and Jacob = <i>Nautilus lobatulus</i> Walker and Jacob, 1798	4	5	9	6	1	4	11	6	11	9	9	1	1	0	0	0	0	0	0	
<i>Marginulina striata</i> d'Orbigny, 1852	0	0	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Melonis barleeanum</i> Williamson = <i>Nonionina barleeanana</i> Williamson, 1858	6	2	14	9	7	13	11	10	2	13	5	4	3							
<i>Melonis pomphiloides</i> Fichtel and Moll, 1798 = <i>Nautilus pomphiloides</i> Fichtel and Moll, 1798	0	0	0	1	1	0	1	0	3	3	0	0	2							
<i>Miliolinella elongata</i> Kruit = <i>Miliolinella circulans</i> Bonnemann var. <i>elongata</i> Kruit, 1955	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Miliolinella subrotunda</i> Montagu, 1803 = <i>Vermiculum subrotundum</i> Montagu, 1803	1	0	3	4	4	9	1	1	2	4	2	3	0							
<i>Neocoorbina terquemii</i> Rzezhak = <i>Discorbina terquemii</i> Rzezhak, 1888	0	0	0	0	0	0	0	1	0	0	0	1	0							
<i>Nonion pauperatus</i> Balkwill and Wright = <i>Nonionina pauperata</i> Balkwill and Wright, 1885	0	0	0	0	0	0	0	0	0	0	0	0	0							
<i>Nonion</i> sp.	0	0	0	0	0	0	0	0	0	0	0	0	0							
<i>Nonionella iridea</i> Héron-Alen and Earland, 1932	0	0	0	2	1	1	0	1	0	1	0	1	0	2	0					
<i>Nonionella turgida</i> Williamson = <i>Rotalina turgida</i> Williamson, 1858	1	0	2	1	0	1	0	0	0	0	0	0	0	0	0					
<i>Nuttallides umboniferus</i> Cushman = <i>Pulvinulina umbonifera</i> Cushman, 1933	1	1	2	7	3	3	7	0	3	1	0	0	0	0	1					
<i>Oridorsalis umbonatus</i> Reuss = <i>Rotalina umbonata</i> Reuss, 1851	0	0	0	0	0	0	0	0	0	2	0	0	0	0	0					
<i>Paratrochammina challenger</i> Brönnimann and Whittaker, 1988	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Paratrochammina murrayi</i> Brönnimann and Zaninetti, 1984	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Planulina auriminensis</i> d'Orbigny, 1826	2	0	1	2	0	2	3	0	0	1	0	0	0	0	0					
<i>Porosononion ex gr. granosum</i> d'Orbigny = <i>Nonionina granosa</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Pseudospionides falsobuccarii</i> Rouvilleis = <i>Ammonia falsobuccarii</i> Rouvilleis, 1974	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0					
<i>Pseudodonodaria aequalis</i> d'Orbigny = <i>Nodosaaria aequalis</i> d'Orbigny, 1908	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Pseudotrioculina oblonga</i> Montagu = <i>Vermiculum oblongum</i> Montagu, 1803	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Pullenia bulloides</i> d'Orbigny = <i>Nonionina bulloides</i> d'Orbigny, 1846	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Pullenia quadriloba</i> Reuss = <i>Pullenia compressiscutula</i> Reuss var. <i>quadriloba</i> Reuss, 1867	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Pullenia quinquelobata</i> Reuss = <i>Nonionina quinqueloba</i> Reuss, 1851	0	0	0	1	0	0	0	1	0	0	0	0	0	0	0					
<i>Pyrgo anomala</i> Schlumberger = <i>Biloculina anomala</i> Schlumberger, 1891	0	0	1	0	0	2	1	0	1	0	1	0	2	4	0					
<i>Pyrgo depressa</i> d'Orbigny = <i>Biloculina depressa</i> d'Orbigny, 1826	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0					
<i>Pyrgo elongata</i> d'Orbigny = <i>Biloculina elongata</i> d'Orbigny, 1826	0	0	0	0	1	0	0	1	0	0	0	0	0	0	0					
<i>Pyrgo inornata</i> d'Orbigny = <i>Biloculina inornata</i> d'Orbigny, 1839	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0					
<i>Pyrgoella sphaera</i> d'Orbigny = <i>Biloculina sphaera</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Quinqueloculina padana</i> Percovich, 1954	0	0	1	2	0	0	3	0	0	1	2	1	0	0	0					
<i>Quinqueloculina seminula</i> Linne = <i>Serpula seminulum</i> Linne, 1758	14	6	7	6	5	4	13	4	8	9	0	2								
<i>Quinqueloculina viennensis</i> Le Calvez and Le Calvez, 1958	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Quinqueloculina pseudobuchiana</i> Luczkowska, 1974	1	4	2	2	3	2	2	1	0	0	0	0	0	0	0					
<i>Rectuvigerina elongatastrata</i> Colom = <i>Angulogerina elongatastrata</i> Colom, 1952	0	0	0	0	0	0	0	1	0	0	0	0	0	0	0					
<i>Robertinoides subcylindrica</i> Brady = <i>Bulimia subcylindrica</i> Brady, 1831	1	3	2	2	5	4	9	3	0	4	1	0	0	0	0					
<i>Rosalina globularis</i> d'Orbigny, 1826	0	0	0	0	0	0	0	0	0	0	1	0	0	0	0					
<i>Rosalina vilardoiana</i> d'Orbigny, 1839	0	1	2	0	1	0	1	1	2	2	0	0	0	0	0					
<i>Sigmoliniella sigmoides</i> Brady = <i>Planispirina sigmoides</i> Brady, 1884	0	0	0	0	0	1	1	0	1	0	1	0	0	0	0					
<i>Sigmoliniella tenuis</i> Czjzek = <i>Quinqueloculina tenuis</i> Czjzek, 1848	0	0	0	1	1	0	1	0	1	0	1	0	0	0	0					
<i>Sigmolopsis schlumbergeri</i> Silvestri = <i>Sigmolina schlumbergeri</i> Silvestri, 1904	4	5	2	5	0	0	7	5	6	1	2	0	0	0	0					
<i>Siphogenerina columellaris</i> Brady = <i>Uvigerina columellaris</i> Brady, 1881	0	0	2	0	0	0	0	0	0	0	0	0	0	0	0					
<i>Siphonina reticulata</i> Czjzek = <i>Rotalia reticulata</i> Czjzek, 1848	0	0	2	1	0	1	0	0	0	0	0	0	0	0	0					

Appendix F. Species list and quantitative data of planktonic foraminifera from the Alboran Sea (Chapter 4). Generic names are in accordance with Loeblich and Tappan (1987).

sample number	M25	M35	M45	M55	M65	M75	M84	M94	M98	M100	M102	M107	M115	M126
sum counts	994	1216	1226	1115	1122	1068	1066	1103	1027	816	1058	854	918	946
<i>Bella digitata</i> Brady = <i>Globigerina digitata</i> Brady, 1879	0	0	0	0	0	1	1	1	1	2	1	0	0	0
<i>Globigerina bulloides</i> d'Orbigny, 1826	97	88	86	71	78	51	34	38	37	24	51	31	46	48
<i>Globigerinella calida</i> Parker = <i>Globigerina calida</i> Parker, 1926	4	3	3	4	4	3	3	2	2	6	1	2	0	2
<i>Globigerinella siphonifera</i> d'Orbigny = <i>Globigerina siphonifera</i> d'Orbigny, 1839	0	7	15	13	8	13	12	8	8	3	6	0	4	1
<i>Globigerinita glutinata</i> Egger = <i>Globigerina glutinata</i> Egger, 1893	205	333	231	200	185	201	198	186	123	113	168	117	141	110
<i>Globigerinita uvula</i> Ehrenberg = <i>Pyloidea uvula</i> Ehrenberg, 1861	11	26	7	17	10	12	13	20	16	10	11	12	21	10
<i>Globigerinoides elongatus</i> Brady, 1879	0	0	0	0	0	0	2	0	0	1	0	0	0	0
<i>Globigerinoides elongatus</i> d'Orbigny = <i>Globigerina elongata</i> d'Orbigny, 1826	15	14	25	18	8	25	20	13	13	6	10	12	6	11
<i>Globigerinoides immaturus</i> Le Roy = <i>Globigerinoides sacculiferus</i> Brady var. <i>Immatura</i> Le Roy, 1939	4	5	9	5	5	10	3	4	6	0	0	0	0	0
<i>Globigerinoides ruber</i> d'Orbigny = <i>Globigerina rubra</i> d'Orbigny, 1839	47	74	89	93	115	106	122	89	75	56	67	58	31	24
<i>Globigerinoides sacculifer</i> Brady = <i>Globigerina sacculifera</i> Brady, 1877	9	10	6	5	4	7	2	0	1	1	8	0	0	0
<i>Globorotalia hirsuta</i> d'Orbigny = <i>Rotolina hirsuta</i> d'Orbigny, 1839	2	3	0	0	0	0	0	0	0	0	1	1	0	2
<i>Globorotalia inflata</i> d'Orbigny = <i>Globigerina inflata</i> d'Orbigny, 1839	270	328	276	266	236	209	144	80	41	34	42	31	47	53
<i>Globorotalia scitula</i> Brady = <i>Pulvinulina scitula</i> Brady, 1882	9	4	4	1	0	3	2	1	0	0	1	0	1	12
<i>Globotuborotalita rubescens</i> Hofker = <i>Globigerina rubescens</i> Hofker, 1956	19	34	42	35	51	47	38	22	19	4	14	11	4	7
<i>Hastigerina pelagica</i> d'Orbigny = <i>Nonionina pelagica</i> d'Orbigny, 1839	0	0	0	0	0	0	0	0	0	0	0	0	0	2
<i>Hastigerinopsis riedeli</i> Rögl and Bolli = <i>Hastigerinella riedeli</i> Rögl and Bolli, 1973	0	0	2	1	0	0	0	0	0	0	0	0	2	0
<i>Neglobiquadrina dutertrei</i> d'Orbigny = <i>Globigerina dutertrei</i> d'Orbigny, 1839	0	0	1	0	0	0	0	0	0	2	16	0	4	9
<i>Neglobiquadrina incompta</i> Cifelli = <i>Globigerina incompta</i> Cifelli, 1961	25	41	69	69	99	105	168	239	307	230	320	245	287	372
<i>Orbulina universa</i> d'Orbigny, 1839	7	7	7	3	8	5	4	0	1	1	4	4	3	0
<i>Pulnietina obliquiloculata</i> Parker and Jones = <i>Pulnella sphaeroidea</i> d'Orbigny var. <i>Obliquiloculata</i> Parker and Jones, 1865	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Tenuitella anfracta</i> Parker = <i>Globorotalia anfracta</i> Parker, 1967	0	0	1	2	1	4	0	0	0	2	0	1	1	0
<i>Tenuitella iota</i> Parker = <i>Globigerinita iota</i> Parker, 1962	11	6	5	3	5	0	2	0	1	1	0	0	0	0
<i>Tenuitella parkerae</i> Brönnimann and Resig, 1971	0	0	0	0	0	0	0	0	2	0	1	1	3	0
<i>Truncorotalia crassiformis</i> Galloway and Wissler = <i>Globigerina crassiformis</i> Galloway and Wissler, 1927	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Truncorotalia truncatulinoides</i> d'Orbigny = <i>Rotolina truncatulinoides</i> d'Orbigny, 1839	10	9	7	12	7	2	3	1	3	3	2	3	2	9
<i>Turborotalita cristata</i> Heron-Allen and Earland = <i>Globigerina cristata</i> Heron-Allen and Earland, 1929	1	0	3	1	2	0	0	0	1	0	2	0	0	1
<i>Turborotalita humilis</i> Brady = <i>Truncatulina humilis</i> Brady, 1884	20	21	22	7	13	12	5	1	4	13	4	4	1	6
<i>Turborotalita quinqueloba</i> Nai land = <i>Globigerina quinqueloba</i> Nai land, 1938	224	200	315	289	283	251	287	397	359	291	344	315	304	271

Appendix F (continued)

sample number sum counts	D0	D10	D15	D20	D25	D30	D35	D40	D45	D50	D55	D60	D65	D70	D75	D70	D150
<i>Bellia digitata</i> Brady = <i>Globigerina digitata</i> Brady, 1879	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globigerina bullioloides</i> d'Orbigny, 1826	165	169	125	73	60	110	50	75	49	87	54	38	34	29	47	36	
<i>Globigerinella calida</i> Parker = <i>Globigerina calida</i> Parker, 1926	8	6	7	2	4	2	7	4	2	4	4	2	5	0	1	3	
<i>Globigerinella siphonifera</i> d'Orbigny = <i>Globigerina siphonifera</i> d'Orbigny, 1839	4	4	8	12	7	13	5	5	4	1	1	1	3	1	4	5	
<i>Globigerinella glutinata</i> Egger = <i>Globigerina glutinata</i> Egger, 1893	89	59	171	132	128	146	126	154	93	120	106	140	176	221	162	182	
<i>Globigerinella uvula</i> Ehrenberg = <i>Pyloidoxia uvula</i> Ehrenberg, 1861	16	21	38	22	20	39	28	14	0	19	20	10	18	20	21	19	
<i>Globigerinoides conglobatus</i> Brady, 1879	0	3	1	1	0	0	0	0	0	0	0	0	0	0	0	0	0
<i>Globigerinoides elongatus</i> d'Orbigny = <i>Globigerina elongata</i> d'Orbigny, 1826	16	15	26	31	27	14	12	10	11	1	5	3	18	5	13	0	
<i>Globigerinoides immaturus</i> Le Roy = <i>Globigerinoides sacculiferus</i> Brady var. <i>Immatura</i> Le Roy, 1939	7	11	16	14	3	7	3	3	10	1	4	0	6	3	7	10	
<i>Globigerinoides ruber</i> d'Orbigny = <i>Globigerina rubra</i> d'Orbigny, 1839	72	72	82	78	80	70	69	71	63	27	23	8	84	61	61	40	
<i>Globigerinoides sacculifer</i> Brady = <i>Globigerina sacculifera</i> Brady, 1877	11	7	5	9	5	12	0	1	3	1	0	1	1	0	2	3	
<i>Globorotalia hispida</i> d'Orbigny = <i>Rotalina hispida</i> d'Orbigny, 1839	6	3	1	4	1	2	2	0	1	0	1	0	1	0	0	0	5
<i>Globorotalia inflata</i> d'Orbigny = <i>Globigerina inflata</i> d'Orbigny, 1839	327	293	272	370	304	231	98	172	191	24	33	8	129	42	105	91	
<i>Globorotalia scitula</i> Brady = <i>Pulvinulina scitula</i> Brady, 1882	2	26	8	4	1	9	8	7	1	8	7	3	1	1	2	4	
<i>Globoturborotalita rubescens</i> Hofker = <i>Globigerina rubescens</i> Hofker, 1956	7	3	8	3	1	4	0	13	10	6	6	6	7	9	19	7	
<i>Hastigerina pelagica</i> d'Orbigny = <i>Nonionina pelagica</i> d'Orbigny, 1839	0	0	1	0	0	0	0	1	0	0	0	0	0	0	0	0	
<i>Hastigerinopsis ridelei</i> Rögl and Bolli = <i>Hastigerinella ridelei</i> Rögl and Bolli, 1973	0	0	0	0	0	0	0	0	0	0	0	0	1	0	0	1	
<i>Negloboquadrina duferrei</i> d'Orbigny = <i>Globigerina duferrei</i> d'Orbigny, 1839	3	0	5	5	2	1	2	3	3	0	3	0	0	4	3	2	
<i>Negloboquadrina incompta</i> Cifelli = <i>Globigerina incompta</i> Cifelli, 1961	78	102	100	74	121	105	210	126	245	426	303	136	113	191	100	117	
<i>Negloboquadrina pachyderma</i> Ehrenberg = <i>Aristospira pachyderma</i> Ehrenberg, 1861	3	7	12	16	7	9	8	13	7	6	5	13	13	10	9	16	
<i>Orbulina universa</i> d'Orbigny, 1839	7	11	9	13	5	9	2	7	7	4	0	2	2	3	2	7	
<i>Pulleniatina obliquiloculata</i> Parker and Jones = <i>Pullenia sphaeroidea</i> d'Orbigny var. <i>Obliquiloculata</i> Parker and Jones, 1865	0	0	0	0	0	0	0	0	1	0	0	0	0	1	0	0	
<i>Tenuitella anfracta</i> Parker = <i>Globorotalia anfracta</i> Parker, 1967	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Tenuitella parkerae</i> Brönnimann and Resig, 1971	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Truncorotalia crassaformis</i> Galloway and Wissler = <i>Globigerina crassaformis</i> Galloway and Wissler, 1927	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	0	
<i>Truncorotalia truncatulinoides</i> d'Orbigny = <i>Rotalina truncatulinoides</i> d'Orbigny, 1839	11	6	4	13	7	7	1	10	5	1	0	6	2	2	3		
<i>Turborotalita cristata</i> Heron-Allen and Earland = <i>Globigerina cristata</i> Heron-Allen and Earland, 1929	1	1	2	6	2	3	7	1	0	1	2	1	0	1	0	0	
<i>Turborotalita humilis</i> Brady = <i>Truncatulina humilis</i> Brady, 1884	12	2	28	12	15	18	12	11	4	9	5	4	10	13	7	13	
<i>Turborotalita quinqueloba</i> Nautland = <i>Globigerina quinqueloba</i> Nautland, 1938	204	221	149	198	271	251	292	218	70	214	223	223	195	217	160	160	

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Oligocene Planktonic Foraminifera Working Group Meeting: 23 – 25 June 2009, Fribourg, Switzerland
Cold-Water Carbonate Reservoir System in Deep Environments (COCARDE) – A pilot Industry - Academia Partnership in Marine Research Drilling – Magellan Workshop Series, 21-24 January, Fribourg, Switzerland.
Cold-water Coral Ecosystems in a Changing World; 17 – 18 January 2008, Gent, Belgium.
Oligocene Planktonic Foraminifera Working Group Meeting; 13 – 16 December 2007, Tübingen, Germany.
Exploring Escarpment Mud Mound Systems and Mud Volcanoes with New European Strategies for Sustainable Mid-Depth Coring - Magellan Workshop Series, 26 – 29 April 2007, Murten, Switzerland.
Workshop on Mesozoic Planktonic Foraminiferal Taxonomy; 10 September 2006, Natal, Brasil.

Publications:

published:

Margreth, S., Rüggeberg, A., Spezzaferri, S., 2009. Benthic foraminifera as bioindicator for cold-water coral reef ecosystems along the Irish margin. *Deep-Sea Research I*, 56, p. 2216 – 2234.
Machguth, H. and Margreth, S. 2005. Baikal Exkursion 2004: Ein wissenschaftlicher Ueberblick zum Baikal-Raum und Reisebericht, in Schriftenreihe physische Geographie, Bodenkunde und Biogeographie 46, Universität Zürich, 2005, pp. 143, ISBN 3 85543 242 2

in review:

Margreth, S., Rüggeberg, A., Spezzaferri, S., submitted. Benthic Foraminifera associated to Norwegian Cold-Water Coral Reefs. *J. Foraminifer. Res.*
Margreth, S., Gennari, G., Rüggeberg, A., Comas, M.C., Pinheiro, L.M., Spezzaferri, S., submitted. Development of cold-water coral ecosystems on mud volcanoes in the West Alboran Basin: paleoceanographic implications. *Mar. Geol.*
Raddatz, J., Rüggeberg, A., Margreth, S., Dullo, W., and IODP Expedition 307 Scientific Party, submitted. Paleoenvironmental reconstruction of deep-water carbonate mound initiation in the Porcupine Seabight, NE Atlantic. *Mar. Geol.*

Reviewer for:

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Conference contributions:

Margreth, S., Spezzaferri, S., Gennari, G., Rüggeberg, A., Stalder, C., Pignat, G., 2010. Biogeography of foraminifera from cold-water coral ecosystems: from active (Northern European Margin) to buried reefs (Western Mediterranean Sea). *Forams 2010 - International Symposium on Foraminifera*, Bonn, Germany, 137-138.
Spezzaferri, S., Margreth, S., Rüggeberg, A., Gennari, G., 2010. Benthic foraminifera associated to cold-water coral ecosystems: towards the assessment of regional bioindicators. The colors of Cretaceous and Paleogene Oceans, Verbania, Italy.
Margreth, S., Rüggeberg, A., Gennari, G., Spezzaferri, G., 2010. Identifying cold-water coral ecosystem by using benthic foraminiferal indicators: from active reefs to the geological record. *European Gesciences Union, General Assembly 2010*. European Gesciences Union, Wien, Austria. *Geophysical Research Abstracts*, EGU2010-4167-1, 2010.
Margreth, S., Rüggeberg, A., Gennari, G., Spezzaferri, G., 2010. Foraminiferal indicator species for cold-water coral ecosystem dynamics along the European continental margin. *SwissSed Meeting*, Fribourg, Switzerland, 37.
Raddatz, J., Rüggeberg, A., Liebetrau, V., **Margreth, S.**, Eisenhauer, A., Dullo, W.-C., and IODP Expedition 307 Scientific Party, 2009. Paleoenvironmental reconstruction of deep-water carbonate mound initiation in the Porcupine Seabight, NE Atlantic. *Eos Trans. AGU*, 90(52), Fall Meet. Suppl., San Francisco, USA. Abstract PP13E-07.
Margreth, S., Gennari, G., Spezzaferri, S., Rüggeberg, A., 2009. Cold-water coral ecosystem dynamics and foraminiferal distribution and dispersal along the European Atlantic and Mediterranean margins. *13th Congress of Regional Committee on Mediterranean Neogene Stratigraphy RCMNS*, Naples, Italy. Earth System evolution

- and the Mediterranean area from 23 Ma to the present. *Acta Naturalia de l'Ateneo Parmense*, 45 (1/4), 219.
- Gennari, G., **Margreth, S.**, Spezzaferri, S., Comas, M.C., Pinheiro, L.M., Rüggeberg, A., 2009. Mud volcanoes and cold-water corals in the West Alboran Basin: new insights from the SAGAS 2008 (TTR-17 Leg 1) cruise. SwissSed Meeting, Fribourg, Switzerland, 13-14.
- Raddatz, J., Rüggeberg, A., **Margreth, S.**, Dullo, W. C. and IODP Expedition 307 Scientific Party, 2009. Paleoenvironment of cold-water coral initiation in the NE Atlantic: Implications from a deepwater carbonate mound drilling core. European Geosciences Union, General Assembly 2009. European Geosciences Union, Wien, Austria, Geophysical Research Abstracts, Vol. 11, EGU2009-2871-1, 2009
- Margreth, S.**, Gennari, G., Spezzaferri, S., Comas, M.C., Pinheiro, L.M., Rüggeberg, A., 2009. Cold-water corals and mud volcanoes in the West Alboran Basin, International Conference and TTR-17th Post-Cruise Meeting of the Training-through-Research Programme Granada, Spain, 43-44.
- Margreth, S.**, Spezzaferri, S., Tamburini, F., Rüggeberg, A., de Haas, H., 2008. Cold water coral mounds from the Porcupine area: On-off mound characterization based on planktonic and benthic foraminifera and phosphorus content in the sediments, AAPG International Conference and Exhibition, Cape Town, South Africa.
- Margreth, S.**, Spezzaferri, S., Tamburini, F., de Haas, H., Party, P.C.S., 2008. Benthic foraminifera as bioindicators for active cold-water coral mounds: Results from the Porcupine and Rockall Banks in the North Atlantic. The micropalaeontological society's foraminifera and nannofossil group's joint spring meeting, Tübingen, Germany, 35.
- Margreth, S.**, Spezzaferri, S., Tamburini, F., de Haas, H., 2008. Cold water coral mounds and foraminifera from the Porcupine and Rockall Banks in the North Atlantic: Results from the Pelagia Cruise. SwissSed Meeting, Fribourg, Switzerland, 30.
- Margreth, S.**, Rüggeberg, A., Tamburini, F., Spezzaferri, S., 2008. Benthic and planktonic Foraminifera and other proxies for determine environmental parameters in cold-water coral ecosystems along the European margin. Deepsea Coral Symposium 2008, Wellington, New Zealand, 13.09.
- Wheeler, A., Freiwald, A., Hebbeln, D., Swennen, R., van Weering, T.C.E., de Haas, H., Dorschel, B., Fink, H., Joseph, N., **Margreth, S.H.**, Pirlet, H., Monteys, X., Thierens, M., 2008. Pre-site drilling survey and carbonate mound spatial analysis results: CARBONATE year one results. Hermes 3rd Annual Meeting, Algarve, Portugal, 76.
- Joseph, N., Wheeler, A., Freiwald, A., Hebbeln, D., Swennen, R., van Weering, T.C.E., de Haas, H., Dorschel, B., Fink, H., **Margreth, S.**, Pirlet, H., Monteys, X., Thierens, M., 2008. The CARBONATE Project - Mid Latitude Carbonate Systems: Complete Sequences from Cold-Water Coral Carbonate Mounds in the N.E. Atlantic. Jahrestagung der Paläontologischen Gesellschaft, Erlangen, Germany, 94.
- De Haas, H., Dorschel, B., Fink, H., Joseph, N., **Margreth, S.**, Pirlet, H., Thierens, M., Van Weering, T.C.E., Wheeler, A. (2008). The R.V. Pelagia pre-drilling site survey at the Rockall and Porcupine cold water coral mounds provinces, European Atlantic margin. The CARBONATE project, ESF EuroMARC program. European Geosciences Union, General Assembly 2008. European Geosciences Union, Wien, Austria. Geophysical Research Abstracts, Vol. 10, EGU2008-A-06379, 2008.
- De Haas, H., Dorschel, B., Fink, H., Joseph, N., **Margreth, S.**, Pirlet, H., Thierens, M., van Weering, T.C.E., Wheeler, A., 2008. The R.V. Pelagia pre-drilling site survey at the Rockall and Porcupine cold water coral mounds provinces, European Atlantic margin. The CARBONATE project, ESF EuroMARC program. EuroMARC joint CARBONATE/CHECREEF Workshop, Sant'Alessio, Italy, 2.
- Margreth, S.**, Tamburini, F., Grobety, B., Coric, S., Spezzaferri, S., Bernasconi, S., 2007. The transition from Marine Isotope Stage 6 to 5 at ODP Hole 1198A (leg 194-Marion Plateau, Australia): micropaleontology and geochemistry. European Geosciences Union, General Assembly 2007. European Geosciences Union, Wien, Austria. Geophysical Research Abstracts, Vol. 9, EGU2007-A-01522, 2007.
- Margreth, S.**, Tamburini, F., Grobety, B., Coric, S., Spezzaferri, S., Bernasconi, S., 2007. Carbon mineralogy, phosphorus and oxygen isotope records from the Late Pleistocene at ODP Hole 1198 A (Leg 194 Marion Plateau, Australia). SwissSed Meeting, Fribourg, Switzerland, 29.
- Cappellacci, S., **Margreth, S.**, Tamburini, F., Grobety, B., Coric, S., Spezzaferri, S., 2006. Pleistocene glacial-interglacial dynamics at ODP Site 1198 (Australia): Micropaleontological and geochemical evidence. Forams 2006 - International Symposium on Foraminifera, Natal, Brasil, 240-241.
- Margreth, S.**, Tamburini, F., Grobety, B., Coric, S., Spezzaferri, S., Bernasconi, S., Cappellacci, S., 2006. Pleistocene glacial-interglacial dynamics along the Australian margin: ODP Leg 194, Site 1198. Swiss Geoscience Meeting, Bern, Switzerland, 125-126.

- Blass, A., Bigler, C., Bluszcz, P., **Margreth, S.**, Schmid, A., Trachsel, M., Troxler, A., von Gunten, L., Grosjean, M., Sturm, M., 2006. Paleo-climate research in the Engadin (Switzerland) - the state of the art. Swiss Geoscience Meeting, Bern, Switzerland, 25-26.
- Sturm, M., Gählman, V., Blass, A., Bühl, R., Kulbe, T., Livingstone, D.M., **Margreth, S.**, Renberg, I., 2005. Seasonal flux of particles and nutrients in mono-pagic lakes of Northern Scandinavia and the Alps. European Geosciences Meeting. European Geosciences Union, Wien, Austria. Geophysical Research Abstracts, Vol. 7, EGU05-A-05713, 2005.
- Blass, A., Buehler, R., Grosjean, M., **Margreth, S.**, Sturm, M., 2005. The sedimentation of the last few centuries in three proglacial lakes, Upper Engadine, Switzerland. In: Haas, H., Ramseyer, K., Schlunegger, F. (Eds.), Sediment 2005; The Sediment Factory. Deutsche Gesellschaft für Geowissenschaften. Hannover, Germany, 35.