

# Modern deep-water agglutinated foraminifera from IODP Expedition 323, Bering Sea: ecological and taxonomic implications



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**Abstract:** Despite the importance of the Bering Sea for subarctic oceanography and climate, relatively little is known of the foraminifera from the extensive Aleutian Basin. We report the occurrence of modern deep-water agglutinated foraminifera collected at seven sites cored during Integrated Ocean Drilling Program (IODP) Expedition 323 in the Bering Sea. Assemblages collected from core-top samples contained 32 genera and 50 species and are described and illustrated here for the first time. Commonly occurring species include typical deep-water *Rhizammina*, *Reophax*, *Rhabdammina*, *Recurvoides* and *Nodulina*. Assemblages from the northern sites also consist of accessory *Cyclammina*, *Eggerelloides* and *Glaphyrammina*, whilst those of the Bowers Ridge sites consist of other tubular genera and *Martinottiella*. Of the studied stations with the lowest dissolved oxygen concentrations, the potentially Bering Sea endemic *Eggerelloides* sp. 1 inhabits the northern slope, which has the highest primary productivity, and the potentially endemic *Martinottiella* sp. 3 inhabits Bowers Ridge, which has the lowest oxygen concentrations but relatively low annual productivity. *Martinottiella* sp. 3, with open pores on its test surface, has previously been reported in Pliocene to Recent material from Bowers Ridge. Despite relatively small sample sizes, ecological constraints may imply that the Bering Sea experienced high productivity and reduced oxygen at times since at least the Pliocene. We note the partially endemic nature of the agglutinated foraminiferal assemblages, which may at least in part be due to basin restriction, the geologically long time period of reduced oxygen, and high organic carbon flux. Our results indicate the importance of gathering further surface sample data from the Aleutian Basin.

**Keywords:** deep-water agglutinated foraminifera, Bering Sea, modern ecology, productivity, oxygen minimum zone

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The Bering Sea extends over a region comparable in size to the Mediterranean, yet the modern agglutinated foraminifera are still virtually unstudied. The interaction of strong currents, upwelling high nutrient water masses, sea ice and strong winds causes high surface water productivity which supports a diverse ecosystem (Stabeno *et al.* 1999) and an expanded oxygen minimum zone (OMZ). Relatively recent palaeoceanographic work indicates that the Bering Sea may have been characterized by high productivity and low oxygen since at least the Pliocene (Expedition 323 Scientists 2011; Kaminski *et al.* 2013) and, therefore, is an ideal place to study the long-term impact of severe hypoxia and high organic carbon flux on benthic organisms, in particular the less well-studied agglutinated foraminifera which are a diverse group particularly tolerant to ocean acidification due to their non-calcareous tests. Observational studies have recorded an expansion of tropical OMZs in the Pacific Ocean and Atlantic Ocean over the last 60 years, which is likely to continue with future increased atmospheric CO<sub>2</sub> emissions and oceanic sequestration (Stramma *et al.* 2008, 2010; Hofmann & Schellnhuber 2009). Studies of OMZ benthic ecology are, therefore, of particular interest (Gooday & Jorissen 2012). Although there have been several studies of modern benthic foraminifera from within OMZs world-wide (e.g. Hermelin & Shimmield 1990; Sen Gupta & Machain-Castillo 1993; Kaminski *et al.* 1995; Kaiho 1999; Gooday *et al.* 2000; Schumacher *et al.* 2007), there remains a lack of information from the Bering Sea.

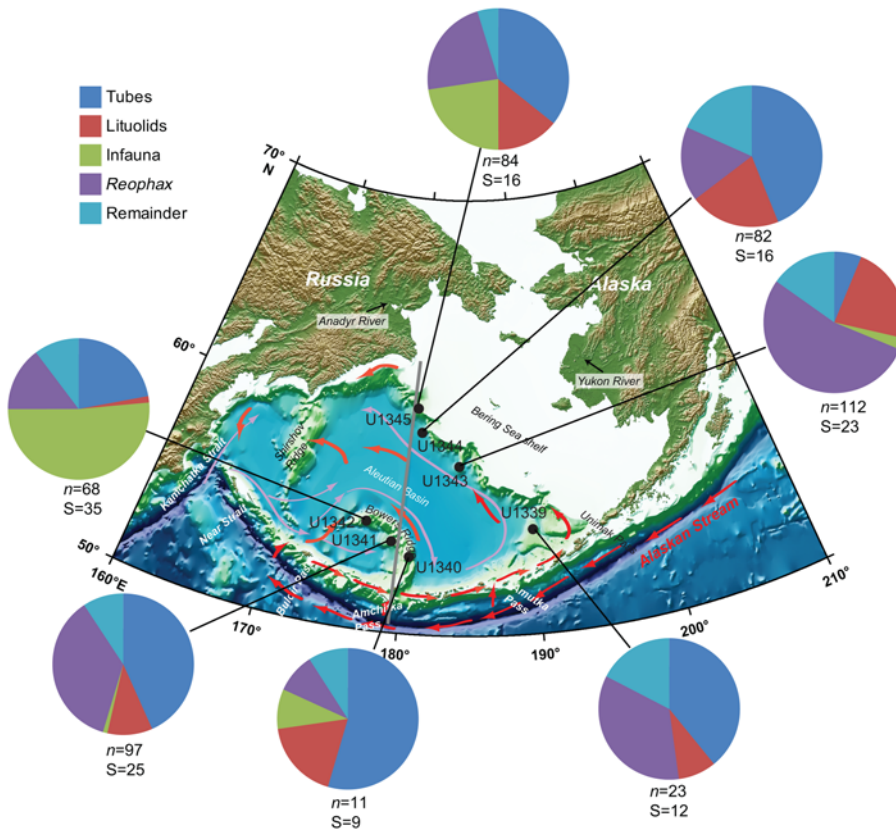
On account of the Bering Sea's high sedimentation rate along the slope, restricted deep-water circulation, low oxygen conditions and its partial isolation from the Pacific by the Aleutian Islands volcanic arc, the Bering Sea slope sites may be a good modern analogue to

the type of high sedimentation-rate deep-sea environments in the Cretaceous to Palaeogene Alpine–Carpathian and North Atlantic basins containing rapidly deposited orogenic-derived sediments called flysch. Under such conditions agglutinated foraminifera are an extremely important component of the benthic fauna, and fossil assemblages from the flysch basins are often comprised exclusively of agglutinated benthic foraminifera (e.g. Gradstein & Berggren 1981; Kender *et al.* 2005; Waskowska-Oliwa 2008; Setoyama *et al.* 2011).

In this study we fully document the agglutinated foraminifera in the deep (>800 m water depth) Bering Sea, in order to assess the degree of endemism in this restricted basin and to assess the possible ecological controls on agglutinated foraminiferal abundance.

## Bering Sea oceanography

Approximately half of the modern Bering Sea comprises a shallow (0–200 m) neritic environment, the remainder a vast plain c. 4 km deep broken by the Bowers and Shirshov ridges (Fig. 1). The northern continental shelf is covered seasonally by sea ice, with little ice presently being formed over the deep SW areas. The Bering Sea is one of the most highly biologically productive regions in the world, exporting some 687 000 tons of carbon per year (Sambrotto *et al.* 1984; Stabeno *et al.* 1999). 'Old' deep water, characterized by low oxygen concentrations, high nutrients (e.g. phosphate and nitrate) and high dissolved CO<sub>2</sub>, flows into the Bering Sea at depth from the North Pacific. It cycles counter-clockwise around the Bering Sea Basin, upwelling particularly over the continental shelf feeding the so-called 'Green Belt' (Springer *et al.* 1996). As large



**Fig. 1.** Bathymetric map of the Bering Sea (Expedition 323 Scientists 2010), showing IODP Expedition 323 sites reported in this study, and the number of specimens ( $n$ ) and number of taxa ( $S$ ) in major agglutinated foraminiferal assemblage components from core-top samples. Also indicated are the major surface (red arrows) and deep (purple arrows) ocean currents. Grey line corresponds to the section shown in Figure 3.

fluxes of organic carbon make their way to the seafloor in parts of the Bering Sea, particularly along the slope and over the shelf in spring (Fig. 2), intense oxygen demand expands the OMZ (Fig. 3), which impacts the composition of benthic foraminiferal communities and the chemistry of ocean water (Expedition 323 Scientists 2011). Significant exchange of Pacific deep water occurs through the Kamchatka Strait (maximum depth of 4420 m), and of low oxygen intermediate water through the Commander-Near Strait at 2000 m (Coachman *et al.* 1999). Very small amounts of bottom water are formed in the Bering Sea today (Warner & Roden 1995) and, as a result, the deep Bering Sea has an expanded OMZ in comparison with the northern Pacific.

### Previous studies of benthic foraminifera

Modern benthic foraminifera have been reported from Rose Bengal-stained core-top samples collected on the Bering Sea shelf at water depths less than 200 m (Anderson 1963). This study reported the occurrence of agglutinated foraminifera, which sometimes dominate the foraminiferal assemblages in the deeper shelf basins. Anderson (1963) reported that the proportion of agglutinated foraminifera may reach 90% of the total foraminiferal fauna on the central Bering Sea shelf. However, the modern deep-water agglutinated foraminifera from the deeper Aleutian Basin, within and below the OMZ, have yet to be documented. Khusid *et al.* (2006) studied the benthic foraminifera from a 660 cm long core collected at 3060 m depth on Bowers Ridge. In this core, the agglutinated foraminifera were found mainly in the core top and to a depth of 20 cm. The late Holocene agglutinated foraminifera comprised 83–99% of the fauna at this location, and consisted of *Rhabdammina*, *Hormosina*, *Ammolagena*, *Cribrostomoides* and *Karreriella*. However, neither Anderson (1963) nor Khusid *et al.* (2006) provided any descriptions or illustrations of the agglutinated foraminifera.

The agglutinated foraminifera from the North Pacific and Siberian Arctic have been more intensively studied than the fauna

from the Bering Sea. In this study we made use of the taxonomic monographs of Cushman (1910, 1921), Saidova (1975), Matoba & Fukusawa (1992) and Zheng & Fu (2001) on North Pacific foraminifera; the work of Vázquez Riveiros & Patterson (2007) on the foraminifera from the North Pacific Fjords; as well as studies on Arctic foraminifera by Cushman (1944), Wollenburg (1992, 1995) and Lukina (2001). The distribution of foraminifera along the North Pacific continental margins was compiled by Culver & Buzas (1985, 1987). Szarek (2001, unpublished PhD thesis, 'Biodiversity and biogeography of recent benthic foraminiferal assemblages in the south-western South China Sea (Sunda Shelf)', Christian-Albrechts University, Kiel) provides an excellent taxonomic section and useful distributional data for Bering Sea fauna *Reophax bradyi* and *R. oviculus* in the South China Sea. The current study aims to bridge a geographical gap in our knowledge of the distribution of North Pacific–Arctic agglutinated foraminifera, by providing descriptions of species recovered from the IODP Expedition 323 coring sites.

### Methods and materials

Samples were collected and prepared on board the JOIDES *Resolution* drillship from each site (U1339–45) during IODP Expedition 323, Bering Sea, in June/July 2009. Samples (quantities of sediment) were collected from the first cores recovered at each new hole (typically several holes were cored at each IODP site, within a distance of <20 m), by emptying the core barrel of unconsolidated core-top sediment into a cleaned receptacle. Samples ranged widely in size, but were generally >100 g in weight. Sediment composition varied between sites, but was varyingly dominated by diatoms and fine clays and silts with only rare coarser sand-sized particles. Two samples were then immediately stained in a Rose Bengal solution for >24 h to ascertain the living component. Samples were carefully washed over a >63  $\mu\text{m}$  mesh sieve with deionized water. Sample residues were oven dried at <40°C and agglutinated foraminifera picked out and placed in



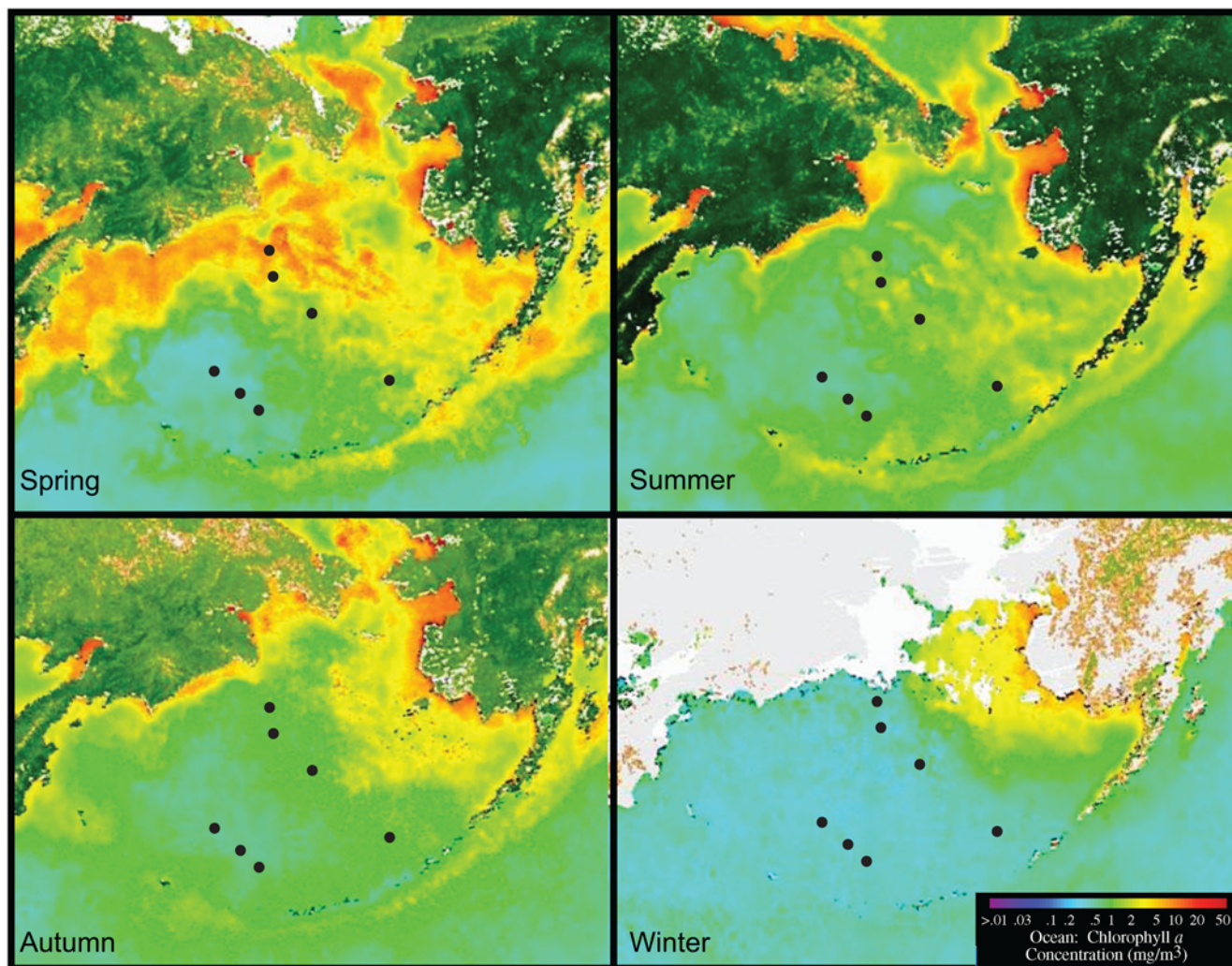


Fig. 2. Seasonal estimates from the years 1998–2003 of oceanic chlorophyll-*a* concentration ( $\text{mg m}^{-3}$ ) from satellite imagery SeaWiFS (<http://oceancolor.gsfc.nasa.gov/SeaWiFS>). Also shown are the positions of the seven IODP Expedition 323 sites (see Fig. 1 for labels).

cardboard reference slides. Specimens were imaged using a JSM-5900LV SEM at King Fahd University of Petroleum and Minerals in Dhahran, and a LEO 535VP SEM at the British Geological Survey in Keyworth. The proportion of faunal groups shown in Figure 1 was calculated for each site by combining the faunal counts of all samples from that site. Correspondence Analysis (CA), a reciprocal averaging algorithm, was carried out (using the software of Hammer *et al.* 2005) on the dataset to statistically ascertain the relationships between samples, species and selected environmental parameters (Figs 4 and 5), as described in Hammer & Harper (2006). CA in

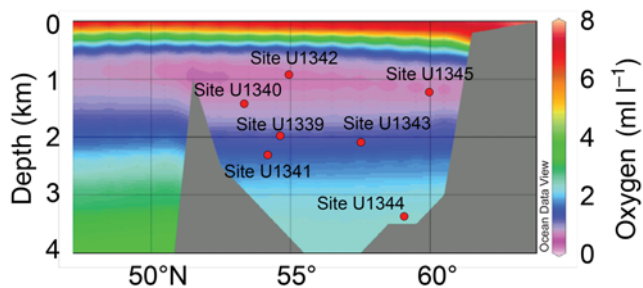
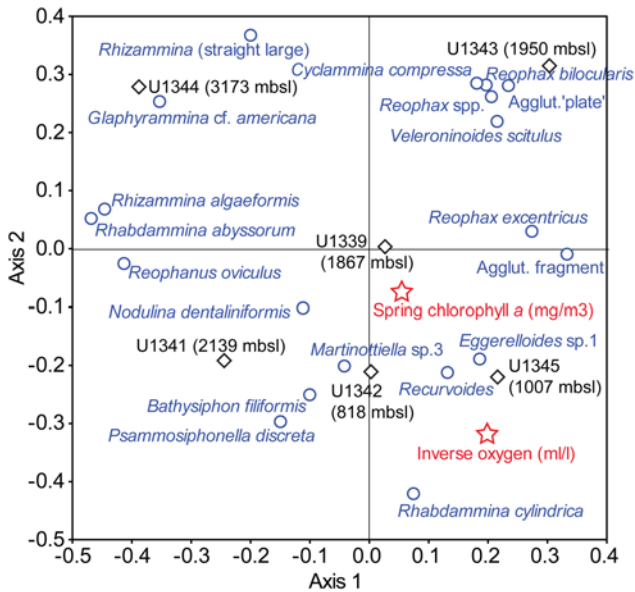


Fig. 3. Bering Sea north–south section (see approximate line in Fig. 1; Bowers Ridge not shown here) showing dissolved oxygen concentrations ( $\text{ml l}^{-1}$ ; Schlitzer 2000) and the position of IODP Expedition 323 sites analysed in this study (Expedition 323 Scientists 2010).

Figure 4 was carried out on a modified dataset, in order to incorporate environmental information with widely varying numerical values compared to species counts. Species counts were summed for each site and Site U1340 was removed because of low counts. Species that had an occurrence of  $<10$  in the total dataset were removed to avoid artificially placing emphasis on low counts. The amount of dissolved bottom water oxygen (estimated from World Ocean Circulation Experiment data, Fig. 3) and spring primary productivity (estimated from satellite chlorophyll-*a* data, Fig. 2) was recorded for each site (Table 1) and included in CA. All species counts and environmental parameters were subsequently normalized, and 2 added to each value to lift all numerical values above 0 (a requirement for CA). CA in Figure 5 was carried out on an unmodified dataset of all samples separately, in order to ascertain the degree of similarity between samples of different sites.

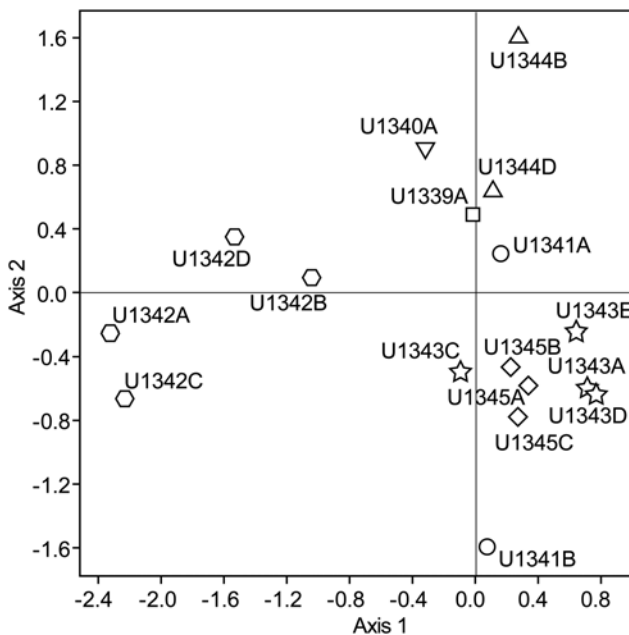
## Results

A total of 524 specimens of agglutinated benthic foraminifera were recorded, identified and described from 19 core-top samples taken at all seven IODP Expedition 323 sites (Figs 1 and 4; Table 2). The samples were dominated by calcareous benthic foraminifera, which were reported by Expedition 323 Scientists (2011) and reproduced in Table 3. Preservation of agglutinated taxa is generally excellent (see Figs 6–10), which is in marked contrast to the agglutinated-poor Pliocene–Recent palaeontological assemblages recovered



**Fig. 4.** Correspondence Analysis (CA) for dataset (including chlorophyll and inverse oxygen estimates; stars), showing species (circles) and sample (diamonds) scores for axis 1 against axis 2. Bottom water oxygen values were inverted, so that proximal species and samples exhibit low oxygen. Only species with >10 specimens are included (see 'Methods' for further details of data analysis). Mbsl, metres below sea-level.

from down-core samples (Expedition 323 Scientists 2011; Kaminski *et al.* 2013), indicating post-mortem dissolution of the organic cement likely occurred. Abundance is >70 specimens at most sites, apart from Sites U1339 and U1340 where abundances are low due to the small volume of core-top samples collected. CA (Fig. 5) indicates that there is generally greater similarity between samples from one site than between samples from different sites, as



**Fig. 5.** Correspondence Analysis (CA) for dataset, showing sample scores for axis 1 against axis 2. Samples plotting close together exhibit similar species compositions. The majority of samples plot close to other samples from the same site, showing the distinctiveness of assemblages from each site. Samples U1341B, U1343C and U1343D have very low abundances, explaining why they plot further away from the other samples of those two sites.

the majority of samples cluster near those of the same site. Samples U1341B and U1343C plot further away, which can be explained by their particularly low abundances (see Table 2). The generally low-diversity agglutinated assemblages (9–35 taxa per site) are predominantly composed of tubular suspension-feeders (e.g. *Rhabdammina*, *Rhizammina* and *Bathysiphon*), epifaunal lituolids (e.g. *Recurvoides*, *Cyclammina*), opportunistic infauna (e.g. *Reophax* and *Hormosinella*) and infauna (e.g. *Eggerelloides* and *Martinottiella*) in varying proportions (Fig. 1). CA indicates that some species are more prevalent at certain sites (i.e. plot in close proximity on Fig. 4); that Sites U1339, U1342 and U1345 are most associated with high chlorophyll-*a* concentrations (a proxy for primary productivity); and that Sites U1342 and U1345 are most associated with low bottom water dissolved oxygen. Two samples were stained with Rose Bengal (at Sites U1342 and U1345). These samples contained a small proportion of living individuals (Table 2), confirming that the IODP cores recovered samples of modern/sub-modern age.

## Discussion

### Endemism

Although a sizeable proportion of Bering Sea agglutinated foraminifera have been recorded in the Pacific Ocean (Jones 1994), there are several species in the core-top samples (this study) and in the Pliocene (Kaminski *et al.* 2013) that appear to be endemic (e.g. *Eggerelloides* sp. 1; *Glaphyrammina cf. americana*; *Martinottiella* sp. 1; *Martinottiella* sp. 2; *Martinottiella* sp. 3; *Karrieriella* sp. 1; *Bathysiphon* sp., *Hormosinelloides* sp.) and confirm the semi-isolated nature of the microfauna in the Bering Sea. Of the 131 agglutinated species recorded by Culver & Buzas (1985) from the North Pacific Margin (at 138 localities) only 13 are present in our samples. This low number of species in common suggests that many taxa present along the Alaskan margin are excluded from our study locations in the Bering Sea. In our current study of the agglutinated foraminifera, 22% of the species are left in open nomenclature and do not yet appear to have been described. In the Pleistocene calcareous benthic assemblage studied by Setoyama & Kaminski (2015) at Site 1341, 23% of the taxa were identified tentatively or left in open nomenclature. In contrast Culver & Buzas (1985) reported only a few species in open nomenclature. Geographical barriers for faunal interchange between the Bering Sea and North Pacific include the restricted Aleutian passes (Fig. 1), although the western passes are deep (>4 km) and the majority of Bering Sea benthic species recorded in our study are cosmopolitan. It is therefore possible that environmental conditions in the isolated Bering Sea, such as high productivity and reduced bottom water oxygen, have allowed for the adaptation of certain new species or varieties.

Considering the long stratigraphic ranges of the majority of benthic foraminifera (e.g. Kaminski & Gradstein 2005; Holbourn *et al.* 2013), and their relatively slow genetic evolution compared with planktonics (Pawlowski *et al.* 1997; Gooday & Jorissen 2012), the occurrence of endemic species is consistent with a Bering Sea that may have been isolated for a considerable length of time. This is not unique for semi-isolated deep-water basins, and one such example is the high-latitude Norwegian Sea during the Eocene, when it was separated from the North Atlantic by the Greenland–Scotland Ridge. The deep-water agglutinated foraminiferal assemblages that developed during the Eocene and Oligocene in this basin contain a number of endemic species that have not been found in the northern Atlantic (Gradstein & Kaminski 1989; Kaminski & Gradstein 2005). The Oligocene deep-water agglutinated foraminiferal assemblage at Site 985A on the Iceland Plateau contains 27% endemic species (Kaminski & Austin 1999). Agglutinated

**Table 1.** Location, water mass properties and average sedimentation rate data of the IODP Expedition 323 sites analysed in this study

Site	Latitude	Longitude	Water depth (mbsl)	Ave. sedimentation rate (cm ka <sup>-1</sup> )	Estimated bottom water oxygen (ml l <sup>-1</sup> )	Bottom water temp. (°C)	Bottom water salinity (psu)	Spring chlorophyll- <i>a</i> (mg m <sup>-3</sup> )	Winter chlorophyll- <i>a</i> (mg m <sup>-3</sup> )
323-U1339A	54° 40.2001' N	169° 58.9017' W	1866.7	28.0	1.1	2.0	34.7	2.00	0.60
323-U1339D	54° 40.1891' N	169° 58.8909' W	1868.1	28.0	1.1	2.0	34.7	2.00	0.60
323-U1340A	53° 24.0008' N	179° 31.2973' W	1294.7	14.5	0.7	2.5	34.4	0.70	0.37
323-U1341A	54° 2.0025' N	179° 0.4999' E	2139.6	14.5	1.5	1.9	34.7	0.50	0.37
323-U1341B	54° 1.9984' N	179° 0.5171' E	2139.6	14.5	1.5	1.9	34.7	0.50	0.37
323-U1341C	54° 2.0010' N	179° 0.5390' E	2139.6	14.5	1.5	1.9	34.7	0.50	0.37
323-U1342A	54° 49.6987' N	176° 55.0027' E	818.3	4.5	0.6	3.0	34.3	0.40	0.37
323-U1342B	54° 49.7004' N	176° 55.0232' E	818.9	4.5	0.6	3.0	34.3	0.40	0.37
323-U1342C	54° 49.7017' N	176° 55.0232' E	818.8	4.5	0.6	3.0	34.3	0.40	0.37
323-U1342D	54° 49.6987' N	176° 55.0027' E	818.2	4.5	0.6	3.0	34.3	0.40	0.37
323-U1343A	57° 33.3993' N	175° 48.9659' W	1950.9	35.0	1.2	2.0	34.7	1.40	0.39
323-U1343B	57° 33.4156' N	175° 48.9951' W	1950.9	35.0	1.2	2.0	34.7	1.40	0.39
323-U1343C	57° 33.3982' N	175° 49.0275' W	1952.6	35.0	1.2	2.0	34.7	1.40	0.39
323-U1343D	57° 33.3817' N	175° 48.9971' W	1954.1	35.0	1.2	2.0	34.7	1.40	0.39
323-U1343E	57° 33.3814' N	175° 48.9974' W	1956.0	35.0	1.2	2.0	34.7	1.40	0.39
323-U1344A	59° 3.0005' N	179° 12.2011' W	3171.8	45.0	2.3	1.7	34.7	3.50	0.40
323-U1344B	59° 3.0112' N	179° 12.2051' W	3173.0	45.0	2.3	1.7	34.7	3.50	0.40
323-U1344C	59° 3.0116' N	179° 12.2052' W	3172.7	45.0	2.3	1.7	34.7	3.50	0.40
323-U1344D	59° 3.0224' N	179° 12.2030' W	3174.1	45.0	2.3	1.7	34.7	3.50	0.40
323-U1345A	60° 9.1917' N	179° 28.2036' W	1007.4	29.0	0.6	2.5	34.4	8.00	0.50
323-U1345B	60° 9.2003' N	179° 28.2127' W	1007.5	29.0	0.6	2.5	34.4	8.00	0.50
323-U1345C	60° 9.2097' N	179° 28.2229' W	1008.8	29.0	0.6	2.5	34.4	8.00	0.50
323-U1345D	60° 9.2175' N	179° 28.2283' W	1008.3	29.0	0.6	2.5	34.4	8.00	0.50

Expedition 323 Scientists (2010). Location, water mass properties are estimated from Figs 1, 2 and WOCE data

**Table 2.** Counts of all agglutinated foraminifera in core-top samples from IODP Expedition 323 sites

Species	U1339A	U1340A	U1341A	U1341B	U1342A	U1342A- stained	U1342B	U1342C	U1342D	U1343A	U1343C	U1343D	U1343E	U1344B	U1344D	U1345A	U1345B	U1345B- stained	U1345C
Agglutinated fragments	9	1	3	2	1		1		1	7		5	9			9	3		11
<i>Ammodiscus</i> sp.			1																
<i>Archimerismus subnodosus</i>			2				1							4					2
<i>Astrorhiza granosa</i>																	1		
<i>Bathysiphon filiformis</i>			7						1							1			3
<i>Bathysiphon</i> sp. 'coarse'			3																
<i>Cribrostomoides subglobosus</i>											1								
<i>Cyclamina compressa</i>										12				1					
<i>Dendrophyra arborescens</i>										2	1								
<i>Eggerelloides</i> sp. 1					1			1			3					1	4		14
<i>Evolutinella rotulata</i>		1																	
<i>Glaphyrammina</i> cf. <i>americana</i>	1												1	15	2				
<i>Hormosinella distans</i>			2																
<i>Hormosinelloides</i> sp. aff. <i>H. guttifer</i>	1	1	1											6					
<i>Hyperammina</i> spp.	3					1				2									
<i>Karrerella</i> sp. 1				1															
<i>Lagenammina</i> sp. 'spicules'		1																	
<i>Lagenammina</i> spp.			1							1			1						
Large agglutinated 'plate'	2									3		1	3	2		2			1
<i>Marsipella elongata</i>			1													1			1
<i>Martinotiella</i> sp. 3		1			19	1	4	8	1										
<i>Nodulina dentaliniformis</i>	5		9		1		1			4									
<i>Nothia</i> sp. 'large spicules'		1					1								1				
? <i>Nothia</i> sp. 'diatoms'															2		2	1	
<i>Psammosiphonella discrete</i>			4	6						1						1			4
<i>Psammosphaera fusca</i>	2	1								1	1		1	3					
<i>Recurvoides</i> spp.	1		8	1						4			2			2	2	4	6
<i>Reophanus oviculus</i>			11											7					
<i>Reophax</i> aff. <i>brevis</i>			1																
<i>Reophax agglutinatus</i>	1				1	1	1		1					1					
<i>Reophax bilocularis</i>							1			13				3		2			1
<i>Reophax duplex</i>													1						
<i>Reophax excentricus</i>			4							11			3				5		8

<i>Reophax pauciloculatus</i> (cf. <i>pilulifer</i> )			5		1	1													
<i>Reophax pilulifer</i>	1					1							1						
<i>Reophax scorpiurus</i>			2						2				4				1		
<i>Reophax</i> sp. B 'tufty spicules'			2																
<i>Reophax spiculifer</i>			1													1			
<i>Reophax</i> spp.			1						18		1	2				2			3
<i>Rhabdammina abyssorum</i>			5					1							2	4			
<i>Rhabdammina cylindrica</i>	3	2	4		1			5							2	3	1		3
<i>Rhabdammina</i> sp. 'smooth spicules'		1															1		2
<i>Rhabdamminella cylindrical</i>			1																
<i>Rhizammina algaeformis</i>	1		12			1		4			1			1	20		2	1	2
<i>Rhizammina</i> sp. 'straight large'	2	2							2			1	5	1			1		
<i>Saccorhiza ramosa</i>			3																
Soft saccamminid						3		2											
<i>Subreophax splendidus</i>			1																
<i>Thurammina albicans</i>									1										
<i>Tolypammina vagans</i>			1																
<i>Trochammina</i> sp.										1									
<i>Veleroninoides scitulus</i>		1	1						7							1			
Forams per sample	32	13	97	10	25	9	21	11	4	91	9	7	28	49	35	22	25	7	61
Species per sample	13	11	28	4	7	7	10	4	4	17	7	3	11	12	9	10	11	4	14

Specimens stained with Rose Bengal (live fauna) are indicated, which constitute 9 stained specimens in sample 323-U1342, and 7 stained specimens in sample 323-U1345.



**Table 3.** Semi-quantitative abundances of calcareous foraminifera in core-top samples from IODP Expedition 323 sites

Species	U1339D	U1340A	U1341A	U1341B	U1341C	U1342C	U1342D	U1343A	U1343B	U1343C	U1343D	U1343E	U1344A	U1344B	U1344C	U1344D	U1345D
Preservation	G	G	G	VG	VG	VG	VG	G	VG	G	G	G	G	VG	M	G	G
<i>Alabaminella weddellensis</i>		P															
<i>Bolivina</i> sp.	F				F												A
<i>Brizalina</i> cf. <i>spathula</i>				F		F	A	P			P	P					F
<i>Brizalina earlandi</i>		R		R		A	F	R		P	F	P			R		
<i>Brizalina pygmaea</i>				P	P	A	A	P									
<i>Bulimina</i> aff. <i>exilis</i>		P		A		A	F	R	F	R	A	P					D
<i>Bulimina</i> sp.		P			D			P									
<i>Cancris</i> cf. <i>phillipinensis</i>						F											
<i>Cassidulina</i> sp.							D	F									
<i>Cassidulinoides tenuis</i>									P								
<i>Elphidium</i> cf. <i>battalis</i>	F							P				R		R			R
<i>Epistominella pulchella</i>					R												
<i>Globobulimina auriculata</i>				P	F			P			P			P			
<i>Globobulimina pacifica</i>	R		P	F			P	R	R	R		F	P	R			R
<i>Globocassidulina</i> sp.		P	P							P							
<i>Gyroidinoides soldanii</i>														P			
<i>Islandiella norcrossi</i>	R	P				F		P		P		P	R				R
<i>Nodosaria</i> spp.									P								
<i>Nonionella labradorica</i>	F							R	F	R	R	A		F			
<i>Nonionella turgida</i>				R										P			
<i>Nonionella turgida digitata</i>								R	A	P	A	P			P		
<i>Procerolagena</i> cf. <i>gracillima</i>										P	R						
<i>Pullenia bulloides</i>												P					
<i>Pygmaesestron</i> cf. <i>hispida</i>												P					
<i>Pyrgo</i> sp.																	
<i>Quinqueloculina</i> sp.																	
<i>Stainforthia</i> aff. <i>fusiformis</i>		P					R	P									
<i>Triloculina</i> cf. <i>trihedra</i>				P													
<i>Uvigerina auberiana</i>			P		R									F			
<i>Uvigerina</i> cf. <i>peregrina</i>	A		P					R	P	P		P					F
<i>Valvulinera</i> sp.							R			P							

From Expedition 323 Scientists (2011). D, dominant; A, abundant; F, few; R, rare; P, present; VG, very good; G, good; M, medium.





**Fig. 6.** (1) *Astrorhiza granulosa* (Brady, 1879), Hole U1342B. (2) *Dendrophyra* sp., Hole U1343A. (3) *Nothia* sp. 'large spicules', Hole U1341A. (4) ? *Nothia* sp. 'diatoms', Hole U1345B. (5) *Marsipella elongata* Norman, 1878, Hole U1342A. (6–8) *Rhabdammina* spp.: 6, Hole U1342A; 7–8, Hole U1344. (9) *Bathysiphon* sp. 'coarse', Hole U1341A. (10) *Bathysiphon filiformis* G.O. & M. Sars, 1872, Hole U1345C. (11) *Psammosiphonella discreta* (Brady, 1881), Hole U1341A. (12–13) *Rhabdamminella cylindrica* (Brady, in Tizard & Murray, 1882): 12, Hole U1340A; 13, Hole U1339A. (14a, b) *Rhizammina algaeformis* Brady, 1879, Hole U1344. (15) *Rhizammina* sp. 'straight large'. Hole U1342B. Scale bar 200 µm.

foraminifera were studied by Matoba & Fukusawa (1992) in the semi-enclosed Sea of Japan, and 26% of the reported taxa were not identified at species level.

### Ecological implications

Benthic foraminifera are impacted by several ecological forcing factors, which include organic carbon type and flux, bottom water oxygenation, bottom water sediment heterogeneity and hydrodynamics, temperature and corrosiveness (Jorissen *et al.* 1995, 2007; Levin *et al.* 2001). Bering Sea deep-water (>800 m water depth) ecology is primarily affected by two of these ecological forcing factors: dissolved oxygen concentrations and organic carbon flux (Expedition 323 Scientists 2011), which is related to primary productivity (quantity, type and duration) and remineralization of particulate organic carbon as it is transported to depth (Arndt *et al.* 2013). Other factors that might affect benthic assemblages in the Bering Sea are the high sedimentation rates along the slope (Table 1). Oxygen levels in the deep Bering Sea are very low (*c.* 2.0–0.2 ml l<sup>-1</sup>), and primary productivity highly variable (Figs 2 and 3), which can be expected to impact upon assemblage composition (e.g. Jorissen *et al.* 1995, 2007; Kaminski *et al.* 1995; Sun *et al.* 2006). Indeed, oxygen concentrations in the bottom water have been hypothesized as significantly reduced compared with the open Pacific (Fig. 3) since at least the Pliocene, when laminations are pervasive in Bering Sea sediment cores and occasional calcareous faunas are dominated by deep infaunal taxa (Expedition 323 Scientists 2011).

The diverse calcareous and agglutinated foraminifera reported in the core-top material (Tables 2 and 3) are somewhat typical for low oxygen and high productivity environments, as described from the Santa Catalina Basin (Kaminski *et al.* 1995), Santa Barbara Basin (Moffitt *et al.* 2014), the Arabian Sea (Gooday *et al.* 2000; Schumacher *et al.* 2007) and OMZs elsewhere (see Sen Gupta & Machain-Castillo 1993). Assemblages within the core of the OMZ are typically dominated by calcareous infaunal taxa that exhibit elongated tapered tests, which are predominantly *Bulimina*, *Brizalina* and *Bolivina* in the Bering Sea (Table 3). The relatively less-specialized agglutinated foraminifera usually occur in higher abundances above and below the core of the OMZ (Kaminski *et al.* 1995; Schumacher *et al.* 2007). Our samples are dominated by calcareous foraminifera, which is consistent with high productivity and low oxygen settings in the Okhotsk Sea (Bubenshchikova *et al.* 2008). The most commonly occurring cosmopolitan agglutinated species in our material (Fig. 4) are wide-ranging and described from diverse environments. *Rhizammina algaeformis* and *Nodulina dentaliniformis* are well-known cosmopolitan species ranging from neritic to abyssal depths. *Nodulina dentaliniformis* has been recorded from relatively shallow water in the Arctic (Lukina 2001) and Antarctic (Majewski 2005) where it may be tolerant of changes in salinity. *Reophax excentricus* and *Reophax bilocularis* are cosmopolitan open ocean species, and were recorded as part of assemblages within the OMZ of the Santa Catalina Basin (Kaminski *et al.* 1995). *Reophax bilocularis* was also recorded in high proportions along the slope beneath the Arabian Sea OMZ associated with high sedimentation rates (Hermelin & Shimmield 1990) and occurs in relatively high abundances at Site U1343 along the slope, where sedimentation rates are higher (Table 1). *Cyclammina compressa* is a less well-known bathyal to abyssal species originally described from the Philippines and also recorded offshore North Carolina (Gooday *et al.* 2001), and has close morphological affinities with the cosmopolitan and wide-ranging *C. cancellata* (see Jones 1994).

Although we obtained no faunal density data (because of the limited availability of equipment onboard the JOIDES *Resolution*), with such strong ecological gradients between sites it is possible to

speculate on the ecology of some of the more abundant key species (Fig. 4). We caution that our estimates of modern ecological parameters (i.e. dissolved bottom water oxygen and primary productivity, Figs 2 and 3; Table 1) at each site are only approximations, as there were no *in situ* water mass measurements made during Expedition 323. The line section of Figure 3 does, however, pass in close proximity to all sites apart from U1339 (see line in Fig. 1). It should also be considered that our benthic faunas probably represent several decades at least, and that the primary productivity proxy chlorophyll-*a*, with its own proxy uncertainties (see Sun *et al.* 2006), represents the years 1998–2003. In addition, primary productivity has only an indirect impact on benthic faunas, as much of the organic carbon is remineralized on its way to the seafloor (see Arndt *et al.* 2013). However, we consider that the largest changes in these parameters between sites will be semi-quantitatively resolved by our estimates. *Glaphyrammina cf. americana* is restricted to Site U1344 (central slope), where the highest bottom water oxygen, highest sedimentation rate and deepest water depth is recorded (Table 1). The deep-water setting probably experiences lower organic carbon fluxes compared to the other slope sites (as organic carbon is remineralized in the water column) and so this species may be adapted to more oligotrophic environments with relatively elevated oxygen levels. At Site U1345 (northern slope), where there is the highest year-round chlorophyll-*a* concentration (and assumed organic carbon flux) and low bottom water oxygen (Figs 2 and 3), the endemic species *Eggerelloides* sp. 1 occurs in relative high abundance (Fig. 4; Table 2) and therefore this species may be adapted to high organic carbon flux in low oxygen settings. The morphologically similar species from the North Pacific, *Eggerelloides advenum*, has been associated with intense eutrophication in Osaka Bay (Tsujimoto *et al.* 2006). At Sites U1342 and U1340 (Bowers Ridge), the only other sites situated in the core of the OMZ (Fig. 1), the endemic species *Martinottiella* sp. 3 is observed (Fig. 4; Table 2). Due to its distribution, we speculate that this species may be adapted to low oxygen environments (e.g. below *c.* 1 ml l<sup>-1</sup>; Table 1), but not high organic carbon flux (as it does not occur along the northern slope but in the more oligotrophic south-central Bering Sea). Kaminski *et al.* (2013) were the first to observe the highly perforate tests of *Karreriella* and *Martinottiella* from the Bering Sea (see Fig. 10:1–10:5) and suggested this feature may have been an adaptation to severely hypoxic conditions, which is supported by the species modern distribution recorded here. *Martinottiella* sp. 3 is larger and more robust than the otherwise morphologically similar species *Martinottiella* sp. 1, recorded from the Pliocene of Bowers Ridge (Kaminski *et al.* 2013), which may have been its evolutionary ancestor. The modern-day distribution of the morphologically similar (although possibly lacking perforations) *Martinottiella* sp. (cf. *M. communis*) is world-wide, including within the OMZ of the Santa Catalina California Borderland basin (Kaminski *et al.* 1995), highly productive areas of the South China Sea (Jian *et al.* 1999), the OMZ of the equatorial East Pacific (Culver & Buzas 1987) and East New Zealand, South Pacific (Hayward *et al.* 2001). It is sometimes associated with high organic carbon flux and low oxygen settings, but its modern-day ecology is yet to be fully resolved. *Martinottiella* spp. is dissolution resistant and survives taphonomic loss, so this taxon can be used as a palaeoenvironmental indicator. Echols (1973), Expedition 323 Scientists (2011) and Kaminski *et al.* (2013) recorded *Martinottiella* sp. (cf. *M. communis*) from several locations in the Bering Sea (including Bowers Ridge) from the Pliocene to Recent. Due to the new distributional data reported here, we suggest that the significance of the Pliocene occurrences of *Martinottiella* may be an indication of low oxygen conditions at times since at least the Pliocene. Our study highlights, however, the importance of obtaining more bottom water and surface sample material from the Aleutian Basin for further study.

We cannot be sure of *Eggerelloides* sp. 1 and *Martinottiella* sp. 3 living depth preferences within the sediment. However, many authors have attempted to ascertain palaeoecology from ancient sediments by placing agglutinated foraminifera into groups on the basis of their morphology; these ‘morphogroups’ are thought to be indicative of their ecology (e.g. Nagy 1992; van den Akker *et al.* 2000; Kaminski *et al.* 2005; Kender *et al.* 2008a, b; Nagy *et al.* 2009; Nikitenko *et al.* 2013). These are based on studies of modern foraminifera and bottom water properties, such as living depth, productivity and ecological disturbance (Jones & Charnock 1985; Kaminski *et al.* 1995). Both *Eggerelloides* and *Martinottiella* are elongated and tapered in shape and would be assigned to the ‘morphogroup 4b’ of Kaminski & Gradstein (2005), a group regarded as infaunal and tolerant of low oxygen conditions. This approach to reconstructing palaeoenvironments using morphogroups should be taken with caution and may be an oversimplification (e.g. Sen Gupta & Machain-Castillo 1993; Jorissen *et al.* 2007), particularly as *Eggerelloides* and *Martinottiella* do not co-occur in the same samples (Table 2). However, the distribution of this group within the Bering Sea OMZ sites (U1340, U1342 and U1345; see distribution in Fig. 1) supports the interpretation of an infaunal living habit.

## Conclusions

We document the occurrence of 50 modern agglutinated foraminiferal taxa at IODP Expedition 323 sites in the Bering Sea, and provide the first descriptions and illustrations. The 19 core-top samples at seven sites, U1339, U1340, U1341, U1342, U1343, U1344 and U1345, contain abundant agglutinated foraminifera in varying proportions, and calcareous benthic foraminifera previously reported (Expedition 323 Scientists 2011), many of which are typical for reduced oxygen and high productivity environments. The agglutinated foraminifera consist of several abundant and ecologically wide-ranging cosmopolitan taxa, and also a number of taxa (e.g. *Glaphyrammina*, *Martinottiella*, *Eggerelloides*, *Bathysiphon*, *Hormosinelloides* and *Karreriella*) that differ in morphology from their counterparts and are here left in open nomenclature. The agglutinated foraminiferal fauna of the deep Bering Sea is thus partially endemic, suggesting that geographical restriction, combined with high productivity and low oxygen environmental conditions, may have persisted within the Bering Sea for a geologically extended period of time, considering the low evolutionary rate of the group. Three of the more abundant endemic species may be ecologically restricted. *Glaphyrammina* cf. *americana* occurs largely at the slope Site U1344, which is the deepest (with likely low organic carbon flux), highest bottom water oxygen and highest sedimentation rate site sampled. *Eggerelloides* sp. 1 occurs in high abundance at the northern slope Site U1345, which experiences the highest seasonal productivity (and possibly organic carbon flux) of the seven sites. *Martinottiella* sp. 3 is restricted to the OMZ of Bowers Ridge, where there is currently relatively low annual productivity, suggesting this species may be a good indicator for reduced oxygen conditions but not elevated organic carbon flux. The occurrence of *Martinottiella* throughout the Bering Sea sporadically over the past c. 4 Ma (Echols 1973; Expedition 323 Scientists 2011; Kaminski *et al.* 2013) therefore adds evidence to the hypothesis that the Bering Sea has had a pronounced OMZ since at least the Pliocene. However, due to the relatively low number of samples, our study highlights the importance of collecting more data from the currently under-sampled deep Aleutian Basin. In addition, two species of agglutinated foraminifera (*Karreriella* sp. 1 and *Martinottiella* sp. 3) were found to contain micro-pores that are open at the test surface, a morphological feature that is possibly indicative of hypoxia (Kaminski *et al.* 2013).

## Systematic palaeontology

In this section species are arranged in taxonomic order according to the classification of Kaminski (2014). Descriptions and comments are provided and important references for understanding each species morphology and distribution are cited. For taxonomic determinations the monographs of Cushman (1910, 1921, 1944), Cushman & McCulloch (1939), Pflieger (1952), Vilks (1969), Saidova (1975), Wollenberg (1992, 1995), Jones (1994), Kaminski & Gradstein (2005) and Vázquez Riveiros & Patterson (2007) were mainly used, and direct comparisons were made with specimens preserved in the HMS *Challenger* Collections at the Natural History Museum (London). The specimen microslides have been deposited in the collections of the European Micropalaeontological Reference Centre at Micropress Europe, Kraków Poland.

Subclass **Monothalamana** Pawlowski, Holzmann & Tyszka, 2013

Genus *Astrorhiza* Sandahl, 1858

*Astrorhiza granulosa* (Brady, 1879)  
(Fig. 6:1)

1879 *Marsipella granulosa* Brady: 38, pl. 3, figs 8–9.

1881 *Astrorhiza granulosa* (Brady); Brady: 48.

1884 *Astrorhiza granulosa* (Brady); Brady, 234, pl. 20, figs 14–23.

2000 *Astrorhiza granulosa* (Brady); Gooday & Smart: 107, pl. 4, figs 1–7.

**Remarks.** A single specimen was found in Hole U1342A. It is triangular in outline, with a coarsely agglutinated wall. It corresponds well with specimens illustrated by Brady (1884, pl. 20). Gooday & Smart (2000) showed that the species has a two-layered wall and agglutinates juvenile planktonic foraminiferal tests to construct its outer layer. Our specimen from Hole U1342B only uses mineral grains.

Genus *Marsipella* Norman, 1878

*Marsipella elongata* Norman, 1878  
(Fig. 6:5)

1878 *Marsipella elongata* Norman: 281, pl. 16, fig. 7.

1884 *Marsipella elongata* Norman; Brady: 264, pl. 24, figs 10–19

2008 *Marsipella elongata* Norman; Kaminski *et al.*: 64, pl. 2, figs 1–4.

**Description.** Test up to 3 mm in length, elongated, tubular, cylindrical, or tapering at both ends, may be slightly twisted or sinusoidal. Wall thin, of agglutinated quartz grains and sponge spicules, firmly cemented. Spicules are concentrated near the tapering ends of the tube. Apertures at the open ends of the tube.

**Remarks.** The type specimen in the Norman Collection is from Porcupine Station 87 in the North Atlantic (59° 35' N, 9° 11' W; 767 fathoms) and is preserved in the NHM, London, in slide 1915.4.1.852. The specimens from Hole U1345C conform well with the type specimen.

Genus *Rhabdammina* Sars in Carpenter, 1869

*Rhabdammina* spp.  
(Fig. 6:6–6:8)

**Remarks.** We used this name to describe fragments of straight tubes, not precisely determined. A distinctive specimen recovered in



Hole U1343A agglutinates grains of very unequal size.

*Rhabdammina* 'smooth spicules'

**Remarks.** A single specimen was found in Hole U1340A. This species has a very thick wall with sponge spicules embedded in a matrix of finer agglutinated particles. The spicules are aligned at right angles to the long axis of the test.

*Bathysiphon* Sars, 1872

*Bathysiphon* sp. 'coarse'  
(Fig. 6:9)

**Remarks.** A large specimen was recovered at Site U1341. The test is over 3 mm in length, arched and the early part of the test is tapered. The wall is coarsely agglutinated and constructed of grains of uneven dimensions. It most closely resembles *Bathysiphon rufus* de Folin, but the species in the NE Atlantic can reach a much larger size, up to 14 mm in length (Gooday 1988a). Gooday (1988b) designated a lectotype of *B. rufus* from the de Folin Collection, but the type locality is unknown.

*Bathysiphon filiformis* G.O. & M. Sars in Sars, 1872  
(Fig. 6:10)

1872 *Bathysiphon filiformis* G.O. & M. Sars in Sars: 251.  
1988b *Bathysiphon filiformis* G.O. & M. Sars in Sars; Gooday: 97, figs 1–3 (fig. 1a is the neotype).

**Remarks.** Test consists of a straight unbranched elongated tube, broken at both ends. Wall very finely agglutinated, thick, with a smooth finish. Gooday (1988b) credited the authorship of the species to both authors – G.O. & M. Sars – and designated a neotype from the Brady Collection (ex Norman collection, Hardanger Fjord, Norway). The specimen is preserved at the NHM, London, in slide BNHM 1887.8.31.1. The specimens from Hole U1341A and U1345C are fragmentary, but conform well to the types.

Genus *Nothia* Pflaumann, 1964

*Nothia* sp. 'large spicules'  
(Fig. 6:3)

**Remarks.** A thin-walled tube constructed of fine mineral grains with agglutinated sponge spicules attached to the surface of its test. The sponge spicules project at a 45° angle away from the test. Only broken fragments were found.

?*Nothia* sp. 'diatoms'  
(Fig. 6:4)

**Remarks.** We used this designation for fragments of a flattened thin-walled tube that contains a high proportion of diatoms incorporated into the wall. Specimens from Hole U1345B use a mixture of mineral grains and centric diatoms, while the specimens from Hole U1344D use mostly diatoms to construct their test. Fossil species of *Nothia* (e.g. *Nothia excelsa* Grzybowski) are commonly flattened, implying that their wall was flexible.

Genus *Psammosiphonella* Avnemelech, 1952

*Psammosiphonella discreta* (Brady, 1881)  
(Fig. 6:11)

1881 *Rhabdammina discreta* Brady: 48.

1884 *Rhabdammina discreta* Brady; Brady: 268, pl. 22, figs 8–10.  
1952 *Psammosiphonella discreta* (Brady); Avnemelech: 65.  
2005 *Psammosiphonella discreta* (Brady); Kaminski & Gradstein: 117, pls 5–6, figs 1–8.

**Description.** Test tubular, round in cross-section, straight, of even diameter or with slight constrictions. The inner surface of the tube is even, not constricted. Wall thick, composed of mineral grains, mostly quartz with some dark mafic grains, with organic cement. Apertures at the open ends of the (broken) tube.

**Remarks.** The type specimens of *P. discreta* are from Porcupine station no. 4, 808 fathoms water depth in the North Atlantic (BMNH ZF 4863–4865). A lectotype from this sample, corresponding to the specimen illustrated by Brady (1884, pl. 22, fig. 8), was designated by Kaminski & Gradstein (2005). In the area of the Philippines, Cushman (1921) listed it from 221 to 985 fathoms depth. These specimens are much larger (up to 18 mm length) and are comprised entirely of quartz grains. Our specimens are broken into small fragments and agglutinate some dark grains, giving the test a 'salt and pepper' appearance.

Genus *Rhabdamminella* de Folin, 1887

*Rhabdamminella cylindrica* (Brady, in Tizard & Murray, 1882)  
(Fig. 6:12, 6:13)

1882 *Marsipella cylindrica* Brady, in Tizard & Murray: 714.  
1884 *Marsipella cylindrica* Brady, in Tizard & Murray; Brady: 265, pl. 24, figs 20–22.  
1987 *Rhabdamminella cylindrica* (Brady, in Tizard & Murray); Loeblich & Tappan: 23, pl. 14, figs 2, 3.  
2008 *Rhabdamminella cylindrica* (Brady, in Tizard & Murray); Kaminski *et al.*: 65, pl. 3, figs 3–5 (fig. 3 is the lectotype).

**Description.** Test an elongated slender tube of constant diameter, may be slightly arcuate. Wall constructed of firmly cemented acicular sponge spicules, aligned more or less parallel to the long axis of the test in more or less irregular overlapping tiers; aperture at the open ends of the tube.

**Remarks.** *Rhabdamminella* differs from *Marsipella* in being fully composed of siliceous sponge spicules along the whole length of the test. Twelve specimens of '*Marsipella*' *cylindrica*, including the specimens figured by Brady (1884), are preserved in the Brady Collection in the NHM, London in Slide ZF1811. A lectotype, corresponding to the specimen figured by Brady (1884, pl. 24, fig. 21), was selected and illustrated by Kaminski *et al.* (2008). Specimens from Holes U1339A and U1340A have sponge spicules that are not so perfectly aligned as in the type specimens in the Brady Collection.

Genus *Dendrophyra* Wright, 1861

*Dendrophyra* sp.  
(Fig. 6:2)

**Remarks.** Small thin-walled fragments, displaying dichotomous or trichotomous branching.

Genus *Rhizammina* Brady, 1879

*Rhizammina algaeformis* Brady, 1879  
(Fig. 6:14)

1879 *Rhizammina algaeformis* Brady: 38, pl. 4, figs 16, 17.



1884 *Rhizammina algaeformis* Brady; Brady: 274, pl. 28, figs 1–11.  
1990 *Rhizammina algaeformis* Brady; Schröder-Adams *et al.*: 35, pl. 1, figs 6–7.

**Description.** Test small, round in cross-section, occasionally branching dichotomously. Test wall is thin and comprised mostly of fine sand grains with occasional short fragments of sponge spicules or diatom frustules loosely attached to the surface of the test.

**Remarks.** Unlike the type specimens housed at the NHM, London, the specimens from the Bering Sea do not attach any planktonic foraminifera to the exterior of their test. Instead, the species attaches occasional sponge spicules or centric diatoms to its test surface. Our specimens more closely resemble those figured by Schröder-Adams *et al.* (1990) from the Axel Heiberg Shelf, Arctic Ocean.

*Rhizammina* sp. 'straight large'  
(Fig. 6:15)

**Description.** Test small, round in cross-section, with the test wall comprised largely of biosiliceous fragments.

Genus *Lagenammina* Rhumbler, 1911

*Lagenammina* sp. 'spicules'  
(Fig. 7:1)

**Remarks.** Test flask-shaped, tapering toward the aperture. Wall consists largely of biogenic siliceous particles, with long sponge spicules extending radially away from the test. A single specimen was found in Hole U1340A.

*Lagenammina* sp.

**Remarks.** A specimen from Hole U1341A is comprised mostly of biogenic siliceous particles, including centric diatoms. The species of *Lagenammina* made of quartz grains (e.g. *L. atlantica* and *L. arenulata*), which are so common in the Arctic, have not been found in our material from the Bering Sea.

Genus *Psammosphaera* Schultze, 1875

*Psammosphaera fusca* Schultze, 1875  
(Fig. 7:2)

1875 *Psammosphaera fusca* Schultze: 113, pl. 2, fig. 8a–f.

**Description.** Test free or attached to a single large sand grain, varying in size, consisting of a single spherical chamber. Wall agglutinated, of a single layer of coarse sand grains, cemented together in a matrix of finer agglutinated particles, without any inner organic layer. Small pores between the loosely agglutinated sand grains serve as apertures.

**Remarks.** The species *P. fusca* uses a combination of larger and smaller agglutinated particles, sometimes selecting a single larger grain that may serve as an attachment surface. The large agglutinated grains may be angular or sub-rounded. Space between grains is filled in by a matrix of much finer agglutinated particles. The size of the recovered specimens is variable. Smaller specimens build their test out of only a few (<10) grains. Already in 1948, Cushman (1948, p. 9) noted 'All the Arctic specimens I have seen are small with rather large, angular quartz grains, and quite different from some of the specimens that are figured and referred to this species'. We observed this species at Sites U1339, U1340, U1343, and U1344.

Large agglutinated 'plate'

**Remarks.** In Hole U1345C, platy fragments of a larger agglutinated foraminifera were found. These fragments consist of equidimensional sand grains that include some mafic grains, cemented in a groundmass of much finer quartz grains. The grains are orientated in such a manner that the flat surfaces of the grains face the interior of the chamber. The agglutinated plates show little or no curvature, which means the whole specimen must have been quite large. These specimens are similar to the taxon that was named '*Astrorhizinella*' by Saidova (1970). Loeblich & Tappan (1987) included this taxon in their list of genera with uncertain status, with the remark that it could be a foraminifera, xenophyophorean or inorganic.

Subclass **Turbothalamana** Pawlowski, Holzmann & Tyszka, 2013

Genus *Saccorhiza* Eimer & Fickert, 1899

*Saccorhiza ramosa* (Brady, 1879)  
(Fig. 7:3)

1879 *Hyperammina ramosa* Brady: 33, pl. 3, figs 14, 15.

2011 *Saccorhiza ramosa* (Brady); Kaminski & Cetean: 62, pl. 1, figs 11–12 (lectotype).

**Description.** Test with smaller diameter than other tubular species. Proloculus or branching not observed. Wall predominantly of agglutinated quartz grains, with yellowish-brown organic cement, with a large component of outwardly projecting sponge spicules that give the surface a hirsute appearance.

**Remarks.** The specimens from Hole U1341A select sponge spicules that project out from the agglutinated tube at angles from 60 to 90°.

Genus *Archimerismus* Loeblich & Tappan, 1984

*Archimerismus subnodosus* (Brady, 1884)  
(not figured)

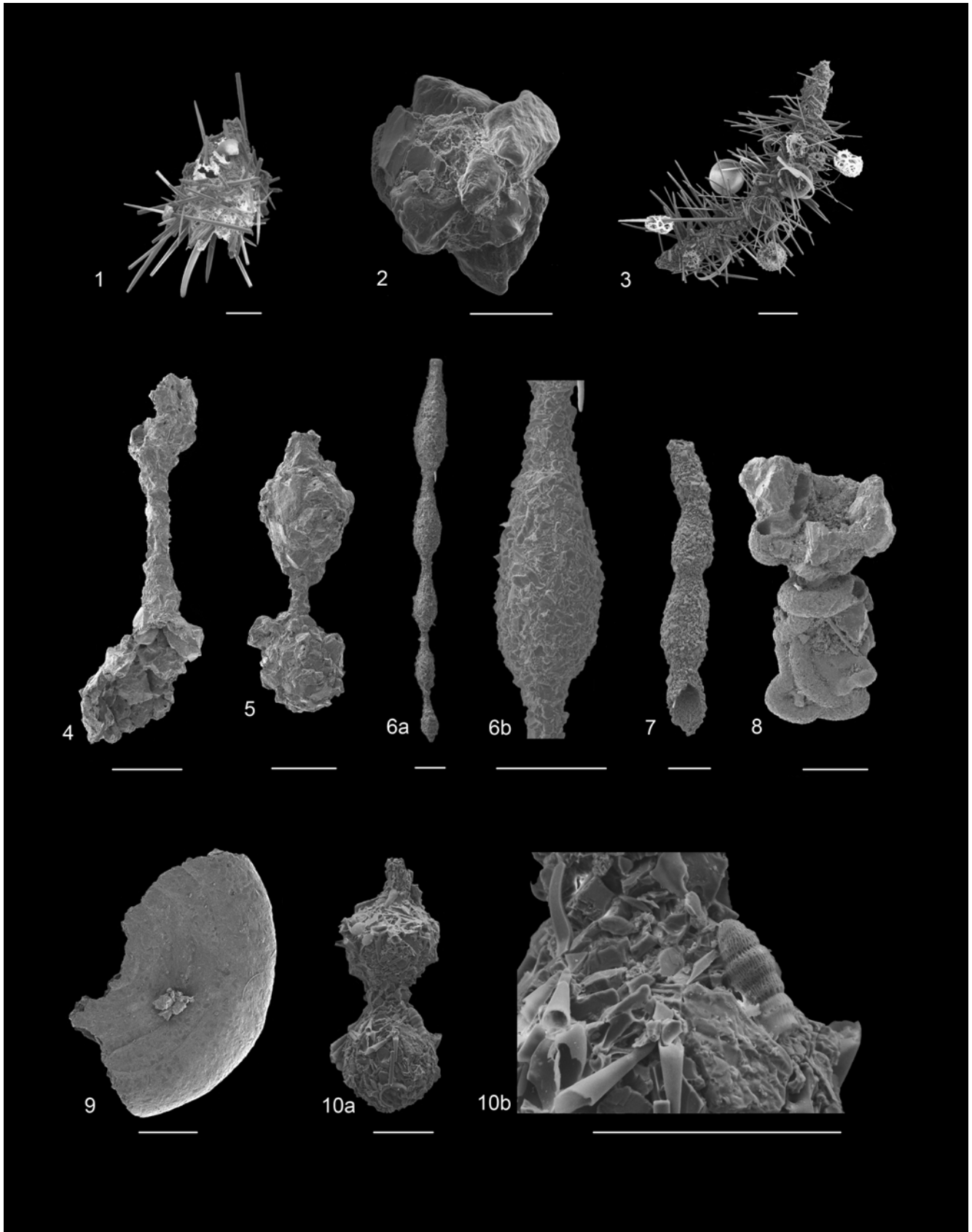
1884 *Hyperammina subnodosa* Brady: 259, pl. 23, figs 11–14.

1984 *Archimerismus subnodosus* (Brady); Loeblich & Tappan: 1161.

2011 *Archimerismus subnodosus* (Brady); Kaminski & Cetean: 63, pl. 1, figs 13–15 (lectotype).

**Description.** Test free, very large (>2 mm), tubular or cylindrical, open only at one end, proloculus with approximately the same diameter as the tubular chamber. The fragments of the tubular chamber are several mm in length and may display constrictions or taper slightly toward the aperture. Wall is several grains thick, of fine sand with occasional larger sand grains, occasional diatom frustules and dark mafic grains, with little cement, grey in colour. The test appears to be comprised of two layers: the interior layer is much thinner, with smaller sand grains lining the central cavity, which has a yellowish-brown inner organic lining. Aperture terminal, constricted, round in outline, may be partially obstructed by agglutinated particles.

**Remarks.** Three slides of type specimens of *Hyperammina subnodosa* are preserved in the Carpenter Collection in the NHM, London, all from Valorous Station 2, 100 fathoms. The lectotype, designated by Kaminski & Cetean (2011), is the specimen illustrated by Brady (1884) in plate 23, figure 11, and is preserved in slide BMNH 1886.4.16.94. These specimens are quite large (>1 mm) for an agglutinated foraminifera. The diameter of the tubular chamber is variable in our specimens, giving the impression of pseudochambers. These pseudochambers sometimes taper toward the aperture and may be several mm in length. However,



**Fig. 7.** (1) *Lagenammina* sp. 'spicules', Hole 1340A. (2) *Psammosphaera fusca* Schultze, 1875, Hole U1343E. (3) *Saccorhiza ramosa* (Brady, 1879), Hole U1341A. (4, 5) *Hormosinella distans* (Brady, 1881), Hole U1341A. (6a, b) *Reophanus oviculus* (Brady, 1879), Hole U1344. (7) *Subreophax splendidus* (Grzybowski, 1898), Hole U1341A. (8) *Tolypammina vagans* (Brady, 1879), Hole U1341A. (9) *Ammodiscus* sp., Hole U1341A. (10a, b) *Hormosinelloides* sp. aff. *H. guttifer* (Brady, 1884), Hole U1344. Scale bar 200  $\mu$ m.

the wall of the tubular chamber is continuous from one pseudochamber to the next; therefore, these are not true chambers. The interior layer contains smaller agglutinated grains than the outer layer, giving the interior a smooth appearance. This is the type species of the genus *Archimerismus* Loeblich & Tappan, 1984, which differs from *Hyperammina* in the partial subdivision of the test to form pseudochambers. The species is found at Sites U1344 and U1345. Gooday *et al.* (2005) noted mass occurrences of the species at deeper stations in the outer reaches of the West Spitsbergen fjords.

Genus *Hormosinella* Stschedrina, 1969

*Hormosinella distans* (Brady, 1881)  
(Fig. 7:4, 7:5)

1881 *Reophax distans* Brady: 50.  
1884 *Reophax distans* Brady; Brady: pl. 31, figs 18–22.  
2005 *Hormosinella distans* (Brady); Kaminski & Gradstein: 246, pl. 45, figs 1–11.  
2011 *Hormosinella distans* (Brady); Kaminski & Cetaan: 63, pl. 1, figs 16–17 (lectotype).

**Description.** Proloculus round, followed by ovoid pseudochambers connected by thin stolons. Wall thin. Chambers taper toward the aperture.

**Remarks.** The subspecies of *H. distans* were discussed by Kaminski & Gradstein (2005). The type specimens are from Challenger Station 300, (33° 42' S, 78° 18' W), north of Juan Fernández Island, South Pacific, 1375 fathoms. A lectotype was designated by Kaminski & Gradstein (2005) and is preserved in the Brady Collection in Slide BMNH ZF 2271. Cushman (1910) recorded the species from two Albatross stations in the North Pacific and from the Bering Sea at 1771 fathoms.

Fragmentary coarsely agglutinated specimens were recovered at Site U1341, while specimens with as many as five chambers were found at Site U1344.

Genus *Reophanus* Saidova, 1970

*Reophanus oviceulus* (Brady, 1879)  
(Fig. 7:6a–b)

1879 *Hormosina oviceula* Brady: 61, pl. 4, fig. 6.  
1884 *Hormosina oviceula* Brady; Brady: 327, pl. 39, figs 7–9.  
1987 *Reophanus oviceulus* (Brady); Loeblich & Tappan: 61, pl. 46, fig. 10.  
2011 *Reophanus oviceulus* (Brady); Kaminski & Cetaan: 63, pl. 2, figs 1–3.

**Description.** Test large, exceeding 4 mm in length, uniserial, rectilinear, unilocular in appearance because the elongated ovate chambers are separated by their respective necks, each new chamber attaching to the upper margin of the previous apertural lip, so that the test is fragile and tends to break into individual chambers. Wall finely agglutinated with several layers of very fine quartz grains, with yellowish-brownish cement which is more prominent on the necks of the chambers, without an inner organic lining. Aperture rounded, terminal on a distinct neck, with somewhat flared lip.

**Remarks.** The type specimens of *Hormosina oviceula* are preserved in the Brady Collection in the NHM, London. The lectotype is the specimen illustrated by Brady (1884) in plate 39, figure 7 (Kaminski & Cetaan 2011). This specimen was designated the 'holotype' by

Loeblich & Tappan (1987) and is preserved in slide BMNH ZF1588. The specimen is from Challenger Station 241, North Pacific at 2300 fathoms. It is the type species of the genus *Reophanus* Saidova, 1970.

This species is common in Hole U1341A and specimens consist of up to five chambers. The chambers (especially the proloculus) in our specimens are slightly more elongated than Brady's specimens from the North Pacific, but otherwise they conform very well to the original description of the species.

Genus *Subreophax* Saidova, 1975

*Subreophax splendidus* (Grzybowski, 1898)  
(Fig. 7:7)

1898 *Reophax splendida* Grzybowski: 278, pl. 10, fig. 16.  
1993 *Subreophax splendidus* (Grzybowski); Kaminski & Geroch: 251, pl. 3, figs 11a–12b.

**Description.** Test comprised of a meandering series of uniserial pseudochambers. Pseudochambers are oval, elongated in the direction of growth and are connected by wide stolons. Wall thin, medium to coarse. Aperture wide, terminal.

**Remarks.** This species was originally described from the Palaeogene of the Polish Carpathians (Grzybowski 1898). The lectotype was designated by Kaminski & Geroch (1993). It differs from *Subreophax aduncus* (Brady) in possessing a more coarsely agglutinated wall and chambers that are elongated in the direction of growth. A single specimen was found in Hole U1341A.

Genus *Ammodiscus* Reuss, 1862

*Ammodiscus* sp.  
(Fig. 7:9)

**Remarks.** A large coarsely agglutinated fragment of a specimen consisting of >6 planispiral whorls was found in Hole U1341A.

Genus *Tolypammina* Rhumbler, 1895

*Tolypammina vagans* (Brady, 1879)  
(Fig. 7:8)

1879 *Hyperammina vagans* Brady: 33, pl. 5, fig. 3.  
1884 *Hyperammina vagans* Brady; Brady: 260, pl. 34, figs 1–5.  
1921 *Tolypammina vagans* (Brady); Cushman: 55, pl. 4, figs 2, 3; pl. 7, figs 1, 2.

**Description.** Test attached, tubular, of constant diameter, streptospirally coiled, finely agglutinated with a thin wall.

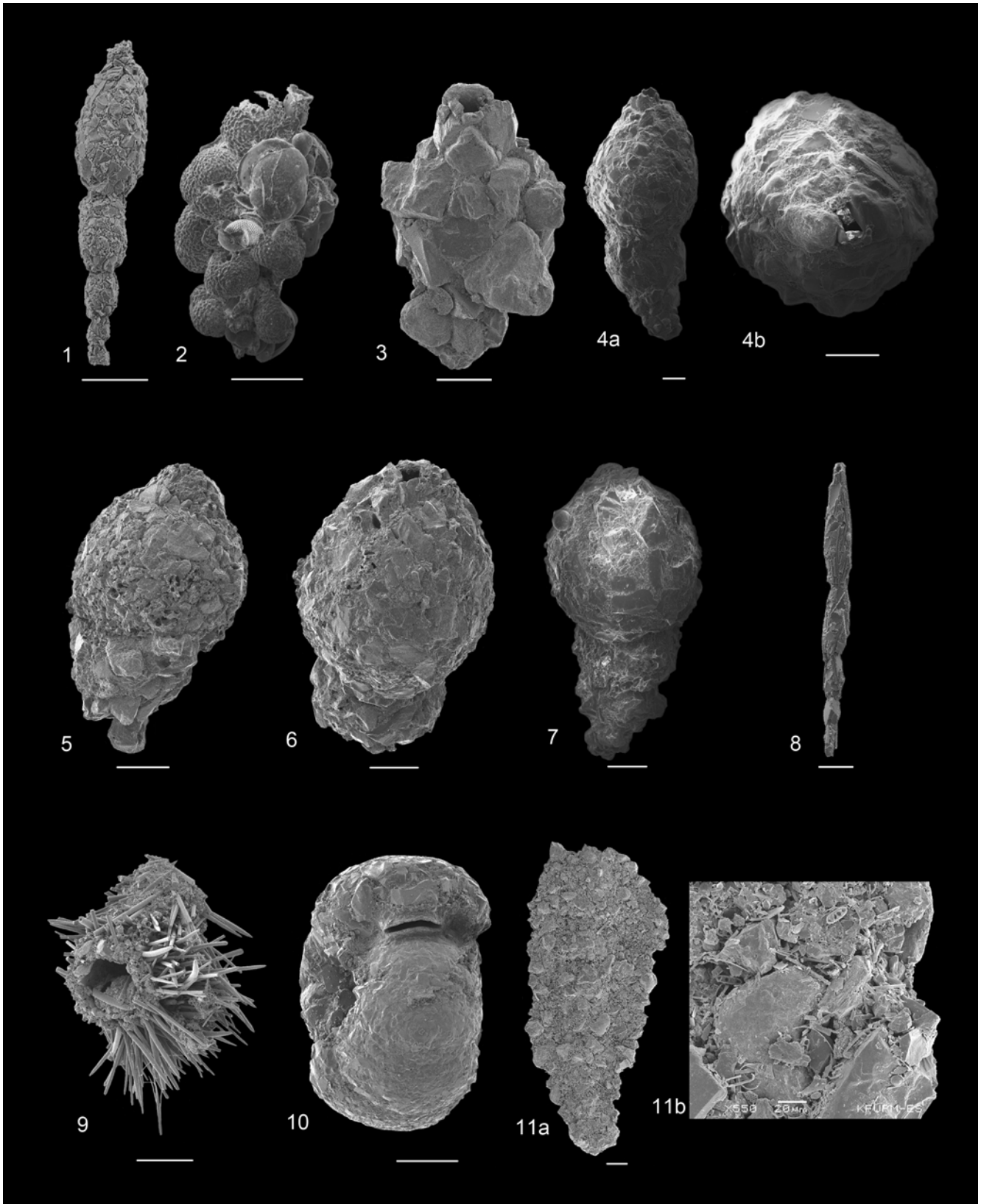
**Remarks.** Brady (1879) originally reported this species from the South Atlantic and from the North Pacific. Brady (1884) illustrated specimens that grew free or attached to shell fragments. Cushman (1921) reported that the species often attaches itself to other agglutinated foraminifera. Schröder (1986) reported that the species grows attached at bathyal depths in the North Atlantic, but is free-living at abyssal depths.

A large specimen was recovered in Hole U1341A. Our specimen was probably free-living and most closely resembles the specimens illustrated by Cushman (1921, pl. 4) from the Philippine Seas.

Subclass *Globothalamana* Pawlowski, Holzmann & Tyszka, 2013

Genus *Hormosinelloides* Zheng, in Zheng & Fu, 2001





**Fig. 8.** (1) *Nodulina dentaliniformis* (Brady, 1881), Hole U1341A. (2) *Reophax agglutinatus* Cushman, 1913, Hole U1342D. (3) *Reophax bilocularis* Flint, 1899, Hole U1342B. (4) *Reophax bradyi* Brönnimann & Whittaker, 1980, Hole U1343A. (5, 6) *Reophax excentricus* Cushman, 1910, Hole U1341A. (7) *Reophax pilulifer* Brady, 1884, Hole U1342A. (8) *Reophax spiculifer* Brady, 1879, Hole U1344D. (9) *Reophax* sp. 2 'tufty spicules', Hole U1341A. (10) *Veleroninoides scitulus* (Brady, 1881), Hole U1343A. (11) *Glaphyrammina* cf. *americana* (Cushman, 1910), Hole U1344B. Scale bar 200  $\mu$ m unless labelled otherwise.

*Hormosinelloides* sp. aff. *H. guttifer* (Brady, 1884)  
(Fig. 7:10a–b)

1910 *Reophax guttifer* (Brady); Cushman: 88, fig. 123.  
1969 *Reophax guttifer* (Brady); Vilks: 44, pl. 1, fig. 10.



**Description.** Test free, small, consisting of two or three pyriform chambers. Aperture terminal, with a distinct neck.

**Remarks.** The specimens from Site U1344 are consistently made up of only few chambers and the wall is made of mineral grains as well as fragments of biosiliceous particles, such as sponge spicules and radiolarians (Fig. 7:10b). The type specimens of *H. guttifer* from Challenger Station 323 (South Atlantic, east of Buenos Aires at 1900 fathoms) are larger and consist of 8 or more chambers (Kaminski & Cetaan 2011, pl. 2, figs 10–12). The chambers of the type specimens in the Brady Collection are slightly more pyriform than in our specimens, but they share the habit of incorporating small biogenic particles, such as radiolarians and sponge spicules, into their wall. Our specimens more closely resemble the Arctic variety of this species, which is smaller and has fewer chambers. The specimen illustrated by Vilks (1969) from Hecla Bay, Arctic Canada, is only a two-chambered individual.

Genus *Nodulina* Rhumbler, 1895

*Nodulina dentaliniformis* (Brady, 1881)  
(Fig. 8:1)

1881 *Reophax dentaliniformis* Brady: 49.  
1884 *Reophax dentaliniformis* Brady; Brady: 293, pl. 30, figs 21–22.  
1980 *Hormosina dentaliniformis* (Brady); Brönnimann & Whittaker: 265, figs 8–11.  
2011 *Nodulina dentaliniformis* (Brady); Kaminski & Cetaan: 65, pl. 2, figs 19–22.

**Description.** Test uniserial and rectilinear, similar to *Reophax* but with a straighter axis, more symmetrical, regular, with gradually enlarging chambers and nearly horizontal sutures. Wall coarsely agglutinated of a single layer of quartz grains. Aperture rounded, at the end of a short tubular neck.

**Remarks.** Brönnimann & Whittaker (1980) selected a lectotype for the type species from the Brady Collection. The type locality is Challenger Station 300, north of Juan Fernández (1375 fathoms). This six-chambered specimen (labelled *Hormosina dentaliniformis*) is housed in slide BMNH 3990. Anderson (1963) listed the species in a single sample from the inner Bering Sea shelf. Our specimens from Hole U1341A possess up to five chambers and closely resemble those in the Brady Collection.

Genus *Reophax* de Montfort, 1808

*Reophax agglutinatus* Cushman, 1913  
(Fig. 8:2)

1913 *Reophax agglutinatus* Cushman: 637, pl. 79, fig. 6.  
1921 *Reophax agglutinatus* Cushman; Cushman: 73, pl. 14, figs 2a, b.  
1939 *Reophax agglutinatus* Cushman; Cushman & McCulloch: 59, pl. 3, figs 1–3.

**Description.** Test free, uniserial, comprised of two chambers, with the second one much larger than the first. Wall very coarsely agglutinated, comprised of quartz grains with an admixture of small planktonic and benthic foraminiferal tests and siliceous sponge spicules that extend outward from the test. Aperture at the end of a tapering neck.

**Remarks.** Differs from *Reophax bilocularis* in possessing small planktonic foraminifera incorporated into its wall and its more robust neck. Anderson (1963) listed the species in a single sample

from the outer Bering Sea shelf. The large specimen from Hole U1342D is the typical form.

*Reophax bilocularis* Flint, 1899  
(Fig. 8:3)

1899 *Reophax bilocularis* Flint: 273, p. 17, fig. 2.  
1920 *Reophax bilocularis* Flint; Cushman: 10, pl. 3, figs 3, 4.

**Description.** Test free, uniserial, comprised of two chambers, with the second one much larger than the first. Wall very coarsely agglutinated, comprised of angular and rounded quartz grains with an admixture of some dark minerals, cemented with a matrix of much finer agglutinated grains. Aperture on a produced neck that consists of much finer agglutinated grains than the chamber wall.

**Remarks.** Several good specimens were found in Holes U1343A and U1344B. In the North Atlantic, *Reophax bilocularis* has a habit of picking up small planktonic foraminiferal tests in addition to mineral grains. This feature is not observed in the specimens from the Bering Sea.

*Reophax bradyi* Brönnimann & Whittaker, 1980  
(Fig. 8:4a–b)

1980 *Reophax bradyi* Brönnimann & Whittaker: 264, figs 13–16.  
1994 *Reophax bradyi* Brönnimann & Whittaker; Jones: 31, pl. 18, fig. 16; p. 37, pl. 30, fig. 12.

**Description.** Test free, uniserial, arched or slightly meandering, consisting of up to 5 chambers, increasing in size gradually. Chambers are round or slightly elongated. Wall very coarsely agglutinated, aperture terminal, without a neck.

**Remarks.** Schröder-Adams *et al.* (1990) illustrated specimens as *R. scorpiurus* from the Canadian Arctic that likely belong in this species.

*Reophax excentricus* Cushman, 1910  
(Fig. 8:5, 8:6)

1910 *Reophax excentricus* Cushman: 92, fig. 134.  
1939 *Reophax excentricus* Cushman; Cushman & McCulloch: 60, pl. 3, figs 4–9.

**Description.** Test fusiform, arched, with three chambers increasing rapidly in size. Wall coarsely agglutinated, of quartz and mafic grains. Aperture on a short neck, eccentrically placed on the last chamber.

**Remarks.** *Reophax excentricus* Cushman has its type locality from the stomachs of holothurians dredged at Albatross station D3603 in the Bering Sea at 1773 fathoms (Cushman 1910). This species is relatively common in the Pacific Ocean – it has also been reported off Oregon and the Alaskan Peninsula (Culver & Buzas 1985) and characterizes depths of 200–2000 m off the Pacific coast of Central America (Culver & Buzas 1987). It is regarded to be an opportunistic species (Kaminski *et al.* 1988).

*Reophax pilulifer* Brady, 1884  
(Fig. 8:7)

1884 *Reophax pilulifer* Brady: 292, pl. 30, figs 18–20.  
2005 *Reophax pilulifer* Brady; Kaminski & Gradstein: 272, pl. 53, figs 1–9 (fig. 1 is the lectotype).

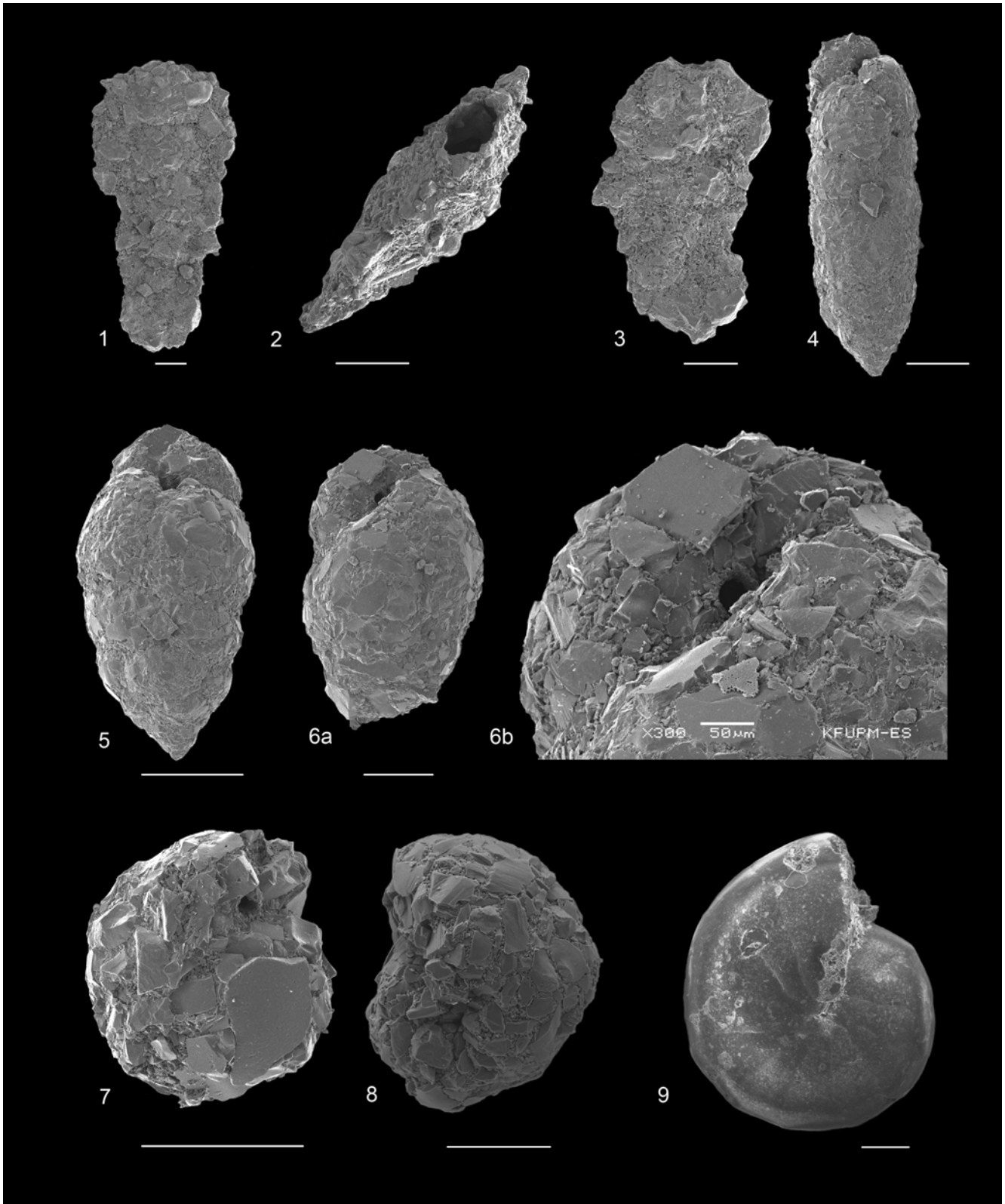


Fig. 9. (1–3) *Glaphyrammina* cf. *americana* (Cushman, 1910), Hole U1344B. (4–6b) *Eggerelloides* sp. 1, Hole U1345C. (7, 8) *Recurvoides* sp.: 7, Hole U1343E; 8, Hole U1345B. (9) *Cyclammina compressa* Cushman, 1917, Hole U1342A. Scale bar 200 μm unless labelled otherwise.

**Description.** Test robust, straight or curved, with 3 to 5 rapidly enlarging chambers. Chambers are globular and only slightly envelop preceding chambers. Wall coarse, comprised of a single layer of large sand grains in a matrix of finer grains, with organic cement. Aperture a round opening, situated on a low apertural shoulder, but without a neck.

**Remarks.** Kaminski & Gradstein (2005) designated a lectotype from the Carpenter Collection in the NHM, London. The type locality is Porcupine Station 31 in the North Atlantic, 1360 fathoms water depth. Cushman (1921) recorded it from 13 Albatross stations in the Philippine Sea between 208 and 1560 fathoms, but noted that these specimens are smaller than Atlantic specimens and ‘are not



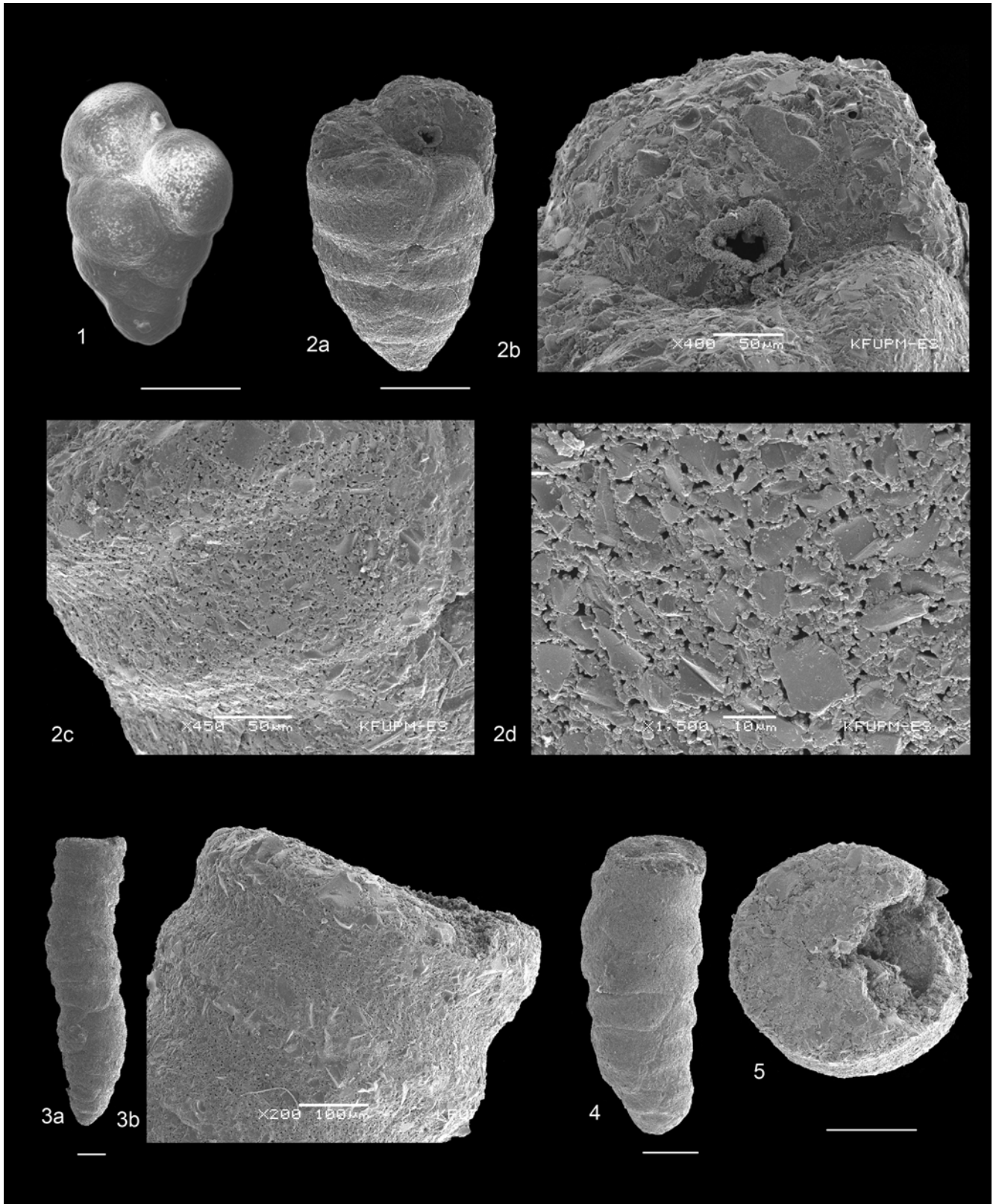


Fig. 10. (1) *Karreriella* sp. 1, Hole U1341B. (2a–5) *Martinottiella* sp. 3, Hole U1342A. Scale bar 200 µm unless labelled otherwise.

typical'. Specimens from the Bering Sea are smaller than the types, and are coarsely agglutinated. Large concentrations of *R. pilulifer* were observed in the St Anne Trough, Kara Sea, Siberian Arctic (Korsun *et al.* 1988).

*Reophax scorpiurus* de Montfort, 1808  
(not figured)

1808 *Reophax scorpiurus* de Montfort: 331, text-fig. 130.  
1980 *Reophax scorpiurus* de Montfort; Brönnimann & Whittaker: 261, figs 1 – 7, 12, 17 (figs 2, 5 show the neotype).

**Remarks.** Smaller than most other species, it displays a characteristic arched test and elongated final chamber, with a

produced aperture. Brönnimann & Whittaker (1980) designated a neotype for *R. scorpiurus* from the SE Adriatic. The neotype and paraneotypes are from a shallow-water sample collected by H. Sidebottom off Corfu, and are preserved in the collections of the NHM, London in slide ZF3985.

*Reophax spiculifer* Brady, 1879  
(Fig. 8:8)

1879 *Reophax spiculifera* Brady: 54, pl. 4, figs 10–11.  
1994 *Reophax spiculifer* Brady; Jones: 38, pl. 31, figs 16–17.

**Description.** Test slender, uniserial and rectilinear. Chambers are strongly elongated and taper toward the aperture. Wall comprised of quartz grains and fragments of sponge spicules that are aligned sub-parallel to the long axis of the test. Spicules do not protrude from the chambers. Aperture terminal, round, without a neck.

**Remarks.** In his description of the species, Brady (1879, p. 55) remarked ‘This is one of the many species of Foraminifera that give evidence of considerable selective power in respect to the material employed for the construction of their tests.’ Specimens from Hole U1344D display a greater admixture of mineral grains in the test wall, but are otherwise similar to Brady’s specimens.

*Reophax* sp. 2 ‘tufty spicules’  
(Fig. 8:9)

**Remarks.** Broken fragments of a *Reophax* species that agglutinates numerous sponge spicules more or less normal to the test surface were found in Hole U1341A. The morphology of the whole specimen is unknown.

Genus *Evolutinella* Mjatluk, 1971

*Evolutinella rotulata* (Brady, 1881)  
(not figured)

1881 *Haplophragmium rotulatum* Brady: 50.  
1994 *Evolutinella rotulata* (Brady); Jones: 40, pl. 34, figs 5–6.

**Remarks.** A single specimen was found in Hole U1340A.

Genus *Veleroninoides* Saidova, 1981

*Veleroninoides scitulus* (Brady, 1881)  
(Fig. 8:10)

1881 *Haplophragmium scitulum* Brady: 50.  
1884 *Haplophragmium scitulum* Brady; Brady: 308, pl. 34, figs 11–13.  
1994 *Veleroninoides scitulus* (Brady); Jones: 41, pl. 34, figs 11–13.

**Description.** Test planispiral, biumbilicate, evolute with deep umbilici, with numerous low chambers in about two to three whorls, increasing very little in size as added. Periphery broadly rounded, margins slightly lobulate, with chambers wider than high when viewed from the periphery. Later sutures slightly depressed, nearly radial. Wall medium–finely agglutinated, surface smoothly finished. Aperture interio-areal, a low oval to elliptical opening near the base of the apertural face, bordered by a thin lip.

**Remarks.** The type specimens of *V. scitulus* in the Brady Collection are from ‘Knight Errant’ Station 7 at 530 fathoms in the Faroe

Channel. Our specimens from the Bering Sea are smaller and more coarsely agglutinated than the type specimens.

Genus *Glaphyrammina* Loeblich & Tappan, 1984

*Glaphyrammina* cf. *americana* (Cushman, 1910)  
(Figs 8:11a, b and 9:1–9:3)

1910 *Ammobaculites americanus* Cushman: 117, text-figs 184–185.  
1994 *Glaphyrammina americana* (Cushman); Jones: 40, pl. 34, figs 1–4.

**Description.** Test free, broad and flattened, thin and fragile. Early portion planispirally enrolled and partially to completely evolute, sutures poorly visible, later portion uncoiled and may have a few rectilinear chambers with horizontal depressed sutures. Wall coarsely agglutinated, with larger grains in a groundmass of smaller grains. Larger pointed quartz grains project out from the periphery in the coiled portion of the test. Aperture a terminal elongate narrow opening, extending across nearly the breadth of the chamber, not produced.

**Remarks.** Loeblich & Tappan (1984) remarked that sutures in *Glaphyrammina* may simply represent external indication of periodic growth, as the interior is hollow and undivided by septa, possibly with organic partitions. The holotype of *Ammobaculites americanus* Cushman, 1910 is from Albatross Station 3419, collected at 772 fathoms off the west coast of Mexico. Our specimens differ from Cushman’s types in possessing a well-developed uniserial portion and a very coarsely agglutinated test with the odd larger mafic grain embedded in the wall, making it difficult to observe the sutures. This species is common in Hole U1344B.

Genus *Cribrostomoides* Cushman, 1910

*Cribrostomoides subglobosus* (Cushman, 1910)  
(not figured)

1910 *Haplophragmoides subglobosum* (M. Sars); Cushman: 105, text-figs 162–164.  
1994 *Cribrostomoides subglobosus* (Cushman); Jones: 40, pl. 34, figs 11–13.  
2005 *Cribrostomoides subglobosus* (Cushman); Kaminski & Gradstein: 392, text-fig 92 (lectotype), pl. 92, figs 1a–3b.

**Remarks.** The correct citation of the authorship of this species was discussed by Kaminski & Gradstein (2005), who provided a detailed description. A rather coarsely agglutinated variety of this species was observed in Hole U1343C.

Genus *Recurvoides* Earland, 1934

*Recurvoides* sp.  
(Fig. 9:7, 9:8)

**Remarks.** A small, almost spherical, coarsely agglutinated species of *Recurvoides* with a round to oval aperture in the lower part of the apertural face.

Genus *Eggerelloides* Haynes, 1973

*Eggerelloides* sp. 1  
(Fig. 9:4–9:6b)

**Description.** Test elongate, fusiform, initially coiled in a high trochospire of about 5 whorls with four chambers per whorl, then



reducing to three chambers per whorl. Chambers are somewhat inflated, sutures depressed. Wall is made of fine quartz with occasional larger grains, with orange-brown organic cement. The initial part of the test has a more intense colour owing to the abundance of organic cement. Aperture a high interiomarginal arch, umbilical in position, surrounded by a thin lip.

**Remarks.** We place this species into *Eggerelloides* based on its high, loop-shaped aperture, though its internal structure (or toothplate) was not observed. *Rhumlerella* differs in possessing a low interiomarginal aperture. The North Pacific form of *Eggerelloides advena* (Cushman), as depicted by Tsujimoto *et al.* (2006) and by Vázquez Riveiros & Patterson (2007), differs in possessing a slender test and more well-developed triserial part. In our samples, the species is most abundant at Site U1345.

Genus *Cyclammina* Brady, 1879

*Cyclammina compressa* Cushman, 1917  
(Fig. 9:9)

1917 *Cyclammina compressa* Cushman: 653.

1921 *Cyclammina compressa* Cushman; Cushman: 85, pl. 16, figs 2a, b.

**Description.** Test small for the genus, coiling planispiral, involute, with 12–13 chambers in the final whorl. Sutures radial, nearly straight, depressed slightly. Umbilicus flush or only depressed slightly. Periphery subacute. Wall consisting of an inner alveolar layer and a much thinner imperforate epidermal (outermost) layer, finely agglutinated with a smooth finish. Alveoles are unidimensional and equally spaced on the outer chamber wall. Aperture a broad interiomarginal slit with a thin upper lip of finely agglutinated particles. Areal supplementary apertures are not observed and, if present, are only small openings between the larger agglutinated grains in the apertural face.

**Remarks.** Cushman (1917, 1921) differentiated *Cyclammina compressa* from *C. cancellata* Brady by its smaller size, open and depressed umbilicus, evolute coiling and acute periphery. The syntypes preserved in the Cushman Collection (CC420) are from Albatross Station 5470, east of Luzon, at 540 fathoms. The specimens are over 3 mm in diameter, and have an average of 14 chambers in the final whorl. Banner (1970) regarded this species to be conspecific with *C. cancellata* Brady based on the structure of the hypodermis and alveoles. Kaminski & Gradstein (2005) accepted Banner's synonymy of the two species.

In the Bering Sea, typical *Cyclammina cancellata* has not been found – the specimens recovered in Hole U1343A are best ascribed to *C. compressa*, which differs from typical *Cyclammina cancellata* in its smaller dimensions, more compressed lateral sides, straighter sutures and in lacking distinct supplementary areal apertures. The Bering Sea specimens are even smaller than Cushman's specimens from the Philippines, and do not have any visible areal supplementary apertures, whereas these are visible in Cushman's (1921) illustrations. Additionally, the umbilicus is not as depressed as Cushman depicted it. Cushman (1921, p. 86) remarked: 'This species seems to be in deeper and colder water than the preceding [*C. cancellata*]'.

Genus *Karreriella* Cushman, 1933

*Karreriella* sp. 1  
(Fig. 10:1)

2013 *Karreriella* sp. 1; Kaminski *et al.*: 339, fig. 3a–f.

**Description.** Test free, elongate, initially trochospiral with up to five chambers per whorl, later reduced to twisted triserial and finally becoming biserial in just the final one or two pairs of chambers. Chambers in the terminal biserial part are globular, with depressed sutures. Wall finely agglutinated, with a smooth outer surface. Aperture areal, a rounded opening slightly above the base of the apertural face in the triserial stage, becoming more areal and increasingly oval in the biserial adult stage, produced and surrounded by a distinct lip.

**Remarks.** A single specimen was found in Hole U1341B. It most closely resembles the species described as '*Karreriella* sp. 1' by Kaminski *et al.* (2013) from the Pliocene of Hole U1341B.

Genus *Martinottiella* Cushman, 1933

*Martinottiella* sp. 3  
(Fig. 10:2–10:5)

**Description.** Test elongate, cylindrical, initial 6–7 whorls coiled in a high trochospire with four chambers per whorl, later reduced to uniserial. Wall finely agglutinated, canaliculate with pores open to the exterior. Aperture areal in the coiled stage, terminal and central in the uniserial stage, on a short tubular neck in the centre of the flattened terminal face.

**Remarks.** This species is common in the samples from Site U1342. It differs from the type species *M. communis* (d'Orbigny) in possessing open pores on the external chamber walls and a round (rather than oval or slit-like) aperture. The open pores are concentrated in the middle of the chamber wall – fewer pores are observed along the sutures (Fig. 10:3b). The apertural face in the adult stage is flat (Fig. 10:5).

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