ABSTRACT

Middle Pliocene to Recent sediments of Site 533, Leg 76, contain a benthic foraminiferal fauna characteristic of the bathyal zone. Few abyssal species are present. Reworked forms of the shelf and the upper slope are rare. The Matuyama/Brühnes boundary occurs above Sample 533-14.CC. Sample 533-1-2, 18-20 cm corresponds to the last glacial stage. Similar faunas—and probably analogous environmental conditions—are present in some earlier levels (Pleistocene and upper Pliocene). The Holocene fauna is characterized by a low specific diversity and the abundance of Globocassidulina moluccensis, which is not present before.

INTRODUCTION, PROCEDURES

Forty samples ranging in age from middle Pliocene to Recent were examined from Site 533 (see Fig. 1 for location). Three stratigraphic intervals were studied in more detail, the number of samples studied by core varying from one to three. In addition, eleven core catchers, located between the first and the second interval, were also examined in order to obtain an estimation of the Pleistocene faunal variation.

The list of the samples studied is given in Table 1. The Pliocene/Pleistocene boundary (FAD—first appearance datum—of Globorotalia truncatulinoides) is located between Sections 1 and 2 of Core 38. A more precise zonation established from planktonic species (M. Moullade, this volume) is indicated in Figure 2: the sediments studied belong to stratigraphical zones PL3 (test interval 3), PL5, PL6 and the base of N22 (test interval 2), and N22/23 (test interval 1). The correspondence between Pleistocene samples and Ericson's climatic zones is also indicated in Figure 2.

The number of foraminifers present in each sample varies considerably and is generally low. As far as possible, I have used tests larger than 160 µm. In two samples, however, the sediment was sieved with a 200 µm mesh and the finer fraction was lost. On the other hand, in some core catchers, the size fraction between 160 and 63 µm was retained. This residue included many tiny forms—mainly Bolivina—which did not appear in the coarser sediments.

For these reasons, the results obtained from the different samples are not exactly comparable. Thus in Figure 2 the frequencies indicated are only approximations. Similarly, when the number of specimens in a sample was too low, the species present were noted by “+” without any frequency estimation. This is the case for Hole 533A samples.

FAUNAL ANALYSIS

Abundance

The number of benthic foraminifers larger than 160 µm is very different from one sample to another, ranging from 3 to 203, as indicated in Figure 2.

The calculated number of specimens per gram of sediment ranges from two to ten in test interval 1; it is somewhat higher during early Pleistocene and late Pliocene, with a maximum of 22.8 in Sample 533-38-X, 17–19 cm; in all the other samples but one, it is less than 2.6 (Section 533-38-X is located between 533-37,CC and 533-38-1, 149.5–154 m sub-bottom.)

There is no trace of test solution, as might be expected from the paleo-water depth.

Specific Composition

About 80 species were encountered; 73 of them were determined at the specific level and listed in the Appendix with taxonomic references and stratigraphical and ecological indications. The different species of some genera (Bolivina, Fissurina, Lagena, Lenticulina, and Oolina), occurring sporadically and with few specimens, were not separated. The number of species per sample varies from 2 to 42.

Species Distribution

Figure 2 shows the occurrence of the main species with, when possible, an estimation of their frequencies. They are listed in the order of their vertical distribution, with the exception of the presumed reworked forms.

Some species occurring sporadically are not indicated in the Figure, such as Ammoscalaria tenuimargo (533-37-2, 25–27 cm), Cornuspira involvens (533-21,CC), Dentalina inornata bradyensis (533A-29,CC), D. submaciata (533-38-X, 17–19 cm), Epistominella exigua (533-38-2, 15–17 cm), Globocassidulina subglobosa (533A-29-3, 40–42 cm), G. crassa (533-37-1, 47–49 cm), Laticarinina pauperata (533-40,CC), Nuttalites rugosus-convexus (533-38-X, 17–19 cm), Osangularia cultur (533-37-1, 47–49 cm), and W. lilacina (533-38-1, 149.5–154 m sub-bottom.).

1, 47-49 cm), Stilostomella abyssorum (533-39-3, 33-35 cm), Stilostomella bradyi (533-6, CC).

The possible interpretations of the species distribution are discussed later. From bottom to top of the sequence, we can note the following facts:

Some species are limited to the Pliocene: Dentalina intorta, Nodosaria ovicula, Pseudonodosaria laevigata, and Dorothia sp.

Several others occur in the Pliocene and in a part of the Pleistocene: different species of Lenticulina, and Ellipsoplymorphismina brevis, Ellipsoidina ellipsoides, Orthomorphina fistuca, Stilostomella ex gr. lepidula, Plectofrondicularia advena.

The family Miliolidae is present throughout the section, but the genus Pyrgo is frequent, especially during the Pleistocene.

Many species show a large stratigraphic distribution. They are generally abundant, and some of them become dominant in certain samples, forming 20 to 50% of the assemblage, such as: Oridorsalis umbonatus, Cibicides wuellestorfi, the species of the genus Melonis, Uvigerina, Globobulimina, and Pullenia.

Few species are restricted to the Pleistocene: Cassidulina carinata, Chilostomella oolina, Francesita advena, Martinotiella communis, Siphotextularia catenata, Stainforthia complanata, Valvulineria laevigata.
Table 1. List of samples studied.

<table>
<thead>
<tr>
<th>Sample (interval in cm)</th>
<th>Age</th>
</tr>
</thead>
<tbody>
<tr>
<td>Test interval 1</td>
<td></td>
</tr>
<tr>
<td>533-1-1, 2.5-3.5</td>
<td>Recent and</td>
</tr>
<tr>
<td>533-1-1, 18-20</td>
<td>uppermost</td>
</tr>
<tr>
<td>533-2-2, 25-27</td>
<td>Quaternary</td>
</tr>
<tr>
<td>Intermediate sampling</td>
<td></td>
</tr>
<tr>
<td>533-6.CC</td>
<td>Quaternary</td>
</tr>
<tr>
<td>533-7.CC</td>
<td></td>
</tr>
<tr>
<td>533-9.CC</td>
<td></td>
</tr>
<tr>
<td>533-14.CC</td>
<td></td>
</tr>
<tr>
<td>533-20.CC</td>
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</tr>
<tr>
<td>533-21.CC</td>
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<td>533-23.CC</td>
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<td>533-25.CC</td>
<td></td>
</tr>
<tr>
<td>533-26.CC</td>
<td></td>
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<tr>
<td>533-33.CC</td>
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</tr>
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<td>533-36.CC</td>
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<td>Test interval 2</td>
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<tr>
<td>533-37-1, 47-49</td>
<td>lower</td>
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<tr>
<td>533-37-2, 25-27</td>
<td>Quaternary</td>
</tr>
<tr>
<td>533-38-X, 17-19</td>
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</tr>
<tr>
<td>533-38-1, 75-77</td>
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<td>533-38-2, 15-17</td>
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<td>533-38-2, 106-108</td>
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<td>533-38-3, 25-27</td>
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<tr>
<td>533-39-1, 48-50</td>
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</tr>
<tr>
<td>533-39-3, 33-35</td>
<td>upper</td>
</tr>
<tr>
<td>533-40-1, 38-40</td>
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<tr>
<td>533-40-2, 36-38</td>
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</tr>
<tr>
<td>533-40.CC</td>
<td></td>
</tr>
<tr>
<td>533-41-1, 47-49</td>
<td></td>
</tr>
<tr>
<td>533-41-2, 42-44</td>
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<tr>
<td>533-41-3, 27-29</td>
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</tr>
<tr>
<td>Test interval 3</td>
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</tr>
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<td>533A-26.CC</td>
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</tr>
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<td>533A-27-2, 88-90</td>
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</tr>
<tr>
<td>533A-27-4, 120-122</td>
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</tr>
<tr>
<td>533A-27-6, 100-102</td>
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<tr>
<td>533A-27.CC</td>
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<td>533A-28-2, 34-36</td>
<td>Pliocene</td>
</tr>
<tr>
<td>533A-28.CC</td>
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<tr>
<td>533A-29-2, 88-90</td>
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<tr>
<td>533A-29-3, 40-42</td>
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</tr>
<tr>
<td>533A-29-4, 48-50</td>
<td></td>
</tr>
<tr>
<td>533A-29.CC</td>
<td></td>
</tr>
</tbody>
</table>

Three species occur only in the uppermost stratigraphic level (533-1-1, 2.5–3.5 cm): Globocassidulina moluccensis, Karreriella apicularis, Rhabdammina linearis.

**DISCUSSION**

**Vertical Distribution and Stratigraphic Implications**

The majority of the species encountered in the two holes have a large time range and cannot be used for stratigraphic indexing. In most cases, especially when the species are rare, their absence may be due to the small size of the samples. It is also difficult to know whether the occurrences of the observed species depend on stratigraphic, ecologic, or other features.

An important fact that must be emphasized is the disappearance in the late Pleistocene of a group of species—Ellipsoidina ellipsoideis, Orthomorpha fistucula, Ellipsoidomorphina brevis, Plectofondicularia advena, Stilostomella ex gr. lepidula. The first of these forms is described from the Miocene, but there are few indications about its level of extinction. The others are known to be extinct at the beginning of the Brunhes epoch (Lutze, 1979). In Hole 533, these forms disappear between Samples 533-37-2, 25-27 cm and 533-14,CC. Three species only occur in the upper levels (test interval 1). This occurrence is difficult to explain: all three are known to be present in recent sediments from different seas in bathyal or abyssal zones. Their restricted occurrence is probably not a stratigraphic fact but may be a response to local ecological conditions.

**Bathymetric Interpretation of the Fauna**

Several recent publications deal with the bathymetric zonation of benthic foraminifers and give a review of older works, for example, for the Atlantic Ocean, Pheger et al. (1953), Pujo-Lamy (1973), Berggren and Haq (1976), Pflum and Frerichs (1976), Lutze (1978), Schnitter (1979), and so on. Results from the Pacific Ocean (Ingle, 1980; Ingle et al., 1980; Boltovskoy, 1981) and the Indian Ocean (Corliss, 1979) were also compiled. Although there are some differences in the detail of the species distribution in different areas, most of the forms found in Holes 533 and 533A have a similar depth range. The majority of them have their upper depth limit below 1000 m (middle bathyal), and most become more abundant in lower bathyal zones, even in abyssal zones (cf., Appendix), for example, Cibicides kullenbergi, Eggerella bradyi, Epistomestumidulus, Epistominella exigua, Francesia advena, Gyroidina soldanii, Karreriella apicularis, Melonis pompilioides, Oridorsalis umbonatus, Quinqueloculina venusta, Uvigerina canariense, U. senitiosa.

Thus the assemblage composition seems to be in agreement with the present water depth of the drilling site.

**The Displaced Faunas**

In the fraction coarser than 200 µm, only two species can be considered as displaced. They are Cibicides lotalatus (533-33,CC) and Textularia sp. (533-36,CC).

In the residue between 200 and 160 µm, the reworked species are a little more abundant, but the number of such specimens in each sample never exceeds two or three. The concerned forms are Ammonia beccarii, Asstronoion sp., Buliminella elegantissima, Elphidium sp., Neoconorbina sp., Nonionella cf. stella, and Rosalina sp.

All these forms are living today in the neritic zone of the different seas; owing to their low frequencies, the displacements of sediments from the shelf seem to have been of little importance during the span of time investigated.

The fraction between 160 and 63 µm contains many specimens of Bolivina belonging to three species: B. seminuda, B. ex gr. spathulata and Bolivina sp. The two former species have a large bathymetric distribution and
<table>
<thead>
<tr>
<th>Sub-bottom depth (m)</th>
<th>Hole 533</th>
<th>Hole 533A</th>
</tr>
</thead>
<tbody>
<tr>
<td>0.025</td>
<td>27</td>
<td>147</td>
</tr>
<tr>
<td>1.1-2.5</td>
<td>166</td>
<td>369</td>
</tr>
<tr>
<td>2.6-3.5</td>
<td>379</td>
<td></td>
</tr>
</tbody>
</table>

**Zonation from planktonic species (M. Moullade, this volume)**

**Ericson's climatic zones**

**Sample (core-section, interval in cm)**

**Benthic foraminifers**

<table>
<thead>
<tr>
<th>Species</th>
<th>no. of benthic (160 µm/g sediment)</th>
<th>no. of benthic counted</th>
<th>no. of species</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cassidulina molecule</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Karreriella apiculans</td>
<td>8.07</td>
<td>19</td>
<td>2</td>
</tr>
<tr>
<td>Rhabadammina linearis</td>
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<td>22</td>
<td>2</td>
</tr>
<tr>
<td>Valvulineria laevigata</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Cassidulina laevigata</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Chilostomella oolina</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Francesco advena</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Martinotiella communis</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Siphonotonotaria celerata</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Pseudestomella spp.</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Alabarnia decorata</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Epiderella bradyi</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Karreriella bradyi</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Epionides pusillus</td>
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</tr>
<tr>
<td>Epionides polius</td>
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<td>2</td>
</tr>
<tr>
<td>Sphaeroidina bulboides</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>g. Frasirna</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>g. Okina</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>g. Lagenia</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Marginulina obesa</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Nodosaria obliquata</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Lemniculina spp.</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Globobulimina affinis</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Globobulimina pyrula</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Uvigerina semicostosa</td>
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<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Uvigerina bellissima</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Uvigerina ex gr. papengine</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Uvigerina aquilana</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Uvigerina canariense</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Melonis pompiloides</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Melonis barleeanum</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Melonis formosum</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
<tr>
<td>Cibicides weillistorf</td>
<td>6.07</td>
<td>17</td>
<td>2</td>
</tr>
</tbody>
</table>
Figure 2. Distribution of the main benthic foraminifers in selected samples from Holes 533 and 533A.
are sensitive to other ecological factors than depth, especially $O_2$ contents (Boltovskoy and Wright, 1976). As they are present in all the sediments in which the finest fraction was preserved, we can assume that they are present throughout the core and they cannot yield any information on possible paleoecological changes.

**Other Factors**

In several levels, some taxa become dominant in the assemblage. As these species have generally the same bathymetric distribution, the causes of their abundance must be searched in the variation of other factors than depth. These forms are *Globobulimina*, *Melonis*, *Pullenia*, *Uvigerina*, and also *Cibicides wuellerstorfi* and *Oridorsalis umbonatus*.

Within the bathyal and abyssal zones, several authors have emphasized the fact that these species are more or less associated with the properties of the different bottom-water masses, mainly temperature, salinity, oxygen, and carbonate contents.

Steele (1973), Schnitker (1974), Lohmann (1978), and Gofas (1978) have shown that deep-sea benthic foraminifers may have responded to environmental changes by bathymetric or geographic migrations related to deep-water circulations. The case of *Melonis pompeioides* was especially studied on the California borderland (Blake and Douglas, 1980).

For the western North Atlantic, Schnitker (1980) presented a model of correlations between the displacement of bottom-water masses during the last 24,000 yr. and the abundance and distribution of some “index species.” He stated that in recent time, as in interstadial periods, *Epistominella exigua*, a dominant fauna, accompanied by *Cibicides wuellerstorfi* characterized the “North-Atlantic deep water” that occupied the northern part of the North Atlantic (32°N latitude), whereas an *Osangularia umbonifera* fauna, corresponding to the “Antarctic bottom water,” occurred in the southern part of the North Atlantic. At the same time, *Uvigerina peregrina* was restricted to shallower areas, along the North American continental margin and on the Mid-Atlantic Ridge.

During glacial periods, the displacement southward of the polar front induced changes in the distribution and circulation of water masses, and the *Uvigerina* fauna invaded most of the North Atlantic, especially the area where Site 533 is located. *Pullenia* faunas seem to characterize oxygen-depleted waters in some limited intervals.

In the Bay of Biscay, Schnitker (1979) also found that *Uvigerina peregrina* (associated with *Pyrgo murrhina*) characterizes glacial stages, whereas *C. wuellerstorfi*, *Epistominella exigua*, *Oridorsalis umbonatus*, *Melonis pompeioides*, *M. barleeanus*, *Pullenia bulloides*, *P. quinqueloba*, and so on become dominant in interglacial and postglacial times.

*Globobulimina affinis* and *G. pyrula* are listed neither in glacial nor in interglacial Atlantic faunas. Nevertheless, these species seem tolerant to low $O_2$ contents (Mullineaux and Lohmann, 1981).

In order to compare the distribution of all these taxa in cores of Holes 533 and 533A, their variations are summarized graphically in Figure 3.

The following remarks can be proposed:

The species considered are very sparse in the third test interval (middle Pliocene), and no indication can be proposed.

Two of the “index species” of Schnitker (1980) are present: *Uvigerina peregrina*, whose extension in this area characterizes glacial stages, and *Cibicides wuellerstorfi*, which is related to the presence of North Atlantic bottom waters during interstadial periods. *Epistominella exigua*, the main characteristic of this water mass, is present in one sample only (533-37-2, 25–27 cm), but this scarcity may be due to the small size of this species. The absence of *Osangularia umbonifera* throughout the drilling may indicate that Antarctic bottom waters have never occupied this area during the periods investigated. This fact may also be due to the bathymetric position of the site.

Two sorts of assemblages can be distinguished: (1) An assemblage in which *Uvigerina peregrina* forms more than 10% of the fauna, often more than 30%. This probably glacial assemblage is also characterized by low specific diversity (generally less than 12 species per sample). Such a fauna occurs in Samples 533-1-1, 18–20 cm; 533-6, CC; 533-38-3, 25–27 cm; 533-39-3, 33–35 cm; 533-41-1, 47–49 cm, 533-41-2, 42–44 cm; and 533-41-3, 27–29 cm. (2) In the other levels, the specific diversity is generally higher (up to 42 species per sample), and the dominant species are *Melonis, Pullenia, Oridorsalis umbonatus*, and *Cibicides wuellerstorfi*. These levels may be interpreted as interglacial intervals. In some of these samples, the abundance of *Globobulimina* and perhaps also of *Pyrgo murrhina* may indicate a decrease in the oxygen contents. But there is no striking relationship between the presence of pyrite in some sediments and the abundance of these two species.

In the uppermost (postglacial) sample (533-1-1, 2.5–3.5 cm), the fauna, and thus the environmental conditions, seem very peculiar and different from those of the preceding interglacial assemblages; some interglacial species are present but not abundant. The specific diversity is relatively low (10 species) and the only frequent species is *Globocassidulina moluccensis*, which was never present in lower levels. Burke (1981) thinks that this species could be influenced by carbonate deficiency, but in Core 533-1, it is impossible to verify this assumption. Two other species (*Karreriella bradyi* and *Rhabdamina linearis*) are also restricted to this upper level.

**CONCLUSIONS**

Holes 533 and 533A contain from bottom to top a lower bathyal fauna; in some levels some middle bathyal or abyssal species appear.

There are few species reworked from the shelf or from the upper slope.

Distribution of *Uvigerina peregrina*, on the one hand, and of *Cibicides wuellerstorfi*, *Melonis, Pullenia, Oridorsalis*, and so on, on the other hand, are interpreted as reflecting movements of bottom-water masses during glacial and interglacial periods. There is a glacial fauna in Sample 533-1-2, 18–20 cm (last glacial stage). Other levels seem to indicate analogous environmental conditions during the Pleistocene and late Pliocene.
BENTHIC FORAMINIFERS

Table 3. Distribution of dominant benthic foraminifers in selected samples from Holes 533 and 533A.

<table>
<thead>
<tr>
<th>Sample (interval in cm)</th>
<th>Species</th>
</tr>
</thead>
<tbody>
<tr>
<td>533A</td>
<td>Pyrgo murrhina</td>
</tr>
<tr>
<td>533</td>
<td>Globobulimima pyrula</td>
</tr>
<tr>
<td>533</td>
<td>Globobulimima affinis</td>
</tr>
<tr>
<td>533</td>
<td>Uvigerina peregrina</td>
</tr>
<tr>
<td>533</td>
<td>Pullenia bulloides</td>
</tr>
<tr>
<td>533</td>
<td>Cibicides wuellestorfi</td>
</tr>
<tr>
<td>533</td>
<td>Melonis barleeanus</td>
</tr>
<tr>
<td>533</td>
<td>Melonis pomplioioides</td>
</tr>
<tr>
<td>533</td>
<td>Pullenia quinqueloba</td>
</tr>
<tr>
<td>533</td>
<td>Oridorsalis umbonatus</td>
</tr>
</tbody>
</table>

Legend:
- <5%, 20–30%, and >40%
- 5–10%
- 10–20%

Figure 3. Distribution of dominant benthic foraminifers in selected samples from Holes 533 and 533A.

The recent assemblage is different in many ways from the previous interglacial faunas and may reflect peculiar conditions established during the Holocene.

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REFERENCES


**APPENDIX**

**Benthic Species from Holes 533 and 533A:**

**Taxonomic and Paleoecological Notes**

Seventy-three species are listed alphabetically. For all species, reference is made to the original description (see Ellis and Messina's Catalogue of Foraminifera if the original work is not cited in the References). Additional references and remarks are added when necessary for more specific identification. The stratigraphic, bathymetric, or ecological distribution of the most important forms are given from the literature mentioned in the text.

**Alabamina decorata** (Pfieger and Parker) = *Pseudoparella(t) decorata* Pfieger and Parker, 1951, p. 15, figs. 4–5.

**Distribution.** Upper depth limit—about 90 m in the Gulf of Mexico.

**Site 533.** Rare: upper Pliocene; Samples 533-38-2, 106–108 cm and 533-27-4, 26–38 cm.

**Distribution.** Widespread neritic species. A little more frequent than *Ammonia beccarii* (Linné).

**Site 533.** Probably reworked from the shelf. One specimen in Sample 533-6-CC.

**Ammonia beccarii** (Linné) = *Nautilus beccarii* Linné, 1767, p. 710.

**Distribution.** Widespread neritic species.

**Site 533.** Probable reworking from the shelf. One specimen in Sample 533-6-CC.


**Systematic remarks.** There is some doubt about the taxonomic appanance of the two fragments found in one sample only.

**Distribution.** Recorded in middle bathyal, lower bathyal, and abyssal zones.

**Site 533.** One sample: 533-37-2, 25–27 cm.

**Distribution.** The different forms are reported from the outer shelf to bathyal environments.

**Holes 533 and 533A.** Frequent in the finest fraction (160–163 µm).

**Bolivina semiinuda** Cushman, 1911, p. 34, fig. 55.

**Distribution.** It is generally reworked deeper than the preceding species in the Gulf of Mexico, occurring in a lower bathyal zone.

**Holes 533 and 533A.** Frequent in the finest fraction (160–163 µm).

**Buclinimella elegansissima** D'Orbigny, 1839.

**Distribution.** Neritic zone of the different oceans.

**Site 533.** Present in two samples (533-22, CC and 533-33, CC), probably reworked.

**Cassidulina carinata** (Silvestri) = *Cassidulina laevigata* var. carinata Silvestri, 1896.

**Distribution.** Widely distributed in upper and middle bathyal zones. Site 533. Sparse during the Pleistocene.

**Chilostomella oolina** Schwager, 1878.

**Distribution.** Bathyal and abyssal zones of all the oceans.

**Site 533.** Sparse in the Pleistocene and upper Pliocene.

**Cibicides kullenbergi** Parker in Pfieger, Parker, and Peirson, 1933, p. 49, pl. 11, figs. 7–8.

**Distribution.** Described from the Atlantic ocean, where it is “widespread...and a true deep sea form” it occurs also in the Pacific and Indian oceans and in the Mediterranean. *C. kullenbergi* seems also to be a “warm” species in the Atlantic.

**Site 533.** Sparse in the Pleistocene, generally associated with *C. robertsonianus* and often with *C. wuellestorfi*.

**Cibicides lobatus** (Walker and Jacob) = *Nautilus lobatus* Walker and Jacob, 1798.

**Distribution.** Neritic species.

**Site 533.** In two samples, with one specimen only. Probably re-worked.

**Cibicides robertsonianus** (Brady) = *Truncoculina robertsonian Brady, 1881, 1884, p. 664, pl. 95, fig. 4A–C.

**Distribution.** Bathyal species in the different seas.

**Site 533.** A little more frequent than *C. kullenbergi*.

**Cibicides wuellestorfi** (Schwager) = *Anomalia wuellestorfi* Schwager, 1866.

**Distribution.** Frequent from the middle bathyal to abyssal zones; it is generally considered as an interglacial index in the North Atlantic.

**Site 533.** Frequent and sometimes abundant in Pleistocene and Pliocene levels (see text).

**Cornuspira involvens** = *Operculina involvens* Reuss, 1851.

**Site 533.** Sample 533-21, CC.

**Dentalina inornata** (D'Orbigny) *Bradynensis* (Dervieux) = *Nodosaria inornata* (D'Orbigny) *bradyensis* Dervieux, 1894.

**Distribution.** In the Gulf of Mexico, this species occurs in the bathyal and the upper part of the middle bathyal.

**Hole 533A.** In one sample only: 533-29, CC—middle Pliocene.

**Dentalina inornata** (Dervieux) = *Nodosaria inornata* Dervieux, 1894, p. 62, figs. 27–31.

**Distribution.** Gulf of Mexico: lower bathyal to abyssal zone.

**Site 533.** In upper Pliocene: Samples 533-39-3, 33–35 cm and 533 41-3, 27–29 cm.

**Dentalina subencrata** Parr, 1950, p. 329, pl. 12, fig. 1.

**Distribution.** Widespread deep species.

**Site 533.** Sample 533A-38-X, 17–19 cm.

**Eggerella bradyi** (Cushman) = *Veilluitella bradyi* Cushman, 1911, p. 54, fig. 87.

**Distribution.** Bathyal and abyssal world-wide spread species.

**Site 533.** Frequent but never abundant in Pleistocene and upper Pliocene.

**Ellipsidina ellipsoidea** Seguenza, 1859, Brady, 1868, p. 338, pl. 13, figs. 1–12.

**Remarks.** Two large specimens occur in Hole 533A (533A-35-2, 25–27 cm and 533A-38-1, 75–77 cm) and a broken one in Sample 533A-40-1, 38–40 cm. They have an overlapping last chamber and seem identical with the figuration of Seguenza Miocene species.

The specimens also resemble *Ellipsoglandulina antillana* (Bermudez, 1939, p. 247, pl. 33, figs. 1–2) but show no trace of the early chambers described by this author.

**Site 533.** This species is present in the upper Pliocene and in the lowermost Pleistocene (up to Sample 533A-37-2, 25–27 cm).

**Ellipsopolyomphora brevis** (Schwager) = *Pleurostomella brevis* Schwager, 1866.

**Distribution.** The type is Pliocene. Brady (1884) records this species at 230 m depth in recent(?) sediments from Ki Island. But according to Lutze (1979), this species disappears at the beginning of the Brunhes Epoch.

**Site 533.** In Sample 533-35-2, 25–27 cm.

**Epistominella exigua** (Brady) = *Pulvinulina exigua* Brady, 1884, p. 693, pl. 103, figs. 13–14.

**Distribution.** There are some discrepancies between the interpretations proposed by various authors concerning the relations of this species with depth, temperature, and other factors. Schnitker (1980) considers this form an “index species” characterizing the Arctic bottom
water and the lower North Atlantic deep water, which occupied the western part of the North Atlantic Basin during interglacial and postglacial times.

Site 533. *E. exigua* is rare, probably because this species is too small for the mesh sieve used (160 μm); found in one sample: 533-38-2, 15–17 cm.

**Eponides polius** Plieger and Parker, 1951, p. 21, pl. 11, figs. 1–2.

**Distribution.** Deep species.

Site 533. Sparse and never abundant (<5%) in some Pleistocene and upper Pliocene levels.

**Eponides pusillus** Parr, 1950, p. 360, pl. 14, fig. 16.

**Distribution.** Widespread in different seas.

Site 533. Always <5% in some upper Pleistocene and Pliocene levels.

**Francesia advena** (Cushman) = *Virgilina^) advena* Cushman, 1922, p. 120, pl. 25, figs. 1–3.

**Distribution.** Abyssal zone.

Site 533. In two Pleistocene samples (<5%).

**Globobulimina affinis** (D’Orbigny) = *Buliminina affinis* D’Orbigny, 1839.

**Globobulimina pyrula** (D’Orbigny) = *Buliminina pyrula* D’Orbigny, 1846.

**Distribution.** In the Gulf of Mexico, the upper depth limit of *G. pyrula* is about 1400 m. Burke (1981) considers this species independent from depth but corresponding to carbonate deficiency.

Site 533. It is present only in the upper sample (533-1-1, 2.5–3.5 cm).

**Globobulimina subglobosa** (Brady) = *Cassidulina subglobosa* Brady, 1881, 1884, p. 430, pl. 54, fig. 17.

**Distribution.** Widespread in the bathyal zone, this species is controlled by temperature, oxygen content, and, perhaps, salinity.

**Hole 533A:** Sample 533A-29-3, 40–42 cm.

**Gyroidina soldanii** (D’Orbigny) = *Rotulia soldanii* D’Orbigny, 1826.

**Gyroidina neosoldanii** Broten, 1836.

**Holes 533 and 533A.** The former is present in two levels in the upper Pleistocene, the latter is more frequent and occurs in the middle Pleistocene, upper Pleistocene, and Pliocene.

**Hoglundina elegans** (D’Orbigny) = *Rotula* (Turbinulina) elegans D’Orbigny, 1826.

**Distribution.** World-wide species with large depth range from the upper to the lower bathyal zone.

Site 533. Frequent in the upper Pleistocene and Pleistocene levels.

**Karreriella apicularis** (Cushman) = *Gaudryina apicularis* Cushman, 1911, pp. 69–70, fig. 110.

**Distribution.** It is generally reported deeper than 4000 m. In the Gulf of Mexico, this species is present from the upper bathyal to abyssal zones.

Site 533. In the upper level only (Sample 533-1-1, 2.5–3.5 cm).

**Karreriella bradyi** (Cushman) = *Gaudryina bradyi* Cushman, 1911 p. 67, fig. 107.

**Distribution.** As *Eggerella bradyi*.

Site 533. Frequent but never abundant in uppermost Pleistocene and lower Pleistocene.

**Laticarinina pauperata** (Parker and Jones) = *Pulvinulina repanda* F. and M. var. *menardii* D’Orbigny subvar. *pauperata* P. and J. 1866.

**Distribution.** Bathyal species, occurring specially in warm intervals of cores (Lutze, 1979).

Site 533. One occurrence: Sample 533-40, CC.

**Marginalina obesa** = *Marginalina glabra* Flint (non D’Orbigny), 1899

**Distribution.** Bathyal species, occurring specially in warm intervals of cores (Lutze, 1979).

Site 533. One occurrence: Sample 533-40, CC.

**Marginalina glabra obesa** Cushman, 1923, p. 128, pl. 37, fig. 1.

Site 533. Rare, in upper Pliocene and lowermost Pleistocene.

**Martiniella communis** (D’Orbigny) = *Clavulina communis* D’Orbigny, 1846.

**Distribution.** Large stratigraphic range (from Miocene to Recent) and large distribution in recent seas, generally bathyal (upper bathyal).

Site 533. Frequent and somewhat abundant (up to 11%) in lower Pleistocene samples.

**Melonis formosum** (Segnem) = *Nonionina formosa* Segnem 1880.

**Remarks.** *M. formosum* is a moderately compressed species without lirate sutures and clear rim around the umbilicus. These characters differentiate it from *M. barleeanum*.

**Distribution.** Described in Miocene sediments of Calabria, this species occurs also in Pliocene, Pleistocene, and Recent sediments.

**Holes 533 and 533A.** Frequent and sometimes abundant (up to 40%) in all the stratigraphic sections investigated (middle Pliocene, upper Pliocene, Pleistocene, and Recent).

**Melonis barleeanum** (Williamson, 1858).

**Remarks.** It is questionable whether or not *M. parkeri* described from the Gulf of Biscay (Berthois and Le Calvez, 1959, p. 363, pl. 1, figs. 13–14) is synonymous. According to Rovillolos (1970, pp. 200–202), *M. parkeri* has an imperforate apertural face, whereas *M. barleeanum* has an imperforate apertural face, p. 199). This taxonomic problem needs further study.

**Distribution.** *Melonis barleeanum* is generally considered a bathyal species. Nevertheless, it seems to have a larger bathymetric range than *M. pompilioides* in the Mediterranean Sea, *“M. parkeri”* occurs in muddy sediments from outer shelf to lower bathyal zones. In the Atlantic, it is sometimes considered a warm species (interglacial and postglacial).

**Holes 533 and 533A.** Frequent (from middle Pliocene to Recent).

**Melonis pompilioides** (Fichtel and Moll) = *Nautilus pompilioides* Fichtel and Moll, 1798.

**Distribution.** In Recent sediments, this species is generally found in the lower bathyal zone, but in the past, it may have responded to climatic changes by bathymetric migrations.

Site 533. Frequent and often abundant from upper Pleistocene to Recent.

**Nodosaria obliquata** (Batsch) = *Nautilus (Orthoceras) obliquatus* Batsch, 1791.

Site 533. Frequent in upper Pliocene and Pleistocene.

**Nodosaria ovicula** D’Orbigny, 1826.

**Hole 533A.** 10% in Sample 533-29-CC (middle Pliocene).

**Nonionella aff. N. miocenica** Cushman var. *stella* Cushman and Mayer, 1930, p. 56, pl. 7, fig. 17.

**Distribution.** Neritic species.

Site 533. This species occurs with few specimens in three Pleistocene levels. It is probably displaced from the shelf.

**Nuditulis rugosa-convexa** (Parker) = *Epistominella rugosa-convexa* Parker, 1858, p. 273, pl. 4, figs. 21–23.

**Remarks.** The specimen seems analogous to the Mediterranean ones found by Parker and reported also by Wright (1979).

Site 533. Sample 533-38X, 17–19 cm.

**Oridoraslis umbonatus** (Reuss) = *Rotalina umbonata* Reuss, 1851.

**Distribution.** This species is frequent in all oceans (middle bathyal to abyssal).

**Holes 533 and 533A.** Frequent and sometimes abundant.

**Orthomorphina fistuca** (Schwager) = *Nodosaria fistuca* Schwager, 1866.

**Distribution.** This species is considered extinct at the base of the Brunhes epoch (Lutze, 1979).

**Site 533.** Upper Pliocene and lower Pleistocene, up to Sample 533-33, CC.

**Osangularia cultur** (Parker and Jones) = *Planorbulina cultur* Parker and Jones, 1865.

**Distribution.** Upper depth limit in middle bathyal zone.


**Plectofondululicula advena** (Cushman) = *Fondululicula advena* Cushman, 1922, p. 141, pl. 20, figs. 1–2.

**Distribution.** According to Lutze, this Pliocene species disappears at the beginning of the Brunhes epoch.
Site 533. Abundant in some upper Pliocene levels (more than 40\% in Samples 533-38-3, 25-27 cm and 533-40,CC); last occurrence: Samples 533-14,CC.

**Pleurostomella alternans** Schwager, 1886.

**Pleurostomella cf. acuminata** Cushman, 1922, p. 50, pl. 19, fig. 6.

**Distribution.** Deep-water form.

Site 533. The two species of **Pleurostomella** occur sporadically in the upper Pliocene and lower Pleistocene.

**Pseudonodosaria laevigata** (D'Orbigny) = **Nodosaria (Glandulina)** laevigata D'Orbigny, 1826.

**Pullenia bulloides** (D'Orbigny) = **Nonionina bulloides** D'Orbigny, 1846.

**Pullenia quinquemula** (Reuss) = **Nonionina quinquemula** Reuss, 1851.

**Distribution.** The two species of **Pullenia** are deep forms.

**Holes 533 and 533A.** Frequent and often abundant; these species are present in almost all the samples; they are interpreted as interglacial and postglacial forms (see text).

**Pyrgo comata** (Brady) = **Biloculina comata** Brady, 1884, p. 144, pl. 3, fig. 9.

**Pyrgo lucernula** Schwager = **Biloculina lucernula** Schwager, 1886.

**Pyrgo murchisoni** (Schwager) = **Biloculina murchisoni** Schwager, 1886.

**Distribution.** The different species of **Pyrgo** are deep forms (bathyal zone).

Site 533. The genus **Pyrgo** is frequent in the uppermost Pliocene and in the Pleistocene.

**Quinqueloculina venesta** Karrer, 1868.

**Distribution.** Bathyal and abyssal zones.

Site 533. Present (10%) in one upper Pliocene sample (533-41-2, 42-44 cm), more common in Pleistocene samples.

**Quinqueloculina viennensis** Le Calvez and Le Calvez, 1958, p. 187, pl. 5, figs. 42, 44, 45 = **Millotina cuveriana** Brady, 1884 (non D'Orbigny), pl. 5, fig. 12.

**Distribution.** From the Pacific (Brady) and Mediterranean (Le Calvez and Le Calvez) from the shelf and the upper slope.

Site 533. In Samples 533-21,CC and 533-23,CC, probably reworked.

**Rhabdammina linearis** Brady, 1879.

**Distribution.** Deep species.

Site 533. In the topmost sample (533-1-1, 2.5-3.5 cm).

**Sigmoilopsis schlumbergeri** (Silvestri) = **Sigmoilopsis schlumbergeri** Silvestri, 1904.

**Distribution.** Bathyal species.

Site 533. Uppermost Pliocene and Pleistocene.

**Siphotextularia catenata** (Cushman) = **Textularia catenata** Cushman, 1927b, p. 159, pl. 3, fig. 14.

**Remarks.** Cushman described this species from the Pacific Ocean and further reported it from the Atlantic Ocean (1918-31, Pt. 3). 

According to Pfleger, Parker and Peirson, 1953, the species is frequent in upper and middle bathyal, but it had a greater extension during the last glaciation.

**Holes 533 and 533A.** Present in all stratigraphical sections investigated; temporarily abundant in some upper Pliocene and Pleistocene samples, which have been interpreted as glacial periods.

**Uvigerina angulosa** Williamson, 1900, p. 168, pl. 42, fig. 12.

**Distribution.** In the western North Atlantic, the species is now located in upper and middle bathyal, but it had a greater extension during the last glaciation.

**Holes 533 and 533A.** Present in all stratigraphical sections investigated; temporarily abundant in some upper Pliocene and Pleistocene samples, which have been interpreted as glacial periods.

**Uvigerina senticosa** Cushman 1927b, p. 159, pl. 3, fig. 14.

**Distribution.** Deep species. The “deepest uvigerinid index” (Pfleur and Frerichs, 1976). 

Site 533. In upper Pliocene only (15-25%).

**Uvigerina umbilica** Cushman, 1886.

**Remarks.** Little form with very attenuate costae at the base of the chambers. 

Brady (1884, p. 574) thought that this character was of little importance and put **U. umbilica** in synonymy with **U. canariensis.** In fact, some specimens have very attenuate costae and are difficult to attribute to one species or the other—this is the case for the specimens found in Sample 533-39-3, 33-35 cm.

**Distribution.** Miocene and Pliocene.

**Holes 533 and 533A.** In middle and upper Pliocene.

**Valvulinaria laevigata** Pfleger and Parker, 1951, p. 25, pl. 13, figs. 11-12.

**Distribution.** Bathyal species.

Site 533. Rare (in three Pleistocene samples).

**REFERENCES**


