3. DIATOM BIOSTRATIGRAPHY OF LEG 108 SEDIMENTS: EASTERN TROPICAL ATLANTIC OCEAN¹

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ABSTRACT

Diatoms are present in middle Eocene, upper Oligocene-lower Miocene, and Pliocene through Holocene sediments recovered during Leg 108 in the eastern tropical Atlantic Ocean. The diatom assemblage is dominated by species characteristic of a low-latitude, pelagic environment. A comparison of the diatom assemblage preserved at each site allows partial recognition of the 1985 diatom zonations of Barron, Fenner, and Baldauf. In addition, the abrupt occurrence of biosiliceous sediment suggests that present-day equatorial circulation and upwelling began during the late early Pliocene, approximately 3.5 Ma.

INTRODUCTION

Ocean Drilling Program (ODP) Leg 108 provided an opportunity to analyze Cretaceous and Eocene through Holocene sediments from the eastern tropical Atlantic Ocean. The 11 sites occupied during Leg 108 (Table 1 and Fig. 1) are situated in a variety of oceanographic environments and exhibit varying degrees of abundance and preservation of siliceous microfossils. For example, rare diatoms occur in the sediment recovered from Site 657 situated beneath the edge of a zone of high surface-water productivity, and few to common diatoms occur at Site 658 positioned near the center of the productivity cell.

Previous biostratigraphic studies of Paleogene through Quaternary diatoms for the low latitudes consist of studies primarily completed for the equatorial Pacific (Burckle, 1972, 1977, 1978; Burckle and Opdyke, 1977; Burckle and Trainer, 1979; Barron, 1980, 1983, 1985a; Harwood, 1982; Sancetta, 1983; Baldauf, 1985). By comparison, few studies have been completed for similar age sediments from the low-latitude Atlantic. A preliminary examination of Miocene through Holocene diatoms recovered at Deep Sea Drilling Project (DSDP) Site 332 was completed by Schrader (1977). Eocene through lower Miocene diatoms from the equatorial Atlantic regions have been studied by Fenner (1978, 1984, 1985).

The diatom assemblages preserved in Leg 108 sediments allow partial recognition of the low-latitude diatom zonations of Barron (1985a) and Fenner (1985), and the North Atlantic diatom zonation of Baldauf (1985). The Eocene diatom zones employed during this study follow the zonation defined by Fenner (1985) for the equatorial Pacific and Atlantic. The Oligocene through Miocene diatom zones follow the diatom zonation proposed by Barron (1985a, 1985b) for the equatorial Pacific; the Pliocene through Holocene zones follow the zonation defined by Baldauf (1985) for the North Atlantic (Figs. 2, 3, and 4).

The diatom assemblages observed in sediments recovered from Leg 108 consist of species typical of the low latitudes, with most age-diagnostic species for the late Pliocene and Quaternary present. However, several of the marker species either have a sporadic stratigraphic occurrence or have stratigraphic ranges that are diachronous with their ranges in the equatorial Pacific and Atlantic oceans. In fact, several of the diatom species, such as *Nitzschia reinholdii*, have stratigraphic ranges more similar to those in the middle- and high-latitude North Atlantic than those in the low-latitude Pacific.

In addition, several species, such as *Rhizosolenia praeber*gonii, Nitzschia jouseae, and Thalassiosira convexa, have sporadic stratigraphic occurrences that inhibit the determination of the exact placement of the zonal boundaries defined by these species. As a result, we are often required to follow Baldauf (1984, 1987) by replacing the *Rhizosolenia praeber*gonii Zone and the uppermost portion of the Nitzschia jouseae Zone of Burckle (1977) with the Nitzschia marina Zone of Baldauf (1984; Figs. 2, 3, and 4). This latter zone is defined as the interval containing Nitzschia marina between the last occurrence (LO) of N. jouseae and the first occurrence (FO) of Pseudoeunotia doliolus. Baldauf (1987) divided this zone into two subzones based on the LO of T. convexa.

Although the chronostratigraphy used during Leg 108 follows that of Berggren et al. (1985a, 1985b), this chronology does not incorporate a diatom zonation. Therefore, a direct correlation of the FOs or LOs of diatom species to a magnetostratigraphic scale follows that of Barron (1985a, 1985b). Age estimates are based on stratigraphic data obtained from previously drilled DSDP/ODP sites in the low latitudes for those biostratigraphic events that lack direct ties with the geomagnetic polarity scale.

METHODS

Shipboard and subsequent shore-based sample preparation followed the method described in Baldauf (1984) with one exception. Because of the acidity of tap water on board *JOIDES Resolution*, decanting was generally not continued until a pH of 7 was reached. Strewn slides of acid-cleaned material were prepared on 22- \times 40-mm cover glasses and mounted on 25- \times 75-mm glass slides using Hyrax mounting medium.

Strewn slides were examined with a Zeiss compound microscope. At least 450 fields of view (0.5-mm diameter) were examined at $500 \times$, with species identifications confirmed when necessary at $1250 \times$. Species were considered abundant when two or more were present in one field of view at $500 \times$, common if one specimen was encountered in two fields of view, few if one specimen was observed in one horizontal traverse, and rare if less than one per traverse. Criteria for distinguishing whole from partial diatoms follow Schrader and Gersonde (1978).

Preservation was considered good if more than 95% of the diatoms were whole and valves showed virtually no signs of partial dissolu-

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Table 1. Latitude, longitude, and water depth of Sites 657 through 668.

Site	Position	Water depth (m)
Site 657	21°19.89'N, 20°56.93'W	4222
Site 658	20°44.95'N, 18°34.87'W	2263
Site 659	18°04.63'N, 21°01.57'W	3071
Site 660	10°00.809'N, 19°23.166'W	4012
Site 661	9°26.81'N, 19°23.166'W	4012
Site 662	1°23.41'S, 11°44.35'W	3824.3
Site 663	1°11.87'S, 11°52.71'W	3708.1
Site 664	0°06.44'N, 23°16.5'W	3816.5
Site 665	2°57.07'N, 19°40.07'W	4752.3
Site 666	3°29.84'N, 20°10.03'W	4527.3
Site 667	4°34.15'N, 21°54.68'W	3535.2
Site 668	4°46.12'N, 20°55.62'W	2074

tion, reprecipitation, or fracturing. Moderate preservation consisted of 30%-95% whole valves, with moderate breakage and slight dissolution, and some fragile specimens still complete. Also, girdle bands were generally intact. If less than 30% of the diatoms were whole, preservation was regarded as poor. Most diatoms showed extensive breakage, partial dissolution, and pitting. Delicate structures were generally not preserved, and fragile species and girdle bands were generally not intact. If no diatoms were encountered, the sample was recorded as barren.

Biostratigraphy

Figure 5 presents the Leg 108 diatom biostratigraphic results. Diatoms were observed in middle Eocene, upper Oligocene-lower Miocene, and Pliocene through Holocene sediments. The abundance and quality of frustule preservation varies at each site. With the exception of rare and sporadic occurrences, samples examined from Sites 659, 661, and 666 are generally barren of diatoms.

SITE 657

Site 657 (Table 1 and Fig. 1) was proposed as representative of the nonupwelling component of a series of sites (657–659) to examine the late Neogene history of the northern trade winds and upwelling off shore from Africa. Two holes were cored at this site: Hole 657A recovered 16 advanced hydraulic piston corer (APC) and 3 extended core barrel (XCB) cores to 178.2 m below seafloor (mbsf), and Hole 657B recovered 19 APC cores to 166.1 mbsf.

With the exception of rare samples, diatoms are generally absent from the samples examined from each hole. The assemblage observed consists of a mixture of species characteristic of pelagic, coastal, and freshwater environments (with the latter two indicating eolian transport and/or redeposition). The low abundance of diatoms and the rare occurrence of biostratigraphic indicators limit the stratigraphic usefulness of diatoms at this site.

The occurrence of *Pseudoeunotia doliolus* without *Nitzschia reinholdii* in Samples 108-657A-3H-2, 42-44 cm, and 108-657A-3H-CC and in samples examined from Cores 108-657B-1H through -3H indicates placement of these samples into the late Pleistocene through Holocene *Pseudoeunotia doliolus* Zone of Burckle (1977). The rare and sporadic occurrence of diatoms stratigraphically below Cores 108-657A-3H and 108-657B-3H prevents placement of the *Pseudoeunotia doliolus/Nitzschia reinholdii* zonal boundary and any other stratigraphic constraint of the sedimentary sequence.

SITE 658

Site 658 (Table 1 and Fig. 1) was proposed as representative of the "upwelling" component of a series of sites (657–659) to examine the late Neogene history of the northern trade winds and upwelling off shore from Africa. Three holes (658A-658C) were cored at this site to depths of 157.9, 163.8, and 72.9 mbsf, respectively.

Diatoms are present in most samples examined from Holes 658A and 658B. Although the diatom abundance and preservation varies from sample to sample, the diatom assemblage generally consists of few, poorly preserved specimens with occasional samples containing common to abundant, moderately well-preserved specimens. The diatom assemblage present is dominated by pelagic species indicative of high surfacewater productivity. Marine benthic diatoms, freshwater diatoms, and opal phytoliths (from eolian sources) were also observed throughout the recovered sequences.

Age-diagnostic species are present, but they often have sporadic ranges that result in tentative zonal assignments. The LO of *Nitzschia reinholdii* is assigned to Samples 108-658A-8H-CC and 108-658B-8H-4, 92-94 cm. The occurrence of *Pseudoeunotia doliolus* stratigraphically above this LO allows Samples 108-658A-1H-CC to 108-658B-7H-CC and 108-658B-1H-CC to 108-658B-8H-1, 92-94 cm, to be placed in the *Pseudoeunotia doliolus* Zone of Burckle (1977).

The LO of N. reinholdii has been dated by Burckle (1977) for the tropical Pacific and has an estimated age of 0.65 Ma. In the middle- and high-latitude North Atlantic, this biostratigraphic event has an assigned age of 0.44 Ma (Baldauf, 1987). The LO of N. reinholdii in Sample 108-658B-8H-4, 92–94 cm, is slightly above the LO of the calcareous nannofossil *Pseudoumbilica lacunosa*, which has an estimated age of 0.474 Ma (Berggren, 1985b). This suggests that the LO of N. reinholdii in the eastern tropical Atlantic corresponds to the estimated age of its LO in the middle- and high-latitude Atlantic, and significantly postdates that in the equatorial Pacific.

Samples 108-658B-8H-4, 92–94 cm, through 108-658B-15H-1, 62–64 cm, are placed in the *Nitzschia reinholdii* Zone of Burckle (1977) based on the occurrence of *P. doliolus* with *N. reinholdii*. The FO of *P. doliolus* is tentatively placed in Sample 108-658B-15H-1, 62–64 cm; however, the occurrence of diatoms in this sample is extremely rare. Samples 108-658B-14H-5, 110–112 cm, and 108-658B-15H-5, 110–112 cm, contain no age-diagnostic species limiting the stratigraphic interpretation. In Hole 658A, *Pseudoeunotia doliolus* first occurs in Sample 108-658A-14H-CC and allows the interval from Core 108-658A-10H through -14H to be assigned to the *Nitzschia reinholdii* Zone.

Samples 108-658A-15H-CC through 108-658C-23X-CC and Samples 108-658B-15H-3, 110-112 cm, through the base of the hole (108-658B-18H-CC) are assigned to the *Nitzschia marina* Zone of Baldauf (1984). The occurrence of *Thalassiosira convexa* in Sample 108-658A-18H-CC and in Sample 108-658B-18H-6, 111-113 cm, indicates that the LO of this species can be placed stratigraphically above these samples and suggests placement of these samples into Subzone A of the *Nitzschia marina* Zone of Baldauf (1987).

The occurrence of N. jouseae in Samples 108-658A-24X-CC and 108-658A-26X-CC allows placement of these samples into the Nitzschia jouseae Zone of Baldauf (1984, 1987) and suggests that the base of the Nitzschia marina Zone approximates Sample 108-658A-24X-CC. Placement of the Nitzschia jouseae/Nitzschia marina zonal boundary is, however, tentative because of the sporadic and unreliable stratigraphic occurrence of Nitzschia jouseae at this site. The calcareous nannofossil stratigraphy (Manivit et al., this vol.) suggests that Sample 108-658A-24X-CC is slightly older (2.9 Ma), indicating that this sample is indeed equivalent to the Nitzschia jouseae Zone, with the Nitzschia marina/Nitzschia jouseae zonal boundary occurring stratigraphically higher in the sequence.



Figure 1. Geographic location of Leg 108 Sites 657-668.



3 = G. calida 6 = G. tosaensis

Figure 2. Pliocene to Holocene diatom zonation used during Leg 108. This zonation is a composite zonation of Burckle (1977), Barron (1985a), and Baldauf (1984, 1987).

The sporadic occurrence of *Rhizosolenia praebergonii* in Samples 108-658A-18H-CC, 108-658A-24X-CC through 108-658A-26X-CC, 108-658B-14H-2, 110-112 cm, and 108-658A-16H-CC allows placement of these samples into the *Rhizosolenia praebergonii* Zone of Burckle (1977). Although the sporadic occurrence of this species makes the use of this species unsuitable as a zonal marker, its occurrence suggests that these samples are equivalent to the upper portion of the *Nitzschia jouseae* through the lowermost portion of the *Nitzschia reinholdii* Zones. Non-age-diagnostic species were observed in samples examined from Cores 108-658A-27X through -33X.

SITE 659

Site 659 is located on the Cape Verde Plateau in the eastern Atlantic (Table 1 and Fig. 1). This site, like Sites 657 and 658 is situated to monitor surface-water productivity and to examine the late Neogene history of the northern trade winds and "upwelling" off shore from Africa. Three holes (659A-659C) were cored at this site penetrating 273.8, 202.1, and 196.0 mbsf, respectively.

Very few diatoms were observed in the samples examined from Site 659. The diatom assemblage consists of neritic, freshwater (eolian), and occasional pelagic species. Preservation was generally poor and no age-diagnostic species were observed.

SITE 660

Site 660 is situated near Kane Gap (Table 1 and Fig. 1) in order to monitor late Neogene bottom-current activity and deep-water stagnation. The two holes cored at this site (660A and 660B) penetrated 164.9 and 148.8 mbsf, respectively.

Diatoms were observed in middle Eocene and Quaternary sediments. The Quaternary assemblage consists of few, poorly preserved specimens. The assemblage contains warm and warm-temperate marine species with minor freshwater/ eolian components. The middle Eocene assemblage contains a poorly preserved assemblage of generally pelagic species. Although diversity is low, the assemblage is somewhat similar to that described from DSDP Site 354 (Fenner, 1978).

Few stratigraphically useful species are observed in the Quaternary and Eocene sequences. The occurrence of *P. doliolus* without *N. reinholdii* allows Cores 108-660A-1H and 108-660B-1H to be placed in the *Pseudoeunotia doliolus* Zone of Burckle (1977). Samples examined from Core 108-660A-2H through Sample 108-660A-13H-7, 33-35 cm, and from Core



Figure 3. Miocene diatom zonation used during Leg 108. This zonation is a composite zonation of Burckle (1977), Barron (1985a), and Baldauf (1984, 1987).



Figure 4. Oligocene diatom zonation used during Leg 108. This zonation is a composite zonation of Burckle (1977), Barron (1985a), and Baldauf (1984, 1987).

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Figure 5. Diatom biostratigraphic results from Sites 657–658, 660, 662–665, and 667–668. The sparse occurrence of diatoms at Sites 661 and 666 prevents any reliable stratigraphic interpretations.

108-660B-2H through Sample 108-660B-13H-2, 42-44 cm, either contain rare diatom fragments or are barren of diatoms.

The occurrence of *Triceratium schulzii* and *Triceratium brachiatum* in Sample 108-660A-14H-2, 100-102 cm, suggests a middle Eocene age for this sample. *Triceratium schulzii* is recorded from middle Eocene sediments at DSDP Site 354 (Fenner, 1978), the middle Eocene of the Kellogg Shale (Barron et al., 1985), and the late Eocene of the Norwegian Sea (Schrader and Fenner, 1976). *Triceratium brachiatum* is recorded from middle Eocene sediments of DSDP Site 149 in the Caribbean Sea (Fenner, 1985). Other Paleogene species present in this sample include *Pyxilla prolongata* and *Melosira architecturalis*.

Rare to few diatoms generally exhibiting poor preservation occur in samples examined from Cores 108-660A-15H through -17H and Cores 108-660B-15H and -16H. The sedimentary sequence representing these intervals are dominated by common to abundant fragments of radiolarians; the high degree of fragmentation suggests possible current transport of the sediment.

The occurrence of *T. brachiatum* and *Triceratium incon*spicuum in Sample 108-660A-16H-2, 85-87 cm, *T. brachia*tum in Sample 108-660A-15H-5, 42-44 cm, and *Pyxilla caput* avis in Samples 108-660A-15H-5, 42-44 cm, and 108-660A-16H-CC suggest that this interval is equivalent to the middle Eocene *Pyxilla caput avis* through *Hemiaulus gondolaformis*







Figure 5 (continued).

Zones of Fenner (1985). The occurrence of *Hemiaulus gondolaformis* in Sample 108-660A-16H-CC allows placement of this sample into the *H. gondolaformis* Zone of Fenner (1984).

The occurrence of *Triceratium* cf. *barbadense* in samples from Cores 108-660A-15H through -17H and Cores 108-660B-14H through -17H suggests a middle Eocene age for this interval. No age-diagnostic species were observed in samples examined from Core 108-660A-18H.

SITE 661

Site 661, situated east of Kane Gap (Table 1 and Fig. 1), is the second of a two-site transect (with Site 660) cored to document late Neogene bottom-current activity and deepwater stagnation. Holes 661A and 661B were cored to 296.1 and 81.7 mbsf, respectively. Diatoms were generally absent from samples examined from Site 661. The occasional samples that contain diatoms have an assemblage composed of rare to few, poorly preserved specimens. Specimens observed are characteristic of pelagic, neritic, and freshwater environments. With the exception of rare fragments of *Nitzschia jouseae*, no agediagnostic species were observed. The occurrence of *N. jouseae* in Sample 108-661A-8H-2, 42-44 cm, allows tentative placement of this sample into the *Nitzschia jouseae* Zone of Baldauf (1984). Rare pyritized specimens of primitive forms of *Triceratium* sp., *?Coscinodiscus* sp., and *Hemiaulus* sp. occur in Samples 108-661A-25X-CC and 108-661A-27X-CC. These samples are assigned a Cretaceous age based on the calcareous nannofossil biostratigraphy (Manivit, this vol.).

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Figure 5 (continued).

SITE 662

Site 662 was positioned on the eastern flank of the mid-Atlantic Ridge (Table 1 and Fig. 1) to obtain a Neogene record of equatorial divergence and eolian sedimentation. Holes 662A and 662B were drilled to depths of 188.2 and 200.0 mbsf, respectively.

Diatoms are generally common to abundant in the samples examined. Preservation is variable with the majority of samples containing a moderately well-preserved diatom assemblage. The diatom assemblage is dominated by transitional species indicative of moderate to high productivity. The species diversity and composition varies for each sample and may reflect surfacewater response to glacial and interglacial conditions. Samples dominated by *P. doliolus, Thalassionema nitzschioides*, and fragments of *Ethmodiscus rex* suggest glacial intervals, whereas samples dominated by *Coscinodiscus nodulifer* and *Roperia tesselata* were probably deposited during interglacial conditions. Windblown freshwater diatoms, predominantly *Melosira* spp., also occur in varying numbers.

Biostratigraphically useful Pliocene to Holocene species typical of the tropical regions are present at Site 662. The occurrence of P. doliolus without N. reinholdii allows Cores 108-662A-1H through -2H to be assigned to the Pseudoeunotia doliolus Zone of Burckle (1977). The LO of N. reinholdii in Sample 108-662A-3H-CC and the FO of P. doliolus in Sample 108-662A-15H-CC and 108-662B-7H-5, 42-44 cm, allows Cores 108-662A-4H through -15H to be placed in the Nitzschia reinholdii Zone. Similar to the findings of L. H. Burckle and E. Fourtanier (pers. comm., 1984), P. doliolus appears in the equatorial Atlantic prior to its FO in the eastern equatorial Pacific or middle-latitude North Atlantic. At Site 662, the FO of P. doliolus has an estimated age between 2.1 and 2.2 Ma compared with an age of 1.8 Ma for this event in the equatorial Pacific (Barron, 1980) and the North Atlantic (Baldauf et al., 1987).

Cores 108-662A-15H through 108-662A-18H-5, 110 cm, are assigned to the *Nitzschia marina* Zone based on the occurrence of *N. marina* between the LO of *N. jouseae* in Sample 108-662A-18H-5, 110 cm, and the FO of *P. doliolus* in Sample 108-662A-15H-CC. The sporadic occurrence of *N. jouseae* in Samples 108-662A-18H-5, 110 cm, through 108-662A-22H allows samples from this interval to be placed into the *Nitzschia jouseae* Zone of Baldauf (1984).

SITE 663

Similar to Site 662, Site 663 was positioned on the eastern flank of the mid-Atlantic Ridge (Table 1 and Fig. 1) in the eastern Atlantic to monitor the Neogene history of equatorial divergence and eolian deposition. Two holes (663A and 663B) were cored at this site: Hole 663A penetrated 147.2 mbsf, and Hole 663B penetrated 150.0 mbsf.

The diatom assemblage observed at this site is similar to that observed at Site 662 and is characterized by species typical of a transitional, moderate to high surface-productivity environment. Few to abundant diatoms are present in all samples examined. Preservation is generally moderate with occasional samples exhibiting poor preservation. The diatom assemblage is typified by pelagic and freshwater (eolian) species.

Similar to Site 662, diatom biostratigraphy allows recognition of the upper portion of the *Nitzschia jouseae* through the *Pseudoeunotia doliolus* Zones. Cores 108-663A-1H and -2H are assigned to the *Pseudoeunotia doliolus* Zone of Burckle (1977) based on the occurrence of *P. doliolus* without *N. reinholdii*. Although *N. reinholdii* occurs in Sample 108-663B-1H-CC, it is assumed that the rare specimens observed are reworked. The co-occurrence of N. reinholdii and P. doliolus in the interval from Sample 108-663A-3H-5, 42-44 cm, through Core 108-663A-11H allows the samples to be assigned to the Nitzschia reinholdii Zone. The LO of T. convexa in Sample 108-663B-14H-CC and the last common occurrence of N. jouseae in Sample 108-663A-15H-CC allows Cores 108-663A-12H through the lowermost portion of Core 108-663A-15H to be assigned to the Nitzschia marina Zone of Baldauf (1984). Cores 108-663A-16H and 108-663B-16H are assigned to the Nitzschia jouseae Zone. The placement of the top of this zonal boundary is tentative since rare specimens of N. jouseae are observed occasionally throughout Cores 108-663A-11H through -14H; these specimens are assumed to be reworked.

SITE 664

Site 664 is situated on the mid-Atlantic ridge just north of the Romanche Fracture Zone (Table 1 and Fig. 1). The scientific objectives at this site are similar to those at Sites 662 and 663. Four holes were cored at this site. Hole 664A consisted of a single core from a depth of 28.9 mbsf. Holes 664B through 664D penetrated 247.0, 61.2, and 296.8 mbsf, respectively.

Similar to Sites 662 and 663, diatoms are observed only in the upper Pliocene and Quaternary sediments. Diatom abundance and preservation vary from sample to sample but, in general, both are lower than at Sites 662 and 663.

The presence of *P. doliolus* without *N. reinholdii* in samples from Cores 108-664C-1H and -2H and 108-664D-1H allows these intervals to be placed into the *Pseudoeunotia doliolus* Zone. Both *N. reinholdii* and *P. doliolus* were observed in Sample 108-664A-1H-CC, Cores 108-664B-1H through -5H, Samples 108-664C-3H-CC through 108-664C-7H-CC, and Samples 108-664D-3H-CC through 108-664D-10H-CC. Thus, these intervals are assigned to the *Nitzschia reinholdii* Zone. The FO of *P. doliolus* in Sample 108-664D-9H-CC marks the base of the *Nitzschia reinholdii* Zone. Because of the occurrence of slump deposits and/or a decline in the diatom preservation and abundance, this zonal boundary was not recognized in Hole 664B.

The LO of *N. jouseae* in Samples 108-664B-14H-CC and 108-664D-14H allows the interval between Samples 108-664D-10H-CC through 108-664D-13H-CC to be placed into the *Nitzschia marina* Zone. Specimens of *N. jouseae* occur downsection until Cores 108-664C-23H and 108-664D-17H, suggesting that these intervals are equivalent to the *Nitzschia jouseae* Zone. The base of this zone was not recognized, since the abundance and preservation of diatoms abruptly decline below Core 108-664D-20H. This sharp downhole decline in the preservation of biosiliceous material suggests that the equatorial divergence at Site 664 may have been less intense prior to 3.5 Ma.

SITE 665

Sites 665 through 668 (Table 1 and Fig. 1) were positioned on the Sierra Leone Rise to monitor late Neogene, long-term fluxes in calcium carbonate and calcium carbonate dissolution. Site 665 is located on the southwest margin of this rise. Holes 665A and 665B penetrated 97.9 and 71.7 mbsf, respectively.

Diatoms were only observed in a few samples. The assemblage present is composed of pelagic species, freshwater (eolian) species, and fragments of *Ethmodiscus* sp. With the exception of the uppermost sequence, age-diagnostic species were not observed. The occurrence of *P. doliolus* without *N. reinholdii* in samples examined from the first two cores of Hole 665B allows these cores to be assigned to the *Pseudoeunotia doliolus* Zone. The co-occurrence of these species in Sample 108-665A-2H-CC places this sample in the *Nitzschia reinholdii* zone.

SITE 666

Site 666 is an intermediate member of the Sierra Leone Rise transect. Because of extensive turbidite deposits, only one hole (666A) was cored at this site to a depth of 150.0 mbsf. Rare diatoms occur sporadically in the upper portion of Hole 666A. Species present include specimens of freshwater *Melosira*, the marine species *Coscinodiscus marginatus*, and fragments of *Ethmodiscus* sp. No age-diagnostic species were observed.

SITE 667

Site 667, also an intermediate member of the Sierra Leone Rise transect, is positioned on the western margin of the Sierra Leone Rise (Table 1 and Fig. 1) to monitor Neogene deep-water circulation. Hole 667A recovered 41 cores that penetrated 381.3 mbsf, and Hole 667B recovered 15 cores to a depth of 139.1 mbsf.

Diatoms are generally absent in the samples examined. However, poor to well-preserved and few to common species were observed in Cores 108-667A-1H, -2H, -23H through -31X, and -33X to -39X as well as in Core 108-667B-1H. These intervals are Pleistocene-Holocene and Oligocene-early Miocene in age.

The Nitzschia reinholdii/Pseudoeunotia doliolus zonal boundary is placed between Samples 108-667A-1H-CC and 108-667A-2H-CC based on the sole occurrence of *P. doliolus* in Core 108-667A-1H and the co-occurrence of *P. doliolus* and *N. reinholdii* in Sample 108-667A-2H-CC. Diatoms were generally not observed from samples in the interval from Cores 108-667A-3H through -22H.

The diatoms present in Cores 108-667A-23H through -31X and 108-667A-33X through -39X are similar to those described by Barron (1985a) from the equatorial Pacific. The interval from Sample 108-667A-24X-6, 20-22 cm, through Core 108-667A-26X is tentatively assigned to the early Miocene Rossiella paleacea Zone based on the sporadic occurrence of Coscinodiscus oligocenicus, Synedra jouseana, and Craspedodiscus elegans in this interval.

Samples 108-667A-27X-CC and 108-667A-28X-CC are assigned to the upper portion of Subzone A of the Rossiella paleacea Zone based on the co-occurrence of Bogorovia veniamini, Thalassiosira primalabiata, and Rossiella paleacea. The occurrence of B. veniamini, Coscinodiscus lewisianus var. similis, S. jouseana, and Rocella vigilans without R. paleacea and Rocella gelida in Sample 108-667A-35X-4, 10-12 cm, suggests placement of this sample into the Bogorovia veniamini Zone.

Sample 108-667A-38X-CC is tentatively placed in Subzone A of the *Rocella vigilans* Zone based on the occurrence of common *R. vigilans* and rare specimens of *Cestodiscus muhinae*.

SITE 668

Site 668 was drilled on the crest of the Sierra Leone Rise as a companion site to Sites 665 through 667 (Table 1 and Fig. 1). The primary objective at this site was to recover a continuous upper Neogene section to monitor deep-water circulation in the eastern equatorial Atlantic.

Rare diatoms were observed in the few samples examined from this site. Preservation is generally poor and, with the exception of the occurrence of P. doliolus in samples from Core 108-668A-1H and Sample 108-668B-1H-CC, age-diagnostic species were not observed. The occurrence of P. doliolus without N. reinholdii allows these intervals to be placed into the *Pseudoeunotia doliolus* Zone.

DISCUSSION

Figure 5 shows diatom biostratigraphic results from Leg 108. Sites 659 and 666 are excluded from this diagram because diatoms were generally absent from the samples examined. As a rule, diatom abundances and preservation in the Pliocene-Pleistocene are greater at sites underlying surface waters that today are relatively rich in nutrients and exhibit moderate to high surface-water productivity.

The existence of a permanent cell of upwelling off northwest Africa thus explains why diatom assemblages are richer at Site 658 than at nearby Sites 657 and 659, which lie outside the upwelling cell. Similarly, the strong divergence along the equator, together with advection of nutrient-rich waters in the Benguela Current, leads to the high productivity of equatorial waters and, hence, to the relatively rich diatom assemblages at Sites 662, 663, and 664. The fact that diatoms tend to be more abundant and better preserved at Sites 662 and 663 than at Site 664 is consistent with the east-to-west decrease in nutrient contents and productivity levels of equatorial waters. Leg 108 sites on and near the Sierra Leone Rise (Sites 660, 661, and 665–668) are also under waters of relatively low productivity and therefore are poor in diatoms.

Comparing Pliocene and Quaternary sediments from each of these sites shows that biosiliceous sediments (based on diatom presence and absence) occurs during the interval equivalent to the upper portion of the *Nitzschia jouseae* Zone to the present day. Thus, siliceous deposition similar to the present day has occurred in these regions during the latest early Pliocene (about 3.5 Ma) to the present. Prior to this time (late Miocene and early Pliocene), biosiliceous sediments (diatoms) do not occur at these sites. The transition from siliceous to nonsiliceous sediments at about 3.5 Ma suggests that present-day surface circulation patterns were established at this time (i.e., equatorial divergence and upwelling), increasing both surface productivity and the flux of biogenic silica to the seafloor.

CONCLUSION

Siliceous microfossils, specifically diatoms, are useful as general stratigraphic indicators for the middle Eocene, upper Oligocene-lower Miocene, and upper Pliocene-Holocene sediments recovered during Leg 108 in the eastern Atlantic. Based on diatom biostratigraphy, the stratigraphic intervals containing a siliceous component can be placed into the *Hemiaulus gondolaformis, Bogorovia veniamini, Rossiella paleacea*, and the upper portion of the *Nitzschia jouseae* through *Pseudoeunotia doliolus* Zones.

The stratigraphic markers used to define the Pliocene through Holocene zones are calibrated to magnetostratigraphy in both the equatorial Pacific (Burckle, 1977; Barron, 1985a) and in the middle and high latitudes of the North Atlantic (Baldauf, 1987; Baldauf et al., 1987). Comparisons of the stratigraphic ranges of marker species in the Leg 108 sediment suggest that the species in the eastern tropical Atlantic have stratigraphic ranges similar to those in the middle- and high-latitude North Atlantic rather than to those in the eastern equatorial Pacific.

The diatom assemblage present in Leg 108 sediments consists of pelagic species typical of the low latitudes, pelagic species characteristic of moderate to high surface-water productivity, freshwater species indicative of eolian deposition, and occasional nearshore benthic forms.

Examination of the Pliocene and Quaternary sequences recovered from the 11 sites occupied during Leg 108 indicates that, if present, biosiliceous sediments were generally incorporated into the sediment record during the time equivalent to and younger than the upper portion of the *Nitzschia jouseae* Zone (latest early Pliocene; about 3.5 Ma). This suggests that the present-day pattern of surface productivity in response to equatorial surface circulation (divergence) commenced about that time.

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