

32. CALCAREOUS NANNOFOSSIL PALEOGENE BIOSTRATIGRAPHY, CÔTE D'IVOIRE-GHANA MARGINAL RIDGE, EASTERN EQUATORIAL ATLANTIC¹

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ABSTRACT

Paleogene sections on the Côte d'Ivoire-Ghana Marginal Ridge in the eastern Atlantic Ocean, sampled at three sites during Ocean Drilling Program Leg 159, are shown to have calcareous nannofossil assemblages in distinct packages of (sub)zones separated by hiatuses or barren sediments. The packages are different at each of the sites; all fall within the interval from late Paleocene Zone CP6 to latest Oligocene Subzone CN1a inclusive. Three packages (CP9b–CP11, CP13, and CP19) on the crest of the Marginal Ridge at Site 960, four (CP7–CP9b, CP11, CP13–?CP15, and CP17–CN1a) at nearby Site 959 on the shoulder of the Marginal Ridge, and only two (CP6–CP8a, and CP10) at the deeper Site 961, on the southern tip of the Marginal Ridge, were identified. The Paleogene section at Site 960 is thinner and less complete than at Site 959, but it is thinnest and least complete at Site 961. Conditions during most of the Paleogene were strikingly different near the crest of the Marginal Ridge (most affected by changes in sea level) compared with those on its lower slopes (where bottom currents were probably more effective in removing sediments or preventing sedimentation in an already sediment-starved area).

Hiatuses on the crest of the Marginal Ridge are more frequent than on its shoulder where the nannofossil succession is interrupted by intervals with seemingly barren sediments instead. Two short hiatuses at Site 960 are almost coincident with, and probably related to, major falls in the global sea level of ~125 m during the earliest middle Eocene (also coeval with a hiatus at Site 959 and elsewhere in the Atlantic), and of ~60 m during the latest Oligocene. An intervening hiatus at the same site (middle Eocene–lower Oligocene section virtually being missing) corresponds with a period of strongly fluctuating global sea level, which included three major falls.

INTRODUCTION

Cores recovered during Ocean Drilling Program (ODP) Leg 159 from the Côte d'Ivoire-Ghana Marginal Ridge in the eastern Atlantic Ocean (Fig. 1) were sampled; one sample per 1.5-m core section in most cases. These were studied in order to document the calcareous nannofossil distribution, and to provide biostratigraphic subdivision and a time framework for interpretation of the geological development of the Marginal Ridge.

The Côte d'Ivoire-Ghana Marginal Ridge, a major physiographic feature, lies at the eastern extension of the Romanche Fracture Zone, separating the Deep Ivorian Basin and the Gulf of Guinea Abyssal Plain. It was formed as a consequence of a series of events that started early in the Cretaceous with rifting of the northern South Atlantic, followed by active transform motion initially between the African and Brazilian plates, and subsequently between the evolving, extensional Deep Ivorian Basin and the newly created, bordering oceanic crust to the south. Thermal subsidence and Cenozoic sea-level changes dominated the final stage in the evolution of the Marginal Ridge.

METHODS

Assemblages were analyzed in smear slides prepared using either raw sediment samples (core-catcher samples) or sediment suspensions in distilled water (other samples). Gravitational settling was also used, in a few cases, to concentrate sparse nannofossil assemblages in critical samples. For each sample two slides were prepared, one with double the number of suspension drops used for the other.

Slides were examined with an optical microscope at 1250× magnification. At least one sample per core section was examined. Estimates for the total nannofossil abundance in a slide were determined as follows: D (dominant) = >60% of all particles; A (abundant) = 30%–60% of all particles; C (common) = 10%–30% of all particles; F (few) = 5%–10% of all particles; R (rare) = 1%–5% of all particles; T (trace) = <1% of all particles. Estimates of nannofossil preservation are coded as follows: E (excellent) = pristine preservation with no overgrowth or dissolution; G (good) = slight dissolution and overgrowth, but all taxa are easily identifiable; M (moderate) = obvious signs of dissolution and overgrowth, but most specimens are readily identifiable; P (poor) = strong overgrowth and dissolution, and most specimens are fragmented and difficult to identify. Estimates of the relative abundances of nannofossil species were determined as follows: D (dominant) = >60% of all nannofossils; A (abundant) = 30%–60% of all nannofossils; C (common) = 10%–30% of all nannofossils; F (few) = 5%–10% of all nannofossils; R (rare) = 1%–5% of all nannofossils; vR (very rare) = <1% of all nannofossils.

NANNOFOSSIL DISTRIBUTION AND BIOSTRATIGRAPHY

The distribution of nannofossils in the Paleogene sections on the Côte d'Ivoire-Ghana Marginal Ridge is given in Tables 1–5. Some of the Eocene species are illustrated in Plates 1 and 2. A biostratigraphic summary is presented in Figure 2 and is discussed below.

We have used the zonation of Okada and Bukry (1980), supplemented by a few of Martini's (1971) zonal indicators for greater biostratigraphic resolution. The zonation of Okada and Bukry (1980: the CP Zones/Subzones) was based originally on low-latitude oceanic assemblages (Bukry, 1973, 1975); whereas, the zonation of Martini (1971: the NP zones) was based on mid-latitude assemblages from land-based sections. Perch-Nielsen (1985) incorporated some of Bukry's (1973) (sub)zonal indicators into the definition of some of the NP Zones.

¹Masle, J., Lohmann, G.P., and Moullade, M. (Eds.), 1998. *Proc. ODP, Sci. Results, 159*: College Station, TX (Ocean Drilling Program).

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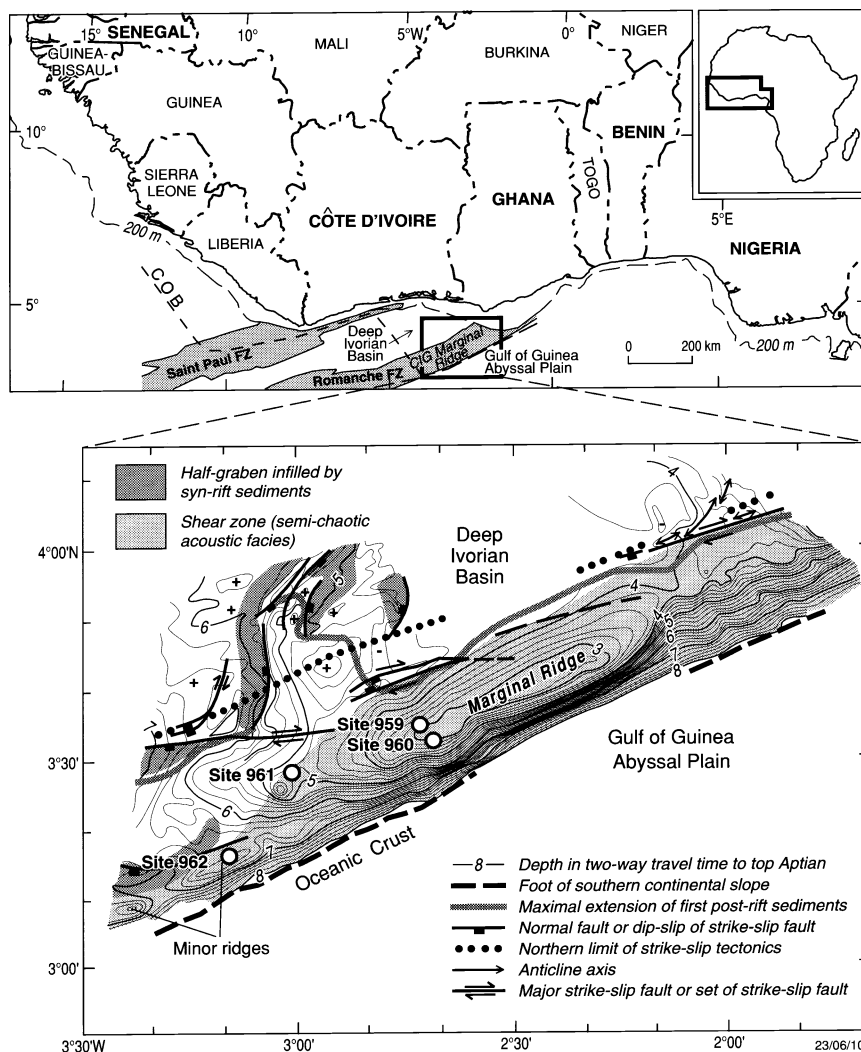


Figure 1. Location of sites drilled during ODP Leg 159 and the neighboring African coast (in part modified after Mascle, Lohmann, Clift, et al., 1996). Site 962 contains no Paleogene calcareous nannofossils.

Hole 959A (Water Depth 2090.7 m)

The basal Samples 159-959A-52X-CC through 43X-CC are barren of calcareous nannofossils. This is attributable in part to the nature of the sediment (clayey diatomite, chert, and massive claystone) and in part to the poor core recovery in this interval. Similar sediment and poor recovery characterize the depth-equivalent part of Hole 959D.

The key species *Sphenolithus distentus* occurs in the interval from Samples 159-959A-42X-CC through 40X-CC, but in association with *Sphenolithus ciperoensis* only in Section 159-959A-40X-CC (Table 1). This indicates Zone CP18 and Subzone CP19a. In Sample 159-959A-42X-CC, forms transitional between *Sphenolithus distentus* and *S. ciperoensis* are considered indicative of the upper part of Zone CP18. The absence of *S. ciperoensis* from the sparse assemblage in Sample 159-959A-41X-CC is probably due to poor preservation, and the assemblage is questionably placed in Subzone CP19a (Fig. 2). The section with assemblages readily assignable to Subzones CP19b and CN1a (Samples 159-959A-40X-1, 106–107 cm, through 32X-5, 129–130 cm; see Table 1) includes several levels that are barren of calcareous nannofossils. These are mostly within Subzone CN1a, being represented by Samples 159-959A-39X-1, 100–101 cm; 34X-4, 102–103 cm; 34X-2, 100–101 cm; 33X-7, 52–53 cm; 33X-6, 97–98 cm; and 33X-3, 100–101 cm.

Dictyococcites bisectus is rare, and several other mid- to high-latitude species (such as holococcolith *Zygrhablithus bijugatus*, and *Pontosphaera pygmaea*) are absent (see Table 1). This is consistent with a dominantly open marine, oceanic setting, at low latitude during the late Oligocene.

In this hole, we recovered the only seemingly complete succession of nannofossil assemblages across the Oligocene/Miocene boundary on the Côte d'Ivoire-Ghana Marginal Ridge (see Fig. 2 for comparison with other holes). Criteria for recognition of this boundary vary between authors (compare, e.g., Martini, 1971; Bukry, 1973; Gartner, 1974; and Martini, 1986), because of differences in types of sediments (oceanic vs. hemipelagic) and in geographic location of sections (e.g., latitudes). In low-latitude oceanic sections, the Oligocene/Miocene boundary is usually approximated by either the end of the acme of *Cyclicargolithus abisectus* or the (slightly older) last appearance datum (LAD) of *Sphenolithus ciperoensis*. As shown in Figure 2, we have both events in the Hole 959A cores, but adopt the former as the evidence for delineating the Oligocene/Miocene boundary. The end of the acme of *C. abisectus*, which marks the top of Subzone CN1a, is inherently difficult to locate precisely and consistently in routine or semiquantitative work; populations of this species and of *Cyclicargolithus floridanus* are usually difficult to separate, particularly within Zone CN1, adding to the difficulty of pinpointing the acme of *C. abisectus*. Nevertheless, we are fairly certain of the identification of this event in Sample 159-959A-32X-5, 129–

Table 2 (continued).

Species	CP18	CP17	CP14b/ ?15a	CP14a	(Sub)Zones	Age
<i>Ericsonia robusta</i>						
<i>Fasciculithus alanii</i>	R					
<i>Fasciculithus involutus</i>						
<i>Fasciculithus lilianae</i>						
<i>Fasciculithus richardii</i>						
<i>Fasciculithus tymparifomis</i>						
<i>Helicosphaera dinesenii</i>	R					
<i>Helicosphaera euphratis</i>		C				
<i>Helicosphaera lophota</i>		F				
<i>Helicosphaera obliqua</i>						
<i>Helicosphaera papillata</i>						
<i>Helicosphaera recta</i>	R					
<i>Helicosphaera seminulum</i>	R					
<i>Helicosphaera truempyi</i>						
<i>Helolithus cantabrigiae</i>						
<i>Helolithus kleinpelli</i>						
<i>Helolithus megastypus</i>						
<i>Helolithus redelti</i>						
<i>Hormibrookina australis</i>						
<i>Lopholithus mochloporus</i>						
<i>Lopholithus nasceus</i>						
<i>Markalius inversus</i>						
<i>Nannoctrina cristata</i>						
<i>Nannoctrina fulgens</i>						
<i>Nannoctrina poppii</i>						
<i>Neochastoygyas junctus</i>						
<i>Neococcolithes dubius</i>						
<i>Placozgyas sigmoides</i>						
<i>Pontosphaera multipora</i>						
<i>Pseudotrifarcta inversus</i>						
<i>Reticulofenestra dictyoda</i>						
<i>Reticulofenestra lockeri</i>						
<i>Reticulofenestra samoderovii</i>						
<i>Reticulofenestra umbilicus</i>						
<i>Rhomboaster birifida</i>						
<i>Sphenolithus anarripopus</i>						
<i>Sphenolithus distentus</i>	F					
<i>Sphenolithus elongatus</i>						
<i>Sphenolithus furcatolithoides</i>	F					
<i>Sphenolithus moriformis</i>						
<i>Sphenolithus predistentus</i>	F					
<i>Sphenolithus primus</i>						
<i>Sphenolithus pseudoradicans</i>	R					
<i>Sphenolithus radicans</i>						
<i>Sphenolithus spiniger</i>						
<i>Toweius emimens</i>						
<i>Toweius magnicrassus</i>						
<i>Toweius pertusus</i>						
<i>Toweius tovae</i>						
<i>Tribrachiatus bramlettei</i>						
<i>Tribrachiatus comortus</i>						
<i>Tribrachiatus orthostylus</i>						
<i>Zygodiscus herlyni</i>						
<i>Zygrrhablithus bijugatus</i>						
(Sub)Zones	CP18	?CP17	CP14b/ ?15a	CP14a		early Oligo.
Age						middle/ ?late Eocene

interval CP14b–CP15. The interval of Cores 159-959D-13R through 9R is barren of calcareous nannofossils.

The first eight cores in this hole are approximately depth equivalents of the last eight cores in Hole 959A, and likewise came from a sequence of diatomite, chert, and claystone, mostly barren of calcareous nannofossils. Only Samples 159-959D-7R-CC and 5R-CC yielded nannofossils and they are assigned to Zones CP17 and CP18 respectively. Identification of Zone CP17 relies on negative evidence (the absence of both *Sphenolithus distentus* and *Reticulofenestra umbilicus*). Because of poor preservation and scarcity of fossils, assignment to this gap zone is somewhat uncertain. Zone CP18 assemblages contains *Dictyococcites scrippsae*, *Helicosphaera recta*, *Sphenolithus distentus*, *Sphenolithus predistentus*, and *S. pseudoradicans*; zonal assignment is more certain than below.

Hole 960A (Water Depth 2048.3 m)

The base of the nannofossil-bearing Paleogene section in this hole is within the 26-cm-long Section 159-960A-20R-CC. A (dark) bluish green claystone sample from this section yielded traces of a poorly preserved assemblage assignable to Subzone CP9b (Table 3), but a sample from a lighter component mixed in with the same claystone produced a diverse assemblage indicative of Late Cretaceous age (Coniacian–Santonian). All samples examined from below Core 159-960A-20R (down to Sample 159-960A-34R-CC) are barren of calcareous nannofossils. Preservation of nannofossils in the palygorskite claystone of Core 159-960A-20R is poor (Table 3), with total

loss in at least two levels (represented by Samples 20R-CC, 8–9 cm, and 20R-1, 9 cm). The few nannofossils in Sample 159-960A-20R-1, 2–3 cm, include *Tribrachiatus orthostylus*, *Discoaster binodosus*, *Chiasmolithus grandis*, and *Sphenolithus radicans*, and suggest Subzone CP9b.

The lithological unit comprising Cores 159-960A-19R through 16R consists of chert, porcellanite, and claystone. Core recovery was very poor, and preservation of calcareous nannofossils is generally poor, with Samples 159-960A-19R-CC and 19R-1, 33–34 cm, being barren; there was no core recovery for Core 159-960A-18R. Assemblages in Sections 159-960A-17R-CC and 16R-CC show contrasting states of preservation: a residual assemblage (a concentration of ortholiths and a few coccoliths) in the former assignable to Zone CP10; and a moderately preserved, more diverse assemblage in the latter assignable to Zone CP11. The key species of Zone CP11, *Coccolithus crassus*, was not found.

The quality of preservation of Subzone CP12a in the poorly recovered Core 159-960A-15R alternates sharply with concomitant species diversity changes. For example, only traces of discoasters (pieces of broken *Discoaster barbadiensis* and ?*D. lodoensis*) occur in Sample 159-960A-15R-1, 39 cm, whereas, a few centimeters above in the better preserved Sample 15R-1, 15–16 cm, several coccolith species (e.g., *Chiasmolithus grandis*, *C. solitus*, *Coccolithus formosus*, and *Reticulofenestra dictyoda*) and a larger number of *Discoaster* species occur.

Several species of the genus *Sphenolithus*, including *S. elongatus*, *S. furcatolithoides*, *S. orphanknollensis*, and *S. radicans*, are particularly notable among the assemblages of Zone CP13 recovered from

Cores 159-960C-17X and 16H. These include species of the largely hemipelagic genus *Transversopontis*.

A barren section of >10 m of glauconitic claystone and radiolarite (comprising all of Core 159-960C-15H and Sample 159-960C-14H-7, 34–35 cm), separates middle Eocene Subzone CP13a (in Sample 159-960C-16H-1, 75 cm) and Subzone CP19a (in Sample 159-960C-14H-6, 34–35 cm). A middle Eocene radiolarian fauna was extracted from Core 159-960C-15H, and assigned to the *Podocyrtis chalara* Zone (Shipboard Scientific Party, 1996a), indicating that the gap in the nannofossil record between Subzones CP13a and CP19a is largely a hiatus. The barren interval is probably restricted to the middle Eocene (see Fig. 2): the radiolarian *Podocyrtis chalara* Zone is correlative with the lower part of Subzone CP14b (see, e.g., correlation chart in Haq et al., 1988).

Similarly to Hole 960A, the youngest Paleogene nannofossils in this hole are assignable to the late Oligocene Subzone CP19b (Table 4), but identification of Subzone CP19a is more certain in this hole. Both *Sphenolithus distentus* and *S. predistentus* are present, and the better preserved assemblages are much more diverse. The solution-prone *Zygrhablithus bijugatus* (known to have preferred higher latitudes) and *Pontosphaera multipora* are present, albeit rarely, among assemblages showing signs of mild dissolution in the upper samples of Zone CP19.

Hole 961A (Water Depth 3292 m)

The nannofossil-bearing Paleogene section in this hole is thin, comprising Core 159-961A-20R through to within the base of Core 159-961A-17R; the underlying thick section (down to Core 159-961A-35R) is barren of calcareous nannofossils. The basal assemblages, in Samples 159-961A-20R-CC and 20R-1, 111–113 cm, are assignable to late Paleocene Zone CP6. These include rare *Prediscosphaera cretacea*, reworked from a Cretaceous source. Sample 159-961A-20R-2, 90–91 cm, within Zone CP6, is barren of calcareous nannofossils.

Zone CP7 nannofossils, in Samples 159-961A-20R-1, 15–16 cm, and 19R-CC, are characterized by a high diversity of the genera *Heliolithus* and *Fasciculithus*. Species of the genus *Fasciculithus* continue in Subzone CP8a above, in samples from Section 159-961A-19R-1, but are absent from overlying cores, indicating the top of the Paleocene in the hole.

Early Eocene Zone CP10 in Sample 159-961A-17R-CC is based on the association of *Cyclicargolithus gammation*, *Chiphragmalithus acanthodes*, *C. calathus*, *Discoaster kuepperi*, and *T. orthostylus*; *Discoaster lodoensis* is absent from this sample, but occurs in the underlying core. Other samples examined from Core 159-961A-17R are devoid of calcareous nannofossils. The radiolarian fauna recovered from this core is suggestive of the combined *Buryella clinata-Phormocyrtis striata striata* Zones of early Eocene age (Shipboard Scientific Party, 1996b). These zones have been shown to correlate with upper Zone CP11 to within Subzone CP12a (see, e.g., chart in Haq et al., 1988). Accordingly, the barren interval (barren samples from Core 961A-17R) above Zone CP10 in Figure 2 is extended to within the lower part of Subzone CP12a.

Core 159-961A-16R is also devoid of calcareous nannofossils, and the first upcore occurrence of nannofossils, in Core 159-961A-15R, is indicative of the early Miocene Subzone CN1c. Radiolarians in Core 159-961A-16R were assigned to the early Miocene *Lychnocanoma elongata* Zone age (Shipboard Scientific Party, 1996b). This indicates a substantial hiatus between lower Eocene and lower Miocene sediments (Cores 159-961A-17R and 16R, respectively).

Eocene Deposition Depth: Comparison with Onshore Sections

Claystones dominated by the mineral palygorskite were recovered in the lower Eocene sediments at Sites 960 and 961 (see Mascle,

Lohmann, Clift, et al., 1996); these are assigned to Subzone CP9b and Zone CP10 at Site 960 (see Tables 3 and 4), and mainly to Zone CP10 at Site 961 (Table 5). The comparatively pure occurrences of this mineral, particularly at Site 960, suggest an authigenic origin, requiring conditions of elevated temperatures and increased salinities (Mascle, Lohmann, Clift, et al., 1996, with references). This indicates shallowing on the Marginal Ridge by uplift or sea-level fall, producing locally restricted, warm, hypersaline conditions (Mascle, Lohmann, Clift, et al., 1996). Where the nannofossils occur in the Eocene sections on the Marginal Ridge, they provide evidence for open marine, oceanic conditions; this is also true for the Paleocene and Oligocene sections

Calcareous nannofossil assemblages assignable to the zonal interval CP9b–CP13b have been documented in land-based sections in neighboring equatorial West Africa. Differences between these assemblages and coeval assemblages in ODP Leg 159 cores are obvious (discussed below) and support the conclusion that at this time deposition on the Côte d'Ivoire-Ghana Marginal Ridge was under oceanic conditions (i.e., in an open marine setting).

Eocene nannofossil assemblages in neritic sections in Nigeria, Benin, and Sénégal (Fig. 1) have been shown to consistently contain taxa indicative of depositional environments akin to continental shelf, or epicontinental or marginal sea (e.g., pentoliths, pontosphaerids, and holococcoliths; see, Perch-Nielsen and Petters, 1981; Akpiti et al., 1982; Toumarkine et al., 1984). Thus, in an outcrop of the Ameke Formation in the southern Benue Trough in Nigeria, assemblages of Subzone CP13b contain a large number of hemipelagic species, with most of the genera *Pontosphaera*, *Pemma*, *Holodiscolithus*, *Micrantholithus*, *Transversopontis*, *Trochoaster*, and *Zygrhablithus* being well represented (Perch-Nielsen and Petters, 1981). Similarly, a large number of species of most of the same genera were reported among diverse assemblages assignable to Zones CP12 and CP13 from a section in Benin (Akpiti et al., 1982). In a bore in the Cap Vert area in Sénégal, assemblages of Zones NP11 and NP12 (Subzone CP9b and Zone CP10) contain *T. pulcher*, *T. pulcheroides*, and *Z. bijugatus* (Toumarkine et al., 1984). In contrast, in the Eocene (and Oligocene) on the Côte d'Ivoire-Ghana Marginal Ridge, none of the pentoliths (e.g., *Pemma* and *Micrantholithus*) are present, and the holococcoliths (e.g., *Holodiscolithus* and *Zygrhablithus*) are restricted to only a few occurrences of *Z. bijugatus*. These are in the middle Eocene Zones CP13 and CP14 at Site 959 (Table 4), where reworking is suspected (e.g., *Discoaster lodoensis* being present), but also include a single occurrence in the upper Oligocene at Site 960. The usually diverse genus *Pontosphaera* (in neritic/land-based sections) is limited to a single species, *P. multipora*. This species occurs mainly in Zones CP13 and CP14 at Site 959 as a very minor component, and with *Transversopontis pulcher* and *T. pulcheroides* in Subzone CP13a at Site 960, also as a very minor component (Tables 2–4). None of the hemipelagic genera/species in the west African Eocene neritic sections was encountered in the lower Eocene of Sites 960 and 961 (Tables 3–5), where concentrations of the mineral palygorskite are present. This evidence, although negative, is in agreement with that from the associated benthic foraminifers that indicate lower bathyal to abyssal depth for the Eocene at Site 961 (see Mascle, Lohmann, Clift, et al., 1996).

INTERVALS WITH NO NANNOFOSSIL RECORD

One striking feature of the Paleogene sections on the Côte d'Ivoire-Ghana Marginal Ridge is the record of repeated episodes of non-deposition and/or condensed sedimentation (Site 960), or deposition of noncalcareous (black claystone/biosiliceous) sediments (Site 959; see Fig. 2). Site 960 on the crest of the Marginal Ridge was sediment starved compared to Site 959 on the shoulder of the Marginal Ridge, and its record of hiatuses (discussed below) is particularly interesting: during the Paleogene (and later) the Marginal Ridge acted as a passive margin—continually subsiding as a result of con-

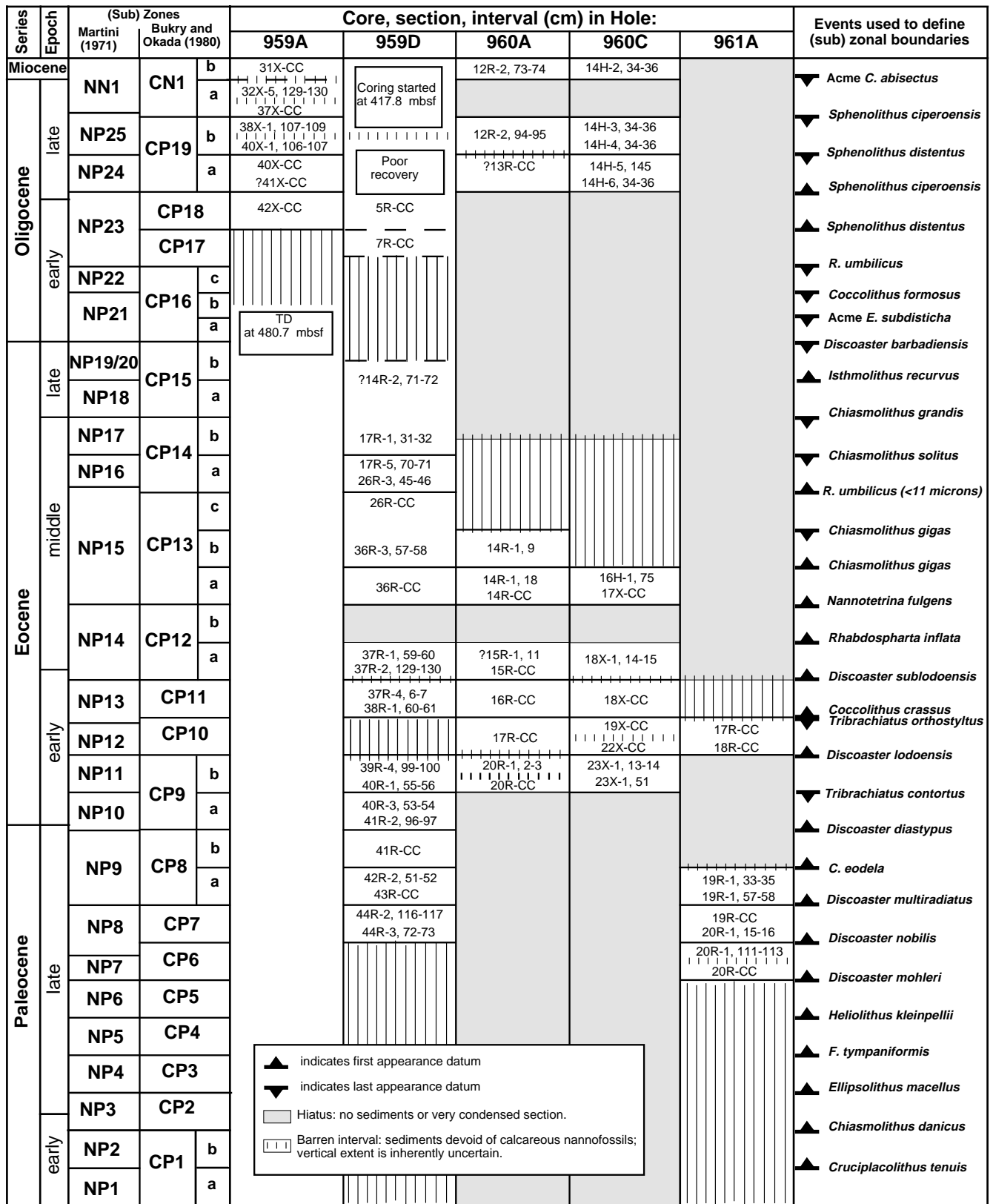


Figure 2. Summary of the Paleogene calcareous nannofossil biostratigraphy on the Côte d'Ivoire-Ghana Marginal Ridge, ODP Leg 159. The vertical extent of the barren middle Eocene sediments at Site 960 is based on a radiolarian fauna in the sediments in Hole 960C (see text). Also, the vertical extent of both the barren lower Eocene sediments in Hole 961A and the overlying hiatus are based primarily on radiolarian fauna in these and overlying sediments (see text).

low-latitude settings, largely affected by eustatic sea-level changes, and in places, by erosive bottom currents. The base of this section appears to become younger up the ridge: it is late Paleocene (CP6 Zone) on the lower slopes (Site 961), later Paleocene (CP7 Zone) on the shoulder of the ridge (Site 959), and early Eocene (Subzone CP9b) higher up on the crest of the ridge (Site 960). This section is discontinuous: on the crest of the ridge it is interrupted principally by hiatuses, whereas on the shoulder of the ridge it is segmented by mainly barren (noncalcareous) sediment packages. The hiatuses on the crest of the Marginal Ridge generally correspond in time with the barren sediments on the shoulder of the Marginal Ridge; however, at both locations the same earliest middle Eocene (CP12b) hiatus occurs, which is evidently widespread in the Atlantic. The timing of individual post-Paleocene hiatuses or barren sediment packages roughly coincides with eustatic falls in sea level, suggesting a close relationship.

Among the major boundaries limiting or subdividing the Paleogene section on the Côte d'Ivoire-Ghana Marginal Ridge, the Cretaceous/Tertiary and Eocene/Oligocene boundaries fall within unsuitable (barren) facies or hiatuses. Cores containing the Paleocene/Eocene and Oligocene/Miocene boundaries from the Marginal Ridge bear nannofossils. The former boundary is conveniently approximated by the highest occurrence of *Fasciculithus* spp.

The nannofossil biostratigraphy in the recently selected GSSP of the base of the Neogene (Lemme-Carrosio Section, Italy) is not yet clear, and its impact, if any, remains to be seen on the recognition of the Oligocene/Miocene boundary by means of nannofossils elsewhere. The end of the acme of *Cyclicargolithus abisectus* (seemingly unrecognizable in the Italian section) is used here for approximation of this boundary. Both *Sphenolithus delphix* and *S. capricornutus*, whose presence emerges from studies of the Lemme-Carrosio Section as possibly good evidence for the Oligocene/Miocene boundary, are already known from older sediments.

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APPENDIX

Calcareous Nannofossils Considered in This Chapter
(In Alphabetical Order of Generic Epithets)

Paleogene Taxa

- Bomolithus conicus* (Perch-Nielsen, 1971) Perch-Nielsen, 1984
Campylosphaera dela (Bramlette and Sullivan, 1961) Hay and Mohler, 1967
Campylosphaera eodela Bukry and Percival, 1971
Chiasmolithus bidens (Bramlette and Sullivan, 1961) Hay and Mohler, 1967
Chiasmolithus danicus (Brotzen, 1959) Hay and Mohler, 1967
Chiasmolithus californicus (Sullivan, 1961) Hay and Mohler, 1967
Chiasmolithus consuetus (Bramlette and Sullivan, 1961) Hay and Mohler, 1967
Chiasmolithus eograndis Perch-Nielsen, 1971
Chiasmolithus expansus (Bramlette and Sullivan, 1961) Gartner, 1970
Chiasmolithus gigas (Bramlette and Sullivan, 1961) Gartner, 1970
Chiasmolithus grandis (Bramlette and Riedel, 1954) Radomski, 1968
Chiasmolithus oamaruensis (Deflandre, 1954) Hay, Mohler and Wade, 1966
Chiasmolithus solitus (Bramlette and Sullivan, 1961) Locker, 1968
Chiasmolithus nitidus Gartner, 1970
Chiphrragmalithus acanthodes Bramlette and Sullivan, 1961
Chiphrragmalithus calathus Bramlette and Sullivan, 1961
Coccolithus crassus Bramlette and Sullivan, 1961
Coccolithus eopelagicus (Bramlette and Riedel, 1954) Bramlette and Sullivan, 1961
Coccolithus formosus (Kamptner, 1963) Wise, 1973
Coccolithus pelagicus (Wallich, 1877) Schiller, 1930
Coccolithus staurion Bramlette and Sullivan, 1961
Coronocyclus nitescens (Kamptner, 1963) Bramlette and Wilcoxon, 1967
Cruciplacolithus frequens (Perch-Nielsen, 1977) Romein, 1979
Cruciplacolithus tenuis (Stradner, 1961) Hay and Mohler, 1967
Cyclicargolithus abisectus (Müller, 1970) Wise, 1973
Cyclicargolithus floridanus (Roth and Hay in Hay et al., 1967) Bukry, 1971
Cyclicargolithus gammation (Bramlette and Sullivan, 1961) Shafik, 1990
Cyclicargolithus luminis (Sullivan, 1964) Bukry, 1971
Cyclicargolithus reticulatus (Gartner and Smith, 1967) Bukry, 1971
Dakylethra punctulata Gartner in Gartner and Bukry, 1969
Dictyococcites bisectus (Hay, Mohler and Wade, 1966) Bukry and Percival, 1971
Dictyococcites scrippsae Bukry and Percival, 1971
Discoaster barbadiensis Tan Sin Hok, 1929 emended Bramlette and Riedel, 1964
Discoaster bifax Bukry, 1971
Discoaster binodosus Martini, 1958
Discoaster calculosus Bukry, 1971
Discoaster cruciformis Martini, 1958
Discoaster deflandrei Bramlette and Riedel, 1964
Discoaster delicatus Bramlette and Sullivan, 1961
Discoaster diastypus Bramlette and Sullivan, 1961
Discoaster druggii Bramlette and Wilcoxon, 1967
Discoaster elegans Bramlette and Sullivan, 1961
Discoaster gemmifer Stradner, 1961
Discoaster kuepperi Stradner, 1959
Discoaster lenticularis Bramlette and Sullivan, 1961
Discoaster lidzi Hay in Hay et al., 1967
Discoaster lodoensis Bramlette and Riedel, 1964
Discoaster mirus Deflandre in Deflandre and Fert, 1954
Discoaster mohleri Bukry and Percival, 1971
Discoaster multiradiatus Bramlette and Riedel, 1964
Discoaster nephados Hay in Hay et al., 1967
Discoaster nobilis Martini, 1961
Discoaster nodifer (Bramlette and Riedel, 1954) Bukry, 1973
Discoaster praebifax Wei and Wise, 1989
Discoaster saipanensis Bramlette and Riedel, 1954
Discoaster salisburgensis Stradner, 1961
Discoaster saundersi Hay in Hay et al., 1967
Discoaster splendidus Martini, 1960
Discoaster strictus Stradner, 1961
Discoaster sublodoensis Bramlette and Sullivan, 1961
Discoaster tani Bramlette and Riedel, 1954
Discoaster wemmelensis Achutan and Stradner, 1969
Ellipsolithus distichus (Bramlette and Sullivan, 1961) Sullivan, 1964
Ellipsolithus lajollaensis Bukry and Percival, 1971
Ellipsolithus macellus (Bramlette and Sullivan, 1961) Sullivan, 1964
Ericsonia subdisticha (Roth and Hay in Hay et al., 1967) Roth in Baumann and Roth, 1969
Ericsonia robusta (Bramlette and Sullivan, 1961) Perch-Nielsen, 1977
Fasciculithus alanii Perch-Nielsen, 1971
Fasciculithus clinatus Bukry, 1971
Fasciculithus involutus Bramlette and Sullivan, 1961
Fasciculithus lilianae Perch-Nielsen, 1971
Fasciculithus richardii Perch-Nielsen, 1971
Fasciculithus tympaniformis Hay and Mohler in Hay et al., 1967
Helicosphaera dinesenii (Perch-Nielsen, 1971) Jafar and Martini, 1975
Helicosphaera euphratis Haq, 1966
Helicosphaera gartneri Theodoridis, 1984
Helicosphaera intermedia Martini, 1965
Helicosphaera lophota (Bramlette and Sullivan, 1961) Jafar and Martini, 1975
Helicosphaera obliqua Bramlette and Wilcoxon, 1967
Helicosphaera papillata (Bukry and Bramlette, 1969) Jafar and Martini, 1975
Helicosphaera recta (Haq, 1966) Jafar and Martini, 1975
Helicosphaera seminulum (Bramlette and Sullivan, 1961) Jafar and Martini, 1975
Helicosphaera truempyi Biolzi and Perch-Nielsen, 1982
Heliolithus cantabriae Perch-Nielsen, 1971
Heliolithus kleinpellii Sullivan, 1964
Heliolithus megastypus (Bramlette and Sullivan, 1961) Romein, 1979
Heliolithus riedelii Bramlette and Sullivan, 1961
Holodiscolithus Roth, 1970
Hornibrookina australis Edwards and Perch-Nielsen, 1975
Isthmolithus recurvus Deflandre, in Deflandre and Fert, 1954
Lophodolichus mochlophorus Deflandre in Deflandre and Fert, 1954
Lophodolichus nascens Bramlette and Sullivan, 1961
Markalius inversus (Deflandre in Deflandre and Fert, 1954) Bramlette and Martini, 1964
Micrantholithus Deflandre, 1950
Nannotetrina cristata (Martini, 1958) Perch-Nielsen, 1971
Nannotetrina fulgens (Stradner, 1960) Achutan and Stradner, 1969
Nannotetrina mexicana (Stradner, 1959) Bukry, 1973
Nannotetrina pappii (Stradner, 1959) Perch-Nielsen, 1971
Neochiastozygus junctus (Bramlette and Sullivan, 1961) Perch-Nielsen, 1971
Neococcolithes dubius (Deflandre, 1954) Black, 1967
Pemma Klumpp, 1953
Placozygus sigmoides (Bramlette and Sullivan, 1961) Romein, 1979
Pontosphaera Lohmann, 1902
Pontosphaera multipora (Kamptner, 1948) Roth, 1970
Pontosphaera pygmaea (Locker, 1967) Bystrická and Lehotayová, 1974
Pseudotriquetrorhabdulus inversus (Bukry and Bramlette, 1969) Bukry, 1981, emended Wise, 1983
Reticulofenestra dictyoda (Deflandre and Fert, 1954) Stradner in Stradner and Edwards, 1968
Reticulofenestra gartneri Roth and Hay in Hay et al., 1967
Reticulofenestra lockeri Müller, 1970
Reticulofenestra samodurovii (Hay, Mohler and Wade, 1966) Roth, 1970
Reticulofenestra umbilicus (Levin, 1965) Martini and Ritzkowski, 1968
Rhabdosphaera inflata Bramlette and Sullivan, 1961
Rhombosphaera bitrifida Romein, 1979
Sphenolithus anarrhophus Bukry and Bramlette, 1969
Sphenolithus capricornutus Bukry and Percival, 1971
Sphenolithus ciproensis Bramlette and Wilcoxon, 1967
Sphenolithus compactus Backman, 1980
Sphenolithus conicus Bukry, 1971
Sphenolithus delphix Bukry, 1973
Sphenolithus dissimilis Bukry and Percival, 1971
Sphenolithus distentus (Martini, 1965) Bramlette and Wilcoxon, 1967
Sphenolithus elongatus Perch-Nielsen, 1980
Sphenolithus furcatolithoides Locker, 1967
Sphenolithus moriformis (Brönnimann and Stradner, 1960) Bramlette and Wilcoxon, 1967
Sphenolithus orphanknollensis (Perch-Nielsen, 1971) Perch-Nielsen, 1985
Sphenolithus predistentus Bramlette and Wilcoxon, 1967
Sphenolithus primus Perch-Nielsen, 1971
Sphenolithus pseudoradians Bramlette and Wilcoxon, 1967
Sphenolithus radians Deflandre in Grassé, 1952
Sphenolithus spiniger Bukry, 1971
Toweius eminens (Bramlette and Sullivan, 1961) Gartner, 1971
Toweius magnicrassus (Bukry, 1971) Romein, 1979
Toweius pertusus (Sullivan, 1965) Romein, 1979
Toweius tovae Perch-Nielsen, 1971
Transversopontis Hay, Mohler and Wade, 1966

Transversopontis pulcher (Deflandre, in Deflandre and Fert, 1954) Hay, Mohler and Wade, 1966
Transversopontis pulcheroides (Sullivan, 1964) Báldi-Beke, 1971
Tribrachiatius bramlettei (Brönnimann and Stradner, 1960) Proto Decima et al., 1975
Tribrachiatius contortus (Stradner, 1958) Bukry, 1972
Tribrachiatius orthostylus (Bramlette and Riedel, 1954) Shamray, 1963
Triquetrorhabdulus auritus Stradner and Allram, 1982
Triquetrorhabdulus carinatus Martini, 1965

Trochoaster Klumpp, 1953
Zygodiscus herlynii Sullivan, 1964
Zygrhablithus Deflandre, 1959
Zygrhablithus bijugatus (Deflandre, in Deflandre and Fert, 1954) Deflandre, 1959

Reworked Cretaceous Taxa

Prediscosphaera cretacea (Arkhangelsky, 1912) Gartner, 1968

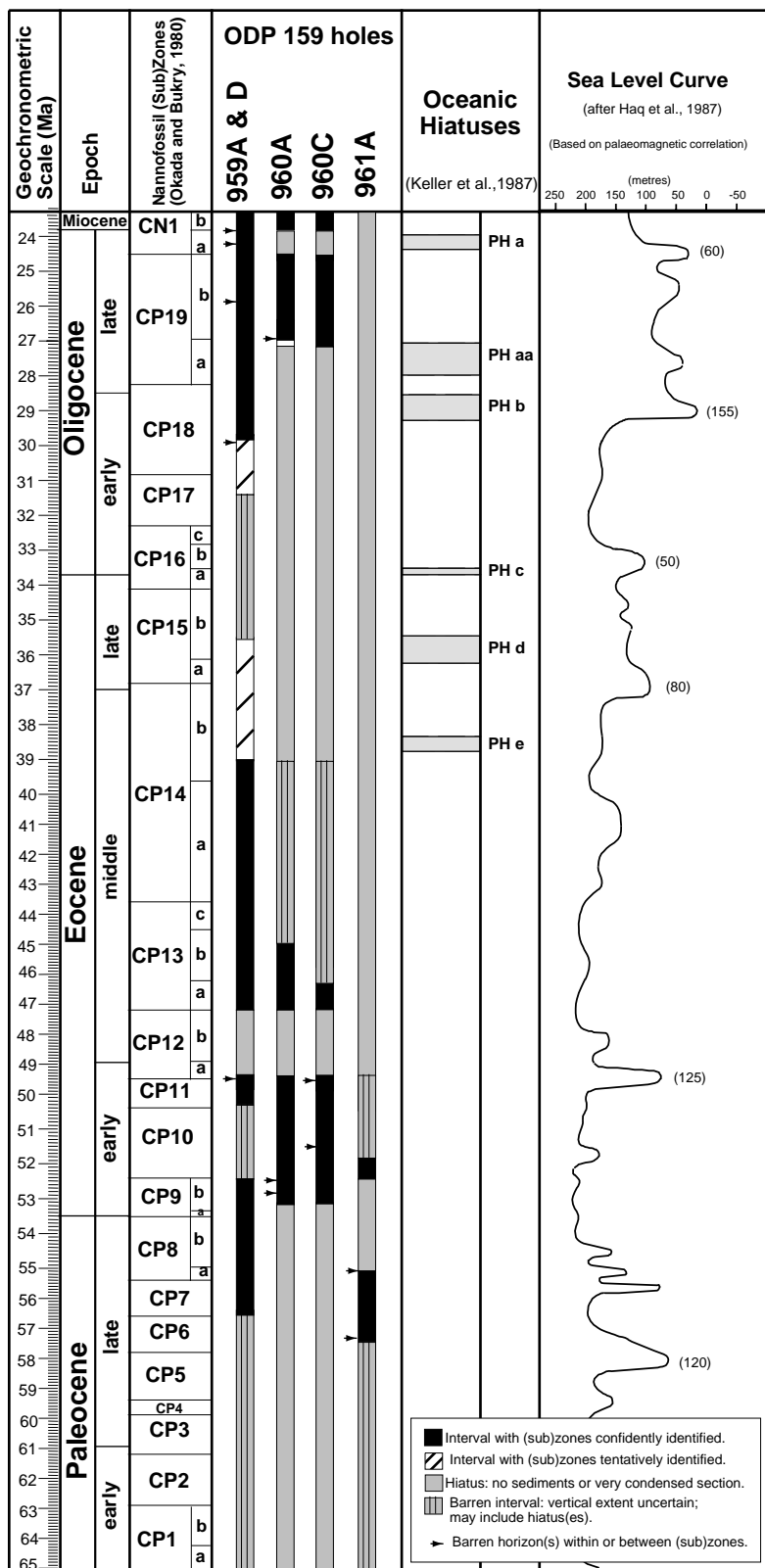


Figure 3. Sea-level curve, oceanic hiatuses and the Paleogene on the Côte d'Ivoire-Ghana Marginal Ridge, ODP Leg 159. In the column of oceanic hiatuses no data are available below the PHe hiatus. In the column of sea-level curve, figures between brackets are total sea level falls in meters. Note the strong correlation between the major drop of global sea level during the late early Eocene of ~125 m and the coeval hiatus at Sites 960 (on the crest of the Marginal Ridge) and 959 (on the shoulder of the Marginal Ridge), also the temporal match of the late Oligocene hiatus at Site 960, the PHa hiatus of Keller et al. (1987), and the drop of ~60 m in global sea level.

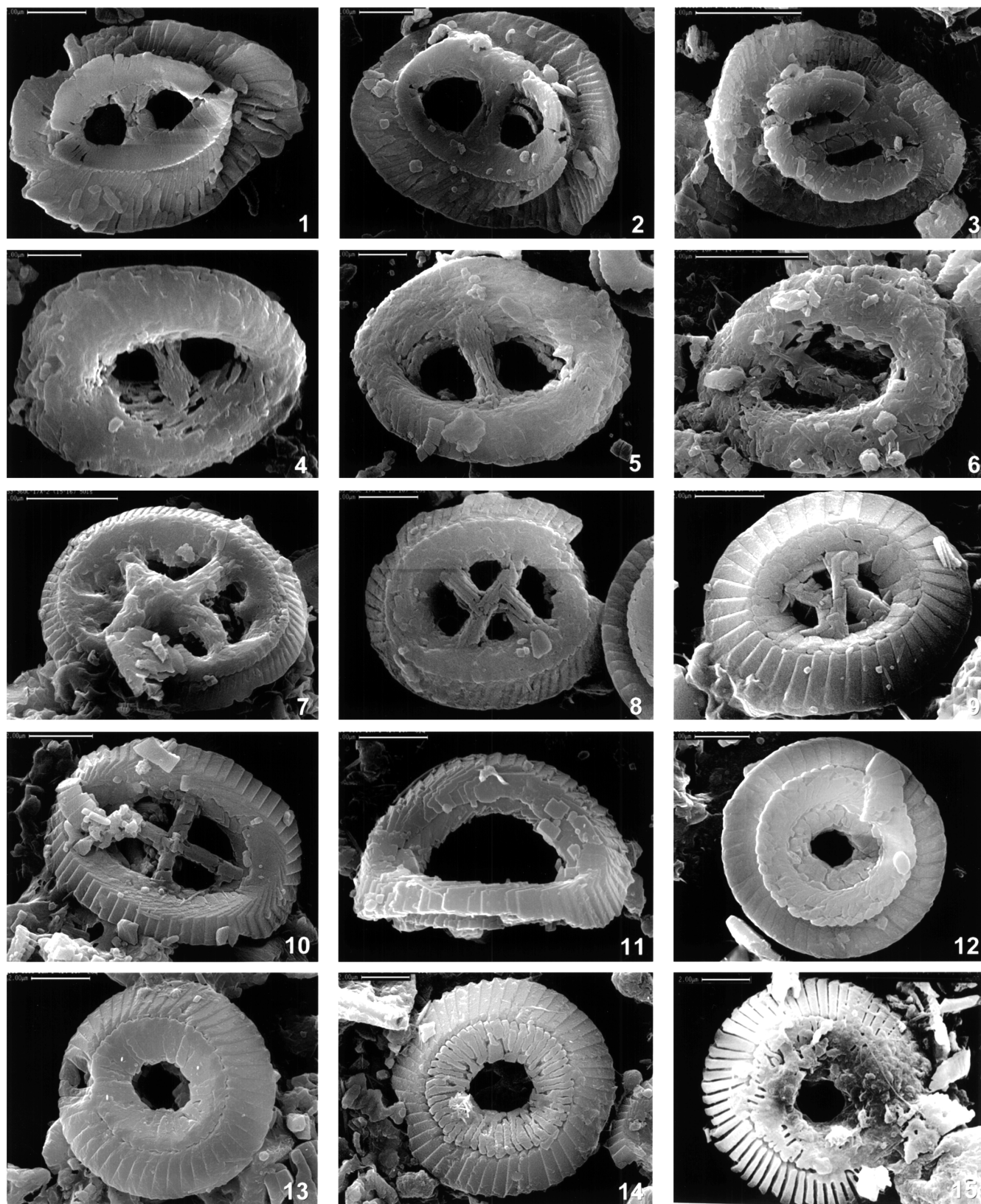


Plate 1. Eocene nanofossils from Site 260. **1, 2, 4, 5.** *Helicosphaera seminulum*, Sample 159-960C-17X-2, 15–16 cm, scale bar = 2 μ m: 1, 2. proximal views; 4, 5. distal views. **3, 6.** *Helicosphaera lophota*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 5 μ m: 3. proximal view; 6. distal view. **7.** *Chiasmolithus grandis* distal view, Sample 159-960C-18X-1, 14–15 cm, scale bar = 5 μ m. **8, 9.** *Chiasmolithus solitus* distal view, Sample 159-960C-17X-2, 15–16 cm, scale bar = 2 μ m. **9.** *Chiasmolithus consuetus* distal view, Sample 159-960C-17X-2, 15–16 cm, scale bar = 2 μ m. **10, 11.** *Campylosphaera dela* distal views, scale bar = 2 μ m: 10. Sample 159-960C-17X-2, 15–16 cm; 11. Sample 159-960C-18X-1, 14–15 cm. **12–15.** *Coccolithus formosus*, Sample 159-960C-18X-1, 14–15 cm; scale bar = 2 μ m: 12. proximal view; 13–15. distal views showing various degrees of dissolution.

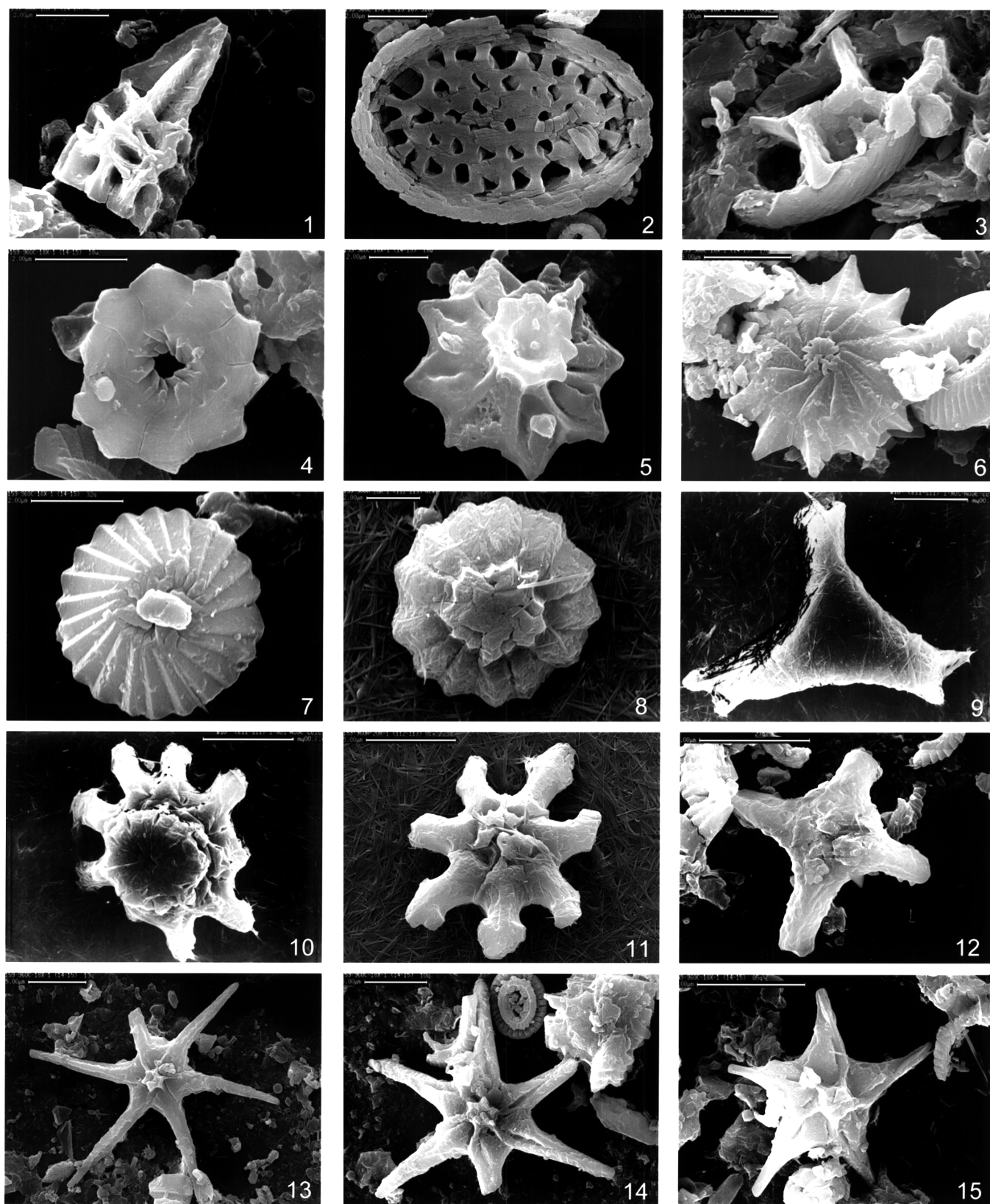


Plate 2. Eocene nannofossils from Site 260. **1.** *Sphenolithus radians*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 2 μ m. **2.** *Pontosphaera multipora* distal view, Sample 159-960C-17X-2, 15–16 cm, scale bar = 2 μ m. **3.** *Neococcolithus dubius* distal view, Sample 159-960C-17X-2, 15–16 cm, scale bar = 2 μ m. **4., 5.** *Discoaster kuepperi*, Sample 159-960C-18X-1, 14–15cm, scale bar = 2 μ m: 4. ?proximal view; 5. ?distal view. **6.** *Discoaster barbadiensis*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 5 μ m. **7.** *Discoaster lenticularis*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 2 μ m. **8.** *Discoaster* sp. aff. *D. mohleri*, Sample 159-960A-20R-1, 112–113 cm, scale bar = 2 μ m. **9.** *Tribrachiatus orthostylus*, Sample 159-960A-20R-1, 112–113 cm, scale bar = 5 μ m. **10., 11.** *Discoaster binodosus*, Sample 159-960A-20R-1, 112–113 cm, scale bar = 5 μ m. **12.** *Discoaster cruciformis*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 5 μ m. **13., 14.** *Discoaster lodoensis*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 5 μ m. **15.** *Discoaster* sp. aff. *D. sublodoensis*, Sample 159-960C-18X-1, 14–15 cm, scale bar = 2 μ m. Note: Sample 159-960A-20R-1, 112–113 cm came from the palygorskite claystone unit at the site.