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**NANOFÓSSEIS CALCÁRIOS DO ALBIANO–CENOMANIANO
NO DSDP *SITE* 364 (BACIA DE KWANZA - ANGOLA): BIOESTRATIGRAFIA E
IMPLICAÇÕES PALEOCEANOGRÁFICAS PARA O ATLÂNTICO SUL**

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Orientador: Prof. Dr. Gerson Fauth

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1 RESUMO

2 O intervalo Albiano–Cenomaniano foi um período de condições climáticas extremas
3 relacionadas a condições de *Greenhouse*, neste intervalo de tempo também são registradas
4 significativas mudanças na paleogeografia e paleoceanografia dos oceanos que causaram
5 evoluções nos ecossistemas marinhos em escala global. Neste estudo foi analisada a
6 assembleia de nanofósseis calcários das amostras recuperadas no *Site 364* (Bacia do Kwanza,
7 Angola), perfurado pelo *Deep Sea Drilling Project*. Foi recuperada uma assembleia com
8 afinidade subtropical-tropical, que fornece um excelente registro bioestratigráfico do intervalo
9 Albiano–Cenomaniano superior, e indica conexão de águas rasas entre as regiões sul e central
10 do Oceano Atlântico, pelo menos até Angola. Os dados mostram que a sedimentação foi
11 predominantemente calcária e pelágica durante o intervalo Albiano–Cenomaniano, exceto na
12 seção basal do *Site 364* onde os depósitos de folhelhos pretos ocorrem intercalados com
13 calcário dolomítico e margas. A dissolução do carbonato de cálcio é um processo
14 significativo registrado no *Site 364* e foi observado na maioria das amostras, seus efeitos são
15 seletivos na assembleia e não foi possível verificar a relação entre as camadas de deposição de
16 folhelhos pretos e a alta fertilidade de nanofósseis calcários. Apesar da dissolução, a
17 assembleia recuperada é relativamente bem diversificada e os táxons suscetíveis à dissolução
18 são observados, mas não muito abundantes. Evidências paleontológicas e geoquímicas
19 sugerem aumento da fertilidade/produktividade da água superficial nos intervalos onde diminui
20 os efeitos da dissolução.

22 APRESENTAÇÃO

23 Durante o Cretáceo ocorreram os estágios finais da fragmentação do Gondwana e no
24 desenvolvimento de oceanos, extensas plataformas carbonáticas e mares epicontinentais que
25 influenciaram na evolução da biota marinha. O intervalo Albiano–Cenomaniano é descrito na
26 literatura como a última expressão de fragmentação e separação das placas Sul-Americana e
27 Africana, portanto é um intervalo de tempo relevante na evolução geológica do Oceano
28 Atlântico. Para este intervalo de tempo, os dados litológicos, geoquímicos e bioestratigráficos
29 têm sido discutidos com o objetivo de caracterizar as condições paleoceanográficas do
30 Oceano Atlântico (e.g., Wagner & Pletsch, 1999; Moulin et al., 2010; Pérez-Díaz & Eagles,
31 2017).

32 Estudos indicam que durante o Albiano, a configuração do Oceano Atlântico seria de
33 mar aberto com livre circulação das massas de água oceânica, adequadas à sobrevivência e
34 distribuição de microrganismos marinhos. Entretanto, também tem sido discutida uma
35 configuração distinta, na qual este oceano teria sido um mar epicontinental com relativa
36 estabilidade tectônica entre as margens continentais da África e América do Sul. Diversos
37 trabalhos atribuem distintos momentos para a livre circulação das massas de água, nos estudos
38 modernos foi inferido que esta fase evolutiva do oceano teria ocorrido durante o intervalo
39 Aptiano–Cenomaniano (e.g., Kellogg & Mohriak, 2001; Azevedo, 2004; Eagles, 2007;
40 Moulin et al., 2010). Não existe um consenso sobre as idades/estágios de deposição e
41 afinidade paleobiogeográfica dos microrganismos marinhos, e a evolução paleoceanográfica
42 do Oceano Atlântico durante o Cretáceo ainda é um tema de amplo debate na comunidade
43 científica.

44 As sequências carbonáticas intercaladas com camadas ricas em matéria orgânica,
45 recuperadas no *Site* 364 perfurado pelo *Deep Sea Drilling Project* (DSDP) na Bacia de
46 Kwanza em Angola, possuem registros das fases evolutivas do Oceano Atlântico. O estudo
47 dos nanofósseis calcários, na seção relacionada ao Cretáceo Inferior do *Site* 364, infere
48 datações relativas para os estratos e apresenta evidências relacionadas a eventos
49 paleoceanográficos. Os índices paleoecológicos deste grupo fóssil foram analisados
50 conjuntamente aos dados geoquímicos de isótopos estáveis de carbono e oxigênio,
51 susceptibilidade magnética, carbono orgânico e carbonato de cálcio com o intuito de
52 reconhecer e entender os eventos paleoceanográficos registrados na seção sedimentar
53 estudada.

55 A hipótese testada neste estudo é a seguinte:

56 (i) O estudo da assembleia de nanofósseis calcários no *Site* 364 pode refinar o
57 posicionamento bioestratigráfico dos estratos, contribuir no entendimento da afinidade
58 paleobiogeográfica deste grupo fóssil e fornecer indícios sobre os eventos paleoceanográficos
59 ocorridos durante o Cretáceo Inferior no Oceano Atlântico.

60 O artigo elaborado foi submetido ao periódico *Paleoceanography and*
61 *Paleoclimatology*, conceito A1 na plataforma Qualis da Capes, que incentiva a publicação de
62 resultados em temas como a bioestratigrafia e a paleoceanografia.

66 **Albian–Cenomanian calcareous nannofossils from DSDP Site 364 (Kwanza Basin -**
67 **Angola): biostratigraphic and paleoceanographic implications for the South Atlantic**

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77 **Key Points:**

- 78 • Paleontological evidence suggests a surface water connection between the Central and
79 South Atlantic oceans during the Albian–Cenomanian
- 80 • Calcareous nannofossil abundance and geochemical signals indicate dissolution,
81 although it did not heavily affect species richness

82 **Abstract**

83 The Albian–Cenomanian was an interval of extreme warmth due to greenhouse climatic
84 conditions, and in which are registered significant changes in the paleogeography and
85 paleoceanography of the oceans, which affected the evolution of marine ecosystems on a
86 global scale. This study analyzed the calcareous nannofossil assemblages from samples
87 recovered from Site 364 (Kwanza Basin, Angola), drilled by the Deep Sea Drilling Project
88 (DSDP) Leg 40. The assemblages were observed to have a subtropical-tropical affinity, which
89 could provide an excellent biostratigraphic record of the Albian–Cenomanian interval, and are
90 indicative of a surface water connection between the Central Atlantic and the South Atlantic
91 oceans at least up to offshore Angola. Sedimentation for the area was predominantly
92 calcareous and pelagic during this interval, with the exception of the basal section, where
93 Albian black shale layers interbedded with dolomitic and marly limestone can be seen.

94 Dissolution is a significant process in Site 364 that can be observed in the majority of the
95 studied samples, though not to the point where species richness was severely impacted, as
96 small amounts of dissolution-susceptible taxa were observed. Dissolution showed a marked
97 decrease in intervals in which paleontological and geochemical data indicated an increase in
98 surface water fertility/productivity. Still, it affected assemblage composition in a way that did
99 not allow for a proper consideration of the relationship between the deposition of black shale
100 layers and the high fertility of calcareous nannofossils.

101 **1. Introduction**

102 Fragmentation of the Gondwana paleocontinent resulted in significant changes of the
103 global paleogeographic configuration, leading to the opening and development of the South
104 Atlantic ocean. This process produced various submarine physiographic features, such as
105 sedimentary basins and plateaus. The stratigraphic evolution of the sedimentary basins in the
106 South Atlantic ocean follows a well-defined pattern, beginning with continental sequences
107 that gradually evolve into marine ones (e.g., Kellogg & Mohriak, 2001; Milani et al., 2007;
108 Beglinger et al., 2012). In the Early Cretaceous, the first marine incursions were recorded by
109 the deposition of evaporites in many sedimentary basins of the Brazilian and African
110 continental margins. Sediments deposited above evaporites mark the moment of transition
111 from a restricted to an open marine environment phase, allowing for the connection between
112 the Central Atlantic and the South Atlantic oceans (Azevedo, 2004; Eagles, 2007; Moulin et
113 al., 2010). During the Albian–Cenomanian the South Atlantic ocean already had open marine
114 conditions, evidenced mainly by the diversity of marine fossils and extensive limestone layers
115 deposited in shallow marine carbonate shelves (Melguen, 1978; Wagner & Pletsch, 1999;
116 Mohriak et al., 2008; Beglinger et al., 2012).

117 The Albian–Cenomanian was a key period for the evolution of marine ecosystems. It
118 was an interval of high atmospheric CO₂ content related to greenhouse climatic conditions, it
119 marked an increase in the eustatic sea-level, and it registered numerous changes in the
120 paleogeography and paleoceanography of the oceans (Poulsen et al., 1999, 2001; Leckie et al.,
121 2002; Huber et al., 2002; Hay, 2008; Torsvik et al., 2009; Haq, 2014). In this global context,
122 numerous episodes of anoxia and dysoxia were registered in the oceans, resulting in the
123 deposition and preservation of organic carbon-rich sediments, black shales, associated with
124 Ocean Anoxic Events (OAEs; Schlanger & Jenkyns, 1976; Arthur & Schlanger, 1979;
125 Erbacher et al., 1996; Leckie et al., 2002; Watkins et al., 2005; Jenkyns, 2010; Wang et al.,

126 2011). During these episodes changes in the physicochemical conditions in the water column
127 were responsible for the enhanced fertility and dissolution of specific calcareous nannofossil
128 species, selectively affecting the planktonic communities (e.g., Roth & Bowdler, 1981;
129 Stradner et al., 1984; Fisher & Hay, 1999; Herrle et al., 2003; Watkins et al., 2005).

130 The geochemical composition of carbonate rocks records global marine signals, and
131 their correlation with paleoecological data from calcareous nannofossils can provide a better
132 understanding of the paleoceanographic changes that occurred in the South Atlantic ocean
133 during the Albian–Cenomanian. One such calcareous nannofossil record was analyzed from
134 the Kwanza Basin (Deep Sea Drilling Project - DSDP) Site 364 from Leg 40. The
135 paleontological record from this site exhibits paleobiogeographic and biostratigraphic
136 discrepancies between its Lower Cretaceous calcareous nannofossil, benthic and planktonic
137 foraminifera assemblages (Bolli et al., 1978; Kochhann et al., 2013, 2014). The main
138 objectives of this paper are to restudy the calcareous nannofossil assemblages recovered from
139 the Albian–Cenomanian of DSDP Site 364 and their biostratigraphy, to complement the data
140 reported during Leg 40 by Proto Decima et al. (1978). The present paleoenvironmental study
141 of this interval also uses the analysis of micropaleontological data, total organic carbon
142 (TOC), calcium carbonate content (CaCO_3), magnetic susceptibility (MS) and stable isotopes
143 ($\delta^{18}\text{O}$ and $\delta^{13}\text{C}$) values.

144 **2. Materials and Methods**

145 **2.1 Site Information**

146 This study analyzed Site 364 (11°34.32'S, 11°58.30'E, 2448 m water depth), which
147 was drilled during the DSDP Leg 40 in the Kwanza Basin, Angola Continental Margin
148 (Figure 1). The studied section comprises the interval from cores 42 to 26 (between 1032.37–
149 715.34 mbsf). During the deposition of this interval, Site 364 was located at a low latitude
150 (24°S to 28°S), with an estimated paleodepth of 1000 m (Roth & Bowdler, 1981; Roth &
151 Krumbach, 1986). Benthic foraminifera assemblages indicate that this interval was deposited
152 at upper bathyal to shallow neritic depths (Holbourn et al., 2001; Kochhann et al., 2014),
153 which is in agreement with recent paleobathymetric models that estimate a depth between
154 neritic to bathyal for the Angola Continental Margin (Pérez-Díaz & Eagles, 2017).

155 Two sedimentary units are described for the studied section: units 7 and 6, from
156 bottom to top (Bolli et al., 1978). Unit 7 is characterized by the occurrence of black shale, and

157 is divided into two subunits (7b and 7a): subunit 7b consists of dolomitic limestone and black
158 shale (1032.37-1009.79 mbsf interval samples); subunit 7a is composed of marly limestone
159 interbedded with black shales (1008.41-968.45 mbsf interval samples). Unit 6 is characterized
160 by the increased occurrence of limestones, and is divided into three subunits (6c, 6b and 6a):
161 subunit 6c is composed of limestone and marly limestone (955.75-825.73 mbsf interval
162 samples); subunit 6b consists of limestone (810.94-768.19 mbsf interval samples); and the
163 subunit 6a contain marly nannofossil chalk, observed between 753.64-717.32 mbsf interval
164 samples (Figure 2).

165 **2.2 Sample Preparations and Analysis**

166 For the study of calcareous nannofossil, 72 smear slides from Site 364 were prepared,
167 following the double slurry method detailed by Watkins and Bergen (2003) and Blair and
168 Watkins (2009). They were analyzed to determinate qualitative preservation, quantitative
169 species abundances, and biostratigraphic ranges. One sample per section was taken, with an
170 average spacing of 1.5 m within the cores. They were examined using a Zeiss Axio Imager
171 A2 microscope, at approximately 1000× magnification. The smear slides are stored in the
172 collection of the Museum of Geological History of Rio Grande do Sul, Unisinos University,
173 Brazil, under the curatorial numbers ULVG 12337 to ULVG 12409.

174 Preservation of calcareous nannofossils was evaluated using qualitative criteria to
175 assess the degree of etching (E) and/or overgrowth (O), where E1 or O1 are relatively good –
176 specimens exhibited little or no dissolution and/or overgrowth; E2 or O2 are moderate -
177 specimens exhibited moderate dissolution and/or overgrowth, and were easily recognizable;
178 and E3 or O3 are poor - specimens exhibited extreme dissolution and/or overgrowth (Roth &
179 Thierstein, 1972; Roth, 1983). For the quantitative analysis, each slide was randomly scanned
180 until 456 specimens had been counted; this number assures accurate abundance estimates at
181 the 95% confidence interval (Chang, 1967; Watkins & Bergen, 2003). These results are
182 shown in the distribution chart (see supporting information Table S1). The biostratigraphic
183 analysis was based on the First appearance datum (FAD) and Last appearance datum (LAD)
184 of markers species, following CC zones of Sissingh (1977) and Perch-Nielsen (1979, 1985).
185 Comparison of these zones to the BC zonation of Bown et al., (1998) and the UC of Burnett et
186 al., (1998), which are shown plotted next to the CC zones.

187 Analyzes of TOC, CaCO₃, MS, $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ were conducted on a number of
188 samples from DSDP Site 364 (supporting information Table S2). TOC and CaCO₃

189 measurements were performed from 51 samples using 0.26 g. For TOC, these sediments were
 190 manually fragmented and acidified using HCl 6N (1:1) for 24 hours. These were determined
 191 through the method described by Huerta-Diaz et al. (2011), using the Leco Elemental
 192 Analyzer model SC-144DR of the Technological Institute of Micropaleontology itt Fossil
 193 (Unisinos University). MS measurements of 49 samples were analyzed and normalized by
 194 mass. These values were obtained in low ($\chi_{lf} = 976$ Hz) and high ($\chi_{hf} = 15\ 616$ Hz)
 195 frequencies, with a MFK1-FA Multifunction Kappabridge magnetic susceptibility analyzer, at
 196 Paleomagnetism Laboratory (USPMag), University of São Paulo. This method resulted in the
 197 evaluation of the concentration of ultrafine ($<0.03\ \mu\text{m}$) superparamagnetic (SP) ferromagnetic
 198 minerals in transition to stable single domain (Bloemendal et al., 1985; Dearing et al., 1996;
 199 Hrouda & Pokorný, 2012). We also calculated the percentage frequency-dependent magnetic
 200 susceptibility χ_{fd} % with the Equation:

$$201 \quad \chi_{fd} \% = (\chi_{lf} - \chi_{hf})/\chi_{lf} \times 100.$$

202 Thirty-one samples were measurement for the $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ on bulk sediments; these
 203 analyses were published by Kochhann et al. (2013), with the isotopic values having been
 204 reported in standard delta notation (δ) relative to the Vienna Pee Dee Belemnite (VPDB).

205 **2.3 Paleocological Analysis**

206 Species and genera of calcareous nannofossils relevant for paleoenvironmental
 207 interpretations were selected based on several studies that highlighted their quantitative and
 208 qualitative relevance in the analysis of Cretaceous paleoceanographic events (Table 1). In
 209 several of these studies, it was interpreted that shifts in the faunal composition of the
 210 assemblages are related to relative changes in surface water fertility, being controlled mainly
 211 by the input of nutrients in the paleoenvironment. These studies also provide information
 212 regarding the diagenesis of the analyzed strata, noting that some species appear to be
 213 particularly resistant to dissolution.

214 For the paleocological analysis, different statistical data-processing techniques were
 215 used, including Pearson correlation coefficient (r), species richness (S), equitability (E) and
 216 Shannon Diversity Index (H). The software package PAST – Paleontological Statistics
 217 (Hammer et al., 2001) was utilized. Pearson correlation is used to quantify the extent to which
 218 two variables are linearly related, with the result being a coefficient (r) between -1 and 1
 219 (Fisher & Hay, 1999). Species richness (S) is simply the number of different species in an
 220 assemblage, which is a rough measure of the relative stability of the ecological conditions

221 when more species occur (Watkins, 1989). Equitability (E) is a measure of the proportional
 222 distribution of species, used to reflect the distribution of the energy flux through the
 223 ecosystem, larger values indicate relatively more stable conditions (Watkins, 1989). The
 224 Shannon Diversity Index (H) is an unbiased measure of diversity based on both species
 225 richness and equitability (Pielou, 1969). For the study of surface water fertility/productivity,
 226 were analyzed the Nutrients Index (NI), based on the studies of Herrle et al., (2003) and
 227 Watkins et al., (2005). This index was established through the relative abundance of two high-
 228 fertility taxa, and *W. barnesiae* abundant under low-fertility conditions. The Nutrients Index
 229 (NI) per sample was established by the relative abundances of two high-fertility taxa
 230 (*Discorhabdus ignotus* and *Biscutum* spp.) and *W. barnesiae* (abundant under low-fertility
 231 conditions), with the formula:

$$232 \quad \text{NI} = [(D_i + B_i) / (D_i + B_i + W_b)] 100$$

233 where $D_i = D. \textit{ignotus}$, $B_i = Biscutum \textit{ spp.}$, and $W_b = W. \textit{barnesiae}$. The specie
 234 *Zeugrhabdotus erectus*, commonly used in NI calculations (Herrle et al., 2003; Watkins et al.,
 235 2005), due to being indicative of high-fertility (Table 1), has a low occurrence in the Site 364
 236 (abundance of 1 %, with an average abundance value of 0.05 %) and was not used in
 237 constructing the NI.

238 3. Results

239 A total of 103 taxa were identified in the 72 studied samples (core 42–26). Calcareous
 240 nannofossil biostratigraphy indicates that the section spans the Albian–Cenomanian (Figure
 241 2). In the basal section of Site 364, between 1032.37–772.34 mbsf (Cores 42–29), the Albian
 242 can be inferred by the *Prediscosphaera columnata* Zone (CC 8), defined at the base by the
 243 occurrence of *P. columnata*, and by the FAD of *Eiffellithus turriseiffelii* at the top. Perch-
 244 Nielsen (1979) used the FAD *Tranolithus orionatus* to subdivide this biozone, with the
 245 stratigraphic interval 1032.37–1008.41 mbsf being defined at the base by the occurrence of *P.*
 246 *columnata* and at the top by the FAD *T. orionatus* (CC 8a subzone, equivalent to the BC 23
 247 zone of Bown et al., 1998), and the stratigraphic interval (1008.41–772.34 mbsf) being
 248 assigned to the CC 8b subzone, between the FAD *T. orionatus* and the FAD *E. turriseiffelii*.
 249 Bown et al., (1998) and many later authors having used the FADs of *Axopodorhabdus*
 250 *biramiculatus* and *Eiffellithus monechiae* to subdivide CC 8b; the stratigraphic interval
 251 1008.41–951.47 mbsf is defined at the base by the FAD *T. orionatus* and at the top by FAD *A.*
 252 *biramiculatus* (Zone BC 24); the interval 951.47–875.96 mbsf was assigned to Zone BC 25

253 defined at the top by the FAD *E. monechiae*; and the interval 875.96–772.34 mbsf is defined
254 by the FAD *E. turriseiffelii* (Zone BC 26). The late Albian to early Cenomanian interval can
255 be inferred by the *Eiffellithus turriseiffelii* Zone (CC 9), identified in the 772.34–726.66 mbsf
256 interval. This zone is defined in this study by the interval from the FAD of *E. turriseiffelii* at
257 the base and by the LAD of *Watznaueria britannica*. The specie *Microrhabdus decoratus* that
258 normally marks the top of this zone was not found. Perch-Nielsen (1979) used the LADs of
259 *Braarudosphaera africana* and *W. britannica* to subdivide this biozone, with the LAD of *B.*
260 *africana* having been identified at 752.37 mbsf and the LAD of *W. britannica* at 726.66 mbsf.
261 The top section of Site 364, ranging from 725.24 to 717.23 mbsf can be inferred as being of
262 late Cenomanian age, based on the consistent occurring of *Cretarhabdus striatus*, *Eiffellithus*
263 *perch-nielseniae*, *Rhagodiscus asper* and *A. biramiculatus*, which are characteristic of the
264 subzone CC 10a or zone UC1b/UC 4a. No evidence of reworking was observed in the species
265 that are considered zonal markers.

266 Only one (912.02 mbsf, core 36) of the 72 studies samples displayed low nannofossil
267 abundance (297 specimens), being which led to it being excluded from the paleoecological
268 analyses. Assemblages are dominated by *Watznaueria barnesiae*, with an abundance range of
269 27–94 %, and an average abundance value of 64 %. The remaining 102 taxa recognized have
270 averages less than 10 %. The preservation of calcareous nannofossil is variable throughout
271 Site 364, with indications of dissolution and recrystallization can be analyzed by means of
272 lithological, micropaleontological and geochemical data.

273 The basal strata of Site 364, ranging between 1032.37 and 968.45 mbsf (cores 42–39,
274 with 21 samples), is referred to as sedimentary unit 7 (Bolli et al., 1978), and is marked by the
275 deposition of dolomitic limestone with black shale intervals (7b subunit) and marly limestone
276 interbedded with black shale layers (7a subunit). In general, the preservation of calcareous
277 nannofossil in sedimentary unit 7 is moderate, with minor overgrowth (O1) and minor to
278 moderate etching (E1-E2), with only three samples (988.42, 987.42 and 975.49 mbsf)
279 showing slightly increased overgrowth (O2) (supporting information Table S1). For
280 sedimentary unit 7, the average species richness is 31 taxa, with a minimum of 15 and a
281 maximum of 44. Shannon Diversity Index values vary from 0.37–2.2, with an average value
282 of 1.42. For Equitability, the average value is 0.41, with values between 0.14–0.60 (Figure 3).
283 The TOC values show the highest values in this interval, ranging from 0.40–8.91 %, with an
284 average value of 3 %. The CaCO₃ varies from 11.08–75.26 %, with an average value of 45.40
285 % (Figure 4). *W. barnesiae* shows high-abundance in this sedimentary unit, comprising

286 between 53–94 % of the assemblage, with an average value of 71 %. On the other hand, the
 287 NI shows the lowest values in this interval ranging from 1–13 and with an average value of 5
 288 (Figure 3). Regarding isotopic measurements, the $\delta^{18}\text{O}$ values vary from -5.47 to -3.05 ‰,
 289 with a mean value of about -4.52 ‰; and the $\delta^{13}\text{C}$ values vary between -2.59 to 1.36 ‰,
 290 averaging at -0.27 ‰ (Figure 4). The analyzed data shows a positive correlation ($r = 0.32$, $p =$
 291 0.53) between $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ stable isotopes to basal strata of Site 364. Results obtained for
 292 MS show consistently low values for sedimentary unit 7, ranging between $1.15\text{--}4.01 \times 10^{-8}$
 293 m^3/kg in low frequency and between $1.02\text{--}3.75 \times 10^{-8}$ m^3/kg in high frequency; which χ_{fd}
 294 varying from -3.22 to 14.56 % (supporting information Table S3).

295 The interval 955.75–717.23 mbsf (cores 38–26, with 51 samples) is referred to as
 296 sedimentary unit 6, and is characterized by a high occurrence of limestones. In this unit, the
 297 calcareous nannofossil assemblage shows preservation variation between minor to moderate
 298 overgrowth (O1-O2) and etching (E1-E2) (supporting information Table S1). This
 299 assemblage is dominated by *W. barnesiae*, which comprises between 27–90 %, with an
 300 average value of 61 %, occurring in low percentages (<40 %) in only five samples (Table 2).
 301 In these five samples, the highest values of NI from studied interval were observed, varying
 302 from 1–67 and with an average value of 17 (Figure 3). This may be related to the relative
 303 abundance of *D. ignotus* (which reaches values of 20 %), and of species of the genus
 304 *Biscutum*, grouped in *Biscutum* spp., reaching 58 % (Table 2). Several studies related the high
 305 abundance of these two taxa as indicators of high carbonate productivity and surface water
 306 fertility, with a preference for nearshore environments (Table 1).

307 For sedimentary unit 6, the average species richness (S) is 33 taxa, varying between 20
 308 and 43. Shannon Diversity Index values vary from 0.62–2.77, with an average value of 1.71.
 309 For Equitability, the average value is 0.49, varying between 0.19–0.77 (Figure 3). TOC values
 310 show a considerable decrease in sedimentary unit 6, varying between 0.55 to 2.89 % and with
 311 an average value of 0.3 % (Figure 4). Sample 894.99 mbsf (core 35) is an exception, with a
 312 value of 2.89 % (Figure 4). On the other hand, CaCO_3 content shows high-values in this
 313 sedimentary unit, ranging between 17.45 and 90.64 % with an average value of 64.22 %. MS
 314 data shows a significant increase when compared to sedimentary unit 7, with values between
 315 1.14×10^{-8} and 7.12×10^{-7} m^3/kg in low frequency; and between 1.04×10^{-8} – 6.91×10^{-7} m^3/kg
 316 in high frequency (Figure 4). The highest values of MS were observed in the interval between
 317 845.92–844.57 mbsf (Core 33), which χ_{fd} varying from 1.35 to 13.27 %. The $\delta^{18}\text{O}$ isotope
 318 values vary between -4.49 and -2.68 ‰, with an average of -3.55 ‰; and the $\delta^{13}\text{C}$ isotope

319 values vary between 1.06–2.94 ‰, with an average value of 2.03 ‰ (supporting information
320 Table S3).

321 4. Discussion

322 4.1 Biostratigraphy

323 The study of the calcareous nannofossil recovered from Site 364 (core 42–26),
324 allowed for the classification of 103 taxa, as well as the identification of three intervals:
325 Albian, late Albian–early Cenomanian and middle–late Cenomanian. Assemblage containing
326 *Prediscosphaera columnata*, *Hayesites albiensis*, *Tranolithus orionatus*, *Axopodorhabdus*
327 *biramiculatus* and *Eiffellithus monechiae* was observed, indicating the presence of zones CC
328 8 or BC 23 to BC 26, of Albian age. The stratigraphic interval between the FAD of
329 *Eiffellithus turriseiffelii* and LAD of *Watznaueria britannica* was attributed to Zones CC 9 or
330 UC 0/UC 1a, of late Albian–early Cenomanian. The middle–late Cenomanian can be inferred
331 due to the consistent occurrences of *Cretarhabdus striatus*, *Eiffellithus perch-nielseniae*,
332 *Rhagodiscus asper* and *A. biramiculatus*, which are indicative of zones CC 10a or UC 1b/UC
333 4a (Figure 2). No evidence of reworking for species that are considered zonal markers was
334 observed, and all zonal marker-species are cosmopolitan. The most conspicuous difference
335 between the biostratigraphic zonation reported by Proto Decima (1978), for DSDP Site 364,
336 and the present study is the identification of the middle–late Cenomanian.

337 The calcareous nannofossil biostratigraphy analysis registered a continuous sequence
338 of Albian bioevents in the interval between cores 42–29, with the biozones CC 8 or BC 23,
339 BC 24, BC 25 and BC 26 having been identified (Figure 2). The Albian can also be inferred
340 by ammonites (Wiedmann & Neugebauer, 1978) and palynomorphs (Morgan, 1978). On the
341 other hand, the biostratigraphic study of Caron (1978) with planktonic foraminifera indicates
342 a late Aptian between the cores 41–35, early-middle Albian between 35–31 cores, and
343 middle-late Albian to 31–29 cores interval. Kochhann et al., (2013) studied this fossil group
344 and inferred the late Aptian for cores 37 to 33, also suggesting a biostratigraphic
345 unconformity within core 31 that separates the early Albian (between cores 33–31) and late
346 Albian (between cores 31–29). Scheibnerová (1978) identified benthic foraminifera that
347 assigned a late Aptian age for cores 41–38, early-middle Albian for cores 35–31, and late
348 Albian to cores 31–29.

349 For the intervals inferred by calcareous nannofossils as being late Albian–early
350 Cenomanian (cores between 29-27) and middle–late Cenomanian (cores between 27-26),
351 Caron (1978) registered, with planktonic foraminifera, the middle–late Albian between cores
352 34-31. However, studies with ammonites (Wiedmann & Neugebauer, 1978), palynomorphs
353 (Morgan, 1978), benthonic foraminifera (Scheibnerová, 1978) and planktonic foraminifera
354 (Kochhann et al., 2013) indicate a late Albian age for this stratigraphic interval. According to
355 Morgan (1978), radiolarians are rare, with poor preservation, and tentatively indicate an
356 Aptian–Albian age.

357 **4.2 Calcareous Nannofossil Preservation, Paleoecology and Paleoceanographic** 358 **Inferences**

359 New data regarding the strata deposited during the Albian–Cenomanian of Site 364
360 was acquired in this study. From the total of 72 analyzed samples, only sample 912.02 mbsf
361 showed low species abundance (297 specimens), possibly due to marly lithological
362 composition, which led to being excluded from the paleoecological inferences to avoid data
363 interpretation errors. The assemblages are mostly dominated by *Watznaueria barnesiae*;
364 several studies noted that a high abundance of *W. barnesiae* is indicative of dissolution,
365 assemblages that contain an abundance of this taxon exceeding 40% are considered strongly
366 altered to the extent that the fossil assemblage no longer reflect the biocoenosis (Thierstein,
367 1981; Roth & Bowdler, 1981; Roth, 1981, 1986; Roth & Krumbach, 1986). The effects of
368 dissolution may be selective, leading to an increase in the percentage of *W. barnesiae* and, at
369 the same time, reducing the overall abundance and richness of the assemblage. Dissolution
370 can also be inferred by low species diversity (Erba, 1992a; Luciani et al., 2001), and the
371 increase in the relative abundance of high-fertility species may result in a lower abundance of
372 *W. barnesiae* (Roth & Krumbach, 1986; Roth, 1989). Moreover, the high abundance of this
373 specie may be related to deep basin deposits and low-fertility conditions (Table 1). The
374 analyzed data shows a high negative correlation ($r = -0.87$) between the Shannon Diversity
375 Index and a high percentage of *W. barnesiae*, suggesting that the assemblage suffered from
376 dissolution (Figure 5). In the 72 studied samples, *W. barnesiae* distribution shows an
377 abundance of more than 40 % in 67 samples, suggesting that dissolution affected the
378 calcareous nannofossil assemblage of Site 364. The magnetic susceptibility values can also be
379 used as indicative of an increase in dissolution, as it would contribute to a higher
380 concentration of magnetic particles. The χ variation in marine deposits can be due to the
381 changes in the concentration of magnetic minerals, which include diamagnetic (for example

382 calcite), paramagnetic (for example clay) and ferromagnetic (for example magnetite). These
383 variations reflect the balance between terrigenous influxes (high MS, Figure 4) and/or
384 carbonate dissolution (low MS, Figure 4). The χ also can represent changes in magnetic
385 mineralogy, which might be related to redox conditions on the seafloor – magnetite vs.
386 hematite – or it might be related to the nature of the weathering in the continental source area
387 (Liu et al., 2012). The susceptibility was calculated on a carbonate-free based (CFB) in order
388 to account for dilution by carbonate (Figure 4, red lines), which is visible the difference
389 between black and red lines. Large values of $\chi_{fd}\%$ (Figure 4) are indicative of a significant
390 proportion of grains near the single domain (SD)/superparamagnetic (SP) boundary
391 (Bloemendal et al., 1985; Thompson & Oldfield, 1986). Chemical reactions that occur during
392 dissolution can be an important determinant of the MS signal in marine sediments (Liu et al.,
393 2012). The results show an inverse relationship between MS and CaCO_3 in the dissolution
394 intervals, which suggest an indicative of climatic controlled dilution of the magnetic signal
395 (e.g., Thompson & Oldfield, 1986; Robinson, 1986; Bloemendal et al., 1988, 1992;
396 Meynadier et al., 1995; Lean & McCave, 1998; Haag, 2000, Lopes et al., 2017).

397 In spite of the abundance of *W. barnesiae* the assemblages are relatively well
398 diversified and show only moderate signs of dissolution and recrystallization. Moreover,
399 dissolution-susceptible taxa were observed, although they were not very abundant. Therefore,
400 some general trends can be identified and are discussed. The rarefaction curve suggests that
401 the majority of the calcareous nannofossil species have been found in this study,
402 demonstrating that the dissolution did not affect species richness, only the abundance of taxa
403 (supporting information Figure S2).

404 In the five samples that the relative abundance of *W. barnesiae* is less than 40 %, it
405 could be observed that the relative abundance of *D. ignotus* and *Biscutum* spp. (indicative of
406 the NI) show the highest values registered in the study (Table 2). Therefore, in samples with
407 low dissolution the recovered assemblage shows high carbonate productivity and surface
408 water fertility. In addition, several studies report that *D. ignotus*, *Biscutum* spp., *Nannoconus*
409 spp. and *Braarudosphaera* spp. have a preference to nearshore paleoenvironments, and the
410 high abundance of *W. barnesiae* is related to deep basin deposits (Table 1).

411 The low-abundance of *Nannoconus* spp. and *Braarudosphaera* spp. (Figure 3) in the
412 studied samples do not support the hypothesis of nearshore deposition except in one sample
413 (749.19 mbsf, core 28), in which the benthic foraminifera assemblages is indicative of a
414 shallow to deep neritic paleoenvironment (Kochhann et al., 2014). In this sample, the low

415 value of CaCO₃ (30.90 %), high values of magnetic susceptibility (between 2.44×10^{-7} –
416 2.30×10^{-7} m³/kg for low and high frequency, respectively), high NI value (67), and an
417 Equitability value of 0.40 (related to the high-abundance of the genus *Biscutum* spp., at 58 %)
418 suggests that these sediments contained a high amount of nutrients, and that they were
419 deposited in a shallow to deep neritic paleoenvironment (Table 2).

420 The relative abundance of *D. ignotus*, *Z. erectus* and *Biscutum* spp. are normally used
421 to determinate the NI. However, *Z. erectus* is relatively rare in Site 364. Several studies have
422 considered *Z. erectus* more susceptible to dissolution than *Biscutum* spp. and *Discorhabdus*
423 *ignotus* (Roth & Krumbach, 1986). According to Fisher and Hay (1999), *Z. erectus* and
424 *Biscutum* spp. are indicative of different levels of increased fertility, with *Z. erectus* being
425 more abundant in high fertility conditions, and *Biscutum* spp. sensitive to very high fertility
426 conditions. On the other hand, Erba (1992a) described *Z. erectus* as being more abundant in
427 eutrophic environments, whereas *Biscutum* spp. has a preference for mesotrophic
428 environments. In general, *D. ignotus*, *Z. erectus* and *Biscutum* spp. are widely used in
429 paleoecological studies for indicating high fertility/productivity (Table 1). In this study,
430 assemblage analysis appears to indicate that *Biscutum* spp. is less susceptible to dissolution
431 than *Z. erectus*, and that *Biscutum* spp. is more abundant in very high fertility conditions, in
432 accordance to Fisher and Hay (1999). Between the five samples in which the relative
433 abundance of *W. barnesiae* is less than 40%, four had a relative abundance of *D. ignotus*
434 (Table 2). In these samples, it could also be observed high values of CaCO₃ (between 78.16–
435 90.03 %), Shannon Diversity Index (between 2.20–2.77 %), the occurrence of planktonic
436 foraminifera and absence of benthonic forms (Kochhann et al., 2013), and low values of
437 magnetic susceptibility (2.60×10^{-8} for low frequency and 2.54×10^{-8} m³/kg for high
438 frequency) (Table 2). This suggests that, for these samples, deposition occurred with high
439 carbonate productivity.

440 During the Cretaceous, many sedimentary basins of the Brazilian and African
441 continental margins show evidence of marine incursions deposited above evaporites, as
442 evidenced by numerous marine fossils and extensive biogenic limestone layers. Several
443 studies that analyze these deposits mark them as the moment of transition from restrict to
444 open marine environment conditions, when the waters of the Central Atlantic and the South
445 Atlantic oceans became connected (e.g., Azevedo, 2004; Eagles, 2007; Moulin et al., 2010).
446 During the Albian–Cenomanian interval, the South Atlantic Ocean already had open sea
447 conditions, with extensive, shallow marine carbonate shelves (Melguen, 1978; Mohriak et al.,

2008; Beglinger et al., 2012; Wagner & Pletsch, 1999). The calcareous nannofossils recovered in this study shows an assemblage with subtropical-tropical affinity, evidencing an influence of the Central Atlantic for this area during the Albian–early Cenomanian. It also gives support to the idea that there was a surface water connection between the Central Atlantic and the South Atlantic oceans, at least up to the Kwanza Basin (north of the Walvis Ridge-Rio Grande Rise). In addition, the absence of high latitude indicator species (Panera, 2011) show that there was no influence of cold surface waters in the area. According to Kochhann et al., (2013), this subtropical-tropical affinity can also be observed in the planktonic foraminifera observed in the same interval, in contrast to the benthic foraminiferal assemblage, which shows an Austral/Transitional province affinity (Scheibnerová, 1978; Kochhann et al., 2014). This paleobiogeographic contrast between the assemblages of planktonic and benthonic microfossils suggests a water column with a well-developed thermal gradient, as reported by Kochhann et al., (2014). These variations in seawater could have been caused by fluctuations in the inflow of surface and deep-water masses, by the influence of wind in surface waters, by subtropical arid circulation, by the progressive subsidence of the basin, and/or by relative oscillations in the sea level (Figure 6) (e.g., Poulsen et al., 2001; Hay, 2008; Hasegawa et al., 2012; Haq, 2014; Pérez-Díaz & Eagles, 2017).

The Cretaceous ocean anoxic events are interesting not only due to the fact that they created particularly favorable conditions for the accumulation and preservation of organic-rich sediments, but also because they provide valuable micropaleontological and geochemical data for paleoceanographic studies. The black shale horizons from the basal strata of Site 364 (between 1032.37–968.45 mbsf; cores 42–39, with 21 samples), referred alongside the dolomitic and marly limestones that surround them as sedimentary unit 7 (Bolli et al., 1978), were deposited during the Albian, as indicated by the presence of zone CC 8 or subzones BC 23 to BC 24, and represent an important record of the paleoceanographic changes that occurred during that time in the South Atlantic ocean. These black shale horizons are characterized by calcareous and pelagic sedimentation with a high percentage of marine organic matter (type II kerogen; Bolli et al., 1978; Tissot et al., 1980; Magniez-Jannin & Muller, 1987; Herbin et al., 1987), being of probable algal origin with low organic diagenesis (Raynaud & Robert, 1978), containing extensive dissolution that is reflected in the low abundance of benthonic foraminifera (Scheibnerová, 1978; Kochhann et al., 2014), and by low primary productivity (Melguen, 1978). The biogenic carbonate fraction of the studied sediments is mainly composed of planktonic and benthonic foraminifera (Melguen, 1978) and

481 calcareous nannofossil. In smear slides, most of the inorganic carbonates consist mainly of
482 calcite, dolomite, and unspecified carbonate particles (Siesser & Bremner, 1978). These
483 horizons present a good potential for the exploration of hydrocarbons, as they were formed in
484 situ by the low temperature degradation of organic matter (Kendrick et al., 1978; Hunt, 1978;
485 Tissot et al., 1980), and have been interpreted as a local response to OAEs events (e.g.
486 Bralower et al., 1994; Kochhann et al., 2013; Naafs & Pancost, 2014). The TOC analysis for
487 this unit shows the highest values for the studied interval, ranging from 0.40 to 8.91 %; on the
488 other hand, the CaCO₃ values registered were the lowest, being in the 11.08–75.26 % range.
489 Geochemical data shows a positive correlation ($r = 0.32$) between $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ stable
490 isotopes in the basal strata of Site 364, suggesting low diagenetic action as reported by
491 Kochhann et al., (2013). The data also shows a negative correlation ($r = -0.75$) between TOC
492 and CaCO₃; therefore, different physicochemical conditions related to black shale
493 preservation/deposition seem to have been responsible for the dissolution of CaCO₃ and the
494 increased TOC content in this interval. According to Bolli et al., (1978) this interval is
495 characterized by the low-deposition of terrigenous sediments, showing a more calcareous and
496 pelagic sedimentation style, with the low values of CaCO₃ being caused by low primary
497 productivity in euxinic conditions (anoxic and sulfidic). These oceanic conditions would have
498 controlled the primary productivity and affected the preservation of carbonates in the Kwanza
499 Basin.

500 In sedimentary unit 7, the calcareous nannofossil assemblage shows minor to
501 moderate preservation and a richness of 44 species, averaging at 31 species. *W. barnesiae*
502 shows high-abundance in this sedimentary unit; on the other hand, NI values are the lowest
503 for the whole interval. The analyzed data shows a high negative correlation ($r = -0.97$)
504 between the Shannon Diversity Index and percentage of *W. barnesiae*, strongly suggesting
505 that dissolution affected the assemblage composition of sedimentary unit 7. In spite of the
506 high-abundance of *W. barnesiae*, the black shale intervals of Site 364 show a moderately
507 preserved, relatively rich assemblage of calcareous nannofossils, with dissolution-susceptible
508 taxa being present, albeit in relatively low quantities. The combination of calcareous
509 nannofossil and geochemistry data alongside the occurrence of dolomite might suggest that,
510 during deposition, the area did not display euxinic conditions. In addition, the dissolution
511 effect on the assemblage composition makes it impossible to discuss the relationship between
512 the black shale layers and the NI. The dissolution observed also prevented the correlation
513 between the Albian black shales from Kwanza Basin and the OAEs described for this time

514 interval. The black shales layers show low magnetic susceptibility and is predominantly
515 controlled by the paramagnetic minerals (Chadima et al., 2006).

516 For the interval between 955.75 and 717.23 mbsf (cores 33–26), no black shale layers
517 were observed (Bolli et al., 1978), and the TOC content is low, with only sample 894.99 mbsf
518 (Core 35) showing a high TOC value (2.89 %). For sample 894.99 mbsf, the high TOC value
519 may be related to carbon-rich organic sediments, although fertility (NI = 13) and MS
520 (5.76×10^{-8} and 5.54×10^{-8} m³/kg, low and high frequency respectively) are not high. In the
521 interval between 850.15–845.90 mbsf (core 33) is observed the highest values of MS being
522 observed, this might be associated with the high clay content recorded for this interval (Bolli
523 et al., 1978).

524 **5. Conclusions**

525 The calcareous nannofossil assemblages of Site 364 (cores 42–26) display a
526 subtropical-tropical affinity, providing an excellent biostratigraphic record of the Albian–late
527 Cenomanian interval of the South Atlantic Ocean. The paleobiogeographic affinity of the
528 nannofossil assemblage, comparable to that of planktonic foraminifera found in the same site,
529 indicate that there was a surface water connection between the Central Atlantic and the South
530 Atlantic oceans at least up to offshore Angola, north of the Walvis Ridge-Rio Grande Rise.

531 These assemblages, alongside geochemical data, show that sedimentation was
532 predominantly calcareous and pelagic during the Albian–late Cenomanian interval of the
533 Kwanza Basin, with the exception of its basal section, where black shale layers interbedded
534 with dolomitic and marly limestone of Albian age can be observed. In this section, the black
535 shale layers are dominated by Type II kerogen (Tissot et al., 1980; Magniez-Jannin and
536 Muller, 1987; Herbin et al., 1987). This, alongside geochemical data and the low values of
537 magnetic susceptibility, suggest that these organic carbon-rich sediments with high TOC
538 values were deposited due to the high fertility of the marine waters at the time. The calcareous
539 nannofossils recovered show moderate signs of dissolution and recrystallization; as a result,
540 they likely do not reflect the original abundance of dissolution-susceptible species and the
541 Nutrients Index values of the black shale layers. In spite of the dissolution, assemblages are
542 relatively well diversified, with dissolution-susceptible taxa having been observed in low
543 abundance. This suggests that the effects of dissolution may have been selective, limiting the
544 abundance but not significantly affecting species richness, and also indicating that, during
545 deposition, the environment did not display euxinic conditions. Also due to the effects of

546 dissolution, the relationship between the deposition of black shale layers and high fertility of
547 calcareous nannofossils could not be properly verified.

548 Nannofloral distribution patterns observed could, however, give a measure of the
549 fertility and productivity of the strata above the black shale layers. They show that there was a
550 decline in the effects of dissolution in the composition of the fossil assemblages and in the
551 geochemical signals for these deposits. Geochemical data and the high relative abundance of
552 *Discorhabdus ignotus* suggest that, during deposition, there was high carbonate productivity,
553 though not necessarily high fertility. The large abundance of the genus *Biscutum* spp. could
554 indicate that this genus is less susceptible to dissolution than *Zeugrhabdotus erectus*, and
555 previous studies show that *Biscutum* spp. is more abundant in very high fertility conditions
556 and in shallow to deep neritic paleoenvironments.

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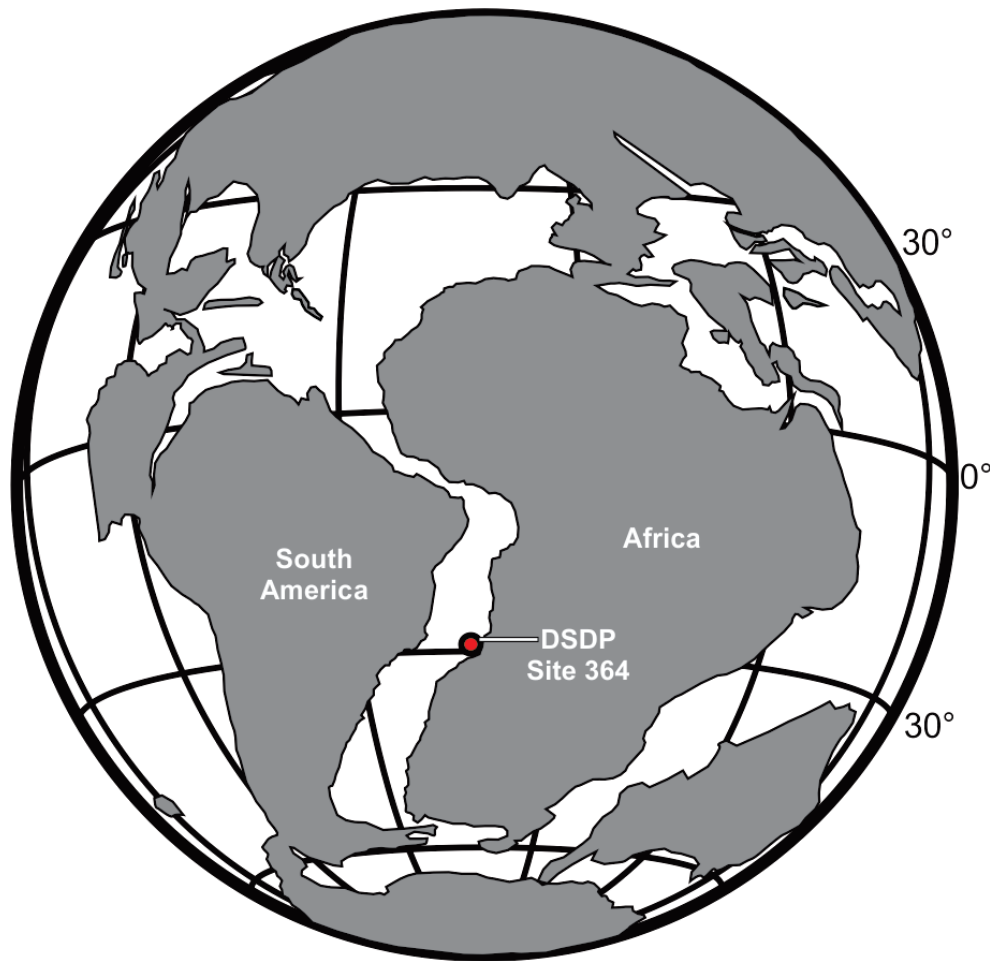
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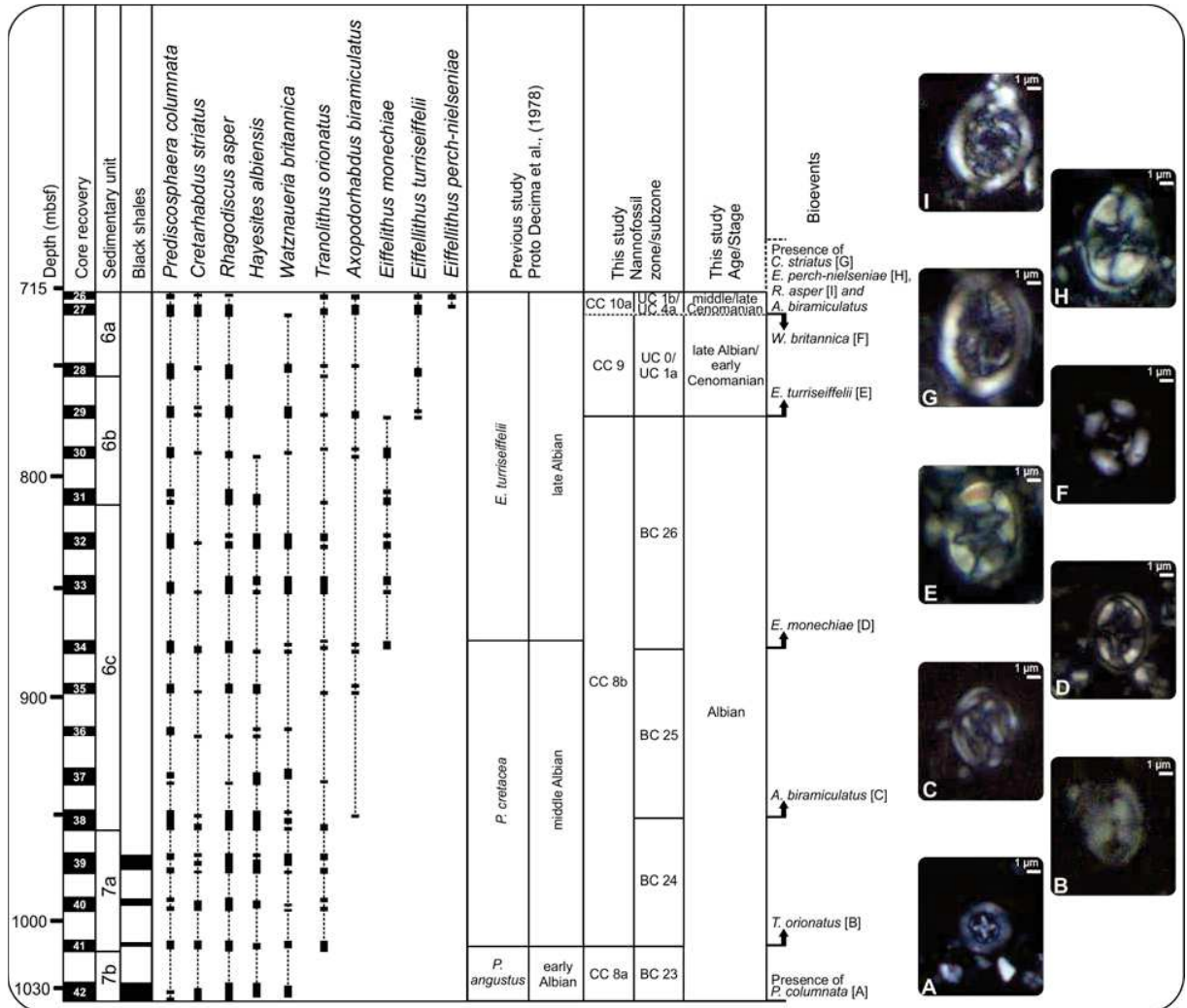
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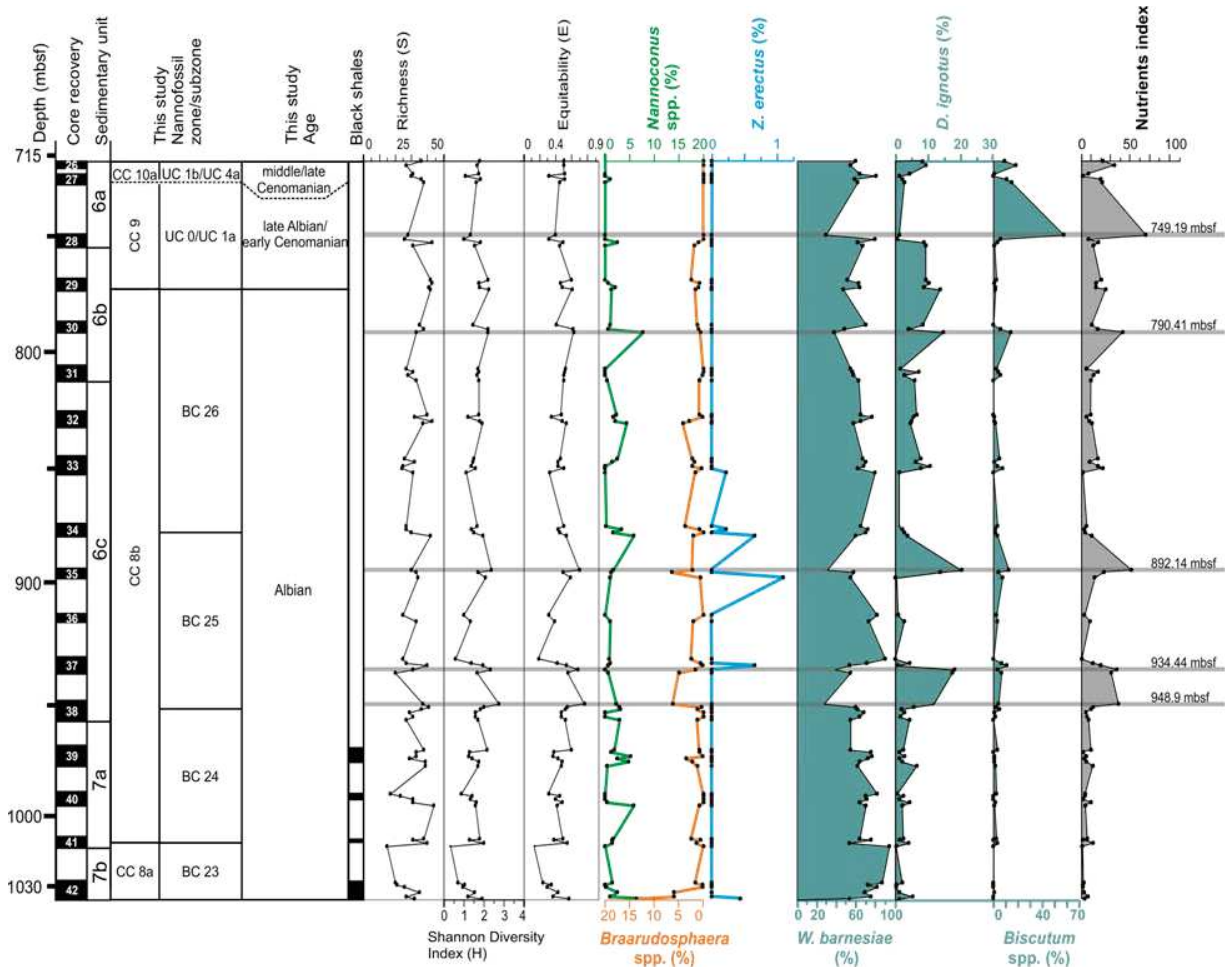
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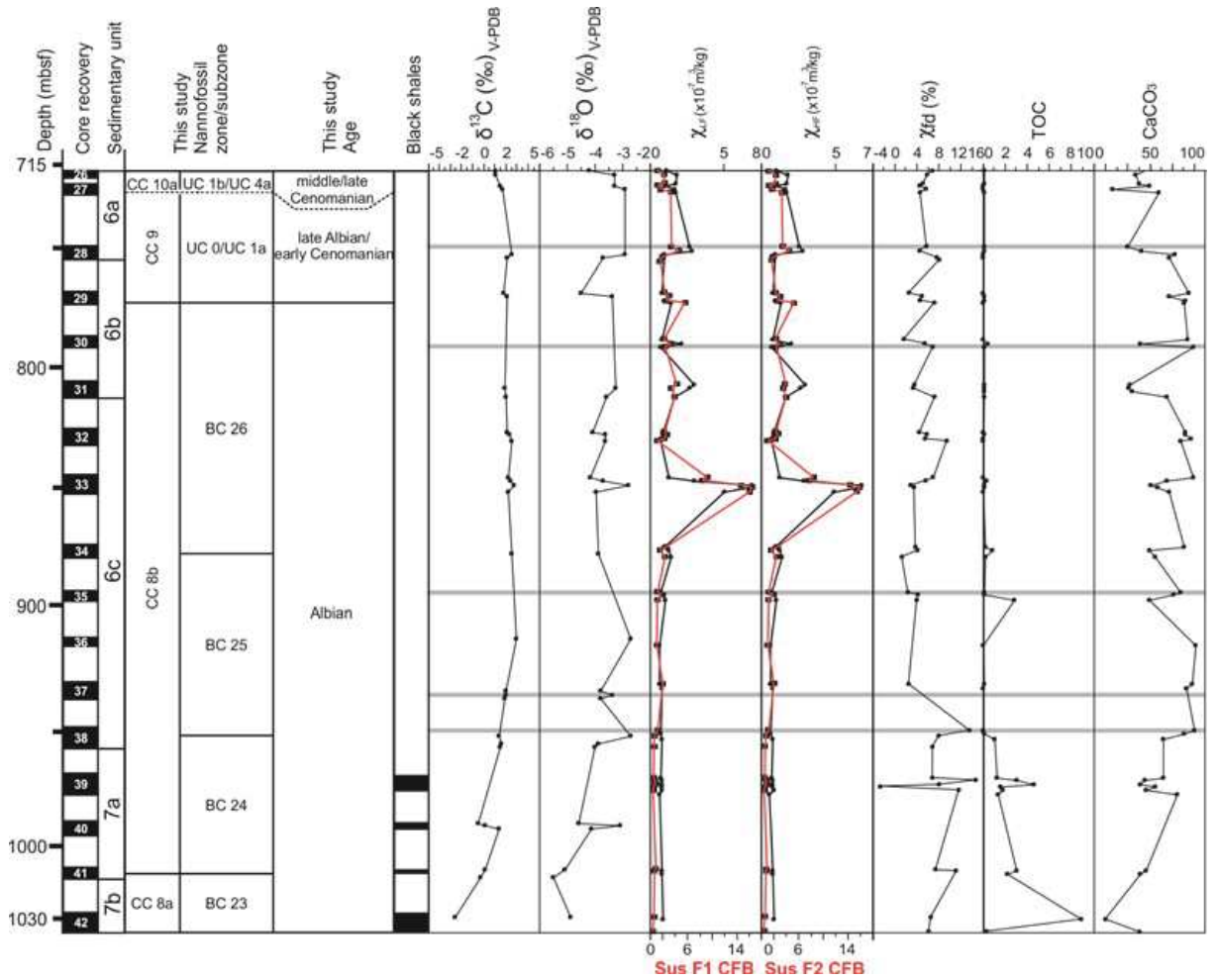
882 **Figure 1.** Paleogeographic reconstruction for 110 Ma. Modified from ODSN Plate Tectonic
883 Reconstruction Service, showing the inferred location of Site 364.



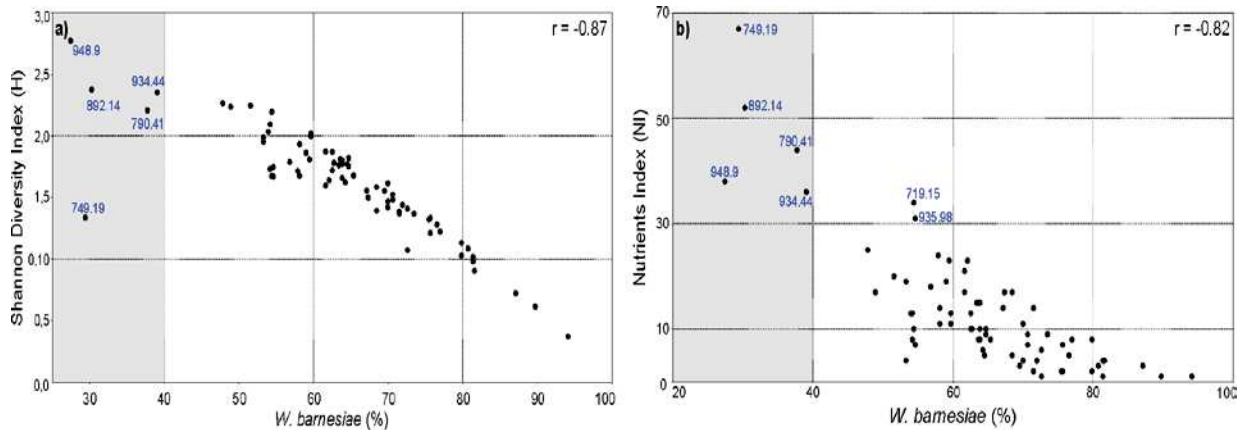
886 **Figure 2.** Biostratigraphic of selected taxa from Site 364. Calcareous nannofossil
 887 biostratigraphy is based on the zonation schemes: CC zones of Sissingh (1977) and Perch-
 888 Nielsen (1979, 1985), BC zonation of Bown et al., (1998) and the UC of Burnett et al.,
 889 (1998). The distribution of black shale layers and sedimentary units is based on data from
 890 Bolli et al., (1978).



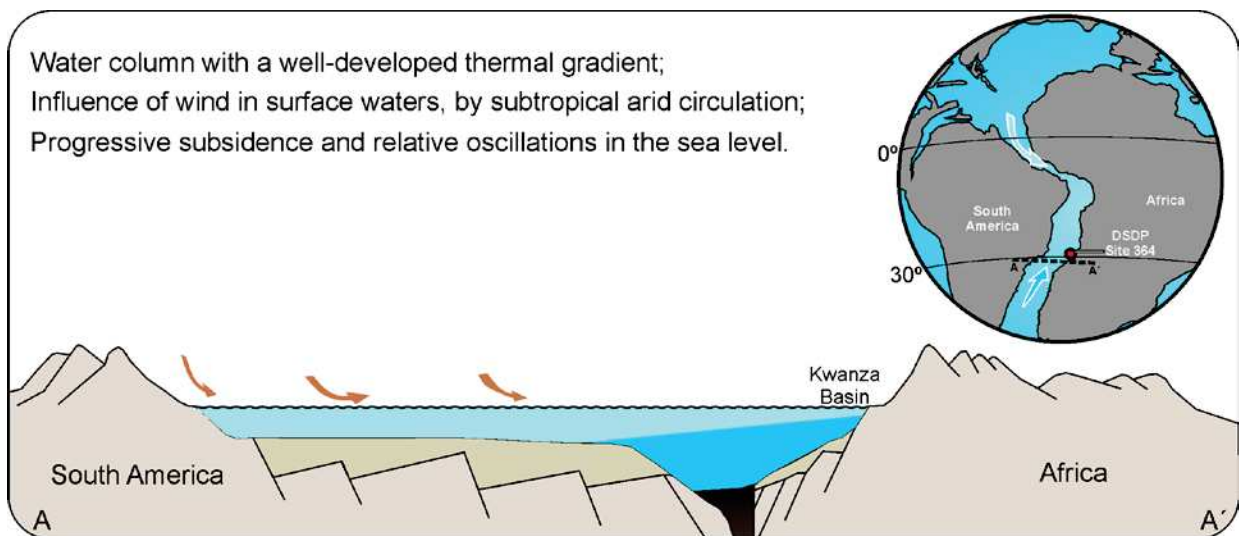
893 **Figure 3.** Values for Richness (S), Shannon Diversity Index (H), Equitability (E),
 894 abundances of major taxa and the Nutrients Index from Site 364. The samples with increased
 895 Nutrients Index and low-abundance of *Watznaueria barnesiae* are indicated by gray horizontal
 896 shading.



899 **Figure 4.** Distribution of carbon ($\delta^{13}\text{C}$) and oxygen ($\delta^{18}\text{O}$) isotope values, magnetic
 900 susceptibility (χ_{lf} , χ_{hf} , χ_{fd} %, and carbonate-free based), TOC and CaCO_3 in the Site 364. The
 901 samples with increased Nutrient Index and low-abundance of *W. barnesiae* are indicated by
 902 gray horizontal shading.



905 **Figure 5.** Array of *Watznaueria barnesiæ* abundance compared to Shannon Diversity Index
 906 (H) and Nutrients Index (NI). a) For the relationship between *W. barnesiæ* (%) and H the
 907 person correlation coefficients (r) is negative and significant, suggesting that the assemblage
 908 suffered from dissolution. b) For the relationship between *W. barnesiæ* (%) and NI the r is
 909 negative and significant, showing high NI values when the percentage of *W. barnesiæ* is
 910 below 40%.



912 **Figure 6.** Paleogeographic inferences for the South Atlantic. The thermal gradient of the
 913 water column is evidenced by the paleobiogeographic contrast between planktonic and
 914 benthonic microfossil assemblages. The arrows illustrate different causes for these variations,
 915 such as wind in surface waters (brown) and fluctuations in the inflow of surface (light blue)
 916 and deep-water (dark blue) masses (Poulsen et al., 2001; Azevedo, 2004; Hay, 2008;
 917 Hasegawa et al., 2012; Haq, 2014; Kochhann et al., 2014; Pérez-Díaz & Eagles, 2017).

919 **Table 1.** Calcareous nannofossil paleoecological indices considered in this study.

Calcareous nannofossil	Paleoecologic preferences						
	Dissolution resistant	Neritic environment	Restricted sea (and/or Epicontinental Basin, Upwelling assemblage, Oceanic Plateau)	Continental Margins (Near-shore)	Offshore (Deep basin and Open ocean)	High carbonate productivity/surface water fertility	Low carbonate productivity/surface water fertility
<i>Biscutum</i> spp. (include <i>Biscutum ubiqem</i> , <i>Biscutum</i> sp. 1 and <i>Biscutum</i> sp. 2)			2, 3, 10, 12, 23	2, 3, 24, 35		2, 3, 8, 9, 11, 12, 14, 16, 20, 21, 22, 24, 26, 27, 29, 31, 32, 33, 36	
<i>Discorhabdus ignotus</i>			12			12, 14, 15, 27	
<i>Zeugrhabdotus erectus</i>			3, 10	2, 3, 24		2, 8, 9, 11, 14, 22, 26, 27, 29, 31, 32, 33	
<i>Watznaeria barnesiae</i>	2, 5, 7, 8, 9, 14, 16, 18, 19, 23, 25, 30, 31				3, 8, 29, 35		9, 16, 19, 22, 24, 27, 28, 31, 32
<i>Nannoconus</i> spp. (include <i>Nannoconus</i> sp.1, <i>N. fragilis</i> , <i>N. truitii</i> subsp. <i>frequens</i> , <i>N. truitii</i> subsp. <i>rectangularis</i> , <i>N. truitii</i> subsp. <i>truitii</i> and <i>N. troelsenii</i>)	17, 29	2, 4, 8, 9, 24	1, 3, 8, 9, 13, 29	3, 6, 9, 13		12, 14, 34	
<i>Braarudosphaera</i> spp. (include <i>B. africana</i> , <i>B. batilliformis</i> , <i>B. hockwoldensis</i> , <i>B. primula</i> , <i>B. pseudobatilliformis</i> and <i>B. regularis</i>)		2, 6, 24	3	3, 6		3	

1. Thierstein (1976); 2. Roth (1981); 3. Roth & Bowdler (1981); 4. Doeven (1983); 5. Roth (1984); 6. Stradner et al., (1984); 7. Perch-Nielsen (1985); 8. Roth (1986); 9. Roth & Krumbach (1986); 10. Roth (1989); 11. Watkins (1989); 12. Premoli-Silva et al., (1989); 13. Busson & Noel (1991); 14. Erba (1992a); 15. Erba (1992b); 16. Erba et al., (1992); 17. Erba (1994); 18. Erba et al., (1995); 19. Williams & Bralower (1995); 20. Eshet & Almogi-Labin (1996); 21. Gardin & Monechi (1998); 22. Fisher & Hay (1999); 23. Mutterlose & Kessels (2000); 24. Street & Bown (2000); 25. Luciani et al., (2001); 26. Gardin (2002); 27. Herrle et al., (2003); 28. Friedrich et al., (2005); 29. Less et al., (2005); 30. Watkins & Self Trail (2005); 31. Kulhanek & Wise (2006); 32. Thibault & Gardin (2006); 33. Hadas & Mutterlose (2007); 34. Browning & Watkins (2008); 35. Linnert et al., (2011); 36. Thibault & Husson (2016).

922 **Table 2.** Data from the five samples with increased Nutrient Index and low-abundance of
 923 *Watznaueria barnesiae*. Planktonic and benthonic foraminifera data, according to Kochhann
 924 et al., (2013, 2014).

Samples (mbsf)	Richness (S)	Shannon Diversity (H)	Equitability (E)	<i>W. barnesiae</i> (%)	<i>D. ignotus</i> (%)	<i>Biscutum</i> spp. (%)	<i>Z. erectus</i> (%)	<i>Nannoconus</i> spp. (%)	<i>Braarudosphaera</i> spp. (%)	Nutrient Index	Susceptibility F1 (m ³ /kg)	Susceptibility F3 (m ³ /kg)	TOC (%)	CaCO ₃ (%)	% Planktonic foraminiferal (Kochhann et al., 2013)	% Benthic foraminiferal (Kochhann et al., 2014)	Preservation Index (Kochhann et al., 2013)
749.19	28,00	1,337	0,4013	29	2	58	0	0	0	67	2,44E-07	2,30E-07	0,11	30,90	97.1	2.9	Good
790.41	33,00	2,209	0,6317	38	15	14	0	8	1	44	2,53E-08	2,36E-08	0,14	90,03	100,00	0,00	Moderate
892.14	30,00	2,375	0,6984	30	20	13	0	2	2	52	2,60E-08	2,54E-08	0,15	78,16	100,00	0,00	Poor
934.44	31,00	2,352	0,6849	39	18	4	0	0	2	36					100,00	0,00	Moderate
948.9	37,00	2,772	0,7676	27	12	5	0	2	6	38	2,08E-08	1,80E-08	0,19	81,59	100,00	0,00	Poor

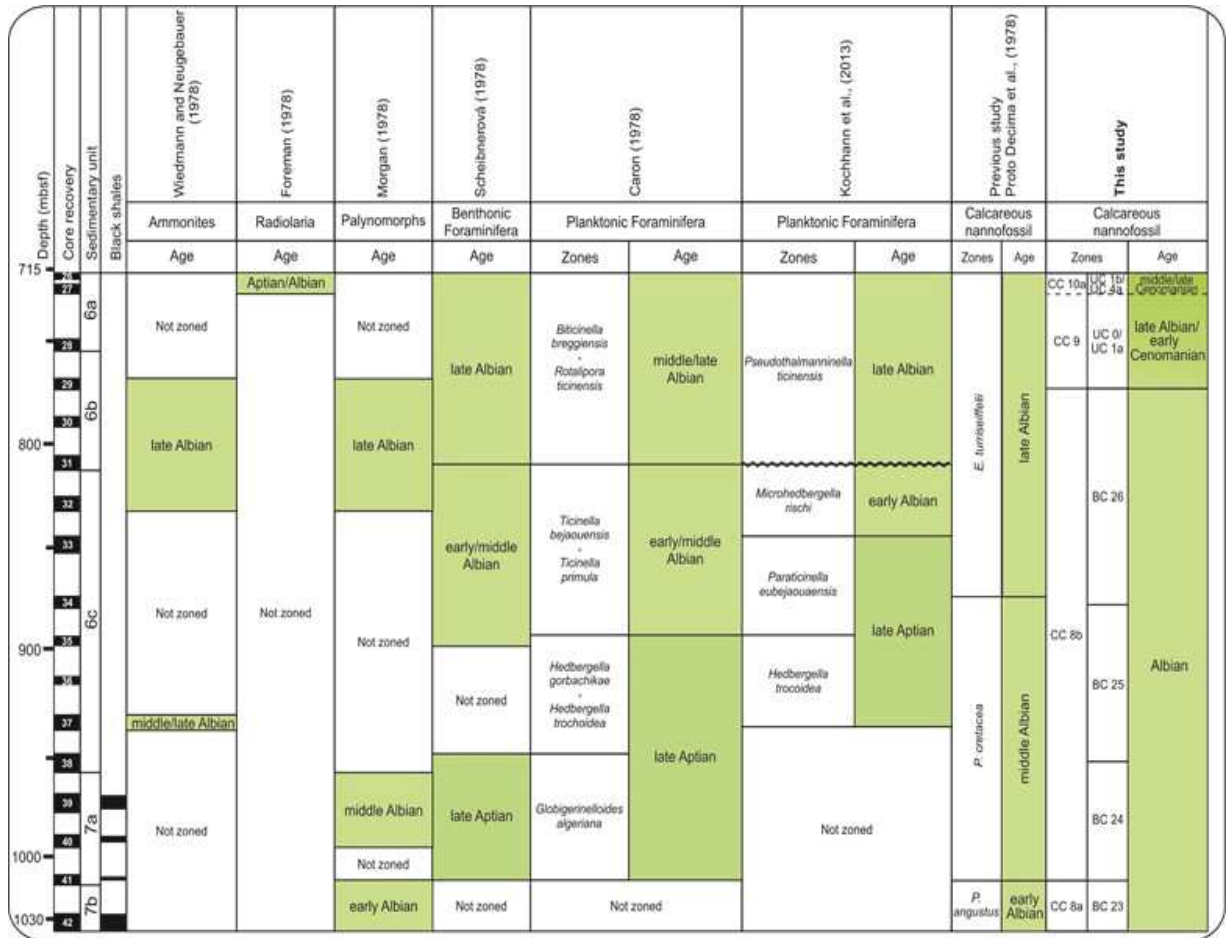


Figure S1. Correlation between the biostratigraphic zonation of calcareous nannofossils, planktonic and benthonic foraminifera, ammonites, palynomorphs and radiolarians for Site 364.

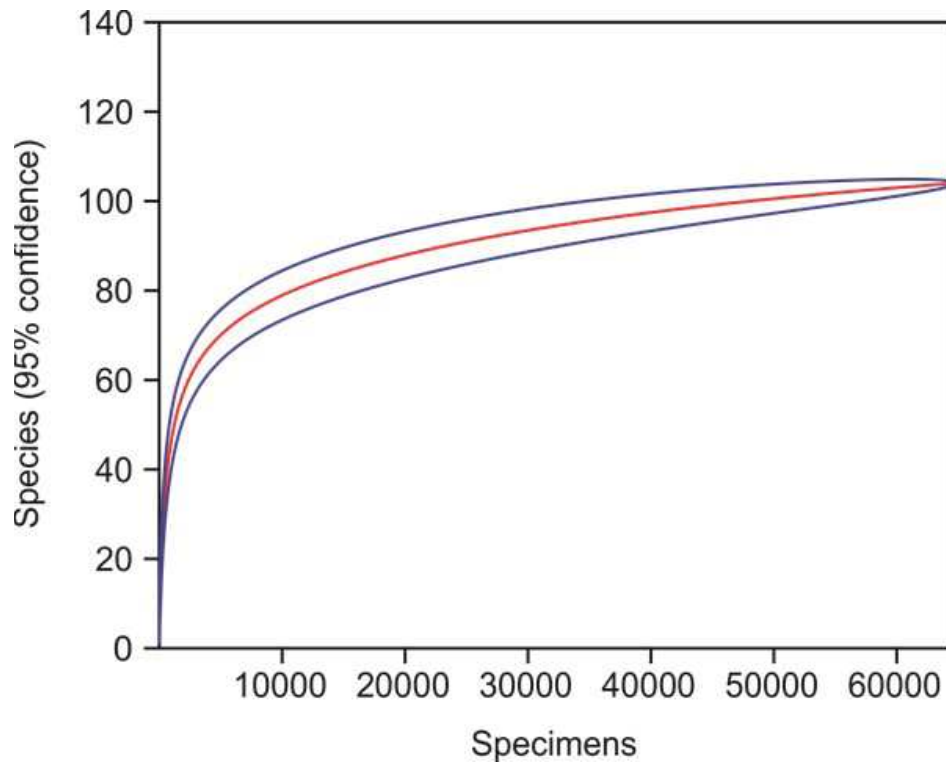


Figure S2. Rarefaction curve, based on 72 samples, show number of specimens against number of species richness recorded in different samples from Site 364. This curve strongly suggests that the majority of the calcareous nannofossil species have been found. Future collecting efforts may occasionally yield new species, but the total number of species is not expected not change substantially.

Table S3. Minimum, maximum and average of calcareous nannofossil and geochemical data, in the basal and top sections studied from Site 364, Kwanza Basin.

Data		Basal Section (1032.37-968.45 mbsf)	Top Section (955.75- 717.23 mbsf)
Sedimentary Unit		7	6
Richness (S)	Minimum	15	20
	Average	31	33
	Maximum	44	43
Shannon Diversity (H)	Minimum	0.37	0.62
	Average	1.42	1.71
	Maximum	2.2	2.77
Equitability (E)	Minimum	0.14	0.19
	Average	0.41	0.49
	Maximum	0.60	0.77
<i>W. barnesiae</i> (%)	Minimum	53	27
	Average	71	61
	Maximum	94	90
<i>D. ignotus</i> (%)	Minimum	0.4	0
	Average	2	6
	Maximum	7	20
<i>Biscutum</i> spp. (%)	Minimum	0	0
	Average	1	6
	Maximum	4	58
Nutrient Index	Minimum	1	1
	Average	5	17
	Maximum	13	67
$\delta^{13}\text{C}_{\text{V-PDB}}$ (‰)	Minimum	-2.59	1.06
	Average	1.36	2.03
	Maximum	-0.27	2.94
$\delta^{18}\text{O}_{\text{V-PDB}}$ (‰)	Minimum	-5.47	-4.49
	Average	-4.52	-3.55
	Maximum	-3.05	-2.68
Susceptibility F1 (m ³ /kg)	Minimum	1,15E-08	1.14E-08
	Average	2,87E-08	1.34E-07
	Maximum	4,01E-08	7.12E-07
Susceptibility F3 (m ³ /kg)	Minimum	1,02E-08	1.04E-08
	Average	2,71E-08	1.28E-07
	Maximum	3,75E-08	6.91E-07
X_{fd} (%)	Minimum	-3.22	1.35
	Average	7.6	5.4
	Maximum	14.56	13.27
TOC (%)	Minimum	0.40	0.055
	Average	3	0.3
	Maximum	8.91	2.89
CaCO ₃ (%)	Minimum	11.08	17.45
	Average	45.40	64.22
	Maximum	75.26	90.64

CONSIDERAÇÕES FINAIS

O estudo dos nanofósseis calcários do Albiano no Site 364 apresenta uma assembleia contendo 103 táxons que são típicos do intervalo Albiano–Cenomaniano. Foi possível observar uma ocorrência contínua de bioeventos descritos para este intervalo de tempo. Portanto, a partir do estudo bioestratigráfico não foram observados indícios de retrabalhamento e hiatos na seção estudada.

A assembleia recuperada também fornece uma importante contribuição ao entendimento da influência das correntes marinhas superficiais, considerando que a assembleia é constituída em sua totalidade por espécies com afinidade subtropical-tropical. Este dado paleobiogeográficos corroboram os estudos de Kochhann et al., (2013, 2014) sobre as influências das águas superficiais da região central do Oceano Atlântico na Bacia de Kwanza, Angola.

A dissolução do carbonato de cálcio é um processo com significativos efeitos na preservação e abundância dos nanofósseis calcários no *Site* 364, os dados sedimentológicos, micropaleontológicos e geoquímicos permitem observar registros dos efeitos da dissolução de modo qualitativo ao longo da seção estudada. O material carbonático sofreu dissolução e forte recristalização na seção basal do testemunho, onde é possível observar graus baixos a médios destes processos nos táxons observados. Os sinais isotópicos sofreram alterações, possivelmente devido a dissolução, assim como o conteúdo de carbonato de cálcio inorgânico oscila entre valores médios e altos (principalmente nos intervalos com dolomita). Os valores de carbono orgânico total (COT) obtidos correspondem de modo satisfatório ao intervalo (unidade sedimentar 7) descrito por Bolli et al., (1978), nesta unidade a dissolução afetou a abundância dos nanofósseis calcários, resultando na baixa preservação de espécies susceptíveis a dissolução. Devido a esta alteração, não foi possível realizar inferências acerca da relação entre os depósitos de folhelhos negros e a alta fertilidade de nanofósseis calcários.

Apesar da dissolução, a assembleia recuperada é relativamente bem diversificada e os táxons suscetíveis à dissolução foram observados, mas não de modo abundante. Em cinco amostras (na unidade sedimentar 6) foi possível realizar inferências paleoceanográficas para a Bacia de Kwanza, nesta unidade a dissolução não foi tão severa como na unidade 7. Nestas cinco amostras foi possível inferir dados de fertilidade e produtividade de carbonatos nas águas superficiais, durante o intervalo Albiano–Cenomaniano do *Site* 364.

Os 103 táxons descritos e identificados neste trabalho, são apresentados nas páginas que seguem.

SYSTEMATIC PALEONTOLOGY: PLATE 1 – HETEROCOCCOLITHS

Order EIFFELLITHALES Rood *et al.*, 1971

Family CHIASTOZYGACEAE Rood *et al.*, 1971

Genus *Tranolithus* Stover, 1966

Tranolithus gabalus Stover, 1966

Description: Medium to large coccoliths; specie have an elliptical outline; the rim is simple and unicyclic; the central opening is spanned transversely by two large blocks.

Tranolithus orionatus (Reinhardt, 1966a) Reinhardt, 1966b

Description: Medium to large coccoliths; specie have an elliptical outline; the rim is simple and unicyclic; the central area have four massive blocks.

Tranolithus minimus (Bukry, 1969) Perch-Nielsen, 1984

Description: Small to medium coccoliths; elliptical outline; the rim is simple and unicyclic; the central area feature a bridge and two large blocks with high birefringence.

Genus *Vekshinella* Loeblich & Tappan, 1963

Vagalapilla angusta (Stover 1966) Roth, 1981

Description: Large to very large coccoliths; elliptical outline; the rim is narrow and bicyclic with the same width, the inner cycle is brighter; the central area have a cross with angles of less than 20° between cross arms and principle axes.

Genus *Staurolithites* Caratini, 1963

Staurolithites laffittei Caratini, 1963

Description: Small to medium coccoliths; elliptical outline; the rim is bicyclic; the central area have a central cross aligned with the axes.

Staurolithites crux (Deflandre & Fert, 1954) Caratini, 1963

Description: Small coccoliths; elliptical outline; the rim is narrow; and the wide central area have a central cross, with high birefringence, aligned with the axes.

Staurolithites sp. 1

Description: Medium coccoliths; elliptical outline; the rim is unicyclic and large; the central area is closed and have a cross.

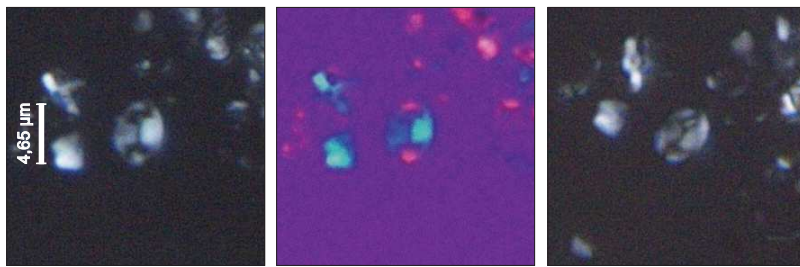
Genus *Glaukolithus* Reinhardt (1964)

Glaukolithus diplogrammus (Deflandre) Reinhardt, 1964

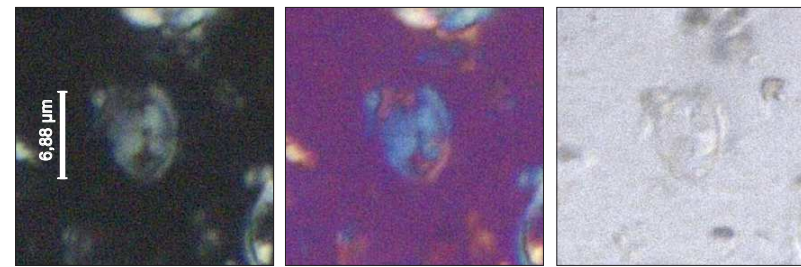
Description: Medium to large coccoliths; elliptical outline; the rim is simple; the central area is broad and have two bars narrow and parallels.

Glaukolithus bicrescenticus (Stover 1966)

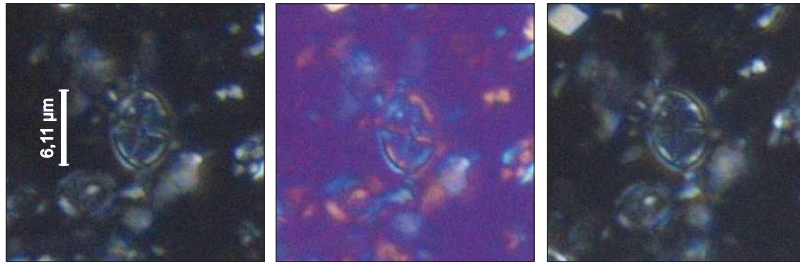
Description: Medium to large coccoliths; elliptical outline; the rim is simple; the central area is narrow and have two bars broad and parallels.



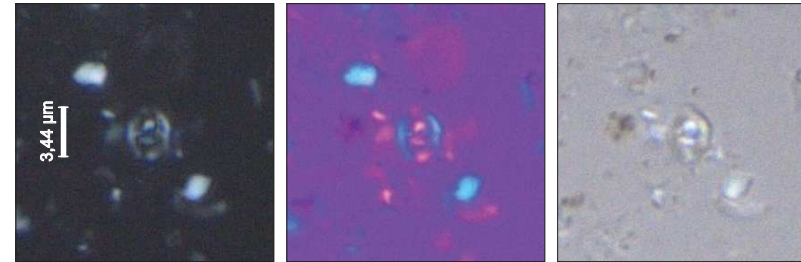
Tranolithus gabalus (DSDP Site 364 / Depth 1.032,37 mbsf)



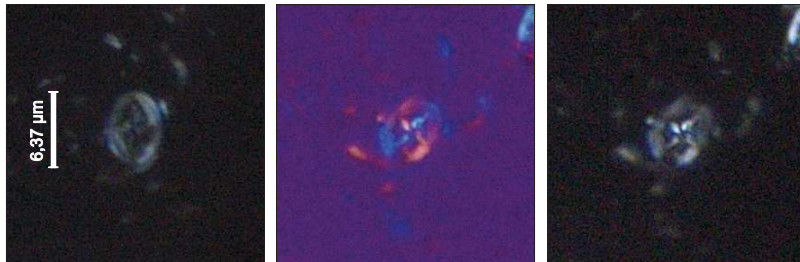
Tranolithus orionatus (DSDP Site 364 / Depth 723,67 mbsf)



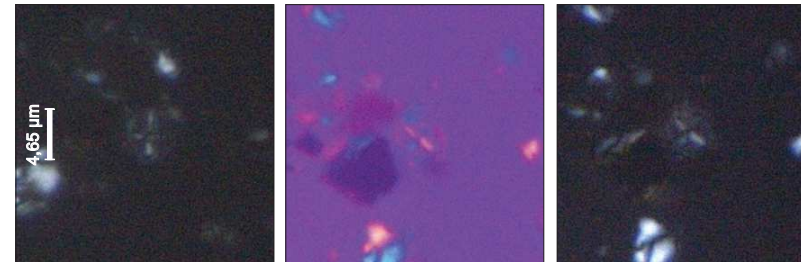
Vagalapilla angusta (DSDP Site 364 / Depth 877,42 mbsf)



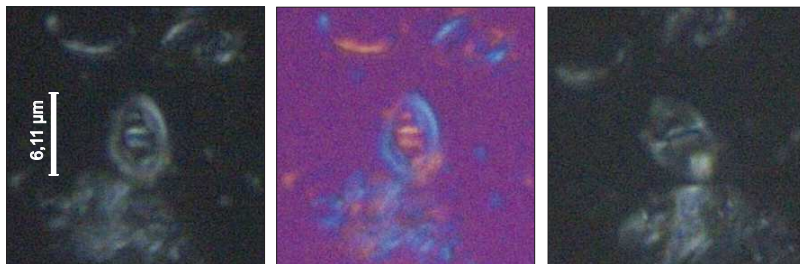
Tranolithus minimus (DSDP Site 364 / Depth 1032,37 mbsf)



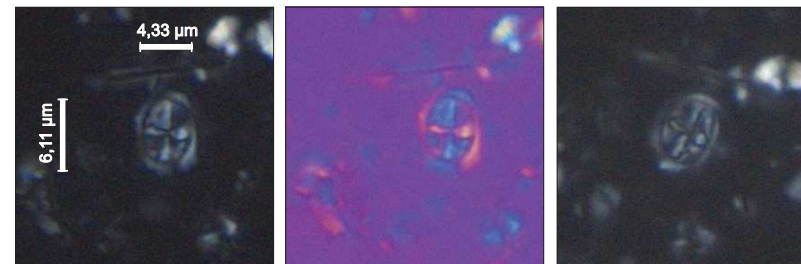
Staurolithites laffittei (DSDP Site 364 / Depth 894,99 mbsf)



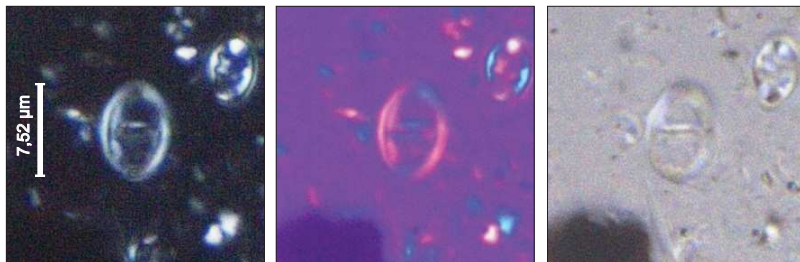
Staurolithites crux (DSDP Site 364 / Depth 1027,24 mbsf)



Glaukolithus bicrescenticus (DSDP Site 364 / Depth 719,15 mbsf)



Staurolithites sp. 1 (DSDP Site 364 / Depth 1.008,41 mbsf)



Glaukolithus diplogrammus (DSDP Site 364 / Depth 807,40 mbsf)



SYSTEMATIC PALEONTOLOGY: PLATE 2 – HETEROCOCCOLITHS

Genus *Zeugrhabdotus* Reinhardt, 1965

Zeugrhabdotus angelozziae Pérez Panera 2012

Description: Medium to large; specie have an elliptical outline; the rim is bicyclic; the central-area spanned by two parallel bars disposed oblique to the main axes of the ellipse.

Zeugrhabdotus howei Bown in Kennedy *et al.*, 2000

Description: Medium coccoliths; elliptical outline; the rim is unicyclic; the central-area have a bar granular.

Zeugrhabdotus embergeri (Noël, 1959) Perch-Nielsen, 1984

Description: Large coccoliths; elliptical outline; the rim is bicyclic; the central-area have a bar granular with diamond ornamentation.

Zeugrhabdotus clarus Bown, 2005

Description: Small to medium coccoliths; elliptical outline; the rim is bicyclic, strongly curved extinction lines cross the bright inner cycle; the central-area have a bar granular.

Zeugrhabdotus erectus (Deflandre in Deflandre & Fert, 1954) Reinhardt, 1965

Description: Small coccoliths; elliptical outline; the rim is unicyclic, the central-area have a bar granular birefringent.

Genus *Chiastozygus* Gartner, 1968

Chiastozygus litterarius (Górka, 1957) Manivit, 1971

Description: Medium to large coccoliths; elliptical outline; the rim is usually unicyclic, the central-area have a crossbars arranged as an x.

Chiastozygus platyrhethus Hill, 1976

Description: Medium to large coccoliths; elliptical outline; the rim is usually unicyclic, the central-area wide have a large central cross.

Genus *Loxolithus* Noël, 1965

Loxolithus armilla (Black in Black & Barnes, 1959) Noël, 1965

Description: Medium to large coccoliths; elliptical outline; the rim is usually unicyclic, strongly curved extinction lines cross the cycle; central-area is broad and void.

Family EIFFELLITHACEAE Reinhardt, 1965

Genus *Helicolithus* Noël, 1970

Helicolithus trabeculatus (Górka, 1957) Verbeek, 1977

Description: Small to medium coccoliths; elliptical outline; the rim is usually bicyclic; the central-area have a symmetrical diagonal cross bars.

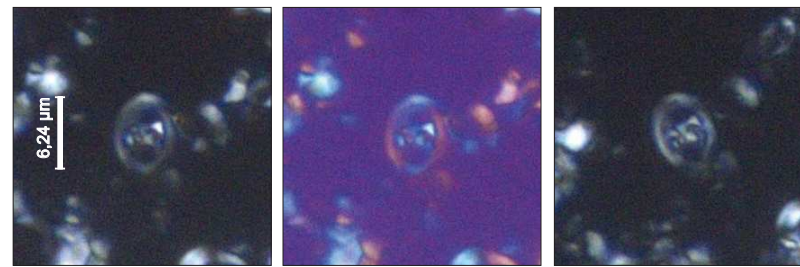
Genus *Tegumentum stradneri* Thierstein in Roth & Thierstein, 1972

Tegumentum stradneri Thierstein in Roth & Thierstein, 1972

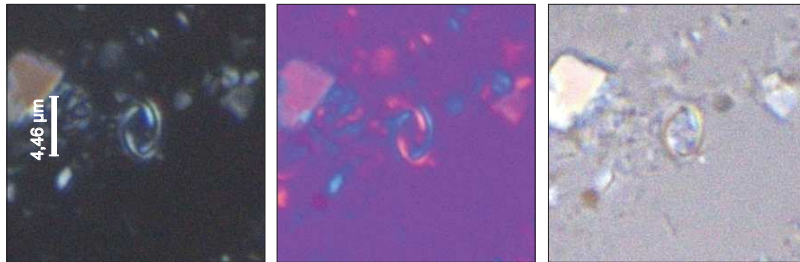
Description: Medium to large coccoliths; specie with a broadly to normally elliptical; the rim is bicyclic, inner cycle is brighter (high birefringence); the central area is large with a diagonal cross bar that appear sigmoidal curved in pointed terminations.



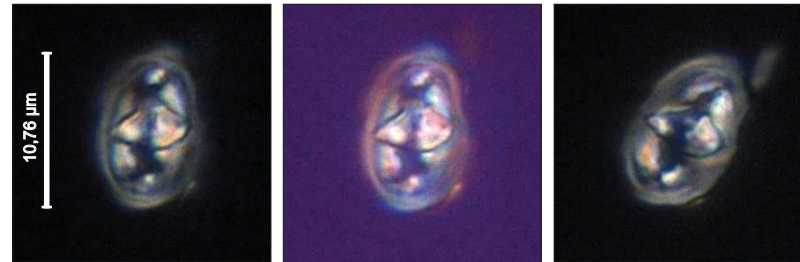
Zeugrhabdotus angelozziae (DSDP Site 364 / Depth 1.006,41 mbsf)



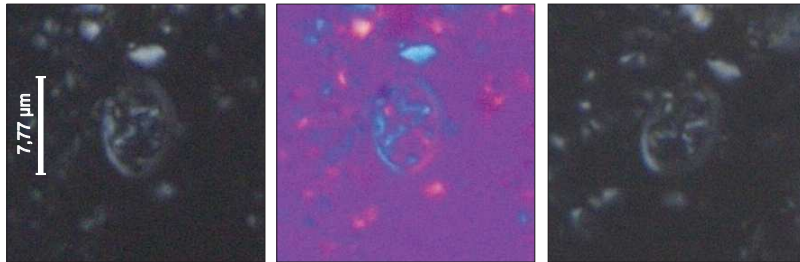
Zeugrhabdotus howei (DSDP Site 364 / Depth 911,25 mbsf)



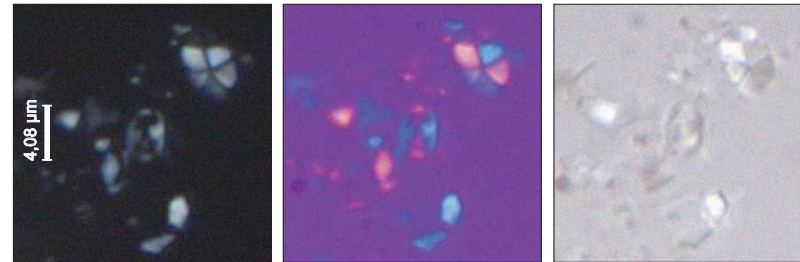
Zeugrhabdotus clarus (DSDP Site 364 / Depth 975,49 mbsf)



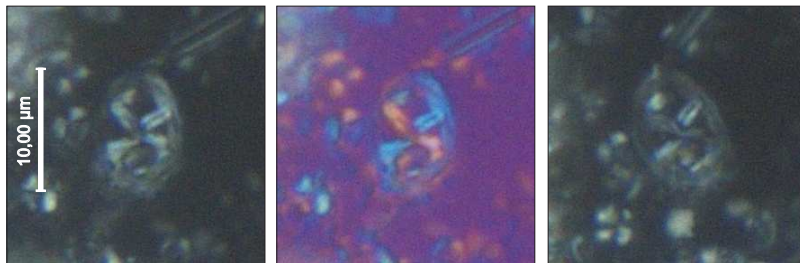
Zeugrhabdotus embergeri (DSDP Site 364 / Depth 877,42 mbsf)



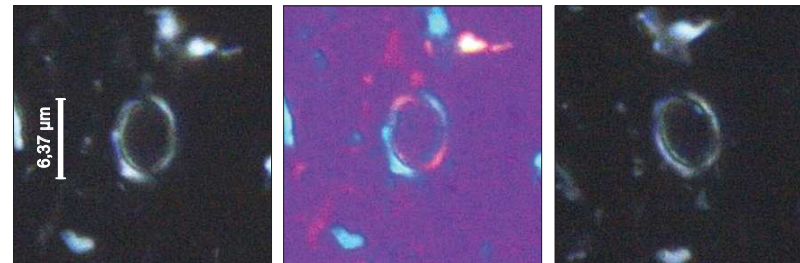
Chiastozygus litterarius (DSDP Site 364 / Depth 1.009,79 mbsf)



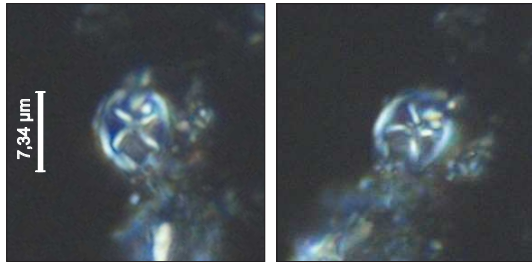
Zeugrhabdotus erectus (DSDP Site 364 / Depth 1032,37 mbsf)



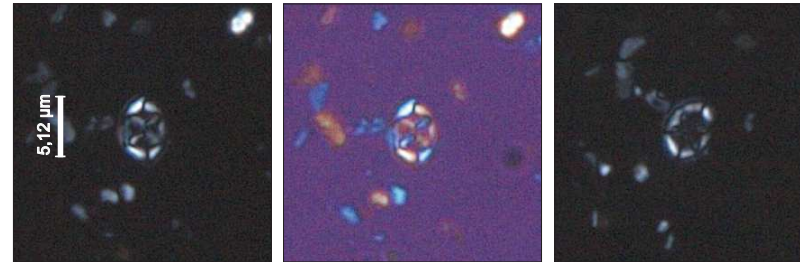
Chiastozygus platyrhethus (DSDP Site 364 / Depth 753,64 mbsf)



Loxolithus armilla (DSDP Site 364 / Depth 932,45 mbsf)



Tegumentum stradneri (DSDP Site 364 / Depth 768,19 mbsf)



Helicolithus trabeculatus (DSDP Site 364 / Depth 768,19 mbsf)



SYSTEMATIC PALEONTOLOGY: PLATE 3 – HETEROCOCCOLITHS

Genus *Eiffellithus* Reinhardt, 1965

Eiffellithus hancockii Burnett, 1997

Description: Small coccoliths; elliptical outline; angular orientation is axial; the central-area is very narrow with an axial cross bar.

Eiffellithus turriseiffelii (Deflandre & Fert, 1954) Reinhardt, 1965

Description: Large to very large coccoliths; elliptical outline; angular orientation is diagonal; the central-area have a diagonal cross bar with arms terminations pointed to blunt and/or forked.

Eiffellithus monechiae Crux, 1991

Description: Medium coccoliths; elliptical outline; angular orientation is axial; the central-area have an axial cross bar with arms terminations pointed to blunt and/or bifurcate. In the diagonal angular orientation the cross bar is subparallel when the ellipse.

Eiffellithus equibramus Watkins & Bergen, 2003

Description: Medium coccoliths; elliptical outline; angular orientation is diagonal; the central-area have a diagonal cross bar with arms terminations pointed to blunt and/or forked. In the diagonal angular orientation the cross bar is subparallel when the ellipse.

Eiffellithus vonsalisiae Watkins & Bergen, 2003

Description: Medium to large coccoliths; elliptical outline; angular orientation is intermediate; the central-area have a diagonal cross bar with arms terminations pointed to blunt and/or forked.

Eiffellithus parvus Watkins & Bergen, 2003

Description: Medium coccoliths; elliptical outline; angular orientation is diagonal; the central-area have a diagonal cross bar with arms terminations pointed to blunt and/or forked.

Eiffellithus praestigium Watkins & Bergen, 2003

Description: Medium coccoliths; elliptical outline; angular orientation is axial; the central-area have an axial cross bar with arms terminations pointed to blunt, forked and/or bifurcate. In the diagonal angular orientation the cross bar is diagonal.

Eiffellithus casulus Shamrock & Watkins, 2009

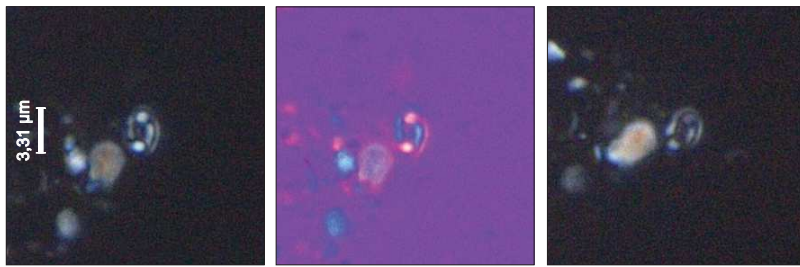
Description: Small to medium coccoliths; elliptical outline; angular orientation is diagonal; the central-area have a diagonal cross bar with arms terminations pointed to blunt and/or forked. The cross bar suture is weak.

Eiffellithus perch-nielseniae Shamrock & Watkins, 2009

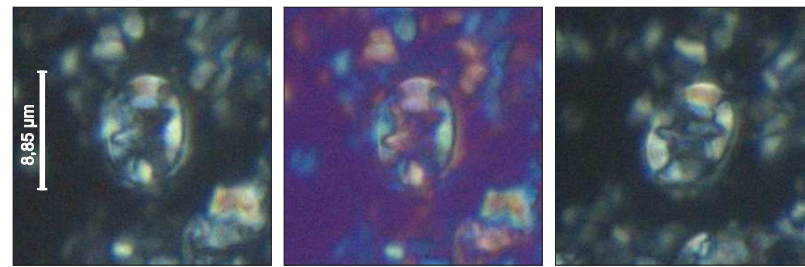
Description: Medium to large coccoliths; elliptical outline; angular orientation is intermediate; the central-area have a diagonal cross bar with arms terminations pointed to blunt, bifurcate and/or forked. The cross bar suture is weak.

Eiffellithus collis Hoffmann, 1970

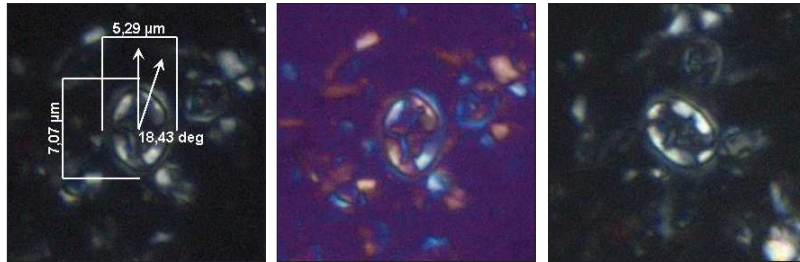
Description: Specimen have 7.45 μ m length, is a medium coccoliths; elliptical outline; angular orientation is diagonal, with 52.55°; the central-area have a diagonal cross bar with arms terminations pointed to blunt. In the diagonal angular orientation the cross bar is axial. Axial ratio is normal (specimen example: 1.4).



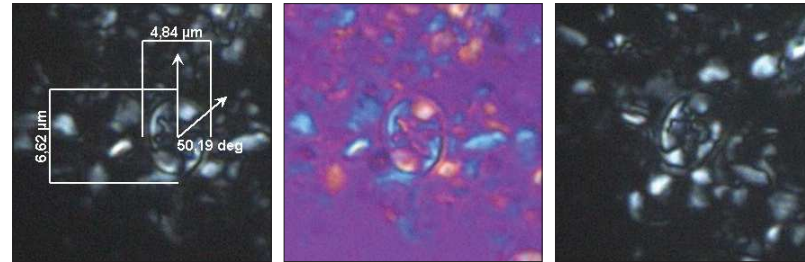
Eiffellithus hancockii (DSDP Site 364 / Depth 1.032,37 mbsf)



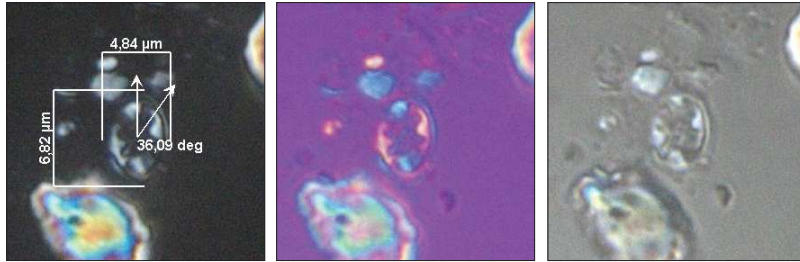
Eiffellithus turriseiffelii (DSDP Site 364 / Depth 772,34 mbsf)



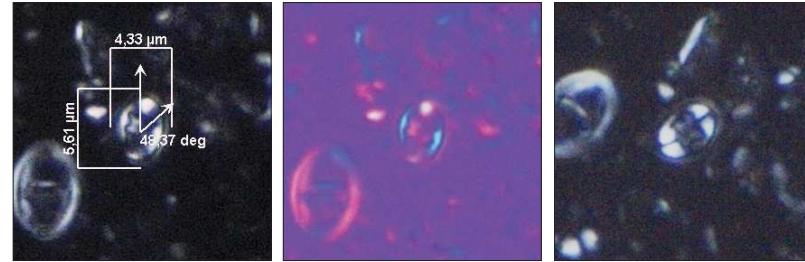
Eiffellithus monechiai (DSDP Site 364 / Depth 875,96 mbsf)



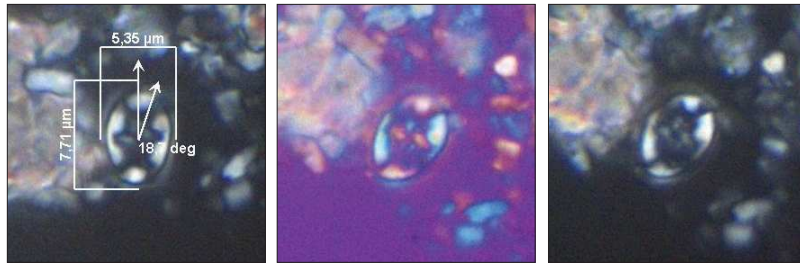
Eiffellithus equibiramus (DSDP Site 364 / Depth 826,40 mbsf)



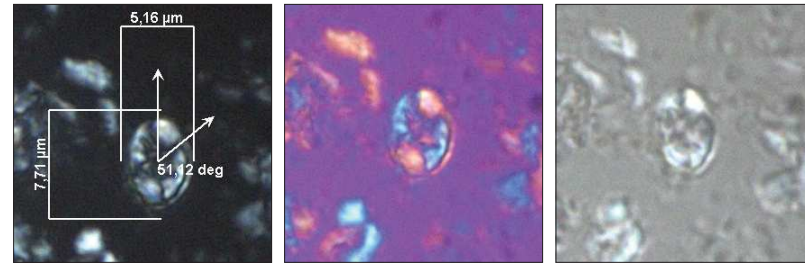
Eiffellithus vonsalisiae (DSDP Site 364 / Depth 826,40 mbsf)



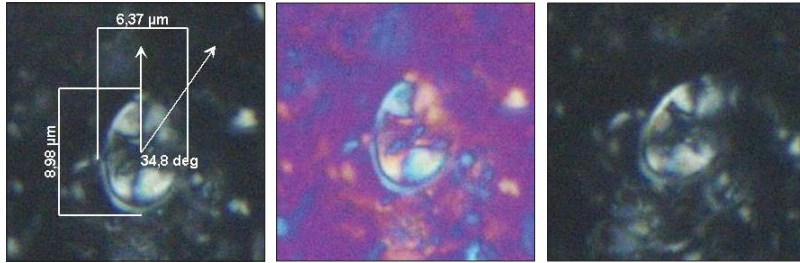
Eiffellithus parvus (DSDP Site 364 / Depth 807,40 mbsf)



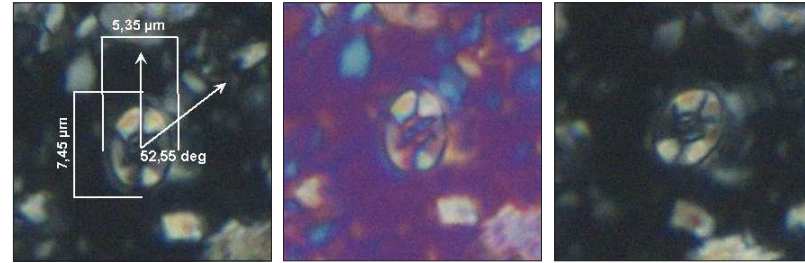
Eiffellithus praestigium (DSDP Site 364 / Depth 829,46 mbsf)



Eiffellithus casulus (DSDP Site 364 / Depth 772,34 mbsf)



Eiffellithus perch-nielseniae (DSDP Site 364 / Depth 719,15 mbsf)



Eiffellithus collis (DSDP Site 364 / Depth 752,37 mbsf)

10 μm

SYSTEMATIC PALEONTOLOGY: PLATE 4 – HETEROCOCCOLITHS

Family **RHAGODISCACEAE** Hay, 1977

Genus *Rhagodiscus* Reinhardt, 1967

Rhagodiscus asper (Stradner, 1963) Reinhardt, 1967

Description: Medium coccoliths; elliptical outline; the rim is unicyclic; the central-area is wide covered by a plate that contains a granulated area and a large hole in the center.

Rhagodiscus angustus (Stradner, 1963) Reinhardt, 1971

Description: Small to medium coccoliths; long-elliptical outline; the rim is unicyclic; the central-area is wide covered by a plate that contains a large hole in the center.

Rhagodiscus achlyostaurion (Hill, 1976) Doeven, 1983

Description: Small to medium coccoliths; elliptical outline; the rim is bicyclic, and the external cycle is greater and with high birefringence; the central-area is wide covered by a plate that contains a large hole in the center with high birefringence.

Order **ARKHANGELSKIALES** Bown & Hampton, 1997 in Bown & Young, 1997

Family **ARKHANGELSKIACEAE** Bukry, 1969 emend Bown & Hampton, 1997 in Bown & Young, 1997

Genus *Broinsonia* Bukry, 1969

Broinsonia signata (Noël, 1969) Noël, 1970

Description: Medium coccoliths; elliptical outline; the rim is usually bicyclic; the central-area is broad and have a central cross with open quadrants, the central cross have arms with a line of extinction that is formed by a suture.

Broinsonia enormis (Shumenko, 1968) Manivit, 1971

Description: Small to medium coccoliths; elliptical outline; the rim is usually unicyclic; the central-area have a central cross with open quadrants.

Family **KAMPTNERIACEAE** Bown & Hampton, 1997 in Bown & Young, 1997

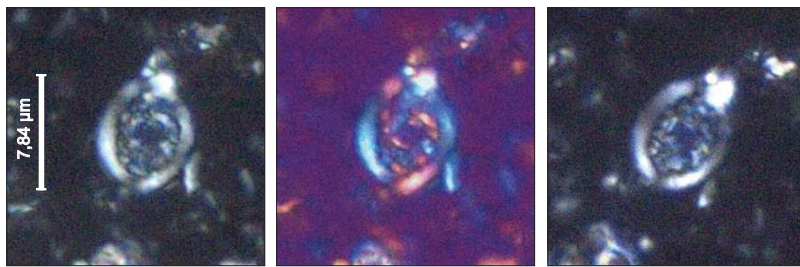
Genus *Gartnerago* Bukry, 1969

Gartnerago stenostaurion (Hill, 1976) Perch-Nielsen, 1984

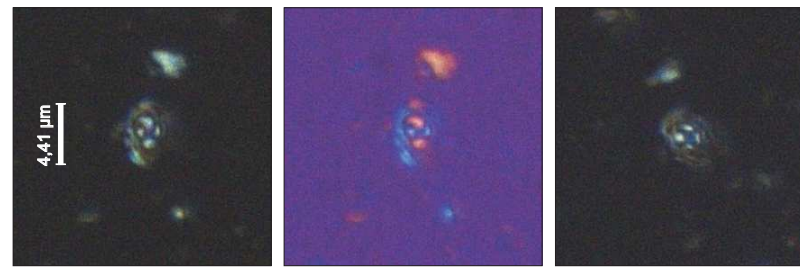
Description: Small to large coccoliths; elliptical outline; the rim is unicyclic and narrow; the central-area is broad and have a plate usually crossed by four radial extinction lines.

Gartnerago sp. 1

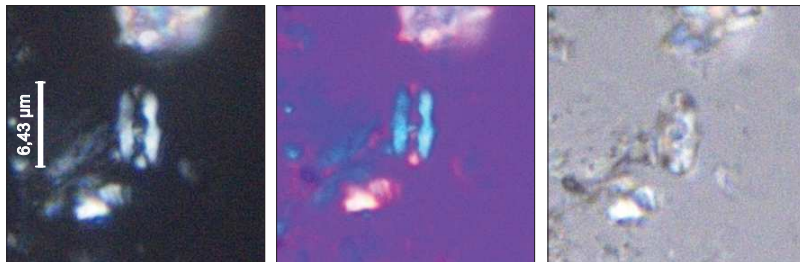
Description: Small to large coccoliths; elliptical outline; the rim is unicyclic and narrow; the central-area is broad and have a plate usually crossed by four radial extinction lines.



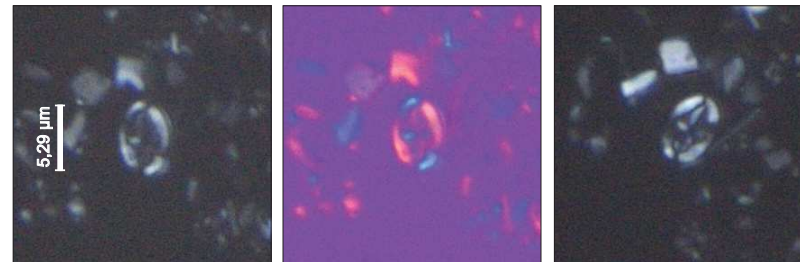
Rhagodiscus asper (DSDP Site 364 / Depth 894,99 mbsf)



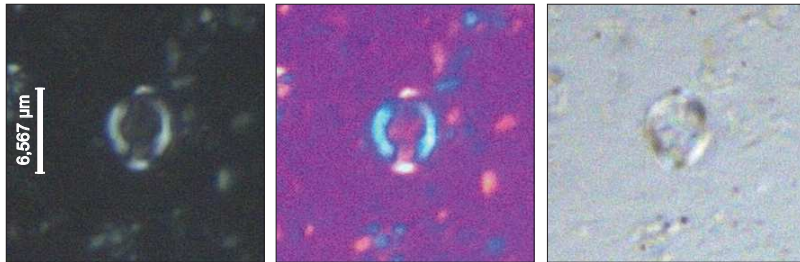
Rhagodiscus achlyostaurion (DSDP Site 364 / Depth 932,45 mbsf)



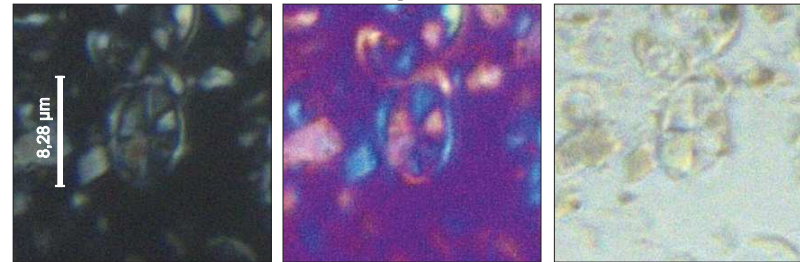
Rhagodiscus angustus (DSDP Site 364 / Depth 1.032,37 mbsf)



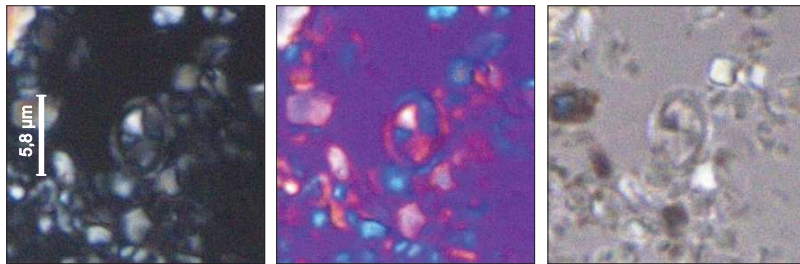
Broinsonia enormis (DSDP Site 364 / Depth 771,17 mbsf)



Broinsonia signata (DSDP Site 364 / Depth 722,75 mbsf)



Gartnerago stenostaurion (DSDP Site 364 / Depth 769,33 mbsf)



Gartnerago sp. 1 (DSDP Site 364 / Depth 934,44 mbsf)

SYSTEMATIC PALEONTOLOGY: PLATE 5 – HETEROCOCCOLITHS

Order STEPHANOLITHIALES Bown & Young, 1997

Family STEPHANOLITHIACEAE Black 1968

Genus *Stoverius* Perch-Nielsen, 1986

Stoverius achylosus (Stover, 1966) Perch-Nielsen, 1986

Description: Small to large; elliptical to circular outline; rim is bicyclic; central-area is spanned by cross bars that meet at a high angle.

Genus *Cylindralithus* Bramlette & Martini, 1964

Corollithion signum Stradner, 1963

Description: Small; hexagonal outline; rim bicyclic with inner cycle with high birefringence; central-area has an axial cross; bar is composed of two elements.

Cylindralithus nudus Bukry, 1969

Description: Small to medium; elliptical; rim is bicyclic; central-area is narrow and open.

Cylindralithus serratus Bramlette & Martini, 1964

Description: Medium; circular; rim bicyclic and serrated; central-area is narrow and open.

Genus *Corollithion* Stradner, 1961

Order PODORHABDALES Rood *et al.*, 1971 emend. Bown, 1987

Family AXOPODORHABDACEAE Wind & Wise in Wise & Wind, 1977

Genus *Axopodorhabdus* Wind & Wise in Wise & Wind, 1977

Axopodorhabdus biramiculatus (Stover 1966) Corbett & Watkins, 2014

Description: medium to large; elliptical; rim is bicyclic with inner cycle with high birefringence; central-area have an axial cross, with 4 almost circular openings to its sides.

Tetrapodorhabdus coptensis Black, 1971

Description: medium to large; elliptical outline; rim is bicyclic; central-area is broad and have a diagonal cross bar, with four almost circular openings to its sides.

Family BISCUTACEAE Black, 1971

Genus *Biscutum* Black in Black & Barnes, 1959

Biscutum ubiquem Brace & Watkins, 2014

Description: small to medium; elliptical; rim is bicyclic, inner cycle is thick; central-area is small and void.

Biscutum sp. 1

Description: Medium; elliptical outline; the rim is unicyclic; the central-area is broad, closed and have two bars transverse.

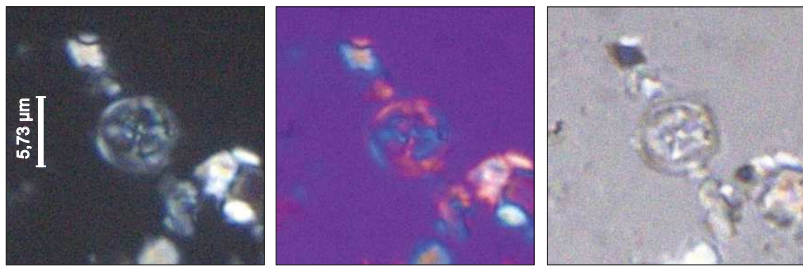
Biscutum sp. 2

Description: Medium; elliptical; rim unicyclic; have 2 bars transverse in the center.

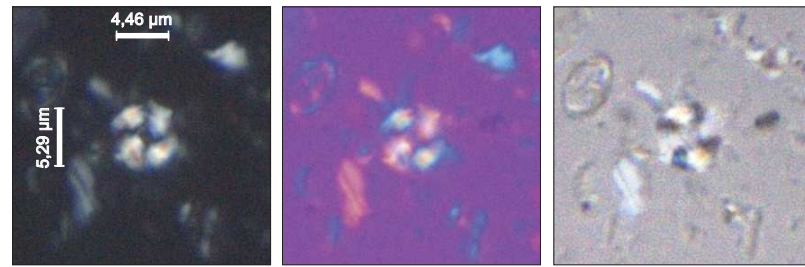
Genus *Discorhabdus* Noël, 1965

Discorhabdus ignotus (Górka, 1957) Perch-Nielsen, 1968

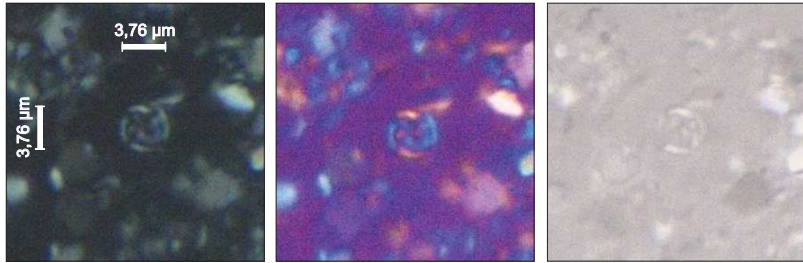
Description: Very small to medium; circular; rim is bicyclic; central-area is closed.



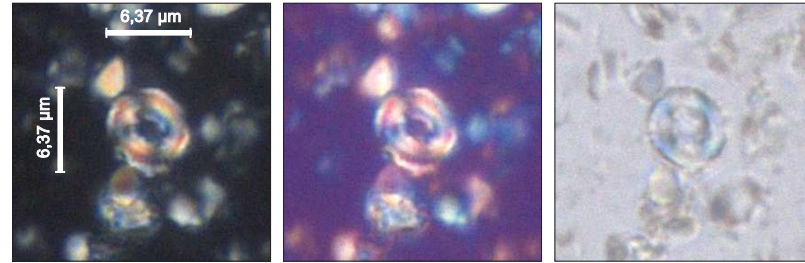
Stoverius achylosus (DSDP Site 364 / Depth 1.029,50 mbsf)



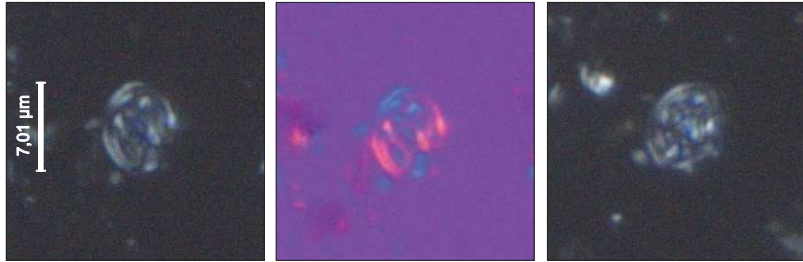
Cylindralithus nudus (DSDP Site 364 / Depth 935,98 mbsf)



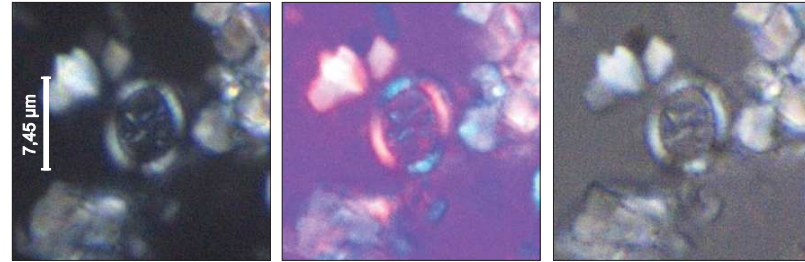
Corollithion signum (DSDP Site 364 / Depth 752,37 mbsf)



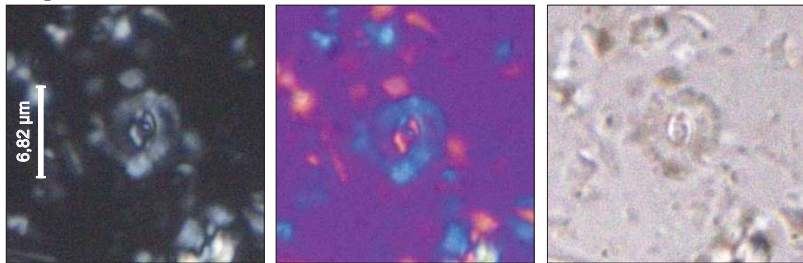
Cylindralithus serratus (DSDP Site 364 / Depth 911,25 mbsf)



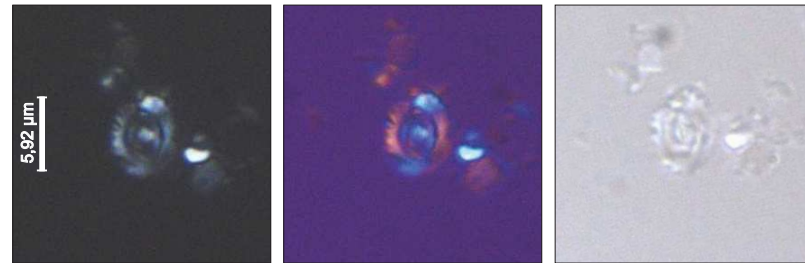
Axopodorhabdus biramiculatus (DSDP Site 364 / Depth 951,47 mbsf)



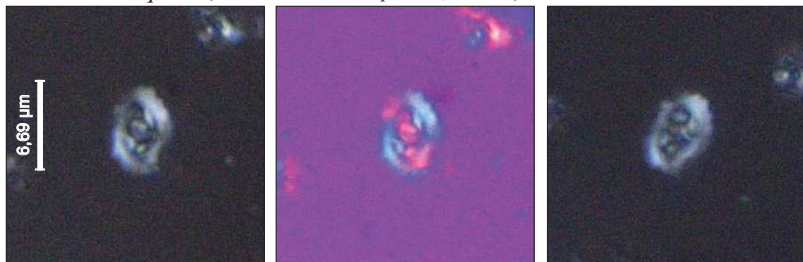
Tetrapodorhabdus coptensis (DSDP Site 364 / Depth 790,41 mbsf)



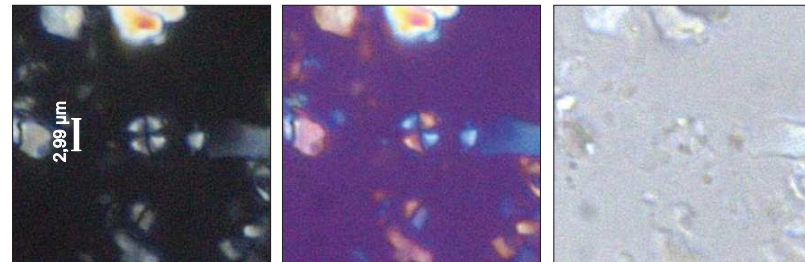
Biscutum ubiqueum (DSDP Site 364 / Depth 911,25 mbsf)



Biscutum sp. 1 (DSDP Site 364 / Depth 935,98 mbsf)



Biscutum sp. 2 (DSDP Site 364 / Depth 810,94 mbsf)



Discorhabdus ignotus (DSDP Site 364 / Depth 892,14 mbsf)

10 µm

SYSTEMATIC PALEONTOLOGY: PLATE 6 – HETEROCOCCOLITHS

Genus *Sollasites* Black, 1967

Sollasites sp. 1

Description: Medium to large; elliptical; rim is unicyclic; central-area is closed with axial plates.

Family **CRETARHABDACEAE** Thierstein, 1973

Genus *Cretarhabdus* Bramlette & Martini, 1964

Cretarhabdus crenulatus Bramlette & Martini (1964)

Description: medium to large coccoliths; elliptical outline; the rim is unicyclic with high birefringence; the central-area is broad with a two lateral bars in each quadrant.

Cretarhabdus surirellus (Deflandre and Fert 1954) Reinhardt, 1970

Description: medium to large coccoliths; elliptical outline; the rim is unicyclic with high birefringence; the central-area is broad with a three lateral bars in each quadrant.

Cretarhabdus conicus Bramlette & Martini, 1964

Description: medium to large coccoliths; elliptical outline; the rim is bicyclic; the central-area is perforated by many pores, and have a cross bar with pointed tips.

Cretarhabdus striatus (Stradner, 1963) Black, 1973

Description: medium to large coccoliths; elliptical outline; the rim is bicyclic; the central-area is wide and have a central cross, and a double marginal plates with numerous pores are arranged in lines which meet at oblique angles at the straight central cross.

Cretarhabdus inaequalis Crux, 1987

Description: Large coccoliths; elliptical outline; the rim is bicyclic; the central-area have an irregular grill.

Genus *Retecapsa* Black, 1971

Retecapsa angustiforata Black, 1971

Description: medium to large coccoliths; elliptical outline; the rim is bicyclic; the central-area is wide and have an axial cross with one broad lateral bar in each quadrant, the 8 central area holes are of relatively equal size.

Genus *Helenea* Worsley, 1971

Helenea chiastia Worsley, 1971

Description: medium to large coccoliths; elliptical to subcircular outline; the rim is bicyclic, with inner cycle high birefringence; the central-area is narrow and have an axial cross.

Genus *Flabellites* Thierstein, 1973

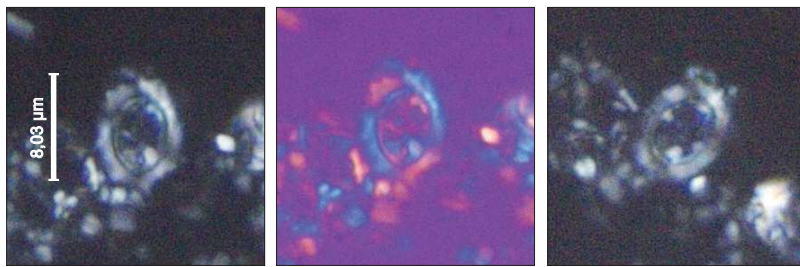
Flabellites oblongus (Bukry, 1969) Crux in Crux *et al.*, 1982

Description: medium to large coccoliths; elliptical outline; the rim is bicyclic; the central-area is granular with a bridge formed by four small calcite crystals arranged in a diagonal cross.

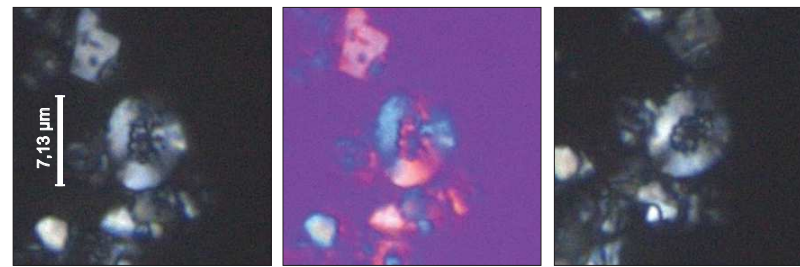
Genus *Grantarhabdus* Black, 1971

Grantarhabdus coronadventis (Reinhardt, 1966) Grün and Allemann, 1975

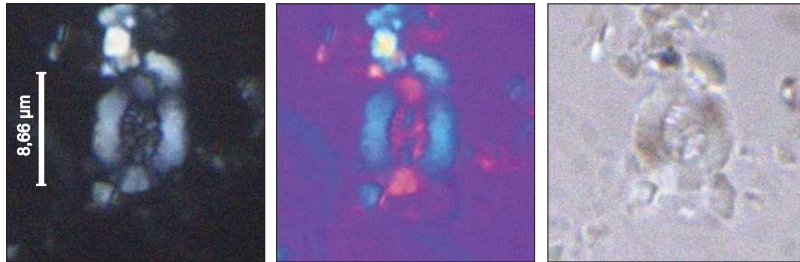
Description: medium to large coccoliths; subcircular outline; the rim is bicyclic; the central-area is have a diagonal cross bars that meet at a high angle.



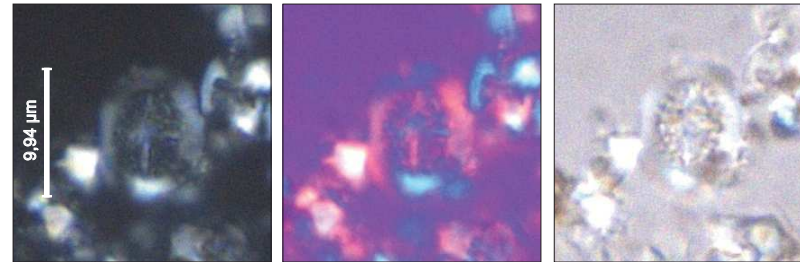
Sollasites sp. 1 (DSDP Site 364 / Depth 934,44 mbsf)



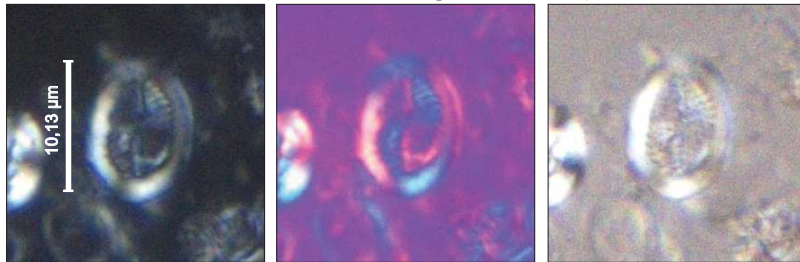
Cretarhabdus crenulatus (DSDP Site 364 / Depth 973,89 mbsf)



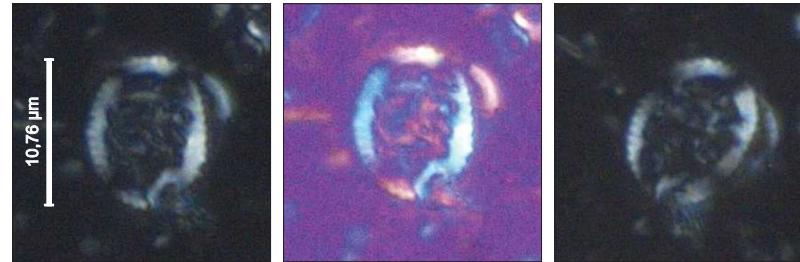
Cretarhabdus surirellus (DSDP Site 364 / Depth 1.031,21 mbsf)



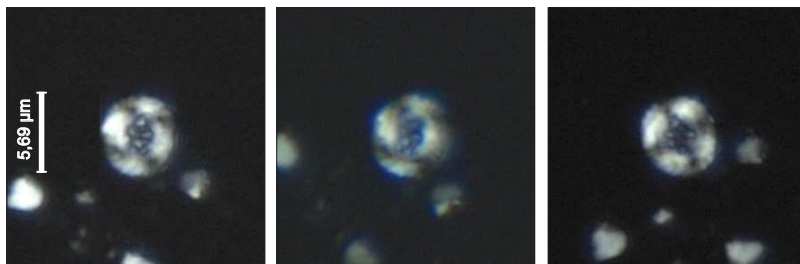
Cretarhabdus conicus (DSDP Site 364 / Depth 951,47 mbsf)



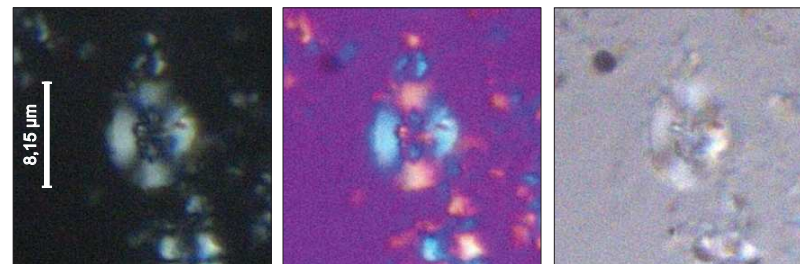
Cretarhabdus striatus (DSDP Site 364 / Depth 788,94 mbsf)



Cretarhabdus inaequalis (DSDP Site 364 / Depth 932,45 mbsf)



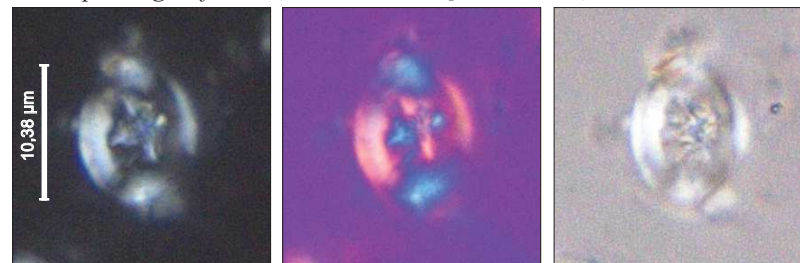
Helenea chiastia (DSDP Site 364 / Depth 726,66 mbsf)



Retecapsa angustiforata (DSDP Site 364 / Depth 726,66 mbsf)



Flabellites oblongus (DSDP Site 364 / Depth 892,96 mbsf)



Grantarhabdus coronadventis (DSDP Site 364 / Depth 790,41 mbsf)

10 μm

SYSTEMATIC PALEONTOLOGY: PLATE 7 – HETEROCOCCOLITHS

Family **PREDISCOSPHAERACEAE** Rood *et al.*, 1971

Genus *Prediscosphaera* Vekshina, 1959

Prediscosphaera columnata (Stover, 1966) Perch-Nielsen, 1984

Description: Medium to large coccoliths; circular to subcircular outline; the rim is bicyclic, distal cycle is broad and the inner cycle very narrow; the central-area have a cross bar.

Prediscosphaera spinosa (Bramlette & Martini, 1964) Gartner, 1968

Description: Medium to large coccoliths; elliptic outline; the rim is bicyclic, distal cycle is broad and the inner cycle very narrow; the central-area have an axial cross bar.

Prediscosphaera cretacea (Arkhangelsky, 1912) Gartner, 1968

Description: Medium to large coccoliths; elliptic outline; the rim is bicyclic, distal cycle is broad and the inner cycle very narrow; the central-area have a diagonal cross bar.

Family **TUBODISCACEAE** Bown & Rutledge, 1997 in Bown & Young, 1997

Genus *Manivitella* Thierstein, 1971

Manivitella pemmatoidea (Deflandre in Manivit, 1965) Thierstein, 1971

Description: Medium to large coccoliths; Ellipse outline; the rim is bicyclic, the inner cycle brighter; the central-area is broad and void.

Order **WATZNAUERIALES** Bown, 1987

Family **WATZNAUERiaceae** Rood *et al.*, 1971

Genus *Watznaueria* Reinhardt, 1964

Watznaueria barnesiae (Black in Black & Barnes, 1959) Perch-Nielsen, 1968

Description: medium coccoliths; elliptical outline; the rim is unicyclic; the central-area is closed or very narrow, with no central area structures.

Watznaueria britannica (Stradner, 1963) Reinhardt, 1964

Description: medium coccoliths; elliptical outline; the rim is unicyclic; the central-area is narrow with spanned by a transverse bar.

Watznaueria biporta Bukry, 1969

Description: medium to large coccoliths; elliptical outline; the rim is unicyclic; the central-area have two well-defined pores aligned in the long axis.

Watznaueria manivittiae Bukry, 1973

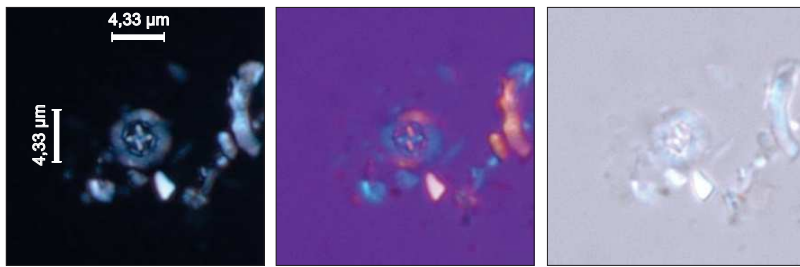
Description: large coccoliths; elliptical outline; the rim is unicyclic; the central-area is closed.

Watznaueria supracretacea (Reinhardt, 1964) Wind & Wise, 1977

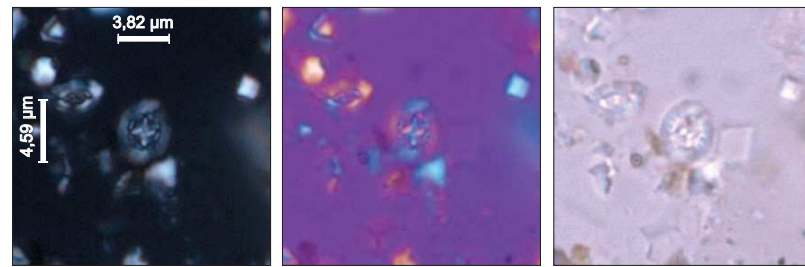
Description: Small to medium coccoliths; elliptical outline; the rim is unicyclic serrate; the central-area is closed with spanned by a transverse bar connecting the sides of the ellipse.

Watznaueria ovata Bukry, 1969

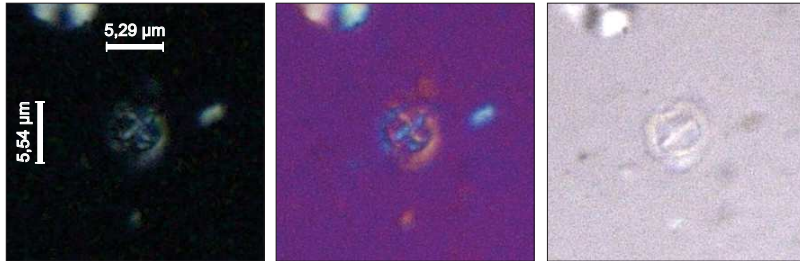
Description: medium; elliptical; the rim is bicyclic; the central-area is broad and void.



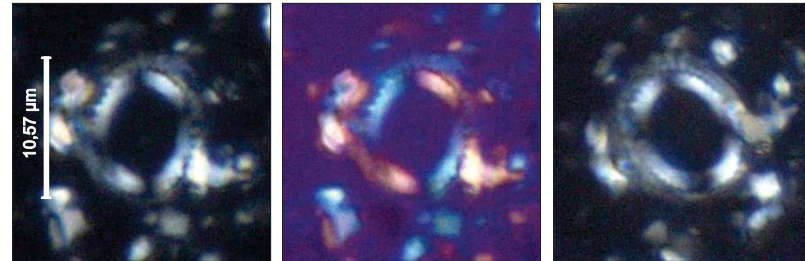
Prediscosphaera columnata (DSDP Site 364 / Depth 892,96 mbsf)



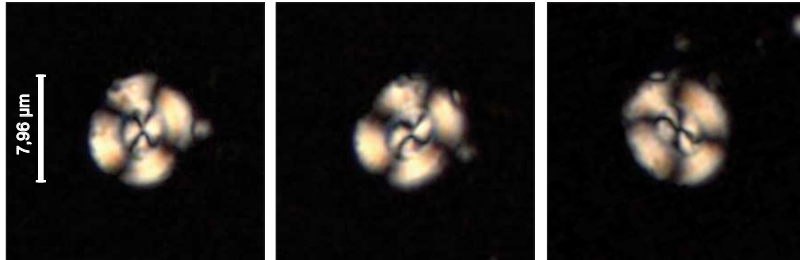
Prediscosphaera spinosa (DSDP Site 364 / Depth 1006,96 mbsf)



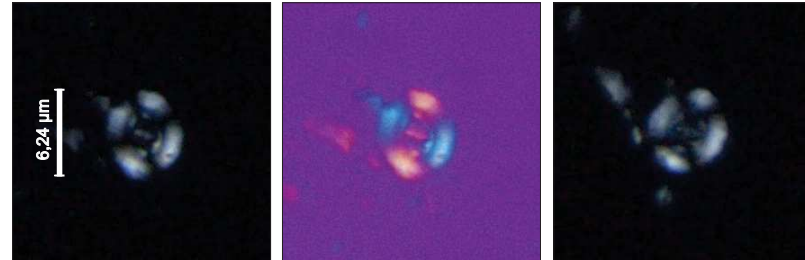
Prediscosphaera cretacea (DSDP Site 364 / Depth 725,24 mbsf)



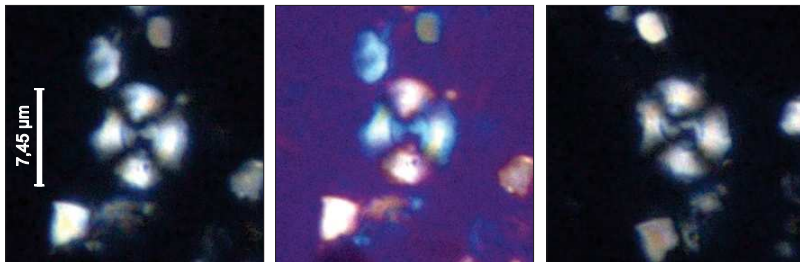
Manivitella pemmatoidea (DSDP Site 364 / Depth 911,25 mbsf)



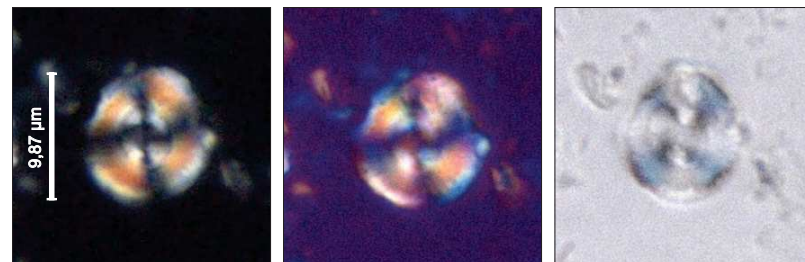
Watznaueria barnesiae (DSDP Site 364 / Depth 829,46 mbsf)



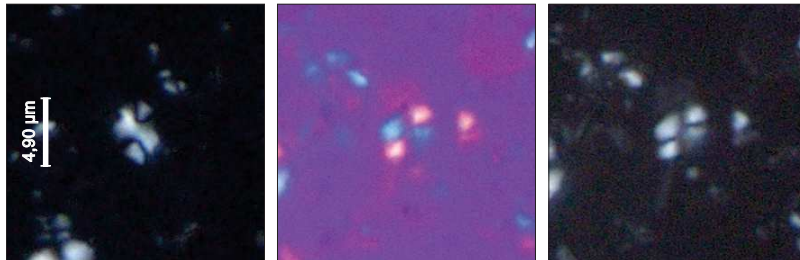
Watznaueria britannica (DSDP Site 364 / Depth 1006,96 mbsf)



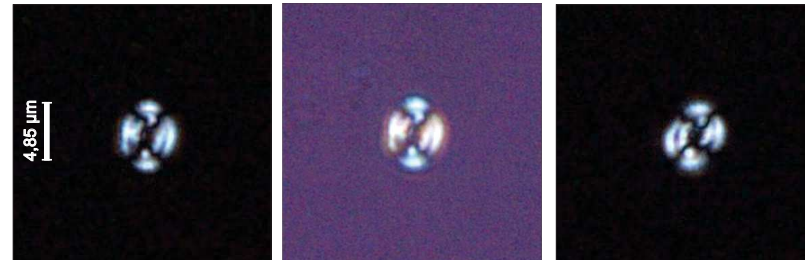
Watznaueria biporta (DSDP Site 364 / Depth 911,25 mbsf)



Watznaueria manivittiae (DSDP Site 364 / Depth 877,42 mbsf)



Watznaueria supracretacea (DSDP Site 364 / Depth 1027,24 mbsf)



Watznaueria ovata (DSDP Site 364 / Depth 719,15 mbsf)

10 μm

SYSTEMATIC PALEONTOLOGY: PLATE 8 – HETEROCOCCOLITHS

Genus *Cyclagelosphaera* Noël, 1965

Cyclagelosphaera rotaclypeata Bukry, 1969

Description: small to medium coccoliths; elliptical outline; the rim is bicyclic; the central-area is filled with radial elements.

Cyclagelosphaera margerelii Noël, 1965

Description: small to medium coccoliths; elliptical outline; the rim is bicyclic; the central-area is closed.

HETEROCOCCOLITHS PLACOLITHS INC SEDIS

Genus *Haqius* Roth, 1978

Haqius circumradiatus (Stover, 1966) Roth, 1978

Description: Medium to large; circular outline; the rim is bicyclic; the central-area is narrow or closed.

NANNOLITHS INC SED - RADIATE MULTIELEMENT

Genus *Hayesites* Manivit, 1971

Hayesites albiensis Manivit, 1971

Description: Small; stellate outline with 6-8 rays, the central-area is narrow containing a circle.

Hayesites irregularis (Thierstein in Roth & Thierstein, 1972) Covington & Wise, 1987

Description: Small; stellate outline with 9-11 rays, the central-area is narrow containing a circle.

Genus *Assipetra* Roth, 1973

Assipetra terebrodentarius (Applegate et al. in Covington & Wise, 1987) Bergen, 1994

Description: Medium to large; globular formed by complexly blocks in radial sutures.

Family **MICRORHABDULACEAE** Deflandre, 1963

Genus *Lithraphidites* Deflandre, 1963

Lithraphidites alatus Roth & Thierstein, 1972

Description: Medium to large; the overall shape is like a closed umbrella.

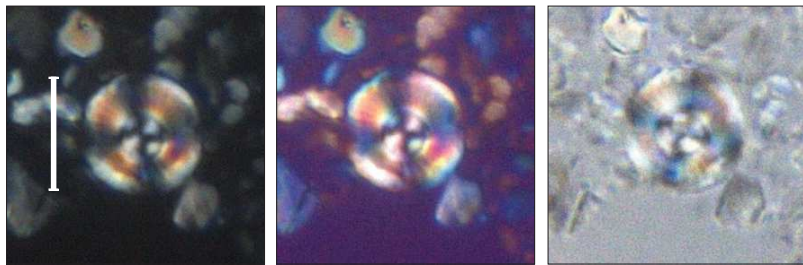
Lithraphidites carniolensis Deflandre, 1963

Description: Medium to large; rod-shaped nannoliths, that tapers towards both ends.

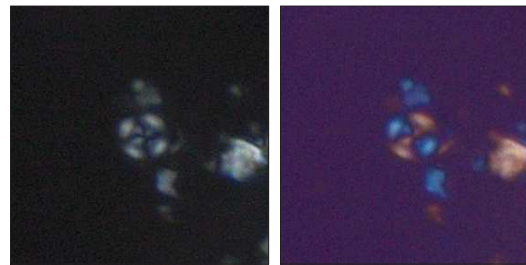
Genus *Microrhabdulus* Deflandre, 1959

Microrhabdus primitivus Troelsen & Quadros, 1971

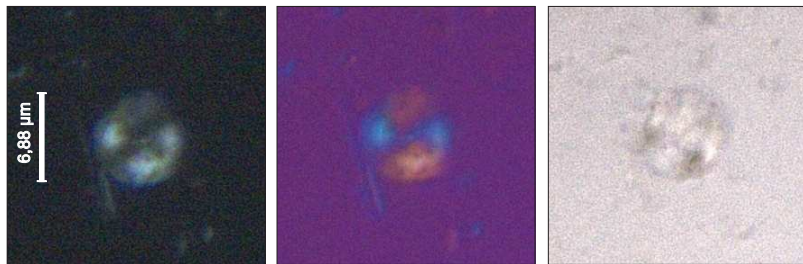
Description: Medium to large; cylindrical shape with nearly parallel sided, shows chequerboard extinction.



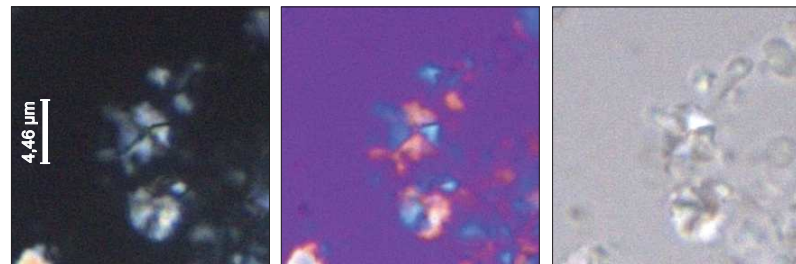
Cyclagelosphaera margerelii (DSDP Site 364 / Depth 912,02 mbsf)



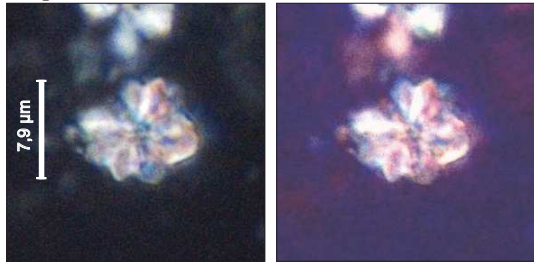
Cyclagelosphaera rotaclypeata (DSDP Site 364 / Depth 912,02 mbsf)



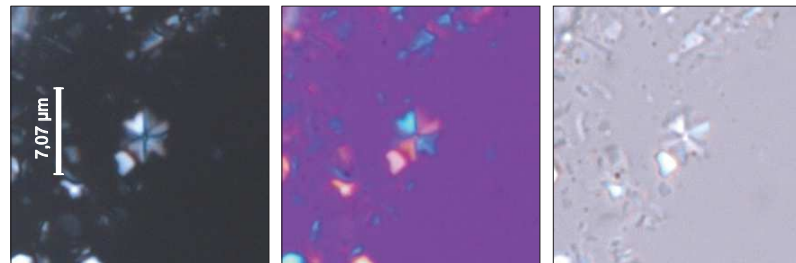
Haqius circumradiatus (DSDP Site 364 / Depth 722,75 mbsf)



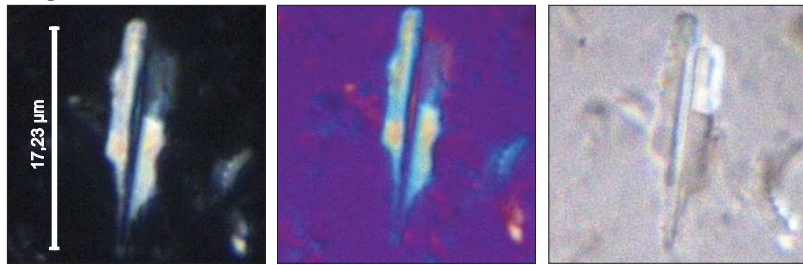
Hayesites albiensis (DSDP Site 364 / Depth 1.031,21 mbsf)



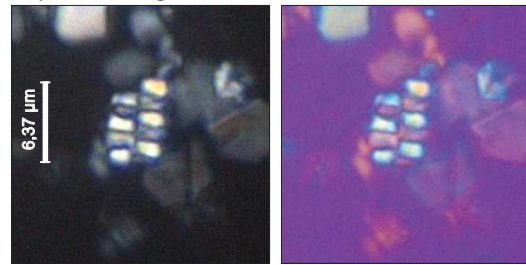
Assipetra terebrodentarius (DSDP Site 364 / Depth 931,70 mbsf)



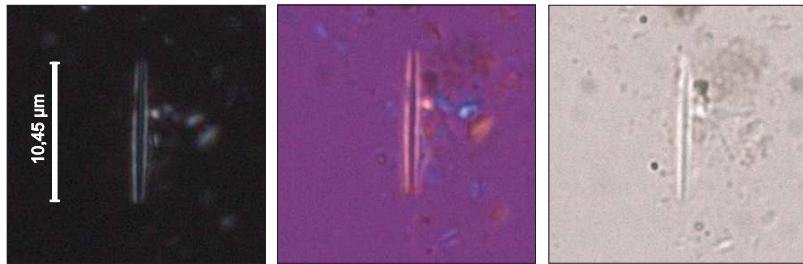
Hayesites irregularis (DSDP Site 364 / Depth 850,15 mbsf)



Lithraphidites alatus (DSDP Site 364 / Depth 932,45 mbsf)



Microrhabdus primitivus (DSDP Site 364 / Depth 828,40 mbsf)



Lithraphidites carniolensis (DSDP Site 364 / Depth 717,23 mbsf)

10 μm

SYSTEMATIC PALEONTOLOGY: PLATE 9 – NANNOLITHS INC SED

Family LAPIDEACASSACEAE Black, 1971

Genus *Lapideacassis* Black, 1971

Lapideacassis sp. 1

Description: Large; cylindrical with tapering outline; curved upper part; central area is diffuse; without apical spine.

Lapideacassis sp. 2

Description: Large; cylindrical with tapering outline; curved upper part; central area is diffuse; without apical spine and protruding ledge just above the base on one side.

Lapideacassis sp. 3

Description: Large; cylindrical with tapering outline, form of bell; curved upper part; central area is diffuse; without apical spine.

Lapideacassis sp. 4

Description: Medium; circular outline, apical view of *Lapideacassis*.

Lapideacassis sp. 5

Description: Large to very large; cylindrical with tapering outline; curved upper part; central area is diffuse; on one side have two apical spines oriented at about 45 degrees.

Lapideacassis glans Black, 1971

Description: Small species; cylindrical with tapering outline, form of bell; curved upper part; central area is diffuse; with protruding ledge just above the base.

Lapideacassis cf. *bispinosa* (Perch-Nielsen & Franz, 1977) Burnett, 1997

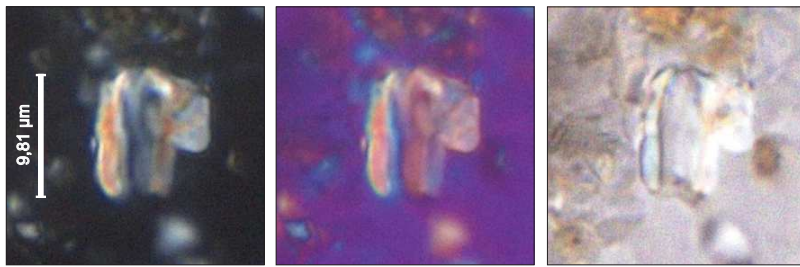
Description: Large to very large; cylindrical with tapering outline; curved upper part; central area is diffuse; have two apical spines at about 45 degrees.

Lapideacassis mariae Black, 1971

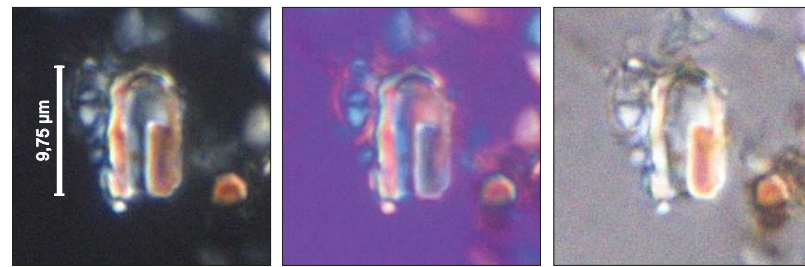
Description: Large species; cylindrical with tapering outline, form of bell; curved upper part; central area is diffuse; without apical spine and protruding ledge just above the base.

Lapideacassis cornuta (Forchheimer & Stradner, 1973) Wise & Wind, 1977

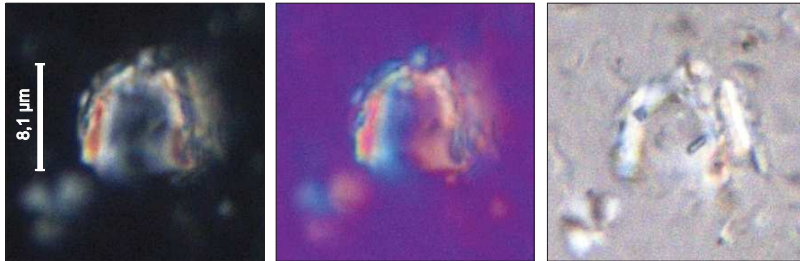
Description: Large to very large; cylindrical with tapering outline; curved upper part; central area is diffuse; have two apical spines at about 45 degrees.



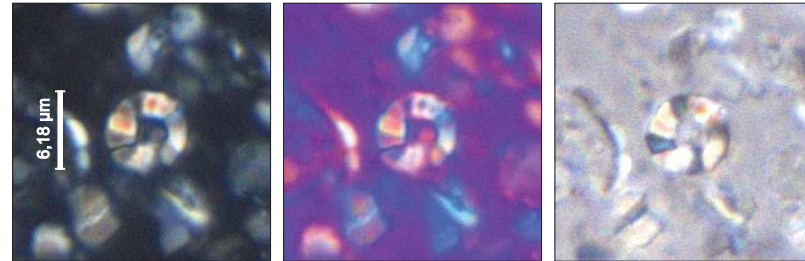
Lapideacassis sp. 1 (DSDP Site 364 / Depth 1.006,96 mbsf)



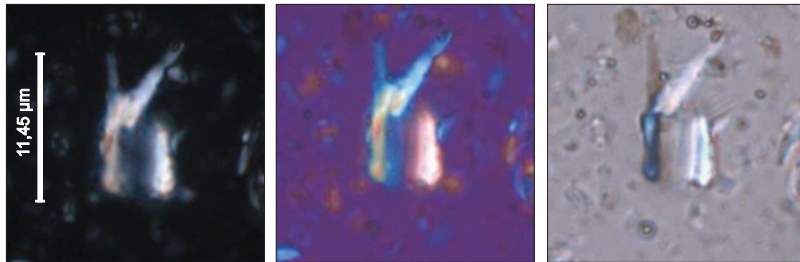
Lapideacassis sp. 2 (DSDP Site 364 / Depth 1006,41 mbsf)



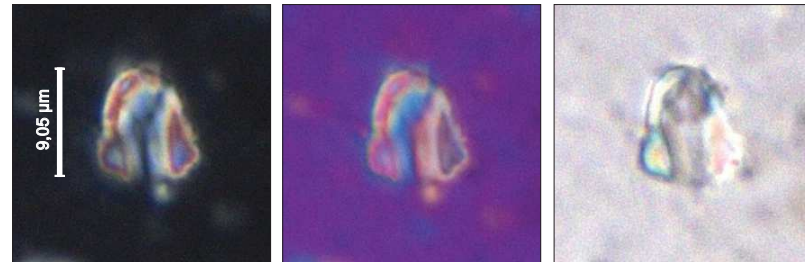
Lapideacassis sp. 3 (DSDP Site 364 / Depth 932,45 mbsf)



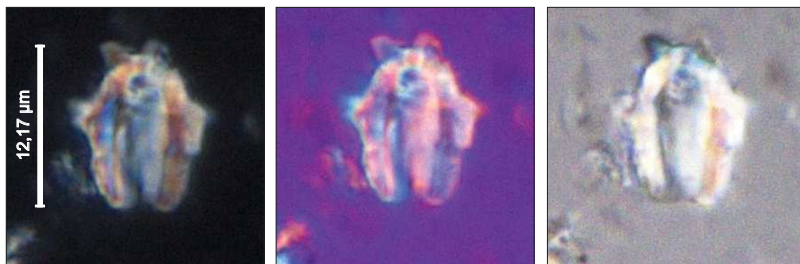
Lapideacassis sp. 4 (DSDP Site 364 / Depth 929,66 mbsf)



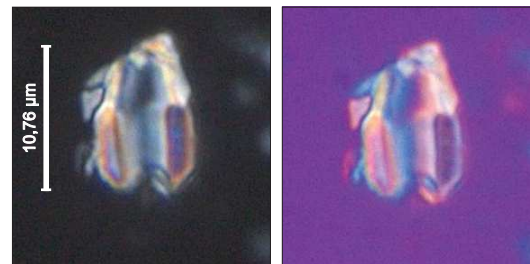
Lapideacassis cf. *bispinosa* (DSDP Site 364 / Depth 932,45 mbsf)



Lapideacassis *glans* (DSDP Site 364 / Depth 932,45 mbsf)



Lapideacassis *cornuta* (DSDP Site 364 / Depth 932,45 mbsf)



Lapideacassis *mariae* (DSDP Site 364 / Depth 971,18 mbsf)

SYSTEMATIC PALEONTOLOGY: PLATE 10 – NANNOLITHS AND BRAARUDOSPHAERALES

Family **POLYCYCLOLITHACEAE** Forchheimer, 1972 emend. Varol, 1992

Genus *Radiolithus* Stover, 1966

Radiolithus undosus (Black, 1973) Varol, 1992

Description: Medium; stellate outline, nannolith constructed from nine segments and have a large diaphragm and small wall.

Radiolithus planus Stover, 1966

Description: Medium; nannolith constructed from nine segments and have a large diaphragm and small wall.

Genus *Eprolithus* Stover, 1966

Eprolithus floralis (Stradner, 1962) Stover, 1966

Description: Medium; nannolith constructed from nine segments and have a medium diaphragm and medium wall.

Genus *Micula* Vekshina, 1959

Order BRAARUDOSPHAERALES Aubry 2013 emend Lees & Bown 2016

Family **BRAARUDOSPHAERACEAE** Deflandre, 1947

Genus *Braarudosphaera* Deflandre, 1947

Braarudosphaera africana Stradner, 1961

Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with long and relatively narrow rays.

Braarudosphaera primula Black, 1973

Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with rounded tips to the segments.

Braarudosphaera hockwoldensis Black, 1973

Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with shallow and narrow rays.

Braarudosphaera pseudobatilliformis Alves *et al.* 2017

Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with one segment asymmetrically enlarged.

Braarudosphaera regularis Black, 1973

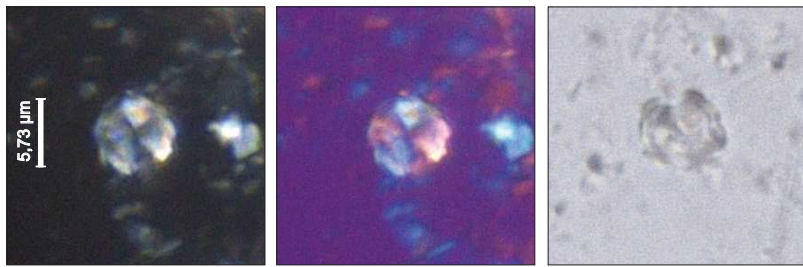
Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with pentagonal outline and bilateral symmetry, i.e. the sutures bisect the edges.

Braarudosphaera batilliformis Troelsen & Quadros, 1971

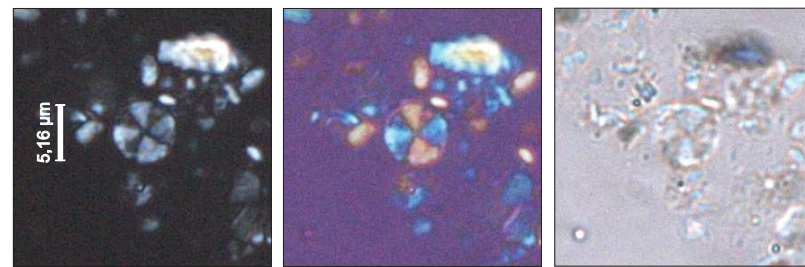
Description: Medium; stellate outline, nannolith constructed from five segments which form a pentalith with one segment bearing a ridge which extends into a protruding spine.

Braarudosphaeraceae sp. 1

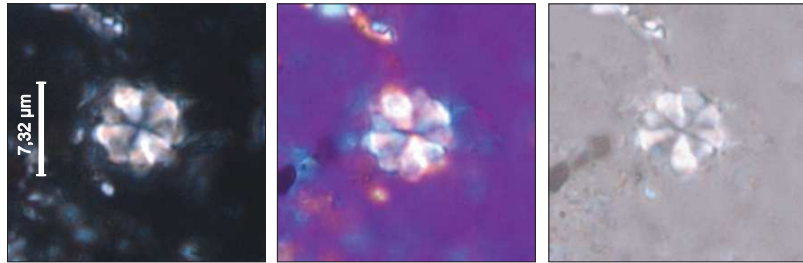
Description: Medium; pentagon outline, nannolith constructed from five triangular segments.



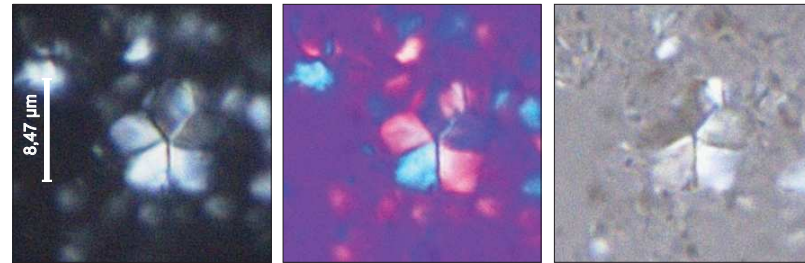
Radiolithus undosus (DSDP Site 364 / Depth 877,42 mbsf)



Radiolithus planus (DSDP Site 364 / Depth 717,23 mbsf)



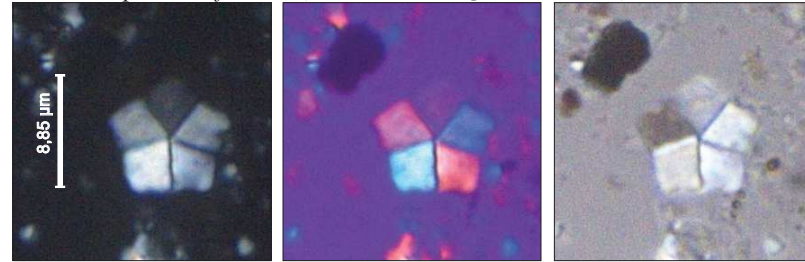
Eproolithus floralis (DSDP Site 364 / Depth 722,75 mbsf)



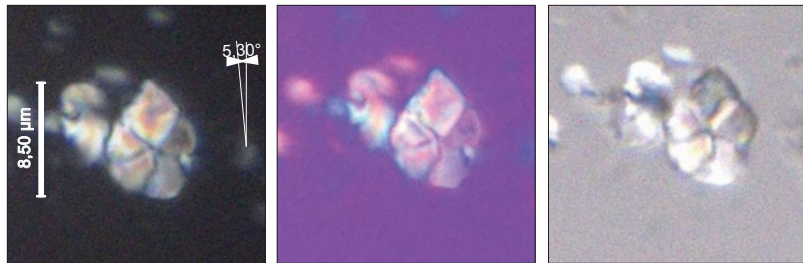
Braarudosphaera africana (DSDP Site 364 / Depth 1.032,37 mbsf)



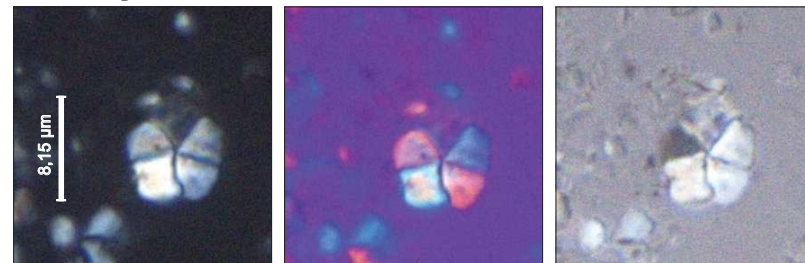
Braarudosphaera primula (DSDP Site 364 / Depth 828,40 mbsf)



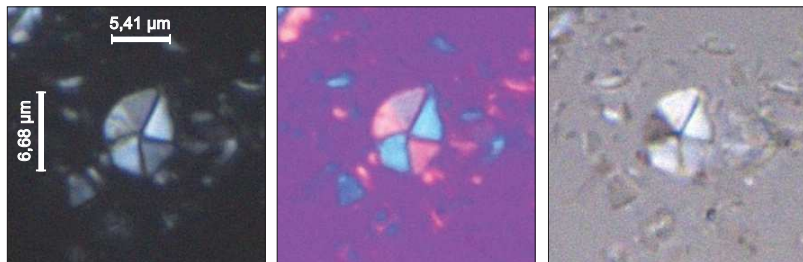
Braarudosphaera hockwoldensis (DSDP Site 364 / Depth 1.032,37 mbsf)



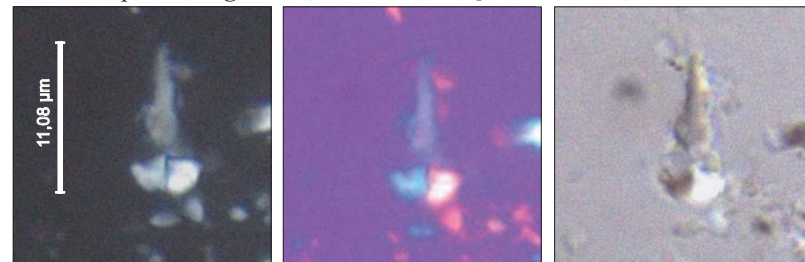
Braarudosphaera pseudobatiliiformis (DSDP Site 364 / Depth 828,40 mbsf)



Braarudosphaera regularis (DSDP Site 364 / Depth 1.032,37 mbsf)



Braarudosphaera sp.1 (DSDP Site 364 / Depth 968,45 mbsf)



Braarudosphaera batiliiformis (DSDP Site 364 / Depth 1.032,37 mbsf)

10 µm

SYSTEMATIC PALEONTOLOGY: PLATE 11 – BRAARUDOSPHAERALES AND CALCAREOUS DINOFLAGELLATES

Order BRAARUDOSPHAERALES Aubry 2013 emend Lees & Bown 2016

Family NANNOCONACEAE Deflandre, 1959

Genus *Nannoconus* Kamptner, 1931

Nannoconus sp. 1

Description: Medium to large; circular outline, apical view of *Nannoconus*.

Nannoconus sp. 2

Description: Medium to large; elliptical outline, oblique angle of *Nannoconus*.

Nannoconus truittii truittii Brönnimann, 1955

Description: Lengths 6-12 μm , medium to very large; width 6-12 μm ; axial channel > thickness of the walls; length > width or length = width; pole termination is only one rounded edges.

Nannoconus fragilis Deres and Achéritéguy, 1980

Description: Lengths 6-8 μm , medium to large; width 7-9 μm ; axial channel > thickness of the walls; length < width; pole termination is only one rounded edges.

Nannoconus truittii frequens Deres and Achéritéguy, 1980

Description: Lengths 11-13 μm , very large; width 8-10 μm ; axial channel > thickness of the walls; length > width or length = width; pole termination is only one rounded edges.

Nannoconus truittii rectangularis Deres and Achéritéguy, 1980

Description: Lengths 6-8 μm , medium to large; width 9-11 μm ; axial channel > thickness of the walls; length < width; pole termination is only one rounded edges or truncated.

Nannoconus troelsenii Alves *et al.*, 2017

Description: medium to large; circular form, central area is closed.

Nannoconus planus Stradner, 1963

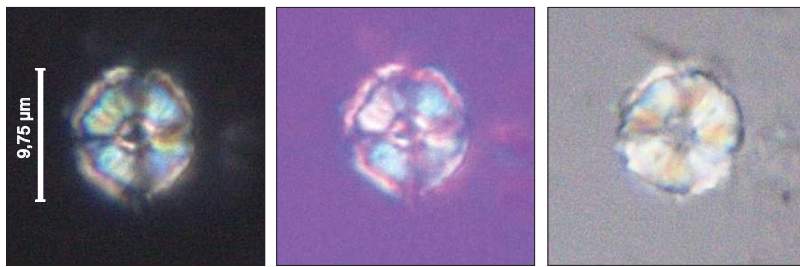
Description: Medium to large; axial channel = thickness of the walls; length > width.

Order CALCAREOUS DINOFLAGELLATES

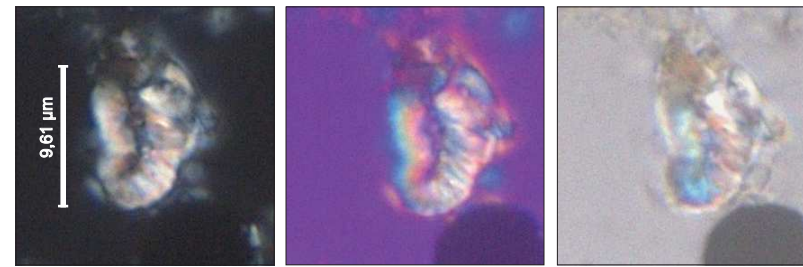
Thoracosphaera Kamptner 1927

Thoracosphaera sp.1

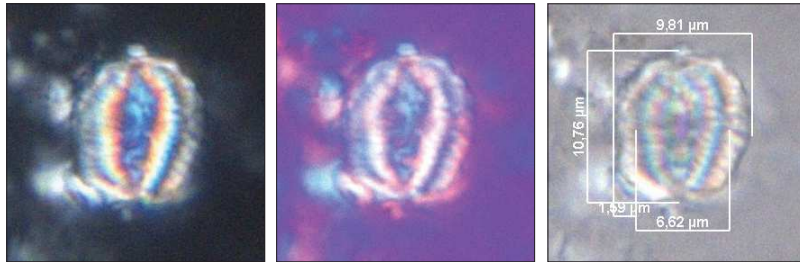
Description: Granular forms, usually circular.



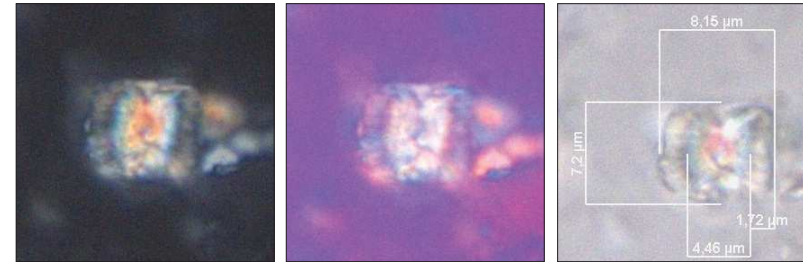
Nannoconus sp. 1 - Apical angle (DSDP Site 364 / Depth 790,41 mbsf)



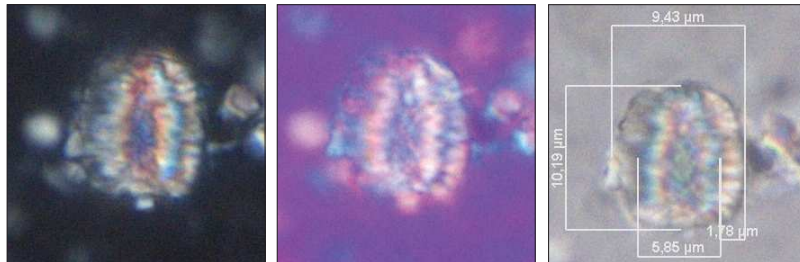
Nannoconus sp. 2 - Oblique angle (DSDP Site 364 / Depth 992,63 mbsf)



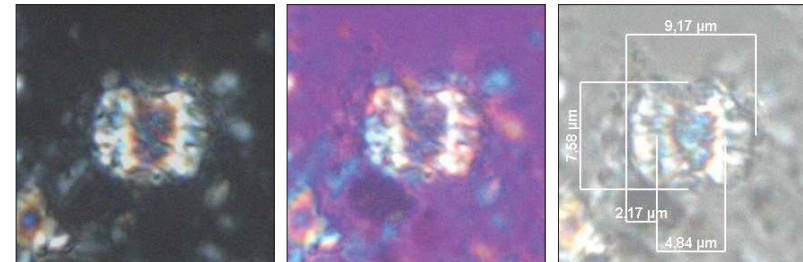
Nannoconus t. truittii (DSDP Site 364 / Depth 1.032,37 mbsf)



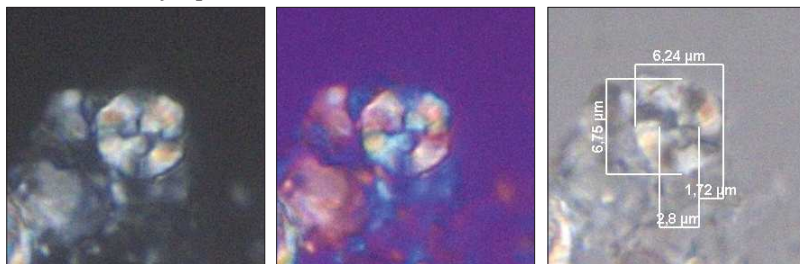
Nannoconus fragilis (DSDP Site 364 / Depth 973,89 mbsf)



Nannoconus t. frequens (DSDP Site 364 / Depth 1.032,37 mbsf)



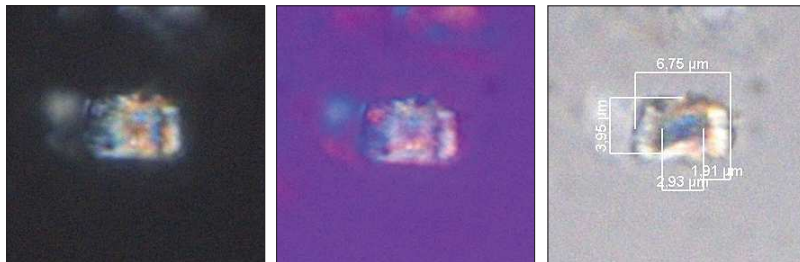
Nannoconus t. rectangularis (DSDP Site 364 / Depth 790,41 mbsf)



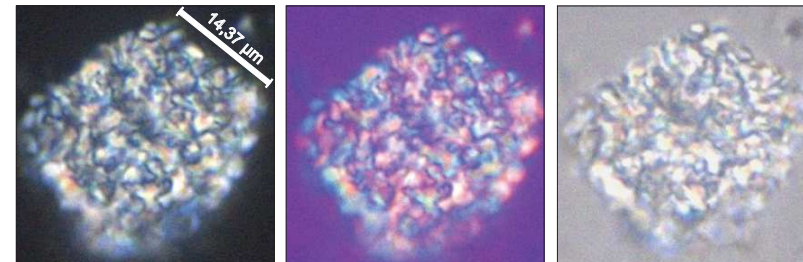
Nannoconus troelsenii (DSDP Site 364 / Depth 826,40 mbsf)



Nannoconus planus (DSDP Site 364 / Depth 973,89 mbsf)



Nannoconus troelsenii (DSDP Site 364 / Depth 968,45 mbsf)



Thoracosphaera sp.1 (DSDP Site 364 / Depth 1.032,37 mbsf)

10 µm