

APTIAN TO CAMPANIAN CALCAREOUS NANNOFOSSIL BIOSTRATIGRAPHY FROM THE BOTTACCIONE SECTION, GUBBIO, CENTRAL ITALY.

FABRIZIO TREMOLADA

Received December 27, 2001; accepted May 3, 2002

Key Words. Calcareous Nannofossils, Cretaceous, Biostratigraphy, Umbria-Marche Basin.

Riassunto. L'analisi biostratigrafica condotta sui Nannofossili calcarei delle Formazioni degli Scisti a Fucoidi e della Scaglia ha permesso di migliorare la risoluzione stratigrafica della sezione del Bottaccione (Bacino Umbro-Marchigiano, Italia Centrale) nell'intervallo Aptiano superiore-Campaniano. Inoltre, caratteristici bioeventi addizionali quali l'ultima comparsa di *Assipetra infracretacea larsonii* e *Rucinolithus terebrodentarius youngii*, la prima comparsa di *Parhabdolithus achlyostaurion* e di esemplari sovradimensionati di *Biscutum constans* potrebbero essere utilizzati per migliorare la stratigrafia a Nannofossili calcarei alle basse latitudini nell'intervallo Aptiano superiore-Albiano.

Abstract. Late Aptian-Campanian biostratigraphic investigations performed on calcareous nannofossils provided a refined stratigraphic resolution of the Scisti a Fucoidi and Scaglia Formations in the Bottaccione section (Umbria-Marche Basin, Central Italy). Additional bioevents such as the Last Occurrence (LO) of both *Assipetra infracretacea larsonii* and *Rucinolithus terebrodentarius youngii* and the First Occurrences (FOs) of *Parhabdolithus achlyostaurion* and large *Biscutum constans* were detected and might be used to refine the Late Aptian-Albian biostratigraphy at low latitudes.

Introduction

The sediments exposed in the Bottaccione Gorge have been investigated by several authors owing to the continuous exposure from Upper Jurassic to lower Upper Eocene and the only modest tectonic disturbances. Many studies on lithostratigraphy, biostratigraphy and magnetostratigraphy of the Bottaccione succession were performed in the last century. Renz (1936) proposed an Aptian-Eocene biostratigraphy. The planktonic foraminiferal biostratigraphy was carried out by Luterbacher & Premoli Silva (1962), Premoli Silva (1977) and Premoli Silva & Sliter (1994). Mohler (1966) tried to combine the nannofossil and planktonic foraminifer biostratigraphy. Arthur & Fischer (1977) and Arthur (1979) surveyed and measured the section.

Alvarez et al. (1977), Lowrie et al. (1982) and Napoleone et al. (1983) provided detailed paleomagnetic analyses. In addition, chemostratigraphic studies based on oxygen, carbon and strontium were performed by Renard (1986), Corfield et al. (1991), Jenkyns et al. (1994) and Stoll & Schrag (2000).

Monechi & Thierstein (1985) established a detailed Early Campanian to Late Eocene nannofossil stratigraphy, recently refined by Gardin et al. (2001), calibrated with the magnetostratigraphy of the Bottaccione section. The nannofossil biostratigraphy previously proposed for the Aptian to Santonian interval by Monechi (1981), Erba (1988) and Ghisletti & Erba in Premoli Silva & Sliter (1994) is here refined based on updated taxonomy and bio-chronostratigraphy.

This study documents a high-resolution biostratigraphy using calcareous nannofossils from the Upper Aptian to the lowermost Campanian of the Bottaccione Gorge. The nannofossil events are directly calibrated to the foraminiferal biostratigraphy reported by Premoli Silva & Sliter (1994), since both studies have been conducted on the same samples.

Geological setting and lithology

The Bottaccione section is located close to the town of Gubbio along the road-cut from Gubbio to Scheggia in the Umbria-Marche Basin, Central Italy (Fig.1). Here a fairly continuous pelagic (lower to upper bathyal) sedimentation characterizes the Jurassic to Paleogene interval. Very condensed or incomplete sequences suggest the occurrence of ancient structural highs.

Above basement and Triassic evaporites, the Jurassic and Cretaceous sedimentary succession is mainly composed by calcareous to marly calcareous lithotypes containing nodules and layers of chert.

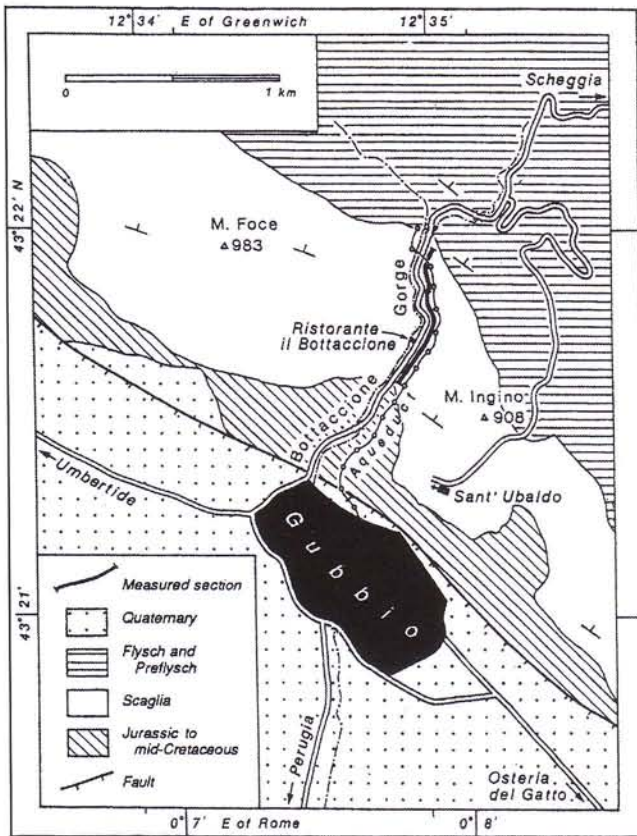


Fig. 1 - Location map of the Bottaccione section (after Premoli Silva & Sliter 1994).

In particular, the succession outcropping at Bottaccione and studied in this paper is referred to Scisti a Fucoidi and Scaglia Formations.

The Scisti a Fucoidi Formation of Early Aptian-Late Albian age consists of alternating marlstones, marly limestones, limestones and organic-rich black shales. The contact between the Scisti a Fucoidi and the overlying Scaglia Formation is transitional; the marly intervals decrease, whereas the limestones become dominant.

The Scaglia Formation (Late Albian-Oligocene) is composed of limestones and marly limestones with lenses or bands of chert. It is possible to subdivide this formation into four members on the basis of lithological characteristics and color (Cresta et al. 1989): Scaglia Bianca (Late Albian-Early Turonian), Scaglia Rossa (Early Turonian-Middle Eocene), Scaglia Variegata (Middle Eocene-Late Eocene) and Scaglia Cinerea (Late Eocene-Oligocene). The most prominent feature of the Scaglia Bianca is the presence of organic-rich black shales and siliceous sediments (Bonarelli Horizon, Oceanic Anoxic Event 2) in the uppermost Cenomanian.

Material and Methods

Biostratigraphic investigations were performed on samples collected from the Scisti a Fucoidi, Scaglia Bian-

ca and Scaglia Rossa. A total of 354 samples of limestones, marly limestones and marlstones were prepared using standard techniques (Monechi & Thierstein 1985); no ultrasonic cleaning or centrifuge concentration were applied in order to retain the original biogenic composition of the sample. Smear slides, prepared with Norland Optical Adhesive, were analyzed using standard light microscope techniques under cross polarizers and transmitted light at 1250X magnification.

Due to calcite dissolution and/or overgrowth, preservation and abundance of calcareous nannofossil species may vary significantly. Thus, to characterize preservation a simple code system (Ocean Drilling Program code) has been adopted and is listed below:

VG (very good) = no evidence of dissolution and/or overgrowth is present; there is no alteration of primary morphological characteristics, and specimens appear diaphanous; specimens are identifiable to the species level.

G (good) = little or no evidence of dissolution and/or overgrowth is present; primary morphological characteristics are only slightly altered; specimens are identifiable to the species level.

M (moderate) = specimens exhibit some etching and/or overgrowth; primary morphological characteristics are sometimes altered; however, most specimens are identifiable to the species level.

P (poor) = most specimens exhibit overgrowth or dissolution; primary morphological characteristics are sometimes destroyed; fragmentation has occurred; species identification is often impaired.

Estimates of the total calcareous nannofossil abundance, compared to that of other biogenic particles and inorganic components, were recorded as follows:

C (common) = >51% of all particles,

F (few) = 11%-50% of all particles,

R (rare) = 1%-10% of all particles,

T (trace) = <1% of all particles, and

B (barren) = no nannofossils are present.

Estimates of the relative abundance of calcareous nannofossil species in the studied assemblages, were determined as follows:

D (dominant) = >51% of the total assemblage,

A (abundant) = 11%-50% of the total assemblage,

C (common) = 2%-10% of the total assemblage,

F (few) = 0.1%-1% of the total assemblage, and

R (rare) = <0.1% of the total assemblage.

Biostratigraphy and discussion

The nannofossil biozonation adopted in this study (Figs 2,3) is that of Roth (1978) and is regarded as standard as summarized in Perch-Nielsen (1985). The NC biozonation provided by Roth (1978) and the improvements proposed by Bralower et al. (1995) are largely

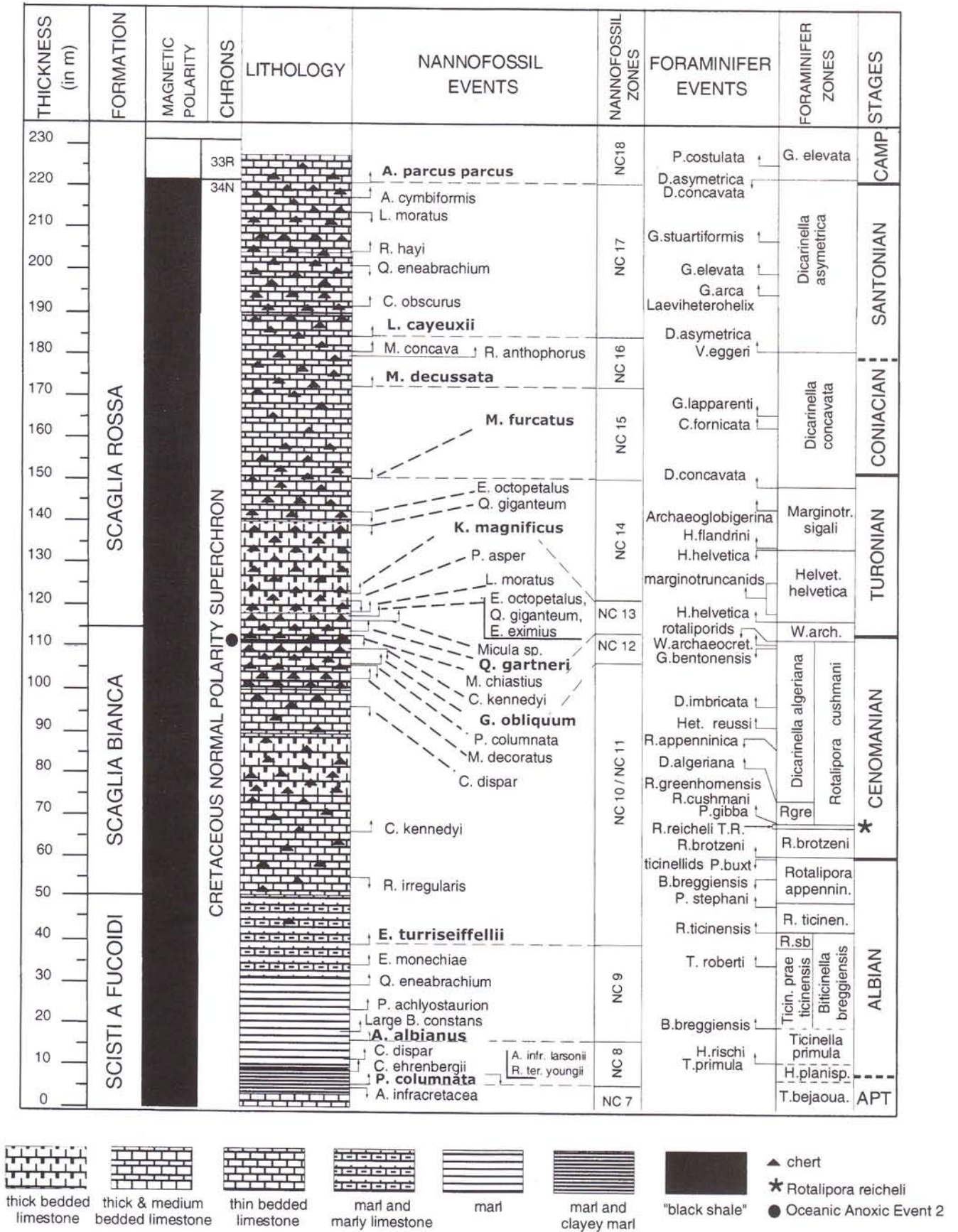


Fig. 2 - Lithostratigraphic log, foraminiferal (after Premoli Silva & Sliter 1994) and nannofossil events recognized at Bottaccione.

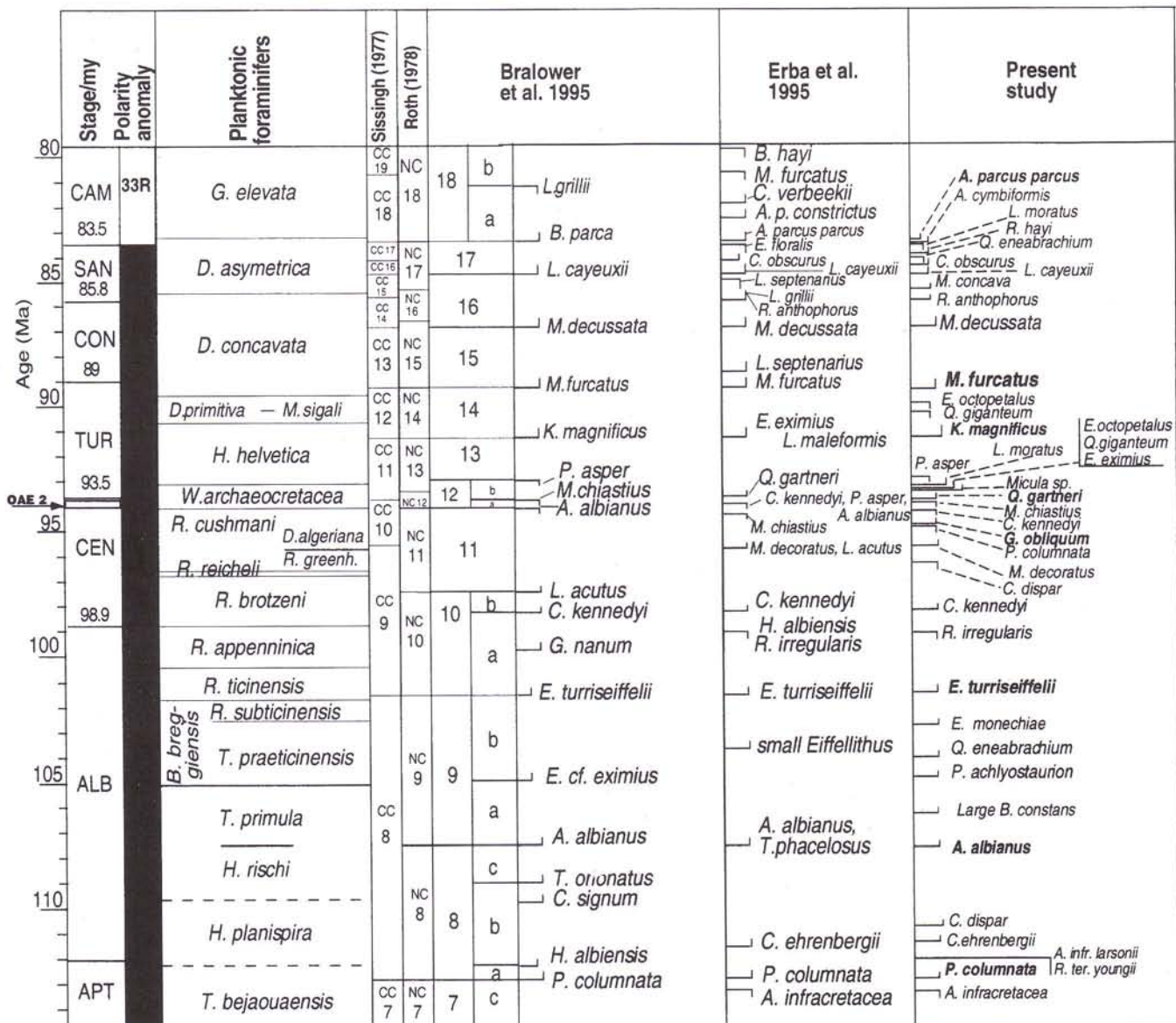


Fig. 3 - Comparison amongst the bioevents detected at Bottaccione and previous biozonations. Planktonic foraminifers from Premoli Silva & Sliter (1999). The gray pattern indicates the time interval investigated.

based on DSDP/ODP results and allow a good correlation between nannofossil and planktonic foraminifer biostratigraphic data.

The Sissingh (1977) scheme was constructed on results from land sections (mostly in Europe), but his planktonic foraminiferal data are often unavailable or different from those observed by Premoli Silva & Sliter (1994) at Bottaccione.

The UC biozonation, recently published by Burnett (1998), does not allow a good calibration with planktonic foraminiferal biozonations and most nannofossil events are recognizable only at higher latitudes or in better preserved material.

Most nannofossil assemblages found are dominated by *Watznaueria barnesae*, a species very resistant to diagenetic modifications (Roth & Krumbach 1986). Indeed, large amounts of *W. barnesae* usually indicate a

low degree of nannofossil preservation.

In general, the Scaglia displays a nannofossil preservation ranging from poor to moderate and the overall nannofossil abundance is low. By contrast, calcareous nannofossils in the Scisti a Fucoidi are just slightly altered, thus allowing the identification at species level and their abundance ranges from few to common.

The bottom of the investigated succession is assignable to the Late Aptian NC7 nannofossil zone based on the occurrence of *Eprolithus floralis* in the lowermost sample investigated (sample Apt 4 - 0.60 m). The NC7/NC8 zonal boundary is placed at the FO of *Pre-discosphaera columnata* (sample Apt 19 - 5.86 m). This zonal boundary is preceded by the LO of *Assipetra infracretacea* (sample Apt 17 - 5.48 m), whereas *A. infr. larsonii* disappears at 7.33 m (sample Apt 22) associated

with the LO of *Rucinolithus terebrodentarius youngii*. *A. infr. larsonii* and *R. ter. youngii* represent the large-sized morphotypes of *A. infracretacea* and *R. terebrodentarius*, respectively and their stratigraphic range is restricted to the Aptian stage (Tremolada & Erba 2002).

The FO of *Cribrosphaerella ehrenbergii* lies at 8.60 m (sample Apt 28) indicating an earliest Albian age. *Calculithes dispar* was first observed in the sample Apt 42 bis (14.75 m) and displays a continuous occurrence although characterized by low abundance. This species was established by Al-Rifa'iy et al. (1990) and there are still limited data concerning its distribution. The LO of *C. dispar* was detected in the Bottaccione section at 97.09 m (sample B 458), of Late Cenomanian age, suggesting a slightly later disappearance than reported by Al-Rifa'iy et al. (1990).

Axopodorhabdus albianus is extremely rare in the samples investigated and its first occurrence is located at 15.74 m (sample Apt 47). The FO of *A. albianus* marks the NC8/NC9 zonal boundary, within the *Ticinella primula* foraminiferal Zone (Premoli Silva & Sliter 1994, 1999). The top of the NC9 nannofossil zone is defined by the first occurrence of *Eiffellithus turriseiffelii* at 37.80 m (sample Apt 89) preceding the FO of *Rotalipora ticiensis* at 43.10 (Premoli Silva & Sliter 1994). The FO of *E. turriseiffelii* is anticipated by the occurrence of *Eiffellithus monechiae* (Bown et al. 1998) at 35.55 m (sample Apt 84). This taxon may represent the *Eiffellithus* cf. *E. eximius* documented by Bralower et al. (1995) or the small *Eiffellithus* reported in Erba et al. (1995).

Within Zone NC9 other events such as the FOs of large *Biscutum constans* at 16.68 m (sample Apt 48) and *Parhabdolitus achlyostaurion* at 23.61 m (sample Apt

64) were recognized. The FO of large-sized specimens of *B. constans* is a peculiar event of the Tethyan Albian documented as *B. magnum* sensu Erba et al. (1999) at the Cismon drillcore (Northeastern Italy, Venetian alps), Erba (1988) at the Piobbico drillcore (Umbria-Marche Basin, Central Italy) and Bellanca et al. (in press) at the Calabianca section (Northwestern Sicily, Southern Italy).

Several authors (e.g. Bralower et al. 1995; Bown et al. 1998) report the FO of *P. achlyostaurion* in the Late Aptian, but at the Bottaccione section, the Cismon drillcore (Erba et al. 1999), the Piobbico drillcore (Erba 1988) and the Calabianca section (Bellanca et al. in press) this event is dated as Middle Albian. This discrepancy is probably due to different criteria in the taxonomic identification or a biogeographic control hampering a widespread and simultaneous first occurrence of *P. achlyostaurion*.

The LO of *Rucinolithus irregularis* lies at 54.65 m (sample B 58) in the Late Albian within the *Rotalipora appenninica* foraminiferal Zone (Premoli Silva & Sliter 1994) and the FO of *Corollithion kennedyi* at 68.74 m (sample B 193), within the *Rotalipora reicheli* Zone (Premoli Silva & Sliter 1994) indicating an Early Cenomanian age.

The FO of the foraminifer *Rotalipora brotzeni* was recognized at 61.20 m (Premoli Silva & Sliter 1994).

C. kennedyi displays a quite continuous presence through the Cenomanian up to 110.78 m (sample B 561), just prior to the deposition of the Bonarelli Horizon. In several sections such as Novara di Sicilia, Northeastern Sicily (Tremolada unp. data) and Eastbourne, United Kingdom (Paul et al. 1999) the LO of *C.*

PLATE 1

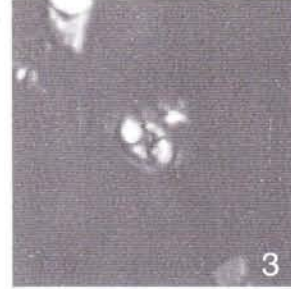
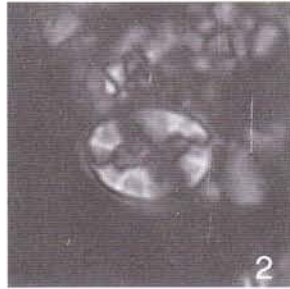
All specimens at x2000 magnification. Fig. 1 - *Eiffellithus eximius*, crossed nicols, sample B 580. Fig. 2 - *Eiffellithus eximius*, crossed nicols, sample 16/164. Figs 3,4 - *Calculithes dispar*, crossed nicols, sample B60, same specimen. Figs 5,6 - *Micula* sp., crossed nicols, sample 16/164, same specimen. Figs 7,8 - *Corollithion kennedyi*, crossed nicols, sample B 558, same specimen. Fig. 9 - *Micula* cf. *decussata*, crossed nicols, sample 16/112. Fig. 10 - *Micula decussata*, crossed nicols, sample 16/114. Fig. 11 - *Rucinolithus irregularis*, crossed nicols, sample B 49. Fig. 12 - *Quadrum eneabrachium*, crossed nicols, sample Apt 73. Fig. 13 - *Rucinolithus hayi*, crossed nicols, sample 16/104R. Fig. 14 - *Biscutum constans*, crossed nicols, sample B 273. Fig. 15 - Large *B. constans*, crossed nicols, sample B 104. Fig. 16 - Large *B. constans*, transmitted light, sample B 123. Fig. 17 - *Aspidolithus parvus parvus*, crossed nicols, sample 16/100R. Fig. 18 - *Calculithes obscurus*, crossed nicols, sample 16/102. Fig. 19 - *Kamptnerius magnificus*, crossed nicols, sample 16/175. Fig. 20 - *Kamptnerius magnificus*, crossed nicols, sample 16/175.

PLATE 2

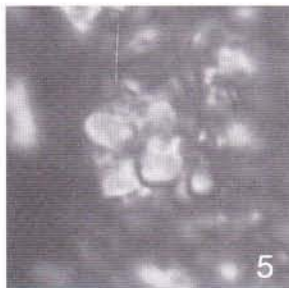
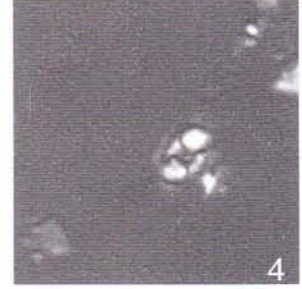
All specimens at x2000 magnification. Figs 1,2 - *Microstaurus chiastius*, crossed nicols, sample B554, same specimen. Fig. 3 - *Microrhabdulus decoratus*, crossed nicols, sample B 535. Fig. 4 - *Prediscosphaera columnata*, crossed nicols, sample Apt 20. Fig. 5 - *Marthasterites furcatus*, transmitted light, sample 16/148. Fig. 6 - *Marthasterites furcatus*, transmitted light, sample 16/145. Fig. 7 - *Marthasterites furcatus*, transmitted light, sample 16/145. Fig. 8 - *Eiffellithus turriseiffelii*, crossed nicols, sample 16/102R. Fig. 9 - *Parhabdolitus achlyostaurion*, crossed nicols, sample B 53. Fig. 10 - *Parhabdolitus achlyostaurion*, crossed nicols, sample B 58. Fig. 11 - *Parhabdolitus achlyostaurion*, crossed nicols, sample B 19. Fig. 12 - *Parhabdolitus achlyostaurion*, crossed nicols, sample B 24. Fig. 13 - *Quadrum gartneri*, crossed nicols, sample B 570. Fig. 14 - *Reinhardtites* cf. *anthophorus*, crossed nicols, sample 16/101. Fig. 15 - *Cribrosphaerella ehrenbergii*, crossed nicols, sample B 471. Fig. 16 - *Assipetra infracretacea larsonii*, crossed nicols, sample Apt 22. Fig. 17 - *Gartnerago obliquum*, crossed nicols, sample B 546. Fig. 18 - *Eiffellithus trabeculatus*, crossed nicols, sample B 471. Fig. 19 - *Rucinolithus terebrodentarius youngii*, crossed nicols, sample Apt 11. Fig. 20 - *Rucinolithus terebrodentarius youngii*, crossed nicols, sample Apt 12.



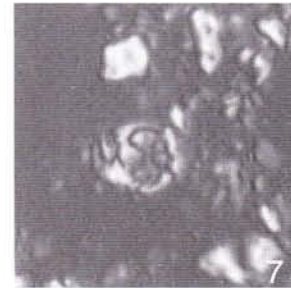
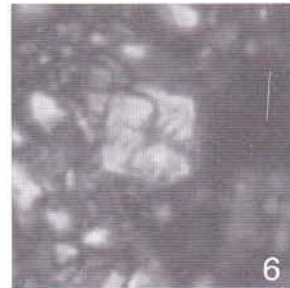
E. eximius



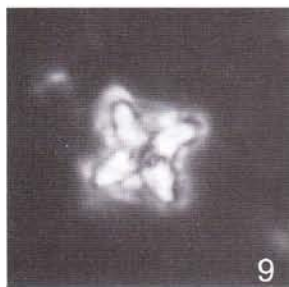
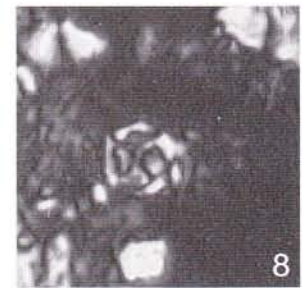
C. dispar



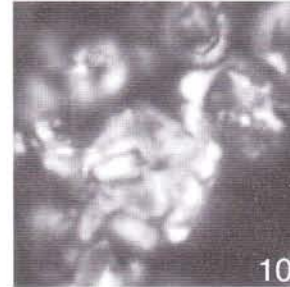
Micula sp.



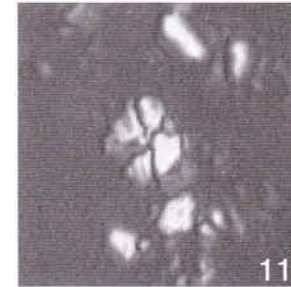
C. kennedyi



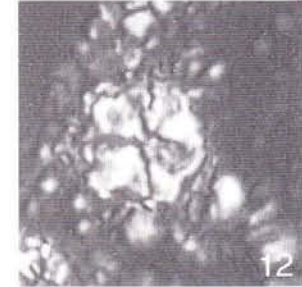
M. cf. decussata



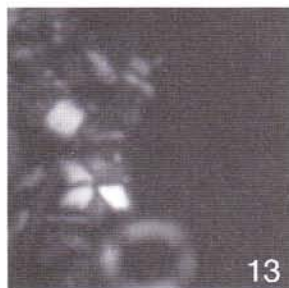
M. decussata



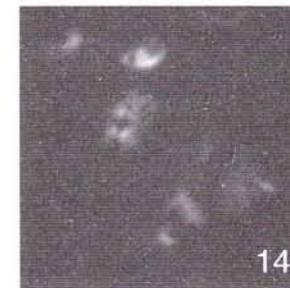
R. irregularis



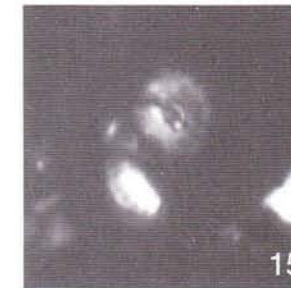
Q. eneabrachium



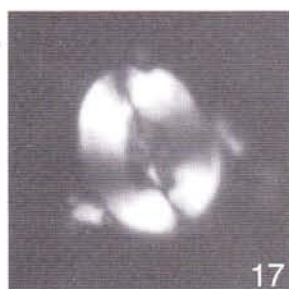
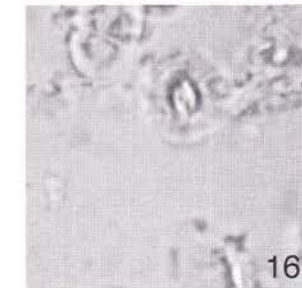
R. hayi



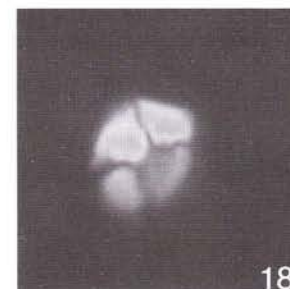
B. constans



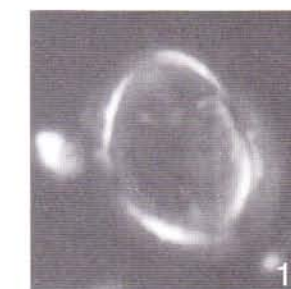
large *B. constans*



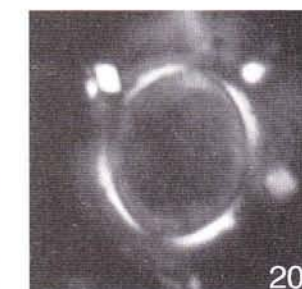
A. parvus parvus

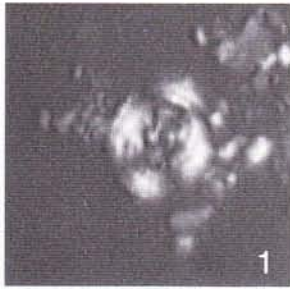


C. obscurus

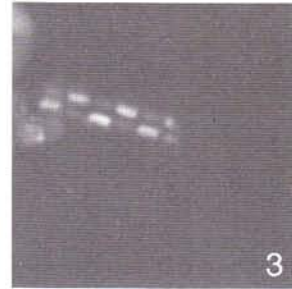
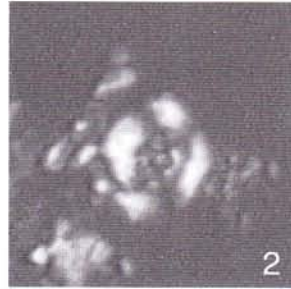


K. magnificus





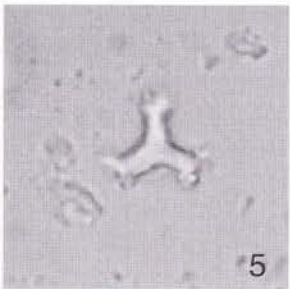
M. chiastius



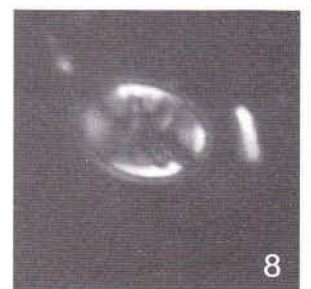
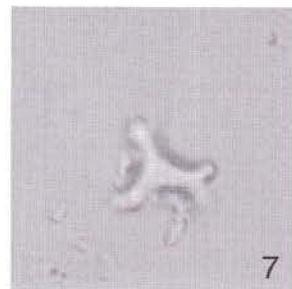
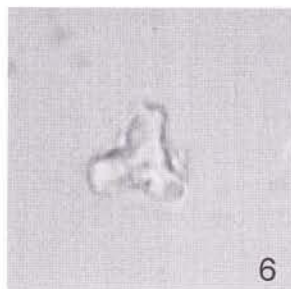
M. decoratus



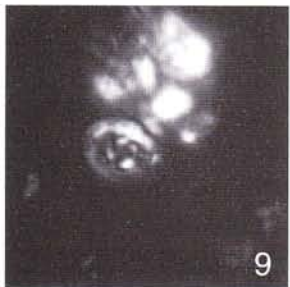
P. columnata



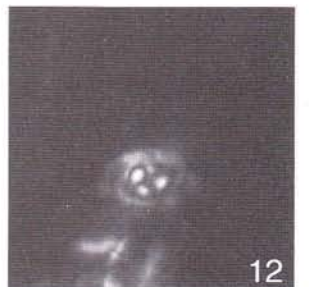
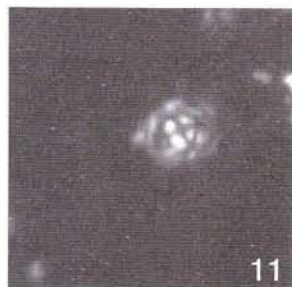
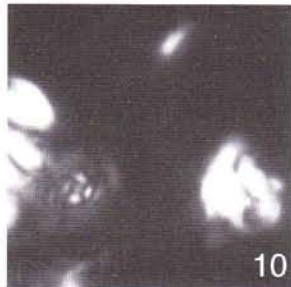
M. furcatus



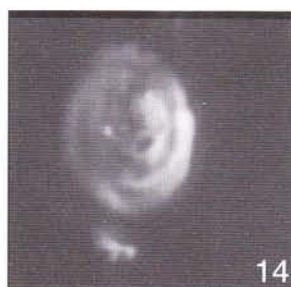
E. turriseiffelii



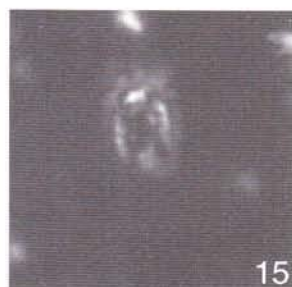
P. achlyostaurion



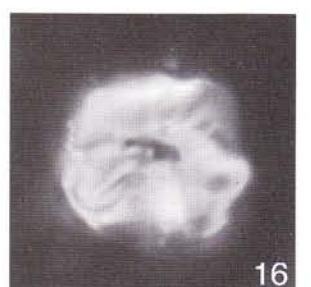
Q. gartneri



R. cf. anthophorus



C. ehrenbergii



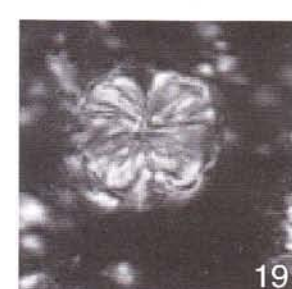
A. infr. larsonii



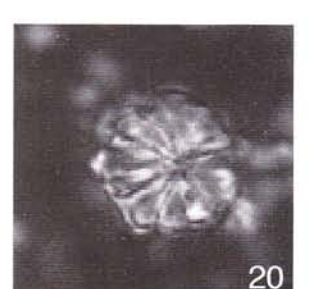
G. obliquum



E. trabeculatus



R. ter. youngii



kennedyi was recognized within the Bonarelli Horizon equivalents. This apparent discrepancy is possibly due to the lack of calcium carbonate in the uppermost Cenomanian black shales at Bottaccione, but Luciani & Cobianchi (1999) detected the LO of *C. kennedyi* just above the Bonarelli Horizon at Antruiles, Northeastern Italy.

The interval immediately preceding the Bonarelli Horizon is marked by several nannofossil events such as the FO of *Microrhabdulus decoratus* at 107.92 m (sample B 533), the LO of *P. columnata* at 108.77 m (sample B 542) and the FO of *Gartnerago obliquum* at 109.35 m (sample B 546), which defines the base of the nannofossil zone NC12. It was not possible to subdivide the zones NC10 and NC11, since the zonal marker *Lithraphidites acutus* shows a too spotty distribution.

The extinction of the foraminiferal genus *Rotalipora* lies at 111 m (Premoli Silva & Sliter 1994) slightly below the Bonarelli Horizon; this event represents the base of the *Whiteinella archaeocretacea* Zone. The top of this zone is placed at the FO of *Helvetoglobotruncana helvetica* (sample 16/177 - 117 m).

The LO of *Microstaurus chiastius* was observed at 112.03 m (sample B 564), whereas the FO of *Quadrum gartneri* (= *Tetralithus pyramidus* of Sissingh 1977 and Roth 1978) lies at 112.59 (sample B 568) and defines the base of the zone NC13. These events are recorded worldwide across the Cenomanian/Turonian boundary. The FO of *Micula* sp., recognized at 112.73 m (sample B 569), precedes the LO of *Parhabdolutus asper* at 114.68 m (sample B 579). *Micula* sp. may be correlatable to *Micula staurophora* as documented by Roth (1978) from the lowermost Turonian.

Kamptnerius magnificus was first observed at 119 m (sample 16/175), although extremely rare, and is used to identify the top of Zone NC13. Within Zone NC13 several bioevents were recognized: the FOs of *Eiffelolithus eximius*, *Quadrum giganteum* and *Eprolithus octopetalus* at 112.94 m (sample B 570), the FO of *Lithastrinus moratus* at 113.07 m (sample B 571) and the FO of *Lucianorhabdus* spp. at 118 m (sample 16/176).

Q. giganteum is usually rare in the Bottaccione section, whereas *E. eximius*, *L. moratus* and *E. octopetalus* display a continuous stratigraphic distribution, even though the poor preservation can sometimes hamper the identification of *E. octopetalus* and *L. moratus*.

The LOs of *Q. giganteum* (138 m - sample 16/160) and *E. octopetalus* (141 m - sample 16/157) were recognized within Zone NC14. The stratigraphic range of *Q. giganteum* and *E. octopetalus* is slightly different from that reported by Varol (1992) who recorded these events in the Early Turonian.

The FO of *Marthasterites furcatus* defines the base of Zone NC15 and this event was observed at 150 m (sample 16/148) postdating the FO of the foraminifer

Dicarinella concavata (sample 16/149 - 149 m). *M. furcatus* is always rare at Bottaccione.

The NC15/NC16 zonal boundary is placed at the FO of *Broinsonia lacunosa*, but this taxon was never detected at Bottaccione. In fact, *B. lacunosa* is a delicate coccolith susceptible to diagenetic modifications and its occurrence is conditioned by the degree of preservation of the investigated material. The FO of *Micula decussata* at 173 m (sample 16/125) was used to replace the FO of *B. lacunosa* and to indicate the base of the Zone NC16 (Bralower et al. 1995).

Reinhardtites anthophorus was first observed at 175 m (sample 16/123). This event precedes the FOs of *Micula concava* at 179 m (sample 16/119), *Lucianorhabdus cayeuxii* at 184 m (sample 16/114) and *Calculithes obscurus* at 191 m (sample 16/107). *M. concava*, *L. cayeuxii* and *C. obscurus* are usually rare and they display a spotty occurrence in the Bottaccione section, whereas *R. anthophorus* shows a rather continuous distribution. *L. cayeuxii* is a cosmopolitan taxon and recorded in low abundances everywhere, but it seems to prefer shallow water environments (Thierstein 1976). In this work I used the FO of *L. cayeuxii* to define the base of Zone NC17 following the modification proposed by Bralower et al. (1995).

The FOs of foraminifers *Dicarinella asymmetrica* and *Ventilabrella egeri* were detected in the same sample at 181 m (Premoli Silva & Sliter 1994).

Within Zone NC17 several bioevents were recognized: the LOs of *Q. eneabrachium* (196 m - sample 16/102) and *L. moratus* (216 m - sample 16/108R) and the FOs of *Rucinolithus hayii* (205 m - sample 149GA) and *Arkhangelskiella cymbiformis* (217 m - sample 16/106R).

The FO of *Aspidolithus parvus parvus* lies at 220 m (sample 16/104R) and defines the NC17/NC18 zonal boundary and the Santonian/Campanian stage boundary. The FO of *A. parvus parvus* is a well established event recognizable at both low- and high-latitudes (Bralower et al. 1995) and correlatable to the end of the normal superchron C34N.

The LO of *D. asymmetrica* is recorded at 221.5 m in the Early Campanian (Premoli Silva & Sliter 1994).

Chronostratigraphy

The integrated stratigraphy proposed in Fig. 2 summarizes the foraminiferal and nannofloral biozonations at Bottaccione. Nannofossil and foraminifer events detected in the samples investigated may be used to approximate the stage boundaries at low latitudes when detailed ammonite investigations are not available. The geomagnetic polarity time scales cannot be integrated with microfossil events over the time interval investigated since the Long Normal Superchron spans

from the Early Aptian to the Campanian (e.g. Alvarez et al. 1977).

The Aptian/Albian stage boundary is not easily recognizable and there are still uncertainties on the ammonite datum to be used (e.g. Br  h  ret et al. 1986; Hart et al. 1996). Kennedy et al. (2000) proposed to use the FO of the ammonite *Leymeriella tardefurcata* to indicate this boundary. Bown (in Kennedy et al. 2000) suggested the FO of the small, circular specimens of *P. columnata* or the LO of *Broinsonia viriosa* to approximate the ammonite-defined Aptian/Albian boundary. Bown (in Kennedy et al. 2000) studied the morphological characteristics of *P. columnata* differentiating three evolutionary stages: elliptical shape (*Prediscosphaera spinosa* of many authors), near-circular shape and circular shape.

However, *B. viriosa* was not found at Bottaccione and it was not possible to differentiate between near-circular and circular morphotypes of *P. columnata*. In fact, the specimens of *P. columnata* recorded in the samples analyzed are affected by overgrowth or etching hampering the study of size/shape variations. The Aptian/Albian stage boundary is usually placed in the lower part of the *Hedbergella planispira* foraminiferal Zone (Premoli Silva & Sliter 1994, 1999; Bralower et al. 1995), between the FOs of *P. columnata* and *C. ebrenbergii*.

A. infr. larsonii and *R. ter. youngii* have their last occurrence within the lower part of the foraminiferal Zone *Hedbergella planispira* and they may be used to approximate the boundary, since these subspecies were only recently established (Tremolada & Erba 2002) and perhaps they need a further documentation of their stratigraphic range, ecologic significance and biogeographic occurrence.

The FO of the planktonic foraminifer *Rotalipora globotruncanoides* (= *R. brotzeni*) best approximates the Albian/Cenomanian boundary (Bralower et al. 1995; Gale et al. 1996; Tr  ger & Kennedy 1996; Premoli Silva & Sliter 1999). The FO of *R. globotruncanoides* lies at 61 m between the LO of *R. irregularis* at 54.65 m and the FO of *C. kennedyi* at 68.74 m in the Bottaccione section.

The Cenomanian/Turonian boundary is defined by the FO of the ammonite *Watinoceras devonense* (Wright & Kennedy 1991; Bengtson 1996). Several studies performed on planktonic foraminifers (Eicher & Diner 1985; Premoli Silva & Sliter 1994, 1999; Bralower et al. 1995; Bengtson 1996) pointed out that the Cenomanian/Turonian boundary lies between the LO of the genus *Rotalipora* and the FO of *Helvetoglobotruncana helvetica*. The FO of the nannofossil *Q. gartneri* is recorded everywhere in the interval between the extinction of the rotaliporids and the FO of *H. helvetica* (e.g. Perch-Nielsen 1983; Watkins 1986; Watkins et al. 1996; Bengtson et al. 1996). Indeed, the FO of *Q. gartneri* may be used to approximate the Cenomanian/Turonian

boundary since it slightly precedes the FO of *W. devonense* at low latitudes, whereas Burnett (1998) reported the first occurrence of *Q. gartneri* from the Lower Turonian (*Mammites nodosoides* ammonite Zone) at high latitudes.

The Turonian/Coniacian boundary can be approximated using the FO of *M. furcatus* (Sissingh 1977; Perch-Nielsen 1983, 1985; Bralower et al. 1995; Watkins et al. 1996), whereas it is not possible to define the Coniacian/Santonian boundary since there are not bioevents directly correlatable to this boundary. The entry of the foraminiferal group of the ventilabrellids may be used to best approximate the Coniacian/Santonian boundary as reported by Salaj (1980, 1984) in Tunisian sections, but this event was detected at the same level as the FO of *Dicarinella asymmetrica* at Bottaccione, thus indicating an Early Santonian age (Premoli & Sliter, 1994).

The Santonian/Campanian boundary is equated to the magnetic chron 34N/33R boundary (e.g. Lommerzhelm & Hambach 1992; Bralower et al. 1995; Gradstein et al. 1995) and correlates with the FO of *A. parvus parvus*.

Conclusions

Several important nannofossil events were recognized in the Bottaccione section, even though the nannofloral preservation is poor. A few zonal markers (e.g. *L. acutus*, *B. viriosa* and *B. lacunosa*) are missing or show a spotty occurrence in the samples analyzed.

The nannofossil and foraminifer events found at Bottaccione are correlated in Fig. 2; this scheme may represent a good integrated stratigraphy for Italian sections and the Tethys area in general. Additional nannofossil events proposed herein such as the FOs of large *B. constans*, *P. achlyostaurion* and the LOs of *R. terebrodentarius youngii* and *A. infracretacea larsonii* may be used also in other regions in order to enhance the resolution of the currently available biozonations.

Acknowledgements. I am grateful to Elisabetta Erba and Isabella Premoli Silva for stimulating discussions regarding microfossils and biostratigraphy. I would like to thank Prof. K. Von Salis, Prof. S. Monchi and Prof. M. Gaetani for reviewing the manuscript and for their helpful comments. This work was supported by grants MURST n   9904152971_001 and MIUR-SAUS n. 2001048975_001 to I. Premoli Silva.

REFERENCES

- Al Rifay I., Varol O., Lemone D. (1990) - Middle to late Albian biostratigraphy of the Cuchillo Formation from Sierra de Sapellò, Mexico. *Newsletters on Stratigraphy*, 21: 187-200, Berlin-Stuttgart.
- Alvarez W., Arthur M.A., Fischer A.G., Lowrie W., Napoleone G., Premoli Silva I., Roggentheim W.M. (1977) - Upper Cretaceous-Paleocene magnetic stratigraphy at Gubbio, Italy. Type section for the Late Cretaceous-Paleocene geomagnetic reversal time scale. *Geol. Soc. Am. Bull.*, 88: 383-389, Boulder.
- Arthur M.A. (1979) - Sedimentologic and geochemical studies of Cretaceous and Paleogene pelagic sedimentary rocks: the Gubbio sequence, Pt. 1, PhD Dissertation, Univ. of Princeton, 1-102.
- Arthur M.A., Fischer A.G. (1977) - Upper Cretaceous-Paleocene magnetic stratigraphy at Gubbio, Italy. Lithostratigraphy and sedimentology. *Geol. Soc. Am. Bull.*, 88: 367-371, Boulder.
- Bellanca A., Erba E., Neri R., Premoli Silva I., Sprovieri M., Tremolada F., Verga D. (in press) - Biostratigraphic and geochemical events in organic-rich pelagic sediments through the Early Aptian "Livello Selli", Northwestern Sicily. *Palaeogeogr. Palaeoclim. Palaeocol.*, Amsterdam.
- Bengtson P. (1996) - The Turonian stage and substage boundaries. *Bulletin Institut Royal des Sciences Naturelles de Belgique*, 6: 69-81, Bruxelles.
- Bown P.R., Rutledge D.C., Crux J.A., Gallagher L.T. (1998) - Lower Cretaceous. In P. R. Bown (ed.), *Calcareous Nannofossil Biostratigraphy*, 86-131, Kluwer Academic Publishers, London.
- Bralower T.J. (1988) - Calcareous nannofossil biostratigraphy and assemblages of the Cenomanian-Turonian boundary interval: implications for the origin of oceanic anoxia. *Paleoceanography*, 3: 275-316, Washington D.C.
- Bralower T.J., Leckie R.M., Sliter W.V., Thierstein H.R. (1995) - An integrated Cretaceous microfossil biostratigraphy. In W.A. Berggren, Kent D.V., Aubry M.P., Hardenbol J. (eds.), *Geochronology, time scales and global stratigraphic correlation*. SEPM Special Publication, 54: 65-79, Tulsa.
- Bralower T.J., Bergen J.A. (1998) - Cenomanian-Santonian calcareous nannofossil biostratigraphy of a transect of cores drilled across the Western Interior Seaway. In *Stratigraphy and Paleoenvironments of the Cretaceous Western Interior Seaway*. SEPM Concepts in Sedimentology and Paleontology, 6: 59-77, Tulsa.
- Bréhéret J.G., Caron M., Delamette M. (1986) - Niveaux riches en Matière organique dans l'Albien Vocontienne; Quelques Caractères du Paléoenvironnement; Essai d'interprétation génétique. *Documents Bureau Recherches Géologiques et Minières*, 110: 141-191, Lyon.
- Burnett J.A. (1998) - Upper Cretaceous. In P.R. Bown (ed.), *Calcareous Nannofossil Biostratigraphy*, 132-199, Kluwer Academic Publishers, London.
- Corfield R.M., Cartlidge J.E., Premoli Silva I., Housley R.A. (1991) - Oxygen and carbon isotope stratigraphy of the Paleogene and Cretaceous limestones in the Bottaccione and Contessa Highway sections, Umbria, Italy. *Terra Nova*, 3: 414-422, Oxford.
- Cresta S., Monechi S., Parisi G. (1989) - Stratigrafia del Mesozoico e del Cenozoico nell'area Umbro-Marchigiana (Mesozoic-Cenozoic stratigraphy in the Umbria-Marche area). *Memorie Descrittive della Carta Geologica d'Italia*, 39: 1-185, Roma.
- Eicher D.L., Diner R. (1985) - Foraminifera as indicators of water mass in the Cretaceous Greenhorn sea, Western Interior. In Pratt L.M., Kauffman, E.G., Zelt F. (eds) *Fine-grained deposits and biofacies of the Cretaceous Western Interior: evidence of cyclic sedimentary processes*. *Society of Economic Paleontologists and Mineralogists, Midyear Field Trip Guidebook*, 4: 60-71, Tulsa.
- Erba E. (1988) - Aptian-Albian calcareous nannofossil biostratigraphy of the Scisti a Fucoidi cored at Piobbico (Central Italy). *Riv. It. Paleont. Strat.*, 97: 455-483, Milano.
- Erba E., Premoli Silva I., Wilson P.A., Pringle M.S., Sliter W.V., Watkins D.K., Arnaud Vanneau A., Bralower T.J., Budd A.F., Camoin G.F., Masse J., Mutterlose J. & Sager W.W. (1995) - Synthesis of stratigraphies from shallow-water sequences at Sites 871 through 879 in the western Pacific Ocean (Leg 144). *ODP Scientific Results Leg 144*, 144: 873-885, College Station TX.
- Erba E., Channell J.E.T., Claps M., Jones C., Larson R.L., Opdyke B., Premoli Silva I., Riva A., Salvini G., Torricelli S. (1999) - Integrated stratigraphy of the Cismon Apticore (Southern Alps, Italy): A "reference section" for the Barremian-Aptian interval at low latitudes. *J. Foraminiferal Res.*, 29: 371-392, Lawrence.
- Gale A.S., Kennedy W.J., Burnett J.A., Caron M., Kidd B.E. (1996) - The Late Albian to Early Cenomanian at Mont Risou, near Rosans (Drôme, SE France): an integrated study (ammonites, inoceramids, planktonic foraminifera, nannofossils, oxygen and carbon isotopes). *Cretaceous Research*, 17: 515-606, London.
- Gardin S., Del Panta F., Monechi S., Pozzi M. (2001) - A tethyan reference section for the Campanian and Maastrichtian stages: the Bottaccione section (Central Italy). Review of the data and new calcareous nannofossil results. In G.S. Odin (ed.), *The boundary between the Campanian and the Maastrichtian stages: characterisation and correlation from Tercis-les-bains to Europe and other continents*. *Developments in Paleontology and Stratigraphy*, 19: 820-833, Elsevier, Amsterdam.
- Gradstein F.M., Agterberg F.P., Ogg J.G., Hardenbol J., Van Veen P., Thierry J., Huang Z. (1995) - A Triassic, Jurassic and Cretaceous Time Scale. In Berggren W.A., Kent D.V., Aubry M.P., Hardenbol J. (eds.), *Geochronology, time scales and global stratigraphic correlation*. SEPM Special Publication, 54: 95-128, Tulsa.
- Hart M., Amedro F., Owen H.G. (1996) - The Albian stage and substage boundaries. *Bulletin Institut Royal des Sciences Naturelles de Belgique*, 6: 45-57, Bruxelles.
- Jenkyns H.C., Gale A.S., Corfield R.M. (1994) - Carbon- and oxygen-isotope stratigraphy of the English Chalk and Italian Scaglia and its palaeoclimatic significance. *Geol. Mag.*, 131: 1-34, Cambridge.

- Kennedy W.J., Gale A.S., Bown P.R., Caron M., Davey R.J., Grocke D., Wray D.S. (2000) - Integrated stratigraphy across the Aptian-Albian boundary in the Marnes Bleues, at the Col de Pre-Guittard, Arnayon (Drôme), and at tartonne (Alpes-de-Haute-Provence), France: a candidate Global Boundary Stratotype Section and Boundary Point for the base of the Albian Stage. *Cretaceous Research*, 21: 591-720, London.
- Lommerzheim A.J., Hambach U. (1992) - Cephalopod- and magnetostratigraphy of the Campanian of the Muensterland Basin, NW Germany. *Cretaceous Symposium Abstracts*, 47-50, Hamburg.
- Lowrie W., Alvarez W., Napoleone G., Perch-Nielsen K., Premoli Silva I., Toumarkine M. (1982) - Paleogene magnetic stratigraphy in Umbrian pelagic carbonate rocks: the Contessa sections. *Geol. Soc. Am. Bull.*, 93: 414-432, Boulder.
- Luciani V. & Cobianchi M. (1999) - The Bonarelli Level and other black shales in the Cenomanian-Turonian of the northeastern Dolomites (Italy): calcareous nannofossil and foraminiferal data. *Cretaceous Research*, 20: 135-167, London.
- Luterbacher H.P., Premoli Silva I. (1962) - Note preliminaire sur une revision du profil de Gubbio, Italie. *Riv. It. Paleont. Strat.*, 68: 253-288, Milano.
- Mohler H.P. (1966) - Stratigraphische Untersuchungen in den Giswiler Klippen (Prealpes Medianes) und ihrer helvetisch-ultrahelvetischen Unterlage. *Beitr. Geologische Karte Schweiz*, 129:15-84, Bern.
- Monechi S. (1981) - Aptian-Cenomanian calcareous nannoplankton from some sections in the Umbria-Marche Apennine. *Riv. It. Paleont. Strat.*, 87: 193-226, Milano.
- Monechi S., Thierstein H.R. (1985) - Late Cretaceous-Eocene nannofossil and magnetostratigraphic correlations near Gubbio, Italy. *Mar. Micropaleont.*, 9: 419-440, Amsterdam.
- Napoleone G., Premoli Silva I., Heller F., Cheli P., Corezzi S., Fischer A.G. (1983) - Eocene magnetic stratigraphy at Gubbio, Italy, and its implications for Paleogene geochronology. *Geol. Soc. Am. Bull.*, 94: 181-191, Boulder.
- Paul C.R.C., Lamolda M.A., Mitchell S.F., Vaziri M.R., Gorostidi A., Marshall J.D. (1999) - The Cenomanian-Turonian boundary at Eastbourne (Sussex, UK): A proposed European reference section. *Paleogeogr. Paleoclim. Palaeocol.*, 150: 83-121, Amsterdam.
- Perch-Nielsen K. (1983) - Recognition of Cretaceous stage boundaries by means of nannofossils. In Birkelund T., Bromley R., Christensen W.K., Håkansson E., Surlyk F. (eds.) Symposium on Cretaceous Stage Boundaries, Abstracts, 152-156, Copenhagen.
- Perch-Nielsen K. (1985) - Mesozoic calcareous nannofossils. In Bolli H.M., Saunders J.B., Perch-Nielsen K. (eds.) - Plankton Stratigraphy, 329-426, Cambridge University press, Cambridge.
- Premoli Silva I. (1977) - Upper Cretaceous-Paleocene magnetic stratigraphy at Gubbio, Italy. II. Biostratigraphy *Geol. Soc. Am. Bull.*, 88: 371-374, Boulder.
- Premoli Silva I., Sliter W.V. (1994) - Cretaceous planktonic foraminiferal biostratigraphy and evolutionary trends from the Bottaccione section, Gubbio, Italy. *Palaeontographia Italica*, 82: 1-85, Pisa.
- Premoli Silva I., Sliter W.V. (1999) - Cretaceous Paleocyanography: Evidence from planktonic foraminiferal evolution. In Barrera E., Johnson C.C. (eds.), The evolution of Cretaceous Ocean/Climatic System. Geological Society of America Special Paper, 332: 301-328, Boulder.
- Renard M. (1986) - Pelagic carbonate chemostratigraphy (Sr, Mg, ¹⁸O, ¹³C). *Mar. Micropaleont.*, 10: 117-164, Amsterdam.
- Renz O. (1936) - Stratigraphische und mikropalaeontologische Untersuchung der Scaglia (Obere Kreide-Tertiar) im zentralen Apennin. *Ecl. Geol. Helv.*, 42: 596-617, Basel.
- Roth P.H. (1978) - Cretaceous Nannoplankton Biostratigraphy and Oceanography of the Northwestern Atlantic Ocean. In Benson W.E., Sheridan R.E. et al. *Initial Report DSDP Leg 44*, 44: 731-759, Washington D.C.
- Roth P.H., Krumbach K.R. (1986) - Middle Cretaceous calcareous nannofossil biogeography and preservation in the Atlantic and Indian oceans: implications for paleoceanography. *Mar. Micropaleont.*, 10: 235-266, Amsterdam.
- Salaj J. (1980) - Microbiostratigraphie du Crétacé et du Paléogène de la Tunisie septentrionale et orientale (Hypostratotypes tunisiens). *Geol. Ustav Dionyza Stura*, 1-238, Bratislava.
- Salaj J. (1984) - Boundaries of Upper Cretaceous hypostratotypes at the profile Djebel Fguria Salah, Tunisia. *Bull. Geol. Soc. Denmark*, 33: 199-201, Copenhagen.
- Sissingh W. (1977) - Biostratigraphy of Cretaceous calcareous nannoplankton. *Geol. Mijnb.*, 56: 37-65, Amsterdam.
- Stoll H.M., Schrag D.P. (2000) - High Resolution Stable Isotope records from the Upper Cretaceous of Italy and Spain: Glacial Episodes in a Greenhouse Planet? *Geol. Soc. Am. Bull.*, 112: 308-319, Boulder.
- Thierstein H.R. (1976) - Mesozoic calcareous nannoplankton biostratigraphy of marine sediments. *Mar. Micropaleont.*, 1: 325-362, Amsterdam.
- Tremolada F., Erba E. (2002) - Morphometric analysis of Aptian Assipetra infracretacea and Rucinolithus terebrentarius nannoliths: Implications for taxonomy, biostratigraphy and paleoceanography. *Mar. Micropaleont.*, 44: 77-92, Amsterdam.
- Tröger K.A., Kennedy W.J. (1996) - The Cenomanian stage. Bulletin Institut Royal des Sciences Naturelles de Belgique, 6: 57-69, Bruxelles.
- Varol O. (1992) - Taxonomic revision of the Polycyclolithaceae and its contribution to Cretaceous biostratigraphy. *Newsletters on Stratigraphy*, 27: 93-127, Berlin-Stuttgart.
- Wright C.W., Kennedy W.J. (1981) - The Ammonoidea of the Plenus Marls and the Middle Chalk. *Palaeontographical Society Publication*, 560: 1-148, London.
- Watkins D.K. (1986) - Calcareous nannofossil paleoceanography of the Cretaceous Greenhorn Sea. *Geol. Soc. Am. Bull.*, 97: 1239-1249, Boulder.
- Watkins D.K., Wise S.W., Pospichal J.J., Crux J. (1996) - Upper Cretaceous calcareous nannofossil biostratigraphy and paleoceanography of the Southern Ocean. In Moguelevsky A., Whatley R. (eds.) Microfossils and Oceanic Environments, 355-381, Aberystwyth.

STAGE	SAMPLE	METER	Total abundance	Preservation	Species
CENOMANIAN	B 482.100.7	F P	R	R	R
	B 478.100.1	F P	R	R	R
	B 474.99.6	C P	R	R	R
	B 471.98.9	C P	R	R	R
	B 465.98.6	F P	R	R	R
	B 464.98.0	C P	R	R	R
	B 463.97.7	C P	R	R	R
	B 458.97.0	C P	R	R	R
	B 455.96.6	F P	R	R	R
	B 451.96.3	F P	R	R	R
	B 447.96.0	F P	R	R	R
	B 445.95.6	C P	R	R	R
	B 439.95.1	F P	R	R	R
	B 434.94.6	F P	R	R	R
	B 431.94.1	F P	R	R	R
	B 428.93.6	F P	R	R	R
	B 422.93.1	F P	R	R	R
	B 419.92.6	F P	R	R	R
	B 416.92.1	F M	R	R	R
	B 412.91.6	F P	R	R	R
	B 409.91.1	F P	R	R	R
	B 404.90.7	F P	R	R	R
	B 394.89.8	F P	R	R	R
	B 391.89.4	F P	R	R	R
	B 388.88.5	F P	R	R	R
	B 385.88.0	F P	R	R	R
	B 382.87.7	F P	R	R	R
	B 378.87.1	F P	R	R	R
	B 371.86.6	C P	R	R	R
	B 368.85.6	C P	R	R	R
	B 354.84.6	R P	R	R	R
	B 348.83.8	R P	R	R	R
	B 344.83.5	F P	R	R	R
	B 336.82.6	C P	R	R	R
	B 321.81.6	F P	R	R	R
	B 317.80.6	C P	R	R	R
	B 307.79.7	F P	R	R	R
	B 298.78.7	F P	R	R	R
	B 296.78.6	F P	R	R	R
	B 287.77.6	F P	R	R	R
	B 278.76.6	F P	R	R	R
	B 273.75.8	C M	R	R	R
	B 272.75.7	F P	R	R	R
	B 266.74.7	F P	R	R	R
	B 262.74.6	R P	R	R	R
	B 257.73.7	F P	R	R	R
	B 244.72.7	F P	R	R	R
	B 227.71.6	F P	R	R	R
	B 216.70.6	F P	R	R	R
	B 211.70.5	F P	R	R	R
	B 211.70.2	F P	R	R	R
	B 204.69.6	F P	R	R	R
	B 197.69.1	F P	R	R	R
	B 193.68.7	C P	R	R	R
	B 184.67.8	F P	R	R	R
	B 172.66.5	F P	R	R	R
	B 169.66.4	F M	R	R	R
	B 162.65.8	R P	R	R	R
	B 161.65.8	F P	R	R	R
	B 154.64.7	F P	R	R	R
	B 150.64.3	F P	R	R	R
	B 145.63.7	F P	R	R	R
	B 144.63.6	R P	R	R	R
	B 142.63.5	F P	R	R	R
	B 136.63.0	F P	R	R	R
	B 134.62.7	F P	R	R	R
	B 133.62.5	F P	R	R	R
	B 128.62.0	F P	R	R	R
	B 126.61.7	F P	R	R	R
	B 125.61.4	F P	R	R	R
	B 122.61.3	F P	R	R	R
	B 117.61.1	C M	R	R	R
	B 113.60.6	C M	R	R	R
	B 104.59.9	C B	R	R	R
	B 103.59.7	F P	R	R	R
	B 95.58.6	F P	R	R	R
	B 93.58.3	C M	R	R	R
	B 90.57.9	C M	R	R	R
	B 88.57.7	F P	R	R	R
	B 87.57.7	F P	R	R	R
	B 86.57.6	C M	R	R	R
	B 84.57.5	C P	R	R	R
	B 82.57.2	C M	R	R	R
	B 79.56.8	F P	R	R	R
	B 78.56.7	F P	R	R	R
	B 77.56.6	C M	R	R	R
	B 74.56.3	F P	R	R	R
	B 68.55.7	R P	R	R	R

Appendix 2. Range chart of calcareous nannofossils detected at Bottaccione from the Upper Albian to the Upper Cenomanian.

STAGE	SAMPLE	METER	Total abundance	CENOMANIAN		TURONIAN	
				Preservation		Preservation	
				R	P	R	P
	B. constans						
	Large B. constans						
	C. litterarius						
	C. surirellus						
	M. chistiatus						
	C. margerelii						
	E. turricifolius						
	L. camileensis						
	M. pennatoides						
	P. achlyostaurion						
	P. angustum						
	R. fenestratus						
	W. bamesae						
	W. communis						
	W. manivatae						
	Z. diplogrammus						
	C. conicus						
	P. embergeri						
	D. rotatorius						
	P. asper						
	D. lehmanni						
	L. floralis						
	Z. xenotus						
	C. ehrenbergii						
	P. splendens						
	C. angustifloratus						
	T. orionatus						
	G. nanum						
	Z. elegans						
	C. dispar						
	V. stradneri						
	Broussonetia sp.						
	B. signata						
	L. acutus						
	Nannococcus cf. truitii						
	V. matalosa						
	N. truitii						
	P. cretacea						
	Gartnerago sp.						
	C. kennedyi						
	Praediscosphaera sp.						
	A. albianus						
	E. trabeculatus						
	P. columnata						
	R. terebrodentarius						
	M. decoratus						
	G. obliquum						
	Q. garneri						
	Micula sp.						
	Lithastrinus 8el.						
	L. moratus						
	Q. enabrachium						
	E. eptapetalus						
	Quadrum sp.						
	E. monechiae						
	E. octopetalus						
	Lucianorhabdus sp.						
	Rucinolithus 7 el.						
	K. magnificus						
	Eprolithus sp.						
	Q. giganteum						
	Corollithus sp.						
	E. eximius						

Appendix 3. Range chart of calcareous nannofossils detected at Bottaccione from the Upper Cenomanian to the Upper Turonian.

