

# FORAMINIFERA AND CALCAREOUS NANNOPLANKTON ASSEMBLAGES FROM THE CENOMANIAN-TURONIAN BOUNDARY INTERVAL OF THE KNOVÍZ SECTION, BOHEMIAN CRETACEOUS BASIN



LENKA HRADECKÁ and LILIAN ŠVÁBENICKÁ

Czech Geological Survey, Klárov 131/3, 118 21 Prague, Czech Republic

(Manuscript received February 16, 1995; accepted in revised form June 14, 1995)

**Abstract:** Boreal foraminifers and calcareous nannofossils from the Cenomanian-Turonian boundary interval of the Bohemian Cretaceous Basin were studied. Detailed analyses of foraminiferal assemblages (benthonic diversity, diversity of species, ratio plankton/benthos) showed the important changes within this interval. The benthonic species are dominant within the foraminiferal assemblages, hence the representatives of genus *Gavelinella* were used for the biozonation. Effects of stressed life conditions on foraminiferal assemblages were recognized within this dysaerobic boundary interval. Similar effects are observed for calcareous nannofossils in the basal Turonian sediments.

**Key words:** Bohemian Cretaceous Basin, Cenomanian-Turonian boundary, *Foraminifera*, calcareous nannofossils.

## Introduction

At present, sedimentologists, paleontologists and geochemists are concerned with study of the Cenomanian-Turonian boundary sediments (Jarvis et al. 1988; Leary & Peryt 1990; Leary et al. 1989; Corfield et al. 1990; Kennedy & Cobbau 1991; Peryt 1991; Peryt & Wyrwicka 1991; Ernst et al. 1984; Walaszczyk 1987). Foraminiferal assemblages show significant changes in abundance and diversity through the Cenomanian-Turonian boundary interval. It was mentioned in many biostratigraphical studies respective to this problematics (Grosheny et al. 1992; Kaiho & Hasegava 1994; Olesen 1991; Robaszynski et al. 1993; Salaj 1987).

Rather sporadic changes in nannofossil assemblages are observed in the Cenomanian-Turonian boundary sediments compared to macrofossils and foraminifers. There are some quite significant blooms of individual taxa around the boundary but their importance is not fully understood (Bralower 1988).

The nannofossil biostratigraphic subdivisions for the Cenomanian-Turonian boundary interval were reviewed by Bralower (1988) who notes that nannofossil events differ in their stratigraphic positions according to published zonations. Many authors (Sissingh 1977; Perch-Nielsen 1985; Peryt & Wyrwicka 1993; Gorostidi & Lamolda 1993; Lamolda et al. 1994 etc.) consider the first appearance of *Quadrum gartneri* Prins & Perch-Nielsen as an indicator for the base of the Turonian stage. Nevertheless, Bralower (1988) found this species already in the uppermost Cenomanian deposits within the *Microstaurus chiastius* Subzone that correlates with *Neocardioceras juddii* ammonite Zone, i.e. with the lower part of the *Whiteinella* archaocretacea Zone. For the recognition of the Ce-Tu bound-

ary, Bralower (item) used the last occurrence of *Microstaurus chiastius* (Worsley) Grün.

Great attention has also been devoted to the Cenomanian-Turonian boundary interval in the Bohemian Cretaceous Basin (Čech & Váně 1989; Čech & Valečka 1991; Pražák 1989; Štemproková-Jírová 1991; Valečka & Skoček 1991; Uličný et al. 1993; Valečka et al. in press). This very important stratigraphical boundary-line is characterized by major changes in biostratigraphic, lithological and geochemical records in marine deposits.

The Cenomanian-Turonian boundary successions in the area of the Bohemian Cretaceous Basin are characterized by the Upper Cenomanian sediments of the Peruc-Korycany Formation (*Metoicoceras geslinianum* Zone) and the Lower Turonian sediments of the Bílá Hora Formation (*Watinoceras coloradoense* Zone). The Peruc-Korycany Formation is a transgressive succession of fluvial, lacustrine and lagoonal deposits (Peruc Member), passing into shallow-marine sandstones in the marginal parts and siltstones to calcareous siltstones in the deeper parts of the basin (Korycany Member).

The character of the Cenomanian-Turonian boundary varies in the Bohemian Cretaceous Basin mostly owing to variability of the Korycany Member facies. Siltstone facies is typically developed in the central part of the basin where it can be divided into two successive parts separated by a horizon rich in decalcified shell debris named "whitish broken fauna horizon" - WBF horizon sensu Pražák (1989). The Bílá Hora Formation is generally represented by marlstones and calcareous claystones.

The locality Knovíz is situated more in the western marginal part of the Bohemian Cretaceous Basin (Fig. 1), but the equivalent of siltstones facies is also developed here in the interval

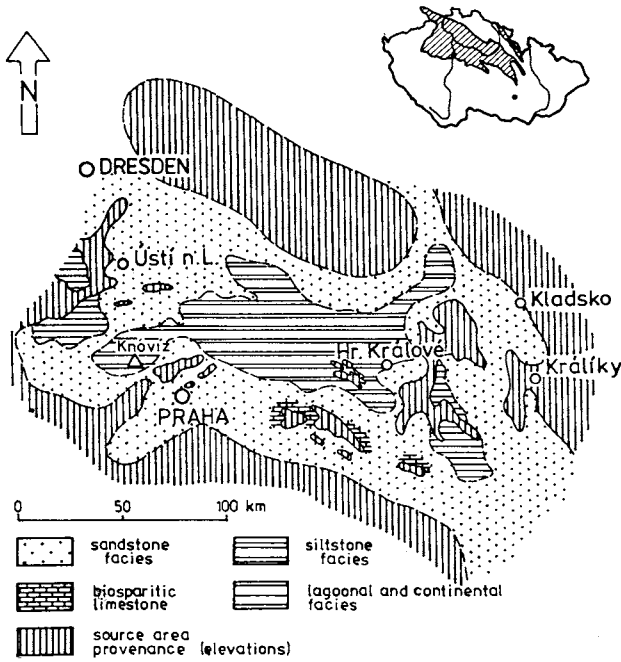


Fig. 1. Paleogeographical reconstruction of the Korycany Member (the Upper Cenomanian, *Metoicoceras geslinianum* Zone), modified after Čech & Valečka (1991).

from 0 to 4 m (Fig. 2). The deposits are represented by silty claystones to claystones with sporadic glauconite content. These sediments are dark in color in the interval from 2 to 4 m (Valečka et al. in press).

Methods

Material from the Knovíz section was collected from a 10 m thick interval reaching from the equivalent of the siltstone facies (Fig. 2) to the overlying beds represented by foraminiferal marlstones and well-bedded spiculite marlstones. The section was sampled with an average of approximately three–four samples per meter.

Foraminifers were isolated from sediment samples of approximately 0.75 kg weight using the usual methods of sieving through 0.063 mm sieves and silk. Diversity was determined using the Simpson equation. The illustrated specimens are stored in the Czech Geological Institute, Prague.

Nannofossils were present in the calcareous sediments of the Bílá Hora Formation only. This fact hindered study of nannofossil assemblage changes in the whole Cenomanian-Turonian boundary interval, and investigation had to be restricted to the basal Turonian deposits.

Nannofossils were studied using a Nikon light microscope with 1,000× magnification. For quantitative analysis, 500 specimens per sample were counted. Nannofossil species observed in this study are listed in the distribution table (Fig. 4).

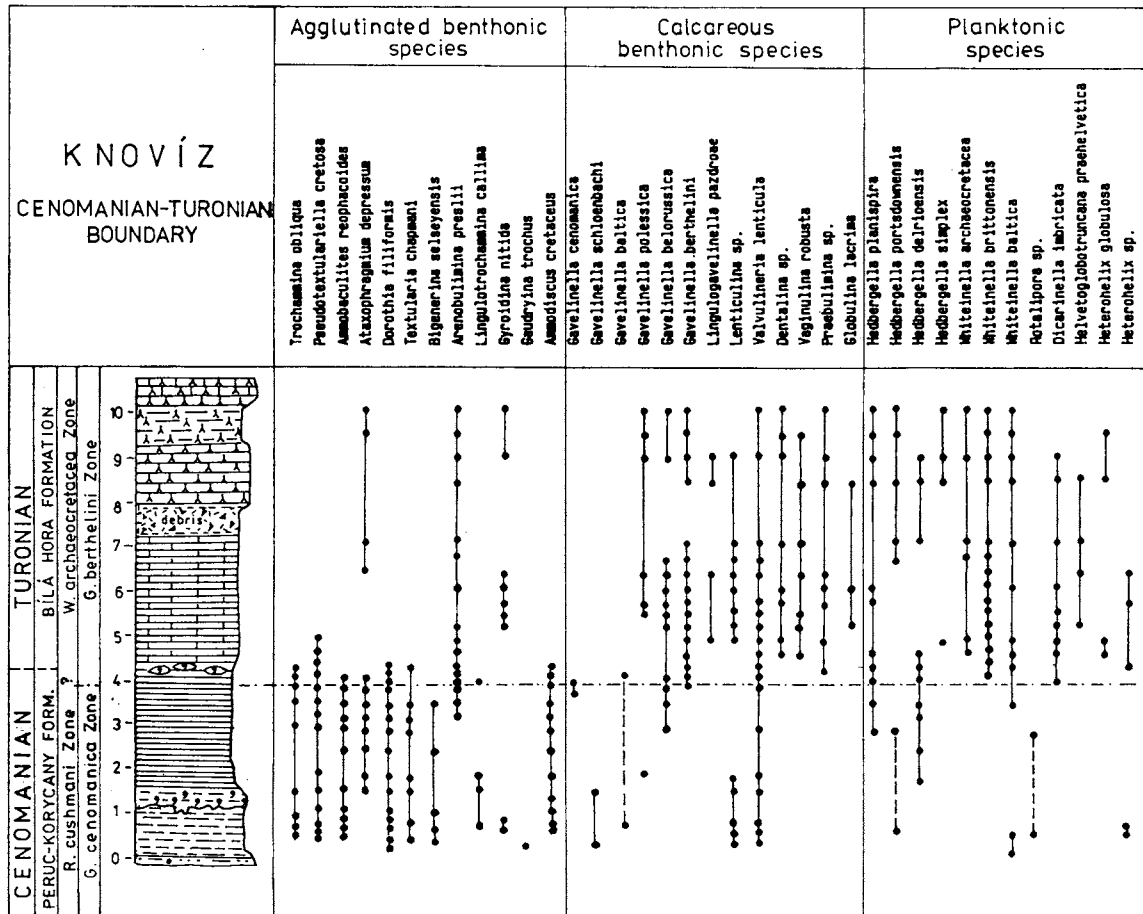


Fig. 2. Vertical distribution of the *Foraminifera* in the sediments of the Knovíz section.

## Results

### Foraminifera

Generally, the foraminiferal associations from these boundary sediments have a typical boreal character. The assemblages are characterized: **1** - by a low abundance of planktonic foraminifers, with small specimens of the genera *Hedbergella* and *Whiteinella* predominating, **2** - by practical absence or very rare presence of keeled forms. In these assemblages, benthic foraminifers predominate, especially agglutinated species (Pl. I, II). In the Knovíz section, the index species *Rotalipora cushmani* is extremely rare, as in the central part of the Bohemian Cretaceous Basin. Only two poorly preserved specimens were found in the sample from 0.4 m and one badly preserved specimen from 2.8 m probably belongs to the genus *Rotalipora* (Fig. 2). On this account, benthic species of the group *Gavelinella* have been used for the foraminiferal zonation of the Upper Cenomanian to lowermost Turonian sediments.

The foraminiferal assemblage taken from the interval 0.4-1.0 m is represented mostly by benthic species. The lowermost part (0.10 m) of the section contained both calcareous and agglutinated species in the mutual ratio 1:1, although towards the top of the bed the agglutinated species prevail (Fig. 3). In the agglutinated association, the following foraminifers dominate: *Haplophragmoides*, *Trochammina obliqua* and *Dorothia filiformis*. Calcareous benthos was represented mainly by the genus *Lenticulina*, *Valvulineria lenticula* and the group of gavelinels. This part is thought to be an equivalent to the lower part of the siltstone facies of the central part of the Bohemian Cretaceous Basin where some calcareous species are present near the bottom of the siltstone facies, but upwards they become extinct and are replaced by agglutinated forms.

In the Knovíz section, the bed of silty glauconite claystone with phosphatic nodules (1.2-1.5 m) is an equivalent to the WBF horizon. This horizon contains extremely rare plankton and even in 1.2 m of thickness no planktonic tests have been found.

The upper part of the siltstone facies in the sense of Pražák (1989) may be equivalent to the claystone bed in the Knovíz

section (1.5-4.0 m). Agglutinated specimens such as *Dorothia filiformis*, *Ataxophragmium depressum*, *Trochammina obliqua*, *Haplophragmoides* etc. were abundant.

Calcareous benthos is rarely represented by occasional findings of *Valvulineria lenticula* and some gavelinels.

Further up calcareous species are more common and are represented mainly by *Gavelinella berthelini*. Planktonic species become a common component of the assemblages.

Geochemical analyses from the Knovíz section (Valečka et al. in press) as well as from the central part of the Bohemian Cretaceous Basin show rather high contents of organic carbon in the interval of the siltstone facies (Uličný et al. 1993). In the Knovíz section, the higher organic carbon content was observed in the range of 2-4 m thickness and may indicate worse life conditions due to oxygen depletion (Fig. 3). The evidence for these dysaerobic conditions is the following (Koutsoukos et al. 1990): **1** - low assemblage diversity, **2** - poorly-developed plankton, dwarfish and juvenile specimens with practically no keeled forms, **3** - extinction of some benthic species.

Some benthic species had an ability to adapt to the worse life conditions. Above all, they were calcareous-hyaline foraminifers mostly free-living deposit-feeding specimens e.g. of semi-epifaunal or shallow infaunal mode of life (partially buried and shallow burrowers in soft muddy substrata) sensu Koutsoukos et al. (1990). Species belonging to this group found in the area of the Bohemia Cretaceous Basin are *Gavelinella berthelini*, *Lingulogavelinella globosa*, *Valvulineria lenticula*, *Praeubulimina*, *Lenticulina*, *Dentalina*, *Globulina lacrima* etc.

Agglutinated foraminifers indicating stressed life conditions are also representatives of an epifaunal or shallow infaunal habitat group (deposit feeders-detritivores or detrital/bacterial scavengers). They belong to the genera *Trochammina*, *Ammodiscus*, *Ammobaculites*, *Textularia* and *Tritaxia*.

### Calcareous nannofossils

The first nannofossils - the rare specimens of *Eprolithus ep-tapetalus* - were found in the sample immediately below the 2nd horizon rich in glauconite (Figs. 4 and 5). For more precise

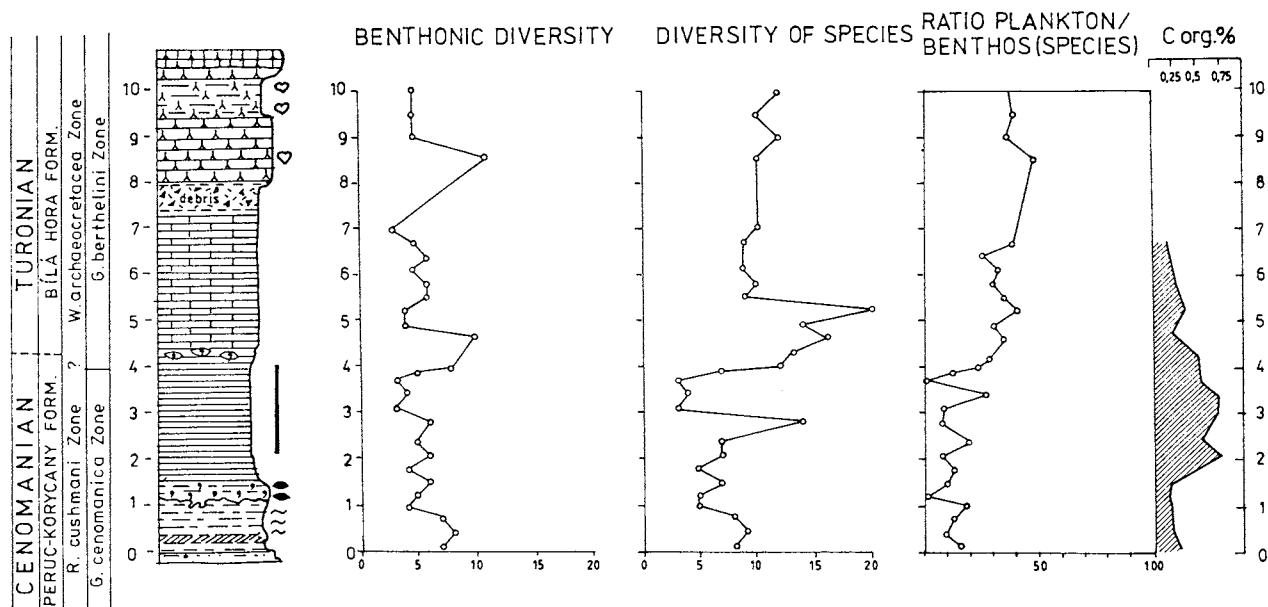


Fig. 3. The development of the foraminiferal assemblage across the Cenomanian-Turonian boundary and correlation with the total  $C_{org}$  content in the Knovíz section.

stratigraphical conclusions, this species is rather unsuitable because its occurrence is known from the Late Cenomanian to the Early Turonian.

Within the 2nd horizon rich in glauconite (at 4.3 m) and in the overlying sediments, the content of CaCO<sub>3</sub> rapidly increases from 0 % and reaches its maximum value 3.95 m above the base of this layer. Simultaneously, an abundant and diverse nanoflora suddenly appears.

In the 2nd horizon rich in glauconite, coccolith preservation is relatively poor and the high percentage content of *Watznaueria barnesae* (more than 70 %) indicates strong nanofossil dissolution (Roth & Krumbach 1986). *Broinsonia enormis*, *Eprolithus floralis*, *Prediscosphaera cretacea*, *Eiffelithus turriseiffelii*, *Glaucolithus diplogrammus* etc. occur frequently, with values over 1 %. An essential component of the assemblage comprises specimens of the family *Polycyclolithaceae*, such as *Quadrum gartneri*, *Q. intermedium*, *Q. sp. cf. Q. giganteum*, *Eprolithus* div. spec. including *E. octopetalus* and *Radiolithus orbiculatus* (Pl. III).

The presence of *Quadrum gartneri* allows us to recognize the standard nannoplankton Zone CC11 sensu Sissingh (1977) and Perch-Nielsen (1985) which documents the Early Turonian age. The same age is also shown by the presence of *Quadrum intermedium* which was reported by Varol (1992) from the short interval within the Early and Middle Turonian. Gorostidi & Lamolda (1993) found *Q. intermedium* already in the basal Turonian sediments associated with the first appearance of *Quadrum gartneri*.

At 4.60 m and in the overlying sediments, the character of the assemblages is rather changing. Calcareous nanofossils are better preserved and species diversity/abundance increases. *Watznaueria barnesae* decreases in abundance to about 35–45 % and this value remains practically unchanged in the rest of the section. Specimens of the genus *Eprolithus* are still common but the quantitative value of *Broinsonia enormis* already drops below 1 %.

The nanofossil assemblages keep this character until 8.50 m, where the species quantity of the whole family *Polycyclolithaceae* rapidly decreases. At 10.00 m, *Eprolithus octopetalus* was already not found. Varol (1992) mentioned *Eprolithus octopetalus* only within the short interval from the Late Cenomanian to the basal Turonian, but Gorostidi & Lamolda (1993) found it only in the latest Cenomanian.

In agreement with Roth & Krumbach (1986) and Bralower (1988), the higher quantity of specimens of the genus *Broinsonia* in the 2nd horizon rich in glauconite (at 4.3 m) may indicate neritic environments and shallow epicontinental seaways. This taxon might be an indicator of numerous factors associated with these environments including shallow waters, reduced salinity, or high fertility. According to Roth & Krumbach (1986), *Eprolithus floralis* is characteristic of high latitudes. Enrichment in this species might, therefore, be indicative of colder and/or fresher water. This would be consistent with the concurrent enrichments in arenaceous foraminifers and ostracods (Diner in Bralower 1988). The idea that both floral anomalies may result from inputs of fresher water is particularly attractive because such influxes, by stabilizing the water column, may have led to the deposition of the black shales which correlate with these floral anomalies.

### Biostratigraphy

The foraminiferal assemblages investigated from thirty samples from the Cenomanian-Turonian boundary interval of the

Knovíz section belong to the *Rotalipora cushmani* Zone and to the lower part of the *Whiteinella archaeocretacea* Zone. Because the index species *Rotalipora cushmani* (Morrow) is extremely rare in the studied section, the upper limit of the *Rotalipora cushmani* total range Zone could be determined using the extinction level of *Gavelinella cenomanica* Brotzen as an additional criterion. This species disappears either earlier or in some areas together with *R. cushmani* (Jarvis et al. 1988; Peryt & Wyrwicka 1991; Uličný et al. 1993). The overlying planktonic Zone *W. archaeocretacea* can be substituted by other species of the genus *Gavelinella* namely by *G. berthelini* (Keller) (Fig. 5).

### *Gavelinella cenomanica* Zone (*R. cushmani* Zone)

*G. cenomanica*, the index species for this zone becomes extinct within its upper limit. It occurs predominately in association with: *Gavelinella polesica*, *G. schloenbachi*, *Margulinella aequivoca*, *Trochammina obliqua*, *Dorothia filiformis*, *Haplophragmoides nonioninoides*, *Ataxophragmium depressum*, *Bigenerina selyensis* and *Ammobaculites reophacoides*.

### *Gavelinella berthelini* Zone (*W. archaeocretacea* Zone)

*G. berthelini* is maximally extended during this zone. Among others, it is accompanied by *Gavelinella belorussica*, *Valvulinella lenticula*, *Lingulogavelinella globosa*.

According to the foraminiferal study, the precise determination of the Cenomanian-Turonian boundary was impossible. The significant and abrupt change in the foraminiferal assemblages in this profile could be traced for about 4 metres. This change probably indicates the boundary-line between two foraminiferal zones. Nevertheless the first appearance of *Dicarinella imbricata* (Momod) and *Helvetoglobotruncana praeahelvetica* (Drujlik) could be indicate the Lower Turonian below the supposed boundary. This change probably indicates the boundary-line between two foraminiferal zones. The first appearance of the nanofossil species *Quadrum gartneri* Prins & Perch-Nielsen was recorded in the sample from 4.3 m and may indicate the base of the Turonian. But this stratigraphical information is debased by nanofossil scarcity in the underlying non-calcareous sediments. A sample from 4.0 m yielded sporadic specimens only of *Eprolithus eptapetalus* which already is known from the Late Cenomanian. According to these data, the Cenomanian-Turonian boundary might be situated within the interval of the 2nd horizon rich in glauconite, i.e. in 4.3 m (Fig. 5).

### Conclusion

The performed analyses may be correlated with the result of studies carried out in the Cenomanian-Turonian boundary sediments from the surrounding areas. Detailed analyses of foraminiferal assemblages showed the important changes within this interval. The indicated changes are probably closely related to the life conditions worsening as a result of oxygen shortage. For the foraminiferal zonation, representatives of genus *Gavelinella* were used (*G. cenomanica* and *G. berthelini*). They are abundant in the assemblages of the boreal bioprovinces.

An abundant and diverse nannoplankton occurs at the Knovíz section only in the calcareous sediments of the Bílá Hora Formation and allows us to recognize the *Quadrum gartneri* Zone

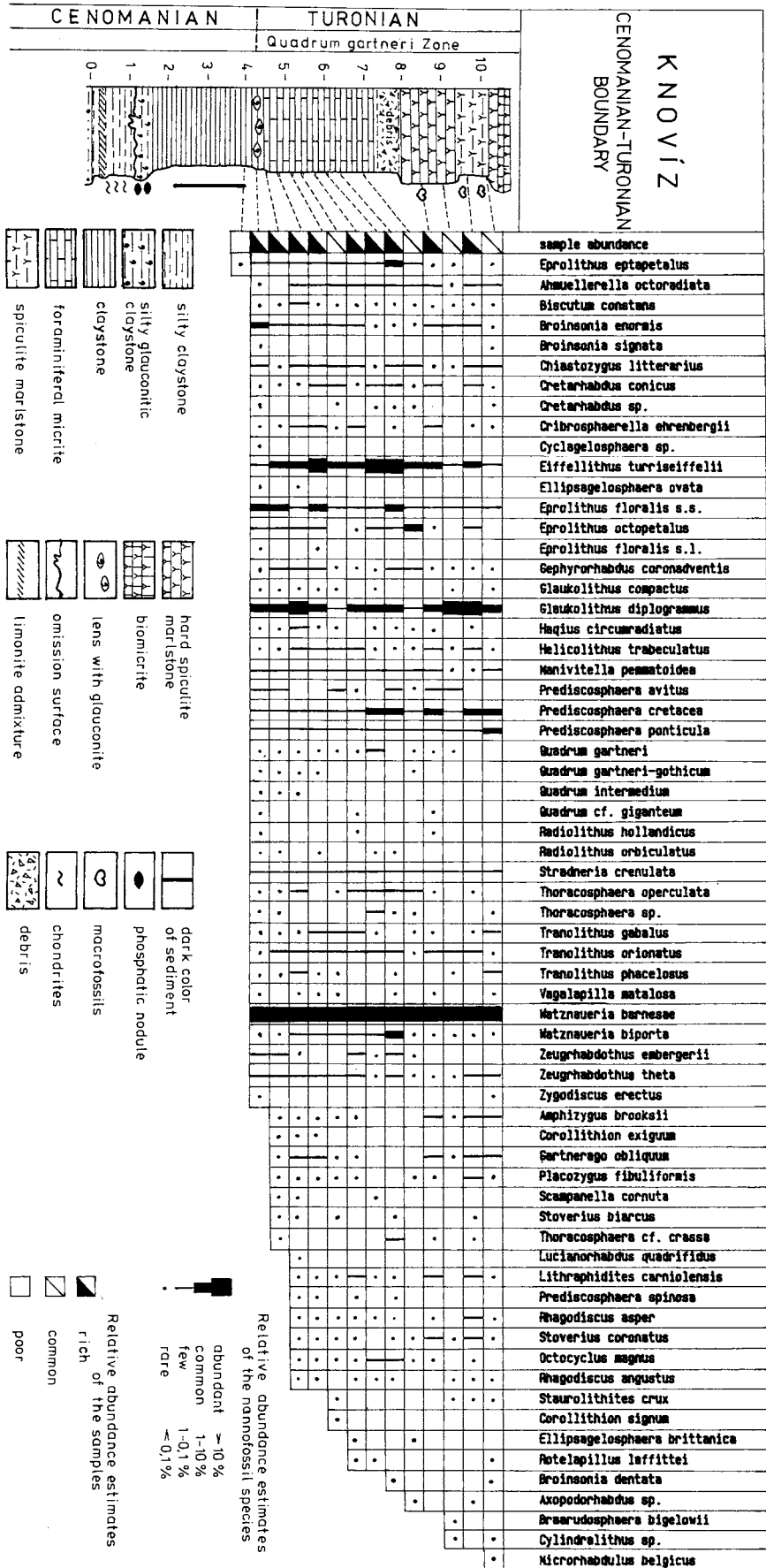
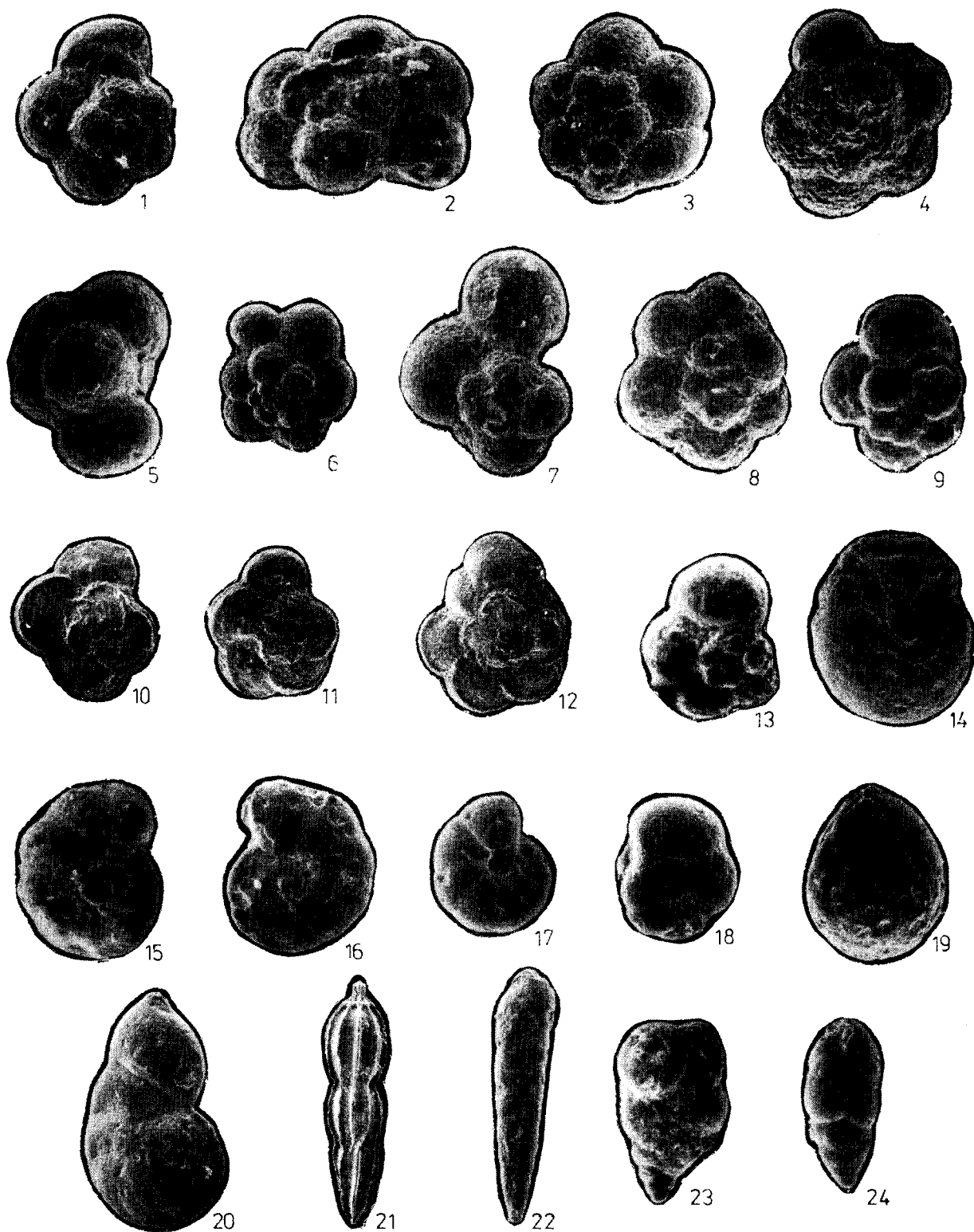


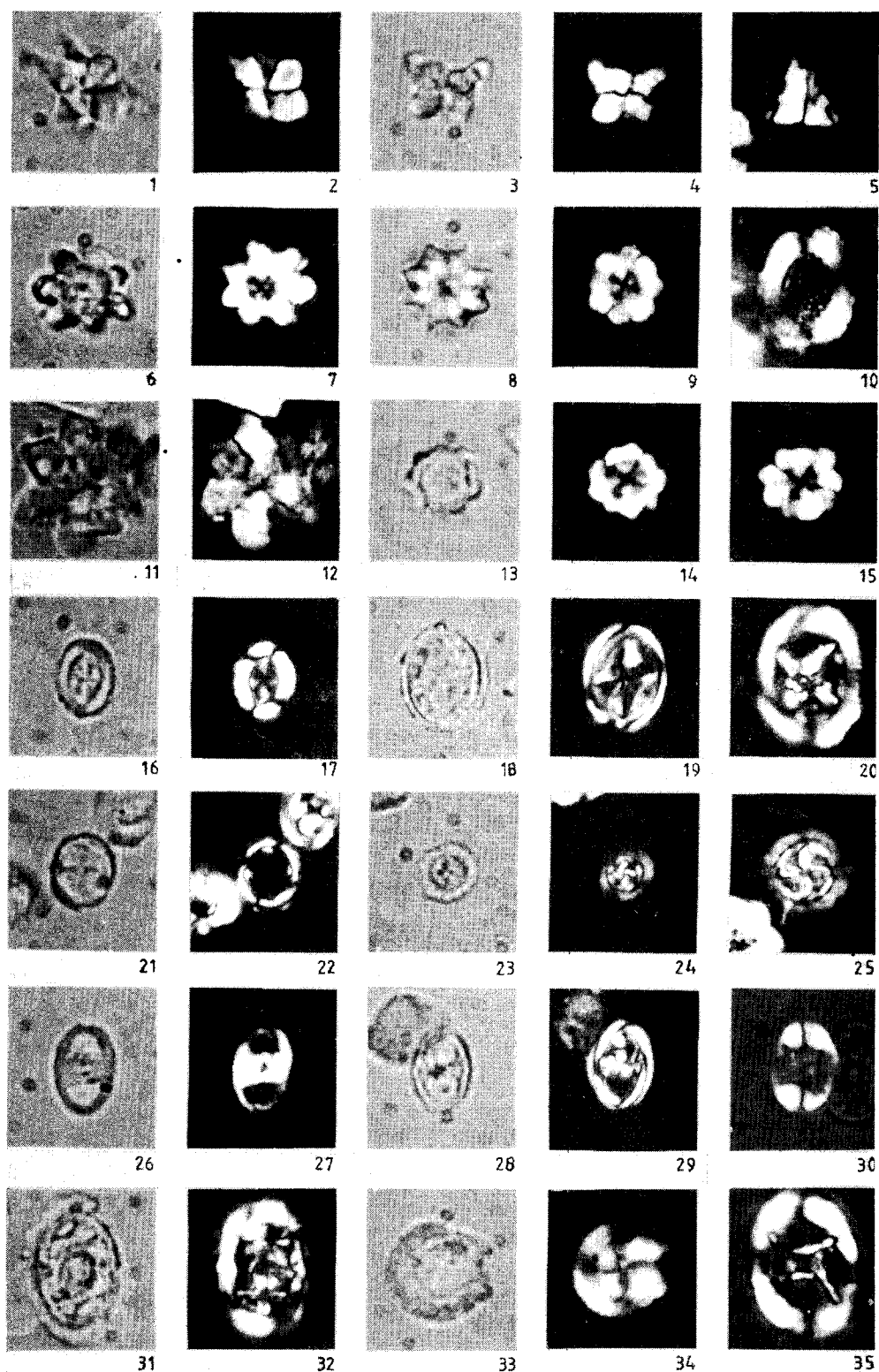
Fig. 4. Vertical distribution of calcareous nannofossils in the sediments of the Knoviz section.



**Plate I:** SEM photomicrographs of Cenomanian and Turonian *Foraminifera* from Knoviz section. **Figs. 1, 2.** *Helvetoglobotruncana praehelvetica* (Trujillo), 5.2 m, 80 $\times$ ; **Fig. 3** *Whiteinella brittonensis* (Loeblich & Tappan), 5.5 m, 80 $\times$ ; **Fig. 4.** *Hedbergella portdownensis* (Williams - Mitchell), 0.7 m, 80 $\times$ ; **Fig. 5.** *Whiteinella paradubia* (Sigal), 2.4 m, 85 $\times$ ; **Fig. 6.** *Whiteinella brittonensis* (Loeblich & Tappan), 5.2 m, 50 $\times$ ; **Fig. 7.** *Hedbergella simplex* (Morrow), 5.5 m, 80 $\times$ ; **Fig. 8.** *Whiteinella brittonensis* (Loeblich & Tappan), 2.4 m, 90 $\times$ ; **Fig. 9.** *Whiteinella baltica* Douglas & Rankin, 5.2 m, 90 $\times$ ; **Figs. 10, 11.** *Dicarinella* sp., 5.2 m, 50 $\times$ ; **Fig. 12.** *Dicarinella imbricata* (Mornod), 5.2 m, 50 $\times$ ; **Fig. 13.** *Hedbergella* sp., 5.2 m, 130 $\times$ ; **Figs. 14, 15.** *Gavelinella berthelini* (Keller), 3.9 m, 90 $\times$ ; **Fig. 16.** *Gavelinella berthelini* (Keller), 9.0 m, 80 $\times$ ; **Fig. 17.** *Valvulineria lenticula* (Reuss), 3.9 m, 90 $\times$ ; **Fig. 18.** *Gavelinella polesseica* Akimec, 9.0 m, 80 $\times$ ; **Fig. 19.** *Globulina lacrima* (Reuss), 3.9 m, 90 $\times$ ; **Fig. 20.** *Lenticulina* sp., 5.2 m, 50 $\times$ ; **Fig. 21.** *Dentalina* sp., 5.5 m, 50 $\times$ ; **Fig. 22.** *Cassidella tegulata* (Reuss), 5.2 m, 70 $\times$ ; **Fig. 23.** *Heterohelix* sp., 5.2 m, 90 $\times$ ; **Fig. 24.** *Praebulimina* sp., 5.5 m, 90 $\times$ . Photomicrographs by Ananda Gabašová, Czech Geological Survey Prague.



**Plate II:** SEM photomicrographs of Cenomanian and Turonian *Foraminifera* from Knoviz section **Figs. 1, 2.** *Haplophragmoides nonioninoides* (Reuss), 1.0 m, 80 $\times$ ; **Fig. 3.** *Trochammina obliqua* Tappan, 2.8 m, 50 $\times$ ; **Fig. 4.** *Haplophragmoides* sp., 1.0 m, 70 $\times$ ; **Fig. 5.** *Lingulotrochammina callima* (Loeblich & Tappan), 1.8 m, 50 $\times$ ; **Fig. 6.** *Ammobaculoides lepidus* Hercogova, 1.8 m, 70 $\times$ ; **Fig. 7.** *Ammobaculites reophacoides* Bartenstein, 1.8 m, 50 $\times$ ; **Fig. 8.** *Ammobaculites reophacoides* Bartenstein, 2.4 m, 70 $\times$ ; **Figs. 9, 10.** *Spiroplectamina scotti* Cushman-Alexander, 1.2 m, 70 $\times$ ; **Fig. 11.** *Bigenerina selseyensis* Heron-Allen-Earland, 3.4 m, 70 $\times$ ; **Fig. 12.** *Gaudryina praepyramidata* Hercogova, 1.5 m, 50 $\times$ ; **Fig. 13.** *Tritaxia compressa* Egger, 1.0 m, 50 $\times$ ; **Fig. 14.** *Textularia foeda* Reuss, 3.4 m, 50 $\times$ ; **Fig. 15.** *Dorothia* sp., 1.8 m, 70 $\times$ ; **Fig. 16.** *Dorothia filiformis* (Berthelin), 1.8 m, 70 $\times$ ; **Fig. 17.** *Pseudotextularia cretosa* (Cushman), 1.8 m, 70 $\times$ ; **Fig. 18.** *Arenobulimina preslii* (Reuss), 3.9 m, 50 $\times$ ; **Fig. 19.** *Ataxophragmium depressum* (Perner), 3.9 m, 50 $\times$ ; **Fig. 20.** *Glomospirella gaultina* (Berthelin), 1.0 m, 50 $\times$ ; **Fig. 21.** *Ammodiscus cretaceus* (Reuss), 1.0 m, 50 $\times$ ; **Fig. 22.** *Arenobulimina brevicona* (Perner), 3.9 m, 90 $\times$ ; **Fig. 23.** *Lituotuba incerta* Franke, 3.1 m, 50 $\times$ ; **Fig. 24.** *Lituotuba* sp., 3.4 m, 50 $\times$ . Photomicrographs by Ananda Gabašová, Czech Geological Survey Prague.



**Plate III:** Knoviz section, sample taken from 4.60 m (with the exception of *Quadrum* cf. *giganteum* that comes from 8.50 m); Figs. 1, 2. *Quadrum intermedium* Varol; Figs. 3, 4. *Quadrum gartneri-gothicum*; Fig. 5. *Lucianorhabdus quadrifidus* Forchheimer; Figs. 6, 7. *Eprolithus eptapetalus* Varol; Figs. 8, 9. *Eprolithus octopetalus* Varol; Fig. 10. *Cretarhabdus* sp.; Figs. 11, 12. *Quadrum* cf. *giganteum* Varol; Figs. 13, 14. *Radiolithus orbiculatus* (Forchheimer) Varol; Fig. 15. *Eprolithus floralis* Stradner; Figs. 16, 17. *Broinsonia enormis* (Shumenko) Manivit; Figs. 18, 19. *Gartnerago obliquum* (Stradner) Noel; Fig. 20. *Gephyrorhabdus coronadventis* (Reinhardt) Hill; Figs. 21, 22. *Stoverius coronatus* (Bukry) Perch-Nielsen; Figs. 23, 24. *Prediscosphaera avitus* (Black) Perch-Nielsen; Fig. 25. *Prediscosphaera ponticola* (Bukry) Perch-Nielsen; Figs. 26, 27. *Tranolithus gabalus* Stover; Figs. 28, 29. *Placozygus fibuliformis* (Reinhardt) Hoffmann; Figs. 30, 35. *Eiffellithus turriseiffelii* (Deflandre) Reinhardt; Figs. 31, 32. *Octocyclus magnus* Black; Figs. 33, 34. *Haqius circumradiatus* (Stover) Roth Magnification: 2,000 $\times$ . Photomicrographs by Lilian Švabeničká.



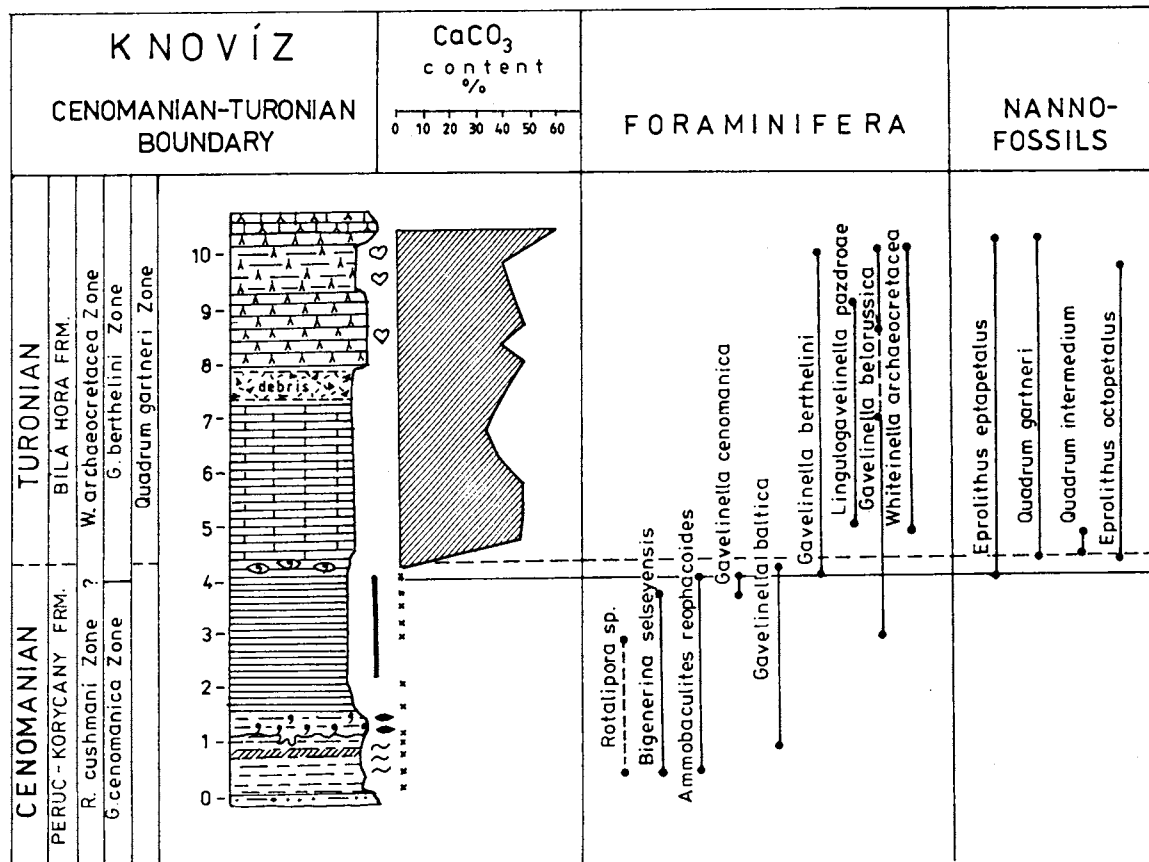


Fig. 5. Knovíz section - distribution of the stratigraphical important *Foraminifera* and calcareous nannofossils and its correlation with the  $\text{CaCO}_3$  content.

(sensu Sissingh 1977 and Perch-Nielsen 1985) from the base of the 2nd horizon rich in glauconite (from 4.30 m) upwards.

The percentage of *Watznaueria barnesae* indicates a decreasing dissolution of coccoliths in an upward direction across the basal Turonian sediments. *Eprolithus floralis* and *Broinsonia enormis* become abundant in the assemblage in the sample taken from the 2nd horizon rich in glauconite. This fact may indicate colder water and shallow epicontinental seaways.

**Acknowledgement:** We are grateful to Dr. Jaroslav Valečka and to Dr. Stanislav Čech (Czech Geological Survey, Prague), who kindly shared their informations with us on the Cenomanian-Turonian boundary in the Bohemian Cretaceous Basin as well as on the Knovíz section. We would like to thank Dr. Adam Gasinski, Dr. Wolfgang Kuhnt and Dr. Jozef Salaj for their additional remarks during the edition of this paper.

## References

- Bralower T.J., 1988: Calcareous nannofossil biostratigraphy and assemblages of the Cenomanian-Turonian boundary interval: Implications for the origin and timing of oceanic anoxia. *Palaeoceanography*, 3, 275-316.
- Corfield R.M., Hall M.A. & Brasier M.D., 1990: Stable isotope evidence for foraminiferal habitats during the development of the Cenomanian-Turonian oceanic anoxic event. *Geology*, 18, 175-178.
- Čech S. & Váně M., 1988: Facies development of the Cenomanian and Lower Turonian at the foothills of the Krušné hory Mts. *Časopis pro mineralogii a geologii*, Praha, 33, 395-410 (in Czech, English summary).
- Čech S. & Valečka J., 1991: Significant transgressions and regressions in the Bohemian Cretaceous Basin. *Manuscript, Czech Geological Survey*, Prague, 1-51 (in Czech).
- Ernst G., Wood C.J. & Hilbrecht H., 1984: The Cenomanian-Turonian boundary problem in NW-Germany with comments on the north-south correlation to the Regensburg Area. *Bulletin of the Geological Society of Denmark*, 33, 103-113.
- Gorostidi A. & Lamolda M.A., 1993: Field-guide excursion to the Maastrichtian-Danian, Zumaya Section, and to the Cenomanian-Turonian, Menoyo Section (Northern Spain). *Excursion Guide of the 5th INA Conference*, Salamanca, 1-18.
- Grosheny D., Tronchetti G. & Schaaf A., 1992: Nouvelles données sur les foraminifères planctoniques et la biostratigraphie du Céno-manién-Turonien dans le S.E. du bassin vocontien (S.E. France). *C.R. Acad. Sci.*, Paris, 315, Série II, 773-776.
- Jarvis I., Carson G.A., Cooper M.K.E., Hart M.B., Leary P.N., Tocher B.A., Horne D. & Rosenfeld A., 1988: Microfossil assemblages and the Cenomanian-Turonian (Late Cretaceous) Oceanic Anoxic Event. *Cretaceous Research*, 9, 3-103.
- Kaiho K. & Hasegawa T., 1994: End-Cenomanian benthic foraminiferal extinctions and oceanic dysoxic events in the northwestern Pacific Ocean. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 111, 29-43.
- Kennedy W.J. & Cobban W.A., 1991: Stratigraphy and interregional correlation of the Cenomanian-Turonian transition of the Western Interior of the United States near Pueblo, Colorado, a potential boundary stratotype for the base of the Turonian Stage. *Newsletters on Stratigraphy*, 24, 1-33.
- Koutsoukos E.A.M., Leary P.N. & Hart M.B., 1990: Latest Cenomanian - earliest Turonian low-oxygen tolerant benthonic foraminifera: a case study from the Sergipe Basin (N.E. Brazil), and the western Anglo-Paris Basin (southern England). *Palaeogeography, Palaeoclimatology, Palaeoecology*, 77, 145-177.

- Lamolda M.A., Gorostidi A. & Paul C.R.C., 1994: Quantitative estimates of calcareous nannofossil changes across the Plenus Marls (latest Cenomanian), Dover, England: implications for the generation of the Cenomanian-Turonian Boundary Event. *Cretaceous Research*, 15, 143-164.
- Leary P.N., Carson G.A., Cooper M.K., Hart M.B., Horne D., Rosenfeld A. & Tocher B.A., 1989: The biotic response to the Late Cenomanian oceanic anoxic event: integrated evidence from Dover, SE England. *Journal Geol. Soc., London*, 146, 311-317.
- Leary P.N. & Peryt D., 1990: The Late Cenomanian oceanic anoxic event in the western Anglo-Paris Basin and SE Danish-Polish Trough: Survival strategies of and recolonization by benthonic Foraminifera. *Historical Biology*, 5, 321-338.
- Olesen J., 1991: Foraminiferal biostratigraphy and paleoecology of the Mancos Shale (Upper Cretaceous), southwestern Black Mesa, Arizona. In: Nations J.D. & Eaton J.G. (Eds.): *Stratigraphy, depositional environments, and sedimentary tectonics of the western margin, Cretaceous Western Interior Seaway*. Geological Society of America, *Special Paper*, 260, 153-164.
- Perch-Nielsen K., 1985: Mesozoic calcareous nannofossils. In: Bolli H.M., Saunders J.B. & Perch-Nielsen K. (Eds.): *Plankton Stratigraphy*. Cambridge Earth Sci. Ser., Cambridge, 329-426.
- Peryt D., 1991: Foraminiferal response to the Late Cenomanian Oceanic Anoxic Event in central Poland. *Géologie Alpine, Mémoire h.s.*, 17, 101.
- Peryt D. & Wyrwicka K., 1991: The Cenomanian-Turonian boundary event in SE Poland. *Cretaceous Research*, 12, 65-80.
- Peryt D. & Wyrwicka K., 1993: The Cenomanian/Turonian boundary event in Central Poland. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 104, 185-197.
- Pražák J., 1989: The Cenomanian-Turonian boundary in the central part of the Bohemian Cretaceous Basin. *Manuscript, Czech Geological Survey, Prague*, 1-34 (in Czech).
- Robaszynski F., Hardenbol J., Caron M., Amédro F., Dupuis Ch., González Donoso J.-M., Linares D. & Gartner S., 1993: Sequence stratigraphy in a distal environment: The Cenomanian of the Kalaat Senan region (Central Tunisia). *Bull. Centres Rech. Explor. - Prod. Elf Aquitaine, Boussens*, 17, 2, 395-433.
- Roth P.H. & Krumbach K.P., 1986: Middle Cretaceous calcareous nannofossil biogeography and preservation in the Atlantic and Indian Oceans: Implications for paleoceanography. *Marine Micropalaeontology*, 10, 235-266.
- Salaj J., 1987: Present trends in the stratigraphy of the Upper Cretaceous units and their application in the Western Carpathians. *Geol. Práce, Spr.*, 86, 197-209 (in Slovak).
- Sissingh W., 1977: Biostratigraphy of Cretaceous calcareous nannoplankton. *Geol. en Mijnb.*, 56, 37-65.
- Štemproková-Jírová D., 1991: Biostratigraphy of planktic Foraminifera from the Cenomanian and Turonian of the Velim (Bohemian Cretaceous Basin, Czechoslovakia). *Acta Univ. Carolinae*, 1-2, 76-81.
- Uličný D., Hladíková J. & Hradecká L., 1993: Record of sea-level changes, O depletion and the  $^{13}\text{C}$  anomaly across the Cenomanian-Turonian boundary, Bohemian Cretaceous Basin. *Cretaceous Research*, 14, 211-234.
- Valečka J. & Skoček V., 1991: Late Cretaceous lithoevents in the Bohemian Cretaceous Basin, Czechoslovakia. *Cretaceous Research*, 12, 561-577.
- Valečka J., Čech S., Hladíková J., Hradecká L. & Švábenická L., in press: Lithological, faunistic and geochemical changes recorded in the Cenomanian-Turonian Knovíz section, NW of Prague (Bohemian Cretaceous Basin). *Věst. Čes. geol. úst.*, Praha.
- Varol O., 1992: Taxonomic revision of the *Polycyclolithaceae* and its contribution to Cretaceous biostratigraphy. *Newsletter Stratigraphy*, 27, 93-127.
- Walaszczyk I., 1987: Mid-Cretaceous events at the marginal part of the Central European Basin (Annapol-on-Vistula section, Central Poland). *Acta Geol. Pol.*, 37, 1-2.