

From dysoxic sea to hypersaline lagoon: paleoenvironmental changes on the Badenian/Sarmatian boundary (borehole MZ102; Vienna Basin)

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Abstract: The main objective of this work was to interpret the environment during a very dynamic paleoenvironmental change from the Badenian to the Sarmatian in the Central Paratethys, as well as to determine ecologic factors, diversity, and the relationship between the foraminiferal morphogroups in the northern part of the Vienna Basin. The samples of this study come from the MZ 102 borehole, which had been drilled near the town of Malacky in the northern part of the Vienna Basin (Slovakia) (Fig.). Based on sedimentology and fossil occurrence, we interpreted a paleoenvironment of the northern part of the Vienna Basin during the middle Miocene. By examining changes in the occurrence of foraminifera (*Articulina sarmatica*, *Cycloforina stomata*, *Tenuitella acme*) and calcareous nannofossils (*Coccolithus miopelagicus*) in the studied samples, we detected the position of the Badenian/Sarmatian boundary. Low-oxic conditions in the uppermost Badenian (Studienka Fm.) and upper Sarmatian (Skalica Fm.) sediments were interpreted, as well as fluctuation from a normal marine to hypersaline shallow water paleoenvironment in the middle and upper Sarmatian sediments (Skalica Fm.). Macrofauna was represented mostly by bivalves and gastropods shells and fish remains. Samples were compared with samples from the nearby MZ 93 borehole to fill in the gaps in the MZ 102 borehole record.

Keywords: Vienna Basin, Foraminifera, Paleoenvironment, Middle Miocene, Calcareous nannoplankton, sedimentology

1. INTRODUCTION

The nutrient content, salinity, type of substrate, and change of oxygen concentration in water at sediment surface may play an important role in the presence and distribution of benthic foraminifera (Sen Gupta, 2003). One of the classical Cenozoic basins for the study of foraminifera in the Central Paratethys is the Vienna Basin, which has been of great interest since the pioneer monograph of d'Orbigny (1846). The Vienna Basin is 200 km long and 55 km wide Neogene basin, oriented SSW-NNE, and nowadays situated in three states: Slovakia, Austria, and the Czech Republic (Kováč et al., 2004). The geological, geodynamic, and tectonic development of the basin has been previously described in various studies (e.g., Ratschbacher et al., 1991a,b; Csontos et al., 1992; Kováč et al., 1994, 1997, 1998, 2016, 2017a; Matenco, 1997; Konečný et al., 2002; Schmid et al., 2008; Ustaszewski et al., 2008; Handy et al., 2010, 2015; Studencka et al., 2016). Central Paratethys has been studied in detail by Senes (1961a,b); Kováč et al. (1994, 1998, 2003, 2016, 2017b); Meulenkamp et al. (1996); Oszczypko (1998, 1999); Popov et al. (2004); Oszczypko & Oszczypko-Clowes (2006); Piller et al. (2007); Gozhyk et al. (2015) and Neubauer et al. (2015).

During the middle Badenian, sea regression took place, which caused changes in water salinity (Harzhauser & Piller, 2004; Koubová & Hudáčková, 2010) and extinction or endemism of

fauna and flora (Harzhauser & Piller, 2007; Palcu et al. 2015). In the upper Badenian/Sarmatian boundary, transgression occurred, causing low-oxic bottom waters with *Bolivina-Bulimina* zonation (Kováč et al., 2017b). At the end of the Pannonian age, the sedimentary environment became hyposaline (Kováč et al., 2008a,b) with changes of foraminifera assemblages to very low diversified to monospecific (Hudáčková et al. 2018, 2019).

The aim of this work was to interpret the origin and environment of collector sediments. Main goal was to interpret the paleoenvironment of the northern part of the Vienna Basin around the Badenian/Sarmatian boundary based on the foraminiferal associations from the studied borehole, and to determine ecologic factors which provoked the change in species diversity. Finally, we compared the functionality and complexity of two methodological approaches for the paleoecological interpretations: foraminiferal species versus foraminiferal morphogroups study.

1.1. Geological settings and methods

Samples come from the MZ 102 borehole, which had been drilled in the northern part of the Vienna Basin, west of the town of Malacky. The natural gas field Malacky-West represents an important geological structure, which in the past produced almost 1.3 billion m³ of gas, while most of the production comes from the sandy horizons of the upper Badenian and Sarmatian deposits (Badenian and Sarmatian sands specified by Nafta a. s.; Fig. 2)

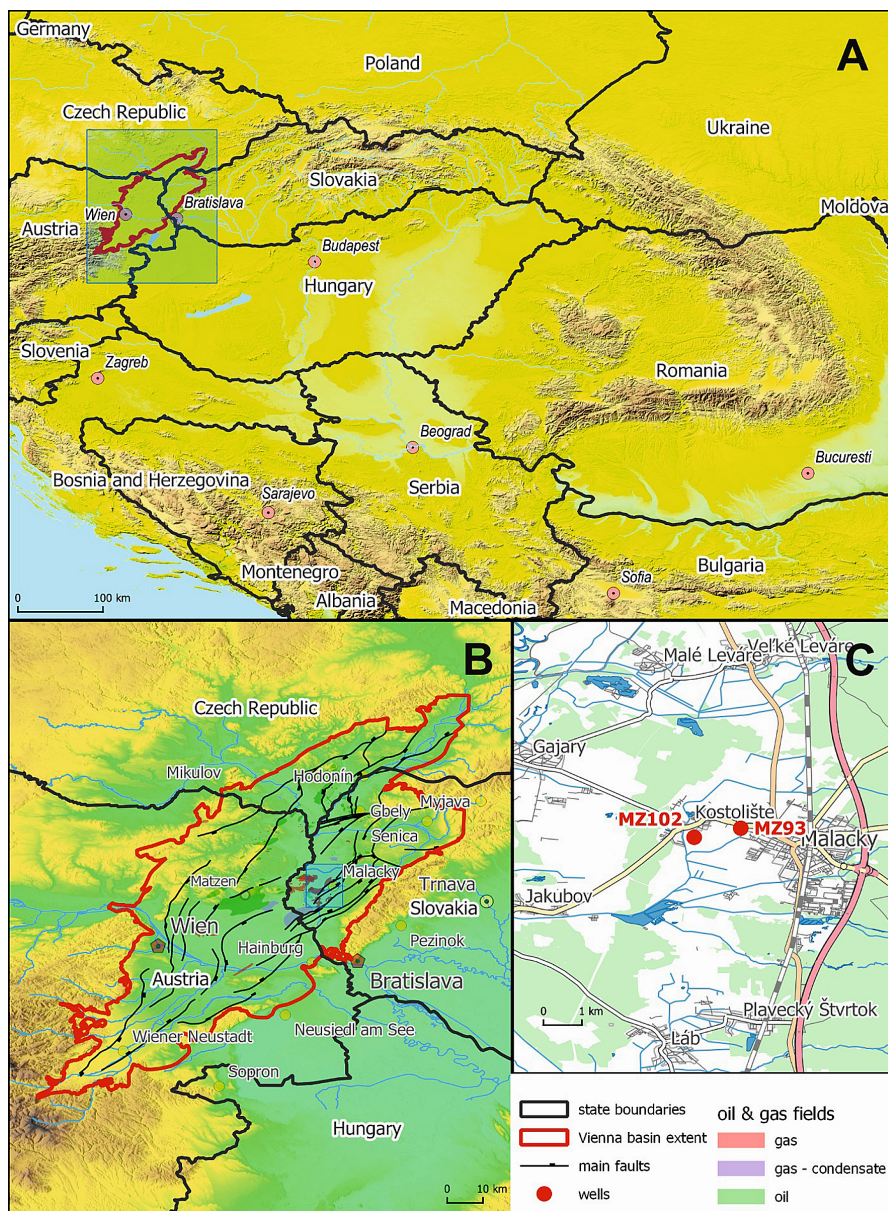


Fig. 1. Localization map (A – Europe; B – Vienna basin, C – boreholes MZ 93 and MZ 102).

(Hudeček & Šály 1995). The depleted gas field is currently being converted into an underground gas storage facility (UGS). The geological structure of the upper Badenian and Sarmatian sediments is well-documented in this area by drilling and 3D seismic data. The MZ 102 borehole was drilled in April 2013 in the SW part of UGS Malacky, the final depth reaching 1215 m.

1.2. Sedimentology

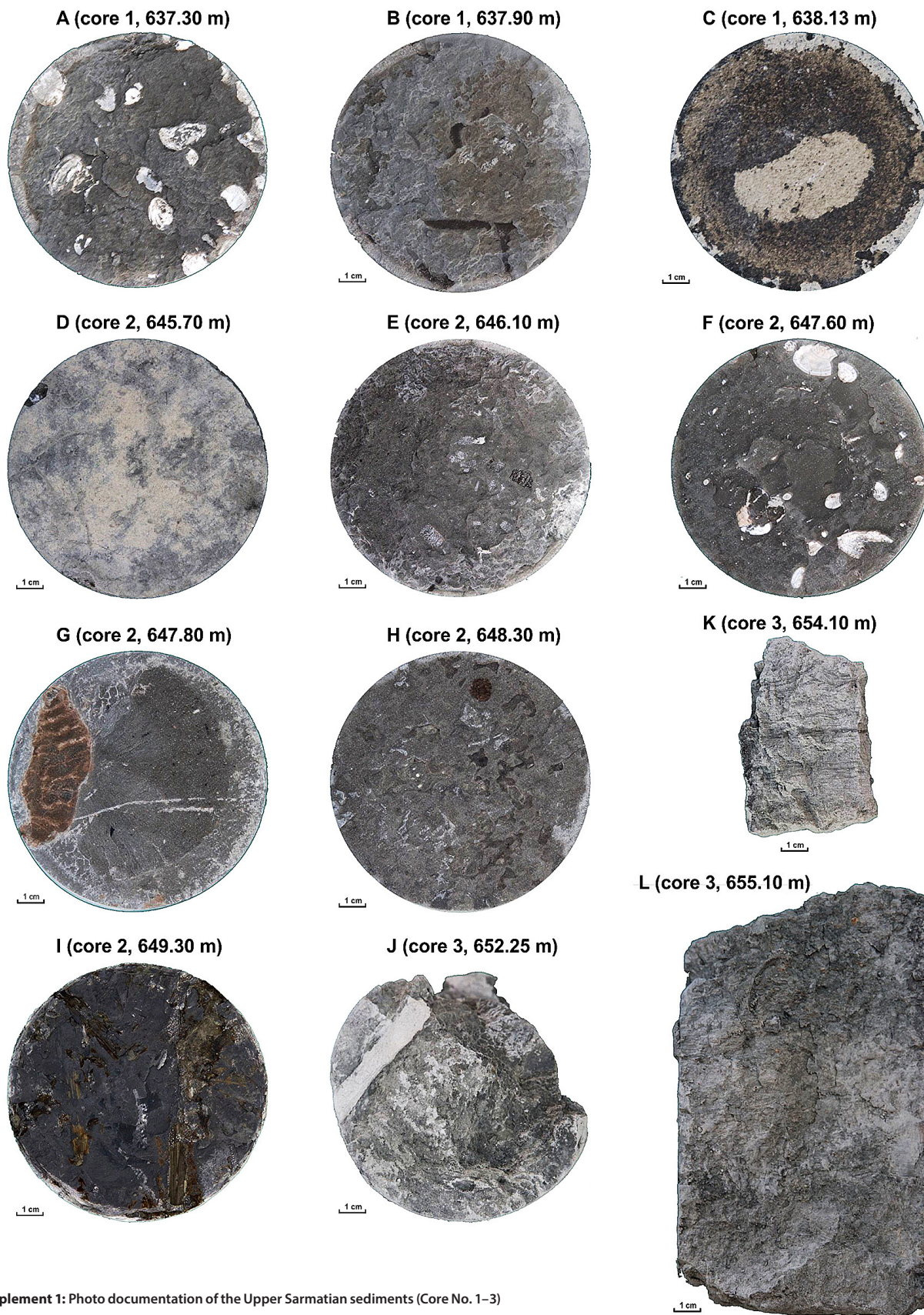
The aim of the borehole was to verify the existence, thickness, and reservoir parameters of UGS objects (1st Badenian – Main sand and 1st Sarmatian sand; see Fig. 2), as well as to specify the geological setting of the upper Badenian, Sarmatian, and Pannonian strata in the Malacky area. A special goal of the borehole was the extraction of rock samples from the drill cores to determine detailed petrophysical parameters of UGS objects and their sealings (Wallner et al. 2014). Both studied boreholes, MZ 102 and

MZ 93, are located near the town of Malacky in the area of the Záhorská nížina plain (Figs. 1, 2). Although samples from borehole MZ 102 were taken from the entire solid rock profile, studied borehole (Figs. 3A–F) data consists of parts that anticipated a Badenian/Sarmatian boundary and the upper Sarmatian: core No. 2 (644–651 m), No. 3 (652.2–659.8 m), No. 4 (1034–1043.1 m), No. 5 (1043.1–1051.6 m) and No. 6 (1051.6–1060.1 m). The core sample predominantly consists of fine-grained, pale to dark grey siltstones with intercalations of well-sorted sandstones. Borehole samples were obtained in the Gbely core repository of the NAFTA Ltd. petroleum company. Visual sedimentological interpretation was refined using a CT scan of the cores. Lithofacies were described according to Bouma et al. (1985), Miall (2006), Nielsen et al. (2007), Talling et al. (2012) and Rossi et al. (2017). Basic borehole logs (Fig. 2; spontaneous potential and resistivity) were evaluated according to Rider & Kennedy (2011).

1.3. Foraminifera and calcareous nannoplankton

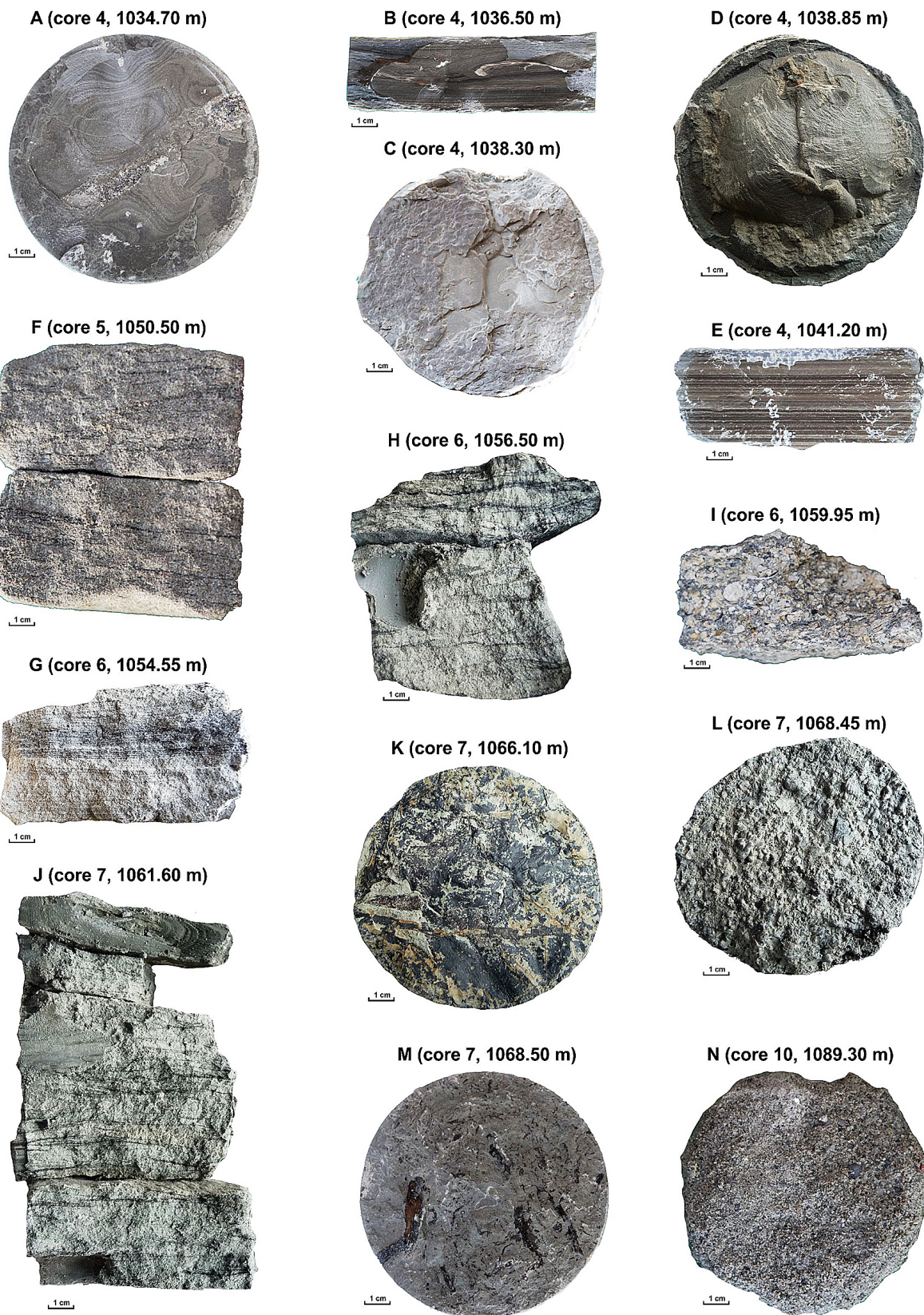
Samples taken from the MZ 102 borehole for Foraminifera and calcareous nannoplankton were performed every 10 cm. Each of the 406 samples were marked in the borehole profile (Fig. 3A–F). The samples were first dried out and separated into 100 g

of borehole core material, diluted by hydrogen peroxide and wet screened on wire screens ranging between 0.071 mm – 1 mm and 1 mm and more; whenever possible, 250 specimens of benthic and planktonic foraminifera were picked out of the residua. Foraminiferal tests were separated by using the binocular stereoscopic microscope Leica MZ and then observed under an electron scanning microscope (SEM) QUANTA FEG 250. The obtained foraminifers were determined into species level based on Loeblich & Tappan (1992), Cicha et al. (1998), and on Luczkowska (1972, 1974), the last two specialized on miliolide foraminifera systematics. Paleoecologic interpretation was obtained from the findings of Boltovskoy & Wright (1976), Jones & Charnock (1985), Murray (1973, 1991, 2006), van der Zwaan et al. (1990), Orpin et al. (1999), Berkeley et al. (2007, 2008), Filipescu (2004) and Patterson et al. (1990). Specimens were grouped into morphogroups according to Langer (1988) and Murray (2006) (Fig. 4) (Tab. 1, Supplement table). Diversity was expressed by



Supplement 1: Photo documentation of the Upper Sarmatian sediments (Core No. 1–3)

A, F: grey, massive to indistinct laminated claystone with abundant molluscs remnants;
 B: grey, slightly laminated claystone with algae remnants; C, D: distinctly laminated claystone with thin carbonate laminae; E: grey, slightly laminated claystone with small plant fragments and coprolites; G, H: distinctly laminated claystone with fish scales and (G) calcareous concretions; I: dark grey claystone with coalified wood fragments; J: grey, massive, bioturbated (sand filled) claystone ; light grey, fine grained laminated , slightly cemented sand; L: grey, massive silty sand.



Supplement 2: Photo documentation of the Upper Badenian and Lower Sarmatian sediments (Core No. 4–10)

A, D: grey, laminated (A) and massive (D), bioturbated (sand filled) claystone, B, E: laminated claystone with (B) diffuse or distinct (E) silty laminae; C: grey, massive claystone with juvenile gastropods; F, G, H, J: light grey, fine grained, horizontally to cross laminated sandstone; organodetritic limestones with fragments of red algae and molluscs; K: grey silty claystone with small sand ripples and coalified plant fragments; L: coarse grained, gravelly sandstone with abundant molluscs and red algae fragments; M: grey claystone with thin silty lenses and laminae and large coalified wood fragments; N: light grey, coarse grained, gravelly sandstone

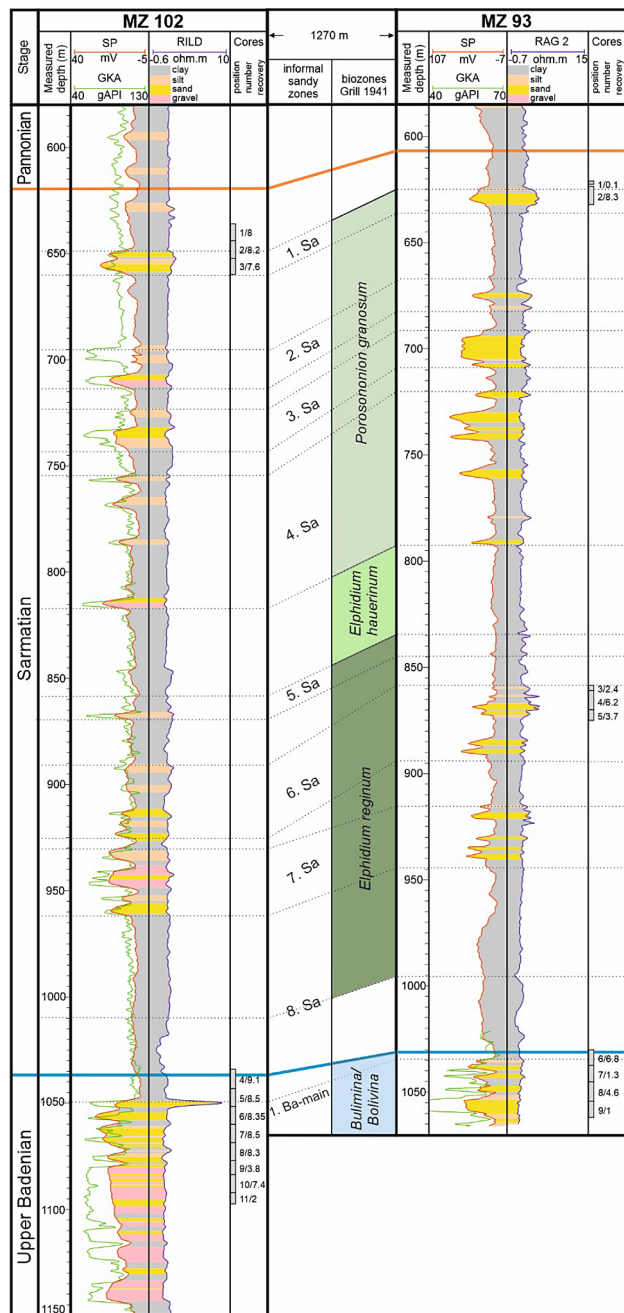


Fig. 2. Lithological scheme of the boreholes MZ 102 and MZ 93 with localization of the studied borehole cores, main sand bodies, and biostratigraphic interpretation.

the Shannon_H index and Fisher alpha formula (Murray, 2006). For specifying the oxygen depletion, the Ammonia – Elphidium index (IAE) was used (Supplement table). The index is given by $IAE = [NA / (NA + NE)] \times 100$, where NA and NE are the numbers of individuals of *Ammonia* and *Elphidium* (Pregmolato et al., 2018). Samples with more than 50 individuals were included in the statistical analysis (Hammer et al., 2001). Data were stored in a Viendat database (edited by Hudáčková & Hudáček, 2001) and analysed with the aid of the programs PAST – Paleontological Statistics, ver. 1.70 (Hammer et al., 2001) and POLPAL (Walanus & Nalepka, 1999). Samples with a predominance of

Tab. 1. Morphogroups of benthic foraminifera adapted according to Langer (1988) and Murray (2006). Epiphytes A, B, C and D, and IN-infauna.

Species	Morphogroup (Langer, 1988) Infauna (Murray, 2006)	Species	Morphogroup (Langer, 1988) Infauna (Murray, 2006)
<i>Adelosina</i> sp.	EP-D	<i>Elphidium hauerinum</i>	IN
<i>Ammonia beccarii</i>	IN	<i>Elphidium macellum</i>	EP-C
<i>Ammonia ex. gr. viennensis</i>	IN	<i>Fissurina laevigata</i>	parasite
<i>Ammonia parkinsoniana</i>	IN	<i>Fissurina mironovi</i>	parasite
<i>Ammonia</i> sp.	IN	<i>Hansenisca soldanii</i>	IN
<i>Ammonia tepida</i>	IN	<i>Haynesina cf. depressula</i>	IN
<i>Articularia articulinoidea</i>	EP-D	<i>Haynesina</i> sp.	EP-C
<i>Articulina multitubulata</i>	EP-D	<i>Heterolepa</i> sp.	EP-B
<i>Articulina problema</i>	EP-D	<i>Hoeglundina elegans</i>	IN
<i>Articulina sarmatica</i>	EP-D	<i>Lobatula lobatula</i>	EP-B
<i>Aubignyna perlucida</i>	IN	<i>Martinottiella</i> sp.	IN
<i>Biasterigerina planorbis</i>	EP-B	<i>Melonis pompolioides</i>	IN
<i>Bolivina cf. dilatata</i>	IN	<i>Miliammina</i> sp.	EP-D
<i>Bolivina dilatata dilatata</i>	IN	<i>Miliolinella</i> sp.	EP-D
<i>Bolivina ex gr. dilatata</i>	IN	<i>Miliolinella subrotunda</i>	EP-D
<i>Bolivina papulata</i>	IN	<i>Nodosaria dina</i>	IN
<i>Bolivina sarmatica</i>	IN	<i>Nodosaria</i> sp.	IN
<i>Buccella frigida</i>	IN	<i>Nonion biporus</i>	IN
<i>Bulimina aff. elongata</i>	IN	<i>Nonion commune</i>	IN
<i>Bulimina elongata</i>	IN	<i>Nonion</i> sp.	IN
<i>Bulimina</i> sp.	IN	<i>Nonionella</i> sp.	IN
<i>Bulimina striata</i>	IN	<i>Porosonion hyalinus</i>	EP-C
<i>Cassidulina laevigata</i>	IN	<i>Porosonion gr. granosum</i>	EP-C
<i>Caucasina gutschulica</i>	IN	<i>Porosonion granosum</i>	EP-C
<i>Cibicides boueanus</i>	EP-B	<i>Pseudotriloculina onsobrina</i>	EP-D
<i>Cibicides dutemplei</i>	EP-B	<i>Pseudotriloculina microdon</i>	EP-D
<i>Cibicides</i> sp.	EP-B	<i>Pullenia bulloides</i>	IN
<i>Cibicides ex.gr. ungerianus</i>	EP-A	<i>Pullenia quinqueloba</i>	IN
<i>Cibroelphidium excavatum</i>	IN	<i>Quinqueloculina akneriana</i>	EP-D
<i>Conorbella erecta</i>	EP-B	<i>Quinqueloculina pseudobuchiana</i>	EP-D
<i>Cycloforina stomata</i>	EP-D	<i>Quinqueloculina selene</i>	EP-D
<i>Dogielina sarmatica</i>	EP-C	<i>Quinqueloculina seminulum</i>	EP-D
<i>Elphidiella arctica</i>	IN	<i>Quinqueloculina</i> sp.	EP-D
<i>Elphidiella cf. minuta</i>	IN	<i>Rosalina austriaca</i>	EP-B
<i>Elphidium aculeatum</i>	EP-C	<i>Rosalina obtusa</i>	EP-B
<i>Elphidium advenum</i>	EP-C	<i>Saccamina sarmatica</i>	IN
<i>Elphidium aff. crispum</i>	EP-C	<i>Siphonaperta lucida</i>	EP-D
<i>Elphidium antonium</i>	IN	<i>Siphonodosaria lepidula</i>	IN
<i>Elphidium cf. fichtellianum</i>	EP-C	<i>Spiroloculina dilatata</i>	EP-D
<i>Elphidium cf. flexuosum</i>	IN	<i>Stilostomella</i> sp.	IN
<i>Elphidium cf. margaritaceum</i>	EP-C	<i>Triloculina</i> sp.	EP-D
<i>Elphidium ex. gr. macellum</i>	EP-C	<i>Valvulineria</i> sp.	IN
<i>Elphidium fichtellianum</i>	EP-C	<i>Varidentella rotunda</i>	EP-D

Fig. 3 A-F. Detailed sedimentology, samples position (circles) and fossil content (symbols) of the studied cores of the borehole MZ 102. Explanation in Fig. 3 F.

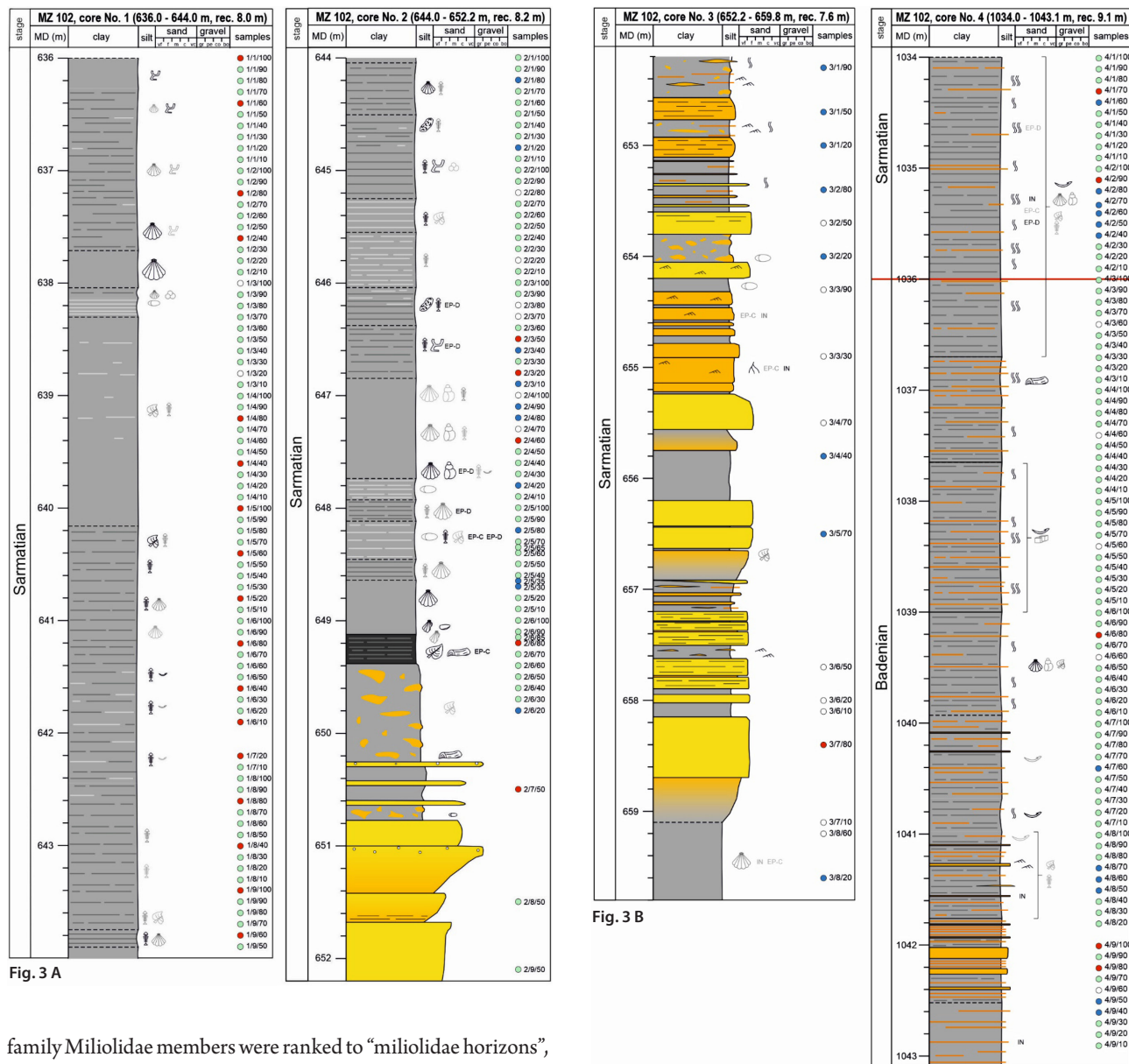


Fig. 3 A

Fig. 3 B

family Miliolidae members were ranked to “miliolidae horizons”, if the amount of miliolids reached more than 15 %.

Calcareous nannofossil samples were prepared by the method of decantation using smear slides. The resulting suspension was mixed with distilled water on a glass slide. The combination created a thick suspension, which after it dried out was ready for observation and compared under the polarized light microscope, Olympus BX 50, at 1250x magnification. Coccoliths were determined into species according to the Nannotax database (Young et al., 2003) and documented by Infinity 2 camera and the QuickPHOTO camera 2.3 program with 10 μ magnification.

2. RESULTS

2.1. Sedimentology

The core samples from the borehole MZ 102 predominantly consists of fine-grained, pale to dark grey siltstones with

intercalations of well-sorted sandstones. The studied core part of the sedimentary record in borehole MZ102 represents sediments of the 1st Badenian - Main sand (cores 11-4, the upper Badenian, and the lower Sarmatian) 1st Sarmatian sand (cores 3-1, terminal Sarmatian) (Fig. 2).

On the well-log curves of the MZ 102 borehole, the Sarmatian sediments begin with a transgressive mudstone (in the northern part of the Malacky field, interrupted by thick 8th Sarmatian sand), which pass locally into the regressive 7th, 6th and 5th Sarmatian sands. The following transgressive – regressive (TR) cycle begins with mudstones overlying the 5th Sarmatian sand, which continue into very distinctive packets of the 4th and 3rd Sarmatian sands (Fig. 2). The sand sedimentation continues with the 2nd Sarmatian sand, (more significantly-developed further south in the Jakubov area) and was terminated by the sands of the 1st Sarmatian sand (with the main depocenter in the Gajary

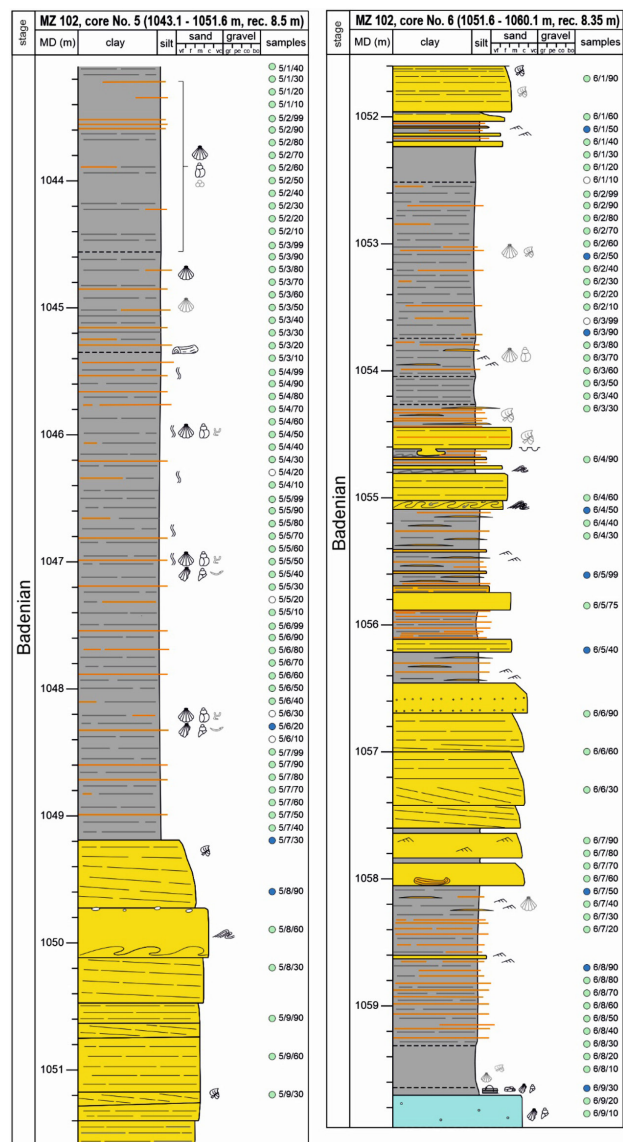


Fig. 3 C

area). Both third order TR cycles are divided into several smaller parasequential cycles.

In cores No. 6-5, the sands are relative well-sorted and, unlike the sediments in cores No. 11-7, they do not contain much gravel or silt. The organodetrritic hardground at the base of core No. 6, which had been formed by small pebbles and fragments of tests and red algae was present (Fig. 2, 3, 4). Mudstones with abundant silty laminae form the upper part of core No. 5 and core No. 4. From a depth of 1039 m upwards, the proportion of pyrite concretions, fish remnants and juvenile shells increases. Core No. 3 and the lower part of core No. 2 are formed by silty claystones, siltstones, and weakly consolidated sandstones. In the upper part of core No. 2, as well as in core No. 1, the above-mentioned sedimentary sequence passes (at a depth of 649.15 m) into overlying fine-grained facies, characterized by massive claystones with abundant mollusc shells alternating with laminated claystones with frequent fish remains. Lamination was often marked by very thin laminae, formed by precipitated CaCO_3 (Fig. 3). Macrofauna was represented mostly by bivalve and gastropod shells, together with fish remains.

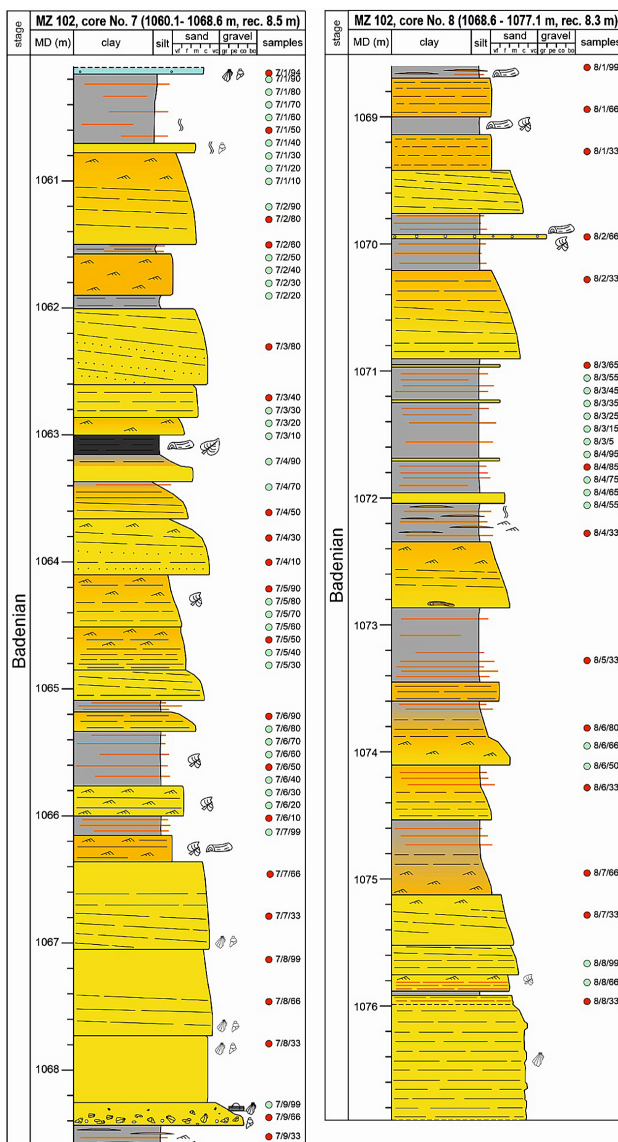


Fig. 3 D

2.2. Calcareous nannoplankton

Calcareous nannofossils were studied in cores No. 6, 4, 3, and 2. Association in the studied samples (30) can be evaluated as low diversified, containing Neogene species and coccoliths reworked from Cretaceous and Paleogene sediments. The samples from depths of 1055.9 m and 1055 m (core No. 6) contain specimens of *Helicosphaera walbersdorfensis*, *Reticulofenestra minuta*, *R. haqii*, *R. pseudumbilicus*, *Coccolithus pelagicus*. Core No. 4 in depth 1042.6 m – 1034.6 m (13 samples) contains association of coccoliths dominated by *Reticulofenestra minuta* species and Cretaceous/Paleogene nannofossils (*Ismolithus recurves*, *Dictyococcites bisectus*, *Discoaster multiradiatus*). The sample from depth 1041.6 m contains abundant associations (*Coronocyclus nitescens*, *Coccolithus miopelagicus*, *C. pelagicus*, *Discoaster exilis*, *D. deflandrei*, *D. variabilis*, *Holodiscus macroporus*, *Helicosphaera carteri*, *H. carteri wallichii*, *H. walbersdorfensis*, *Pontosphaera multipora*, *Reticulofenestra pseudumbilicus*, *R. haqii*, *Sphenolithus heteromorphus*, *Umbilicosphaera jafari*, *U. rotula*). In the sample

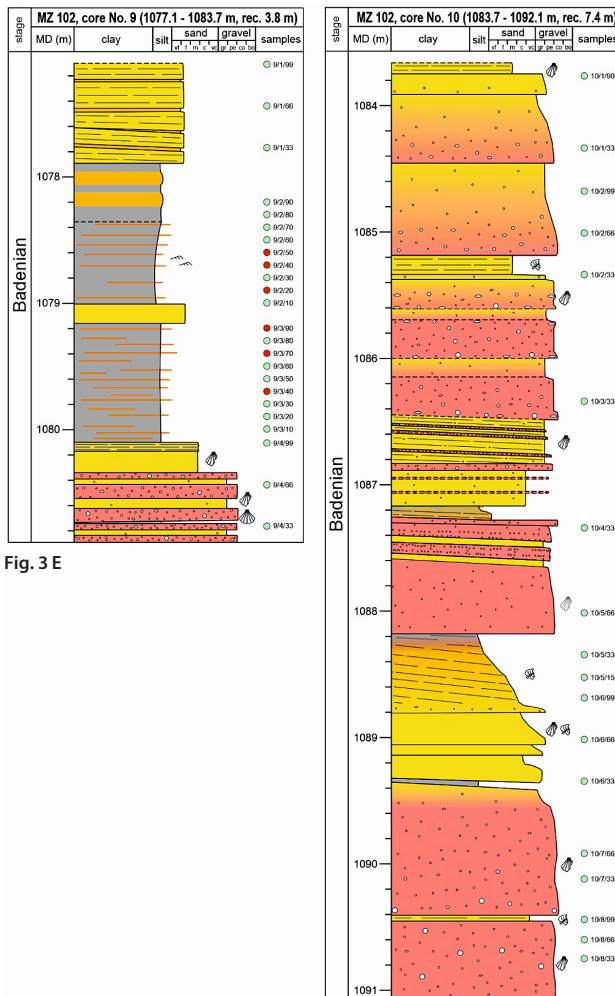


Fig. 3 E

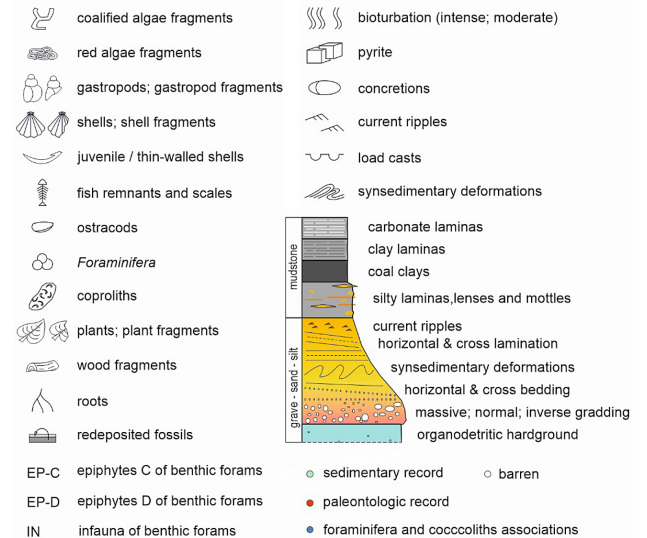
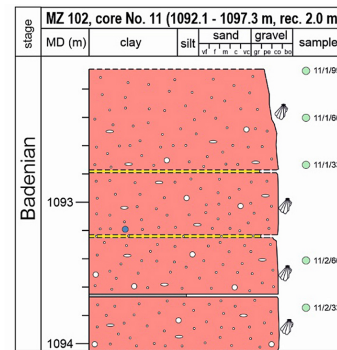


Fig. 3 F

from 1035.8 m, the acme of species *Reticulofenestra minuta* was identified. The core No. 3 (659.8 m – 652.4 m) contains only one sample (depth 655 m) with presence of calcareous nannoplankton with associations of *Braarudosphaera bigelowii parvula*, *Reticulofenestra pseudoumbilicus* and *Sphenolithus heteromorphus* and Cretaceous/Paleogene nannofossils (*Nannoconus steinmannii steinmannii*, *Cyclicargolithus floridanus*). Core No. 2 yields sediments (648.3 and 644.8 m) with the acme of *Sphenolithus abies*, as well as monogeneric associations represented by *Braarudosphaera bigelowii* and *B. bigelowii parvula* together with abundant ascidian spicules (samples in depth 648.8, 648, and 647.2 m).

2.3. Foraminifera

Foraminifera from the MZ 102 borehole were determined into species level (Supplement table) and classified into morphological groups (Tab. 1). Overall, there were 141 systematically processed samples, from which 81 yielded fossil remains, and 39 of them yielded foraminiferal tests. 86 taxa of benthic and 12 taxa of planktonic foraminifera were determined, of which 16 of them stayed in open nomenclature due to poor preservation of the tests (Figs. 5, 6). According to our data, the benthos was more common than plankton in the entire studied part of the





Morphogroup	Features
EP-A 	sesile, attachment surface, flat, concave, or folded over, e. g. <i>Planorbulina</i> , <i>Cibicidoides</i>
EP-B 	temporary motile, apertural faces wide, motion by swimming on pseudopodial network, e. g. <i>Rosalina</i>
EP-C 	motile, pseudopods from canal system and multiple apertural openings, e. g. <i>Elphidium</i>
EP-D 	permanently motile, apertural faces narrowing the bottle-neck apertures, motion by striding on the apertural face in upright position, e. g. <i>Quinqueloculita</i>

Fig. 4. Morphogroups of epiphytes (EP-A, EP-B, EP-C, EP-D) with main behavioural and morphological features and examples from our study (edited by Langer, 1988). For detailed ranking of species into morphogroups, see Tab.2.

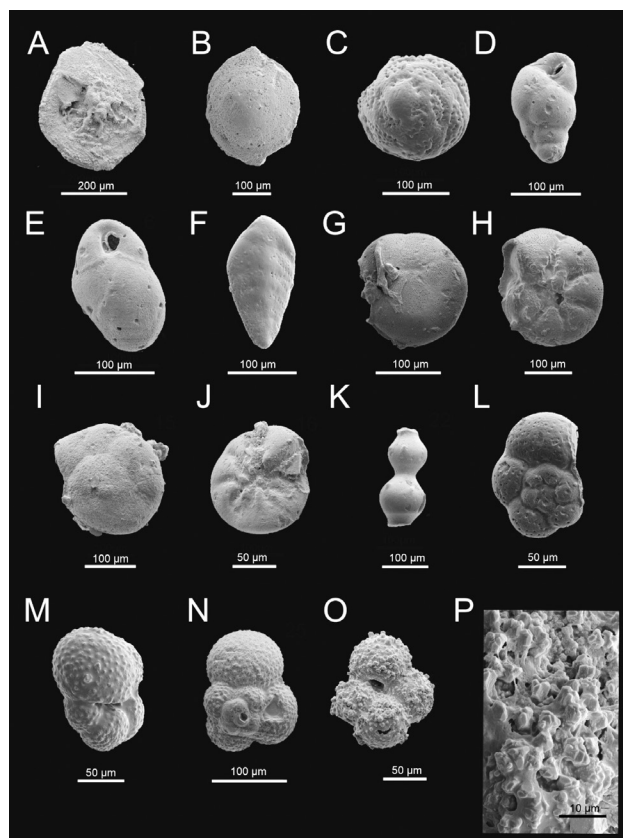


Fig. 5. A, B - *Biasterigerina planorbis* (depth: 1048,8 m); C - *Rosalina* sp. (depth: 1047.8 m); D - *Bulimina elongata* (depth: 1042.9 m); E - *Caucasina gutsulica* (depth: 1041.6 m); F - *Bolivina dilatata* (depth: 1042.9 m); G, H - *Ammonia viennensis* (depth: 655 m); I, J - *Ammonia parkinsoniana* (depth: 659.4 m); K - *Stilostomella* sp. (depth: 1041.6 m); L, M - *Globorotalia* sp. (depth: 1041.6 m); N - *Turborotalita quinqueloba* (depth: 1041.6 m); O - *Tenuitella munda* (depth: 1041.6 m); P - wall detail of *Tenuitella munda* (depth: 1041.6 m).

borehole. Morphogroups were figured by the POLPAL program, in which axis X represents the depth (in meters) and axis Y the percentual proportion of certain foraminiferal morphogroups and other dominated associations (Fig. 7).

2.3.1. Planktonic foraminifera

A low number of small, round planktonic forms (164 individuals), often with signs of size sorting were found in cores 6, 5, and the lower part of core 4 (1059.8 m – 1041.6 m), dominated by association of *Globigerina* sp. div., *Globigerinita uvula*, *Globorotalia* (*Obandyella*) *transsylvanica*, *Trilobatus quadrilobatus*, *Paragloborotalia opima opima*, *Turborotalita quinqueloba*, *Tenuitella* sp. div. indet. and *T. munda*.

2.3.2. Benthic foraminifera

The most common benthic foraminifera obtained from entire borehole profile are species from following genera: *Ammonia* (to 80 % in depth 656.9 m), *Elphidium* (90 % in sample 1041.6 m), *Criboelphidium* (50–90 % in depth interval 1035.8 – 1034.6 m),

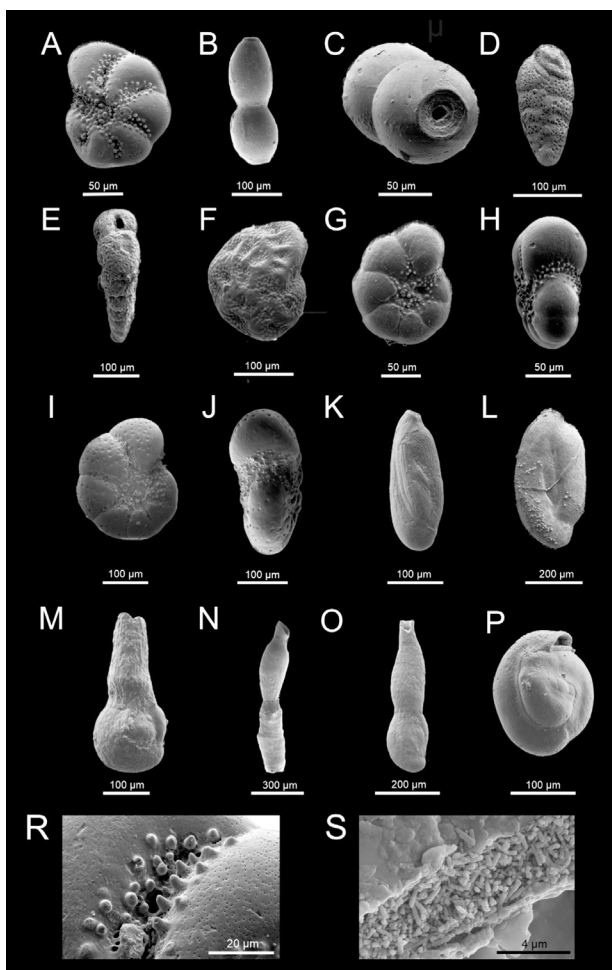


Fig. 6. A - *Criboelphidium excavatum* (depth: 1035.7 m); B - *Nodosaria dina* (depth: 1035.6 m); C - *Articulina sarmatica* (depth: 648); D, E - *Bolivina sarmatica* (depth: 654.5 m); F - *Elphidium margeritaceum* (depth: 648 m); G, H - *Porosonion hyalinus* (depth: 1035.6 m); I, J - *Porosonion granosum* (depth: 649.2 m); K - *Pseudotriloculina consobrina* (depth: 654.5 m); L - *Quinqueloculina seminulum* (depth: 648.3 m); M, N - *Articulina problema* (depth: 648 m); O - *Articulina articulinoidea* (depth: 647.8 m); P - *Miliolinella subrotunda* (depth: 647.8 m); R - wall detail of *Elphidium advenum* (depth: 654.5 m); S - wall detail of *Articulina problema* (depth: 648 m).

Porosonion and *Articulina* (30 % in depth interval 648 – 646.1 m) (Fig. 8). Predominance of miliolids (miliolid horizons, MH; more than 15 %) were found in the depths 1035.6 – 1034.6 m; 648.35 m; 648 – 647.7 m and 646.5 – 646.1 m.

The IAE index value ranged from 0 to 100 with the minima in the interval 648.3 – 646.1 m and with the highest values in 1048.8 – 1041.6 m; 656.9 – 652.4 m and 655 m.

The base of rock sequence (in depth 1059.8 – 1041.6 m) contains few species from morphogroup A/epiphyte (*Cibicides ungerianus* and *Cibicides boueanus*) and morphogroup B/epiphyte – planoconvex shape (*Ammonia viennensis* and *Asterigerinata planorbis*). We observed several flattened epifaunal species with planispiral involute (morphogroup C/epiphyte), such as *Porosonion granosum*, *Haynesina depressula*, *Elphidium macellum* and *Elphidium fichtelianum* (Figs. 7, 8). Occurrence of species *Cassidulina laevigata* and

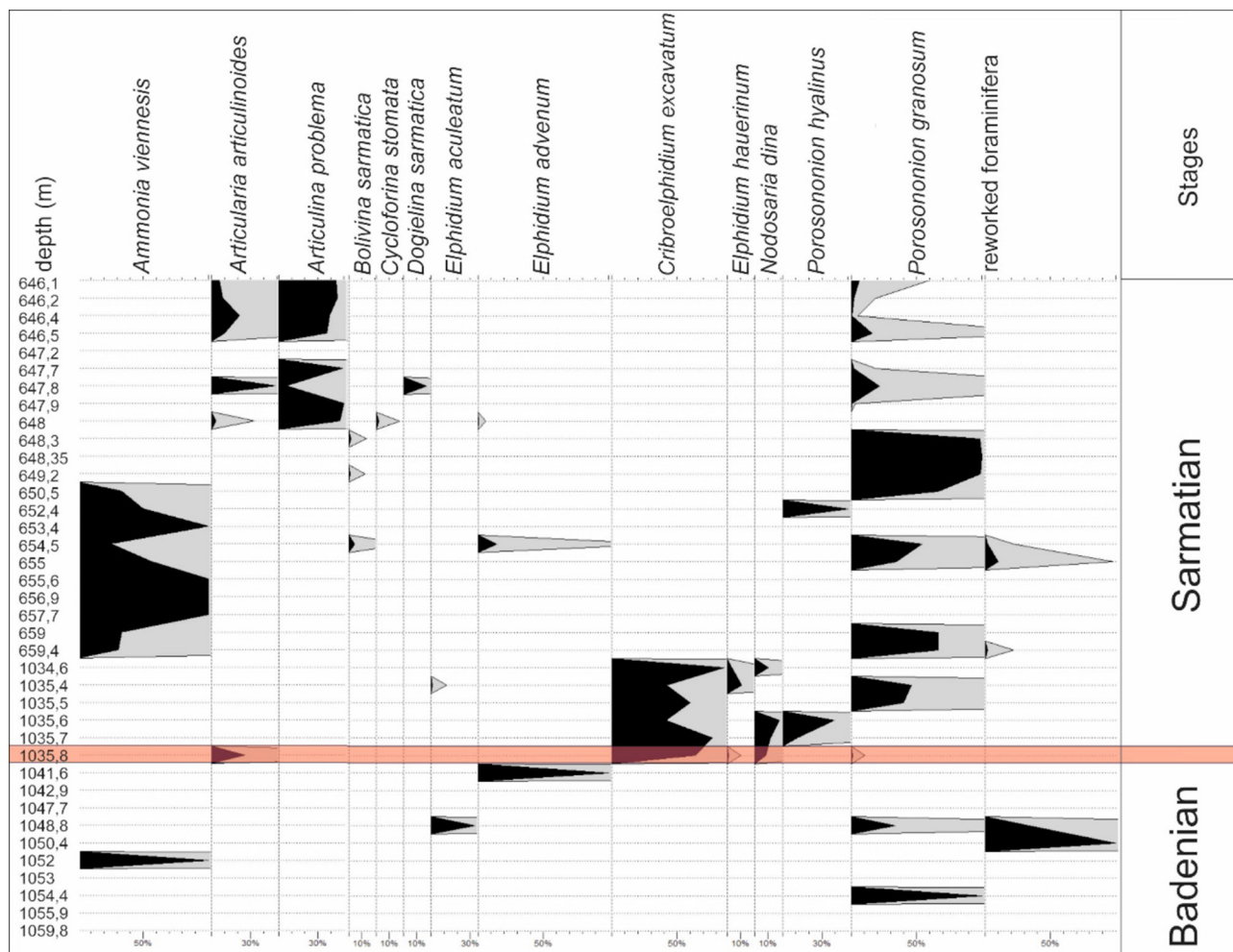


Fig. 7. The dominance of the significant taxa identified in non-barren samples of the MZ 102 borehole. The orange line shows revealed Badenian/Sarmatian boundary.

Globocassidulina crassa with a lenticular shape of the tests was rare. At the depth from 1059.8 to 1041.5 m, the presence of epiphytes was also rare; however, the *Caucasina gutsulica* and *Bulimina elongata* were observed. From the sample taken at 1035.8 m, *Porosonion granosum*, *Haynesina* sp., and *Ammonia* sp. dominate with *Cribroelphidium excavatum* (high abundances) making its first appearance in this part of the borehole. The size sorting of the rounded tests in this sample was observed.

The sample from core No. 3 at a depth of 659.8 m yields poor foraminiferal record of planoconvex trochospiral species (morphogroup B/epiphytes), such as *Biasterigerinata planorbis*, *Ammonia viennensis*, *Ammonia parkinsoniana* and *Aubignyna perlucida*. From a depth of 650.5 m, *Porosonion granosum* highly dominates (morphogroup C/epiphytes) together with miliolids (EP-D morphogroup) *Quinqueloculina seminula*, *Pseudotriloculina consobrina* and *Quinqueloculina rotunda*. From the sample taken at 648 m, the miliolidae foraminifera *Articulina problema*, *Articularia articulinooides* begin to dominate, accompanied by a few species of *Articulina sarmatica*, *Cycloforina stomata*, and *Dogielina sarmatica*. The sediments with this associations end at a depth of 646.1 m, above which the sediments are barren of foraminifera tests.

Infauunal associations dominated at depths between 1053 to 1035.7 m and 659.4 to 649.2 m. Epiphytes of group A and B were dominant in the lower parts of the borehole core; however, epiphytes of group C and D were abundant in the upper part.

2.4. Diversity

Diversity results show values on the Fisher alpha formula with the minimum being 1.3 and the maximum being 5.97, and on the Shannon_H formula with values being the minimum 0.77 and the maximum 2.41. Values were higher in the lower part of borehole MZ 102, yet lower in the upper part. In general, the relationship between the values of estimated diversities of the samples are presented in three groups (Fig. 9). The first group (A) specified the values of the Shannon_H index from 0.77 – 1.66 with the Fisher alpha index from 1.3 to 1.85 in the samples from depths of 646.2; 647.7; 647.9; 648; 648.3; 648.35; 649.2; 1034.6, and 1035.4 meters. These samples contain rich association of miliolids, such as *Articulina problema*, *Varidentella rotunda*, *Pseudotriloculina consobrina*, *Siphonaperta lucida*, *Quinqueloculina seminulum*, and numerous hyaline species, such as *Porosonion granosum* and *Cribroelphidium excavatum*. The second group

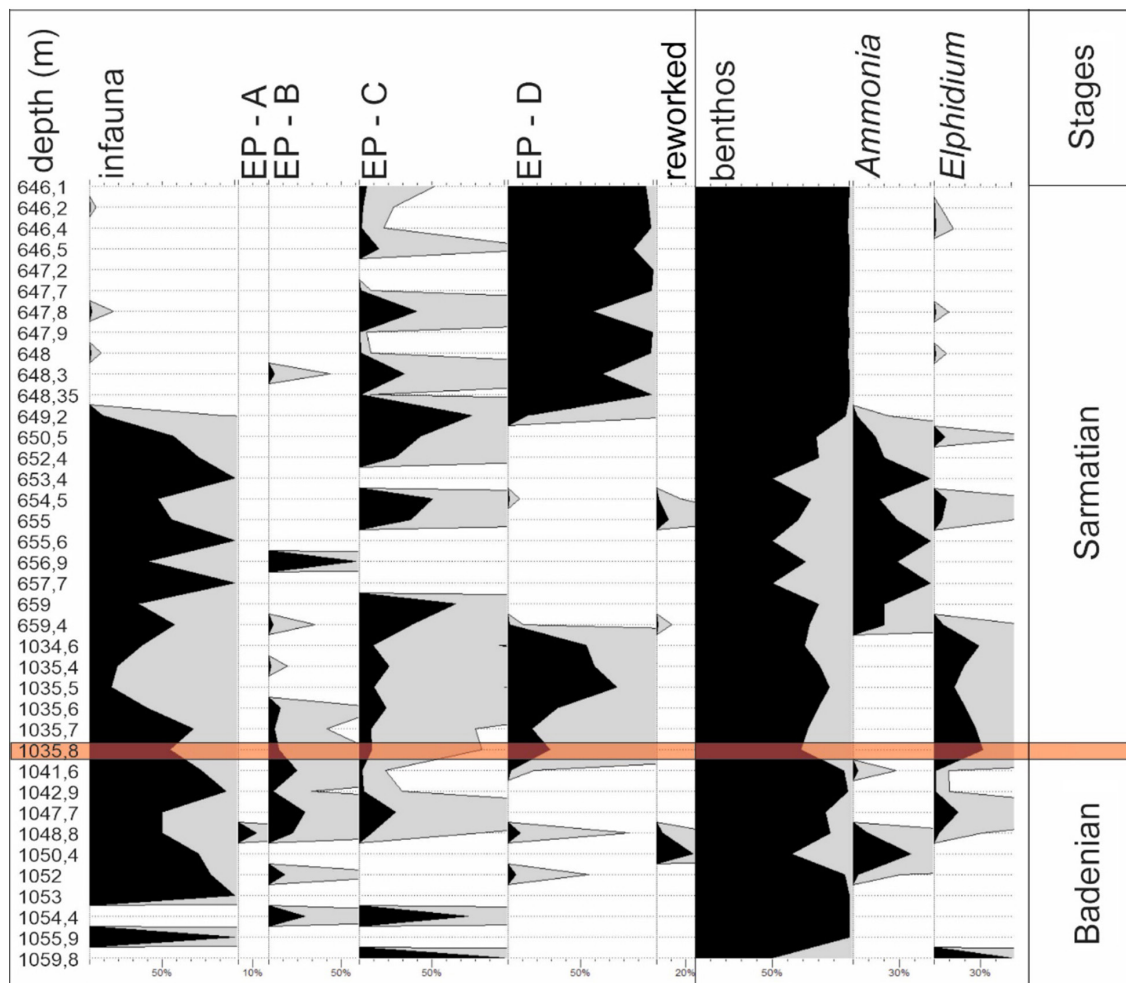


Fig. 8. Morphogroups abundance divided according to Langer (1988), infaunal abundance according to Murray (2006), redeposits, benthos, and Ammonia-Elphidium abundance.

(B) specified associations with the values of the Shannon_H index from 1.76 to 2 and the Fisher alpha index from 3.17 to 2.77. Group B contains samples from the depths of 646.1; 646.4; 646.5; 647.8; 655; 1035.6; 1035.7; 1035.8; 1041.6, and 1042.9 meters, with the possibility of defining a subgroup B-a consisting of samples 655, 1035.6, 1041.6 and 1042.9 meters. It contains rich association of miliolids, such as *Articulina problema*, *Varidentella rotunda*, *Articularia articulinoidea*, *Milionella subrotunda* and *Quinqueloculina* sp. and hyaline species *Ammonia viennesis*, *Porosonion granosum*. In cores 4 and 5 (subgroup B-a), the species *Criboelphidium excavatum*, *Nonion* sp., *Porosonion hyalinus*, and *Bolivina dilatata* dominate, as well as the miliolide *Varidentella rotunda* (Fig. 9). The third group (C) specified values of the Shannon H index from 2.22 – 2.41 with the Fisher alpha index from 5.14 to 5.97 in two samples (654.5 and 659.4 m) with *Porosonion* and *Elphidium* genera.

3. INTERPRETATION AND DISCUSSION

Part of the MZ 102 borehole cores (74 samples) were studied based on sedimentology, as well as foraminiferal associations and species supplemented by a calcareous nannoplankton study.

3.1. Biostratigraphy

Biostratigraphy was solved with the help of planktonic and benthic foraminifers and calcareous nannoplankton. The sediments from cores No. 6 – 4 (1059.8 m to 1041.6 m) of the MZ-102 borehole containing association of planktonic foraminiferal species were associated with the uppermost Badenian. Badenian age of the mentioned sediments was proved by presence of *Globorotalia (Obandyella) transsylvanica* and *Globigerinoides quadrilobatus*. Based on the common occurrence of *Tenuitella* sp. div. and small sized *Globigerina* sp. div. in accordance with the assemblage mentioned in Filipescu & Silye (2008), this interval was correlated with upper Badenian acme - zone *Tenuitellinata* (Filipescu & Silye, 2008). The calcareous nannoplankton index species *Discoaster exilis* confirms the NN6 Zone from the depth of 1041.6 m. We interpret the sediments as being upper Badenian, as well as having the age equivalent of *Bulimina/Bolivina* zone or younger due to signs of tests reworking.

The first occurrence of the Sarmatian index species such as *Nodosaria dina*, *Elphidium hauerinum*, and *Articularia articulinoidea* (Cicha et al., 1975; Luczkowska, 1972) was documented in a sample from the upper part of core No. 4 (1035.8 m); thus, the Badenian/Sarmatian boundary was determined in this part

of the MZ 102 borehole. However, index taxa of the lowermost Sarmatian biozones as the *Anomalinoidea dividens* and *Elphidium reginum* (Grill, 1941, 1943; Luczkowska, 1972; Cicha et al., 1975) were not recorded within the studied samples, and the middle Sarmatian index species *Elphidium hauerinum* was rare. It is important to note, the benthic foraminiferal biozones strongly depend on the type of environment [mentioned mainly in the Sarmatian sediments: Harzhauser & Piller (2004); Koubová & Hudáčková (2010); Zlinská et al. (2010)] and one can speculate, that due to unsuitable conditions they can only be developed in certain parts of the basin with special conditions. We consider these associations may occur in the sediments of the non-cored part of the MZ 102 borehole (1034.6 – 659.8 m). In the nearby MZ 93 borehole (Fig. 2), the lower Sarmatian *Elphidium reginum* Zone was observed in the cores 5, 4 and 3 of the circa 860 – 875 m of depth (Koubová & Hudáčková 2010), and we correlate it with the non-cored part of the MZ 102 borehole (Fig. 2). In the samples from cores No. 3 and 2 (659.8 – 644.2 m) of the MZ 102, the associations with predominance of *Porosonion granosum* were assigned to the upper Sarmatian, *Porosonion granosum* Zone (Grill, 1943) of the Skalica Fm (Vass, 2002). The Sarmatian part of the NN6 Zone, based on *Sphenolithus abies acme*-Zone, was documented in core No. 2 in the depths of 648.3 and 644.8 m.

3.2. Sedimentology

The Upper Badenian sediments in this region have a clearly visible progradational to aggradational pattern of seismic reflexes, while the Sarmatian sediments do not have any significantly developed seismic clinofolds (Kováč et al. 2008a). Sedimentological interpretation of cores No. 6 – 5 revealed that the deltaic sands of the Upper Badenian are developed. The character of

the sorting, as well as the upward fining trend of sediments, indicates a wave, transgressive reworking (Fig. 2). Transgression was documented by a higher content of reworked foraminiferal taxa (Figs. 7, 8). Size sorting during postmortal transport in the latest Badenian was documented in the MZ 93 borehole (depth of ~1060 m) by Koubová & Hudáčková (2010) as well.

From the upper part of core No. 4 (MZ 102) upwards, the fine silty clay with an increasing proportion of pyrite concretions and fish remnants indicates a deepening of the sedimentary environment (Kováč et al. 2008a,b). Increase of the infaunal foraminifera (sample 1035.7 m, Fig. 7), together with rich framboidal pyrite, can prove the low oxygen bottom zone (Jorissen et al, 1995).

The sedimentary analysis of cores No. 3 and 2 indicates sedimentation in the distal part of deltaic lobes and poorly-aerated lagoons and marshes (Figs. 2, 3A, B). In the upper part of core No. 2 and in the core No. 1, the lamination was often marked by very thin lamines formed by precipitated CaCO_3 (Fig. 3A), which could represent the so-called non-glacial varves, sensu Galović & Bajraktarević (2006).

3.3. Paleoenvironment

Paleoenvironmental analysis combine the results from calcareous nannoplankton, foraminifera, and sedimentology. Species-based paleoenvironmental interpretation corresponds with morphogroup distribution (Fig. 4, Tab. 1). In the lowermost part of the borehole (interval from 1059.8 to 1041.6 m), the low-oxygen environment was documented by the absence of the oxyphilic species and the occurrence of the species *Cassidulina laevigata* and *Globocassidulina crassa*, *Caucasina gutsulica*, and *Bulimina elongata* (Fig. 5 D, E), which can tolerate stress conditions from reduced oxygen (Bernhard, 1986; Kaiho, 1994). In this interval, we can also observe several planktonic species of foraminifera

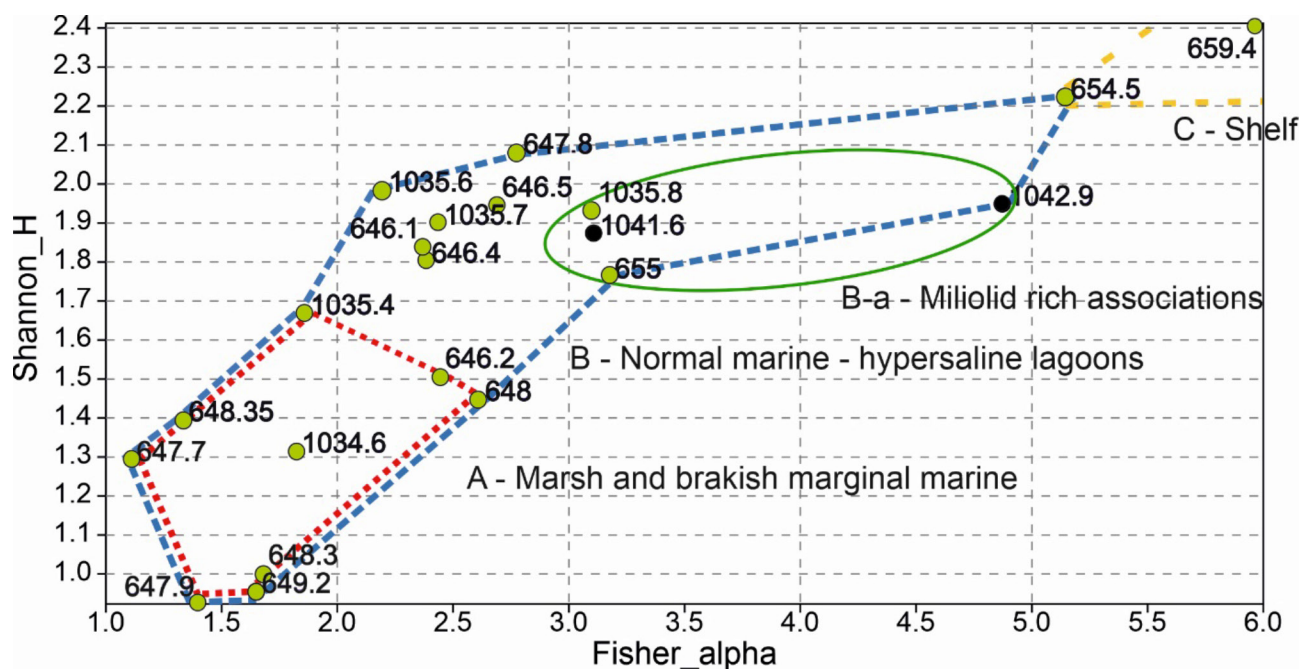


Fig. 9. Space-diversity data (Fisher alpha and Shannon H index) according to Murray (2006). Red dash line: group (A) – marsh and brackish marginal marine, blue dash line: group (B) – normal marine/hypersaline lagoons with green subgroup B-a – miliolid rich associations, yellow dash line: group (C) – shelf.

(*Globigerina* sp., *Globorotalia* sp., *Turborotalita quinqueloba*) together with coccoliths documenting open marine conditions with the possibility of stress conditions in the eutrophic water surface (van der Zwaan et al., 1990, Ćorić & Hohenegger, 2008). The presence of the infaunal species *Bulimina-Bolivina* (cone-shaped and elongated shaped morphogroups), together with members of *Nodosaria* and *Stilostomella* genera (cylindrical uniserial morphogroup), document a deeper shellfal environment with lowered oxygen in the water/sediment interface (Kaiho, 1994; Jorissen et al., 1995; Murray, 2006) and low CaCO₃ content caused by high input of organic material (Bernhard, 1986; Kaiho 1994). This is in accordance with the interpretation of low oxygen levels in upper Badenian sediments in the *Vienna Basin* because of changes of water in the current regime from an anti-estuarine to open marine (Báldi, 2006; Kováčová & Hudáčková, 2009; Kováč et al., 2017b). According to the species-diversity analysis of the benthic foraminifera samples, assemblages from the above-mentioned interval represent a normal marine environment (Fig. 9).

The stressed environment was documented by the prevalence of calcareous nannoplankton species *Reticulofenestra minuta* (depth 1035.8 m) similar to what was mentioned in borehole MZ 93 (Koubová & Hudáčková, 2010). In addition to the species considered to be index fossils for Sarmatian (*Nodosaria dina*, *Articularia articularioides*, 1035.8 m), the size sorting of the rounded tests indicates reworking of Badenian planktonic forms in these horizons. A different type of foraminiferal associations was found in the interval of 1035.8 – 1034.6 m, dominated by *Porosonion granosum*, *Haynesina* sp., and *Ammonia* sp. species and the high abundance of *Criboelphidium excavatum*. The pronounced dominance of epiphytic groups (morphogroups C, D/epiphytes) can document a distinct euphotic zone in interval from 1035.6 to 659 m proved by space-diversity analysis (Murray, 2006) documented in the marginal marine environment (Fig. 7, 9). Epiphytic association in coeval sediments in the Vienna Basin was documented in various localities of Holíč and the Skalica fms. (Fordinál & Zlinská, 1998; Koubová & Hudáčková, 2010; Zlinská et al. 2010, Hudáčková et al. 2021). Almost 400 meters of rock sequence was not cored. However, we can assume a continuing trend of shallow-water sedimentation based on geophysical and sedimentological (CT_P) analysis of the borehole MZ 102 (Sp; RILD curves; Fig. 2) and assemblages from MZ 93 cored part.

The rock record continues with core No. 3 at a depth of 659.8 m, which yields poor foraminiferal fossil record of planiconvex trochospiral-shaped species (morphogroup B/epiphytes), such as *Biasterigerinata planorbis*, *Ammonia viennensis*, *Ammonia parkinsoniana* and *Aubignyna perlucida*. This test morphology was advantageous for maintaining stability on the surface of the substrate (Corliss & Emerson, 1990). The forementioned foraminifera show wide salinity tolerance (Murray, 2006), which may indicate fluctuations in freshwater inflow from the mainland that can support the assumption of the marginal high-energy environment. The planispiral morphology forms, such as *Porosonion granosum*, *Haynesina depressula*, and *Elphidium macellum* are present here and can document enough nutrient supply. In the interval from 656.9 to 652.4, a higher abundance

of *Ammonia* spp. was noticed (with 3 peaks 657.7; 655.6; 653.4 m), which corresponds to the highest values (> 33.3) of the IAE index (Pregolato et al., 2018). We can observe associations dominated by *Ammonia tepida* and *Criboelphidium excavatum* affected by the reduction of oxygen content, very similar to this in Burone & Pires-Vanin (2006) and Burone et al. (2006).

The sedimentary analysis of the borehole core No. 2 (644 – 652.2 m) indicates sedimentation in the distal part of the deltaic lobes, as well as poorly-aerated lagoons and marshes. One important foraminiferal assemblage change was identified from the depth between 650.5 m and 648.3 m, where *Porosonion granosum* highly dominates (morphogroup C/epiphytes, Fig. 8). This, together with miliolids, which is typical in hypersaline conditions (Boltovskoy & Wright, 1976; Murray, 2006), may document strong short-term salinity fluctuation during sedimentation. From 648 m, miliolid foraminifera dominate (*Articulina*, *Articularia*, *Cycloforina*, *Dogielina*) in the samples, what according to Alperin et al., (2011), implies sand sediment of shallow gulf areas with the temperate, hypersaline, and productive conditions. Depending on the spatial diversity distribution (Fig. 9), we interpret them as association from the marginal seas of hypersaline lagoons (Murray, 2006). In Devínska Kobyla (Hyžný et al., 2012; Pivko et al. 2017), shallow water hypersaline foraminiferal association was also observed. Small miliolids were observed only in the topmost sample from the MZ 93 borehole. Based on the correlation of biostratigraphy and sedimentology in the studied area, we can interpret an alternation of salty fens or marshes inhabited by miliolids of the EP-D morphogroup, as well as brackish fens inhabited by the EP-C morphogroup dominated with *Porosonion granosum*. Calcareous nannoplankton associations from depth interval (from 649.2 to 647.2 m) are exclusively represented by *Braarudosphaera bigelowii*, including *B. bigelowii parvula* and abundant ascidian spicules typical for lowered salinity (Stradner & Fuchs, 1979). Similar assemblages were observed in Sarmatian sediments of the Vienna and Danube basins in a marginal environment, with sea level fluctuations and terrestrial draw (Kováč et al., 2005; Kováč et al., 2008a). The sediments with shallow water hyposaline associations end at a depth of 646.1 m, above which sediments are barren of foraminifera tests.

4. CONCLUSIONS

Paleoecological and sedimentological changes of the Vienna Basin sediments were studied in the MZ 102 borehole and correlated with borehole MZ 93, both located near the town of Malacky in the western (Slovak) part of the Vienna Basin.

The Badenian sediments in the MZ 102 borehole are documented based on the foraminifera from the depth of 1059.8 to 1041.6 meters (uppermost Badenian – equivalent of *Bulimina/Bolivina* Zone). The calcareous nannoplankton confirms the NN6 Zone from the depth of 1041.6 m. The first occurrence of the Sarmatian index species was documented in a depth of 1035.8 m thus we interpret it as the Badenian/Sarmatian boundary. From the depth 646.1 to ~ 620 m, the Sarmatian sediments (*Porosonion granosum* Zone) were documented based on

foraminifera and calcareous nannofossils. Interpretation fits well with the results obtained from the M 93 borehole, in which lower Sarmatian was cored. Badenian sediments originated in the deltaic condition with possible reworking with episodes of deeper open marine waters with lowered oxygen on the sea floor. Major paleoecological change has been identified in samples from the depths of 1038.8 m – 1034.6 m, where the sea floor conditions changed from normal marine to low-oxic.

The Sarmatian sediments studied here generally indicate sedimentation in the distal part of the deltaic lobes, as well as poorly-aerated lagoons and marshes that can explain the variability in proportion of the foraminiferal morphogroups. The shallow water environments with significant changes in salt and oxygen content at the bottom with facultative sea grass cover were found in the studied sediments. Major paleoecological changes have been identified in depths from 659.4 m to 650.5 m, where the lowered oxygen level in the water/sediment interface was identified with the three most pronounced excursions of very low level of oxygen in the interval between 656.9 m to 652.4 m. The predominance of miliolids (family Miliolidea) indicates the shallow water, hypersaline, as well as an aerated lagoonal environment from the depth of 648.3 m up to the 641.1 m.

Comparison of the two methodological approaches to the paleoenvironmental analyses – (1) species analysis or (2) morphogroups analysis shows that the morphogroup approach seems to be a quick and sufficiently effective method for paleoenvironmental reconstructions.

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