

Maastrichtian island in the central European Basin—new data inferred from palynofacies analysis and inoceramid stratigraphy

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Abstract The palynological study and palynofacies analysis supported by size analysis of opaque phytoclasts and diversity indexes for particulate organic matter in stratigraphically well-constrained (inoceramid biostratigraphy) sections is applied in monotonous Upper Cretaceous carbonate-siliciclastic sediments of southern Poland. Integrated data allow for the estimation of the proximity of an uplifted area from the studied sections. For the first time in palynofacies analysis, both Simpson's Index of Diversity ($1 - D$) and Shannon's index are used for organic matter distribution providing a comprehensive understanding of a proximal–distal trend in a sedimentary basin. Cluster analysis allowed grouping the samples within proximal to distal shelf zones. The size analysis of opaque woody phytoclasts is used to extrapolate the distance from the source area. Palynofacies analysis indicates the existence of an uplifted area (probably connected with “Kukernitz Island”) in the Holy Cross part of the Danish-Polish Trough during the Maastrichtian. The presented results improve contemporary paleogeographical interpretations for this part of the Late Cretaceous central European Basin. The youngest Maastrichtian deposits of the

Miechów Synclinorium are described, based on inoceramids which are of early Late Maastrichtian age.

Keywords Palynofacies · Inoceramid stratigraphy · Paleogeography · Late Cretaceous · Central European Basin

Introduction

The Maastrichtian deposits of the present-day Miechów Synclinorium territory were accumulated in a shallow epicontinental sea. The monotonous succession of sandy chalk was deposited in a calm low-energy environment with a moderate rate of sedimentation. Although carbonate sedimentation prevailed, a high admixture of quartz grains (Rutkowski 1965; Jurkowska 2016) and the occurrence of well-preserved plant fossils (Cieslinski and Milacovic 1962; Halamski 2013) indicate the presence of an uplifted area in close proximity. Rutkowski (1960) indicated the WWS–EEN direction of terrigenous grains transport based on size measurement of quartz grains in Maastrichtian deposit in the SW part of the Miechów Synclinorium. Świdrowska et al. (2008) pointed out that during the Maastrichtian, terrigenous material was transported to the deeper basin on the SW slope of the Holy Cross segment from the SW without reaching the SE part of the Miechów Synclinorium. Although both studies present the possible direction of terrigenous grain transport, they do not indicate the source area of the material and focus of the whole Maastrichtian stage, without detailed stratigraphy available.

The presence of an uplifted area called “Kukernitz Island” in Ukraine (Pasternak et al. 1987) near the Holy Cross Mountains part of the Danish-Polish Trough during the Turonian–Campanian is well documented, the existence

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of the island during the Campanian–Maastrichtian is mentioned in the literature (Gierliński 2015; Bojanowski et al. 2017), but has not been explored in detail. Understanding the development of the Miechów Synclinorium is critical for the interpretation of the tectonic evolution of extra-Carpathian Poland and important for deciphering the final stages (Campanian or Maastrichtian) of the inversion history of the Danish-Polish Trough (Marcinowski 1974; Walaszczyk 1992; Kutek 1996; Krzywiec 2006).

Initially, the palynofacies analysis principles of Tyson (1993) as well as others revised by Roncaglia and Kuijpers (2006) and Zobaa et al. (2015) have been applied by us to recognize the sedimentary environment of the studied area. However, the unusual distribution pattern of marine phytoplankton decreasing offshore in the samples analyzed revealed that these models are inadequate for the presented case study. The paper presents the application of an integrated palynological study and palynofacies analysis supported by size analysis of opaque phytoclasts and diversity indexes for particulate organic matter in stratigraphically well-constrained (inoceramid biostratigraphy) sections. This method seems to be valuable in determining the distance from land in monotonous carbonate-siliciclastic sediments devoid of unequivocal sedimentological and paleontological data determining depositional environments.

Geological setting

The Miechów Synclinorium is positioned in the southeastern part of the Szczecin-Miechów Synclinorium (Fig. 1a), adjacent to the Mid-Polish Anticlinorium interpreted as an inverted part of the former Danish-Polish Trough (e.g., Pożaryski et al. 1979). The Cretaceous Miechów Synclinorium deposits, which are of Albian to Maastrichtian age (Rutkowski 1965; Marcinowski 1974; Walaszczyk 1992; Jurkowska 2016), overlay the Jurassic substrate unconformably, and are covered in their central and southern parts by Miocene deposits of the Carpathian Foredeep (Pożaryski 1977). The studied sections (Fig. 1b) present an almost complete succession ranging in age from the Lower Maastrichtian to the lower Upper Maastrichtian, and are represented by marly chalk in the lower part and sandy chalk with marly intercalations in the upper part (Rutkowski 1965; Jurkowska 2016). An increasing input of terrigenous quartz continuing within the succession has been noted by Jurkowska (2016). Under an optical microscope, the studied rocks represent packstone with sponge spicules and foraminifera with a significant admixture of terrigenous quartz (Rutkowski 1960; Jurkowska 2016).

The existence of an uplifted area of the Danish-Polish Trough (present Mid-Polish Anticlinorium) called “Kukernitz Island” (Pasternak 1959; Pasternak et al. 1987) or

Łysogóry–Dobruża Land (Samsonowicz 1925; Pożaryski 1962) during the Turonian–Santonian is well documented and recently confirmed by micropaleontological (Dubicka et al. 2014) and ammonites data (Remin and Marshall 2016). Its formation was an effect of Subhercynian tectonic movements. The island probably expanded during the Santonian and Middle and Late Campanian (Świdrowska et al. 2008; Gierliński 2015; Walaszczyk and Remin 2015; Remin and Marshall 2016; Bojanowski et al. 2017). The uplifted areas in the Holy Cross part of the Danish-Polish Trough were an alimentation area for the Miechów Synclinorium, where increasing siliciclastic input was noted from the Late Campanian onwards (Jaskowiak-Schoeneichowa and Krassowska 1988; Jurkowska 2016).

Localities and inoceramid succession

The Maastrichtian succession of the Miechów Synclinorium is accessible through a series of natural and artificial (abandoned quarries) exposures. One of the authors (Jurkowska 2016) described the inoceramid stratigraphy and provided the coordinates of the most thoroughly studied sections. In this paper, new exposures and new stratigraphic data are also presented. The studied succession belong to the *Endocostea typica* and *Trochoceras radius* zones of the Lower and Upper Maastrichtian.

Lower Maastrichtian (Fig. 2)

Endocostea typica Zone

The lowermost part of the *E. typica* Zone is represented by a glauconitic horizon in the SW part of the Miechów Synclinorium (Jurkowska 2016). In the NW part of the studied area, it is represented by marly fossiliferous chalk in the Pelczyska section (Fig. 1b) with many *Endocostea typica* (Whitfield 1877), and small (<3 cm) *Cataceramus subcircularis* (Meek 1876) specimens. The Dziewięcioły, Brynica Mokra, Pińczów, and Kozubów localities (Fig. 1b) were described by Jurkowska (2016) and the rocks are composed of white sandy chalk of the upper part of the *E. typica* Zone with numerous small (3–5 cm) representatives of *C. subcircularis*, *Cataceramus barabini* (Morton 1834), and *Cataceramus glendivensis* (Walaszczyk et al. 2001).

Trochoceras radius Zone

New findings of *Trochoceras radius* specimens from Wola Chrobberska, Strzeżów 2, and Rzędowice (Jurkowska 2016) (Fig. 1b) confirmed that these sections represent the lower part of the *T. radius* Zone. In the Jelcza Wielka (Peryt and Dubicka 2009; Jurkowska 2016) and Michałów

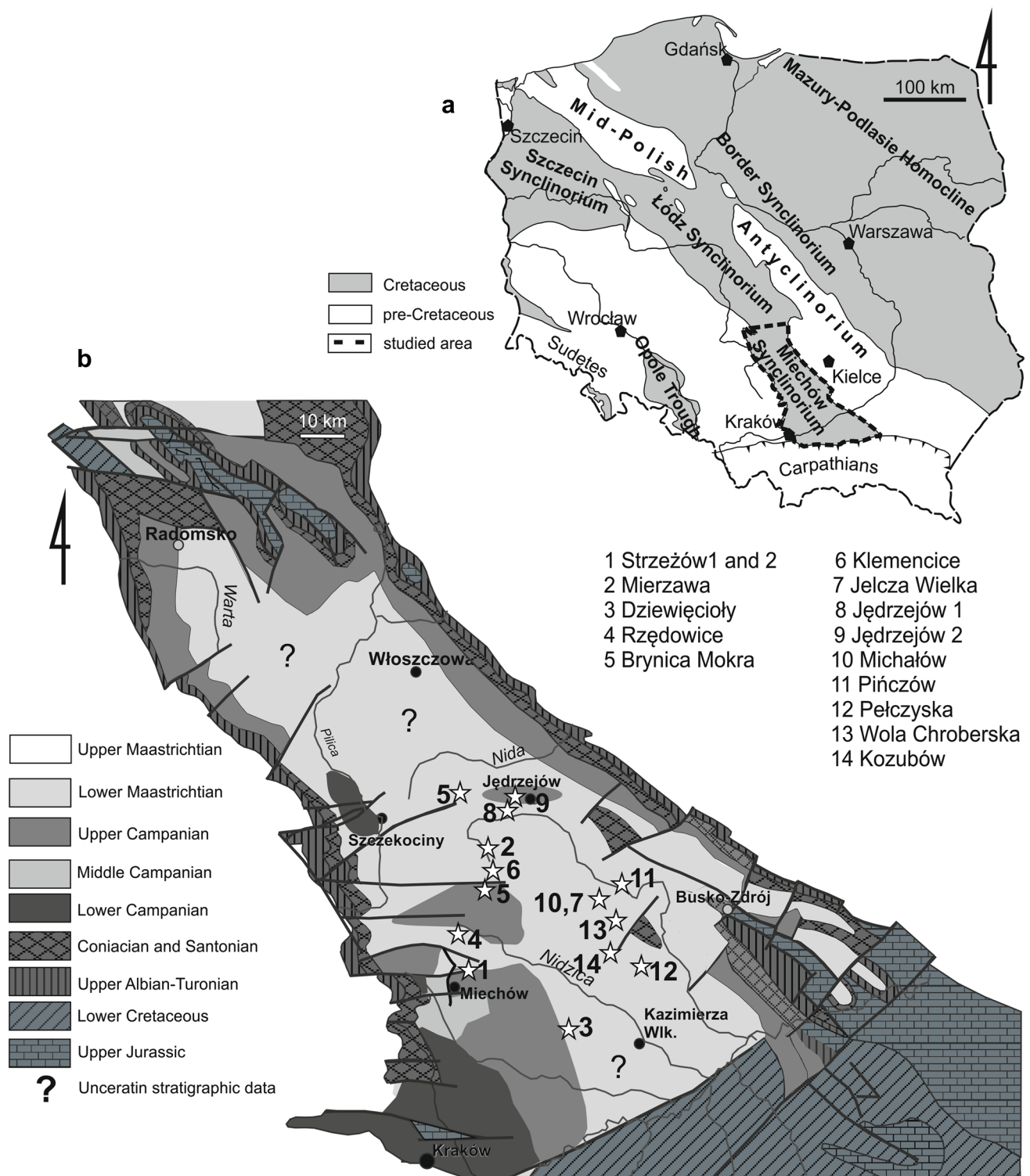


Fig. 1 **a** Tectonic-sketch map of Poland (without Cenozoic cover) (after Pożaryski 1974; changed after Zelazniewicz 2008 and Zelazniewicz et al. 2011; Jurkowska 2016). **b** Geological map of

the Mięchów Synclinorium (modified after Dadlez et al. 2000 and Jurkowska 2016) with studied localities

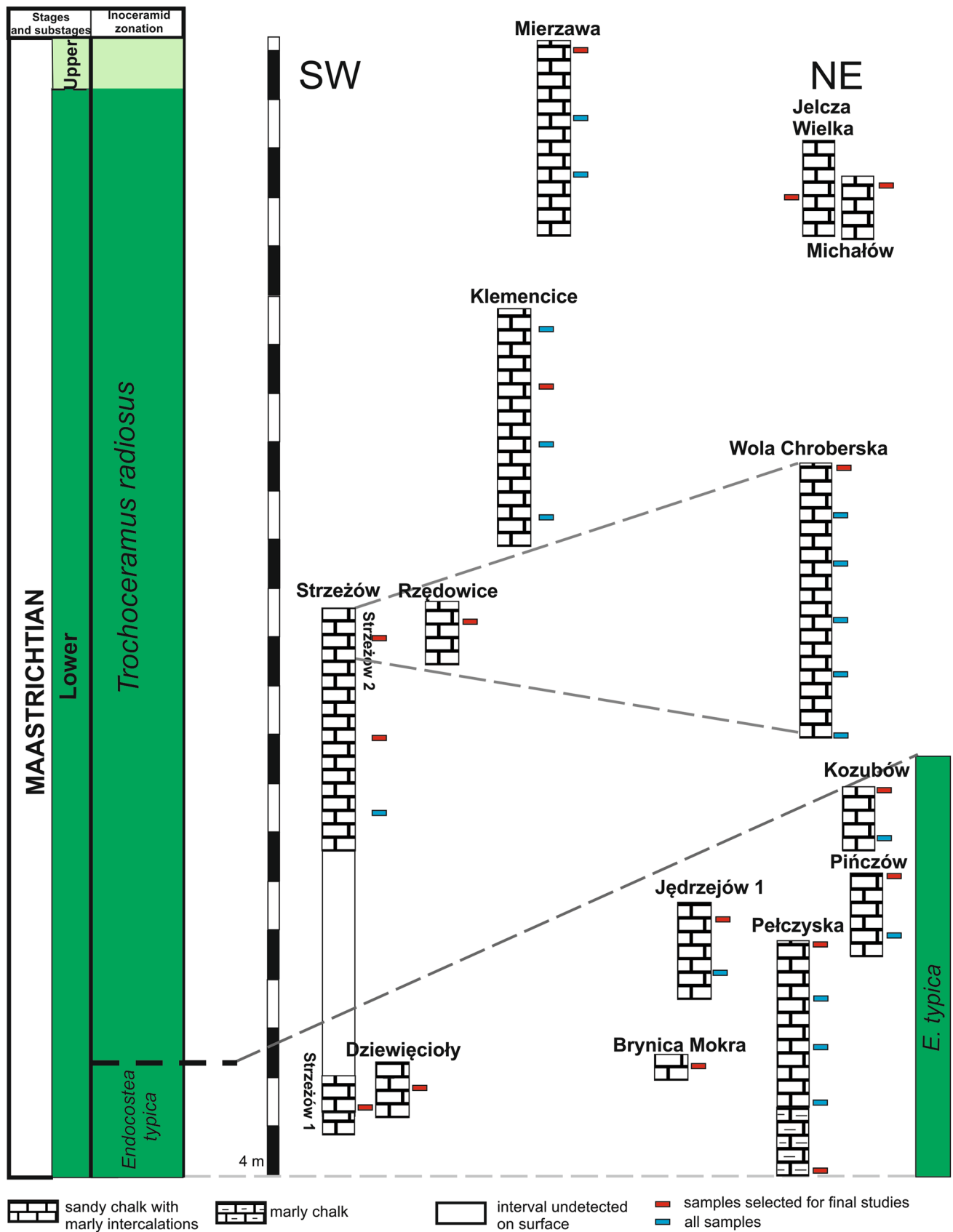


Fig. 2 Geological columns, stratigraphy (correlation of inoceramid zonation applied herein (after Walaszczyk et al. 2009, 2010, 2016) of the Upper Maastrichtian of studied area (Jurkowska 2016; this paper)

(Jurkowska 2016) sections, new findings of *Spyrideoceras tegulatus* (Von Hagenow 1842) forma A indicate the upper part of the *T. radiosus* Zone.

Jędrzejów 1 [N50°37'51.26"E20°17'37.05"] and Klemencice [N50°32'31.89"E20°11'53.47"] (Fig. 1b) are newly studied sections and are exposed at the temporary road cutting from Świątniki to Jędrzejów. In Jędrzejów 1 (Fig. 1b), sandy white chalk with abundant inoceramids is exposed, and specimens of *C. subcircularis* (3–4 cm) of the upper part of the *E. typica* Zone were recognized. In Klemencice, yellow, highly sandy chalk with a large number of *C. subcircularis* (6–9 cm) (Fig. 3a), *C.? glendivensis* (Fig. 3b), *Trochoceras* cf. *thomasi* (Walaszczyk, Kennedy, Klinger, 2009), *Platyceras* cf. *salisburgensis* (Fugger and Kastner, 1885) (Figs. 3g, 4a), and *C. barabini* were noted. Based on the inoceramid fauna, the section represents the upper part of the *T. radiosus* Zone (Walaszczyk et al. 2009, 2010).

Upper Maastrichtian (Fig. 2)

Trochoceras radiosus Zone

The newly studied Mierzawa section [N50°34'5.48"E20°13'2.44"] (Fig. 1b) represents the youngest Upper Cretaceous blue and yellow highly sandy chalk of the Miechów Synclinorium. A rich inoceramid fauna composed of *Spyridoceras tegulatus* (Von Hagenow, 1842) forma A (only in the lowermost part of the section) (Fig. 3e), *Spyridoceras tegulatus* (Von Hagenow, 1842) forma B (see: Walaszczyk et al. 2010) (Fig. 3f), *Cataceramus* cf. *subcircularis* (Meek 1876) (see: Walaszczyk et al. 2010) (Fig. 3c), and a single representative of '*Inoceramus*' cf. *ianjoaensis morgani* (Sornay 1973) (Fig. 3d), indicating the highest part of the *T. radiosus* Zone.

Inoceramid succession

In inoceramid terms, the studied succession comprises an interval from the *E. typica* to the *T. radiosus* Zone. In the *E. typica* Zone, the index taxon was noted only in the lowest part of the zone, while above this only a mass occurrence of small (<4 cm) *C. subcircularis* and *C. barabini* was observed. In the uppermost part of the zone, representatives of *C. subcircularis* and *C. barabini* occur together with *C. glendivensis*. A gradual size increase of *C. subcircularis* is observed within the *E. typica* and *T. radiosus* zones. The *E. typica* Zone could be informally divided into a lower (with index taxon) and upper assemblage (with numerous small representatives of a *Cataceramus* Assemblage).

The index taxon of the *T. radiosus* Zone was noted only in the lower part of the zone, while a few meters above the FO of *T. radiosus*, a single representative of *S. tegulatus* forma A was observed (Jurkowska 2016). Above this, numerous

representatives of *C. subcircularis* (6–9 cm), *C.? glendivensis*, *C. barabini*, *T. ?thomasi*, and *P. cf. salsiburgensis* occur. The *T. radiosus* Zone could be informally subdivided into a lower assemblage with the index taxa (*Trochoceras* Assemblage) and an upper one without them (*Cataceramus* Assemblage) (Walaszczyk et al. 2010). In the highest part of the studied succession, numerous representatives of *S. tegulatus* forma B (which is 4 m above mass occurrence of *S. tegulatus* forma A), *C. cf. subcircularis* (6–9 cm) and a single representative of '*I.*' cf. *ianjoaensis morgani* were noted. The youngest assemblage represents the highest part of the *T. radiosus* Zone.

Materials and methods

The inoceramids were collected bed by bed from each section providing a detailed stratigraphic framework enabling lateral correlations of the sections representing a single inoceramid zone. The sandy chalk was deposited under stable paleoenvironmental conditions with no signs of strong sea bottom currents, erosion, and synsedimentary tectonic movements, and are preferable targets for palynofacies analysis (Tyson and Follows 2000) and allow for the reconstruction of the intact pattern of particulate organic matter in the sedimentary basin. Because it was practically impossible to distinguish isochronous facies transitions according to Walther's law, we decided to analyze most similar common and predominating lithology of sandy and marly chalk in all sections. On the one hand, it minimizes differential diagenetic imprint on organic matter while on the other it ignores the subtle fluctuation of the local environment, which is negligible for the present scale of observation.

Thirteen outcrops (Figs. 1b, 2) representing Lower Maastrichtian to lower Upper Maastrichtian sandy/marly chalk were selected for palynofacies analysis. The samples were collected every 3 m from all sections (Fig. 2). For the final palynofacies analysis, the 14 most representative and least weathered samples from each section have been selected.

The idea to compare samples from several sections, preferably representing the same inoceramid zone and encompassing the widest lateral distance, was essential for the present study. Therefore, only a restricted number of samples met the requirements due to the inaccessibility of other stratigraphically equivalent areas devoid nowadays of outcrops.

One hundred grams of each rock sample was treated with 37% HCl and 40% HF to remove carbonates and silicate minerals. Finally, the organic residue was neutralized on a sieve, the diameter of which was 15 µm, and condensed by heavy liquid (ZnCl_2 , 2 g/cm³) separation. Two slides per sample were examined. Organic matter particles were

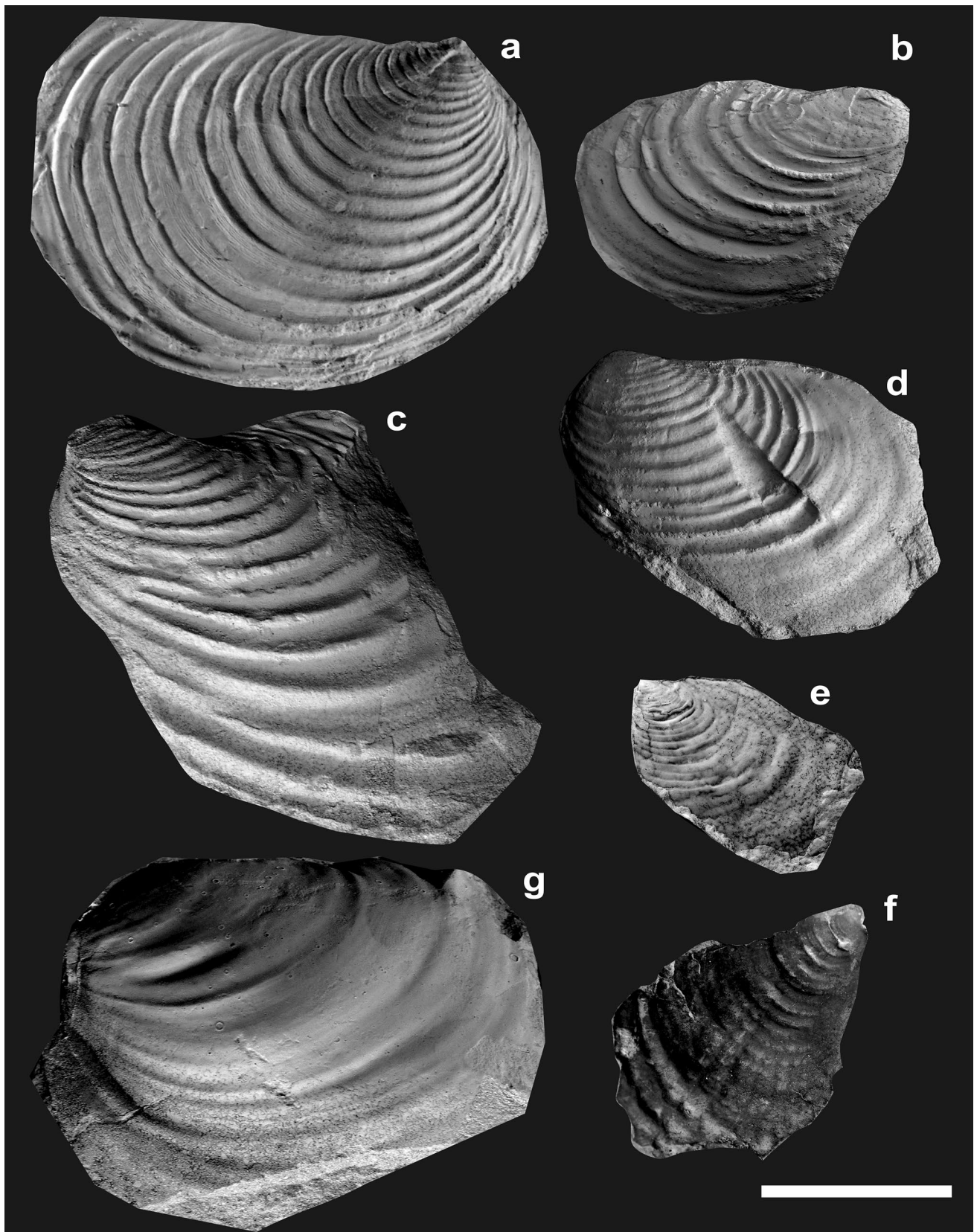


Fig. 3 **a** *Cataceramus subcircularis* (Meek 1876), Klemencice, *T. radiosus* Zone; ZI/68/004. **b** *Cataceramus?* *glendivensis* Walaszczyk, Cobban and Harries, 2001, Klemencice, *T. radiosus* Zone; ZI/68/003. **c** *Cataceramus* cf. *subcircularis* (Meek 1876), Mierzawa, *T. radiosus* Zone; ZI/68/006. **d** *Inoceramus* cf. *ianjonaensis* morgani Sornay, 1973, *T. radiosus* Zone; ZI/68/012. **e** *Spyridoceramus tegulatus* (Von Hagenow 1842) forma B, Mierzawa, *T. radiosus* Zone; ZI/68/009. **f** *Spyridoceramus tegulatus* (Von Hagenow 1842) forma A, Mierzawa, *T. radiosus* Zone; ZI/68/001. **g** *Platyceramus* cf. *salisburgensis* (Fugger and Kästner 1885), Klemencice, *T. radiosus* Zone; ZI/68/015. Scale bar 3 cm

counted to 500 per sample. Microphotographs were taken using a Nikon Eclipse E-600 microscope equipped with phase contrast and a digital camera.

The palynofacies analysis, i.e., the classification of the main morphological components, follows the categories established by Tyson (1993, 1995). Late Cretaceous

dinoflagellate assemblages are analyzed only for qualitative purpose, as the gonyalacoid type of cysts which predominate the samples are not characteristic for specific environmental conditions.

The measurements of opaque phytoclasts have been performed using a $\times 40$ microscope objective and a reticule. One hundred phytoclasts from each sample were counted. The shape of the clasts has been approximated to square or rectangular forms, and their respective edges have been measured. Following Tyson (1993), opaque phytoclasts with a length–width ratio higher than 3.0 have been classified as lath-shaped. Maximum and minimum clast size values are also presented to estimate the sorting ratio and distance from the shoreline. The Simpson's Index of Diversity ($1 - D$) and Shannon's diversity index diagram are provided for organic matter diversity reflecting proximal–distal position of the studied sections along the carbonate shelf.

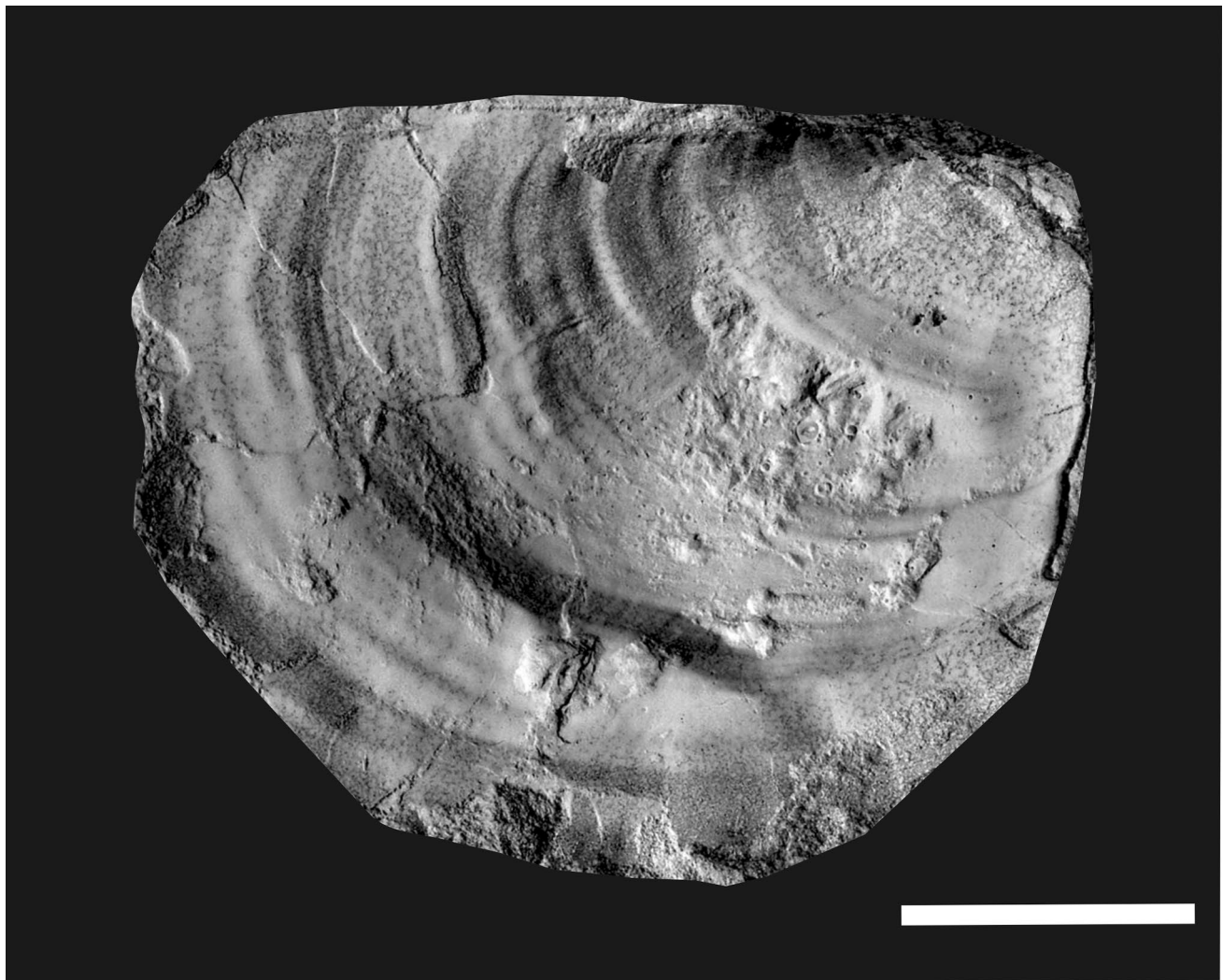


Fig. 4 *Platyceramus* cf. *salisburgensis* (Fugger and Kästner 1885), Klemencice, *T. radiosus* Zone; ZI/68/013. Scale bar 3 cm

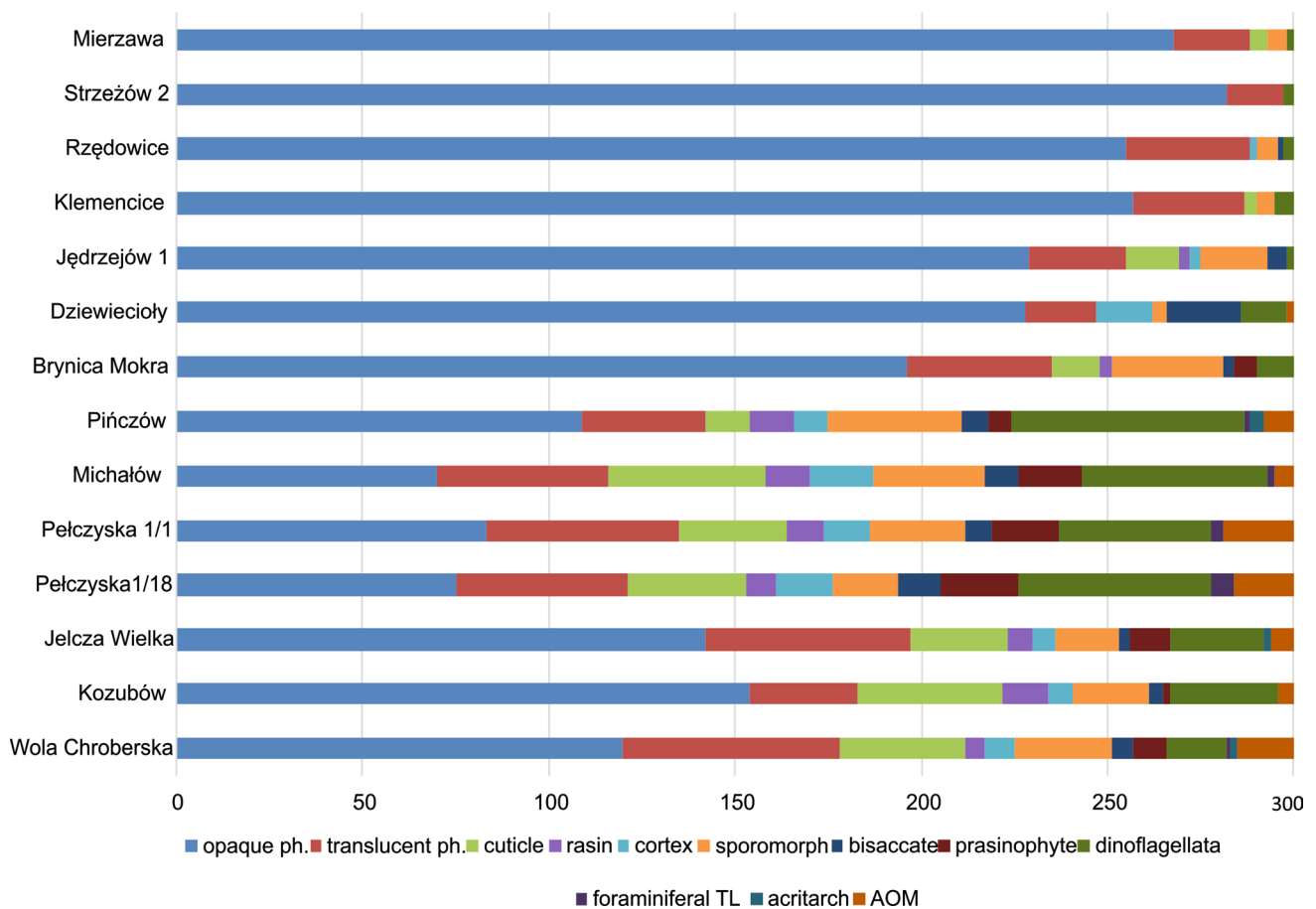


Fig. 5 Quantitative palynofacies data from the studied samples

Palynofacies associations—cluster analysis

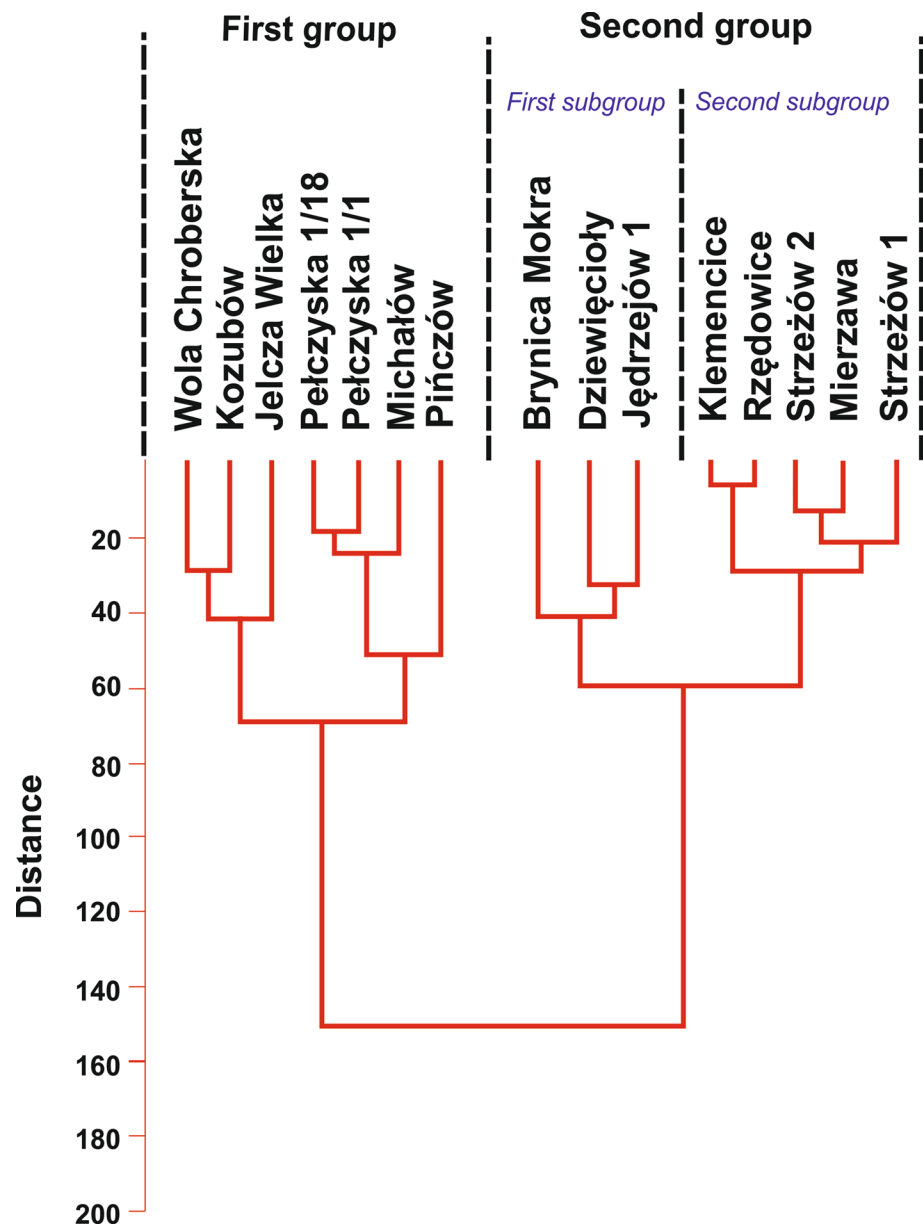
Based on the palynofacies analysis (Fig. 5), two main groups and two subgroups (Fig. 6) related to the facies distribution along the shelf are distinguished within the *E. typica* and *T. radiosus* zones. The groups are created automatically by hierarchical clustering provided by the PAST software package (Hammer et al. 2001). The “unweighted pair-group average” (UPGMA) algorithm has been applied with the Euclidean similarity index to ensure the best results. The Cohen’s kappa coefficient for the cluster analysis was 0.8535, which reflects an almost perfect agreement. Two major groups of palynofacies and one subgroup have been distinguished on the distance level of ca. 145 (Fig. 6). Minor internal difference of the first group could be explained by natural fluctuations of palynofacies components in shore vicinity and close geographical location (Müller 1959; Traverse 1988; Tyson 1995).

The first palynofacies group (Fig. 7a–d) has been recognized in samples from the Pełczyska (1/1 and 1/18), Pińczów, Kozubów, Wola Chrobberska, Jelcza Wielka, and Michałów sections. All samples are characterized by a high abundance and diversity of particulate organic matter. They

are also dominated by land-derived palynological components with a predominance of opaque phytoclast, pollen grains, as well as large and intact cuticle particles, resin, and cortex. Additionally, prasinophyte algae (*Tasmanites* sp., *Pterospermella* sp.) (only in Pełczyska, Kozubów, and Michałów) and chlorococcale algae (*Botryococcus* sp., *Palambages* sp.) (Pełczyska) have been recognized. Marine palynomorphs are moderately abundant, and are mostly represented by dinoflagellate cysts especially in the Pełczyska (1/1 and 1/18), Pińczów and Michałów sections. This leads to an internal diversification of this palynofacies group in terms of clustering (Fig. 8). Moreover, the samples from Pełczyska 1/1 and Pełczyska 1/18 show noticeable admixtures of foraminifera test linings and a noticeable content of amorphous organic matter (AOM), which also predominates the samples from Wola Chrobberska.

The predominance of the phytoclast group with a high content of opaque particles is the key feature of the second group. Within this group, two subgroups have been distinguished. The first subgroup (Figs. 7e, f; 8a–c) is recognized in the sections Brynica Mokra, Dziewięcioły, Jędrzejów 1, and is characterized by features typical for the second group,

Fig. 6 Dendrogram (r-mode) of the palynofacies from outcrops studied showing the three distinctive environmental groups



but a moderate content of translucent phytoclast and sporomorphs is also noted. Cuticle, resin, and single-cortex tissues are present, but with conspicuously declining tendency. Marine plankton content represented by dinoflagellate cyst is noticeable. The samples from Dziewięcioły are unique as they contain large quantities of bisaccate pollen grains.

The second subgroup of the second group (Fig. 8d–f) is encountered in the Strzeżów 1 and 2, Rzędowice, Klemencice, and Mierzawa sections, and is characterized by a predominance of the phytoclast group with high content of opaque particles. Translucent particles are mainly composed of brown structured wood tissue. Dinoflagellate cysts are noted in all samples.

To understand the provenience of the palynofacies grouped by cluster analysis, the widely used plots of Tyson (1993, 1995) and further modification of Roncaglia and Kuijpers (2006) and Zobaa et al. (2015) have been used. Tyson's (1993) ternary diagram revealed the concentration of the studied samples within fields I and III (Fig. Online Resource 1). According to the definition, they represent “highly proximal shelf or basin” and “heterolithic oxic shelf (proximal shelf)” environments, respectively. In this way, the most diverse palynofacies associations from the first group have been placed in the “proximal shelf” environment field and the second group with monospecific high content of opaque particles in the “highly proximal

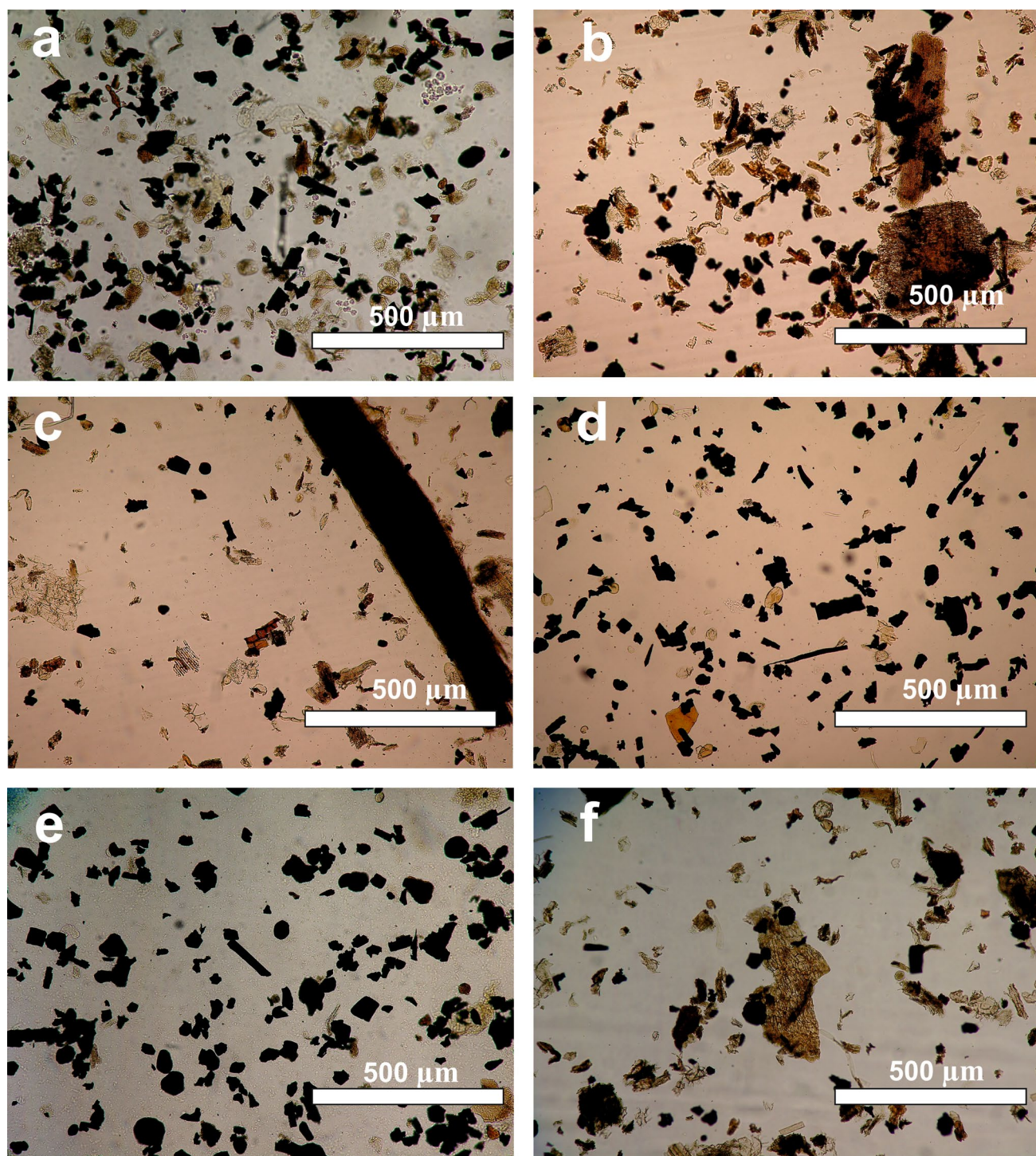


Fig. 7 Microscopic photographs of the most representative palynofacies of studied sections. Scale bar of 500 µm valid for all captions. First group (proximal palynofacies): **a–c** Pełczyska, **d** Pińczów, **e** Jelcza Wielka, **f** Michałów

shelf” field. The positions within the fields in the present study are evidently constrained by the content of marine palynomorphs with a major influence of the abundance dinoflagellate cysts.

Palynofacies—diversity indexes

Due to some discrepancies within the used palynofacies ternary plots (Tyson 1993, 1995; Roncaglia and Kuijpers

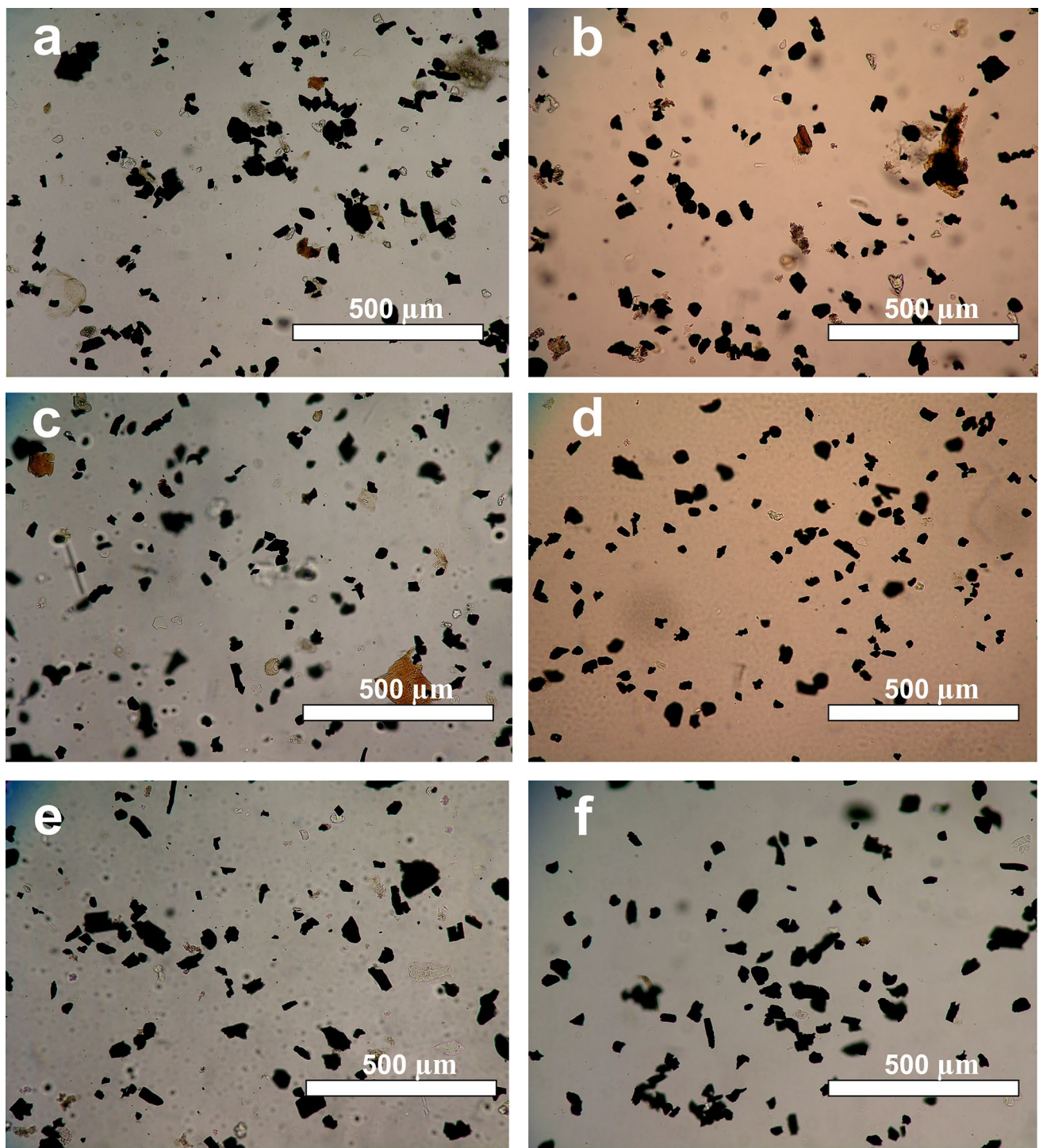


Fig. 8 Microscopic photographs of the most representative palynofacies of studied sections. *Scale bar* of 500 µm valid for all captions. Second group-first subgroup (intermediate palynofacies): **a** Brynica

Mokra, **b** Dziewięcioły, **c** Jędrzejów 1; Second group-second subgroup (distal palynofacies) **d** Strzeżów 1, **e** Mierzawa, **f** Strzeżów 1

2006), especially with respect to the identification of proximal–distal environments, diversity indexes of sedimentary organic matter have been calculated. Widely used diversity indexes in the ecological study are the Simpson's Index of

Diversity ($1 - D$) and the Shannon's diversity index, both of which have been used herein. The results of the diversity analysis are presented in the Fig. 9a, b, where the values of both indexes are plotted for all studied locations in ascending

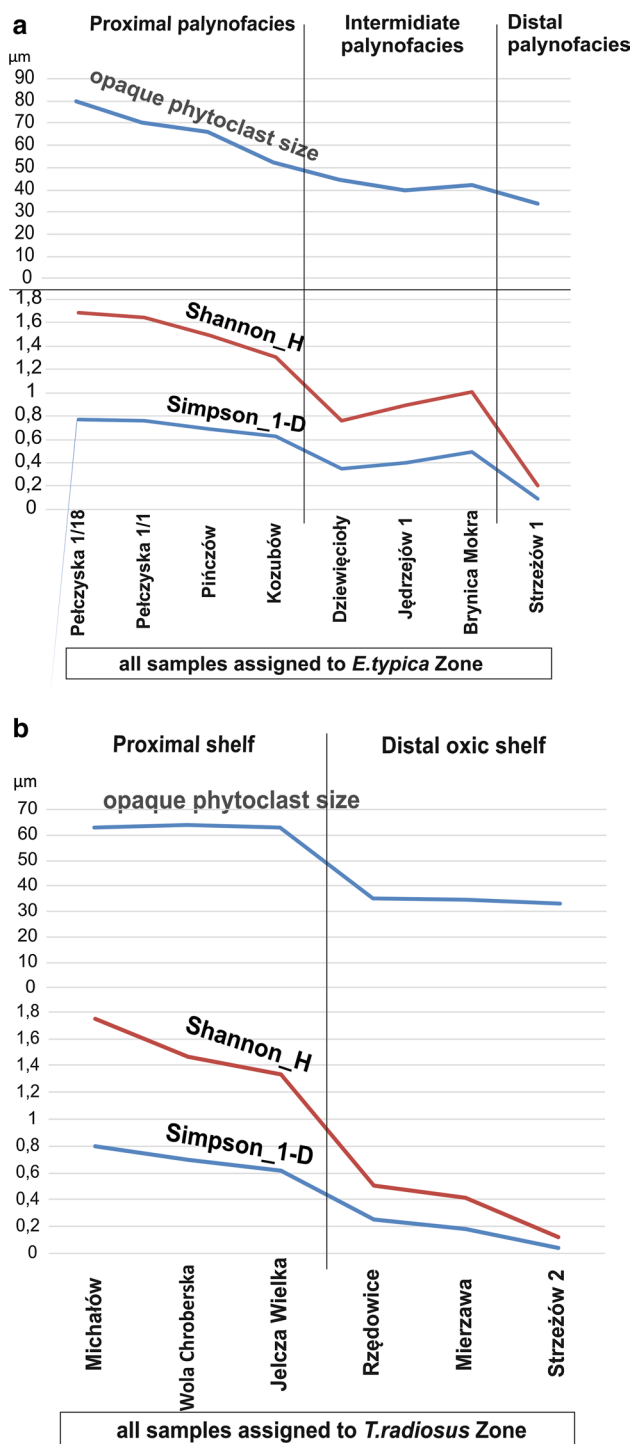


Fig. 9 a, b Lateral variations of opaque phytoclasts sizes and diversity indexes from two transects along distinguished palynofacies zones

order. Simpson's Index spans the values from 0.004 to 0.80, and Shannon's index from 0.12 to 1.75. In both cases, the increasing values of indexes coincide with a higher biodiversity connected to a more diversified palynological content.

The higher diversity of organic particles could also be interpreted as a proximity index to the source area (Müller 1959; Degens and Mopper 1976; Tyson 1993). The ranges of biodiversity constrained by the Simpson and Shannon indexes follow the three groups of palynofacies revealed by the cluster analysis (Fig. 6). The boundaries between the diversity groups are sharply distinguishable by the Shannon index values (Fig. 9a, b). The boundary between less diversified and intermediate palynofacies groups run across 0.507 and 0.681 and between intermediate and most diversified palynofacies across 1.003 and 1.334 of the Shannon index. Subsequently, general diversity indexes for all samples are tested for time intervals of the samples assigned to the Maastichtian *E. typica* and *T. radiosus* zones separately (Fig. 9a, b). For the *E. typica* Zone, three generations of index values are discernible with the boundaries placed between the Kozubów-Dziwięcioły and the Brynica Mokra-Strzeżów 2 sections. These indexes reflect three sedimentary zones within a basin with westwards-declining diversity of organic matter. A similar trend is observed for the *T. radiosus* Zone, however only two diverse populations were noted.

Distance from the shoreline—opaque phytoclasts' size

The size analysis of opaque phytoclasts used for evaluation of the distance (Tyson and Follows 2000) from the shoreline is presented in Fig. 9a, b. The actual distance of 35 km between analyzed areas approximates the original sedimentary distances in the basin with relation to the shoreline (Figs. 4, 6). Mean values of opaque phytoclast sizes from samples collected from all analyzed sites range from 80 μm to about 35 μm (Fig. 9a, b). The study of samples revealed that quantitative variations in size, shape, and preservation of plant debris correlate with actual distance from the shoreline along shelf deposits. The opaque phytoclast size curve for the *E. typica* Zone shows a continuous but relatively steep trend of decreasing values from the Pelczyska to the Strzeżów 1 sections. The maximum value of 80 μm for Pelczyska 1/18 is related to a proximal position in the basin. The curve for the *T. radiosus* Zone, starting with a value of 62 μm value for the Michałów section, reveals two populations of phytoclast size and a flat character. This probably expresses a more distal locality with respect to the terrestrial organic matter input and a more stable dispersal of organic particles offshore. When calculating opaque phytoclast sizes, the clear differences between maximum and minimum size ranges in the Pelczyska and Pińczów sections stood out. This trend decreased distally. According to Tyson (1993), this results from a low sorting value, characteristic for nearshore areas, which is a function of distance from the source area.

According to the size analysis of Tyson and Follows (2000), the 35- μm size of the opaque phytoclast

Fig. 10 Qualitative distribution of dinoflagellate cyst in studied samples

	Brynica Mokra	Dziewięcioły	Jelcza Wielka	Jędrzejów 1	Klemencice	Kozubów	Michałów	Mierzawa	Pełczyńska 1/1	Pełczyńska 1/18	Pińczów	Rzędowice	Strzeżów 2	Wola Chrobberska
<i>Achomosphaera</i> sp.									X	X				
<i>Alterbidinium</i> sp.									X	X				
<i>Apteodinium delicatum</i>											X			
<i>Areoligera coronata</i>							X				X			
<i>Areoligera senonensis</i>							X							
<i>Areoligera</i> sp.	X	X	X								X			X
<i>Biconidinium reductum</i>											X			
<i>Botrycoccus</i> sp.						X	X		X	X				
<i>Cerodinium diebelii</i>									X	X				
<i>Circulodinium distinctum</i>											X			
<i>Cleistosphaeridium</i> sp.			X					X			X		X	X
<i>Dinogymnium acuminatum</i>									X	X				
<i>Dinogymnium longicorne</i>									X					
<i>Florentinia aculeata</i>										X				
<i>Glaphyrocysta</i> sp.	X		X		X			X			X			X
<i>Heterosphaeridium</i> sp.	X			X	X		X	X	X	X	X			X
<i>Hystriochosphaeridium</i> sp.									X	X	X			
<i>Isabelidium</i> sp.									X	X				
<i>Kiokansium polypes</i>							X							
<i>Kiokansium</i> sp.	X	X	X		X	X	X	X			X		X	X
<i>Membranilarnacia</i> sp.											X			
<i>Odontochitina operculata</i>											X			
<i>Odontochitina</i> sp.											X			
<i>Oligosphaeridium complex</i>									X	X				
<i>Oligosphaeridium complex</i>							X							
<i>Operculodinium</i> sp.							X							
<i>Palaeocystodinium</i> sp.									X	X				
<i>Palambages</i> sp.										X				
<i>Palynodinium grallator</i>									X	X				
<i>Pervosphaeridium</i> sp.											X			
<i>Pterospermella</i> sp.						X	X		X	X				
<i>Riculacysta</i> sp.	X													
<i>Spinidinium eggeri</i>									X					
<i>Spiniferites ramosus</i>									X	X				
<i>Spiniferites</i> sp.	X		X		X						X	X	X	X
<i>Spongodinium delitiense</i>									X	X				
<i>Surculosphaeridium longifurcatum</i>									X	X	X			
<i>Tasmanites</i> sp.									X	X				

corresponds to a distance of 10 km from the shoreline. In our study, the same values occur in the sections situated at least approximately 20–35 km from inferred shoreline. This discrepancy is probably due to seafloor configurations and sedimentary basin character for both case studies.

Phytoplankton assemblages

Only a few samples from the studied material yield rich and diversified phytoplankton assemblages (Fig. 10). They are represented by autotrophic and heterotrophic dinoflagellates cysts, phycococci of prasinophytes, and colonies of chlorococcale algae (Fig. 11). The most diverse assemblages occur in the Pełczyńska, Pińczów and Michałów

sections. The Wola Chrobberska, Kozubów, Jelcza Wielka, Brynica Mokra and Dziewięcioły sections are characterized by scarce phytoplankton content, whereas the Jędrzejów, Klemencice, Rzędowice, Strzeżów and Mierzawa sections are practically barren with records of only 2–5 specimens per slide.

Discussion

Cluster analyses of organic matter combined with palynofacies diversity and opaque phytoclast size analyses in stratigraphically well-constrained sections provide valuable data for paleogeographic reconstructions in monotonous Upper

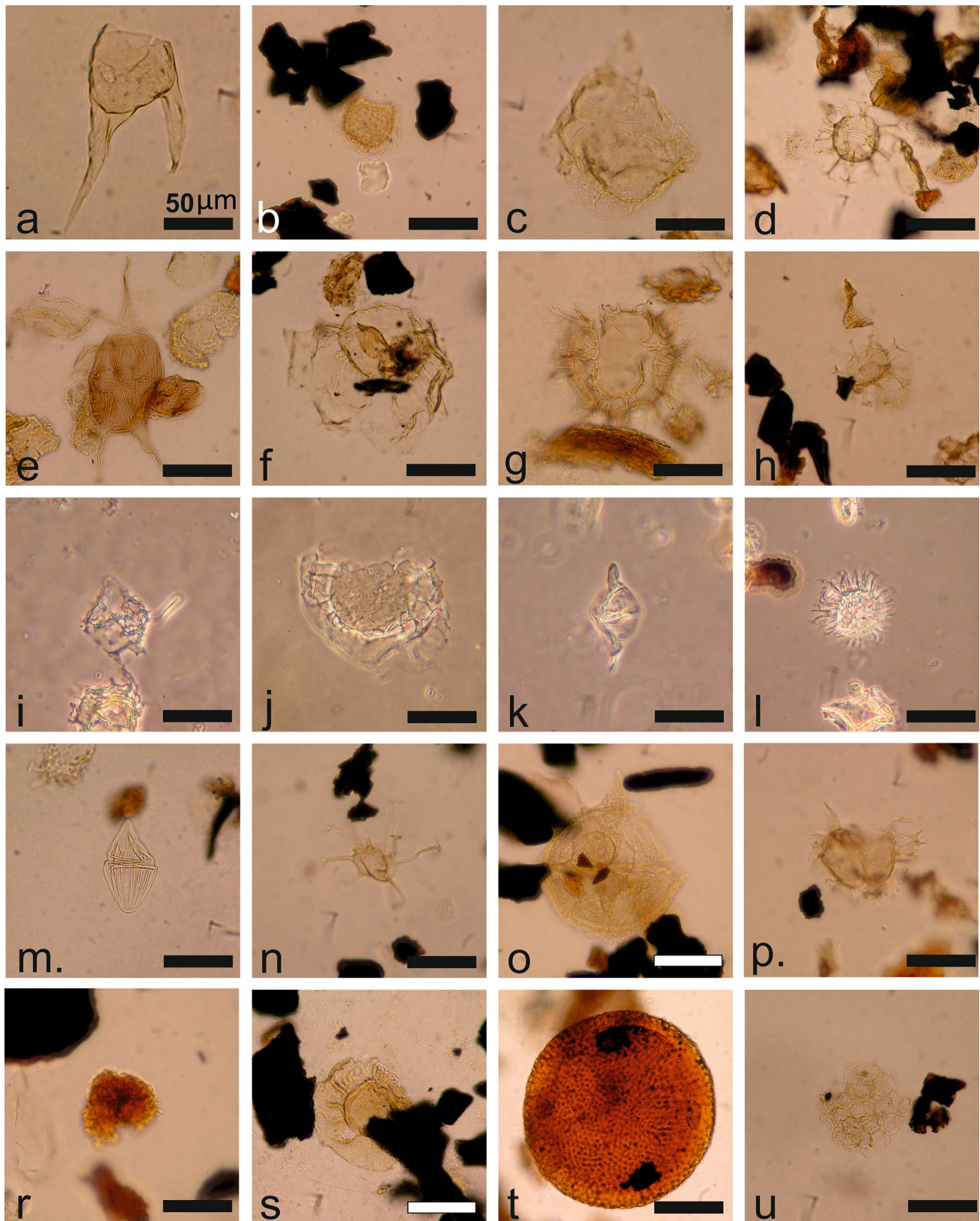


Fig. 11 Photographs of the selected taxa of dinoflagellate cyst and other algae from studied samples. Scale bar of 50 μm valid for all photographs. **a** *Odontochitina operculata* (Wetzel, 1933) Deflandre and Cookson, 1955, Pińczów. **b** *Membranilarnacia* sp Eisenack, 1963, Pińczów. **c** *Spongodinium delitiense* (Ehrenberg, 1838) Deflandre, 1936, Pełczyńska 1/1. **d** *Spiniferites ramosus* (Ehrenberg, 1838) Mantell, 1854, Pełczyńska 1/18. **e** *Cerodinium diebelii* Alberti, 1959) Lentin and Williams, 1987, Pełczyńska 1/18. **f** *Riculacysta amplexa* Kirsch, 1991, Brynica Mokra. **g** *Areoligera senonensis* Lejeune-Carpentier, 1938, Michałów. **h** *Hystriosphæridium tubiferum* (Ehrenberg, 1838) Deflandre, 1937, Pińczów. **i** *Spinidinium eggeri* Kirsch, 1991, Pełczyńska 1/1. **j** *Glaphyrocysta semitecta* (Bujak, 1980) Lentin and Williams, 1981, Brynica Mokra. **k** *Biconidinium reductum* (May, 1980,) Kirsch, 1991, Pińczów. **l** *Heterosphaeridium* sp Cookson and Eisenack, 1968, Kozubów. **m** *Dinogymnium acuminatum* Evitt et al., 1967, Pełczyńska 1/18. **n** *Oligosphaeridium complex* (White, 1842) Davey and Williams, 1966, Michałów. **o** *Apteodinium delicatum* (Davey, 1975) Schrank, 1987, Pińczów. **p** *Palynodinium grallator* Gocht, 1970, Pełczyńska 1/1. **q** *Botrycoccus* sp., Pełczyńska 1/1. **r** *Pterospermella* sp., Pińczów. **s** *Tasmanites* sp., Pińczów. **t** *Palambages* sp., Pełczyńska, 1/1

Cretaceous carbonate-siliciclastic rocks. The models of organic matter distribution which originated from clastic sedimentary systems (Tyson and Follows 2000; Roncaglia and Kuijpers 2006) could not be directly applied to carbonate sedimentary systems (Courtinat and Malartre 2003; Goetz et al. 2008). The dynamic nature of sedimentary basins and carbonate sediments influenced by siliciclastic deposits have to be carefully considered are especially important for creating meaningful paleogeographic reconstructions. According to Zeller et al. (2015), a mixing of carbonate production and siliciclastic input causes variable sedimentation conditions and a high degree of lateral and vertical facies heterogeneity. Generally, carbonate facies development is triggered by the requirements of the carbonate-secreting biota, which are controlled by temperature, salinity, sea-level change, input of siliciclastic sediments, and nutrients (Sanders and Höfling 2000). It seems that the sedimentary conditions of the Late Cretaceous low-energy basin in the Miechów Synclinorium have been predominately driven by blooms of coccolithophores (Saavedra-Pellitero et al. 2014) as well as inputs of siliciclastic deposits of terrestrial origin. These conditions lead to restricted environments unaffected by high energetic sedimentary processes more characteristic for clastic deposits and classic carbonate ramp setting (Courtinat and Malartre 2003). Therefore, it seems that the aforementioned trends within sizes of opaque phytoclasts as well the diversity indexes for particulate organic matter are a reliable proxy of shoreline distance within the studied area. The application of these parameters combined with those from palynofacies and phytoplankton analyses allowed for reconstructions of proximal–distal palynofacies trends to be distinguished (Fig. 12) (Tyson and Follows 2000; Tyson 1993; Batten 1982; Goetz et al. 2008).

The first group of palynofacies characterized by high abundance of particulate organic matter is dominated by land-derived components with a predominance of opaque phytoclast pollen grains as well as large and intact cuticle particles, resin, and cortex. This, combined with a higher diversity of organic particles and maximum sizes of opaque phytoclast, indicates proximal conditions in the Wola Chrobberska and Pełczyńska sections. These sections are conspicuously enriched in AOM, indicating lagoonal conditions or vicinity of marshes, where AOM is well preserved (Rich 1989). Moreover, in Pełczyńska, the occurrence of chlorococcale algae, represented by *Botrycoccus* sp. (Traverse 1988; Wetzel 1983) and *Palambages* sp. (Prauss 2000), confirms the influence of fresh water, at least occasionally. *Botrycoccus* algae are widely used in palynological studies to determine fresh water environments (lacustrine, fluvial, lagoon, and deltaic) (e.g., Batten and Lister 1988; Guy-Ohlson and Norling 1988; Riding et al. 1991; Williams 1992). Subsequently, *Botrycoccus* colonies may be transported by the rivers towards marine shelf facies (Combaz et al. 1977; Caratini et al. 1983; Guy-Ohlson 1986; Prauss 1989) and therefore often occur together with marine algae.

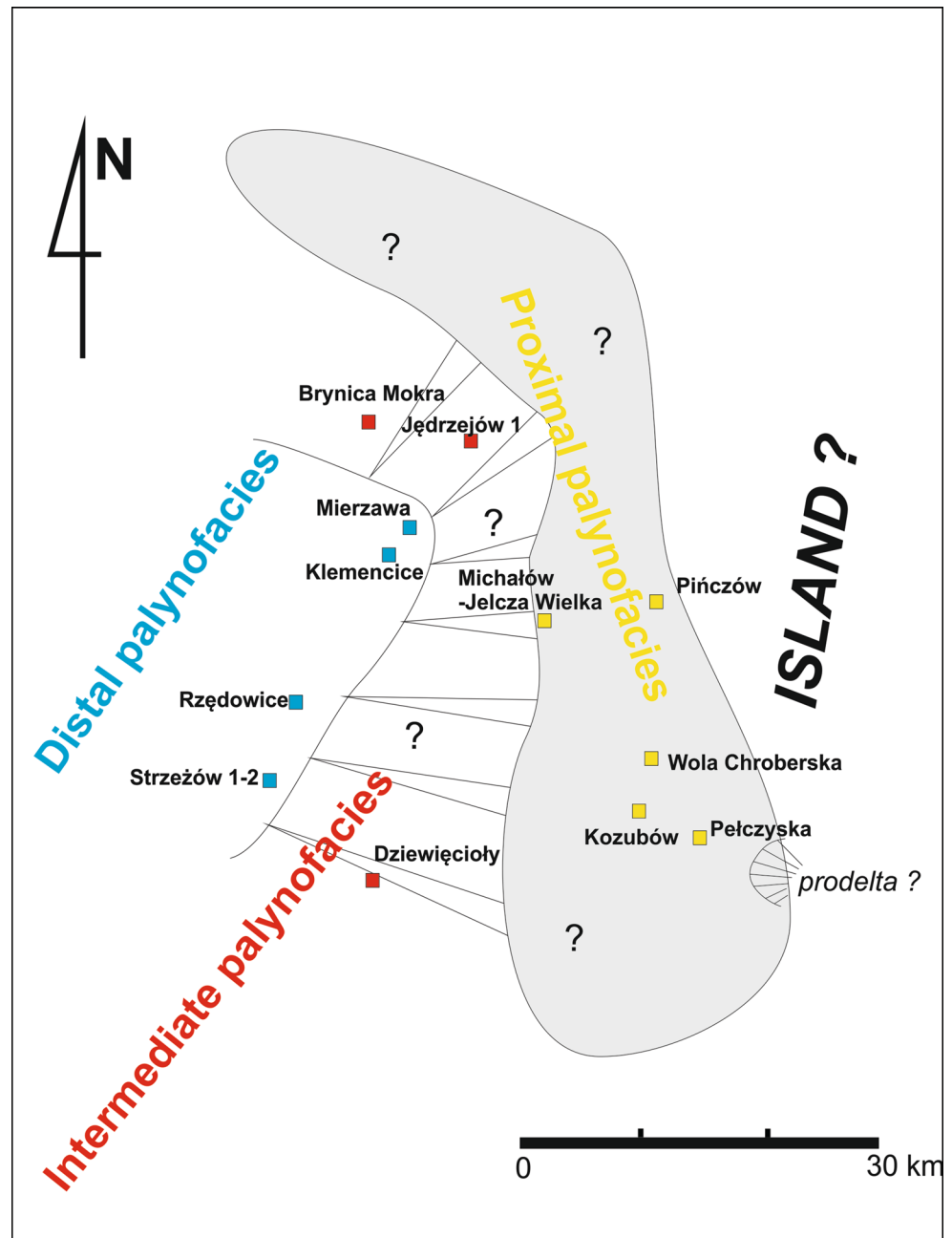
An increased occurrence of prasinophyte algae, represented by *Tasmanites* sp. and *Pterospermella* (Tappan 1980; Bernier and Courtinat 1979; Prauss 1989) with large intact cuticle tissues and needle-shaped phytoclasts, seems to confirm a close distance of these sites to the fluvio-deltaic or lagoonal environments, and probably local water stratification (Tyson 1984, 1993). This is further indicated by the apparent influx of terrestrial palynomorphs in the Kozubów and Michałów sections.

The first subgroup within the second group of palynofacies, predominated by opaque particles with moderate content of translucent phytoclast and sporomorphs indicates intermediate palynofacies. This is further supported by the moderate index diversity and medium value of opaque phytoclast sizes. The absence of intermediate palynofacies in the *T. radiosus* Zone is the result of lacking the outcrops due to the extensive cover of the Maastrichtian rocks by younger deposits. The conspicuous enrichment of baccate pollen grains in the Dziewięcioły section, located in intermediate palynofacies, is probably due to bypass processes (Mildenhall 2003) generated by wind transport from the conifer-rich area of Pełczyńska site located next to the postulated land area.

The second subgroup of the second palynofacies group, composed almost solely of phytoclasts with high content of opaque particles, is characterized by the lowest diversity index and smallest size values of opaque phytoclast, and indicates the distal palynofacies of the studied sections.

The declining tendencies of marine palynomorphs from the proximal, through the intermediate, towards the distal part of the shelf are probably driven by local nutrient

Fig. 12 Palynofacies interpretation of studied area during Maastrichtian time



circulation in the basin. Dinoflagellates produce fossil cysts during periods of unstable conditions in shallow-water environments along continental margins and around oceanic archipelagos (Wall et al. 1977). In normal conditions, the abundance of dinoflagellate cysts is in inverse proportion to the content of terrestrial palynomorphs. However, they can predominate in shelf areas devoid of active fluvio-deltaic influences, in areas poorly covered by vegetation, and in areas of enhanced primary productivity (Tyson 1993). Increased productivity is usually the effect of upwelling currents generated by wind, or nutrient influx by terrestrial runoff (Davey 1970; Davey and Rogers 1975; Hooghiemstra

et al. 1986). It often results in an enrichment in peridinioid dinoflagellate cyst taxa (Wall et al. 1977; Bujak 1984). According to many authors (Downie et al. 1971; Davey 1970; Mutterlose and Harding 1987; Lister and Batten 1988), peridinioid-dominated assemblages are also indicative of nearshore lagoonal, estuarine, or brackish water environments (Pełczyska). Generally, dinoflagellate cyst diversity shows on offshore increase. However, reversed tendencies sometimes occur in estuarine environments (Wall et al. 1977; Morzadec-Kerfourn 1977; Tyler et al. 1982). Impoverished macrofauna assemblages observed by

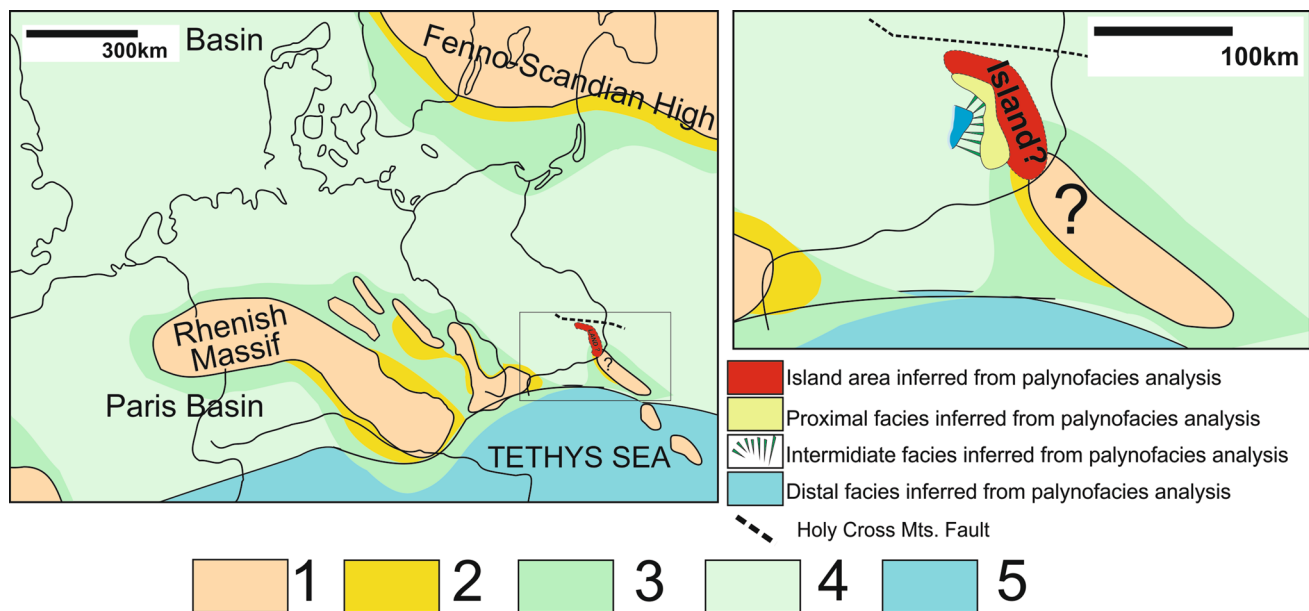


Fig. 13 Simplified Late Cretaceous (75 Ma) paleogeographic map (Naidin 1959; Kauffman 1973; Ziegler 1990; Philip et al. 2000; Świdrowska et al. 2008; Jurkowska et al. 2015 modified; Bojanowski et al. 2017) with regional paleogeographical data inferred from

palynofacies analysis. 1 areas of non-deposition, 2 deltaic, coastal and shallow-marine siliciclastic facies, 3 near-shore facies, 4 shallow-marine facies, 5 deeper-marine facies

Jurkowska (2016) also reflect offshore diminishing diversity and abundance in the Miechów Synclinorium.

The results presented here suggest the occurrence of an island area in close proximity of the sections belonging to the first group, i.e., the proximal palynofacies. The proximal part of this system is indicated by a high diversity of terrestrial organic matter, influence of fresh water input (*Botryococcus*), high productivity probably controlled by terrestrial runoff, which results in high abundance of peridinioid dinoflagellate cysts and larger sizes of opaque phytoclasts. Towards the distal part of the shelf, the number of terrestrial-derived particles and dinoflagellates declines, and the size of wood particles and the diversity indexes are reduced. The distal facies are practically solely represented by opaque phytoclasts assemblages of rounded shapes and small sizes. The low number of dinoflagellate cysts in this part of the system is explained by local trophic conditions.

The island, which was probably connected with the Kukurritz Island (Fig. 13) was located close to the Danish-Polish Trough (present day Mid-Polish Anticlinorium). Generally, two divergent opinions of the time of inversion of the Danish-Polish Trough have been established. Dadlez et al. (1997) and Leszczyński (2012) presented the idea of a multiphase inversion, which may have started during Santonian–Campanian and reactivated older synsedimentary structures. The key data for this interpretation is based on lithofacies pattern and geometry of the Upper Cretaceous succession and thickness analysis. Kutek and Głazek (1972), Świdrowska and Hakenberg (1999), Świdrowska et al. (2008), and Jurkowska

(2016) presented data which indicate that the inversion could not have started before the Maastrichtian. This is based on the increasing thickness of the whole Upper Cretaceous succession in the direction of the Holy Cross Mountains (Kutek and Głazek 1972; Heller and Moryc 1984; Świdrowska and Hakenberg 1999; Świdrowska et al. 2008; Jurkowska 2016). Świdrowska and Hakenberg (1999) also presented lithofacies data of Upper Cretaceous deposits, indicating increasing water depths towards the Holy Cross Mountains. Recently, based on detailed stratigraphy, a more complicated model of inversion has been discussed (Walszczyk and Remin 2015; Jurkowska 2016). New high-resolution seismic data (Krzywiec 2000; Krzywiec et al. 2009) also indicate a more complicated model in which the individual parts of the Danish-Polish Trough have been inverted at different times.

The data presented in this paper indicate that during the early Late Maastrichtian, the sea was still present in the Miechów Synclinorium, and an uplifted area of the Holy Cross Mountains part of the Danish-Polish Trough was emerged and was providing the terrigenous material to the southeastern areas.

Conclusions

The palynological study combined with a palynofacies analysis supported by size analysis of opaque phytoclasts and diversity indexes for particulate organic matter in well stratigraphically constrained monotonous Upper Cretaceous

carbonate-siliciclastic sediments devoid of unequivocal sedimentological and paleontological data is a valuable tool in determining the depositional environments. Despite the monotonous lithology and the lack of macrofaunistic paleoenvironmental indicators, the distribution of sedimentary organic matter suggests the occurrence of proximal to distal shelf facies belts with a well-developed intermediate facies zone. The size analysis of opaque phytoclasts indicates a close distance of the studied area from the shore. This is also confirmed by the increased quantity of chlorococcale algae represented by *Botrycoccus* sp. and *Palambages* sp. and the increase number of heterotrophic *Peridinioid* species and foraminifera test linings in proximal samples. The presented data indicate that during the Maastrichtian an uplifted area existed in the Holy Cross part of the Danish-Polish Trough. The youngest Upper Cretaceous rocks of the Miechów Synclinorium, for the first time described in the present paper, indicate that the sea was still present during the early Late Maastrichtian.

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