

Palaeoenvironmental changes recorded in the oxygen and carbon isotope composition of Kimmeridgian (Upper Jurassic) carbonates from central Poland

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Oxygen and carbon isotope ratios of well-preserved calcitic bivalves from the Lower–lowermost Upper Kimmeridgian of Central Poland (SW margin of the Holy Cross Mountains) have been studied to reconstruct palaeoenvironmental conditions and variations in ancient water chemistry. Low and scattered $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of bivalve shells from shallow carbonate deposits of the Hypselocyclum and the Hypselocyclum–Divisum zone boundary (–3.5 to –1.5, and 2.6 to 4.0‰, respectively) are a result of salinity changes and local variations in the composition of dissolved inorganic carbon (DIC) in conditions of restricted water circulation. A slight increase in bivalve $\delta^{13}\text{C}$ values and more densely clustering of $\delta^{18}\text{O}$ values is observed after the marine transgression at the Divisum–Mutabilis zone boundary. A global decrease of $\delta^{13}\text{C}$ values of marine carbonates is partly recorded in Lower–lowermost Upper Kimmeridgian bulk carbonates from central Poland (from the Radomsko Elevation and the Wieluń Upland). Local negative shifts and data scatter are, however, observed in rocks deposited in a very shallow environment of carbonate platforms during the Platynota and Hypselocyclum chrons. This interval corresponds to the uppermost part of the lowstand systems tract of a major regressive trend, which had started in the Oxfordian.

Key words: Upper Jurassic, stable isotopes, oysters, *Trichites*, carbonate platform, salinity effects.

INTRODUCTION

Lower–lowermost Upper Kimmeridgian (Platynota–Mutabilis zones) strata of central Poland, which were deposited during phases of growth and decay of shallow-water carbonate platforms, are poorly investigated as regards their mineralogical and geochemical compositions. This is due to the restricted outcrop area and problems with dating and correlation of the beds. Recent studies of [Matyja et al. \(2006\)](#), [Matyja \(2011\)](#), [Wierzbowski \(2017\)](#), and [Wierzbowski and Główniak \(2018\)](#) have, however, allowed re-definition of lithostratigraphical units and precise biostratigraphical dating of the Kimmeridgian succession in the Wieluń Upland, the Radomsko Elevation and the SW margin of the Holy Cross Mountains.

The present study is based on accessible Lower–lowermost Upper Kimmeridgian exposures in these regions and archival samples. Well-preserved and stratigraphically well-dated bivalve shells were sampled from the Małogoszcz section (SW margin of the Holy Cross Mountains). Bulk carbonates from the Wieluń Upland, its vicinity, and the Rogaszyn section (the Radomsko Elevation) were additionally investigated ([Fig. 1](#)).

Oxygen and carbon isotope analyses have been undertaken to provide information on the environmental conditions recorded in marine carbonates, including variations in water temperatures, salinity and biological productivity. The results obtained are compared with the coeval geochemical record of other regions to discriminate between local and global factors. Long-term bathymetry and water circulation changes, which occurred in the epicontinental Polish basin during the Late Jurassic, are additionally discussed as an alternative factor controlling the isotope record. The isotope data provided may have broader significance due to the similarity in facies development of Kimmeridgian deposits in various parts of Europe.

GEOLOGICAL SETTING

Deep-water biohermal and bedded Oxfordian–lowermost Kimmeridgian limestones and marls of the sponge megafacies are overlain in central Poland by relatively shallow-water Lower Kimmeridgian carbonate facies ([Kutek, 1968, 1994](#); [Matyja, 1977, 2011](#); [Matyja et al., 2006](#); [Matyja and Wierzbowski, 2014](#); [Wierzbowski, 2017](#)). This change was a result of pronounced shallowing of the depositional basin and progradation of the carbonate platform from east and north-east to west and south-west ([Matyja and Wierzbowski, 2014](#)). The shallow-water carbonate sedimentation started in the latest Planula Chron on the SW margin of the Holy Cross Mountains and the Radomsko El-

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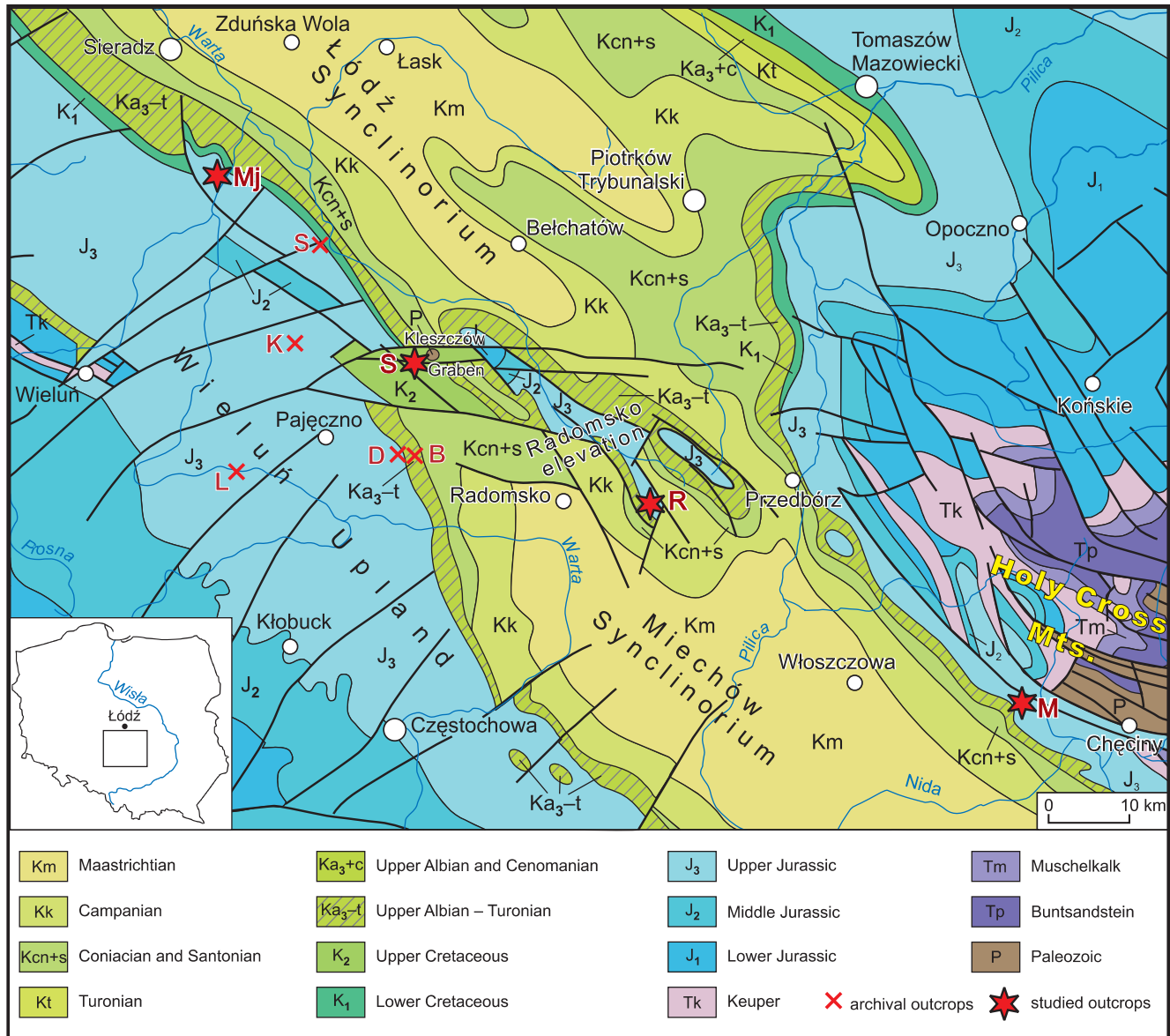


Fig. 1. Geological map of central Poland (after Dadlez et al., 2000) with location of the study area

Outcrops studied: M – Małogoszcz, Mj – Majaczewice, R – Rogaszyn, S – Szczerców; archival outcrops: B – Błota Kruplińskie (cores KP2 and KP3), D – Dubidze (core 6W), K – Kule (84), L – Lisowice (PJ110), S – Sarnów

evation, and in the Platynota Chron in the Wieluń Upland (Matyja, 2011; Matyja and Wierzbowski, 2014; Wierzbowski, 2017; Wierzbowski and Głowniak, 2018).

Moderately shallow and shallow-water Lower Kimmeridgian deposits, which are nowadays restricted to the eastern part of the Wieluń Upland and its vicinity, overlie well-bedded micritic limestones (Wolbrom Limestone Member) and marls (Latośówka Marl Member) of the Pilica Formation belonging to the sponge megafacies. The Lower Kimmeridgian deposits studied comprise:

- bedded chalky limestones with siliceous sponges rich in benthic fauna, and micritic limestones and marls of the Prusicko Formation known from the Wieluń Upland and dated to the Platynota Zone,
- oolitic and chalky limestones with oncolites, and marls of a so-called “oolitic” formation from the Wieluń Upland, as-

signed to the uppermost Platynota Zone and lower and middle parts of the Hypselocyclum Zone,

- marls, marly- and micritic limestones with common bivalves and locally occurring oncolites of the Burzenin Formation, which are assigned to the upper part of the Hypselocyclum and the Divisum Zone in the Wieluń Upland, and to almost the whole Hypselocyclum and the Divisum Zone north of the Wieluń Upland (Wierzbowski, 2017; Fig. 2).

The deposition of the Prusicko Formation in the Wieluń Upland took place on the outer slope of a carbonate ramp, in a relatively shallow environment, below the fair-weather wave base. Major shallowing, observed in the Wieluń Upland, is marked by occurrences of oolitic limestones belonging to the so-called “oolitic” formation, which are exposed in the Szczerców outcrop (Fig. 3). These strata are, in turn, overlain by transgressive deposits of the Burzenin Formation, partly exposed in the

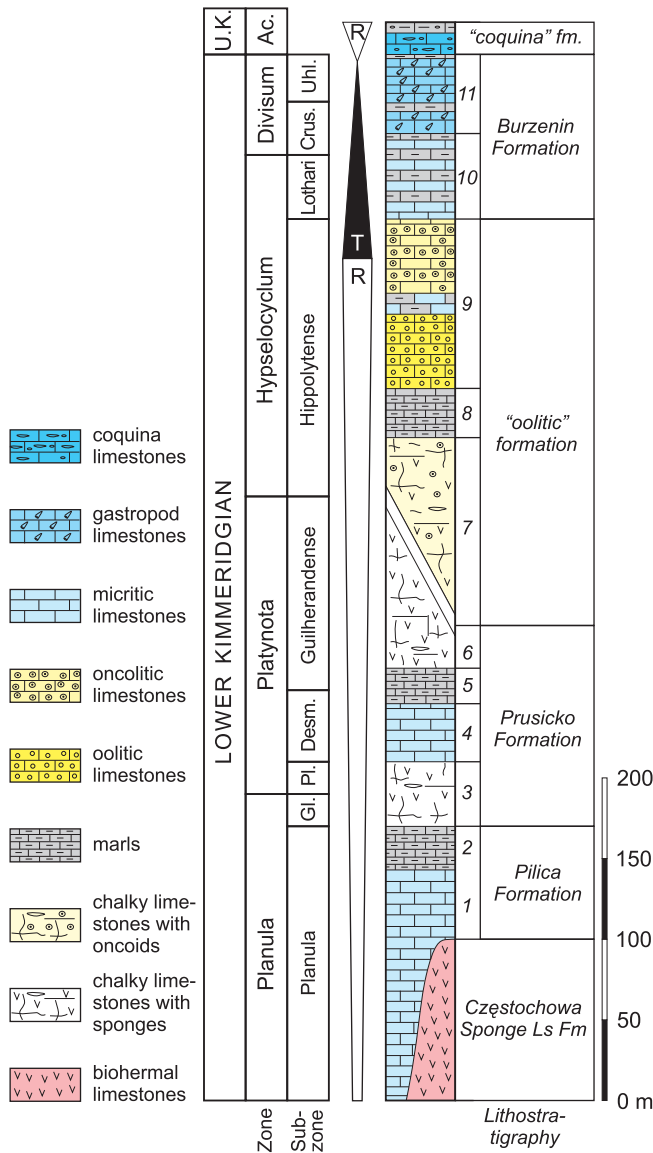


Fig. 2. Generalized lithology, lithostratigraphy, and transgressive-regressive sequences of the Wieluń Upland (Planula to Platynota–Hypselocyclum zone boundary) and its northern vicinity (Platynota–Hypselocyclum zone boundary to Acanthicum Zone; after Wierzbowski, 2017)

Ac. – Acanthicum, Crus. – Crussoliense, Desm. – Desmoides, Gl. – Galar, Pl. – Polygratus, Uhl. – Uhlandi; 1 – Wolbrom Limestone Member, 2 – Latosówka Marl Member, 3 – Kuchary Chalky Limestone Member, 4 – Skowronów Limestone Member, 5 – Góry Marl Member, 6 – Kule Chalky Limestone Member, 7 – unit A, 8 – Kielczygłów Marl Member (unit B), 9 – units C, D, E, F, 10 – Majaczewice Member (units G, H), 11 – Sarnów Gastropod Limestone Member (unit I)

Majaczewice section (Fig. 4), which were deposited in a moderately shallow sea after a decline of the shallow water carbonate platform (Wierzbowski, 2017). Deposition of the Burzenin Formation started earlier in a deeper part of the basin, north of the Wieluń Upland, directly above the moderately deep-water Pilica Formation, where the Prusicko and the “oolitic” formation do not occur. Part of the basal deposits of the Burzenin Formation, north of the Wieluń Upland, i.e. the Brzyków Oncolite Bed dated to the upper part of the Hippolytense Subzone of the Hypselo-

cyclum Zone was, however, formed in a relatively shallow environment (Wierzbowski, 2017). This may be treated as an indicator of the maximum regression surface of the lowstand systems tract, which corresponds to thicker oolite deposits from the south. A major part of the Burzenin Formation from the Wieluń Upland and its vicinity, with abundant condensation surfaces, shows a transgressive character and was deposited starting from the Lothari Subchron of the Hypselocyclum Chron to the end of the Divisum Chron (Wierzbowski, 2017). Younger Acanthicum (=Mutabilis) deposits of the Upper Kimmeridgian in the study area, which are not exposed nowadays, consist of bivalve coquinas (a so-called “coquina” formation) and show a slightly regressive character (Wierzbowski, 2017).

Oncolite-micritic limestones (unit 1) are exposed in the oldest part of the Rogaszyn section on the Radomsko Elevation. They are dated to the Desmoides Subzone of the Platynota Zone of the Lower Kimmeridgian (Kutek, 1968; Wierzbowski and Głowniak, 2018; see also Fig. 5). Younger strata in this section consist of intercalations of marls, micritic and subordinate organodetrital limestones (units 2–8) and represent the upper part of the Platynota Zone (Wierzbowski and Głowniak, 2018). Higher, above an omission surface, occur biodetrital limestones, which gradually pass into oncolitic limestones with oolites (unit 9). The latter strata are assigned to the Hippolytense Subzone of the Hypselocyclum Zone (Wierzbowski and Głowniak, 2018). In the youngest part of the Rogaszyn Quarry, which is poorly exposed nowadays, there occur brittle marls (unit 10) as well as micritic-oolitic limestones and marls (units 11 and 12) of the Lothari Subzone of the Hypselocyclum Zone (Wierzbowski and Głowniak, 2018).

A major part of the Rogaszyn section (units 1–11) consists of shallow water deposits of the carbonate platform. These deposits constitute a lateral equivalent of the “oolite” formation and the Prusicko Formation, known from the Wieluń Upland (Wierzbowski and Głowniak, 2018; Fig. 5). The very small thickness of the carbonate platform deposits in the Radomsko Elevation is, however, unusual, and may result from tectonic movements of local fault blocks (Wierzbowski and Głowniak, 2018). The demise of the shallow-water carbonate platform took place in the latest Hypselocyclum Chron (during the Lothari Subchron) and is marked by the occurrence of deeper water facies of unit 12 (Wierzbowski and Głowniak, 2018), which corresponds to the Burzenin Formation.

The Lower Kimmeridgian succession of the Małogoszcz section at the SW margin of the Holy Cross Mountains begins with the “oolite” formation assigned to the Platynota–Hypselocyclum zones (Kutek, 1968, 1994; Matyja et al., 2006; Matyja, 2011; see also Fig. 6). It consists of two oolitic units (the Lower Oolite and the Upper Oolite) underlain by Pelitic Limestones and divided by the Banded Limestone Member. These units are, in turn, overlain by the Oncolite Layer, the Oolite–Platy Member, as well as marls and marly limestones (Shaly Limestones and Underlying Shales). All the rocks of the “oolite” formation were formed during various phases of growth of a shallow-water carbonate platform (Matyja et al., 2006; Matyja, 2011). A younger part of the Małogoszcz section begins with an encrusted hardground around the Hypselocyclum–Divisum zone boundary. This hardground is overlain by the “coquina” formation, which consists of the Skorków Lumachelle and Upper Platy Limestones. Deposition of the “coquina” formation reflects a deepening of the basin during the Divisum Chron (Matyja et al., 2006; Matyja, 2011). The top of the Małogoszcz section consists of marls and marly shales with intercalations of organodetrital limestones (Top Shales), which are dated to the Upper Kimmeridgian Mutabilis (=Acanthicum) Zone (Kutek, 1968, 1994; Matyja et al., 2006; Matyja, 2011). These deposits

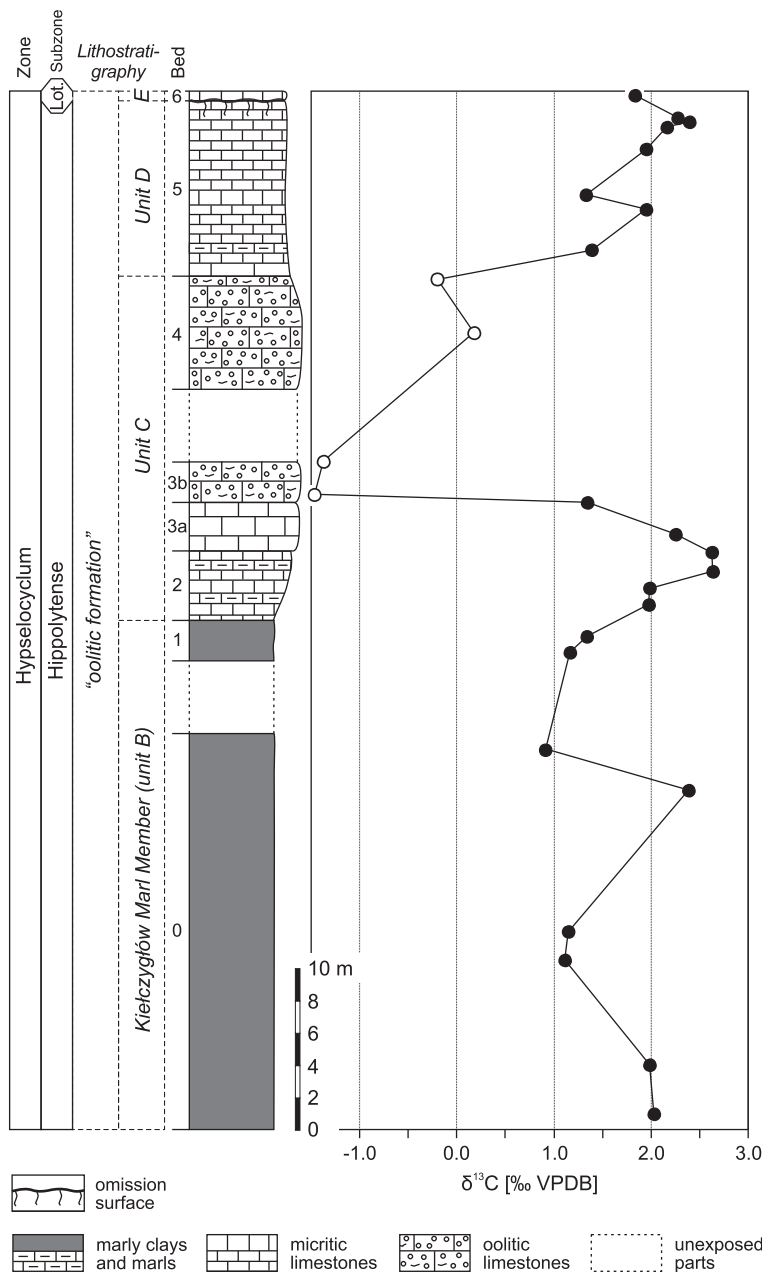


Fig. 3. Bulk carbonate carbon isotope record of the Szczerców section (northern border of the Wieluń Upland)

Bio- and lithostratigraphy are given after Wierzbowski (2017); data points of altered samples (see text) are marked with open circles; Lot. – Lothari Subzone

show a slightly regressive character, and are intercalated, especially in their higher part, with bivalve coquinas.

Although continuous limestone sedimentation prevailed in the Early Kimmeridgian of the study area, limestone beds are intercalated with marly layers. The appearance of the Latośówka Marl Member within the Pilica Formation, the Góry Marl Member within the Prusicko Formation and the Kiełczygłów Marl Member within coeval parts of the “oolitic” and the Burzenin formations as well as unnamed marls within the uppermost part of the Burzenin Formation from the Wieluń Upland and its vicinity, may be climatically or tectonically induced (Wierzbowski, 2017). The marlstone members from the Wieluń Upland correlate with coeval marly beds from the Radomsko Elevation

and the SW margin of the Holy Cross Mountains (cf. Matyja and Wierzbowski, 2006; Matyja, 2011; Wierzbowski, 2017). A deepening event on the Wieluń Upland, which is marked by the demise of carbonate platforms in the Lothari Subchron of the Hypselocyclum Chron, is almost coeval with the occurrence of deeper water facies on the Radomsko Elevation (Wierzbowski, 2017; Wierzbowski and Główniak, 2018). It predates, however, a similar episode known from the SW margin of the Holy Cross Mountains, which is assigned to the Hypselocyclum–Divisum Chron boundary (Kutek, 1994; Matyja et al., 2006; Matyja, 2011). Shallow-water carbonate sedimentation lasted longer at the SW margin of the Holy Cross Mountains probably because of the lesser depth of this part of the epicontinental Polish basin. In all areas studied, deposits of carbonate platforms are overlain by deeper water facies of the uppermost Lower and the Upper Kimmeridgian (Matyja and Wierzbowski, 2014).

MATERIAL AND METHODS

Deltoideum delta (oysters) and *Trichites* (pinnid bivalves), which are abundant in the Małogoszcz Quarry section (Fig. 6), were collected for isotope studies. A few undetermined oyster shells were also collected from the Małogoszcz and the Szczerców sections. Thin-sections prepared from the bivalve shells were studied using a cold cathodoluminescence microscope. Non-luminescent or very weakly luminescent shell fragments (38 samples), away from the hinge or the muscle scar areas, were cleaned manually, using a microdrill, from sediment remains and borings. The samples were ground in an agate mortar. Aliquots of carbonate powders were used for chemical and oxygen and carbon isotope analyses.

65 bulk carbonate samples (limestones, marly limestones, marls and marly clays) were collected manually from exposures at Szczerców, Majaczewice, and Rogaszyn, which are assigned to the Platynota Zone and a lower part of the Hypselocyclum Zone (Figs. 3–5). The sample set has been supplemented with 31 archival samples derived from W.C. Kowalski’s collection, who studied the Lower Kimmeridgian in exposures located north of the Wieluń Upland (cf. Kowalski, 1958) and from A. Wierzbowski’s collection, who mostly studied borehole cores from the northern border of the Wieluń Upland (cf. Wierzbowski, 2017). The majority of supplementary samples come from the vicinity of the Majaczewice section (localities: Góry Wapienne, Burzenin 47, Burzenin 16/9, Burzenin 16/2) and the Szczerców outcrop (boreholes: 37/13.5, PD20B, 130SP, PW408, KT109). Other localities are indicated on Figure 1.

Ca, Mg, Sr, Na, Mn and Fe concentrations in bivalve shells were determined by means of the ICP-OES (Inductively Coupled Plasma Optical Emission Spectrometry) method at the Polish Geological Institute – National Research Institute. 50–100 mg samples were dissolved in 5 wt.% hydrochloric acid. Reproducibility of chemical analyses (2σ S.D.) was controlled by multiple analyses of samples and averages as follows: 0.7% for Ca, 1.6% for Mg, 1.1% for Sr, 4.8% for Na, 2.9% for Mn, and 14% for Fe. Repeated analyses of JLs-1 calcite and JDo-1 do-

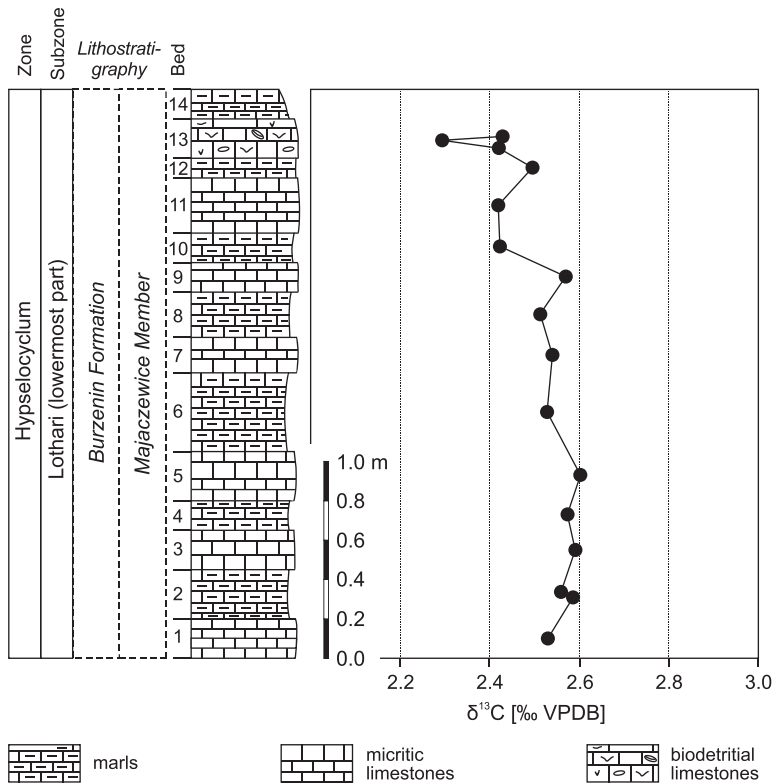


Fig. 4. Bulk carbonate carbon isotope record of the Majaczevice section (north of the Wieluń Upland)

Bio- and lithostratigraphy are given after Wierzbowski (2017)

lomite references (cf. Imai et al., 1996) yielded accuracies of measurements (2σ S.D.) better than 2.2% for Ca, 1.5% for Mg, 1.3% for Sr, 4.5% for Mn, and 12.1% for Fe. The presence of non-carbonate iron compounds may have affected reproducibility and accuracy of Fe analyses. The accuracy of Na analyses cannot be specified because of low sodium concentrations in the references.

Oxygen and carbon isotope analyses of bulk carbonates and bivalve shells were conducted at the GeoZentrum Nordbayern, University of Erlangen-Nuremberg (Germany). Samples were reacted with 100% phosphoric acid at 70°C using a *Gasbench II* connected to a *ThermoFisher Delta V Plus* mass spectrometer. All values are reported in per mil relative to the VPDB scale by assigning $\delta^{13}\text{C}$ values of +1.95‰ to NBS19 and -47.3‰ to IAEA-CO9 and $\delta^{18}\text{O}$ values of -2.20‰ to NBS19 and -23.2‰ to NBS18. Reproducibility of measurements was monitored by replicate analyses of laboratory standards Sol 2 ($n = 24$) and Erl 5 ($n = 18$). Reproducibility for $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ values was 0.09 and 0.10‰ ($\pm 2\sigma$ S.D.) for Sol 2, and 0.12‰ and 0.12‰ ($\pm 2\sigma$ S.D.) for Erl 5, respectively.

To calculate $\delta^{18}\text{O}$ -derived temperatures for calcite the relationship of O'Neil et al. (1969) modified by Friedman and O'Neil (1977), along with the SMOW to PDB scales conversion given by Friedman and O'Neil (1977), was used:

$$1000\ln\alpha_{\text{calcite-water}} = 2.78 \cdot 10^6/T^2 - 2.89 \quad [1]$$

where: $\alpha_{\text{calcite-water}}$ – oxygen isotope fractionation factor between calcite and water, T – temperature in Kelvin.

Temperatures calculated for the measured range of $\delta^{18}\text{O}_{\text{calcite}}$ values using the equation of Friedman and O'Neil

(1977) are almost indistinguishable from temperatures calculated using the equation of Anderson and Arthur (1983):

$$T(^{\circ}\text{C}) = 16.0 - 4.14 \cdot (\delta_c - \delta_w) + 0.13 \cdot (\delta_c - \delta_w)^2 \quad [2]$$

where: δ_c – oxygen isotope composition of carbonate on the PDB scale, δ_w – oxygen isotope composition of water on the SMOW scale.

Palaeotemperatures calculated using both equations are shown on Figure 6 for comparison.

DIAGENETIC ALTERATION OF BIVALVE SHELLS

Oyster shells possess foliated microstructures and are made of calcite (Esteban-Delgado et al., 2008). *Trichites* shells are primarily built of coarse calcite prisms (Chateigner et al., 2002). Although a thin inner, aragonitic layer occurs in the pointed area of modern pinnid shells its remnants are usually not found in fossil *Trichites* (cf. Chateigner et al., 2002; Sturman et al., 2014). The aragonite shell layer was also not observed in the *Trichites* shells studied.

Diagenetic alteration of skeletal calcite can be screened using cathodoluminescence studies and analyses of minor and trace element concentrations. Iron and manganese contents of marine calcite increase as a result of alteration under reducing conditions. A diagenetic decrease in strontium content is, in turn, observed because of its low concentration in freshwater and lesser partitioning of this element in inorganic calcites (Veizer, 1983; Brand and Veizer, 1980; Marshall, 1992; Ullmann and Korte, 2015). In addition, diagenetic Mn^{2+} ions activate bright orange-red cathodoluminescence in calcites, which is typical of altered shell material (Marshall, 1992; Savard et al., 1995). Pristine carbonate shells may, however, show dull luminescence with narrow brighter bands (Barbin, 2000, 2013).

The oyster and *Trichites* shells studied from the Lower Kimmeridgian of central Poland are non-luminescent or show different cathodoluminescence intensities from dull to bright (Fig. 7). As bright, orange-red luminescence is interpreted as a result of diagenetic alteration, only non-luminescent or dull luminescent bivalve shells were selected for chemical and isotope analyses.

The bivalve shells studied from central Poland are characterized by variable minor and trace element concentrations: Mn from below detection limit (1 ppm) to 74 ppm, Fe from below detection limit (20 ppm) to 596 ppm, and Sr from 592 to 860 ppm (Table 1). A strong correlation is observed between Fe and Mn contents in the shells (Fig. 8). Threshold limits of $\text{Mn} \leq 100$ ppm, $\text{Fe} \leq 250$ ppm, and $\text{Sr} \geq 490$ ppm may be accepted as indicative of well-preserved Jurassic calcite bivalve shells according to Wierzbowski and Joachimski (2007), and Wierzbowski et al. (2016). The given limits of manganese and iron concentrations are in line with the data of Jones et al. (1994). The accepted manganese and strontium cut-off limits are also similar to the limits specified for Jurassic bivalves by Anderson et al. (1994), Price and Page (2008), Price and Teece (2010), and Zuo et al. (2019) although the latter authors report higher iron contents.

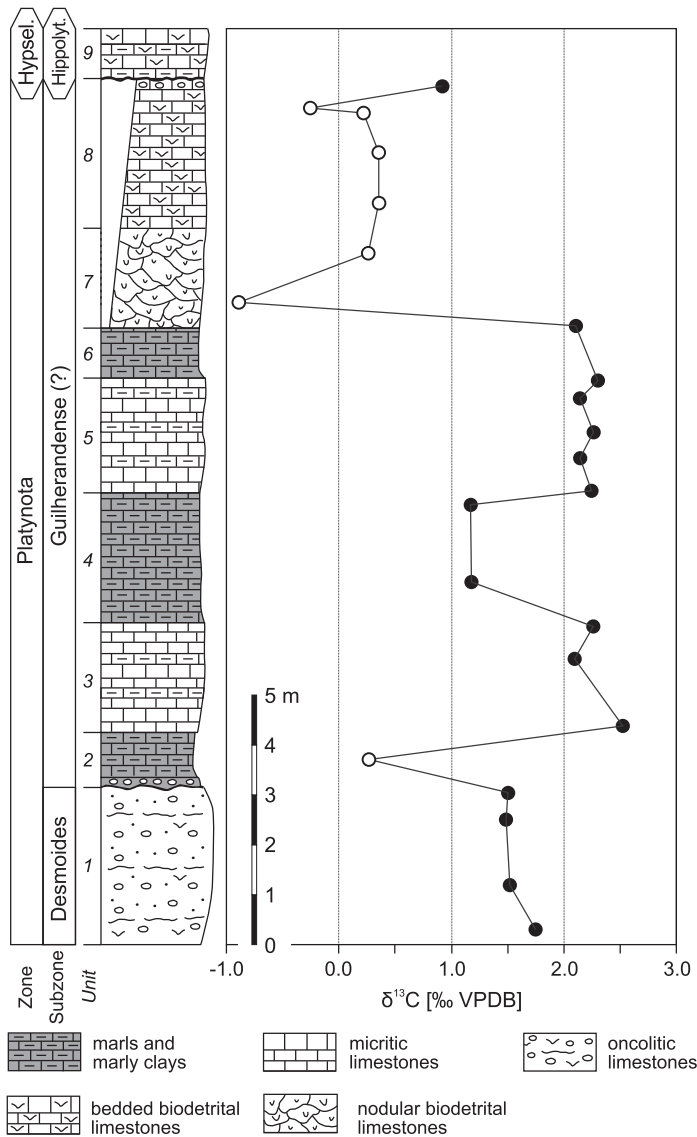


Fig. 5. Bulk carbonate carbon isotope record of the Rogaszyn section (the Radomsko Elevation)

Biostratigraphy and lithological units are given after [Wierzbowski and Główniak \(2018\)](#); data points of altered samples (see text) are marked with open circles; Hypsel. – Hypselocyclus Zone, Hippolyt. – Hippolytense Subzone

Six bivalve shells with Fe contents above 250 ppm were removed from the well-preserved sample set. After removal of these samples the remaining sample set is characterized by concentrations of Mn \leq 27 ppm, Fe \leq 206 ppm, and Sr \geq 631 ppm, and a moderate correlation between Fe and Mn contents ([Fig. 8](#)). All well-preserved bivalve shells (32 samples) come from the Małogoszcz section.

RESULTS

OXYGEN AND CARBON ISOTOPE COMPOSITION OF BIVALVE SHELLS

$\delta^{18}\text{O}$ values of well-preserved bivalve shells from the Małogoszcz section vary between -0.6 and -3.5‰ ([Fig. 6](#) and [Table 1](#)). Oxygen isotope values from the Upper Oolite and the

Skorków Lumachelle (the Hypselocyclus and the lowermost Divisum zones) are characterized by a significant scatter (from -3.5 to -1.5‰). Bivalve $\delta^{18}\text{O}$ values from the Upper Platy Limestones and Top Shales (the Divisum–Mutabilis zone boundary) show a lesser spread (from -3.4 to -2.2‰). Two oxygen isotope data points from two oysters from the upper part of the Skorków Lumachelle (the middle Divisum Zone) are higher (-1.0 to -0.6‰) than all other data. No clear temporal trend is observed within the oxygen isotope dataset.

$\delta^{13}\text{C}$ values of well-preserved bivalve shells from the Małogoszcz section vary between 2.6 and 4.2‰ ([Fig. 6](#) and [Table 1](#)). Carbon isotope values from the Upper Oolite and the Skorków Lumachelle (the Hypselocyclus and the lowermost Divisum zones) are lower and show a larger scatter (from 2.6 to 4.0‰) than the values from the Upper Platy Limestones and Top Shales (the Divisum–Mutabilis zone boundary), which vary between 3.6 to 4.2‰ . A gradual increase in bivalve $\delta^{13}\text{C}$ values upsection is observed, except for two low $\delta^{13}\text{C}$ values from the upper part of the Skorków Lumachelle (the middle Divisum Zone).

CARBON ISOTOPE COMPOSITION OF BULK CARBONATES

Bulk carbonate $\delta^{13}\text{C}$ values range from -1.5 to 2.7‰ in the entire study interval. The most variable carbon isotope values are documented from the Szczerców (-1.5 to 2.6‰) and the Rogaszyn section (-0.9 to 2.5‰ ; [Fig. 9](#)). Lesser variations of $\delta^{13}\text{C}$ values are noted in the samples derived from bore-holes (0.0 to 2.5‰), small exposures of the Lower Kimmeridgian deposits (1.1 to 2.7‰) and from the Majaczewice section (2.3 to 2.6‰). There is no distinct difference in $\delta^{13}\text{C}$ values of co-occurring limestone and marly beds.

Strong correlations between bulk carbonate $\delta^{18}\text{O}$ values and $\delta^{13}\text{C}$ values, which may be linked to diagenetic alteration, is observed in the datasets from the Rogaszyn and the Szczerców section ([Fig. 9](#)). Since diagenetic processes may have affected the original carbon isotope composition of the highly weathered subsurface or oolitic parts of these sections, they are discussed later in the text.

DISCUSSION

OXYGEN AND CARBON ISOTOPE RECORDS OF BIVALVE SHELLS

Modern oysters precipitate shell calcite in the oxygen isotope equilibrium with ambient sea water or very close to it ([Hong et al., 1995](#); [Kirby et al., 1998](#); [Surge et al., 2001, 2003](#); [Titschack et al., 2010](#); [Ullmann et al., 2010](#)). The equilibrium precipitation of oxygen isotopes by Jurassic oysters is documented by the similarity of their $\delta^{18}\text{O}$ values with those of co-occurring benthic and necto-benthic molluscs and brachiopods ([Anderson et al., 1994](#); [Wierzbowski and Joachimski, 2007](#); [Price and Teece, 2010](#); [Mettam et al., 2014](#)). Although slight disequilibrium fractionation of carbon isotopes (of $\sim 1.0\text{‰}$, except shell portions precipitated

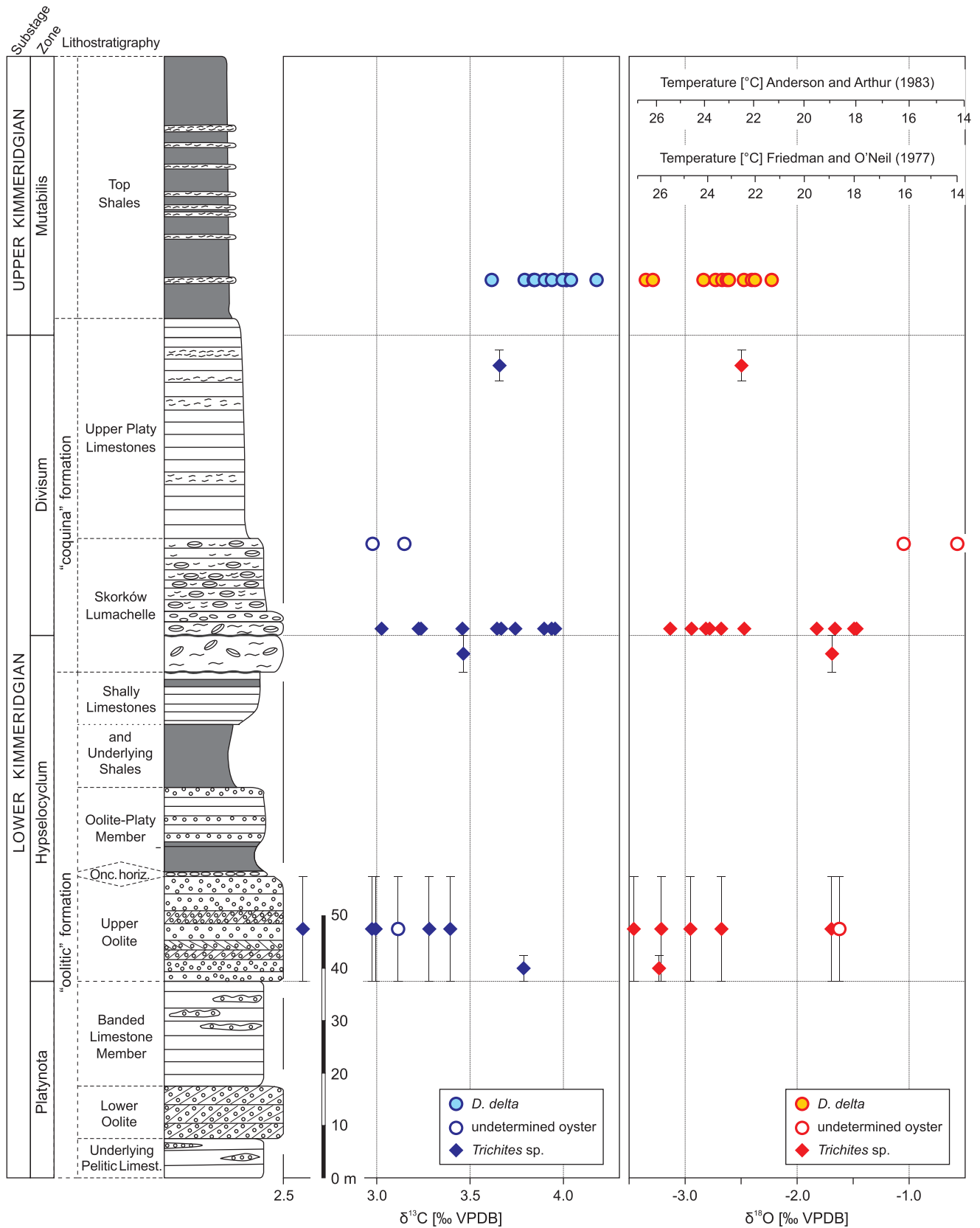


Fig. 6. Oxygen and carbon isotope records of the Małogoszcz section (SW margin of the Holy Cross Mountains)

Bio- and lithostratigraphy are given after Matyja et al. (2006), Matyja (2011) and A. Wierzbowski's (pers comm., 2017)

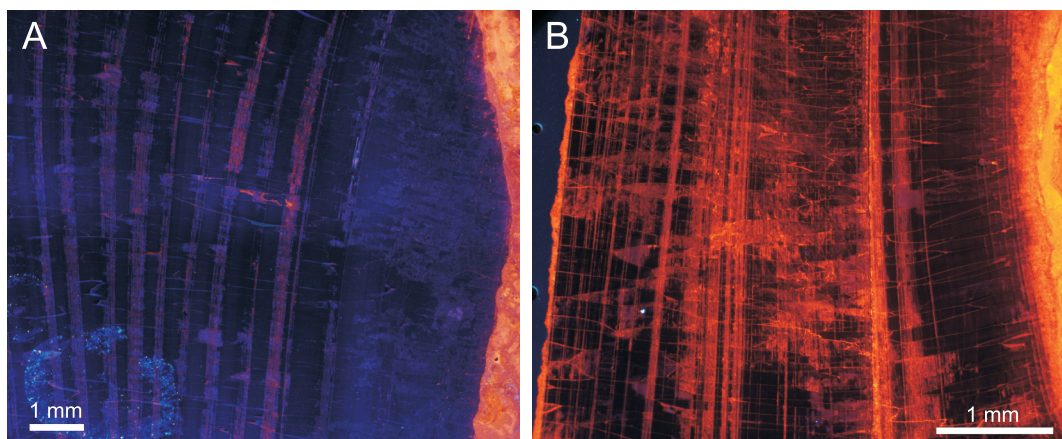


Fig. 7. Cathodoluminescence photomicrographs

A – well-preserved, dull-luminescent *Trichites* shell (sample M80, Upper Oolite, Hypselocyclus Zone, Małogoszcz section); **B** – altered oyster shell, showing moderate to bright luminescence intensities (sample M41, bed 5, Hypselocyclus Zone, Szczerców section)

during cold seasons) is observed in modern oysters (Surge et al., 2001, 2003; Titschack et al., 2010), Jurassic oysters show higher $\delta^{13}\text{C}$ values than those of coeval molluscs (Wierzbowski and Joachimski, 2007; Price and Teece, 2010; Mettam et al., 2014). Accordingly, it may be assumed that the Jurassic oysters exerted relatively small vital effects in carbon isotope composition. In addition, Brigaud et al. (2009), Lathuilière et al. (2015), and Zuo et al. (2019) have recently shown that isotope signatures of Upper Jurassic *Trichites* are similar to those of coeval oysters and may be used as a reliable proxy for ancient water temperatures.

The oxygen isotope composition of marine calcite, precipitated in isotopic equilibrium with ambient sea water, is dependent on the temperature and oxygen isotope composition of ambient water. $\delta^{13}\text{C}$ values of calcite precipitated in isotopic equilibrium vary, in turn, along with primary fluctuations in the composition of dissolved inorganic carbon (DIC).

Relatively scattered $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of bivalve shells from the lower part of the Małogoszcz section i.e., the Upper Oolite and a lower part of the Skorków Lumachelle (the Hypselocyclus and the lowermost Divisum zones; Fig. 6) may indicate considerable variations in water temperatures, salinity and the carbon isotope composition of DIC. Early Kimmeridgian carbonate platforms from the SW margin of the Holy Cross Mountains grew in a shallow basin, above storm wave-base (Matyja, 2011). Significant freshwater inflow from land areas or an enhanced evaporation rate may have caused temporal variations in $\delta^{18}\text{O}_{\text{water}}$ values. River water rich in terrestrial organic matter or oxidation of marine organic carbon during the process of water-mass ageing (cf. Patterson and Walter, 1994) may, in turn, have produced variations in the carbon isotope composition of DIC. More densely clustered $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ data points of bivalves from the upper part of the Małogoszcz section i.e. the Upper Platy Limestones and the Top Shales (the Divisum–Mutabilis zone boundary) probably indicate more stable water temperature and chemistry, after a sea level rise and drowning of the shallow water carbonate platform (cf. Matyja et al., 2006; Matyja, 2011). Although $\delta^{13}\text{C}$ values of two oysters from the upper part of the Skorków Lumachelle are relatively low, a general increase in mean bivalve $\delta^{13}\text{C}$ value (by $\sim 0.5\text{‰}$) towards the top of the Małogoszcz section is observed. This may be linked to a diminishing inflow of freshwater enriched in the light $\delta^{12}\text{C}$ isotope and a higher rate of water exchange and circulation.

Palaeotemperatures calculated from oxygen isotope ratios of bivalves from the Upper Oolite and the lower part of the Skorków Lumachelle (the Hypselocyclus and the lowermost Divisum zones), assuming normal marine salinity and a mean $\delta^{18}\text{O}_{\text{seawater}}$ value of -1‰ VSMOW as typical of an ice-free world (Shackleton and Kennett, 1975), vary between 18 and 27°C (Fig. 6). Two outlier data points from the uppermost part of the Skorków Lumachelle (Divisum Zone) translate into lower temperatures of 14–16°C. Temperatures calculated for the Upper Platy Limestones and the Top Shales (the Divisum–Mutabilis zone boundary) range from 21 to 27°C.

Most of the calculated average water temperatures seem to be overestimated taking into account the palaeolatitude (38°N) of central Poland in the Kimmeridgian (cf. van Hinsbergen et al., 2015) as well as the Late Jurassic palaeoclimatic model of Sellwood and Valdes (2008) and the Early Kimmeridgian temperature ($\sim 24^\circ\text{C}$) reconstructed for tropical regions (Alberti et al., 2017). The assumption of a mean Kimmeridgian surface temperature of $\sim 18^\circ\text{C}$ inferred for the area of present-day Poland after Sellwood and Valdes (2008) allows estimation of the original $\delta^{18}\text{O}_{\text{water}}$ values. Although temperature usually drops downwards in the water column (cf. Manca et al., 2004), freshwater influx occurs more often in very shallow environments, where the thermal stratification is negligible. The measured range of bivalve $\delta^{18}\text{O}$ values (-3.5 to -0.6‰) translates into $\delta^{18}\text{O}_{\text{water}}$ values from -3.0 to -0.1‰ VSMOW using the palaeotemperature equation of Friedman and O'Neil (1977). Water salinity (S_{water}) can, in turn, be calculated on the basis of the universal marine salinity– $\delta^{18}\text{O}_{\text{water}}$ relationship of Railsback et al. (1989), which takes into account effects of freshwater runoff [3] and evaporation [4]. The two given equations were used for salinity calculations below and above the normal marine salinity respectively.

$$\delta^{18}\text{O}_{\text{water}} = \delta^{18}\text{O}_0 + \Delta_{\text{fw}}/S_0 \cdot (S_{\text{water}} - S_0) \quad [3]$$

where: $\delta^{18}\text{O}_0$ – average oxygen isotope composition of Jurassic sea water (assumed as -1‰ VSMOW after Shackleton and Kennett, 1975), Δ_{fw} – difference between $\delta^{18}\text{O}$ values of average sea water and local meteoric water (assumed as -6‰ VSMOW using the modern relationship between annual mean $\delta^{18}\text{O}$ values of precipitation and surface air temperature; cf. Rozanski et al., 1993), and S_0 – average sea water salinity (assumed as 34‰ after Railsback et al., 1989).

Table 1

Position, chemical and isotope data of bivalve shells from the Małogoszcz section

No.	Taxonomy	Unit	Position [m]	Position error [+/-] m	Ca [%]	Fe [ppm]	Mg [ppm]	Mn [ppm]	Na [ppm]	Sr [ppm]	$\delta^{13}\text{C}$ [‰]	$\delta^{18}\text{O}$ [‰]
M1*	oyster	Top Shales	206.5	1.5	38.3	596*	333	74	499	617	–	–
M70	<i>D. delta</i>	Top Shales	171.3	1.2	38.4	104	551	17	1017	709	3.62	–3.35
M59	<i>D. delta</i>	Top Shales	171.3	1.2	39.4	114	500	21	796	675	3.84	–2.63
M57	<i>D. delta</i>	Top Shales	171.3	1.2	39.1	92	499	18	683	690	4.00	–2.61
M56	<i>D. delta</i>	Top Shales	171.3	1.2	38.8	171	486	17	916	642	4.02	–2.23
M54*	<i>D. delta</i>	Top Shales	171.3	1.2	36.6	509*	1422	59	629	592	–	–
M53	<i>D. delta</i>	Top Shales	171.3	1.2	38.6	206	553	22	692	653	3.94	–2.47
M52	<i>D. delta</i>	Top Shales	171.3	1.2	38.6	152	542	14	886	687	3.90	–3.29
M51	<i>D. delta</i>	Top Shales	171.3	1.2	39.2	86	397	10	1018	694	4.04	–2.84
M36	<i>D. delta</i>	Top Shales	171.3	1.2	39.5	71	421	18	751	658	4.02	–2.67
M35	<i>D. delta</i>	Top Shales	171.3	1.2	39.8	74	429	10	651	631	3.79	–2.41
M33	<i>D. delta</i>	Top Shales	171.3	1.2	39.5	112	489	27	707	711	4.18	–2.38
M15	<i>D. delta</i>	Top Shales	171.3	1.2	39.7	77	443	19	734	636	3.85	–2.73
M7	<i>Trichites</i>	Up. Platy Ls	155	3	37.7	71	419	14	1018	721	3.66	–2.50
M69	oyster	Skork. Lum.	121	1.2	39.1	109	797	5	548	649	2.98	–0.57
M9	oyster	Skork. Lum.	121	1.2	39.4	38	569	1	399	684	3.15	–1.05
M73*	oyster	Skork. Lum.	104.8	1.3	38.7	464*	2920	17	1483	860	–	–
M65*	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.7	253*	1723	6	2028	729	–	–
M61*	<i>Actinostreon</i>	Skork. Lum.	104.8	1.3	38.5	558*	1896	24	555	650	–	–
M31	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.6	195	1679	5	2321	786	3.96	–2.94
M28	<i>Trichites</i>	Skork. Lum.	104.8	1.3	39.2	119	1664	4	2405	819	3.94	–2.81
M25	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.6	120	1578	5	2374	740	3.03	–1.66
M24	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.7	73	1678	2	2308	792	3.64	–2.68
M21	<i>Trichites</i>	Skork. Lum.	104.8	1.3	39.3	120	1551	4	2362	776	3.74	–2.78
M20	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.5	74	1702	2	2167	721	3.22	–1.49
M17	<i>Trichites</i>	Skork. Lum.	104.8	1.3	39.3	192	1528	6	2368	769	3.90	–3.13
M12	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.7	61	1371	1	2495	740	3.24	–2.47
M5	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.7	121	1690	3	2232	756	3.46	–1.82
M3	<i>Trichites</i>	Skork. Lum.	104.8	1.3	38.7	47	1779	2	2256	715	3.67	–1.47
M2*	oyster	Skork. Lum.	104.8	1.3	39.4	457*	1427	21	817	637	–	–
M18	<i>Trichites</i>	Skork. Lum.	100	3.5	38.6	170	1728	3	2279	761	3.46	–1.69
M81	<i>Trichites</i>	Up. Oolite	47.5	10	38.9	<20	1305	<1	2342	688	3.28	–1.69
M80	<i>Trichites</i>	Up. Oolite	47.5	10	35.5	<20	1246	1	1834	690	2.97	–3.21
M50	oyster	Up. Oolite	47.5	10	38.9	25	586	2	234	715	3.11	–1.62
M48	<i>Trichites</i>	Up. Oolite	47.5	10	38.6	<20	1284	<1	2568	805	3.39	–2.68
M47	<i>Trichites</i>	Up. Oolite	47.5	10	39.3	<20	1467	5	1928	734	2.60	–2.95
M44	<i>Trichites</i>	Up. Oolite	47.5	10	39.4	<20	1273	2	1315	761	2.99	–3.46
M77	<i>Trichites</i>	Up. Oolite	40	2.5	37.9	<20	1395	<1	2374	752	3.79	–3.23

*– altered samples showing elevated Fe concentrations

$$\delta^{18}\text{O}_{\text{water}} = \delta^{18}\text{O}_0 + m_0 \cdot (S_{\text{water}} - S_0) \quad [4]$$

where: m_0 – evaporative enrichment parameter being the relation between $\delta^{18}\text{O}$ and salinity observed in evaporative basins (assumed as 0.35‰ VSMOW per 1 salinity per mil according to Railsback et al., 1989), and other symbols are the same as in eq. [2].

The calculated sea water $\delta^{18}\text{O}$ values translate into a salinity range of 21 to 37‰. Although the modelling does not allow for possible changes in depth of the basin and temporal climate fluctuations, it points to the possibility of occurrence of episodes of decreased salinity in the basin studied. Low and variable $\delta^{18}\text{O}$ values of Lower Kimmeridgian–lowermost Upper

Kimmeridgian fossils from the SW margin of the Holy Cross Mountains may be regarded as a manifestation of major shallowing of the epicratonic Polish basin, being an effect of prolonged marine regression, which is well-documented starting from the Upper Oxfordian (cf. Matyja, 1977, 2011; Kutek, 1994; Matyja et al., 2006; Matyja and Wierzbowski, 2014; Wierzbowski, 2017). The bathymetry and salinity changes connected with this sea level fall likely masked subtle climatic and environmental fluctuations in central Poland. Although some authors have postulated the presence of such variations in the Kimmeridgian e.g. a cooling at the Oxfordian–Kimmeridgian transition (Abbink et al., 2001; Zou et al., 2019), a warming of deep sea water at the Early–Late Kimmeridgian transition

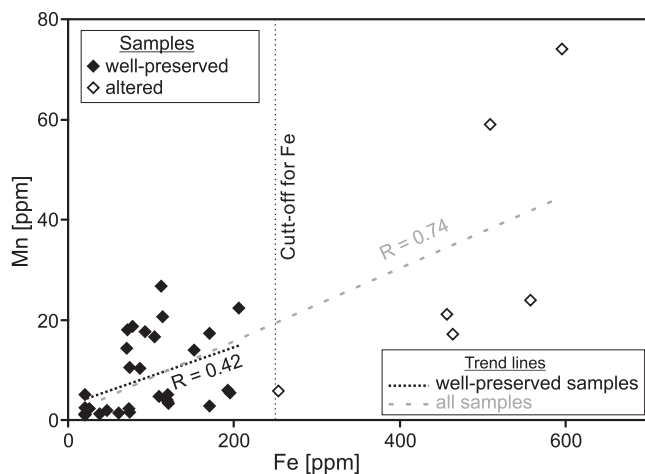


Fig. 8. Cross-plot of Fe and Mn concentrations in the bivalve shells studied

Cut-off limit of 250 ppm of Fe is accepted for well-preserved samples; a strong correlation ($R = 0.74$, $p < 0.01$) is observed in the entire dataset, and a moderate one ($R = 0.42$, $p < 0.05$) in well-preserved samples

(Colombié et al., 2018) or a slight warming throughout almost the whole Kimmeridgian (Zuo et al., 2019), they are not observed in the study area (Fig. 6; see also Wierzbowski, 2015).

A gradual decrease in belemnite and brachiopod $\delta^{18}\text{O}$ values (from -0 to -1‰ VPDB) is observed in central Poland throughout the Oxfordian (Wierzbowski, 2015). This decrease is attributed to a temperature rise and/or increased freshwater runoff during the onset of the marine regression. The oxygen and carbon isotope values of Lower Kimmeridgian bivalves from central Poland (-3.5 to -0.6‰ , and 2.6 to 4.2‰ , respectively) are comparable to those from shallow-marine facies of the Paris Basin, Burgundy, the Jura Mountains, and the Lower Saxony basin (cf. Brigaud et al., 2008; Lathuilière et al., 2015; Colombié et al., 2018; Zuo et al., 2019). Low $\delta^{18}\text{O}$ values of Lower Kimmeridgian fossils from western Europe are likely related to shallowing of epicontinental marine basins. A similar phenomenon and a salinity-related decrease is also reported for the Early Kimmeridgian basin of the Russian Platform (Wierzbowski et al., 2018). Salinity and thermobathymetry effects on fossil $\delta^{18}\text{O}$ values at European localities may also be an alternative explanation of such phenomena as weak latitudinal temperature gradients or thermal homogenization of sea water postulated for the Kimmeridgian (see Alberti et al., 2017; Colombié et al., 2018).

Higher $\delta^{18}\text{O}$ values (-0.5 to 0.5‰) are reported again for the mid-Tithonian oysters from central Poland (Wierzbowski et al., 2016). The mid-Tithonian fossils are derived from open marine deposits belonging to the Pałuki Formation and the lowermost part of the Kcynia Formation (Dembowska, 1979; Kutek and Zeiss, 1997; Matyja and Wierzbowski, 2016; Wierzbowski et al., 2016). This may be related to a successive deepening of the mid-Polish sedimentary basin during a Late Kimmeridgian transgression, which began in the Eudoxus Chron (cf. Kutek, 1994; Matyja and Wierzbowski, 2014; Wierzbowski, 2017). All the data show that the Upper Jurassic oxygen isotope record of central Poland was mainly controlled by sea level changes.

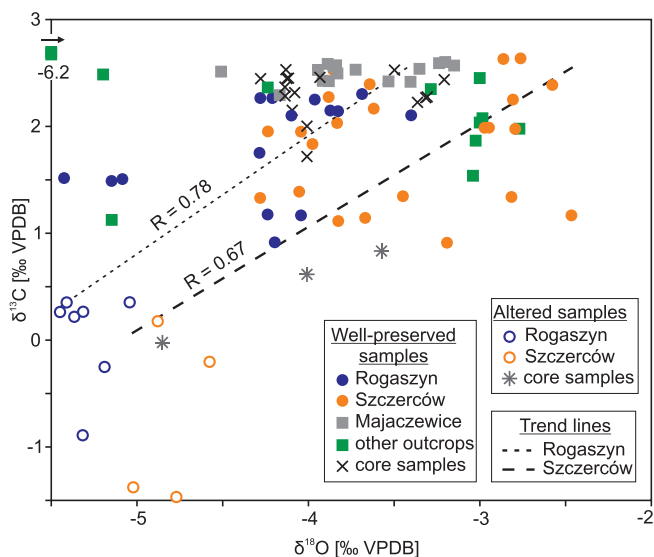


Fig. 9. Cross-plot of $\delta^{13}\text{C}$ versus $\delta^{18}\text{O}$ values of bulk carbonates from the Lower Kimmeridgian sections studied

Strong and statistically significant correlations between $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values are observed in the Rogaszyn ($R = 0.78$, $p < 0.01$) and Szczerców ($R = 0.67$, $p < 0.01$) datasets

CARBON ISOTOPE RECORD OF BULK CARBONATES

The bulk rocks studied are derived from deeper parts of the Kimmeridgian basin of central Poland i.e. the Wieluń Upland, its northern vicinity and the Radomsko elevation. Low $\delta^{13}\text{C}$ values of bulk carbonates (-1.5 to 0.4‰) are observed within the highly weathered and abandoned upper part of the Rogaszyn section, and porous oolitic limestones from the Szczerców section (Figs. 3 and 5). A low $\delta^{13}\text{C}$ value (0.3‰) is also observed in proximity to the oncolite layer in the lower part of the Rogaszyn Quarry. Strong correlation observed between $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ values from these localities (Fig. 9) may be linked to the post-depositional equilibration of rocks, sensitive to diagenesis, with meteoric waters in the subsurface vadose or phreatic zone (cf. Jenkyns and Clayton, 1986; Banner and Hanson, 1990; Marshall, 1992; Huck et al., 2013; Jach et al., 2014; Swart and Oehlert, 2018). Since other parts of the sections studied show more stable $\delta^{13}\text{C}$ values (0.9 to 2.6‰), which are not noticeably correlated with $\delta^{18}\text{O}$ values (Fig. 9), one can assume that their original marine carbon isotope composition is still preserved. Moderately scattered $\delta^{13}\text{C}$ values (0.0 to 2.6‰) are observed within borehole core samples (Fig. 9). The majority of these samples have, however, high $\delta^{13}\text{C}$ values (above 0.9‰) and no correlation between their $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values is observed. Distinctly lower $\delta^{13}\text{C}$ values (0.0 to 0.8‰) are seen in only three samples from a narrow interval (66.8 – 72.0 m) of a borehole core 6W (Table 2; see also Wierzbowski, 2017). As coeval samples from other borehole cores and the Majaczewice section show much higher $\delta^{13}\text{C}$ values (Fig. 10) this interval of the core 6W is interpreted as diagenetically altered.

Table 2

Stratigraphical and isotope data of bulk-carbonates

No.	Lithology	Outcrop	Bed/depth	Zone	Subzone	Position in subzone	$\delta^{13}\text{C}$ [‰]	$\delta^{18}\text{O}$ [‰]
KL 66	ls	Majaczewice	13	Hypselocyclum	Lothari	0.20	2.43	-3.92
KL 76	ls	Majaczewice	13	Hypselocyclum	Lothari	0.20	2.29	-4.17
KL 71	ls	Majaczewice	13	Hypselocyclum	Lothari	0.20	2.42	-3.53
KL 74	mrl	Majaczewice	12	Hypselocyclum	Lothari	0.19	2.50	-3.83
KL 63	ls	Majaczewice	11	Hypselocyclum	Lothari	0.18	2.42	-3.40
KL 61	mrl	Majaczewice	10	Hypselocyclum	Lothari	0.16	2.42	-3.88
KL 60	ls	Majaczewice	9	Hypselocyclum	Lothari	0.15	2.57	-3.15
KL 86	mrl	Majaczewice	8	Hypselocyclum	Lothari	0.13	2.51	-4.51
KL 78	ls	Majaczewice	7	Hypselocyclum	Lothari	0.12	2.54	-3.35
KL 62	mrl	Majaczewice	6	Hypselocyclum	Lothari	0.10	2.53	-3.94
KL 65	ls	Majaczewice	5	Hypselocyclum	Lothari	0.07	2.60	-3.20
KL 77	mrl	Majaczewice	4	Hypselocyclum	Lothari	0.06	2.57	-3.84
KL 59	ls	Majaczewice	3	Hypselocyclum	Lothari	0.04	2.59	-3.24
KL 84	mrl	Majaczewice	2	Hypselocyclum	Lothari	0.03	2.56	-3.85
KL 80	mrl	Majaczewice	2	Hypselocyclum	Lothari	0.02	2.58	-3.89
KL 72	ls	Majaczewice	1	Hypselocyclum	Lothari	0.01	2.53	-3.73
KL 1	ls	Szczerców	6	Hypselocyclum	Lothari	0.04	1.83	-3.98
KL 24	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.98	2.28	-3.88
KL 25	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.98	2.40	-3.64
KL 23	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.98	2.17	-3.62
KL 10	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.96	1.95	-4.24
KL6	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.91	1.33	-4.28
KL 15	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.90	1.95	-4.04
KL 4	ls	Szczerców	5	Hypselocyclum	Hippolytense	0.86	1.39	-4.05
KL 11*	ool ls	Szczerców	4	Hypselocyclum	Hippolytense	0.83	-0.20*	-4.58*
KL 8*	ool ls	Szczerców	4	Hypselocyclum	Hippolytense	0.79	0.18*	-4.88*
KL 5*	ool ls	Szczerców	3b	Hypselocyclum	Hippolytense	0.67	-1.38*	-5.02*
KL 50*	ool ls	Szczerców	3b	Hypselocyclum	Hippolytense	0.64	-1.47*	-4.77*
KL 19	ls	Szczerców	3a	Hypselocyclum	Hippolytense	0.63	1.35	-3.45
KL 12	ls	Szczerców	3a	Hypselocyclum	Hippolytense	0.60	2.25	-2.81
KL 22	ls/mrl	Szczerców	2	Hypselocyclum	Hippolytense	0.58	2.63	-2.86
KL 3	ls/mrl	Szczerców	2	Hypselocyclum	Hippolytense	0.56	2.64	-2.76
KL 20	ls/mrl	Szczerców	2	Hypselocyclum	Hippolytense	0.55	1.99	-2.97
KL 21	ls/mrl	Szczerców	2	Hypselocyclum	Hippolytense	0.53	1.98	-2.79
KL 13	mrl cl	Szczerców	1	Hypselocyclum	Hippolytense	0.50	1.34	-2.82
KL 17	mrl cl	Szczerców	1	Hypselocyclum	Hippolytense	0.49	1.17	-2.47
KL 16	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.40	0.91	-3.19
KL 7	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.36	2.39	-2.58
KL 2	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.23	1.14	-3.67
KL9	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.20	1.11	-3.83
KL 18	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.11	1.99	-2.95
KL 14	mrl cl	Szczerców	0	Hypselocyclum	Hippolytense	0.06	2.03	-3.83
KL 87	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.99	0.91	-4.20
KL 89*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.96	-0.25*	-5.19*
KL 93*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.95	0.22*	-5.37*
KL 94*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.89	0.35*	-5.04*
KL 88*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.82	0.35*	-5.41*
KL 92*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.75	0.26*	-5.45*
KL 91*	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.68	-0.89*	-5.32*
KL 53	ls	Rogaszyn	7/8	Platynota	Guilherandense	0.65	2.10	-3.40
KL 83	ls	Rogaszyn	5	Platynota	Guilherandense	0.57	2.30	-3.69
KL 75	ls	Rogaszyn	5	Platynota	Guilherandense	0.55	2.14	-3.83

Tab. 2 cont.

No.	Lithology	Outcrop	Bed/depth	Zone	Subzone	Position in subzone	$\delta^{13}\text{C}$ [‰]	$\delta^{18}\text{O}$ [‰]
KL 52	ls	Rogaszyn	5	Platynota	Guilherandense	0.50	2.27	-4.28
KL 73	ls	Rogaszyn	5	Platynota	Guilherandense	0.46	2.15	-3.87
KL 54	ls	Rogaszyn	5	Platynota	Guilherandense	0.42	2.25	-3.96
KL 85	mrl	Rogaszyn	4	Platynota	Guilherandense	0.40	1.17	-4.04
KL 51	mrl	Rogaszyn	4	Platynota	Guilherandense	0.29	1.18	-4.24
KL 82	ls	Rogaszyn	3	Platynota	Guilherandense	0.23	2.27	-4.21
KL 69	ls	Rogaszyn	3	Platynota	Guilherandense	0.18	2.10	-4.10
KL 79	ls	Rogaszyn	3	Platynota	Guilherandense	0.09	2.53	-3.85
KL 64*	mrl cl	Rogaszyn	2	Platynota	Guilherandense	0.04	0.26*	-5.31*
KL 68	ls	Rogaszyn	1	Platynota	Desmoides	0.97	1.51	-5.09
KL 67	ls	Rogaszyn	1	Platynota	Desmoides	0.82	1.49	-5.15
KL 81	ls	Rogaszyn	1	Platynota	Desmoides	0.45	1.52	-5.43
KL 70	ls	Rogaszyn	1	Platynota	Desmoides	0.19	1.75	-4.29
KL 34	mrl	core 37/13.5	(69 m)	Hypselocyclus	Lothari	0.69	1.72	-4.01
KL 33	mrl	core PD20B	(175–176 m)	Hypselocyclus	Lothari	0.55	2.32	-4.08
KL 35	mrl	core 130SP	(127–137 m)	Hypselocyclus	Lothari	0.54	2.37	-4.14
KL 29	ls	core PW408	(149.5 m)	Hypselocyclus	Lothari	0.53	2.44	-3.21
KL 27	ls	core PW408	(149.5 m)	Hypselocyclus	Lothari	0.53	2.45	-4.12
KL 31	ls	core PW408	(150 m)	Hypselocyclus	Lothari	0.52	2.45	-4.28
KL 30	ls	core PW408	(152.6 m)	Hypselocyclus	Lothari	0.51	2.45	-4.13
KL 32	ls	core 37/13.5	(81 m)	Hypselocyclus	Lothari	0.5	2.23	-3.36
KL 36	ls	core 37/13.5	(83 m)	Hypselocyclus	Lothari	0.49	2.00	-4.01
KL 28	ls	core KT109	(256.5 m)	Hypselocyclus	Lothari	0.37	2.28	-4.14
KL 26	ls	core 130SP	(173 m)	Hypselocyclus	Lothari	0.17	2.27	-3.32
KL 40*	mrl	core 6W	(66.8–68.1 m)	Hypselocyclus	Lothari	0.16	-0.02*	-4.85*
KL 43*	mrl	core 6W	(66.8–68.1 m)	Hypselocyclus	Lothari	0.15	0.62*	-4.01*
KL 44*	mrl	core 6W	(68.1–72.0 m)	Hypselocyclus	Lothari	0.14	0.83*	-3.57*
KL 46	ls	core KP2	(103.2–103.6m)	Hypselocyclus	Lothari	0.1	2.15	-4.09
KL 45	ls	core KP2	(113.7–115 m)	Hypselocyclus	Lothari	0.07	2.28	-3.31
KL 48	ls	core KP3	(116–117 m)	Hypselocyclus	Lothari	0.02	2.46	-3.93
KL 49	ls	core KP2	(117 m)	Hypselocyclus	Lothari	0	2.53	-3.50
KL 47	ls	core KP2	(117–118 m)	Hypselocyclus	Hippolytense	0.98	2.53	-4.13
KL 97	ls	Góry Wapienne	–	Acanthicum	–	0.25	1.54	-3.05
KL 90	ls	Góry Wapienne	–	Acanthicum	–	0.25	1.87	-3.02
KL 95	ls	Burzenin 47	–	Divisum	Uhlandi	0.5	2.04	-3.01
KL 39	ls	Burzenin 47	–	Divisum	Uhlandi	0.5	1.98	-2.77
KL 98	ls	Sarnów	–	Divisum	Crussoliense	0.5	2.46	-3.01
KL 58	ls	Sarnów	–	Divisum	Crussoliense	0.5	2.35	-3.29
KL 38	ls	Burzenin 16/9	–	Hypselocyclus	Lothari	0.9	2.08	-2.99
KL 37	ls	Burzenin 16/2	–	Hypselocyclus	Lothari	0.75	2.49	-5.20
KL 41	ls	Kule 84	–	Hypselocyclus	Hippolytense	0.25	1.13	-5.15
KL 42	ls	Kule 84	–	Hypselocyclus	Hippolytense	0.25	2.36	-4.24
KL 96	mrl	PJ110	–	Planula	Planula	0.95	2.68	-6.72
KL 55	mrl	PJ110	–	Planula	Planula	0.95	2.70	-6.16

* – altered samples (see text); position in subzone – percentage position in a stratigraphical subzone, ls – limestone, ool ls – oolitic limestone, mrl – marl, mrl cl – marly clay

Although bulk carbonate $\delta^{13}\text{C}$ values, which are considered as well-preserved, show some scatter, a gentle, long-term decreasing trend of the values is observed (Fig. 10). A similar decrease in $\delta^{13}\text{C}$ values of marine carbonates and belemnite rostra is also noted from western Europe (Colombié et al., 2011; 2018) and the Russian Platform (Riboulleau et al., 1998; Zakharov et al., 2005; Wierzbowski et al., 2013, 2018; see Fig.

10). Absolute $\delta^{13}\text{C}$ values of an “upper envelope” of the current dataset are similar to coeval bulk carbonate data from W France, W Switzerland, central Germany and the Tatra Mountains (Colombié et al., 2011, 2018; Jach et al., 2014; Zuo et al., 2018). This indicates that the Kimmeridgian global trend of decreasing carbon isotope values (cf. Price et al., 2016; Zuo et al., 2018) and marine carbon isotope signatures are, at least partly,

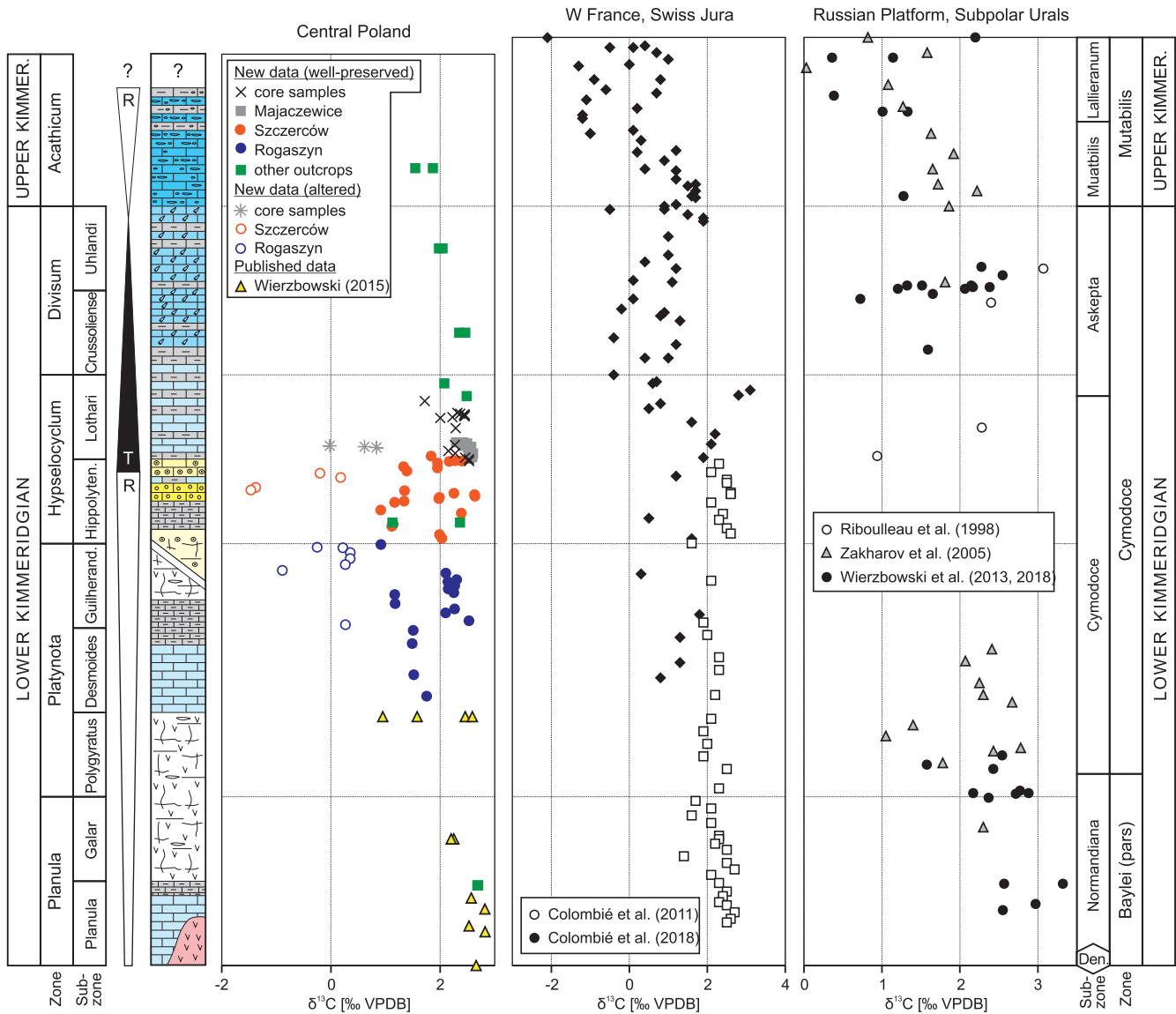


Fig. 10. Bulk carbonate $\delta^{13}\text{C}$ record of central Poland (present and Wierzbowski's 2015 data), and its comparison with the bulk carbonate record of western Europe (after Colombié et al., 2011, 2018) and belemnite data from the Russian Platform and Subpolar Urals (after Ribouilleau et al., 1998; Zakharov et al., 2005; Wierzbowski et al., 2013, 2018)

Stratigraphical correlations between regional zonal schemes are established after Wierzbowski and Rogov (2013), Wierzbowski et al. (2013), and Comment et al. (2015). Generalized lithological and major sequence logs for the Wieluń Upland (Planula to Platynota–Hypselocyclum zone boundary) and its northern vicinity (Platynota–Hypselocyclum zone boundary to Acanthicum Zone) are given after Wierzbowski (2017); the vertical scale of the diagram is based on the biostratigraphical zonal scheme and does not reflect variable thickness of the strata. Lithological symbols as in Figure 2

recorded in central Poland. Local effects connected with fresh-water inflow and water-mass ageing in a very shallow environment probably caused negative shifts within some carbon isotope data of bulk carbonates (Fig. 10). The negative carbon isotope shifts are mostly confined to very shallow, lowstand deposits of the Prusicko and "oolitic" formations encompassing the upper Platynota and the lower Hypselocyclum zones. Similar $\delta^{13}\text{C}$ values of adjoining limestone and marly beds indicates that environmental factors driving contents of carbonate and clastic material are not recorded in the carbon isotope signatures of the rocks. This may be due to a weak relationship between short-term changes in depositional conditions and the carbon isotope composition of DIC of the whole basin or due to early diagenetic processes, which might have caused carbon isotope homogenisation of neighbouring beds.

CONCLUSIONS

Newly presented oxygen and carbon isotope values of well-preserved marine bivalves and bulk carbonates from the Lower–lowest Upper Kimmeridgian of central Poland have allowed documentation of local environmental changes driven by a major regressive-transgressive cycle. $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of bivalve shells from the Małogoszcz section (SW margin of the Holy Cross Mountains), which were deposited in an extremely shallow environment during the Hypselocyclum and the earliest Divisum chrons, show significant scatter because of salinity variations and water-mass ageing. Calculated oxygen isotope temperatures (between 18 and 27°C) may be partly overestimated as a result of decreases in the salinity of

the basin. Partial stabilization of the isotope signatures is noted after the marine transgression, in the lowermost Mutabilis Zone.

Although a global, gentle decrease in marine $\delta^{13}\text{C}$ values throughout the Kimmeridgian is recorded in bulk carbonate data from central Poland (on the Radomsko Elevation and the Wieluń Upland), numerous, low $\delta^{13}\text{C}$ values are found in the interval of maximal shallowing of the basin, which took place during the late Platynota and early Hypselocyclus chrons.

Newly obtained oxygen isotope data of well-preserved fossils, along with published $\delta^{18}\text{O}$ values of Wierzbowski (2015), and Wierzbowski et al. (2016), may be used to document tem-

perature and salinity variations in the epicontinental Polish basin during the major Late Oxfordian–earliest Kimmeridgian regression and the consecutive transgressions, which started, depending on the location, in the latest Hypselocyclus or the Divisum chrons of the latest Early Kimmeridgian. The present and published data suggest that $\delta^{18}\text{O}$ values of Upper Jurassic fossils from central Poland correspond closely to sea level variations.

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