

Material

The present study combines new stable oxygen isotope data from Morocco, Tunisia, Israel, and Jordan with already published results from Egypt (Alberti et al., 2017). The geographic coordinates of the individual study areas are listed in Table S1.

Previous reconstructions of absolute water temperatures for the Jurassic of Europe are dominated by stable oxygen isotope analyses of belemnite rostra (e.g., Dera et al., 2011; Martinez and Dera, 2015; Korte et al., 2015). However, these fossils generally record more positive $\delta^{18}\text{O}$ values than co-occurring bivalve and brachiopod shells. The reason for this discrepancy is not yet completely known, but some authors consider belemnite rostra to be less reliable for temperature reconstructions (compare Mutterlose et al., 2010; Alberti et al., 2012a, b; Price et al., 2015; Hoffmann and Stevens, 2019). More recent studies suggest that the fractionation factor for stable oxygen isotopes in belemnites from seawater was closer to that of abiogenic calcite compared to other marine organisms (Vickers et al., 2020). If this is true, temperatures reconstructed with belemnite rostra would be considerably higher. Due to these problems and because belemnites are relatively uncommon in most of the Middle and Upper Jurassic strata of northern Africa and western Asia, the current study uses calcitic shells of bivalves and rhynchonellid brachiopods.

Only very few fossil shells were available for analyses from the Jurassic of Morocco collected during a field survey by F.T. Fürsich in March-April 1990. These specimens were collected from the ?Toarcian-Aalenian south of Ifri, the Bajocian south of Feggou, the Callovian south of Imouzèrdes-Ida-Outanane, and the Oxfordian east of Smimou. All these outcrops are part of the Atlas Mountains System. The shells include rhynchonellid brachiopods as well as the oysters *Nanogyra nana*, *Exogyra chouberti*, and *Gryphaea* sp. Unfortunately, no precise age assignments beyond stage level were possible.

Jurassic fossils of southern Tunisia were collected for palaeoecologic analyses by S. Holzapfel (1998) and her material is stored in the Bayerische Staatssammlung für Paläontologie und Geologie in Munich, Germany. In the present study, 41 well-preserved rhynchonellid brachiopod shells from seven Callovian sections close to and south of Tataouine were analysed. Each shell could be attributed to a particular horizon in the sections measured by Holzapfel (1998), but the scarcity of ammonites in the succession does not allow very precise age assignments beyond substage level. The sampled horizons belong to the Beni Oussid, Krechem el Miit, and Ghomrassen members of the Tataouine Formation. The Beni Oussid and Krechem el Miit members are characterized by alternations of marl, carbonates, silt- and sandstones, while the lower Ghomrassen Member consists of pack- and grainstones (Holzapfel, 1998). The sampled levels are characterized by a highly diverse fauna (including irregular echinoids and nautiloids) pointing to fully marine conditions on a carbonate platform (Holzapfel, 1998).

Fossils from the Middle to Upper Jurassic of Gebel Maghara in the Sinai of northeastern Egypt were collected for palaeontological studies by A.A. Abdelhady (2014; Abdelhady and Fürsich, 2014, 2015a, b, c). This material is currently stored at the GeoZentrum Nordbayern of the Friedrich-Alexander-Universität Erlangen-Nürnberg, Germany. The stable isotope ($\delta^{18}\text{O}$) composition of 73 well-preserved shells with Bajocian to Kimmeridgian ages was analysed (Alberti et al., 2017). Biostratigraphic studies allowed the assignment of ammonite zones to the individual samples (Abdelhady and Fürsich, 2015a). The stable isotope data is presented here with an improved stratigraphic resolution, but for more details see the earlier study (Alberti et al., 2017).

Thirty-five well-preserved rhynchonellid brachiopods and oysters were collected by M. Alberti and Y. Leshno from the Middle Jurassic and Lower Cretaceous of the Negev Desert of southern Israel during a field survey in November 2017. Two brachiopods could be found in the Upper Bajocian Mahmal Formation of Makhtesh Ramon, 29 shells were collected from the Callovian Zohar and Matmor formations of the Makhtesh Gadol, and four oysters stem from the Lower Cretaceous of Makhtesh Ramon. The Jurassic material could be assigned to earlier defined subunits (Goldberg, 1963; Parnes, 1981; Hirsch, 2005). Unfortunately, the general scarcity of ammonites complicates biostratigraphic dating for most of the Jurassic succession and only the Callovian specimens could be assigned to ammonite zones. While large parts of the Jurassic of southern Israel are nearly unfossiliferous, the Callovian strata of the Makhtesh Gadol contain an abundant fully marine fauna

Table S1 Approximate geographic coordinates of the sampled sections in this study.

| No. | Locality | Latitude | Longitude |
|-------|---|----------|-----------|
| MO-1 | Morocco (~1 km S of Ifri) | N 31.272 | W 6.072 |
| MO-2 | Morocco (~1.5 km S of Feggou; precise locality unknown) | | |
| MO-3 | Morocco (~10 km S of Imouzér-des-Ida-Outanane) | N 30.620 | W 9.400 |
| MO-4 | Morocco (~12 km E of Smimou) | N 31.208 | W 9.579 |
| TU-1 | Tunisia (15 Remada Southeast) | N 32.250 | E 10.545 |
| TU-2 | Tunisia (14 Faljet Jdari / Ed- Dghaghra) | N 32.462 | E 10.426 |
| TU-3 | Tunisia (13 Krechem el Miit) | N 32.617 | E 10.400 |
| TU-4 | Tunisia (12 Bir Remtha) | N 32.735 | E 10.478 |
| TU-5 | Tunisia (11 Ksar Beni Soltane; Tazerdunet) | N 32.790 | E 10.515 |
| TU-6 | Tunisia (5 Foum Tataouine Poste Optique) | N 32.970 | E 10.450 |
| TU-7 | Tunisia (4 Foum Tataouine North) | N 33.045 | E 10.380 |
| EG-1 | Egypt (Gebel Maghara-Homayir) | N 30.647 | E 33.328 |
| EG-2 | Egypt (Gebel Maghara-Arousiah) | N 30.677 | E 33.355 |
| EG-3 | Egypt (Gebel Maghara-Engabashi) | N 30.698 | E 33.387 |
| EG-4 | Egypt (Gebel Maghara-Mowerib) | N 30.682 | E 33.422 |
| IS-1 | Israel (Makhtesh Gadol; gad-11) | N 30.948 | E 35.000 |
| IS-2 | Israel (Makhtesh Gadol; gad-43) | N 30.941 | E 34.983 |
| IS-3 | Israel (Makhtesh Gadol; gad-47) | N 30.941 | E 34.982 |
| IS-4 | Israel (Makhtesh Gadol; gad-48) | N 30.941 | E 34.981 |
| IS-5 | Israel (Makhtesh Gadol; gad-60/61) | N 30.942 | E 34.980 |
| IS-6 | Israel (Makhtesh Gadol; gad-62) | N 30.943 | E 34.980 |
| IS-7 | Israel (Makhtesh Gadol; gad-63) | N 30.943 | E 34.980 |
| IS-8 | Israel (Makhtesh Ramon; ram-5a; 15-01) | N 30.627 | E 34.842 |
| IS-9 | Israel (Makhtesh Ramon; ram-5a; 15-02) | N 30.617 | E 34.836 |
| IS-10 | Israel (Makhtesh Ramon; ram-Cr) | N 30.624 | E 34.842 |
| JO-1 | Jordan (Wadi Nimr) | N 32.108 | E 35.660 |
| JO-2 | Jordan (Arda) | N 32.125 | E 35.654 |
| JO-3 | Jordan (Wadi Huni) | N 32.175 | E 35.705 |
| JO-4 | Jordan (Tal el Dahhab) | N 32.189 | E 35.713 |
| JO-5 | Jordan (King Talal Dam) | N 32.191 | E 35.800 |

including echinoids, brachiopods, and occasional ammonites.

Jurassic fossils in northern Jordan were collected for palaeoecologic and taxonomic studies by F. Ahmad (1999) and his material is stored in the Bayerische Staatssammlung für Paläontologie und Geologie in Munich, Germany. Thirty bivalve shells from five sections northwest of Amman were analysed in the present study. The material can be attributed to horizons with Bajocian, Bathonian, and Callovian ages, but the general lack of ammonites in the sections hinders more precise age assignments beyond stage level (Ahmad, 1999). Lithostratigraphically, the fossils represent the Dahhab, Ramla, Hamam, and Mughanniyya formations.

Illustrations of the analysed species can be found readily in already published articles (Dubar, 1967; Holzapfel, 1998; Ahmad, 1999; Feldman et al., 2001, 2012, 2014; Alberti et al., 2017).

Palaeolatitudes

Palaeolatitudes were reconstructed with the help of the so-called Paleolatitude Calculator developed by van Hinsbergen et al. (2015) and using the palaeomagnetic reference curves of Kent and Irving (2010) for Morocco and Torsvik et al. (2012) for the other study areas. Due to a slight clockwise rotation of Africa, Morocco was situated at a much more northern position compared to Jordan during the Jurassic. Within the present study, palaeolatitudes are usually rounded to integers, but in the tables and figures values with one decimal are given. It has to be noted, however, that the reconstruction of palaeolatitudes is not a very precise science and absolute values can differ considerably among different studies. The given values are therefore mathematical and some degree of error has to be tolerated. Nevertheless, even changes of a few degrees in latitude for the different study sites would not influence the overall shape of the reconstructed latitudinal temperature gradients or the approximated $\delta^{18}\text{O}_{\text{sea}}$ values. Although the exact locality of the sampling site near Feggou in Morocco is unknown, a palaeolatitude of 27.1 °N was determined for the general study area in the Bajocian.

Preservation of fossil shells

A thorough check of the preservational quality of fossils used for stable isotope analyses is a fundamental step in producing reliable temperature data. In this context, cathodoluminescence microscopy has become a standard method to select suitable specimens and sampling areas (e.g., Wierzbowski, 2002, 2004; Ullmann and Korte, 2015; Arabas, 2016). Only shells with large non-luminescent areas pointing to a good preservation were used to reconstruct water temperatures in the present study (Fig. S1A-C). Furthermore, selected fossils were studied with a scanning electron microscope, which was particularly helpful in examining the fine microstructure of rhynchonellid brachiopods (Fig. S1D-F). For some fossils from Jordan (with the lowest palaeolatitudes in the dataset), enough carbonate powder could be extracted to also measure Mg/Ca- and Sr/Ca-ratios as well as manganese and iron contents. Shells with an original, unaltered chemical composition are believed to have very low manganese and iron contents (e.g., Price and Sellwood, 1997; Wierzbowski and Joachimski, 2007; Wierzbowski et al., 2009). Consequently, eight specimens were excluded from temperature reconstructions, because they contained more than 300 µg/g iron and/or more than 100 µg/g manganese (comparable to cut-off-values used by Alberti et al., 2012a, b). Another commonly mentioned indicator for the diagenetic alteration of fossil shells in a collection is a strong, positive correlation between their $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values, since both values generally decrease progressively with increased alteration (compare Hodgson, 1966; Hudson, 1977; Nelson and Smith, 1996). This effect can be seen very well in the sediment and cement samples which commonly show lower $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values than the seemingly well-preserved shells (Fig. S2A). While the number of specimens from Morocco is too small to examine a possible relationship between their $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values, there is no statistically significant correlation in the results from Tunisia ($r_s = 0.22$; $p =$

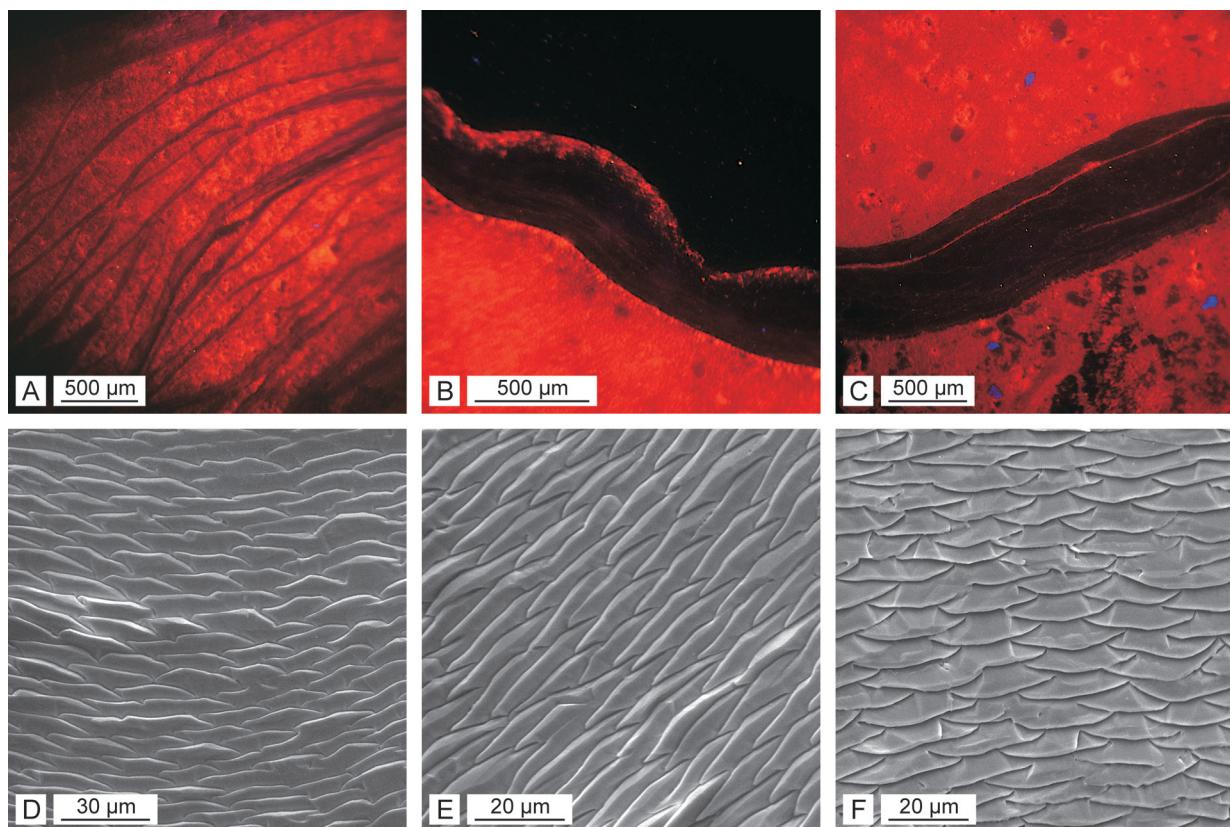
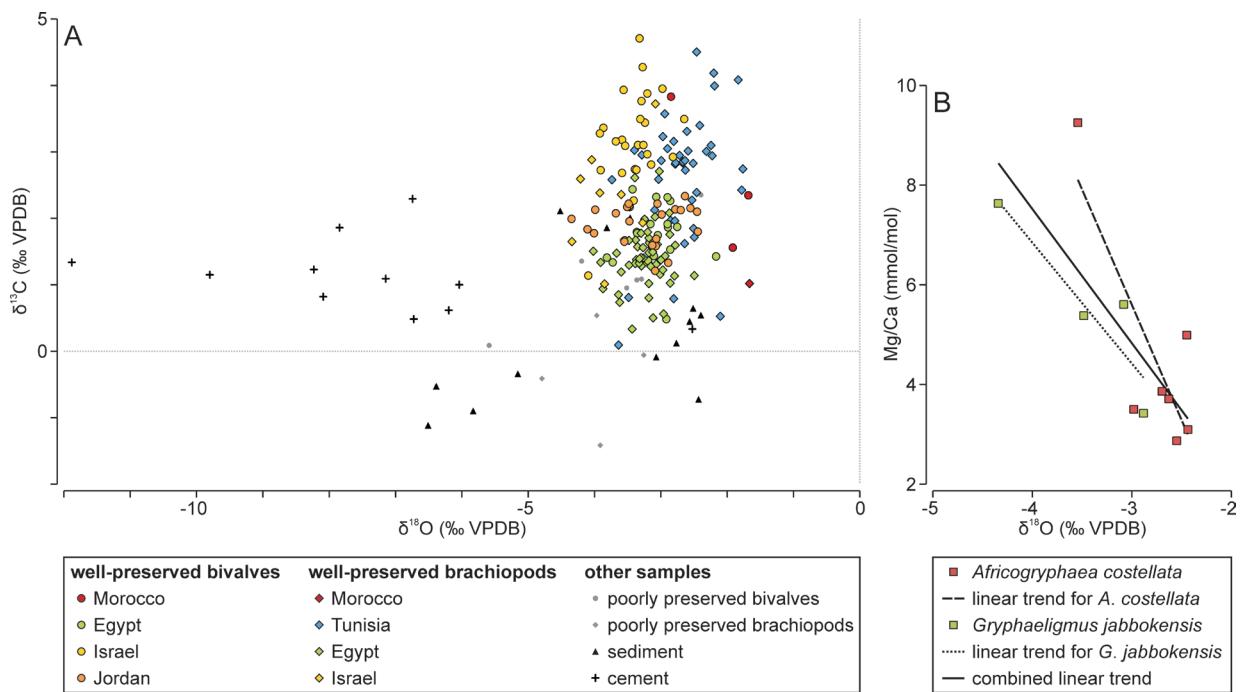


Fig. S1 Cathodoluminescence and scanning electron microscopy. The specimens were cut into slabs with their surface ground. Scanning electron microscopy was conducted after etching of the fossils with 5 % HCl for ca. 40 s. **A.** Poorly preserved gryphaeid oyster with sparry calcite filling voids within the shell (Jurassic, Morocco). **B.** Well-preserved specimen of *Somalirhynchia* cf. *africana* showing a luminescent outer shell margin due to weathering and possibly bioerosion (Callovian, Tunisia). **C.** Luminescent cracks in an otherwise well-preserved shell of *Septirhynchia pulchra* (Callovian, Tunisia). **D-F.** Specimens of *Somalirhynchia* cf. *africana* (D), *Daghanirhynchia angulocostata* (E), and *Somalirhynchia smelliei* (F) exhibiting well-preserved shell microstructure (Callovian, Tunisia).



0.17), Egypt ($r_s = 0.11$; $p = 0.36$), and Jordan ($r_s = 0.17$; $p = 0.44$). The data from Israel show a weak positive correlation ($r_s = 0.51$; $p = 0.002$), which is not conclusive. Nevertheless, since the absolute $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values of shells from Israel are comparable to those of Jordan and Egypt, it can be assumed that they are not altered. Summarizing the studies examining the preservational quality of the analysed shells, it seems plausible that the specimens used for temperature reconstructions are well preserved and show a pristine chemical composition.

Geochemical analyses

The selected fossils were sampled with a hand-held dental drill in the case of thick shells or with a computer-controlled micromill. Areas near the umbo/hinge as well as the muscle scar were avoided for bivalve shells (compare Huyghe et al., 2020). Similarly, rhynchonellid brachiopod shells were preferentially sampled in the center of the shells avoiding the umbo/pedical opening. The retrieved carbonate powder (0.05-0.50 mg) was then analysed using a carbonate preparation device (Kiel IV) connected with a ThermoScientific MAT 253 mass spectrometer at the Leibniz Laboratory for Radiometric Dating and Stable Isotope Research at the Christian-Albrechts-Universität zu Kiel, Germany. The samples were reacted within the preparation device with 100 % H_3PO_4 at 75 °C and the evolved CO_2 gas was analysed using the mass spectrometer. On a daily routine, different laboratory internal carbonate standards and two international carbonate standards were analysed to control the precision of measured $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ values. As documented by the performance of international [NBS19: +1.95 ‰ VPDB (^{13}C), -2.20 ‰ VPDB (^{18}O); IAEA-603: +2.46 ‰ VPDB (^{13}C), -2.37 ‰ VPDB (^{18}O)] and laboratory-internal carbonate standards [Hela1: +0.91 ‰ VPDB (^{13}C), +2.48 ‰ VPDB (^{18}O); HB1: -12.10 ‰ VPDB (^{13}C), -18.10 ‰ VPDB (^{18}O); SHK: +1.74 ‰ VPDB (^{13}C), -4.85 ‰ VPDB (^{18}O)], analytical precision of stable isotope analysis is better than ± 0.08 ‰ ($\pm 1\text{SD}$) for $\delta^{18}\text{O}$ and better than ± 0.05 ‰ ($\pm 1\text{SD}$) for $\delta^{13}\text{C}$. All values are reported in per mil relative to the Vienna Pee Dee Belemnite (VPDB) scale using NBS-19.

Trace element contents and molar Sr/Ca and Mg/Ca ratios of the bivalve shells of Jordan were determined by ICP-Optical Emission Spectrometry (ICP-OES) using a Ciros SOP instrument (Spectro Analytical Instruments, Germany) at the Institute of Geosciences, Christian-Albrechts-Universität zu Kiel. Sample introduction was via a concentric micro-nebulizer (GE SeaSpray 400 μL) combined

with a temperature-controlled cyclonic spray chamber. Typical sample weights for dissolution with 0.3 molar nitric acid were between 0.01 and 0.6 mg, and the final solution ready for analysis was adjusted to a concentration of 40 ± 10 mg/L Ca and 20 ± 10 mg/L Ca for oysters and belemnites, respectively. Every sample was analysed with 5 runs applying truly simultaneous acquisition times to all spectral lines investigated. Raw data in [cps] was exported and processed with external spreadsheet software combining intensity-ratio calibration after de Villiers et al. (2002) and standard-sample-standard bracketing and drift correction procedures after Schrag (1999). Data for oysters were normalized to ECRM-752 with Mg/Ca 3.821 mmol/mol and Sr/Ca 0.174 mmol/mol (Greaves et al., 2008), and data for belemnites were normalized to in-house standard sample belemnite Ja002 with Mg/Ca 16.42 Mg/Ca and Sr/Ca 0.3739 mmol/mol. Average uncertainty as estimated from 5 runs was 1.3 and 1.2 ‰ (1SD) for Mg/Ca and Sr/Ca in oysters, respectively, and 1.6 and 1.3 ‰ (1SD) in belemnites. Corals at 8 mg/L Ca typically yield uncertainties of 0.7–0.9 ‰ (1SD) for Sr/Ca. Duplicate sample analyses after approx. 90 minutes, and samples from the beginning of an analytical session re-analysed at the end of the session after ten or more hours yield average reproducibilities in the same range. Reference materials Coral JCp-1, Tridacna JCt-1, limestones ECRM 752, and CAL-S were used as secondary standards using consensus values from Hathorne et al. (2013) and Greaves et al. (2008).

Detailed description of the stable isotope record

Results of the stable isotope analyses of bivalve and brachiopod shells of northern Africa and western Asia are listed in Table S2.

Analysed shells from Morocco include a specimen of *Gryphaea* sp. from the ?Toarcian-Aalenian ($\delta^{18}\text{O}$: -2.84 ‰; $\delta^{13}\text{C}$: 3.83 ‰), one shell of *Exogyra chouberti* from the Bajocian ($\delta^{18}\text{O}$: -1.91 ‰; $\delta^{13}\text{C}$: 1.56 ‰), a rhynchonellid brachiopod from the Callovian ($\delta^{18}\text{O}$: -1.66 ‰; $\delta^{13}\text{C}$: 1.02 ‰), and one specimen of *Nanogyra nana* from the Oxfordian ($\delta^{18}\text{O}$: -1.68 ‰; $\delta^{13}\text{C}$: 2.34 ‰). Due to the scarcity of the data, no reliable trends through time can be discerned.

The collection from Tunisia includes 41 seemingly well-preserved rhynchonellid brachiopods with Early to Late Callovian ages (Fig. S3). Throughout time, $\delta^{18}\text{O}$ values vary around an overall average of -2.66 ‰ (minimum: -3.73 ‰; maximum: -1.76 ‰). Short-term fluctuations in $\delta^{18}\text{O}$ values can be seen (e.g., the most negative $\delta^{18}\text{O}$ values being reached around the Middle to Late Callovian boundary), but no long-term temporal trend can be found. The $\delta^{13}\text{C}$ values fluctuate more strongly around an overall average of 2.65 ‰ (minimum: 0.09 ‰; maximum: 4.50 ‰). From relatively high $\delta^{13}\text{C}$ values at the beginning of the Callovian (average: 3.82 ‰), values decrease to 0.53 ‰ before more or less steadily rising again towards the Late Callovian (average: 2.66 ‰).

The collection from Egypt consists of 51 well-preserved rhynchonellid brachiopods and 22 bivalves with Bajocian to Kimmeridgian ages. Their stable isotope data has been described already in detail by Alberti et al. (2017), but is presented here again with an updated lithostratigraphic correlation at higher stratigraphic resolution (Fig. S4). The results of the stable isotope analyses show relatively equitable conditions through time with an overall average $\delta^{18}\text{O}$ value of -3.20 ‰ and an average $\delta^{13}\text{C}$ value of 1.50 ‰ (for more information see Alberti et al., 2017).

The stable isotope dataset of Israel includes results of 26 bivalves and 9 brachiopods with Late Bajocian, Callovian, and Early Cretaceous ages (Fig. S5). The $\delta^{18}\text{O}$ record starts with relatively low values in the Late Bajocian (average: -4.06 ‰). The $\delta^{18}\text{O}$ values are slightly higher in the Middle Callovian (Coronatum Zone, average: -3.37 ‰) and remain at similar values in the Late Callovian (Athleta Zone, average: -3.54 ‰; Lamberti Zone, average: -3.36 ‰). Four oysters from the Early Cretaceous show comparable $\delta^{18}\text{O}$ values around an average of -3.58 ‰. The $\delta^{13}\text{C}$ values show a slight increase throughout the studied time interval. Their record starts with comparatively low values in the Late Bajocian (average: 2.01 ‰). Middle Callovian (Coronatum Zone) $\delta^{13}\text{C}$ values are scattered around an average of 2.55 ‰, while Late Callovian values are still higher (Athleta Zone, average: 2.99 ‰; Lamberti Zone, average: 3.63 ‰). The Early Cretaceous oysters show $\delta^{13}\text{C}$ values around an average of 2.88 ‰.

Material from Jordan includes 22 seemingly well-preserved bivalves with Bajocian to Callovian

| sample no. | taxonomy | group | age | bed | locality | coordinates | paleolatitude | $\delta^{13}\text{C}$ | $\delta^{18}\text{O}$ | T (°C) ¹ | $\delta^{18}\text{O}_{\text{sea}} \text{--} \delta^{18}\text{O}_{\text{sea}}$ T (°C) ³ |
|--------------------------|-----------------------------|---------------|-----------------|----------|----------|-------------------|---------------|-----------------------|-----------------------|---------------------|---|
| Morocco | | | | | | | | | | | |
| well-preserved fossils | | | | | | | | | | | |
| MO18-007 | Gryphaea sp. | bivalve | ?Toa.-Aal. | - | MO-1 | 31.272° .-6.072° | 29.3 | 3.83 | -2.84 | 24.0 | -0.17 |
| MO18-005 | Exogyra chouberti | bivalve | Bajocian | - | MO-2 | - | 1.56 | -1.91 | 19.9 | -0.10 | 23.9 |
| MO18-006 | rhyynchonellid | brachiopod | Callovian | - | MO-3 | 30.620° .-9.400° | 27.4 | 1.02 | -1.66 | 18.8 | -0.11 |
| MO18-002-A2B | Nanogyra nana | bivalve | Oxfordian | - | MO-4 | 31.208° .-9.579° | 28.1 | 2.34 | -1.68 | 18.9 | -0.13 |
| sediment & cement | | | | | | | | | | | |
| MO18-003Sed | - | sediment fili | Bajocian | - | MO-2 | - | 27.1 | -0.73 | -2.43 | - | - |
| MO18-005Cem | - | cement fili | Bajocian | - | MO-2 | - | 27.1 | 0.33 | -2.52 | - | - |
| MO18-006Sed | - | sediment fili | Callovian | - | MO-3 | 20.620° .-9.400° | 27.4 | 0.64 | -2.51 | - | - |
| poorly preserved fossils | | | | | | | | | | | |
| MO18-003 | rhyynchonellid | brachiopod | Bajocian | - | MO-2 | - | 27.1 | -0.06 | -3.25 | - | - |
| Tunisia | | | | | | | | | | | |
| well-preserved fossils | | | | | | | | | | | |
| TU17-034-A2B | Septirhynchia pulchra | brachiopod | E.-M. Callovian | r2/26/4 | TU-4 | 32.7735° .10.478° | 13.7 | 4.50 | -2.46 | 22.3 | 0.07 |
| TU17-036 | Septirhynchia pulchra | brachiopod | E.-M. Callovian | r2/26/4 | TU-4 | 32.7735° .10.478° | 13.7 | 2.95 | -3.28 | 26.1 | 0.07 |
| TU17-037 | Septirhynchia pulchra | brachiopod | E.-M. Callovian | r2/26/4 | TU-4 | 32.7735° .10.478° | 13.7 | 3.99 | -2.18 | 21.1 | 0.07 |
| TU17-008 | rhyynchonellid | brachiopod | E.-M. Callovian | 3/22/2 | TU-7 | 33.045° .10.380° | 14.0 | 0.53 | -2.09 | 20.7 | 0.08 |
| TU17-039 | Septirhynchia pulchra | brachiopod | E.-M. Callovian | p5/12/4 | TU-6 | 32.970° .10.450° | 13.9 | 1.72 | -2.49 | 22.4 | 0.08 |
| TU17-040ab | Septirhynchia pulchra | brachiopod | E.-M. Callovian | p5/12/4 | TU-6 | 32.970° .10.450° | 13.9 | 0.79 | -2.80 | 23.9 | 0.08 |
| TU17-002 | Somalirhynchia cf. africana | brachiopod | E.-M. Callovian | s3/17/8 | TU-1 | 32.250° .10.545° | 13.3 | 2.13 | -3.09 | 25.2 | 0.07 |
| TU17-003 | Somalirhynchia cf. africana | brachiopod | E.-M. Callovian | s3/17/8 | TU-1 | 32.250° .10.545° | 13.3 | 3.57 | -2.93 | 24.5 | 0.07 |
| TU17-024 | rhyynchonellid | brachiopod | E.-M. Callovian | s3/17/8 | TU-1 | 32.250° .10.545° | 13.3 | 2.58 | -3.73 | 28.3 | 0.07 |
| TU17-001 | Somalirhynchia cf. africana | brachiopod | E.-M. Callovian | r3/2/7 | TU-4 | 32.733° .10.478° | 13.7 | 2.82 | -2.78 | 23.8 | 0.07 |
| TU17-019 | Somalirhynchia smeliei | brachiopod | E.-M. Callovian | r3/2/8 | TU-4 | 32.733° .10.478° | 13.7 | 3.03 | -3.39 | 26.6 | 0.07 |
| TU17-011-2AB | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 2.72 | -2.62 | 23.1 | 0.07 |
| TU17-012 | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 2.28 | -2.52 | 22.6 | 0.07 |
| TU17-013ab | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 3.10 | -2.24 | 21.3 | 0.07 |
| TU17-014ab | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 3.01 | -2.58 | 22.9 | 0.07 |
| TU17-015ab | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 0.09 | -3.63 | 27.8 | 0.07 |
| TU17-016 | Kallirhynchia africana | brachiopod | Late Callovian | f3/6/1 | TU-2 | 32.462° .10.426° | 13.5 | 2.87 | -2.99 | 24.7 | 0.07 |
| TU17-042ab | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 3.40 | -2.41 | 22.1 | 0.07 |
| TU17-043 | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 2.85 | -2.66 | 23.2 | 0.07 |
| TU17-044-A2B | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 3.31 | -2.60 | 23.0 | 0.07 |
| TU17-045-A2B | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 3.05 | -2.89 | 24.3 | 0.07 |
| TU17-046ab | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 3.23 | -2.96 | 24.6 | 0.07 |
| TU17-047-A2B | Somalirhynchia smeliei | brachiopod | Late Callovian | m5/8/4 | TU-3 | 32.617° .10.400° | 13.7 | 2.87 | -2.63 | 23.1 | 0.07 |
| TU17-004ab | rhyynchonellid | brachiopod | Late Callovian | gt3/24/1 | TU-7 | 33.045° .10.380° | 14.0 | 4.08 | -1.83 | 19.5 | 0.08 |
| TU17-005 | rhyynchonellid | brachiopod | Late Callovian | gt3/24/1 | TU-7 | 33.045° .10.380° | 14.0 | 3.00 | -2.31 | 21.6 | 0.08 |
| TU17-006A | rhyynchonellid | brachiopod | Late Callovian | gt3/24/1 | TU-7 | 33.045° .10.380° | 14.0 | 2.94 | -2.22 | 21.2 | 0.08 |
| TU17-032 | Rhynchonella tazerdunensis | brachiopod | Late Callovian | r3/3/1 | TU-4 | 32.7735° .10.478° | 13.7 | 2.83 | -2.66 | 23.2 | 0.07 |

Table S2 Results of the stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of samples from northern Africa and western Asia (°Water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰, $\delta^{18}\text{O}_{\text{sea}}$ value calculated for the paleolatitude with equation (4), °Water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value calculated with equation (4) for the paleolatitude).

Table S2 (continued) Results of the stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of samples from northern Africa and western Asia (${}^1\text{H}_2\text{O}$ water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1‰ , ${}^2\delta^{18}\text{O}_{\text{sea}}$ value calculated for the palaeolatitude with equation (4), ${}^3\delta^{18}\text{O}_{\text{sea}}$ value calculated with equation (4) for the palaeolatitude).

| sample no. | taxonomy | group | age | bed | locality | coordinates | palaeolatitude | $\delta^{13}\text{C}$ | $\delta^{18}\text{O}$ | $T({}^\circ\text{C})$ | $\delta^{18}\text{O}_{\text{sea}} {}^2\text{T}({}^\circ\text{C})$ |
|---------------------------------|--------------------------------------|---------------|------------------|----------|----------|------------------|----------------|-----------------------|-----------------------|-----------------------|---|
| TU17-033ab | <i>Rhynchonella tazerdunensis</i> | brachiopod | Late Callovian | r/3/3/1 | TU-4 | 32.735°, 10.478° | 13.7 | 2.59 | -3.03 | 24.9 | 0.07 |
| TU17-022-AB | <i>Rhynchonella dieffarae</i> | brachiopod | Late Callovian | gt3/24/6 | TU-7 | 33.045°, 10.380° | 14.0 | 2.74 | -1.76 | 19.2 | 0.08 |
| TU17-023ab | <i>Rhynchonella dieffarae</i> | brachiopod | Late Callovian | gt3/24/6 | TU-7 | 33.045°, 10.380° | 14.0 | 1.62 | -2.64 | 23.1 | 0.08 |
| TU17-026ab | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 0.81 | -3.48 | 27.1 | 0.08 |
| TU17-027ab | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 1.85 | -2.50 | 22.5 | 0.08 |
| TU17-028 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 2.39 | -2.45 | 22.3 | 0.08 |
| TU17-029 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 1.97 | -2.85 | 24.1 | 0.08 |
| TU17-030 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 2.42 | -1.77 | 19.3 | 0.08 |
| TU17-031 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Late Callovian | k5/5/2-1 | TU-5 | 32.790°, 10.515° | 13.8 | 1.96 | -2.78 | 23.8 | 0.08 |
| TU17-017A | <i>Somalirhynchia cf. africana</i> | brachiopod | Late Callovian | r3/3/5 | TU-4 | 32.735°, 10.478° | 13.7 | 2.83 | -2.51 | 22.5 | 0.07 |
| TU17-018ab | <i>Somalirhynchia cf. africana</i> | brachiopod | Late Callovian | r3/3/5 | TU-4 | 32.735°, 10.478° | 13.7 | 4.18 | -2.19 | 21.1 | 0.07 |
| TU17-009 | <i>Somalirhynchia cf. africana</i> | brachiopod | Late Callovian | f3/9/2 | TU-2 | 32.462°, 10.426° | 13.5 | 3.15 | -2.80 | 23.9 | 0.07 |
| TU17-010ab | <i>Somalirhynchia cf. africana</i> | brachiopod | Late Callovian | f3/9/2 | TU-2 | 32.462°, 10.426° | 13.5 | 2.83 | -2.77 | 23.7 | 0.07 |
| TU17-038-AB | <i>Somalirhynchia smeliei</i> | brachiopod | Late Callovian | m5/9/21 | TU-3 | 32.617°, 10.400° | 13.7 | 2.95 | -2.71 | 23.5 | 0.07 |
| <i>sediment & cement</i> | | | | | | | | | | | |
| TU17-045C | - | cement | Late Callovian | m5/8/4 | TU-3 | 32.617°, 10.400° | 13.7 | 1.09 | -7.14 | - | - |
| TU17-045D | - | sediment fill | Late Callovian | m5/8/4 | TU-3 | 32.617°, 10.400° | 13.7 | 0.54 | -2.39 | - | - |
| TU17-006B | - | cement | Late Callovian | gt3/24/1 | TU-7 | 33.045°, 10.380° | 14.0 | 1.00 | -6.03 | - | - |
| TU17-021B | - | cement | Late Callovian | gt3/24/6 | TU-7 | 33.045°, 10.380° | 14.0 | 1.34 | -11.88 | - | - |
| TU17-017B | - | sediment fill | Late Callovian | r3/3/5 | TU-4 | 32.735°, 10.478° | 13.7 | 0.45 | -2.56 | - | - |
| <i>poorly preserved fossils</i> | | | | | | | | | | | |
| TU17-035 | <i>Septirhynchia pulchra</i> | brachiopod | E.-M. Callovian | r2/26/4 | TU-4 | 32.735°, 10.478° | 13.7 | -1.42 | -3.91 | - | - |
| TU17-041A | <i>Septirhynchia pulchra</i> | brachiopod | E.-M. Callovian | p5/12/4 | TU-6 | 32.970°, 10.450° | 13.9 | 0.54 | -3.96 | - | - |
| TU17-041B | <i>Septirhynchia pulchra</i> | brachiopod | E.-M. Callovian | p5/12/4 | TU-6 | 32.970°, 10.450° | 13.9 | -0.42 | -4.78 | - | - |
| <i>Egypt</i> | | | | | | | | | | | |
| H1-1 | <i>Cymatirhynchia quadriflicata</i> | brachiopod | Early Bajocian | H1 | EG-1 | 30.647°, 33.328° | 3.3 | 1.63 | -3.17 | 25.6 | -0.16 |
| M1-1 | <i>Cymatirhynchia quadriflicata</i> | brachiopod | Early Bajocian | M1 | EG-4 | 30.682°, 33.422° | 3.3 | 1.67 | -3.37 | 26.6 | -0.16 |
| A5-1 | <i>Atricoryphphaea costellata</i> | bivalve | Late Bajocian | A5 | EG-2 | 30.677°, 33.355° | 3.3 | 2.43 | -3.42 | 26.8 | -0.16 |
| A6-1 | <i>Daghanirhynchia daghamiensis</i> | brachiopod | Late Bajocian | A6 | EG-2 | 30.677°, 33.355° | 3.3 | 1.61 | -3.05 | 25.0 | -0.16 |
| A6-2 | <i>Daghanirhynchia daghamiensis</i> | brachiopod | Late Bajocian | A6 | EG-2 | 30.677°, 33.355° | 3.3 | 1.75 | -2.86 | 24.2 | -0.16 |
| A7-2 | <i>Conarosia rotundata</i> | brachiopod | Late Bajocian | A7 | EG-2 | 30.677°, 33.355° | 3.3 | 1.28 | -3.14 | 25.5 | -0.16 |
| A9-1 | <i>Conarosia rotundata</i> | brachiopod | Late Bajocian | A9 | EG-2 | 30.677°, 33.355° | 3.3 | 1.14 | -2.49 | 22.5 | -0.16 |
| A9-2 | <i>Conarosia rotundata</i> | brachiopod | Late Bajocian | A9 | EG-2 | 30.677°, 33.355° | 3.3 | 1.57 | -3.35 | 26.4 | -0.16 |
| H3-1 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Early Bathonian | H3 | EG-1 | 30.647°, 33.328° | 3.3 | 1.38 | -3.36 | 26.5 | -0.16 |
| H3-3 | <i>Daghanirhynchia angulocostata</i> | brachiopod | Early Bathonian | H3 | EG-1 | 30.647°, 33.328° | 3.3 | 0.73 | -3.62 | 27.7 | -0.16 |
| A10-1 | <i>Conarosia rotundata</i> | brachiopod | Early Bathonian | A10 | EG-2 | 30.677°, 33.355° | 3.3 | 2.71 | -3.02 | 24.9 | -0.16 |
| A11-1 | <i>Cymatirhynchia quadriflicata</i> | brachiopod | Early Bathonian | A11 | EG-2 | 30.677°, 33.355° | 3.3 | 1.34 | -3.90 | 29.1 | -0.16 |
| H6-1 | <i>Daghanirhynchia daghamiensis</i> | brachiopod | Middle Bathonian | H6 | EG-1 | 30.647°, 33.328° | 3.3 | 1.30 | -3.06 | 25.1 | -0.16 |
| H6-2 | <i>Daghanirhynchia daghamiensis</i> | brachiopod | Middle Bathonian | H6 | EG-1 | 30.647°, 33.328° | 3.3 | 1.60 | -2.77 | 23.8 | -0.16 |
| H6-3 | <i>Daghanirhynchia daghamiensis</i> | brachiopod | Middle Bathonian | H6 | EG-1 | 30.647°, 33.328° | 3.3 | 1.79 | -3.19 | 25.7 | -0.16 |

Table S2 (continued) Results of the stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of samples from northern Africa and western Asia (${}^1\text{H}_2\text{O}$ temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1‰ , ${}^{2\delta^{18}\text{O}_{\text{sea}}}$ value calculated for the palaeolatitude with equation (4), ${}^3\text{H}_2\text{O}$ temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value calculated with equation (4) for the palaeolatitude).

| sample no. | taxonomy | group | age | bed | locality | coordinates | $\delta^{18}\text{O}$ | $\delta^{18}\text{O}_{\text{sea}}$ | $T({}^\circ\text{C})$ | $\delta^{18}\text{O}$ | $\delta^{18}\text{O}_{\text{sea}}$ | $T({}^\circ\text{C})$ |
|------------|------------------------------|------------|------------------|-----|----------|-------------------|-----------------------|------------------------------------|-----------------------|-----------------------|------------------------------------|-----------------------|
| A13-1 | Daghanirynchia daghamiensis | brachiopod | Middle Bathonian | A13 | EG-2 | 30.677° . 33.355° | 3.3 | -3.38 | 26.6 | -0.16 | 30.7 | |
| A14-1 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | A14 | EG-2 | 30.677° . 33.355° | 3.3 | -3.29 | 26.2 | -0.16 | 30.3 | |
| A14-2-ab | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | A14 | EG-2 | 30.677° . 33.355° | 3.3 | 0.94 | -3.87 | 28.9 | -0.16 | 33.1 |
| A14-3 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | A14 | EG-2 | 30.677° . 33.355° | 3.3 | 1.99 | -3.24 | 25.9 | -0.16 | 30.0 |
| A14-4 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | A14 | EG-2 | 30.677° . 33.355° | 3.3 | 1.70 | -3.17 | 25.6 | -0.16 | 29.7 |
| A15-1 | Daghanirynchia daghamiensis | brachiopod | Middle Bathonian | A15 | EG-2 | 30.677° . 33.355° | 3.3 | 1.19 | -3.59 | 27.6 | -0.16 | 31.8 |
| H8-3 | Daghanirynchia daghamiensis | brachiopod | Middle Bathonian | H8 | EG-1 | 30.647° . 33.328° | 3.3 | 1.01 | -3.17 | 25.6 | -0.16 | 29.7 |
| A17-1 | Burmimyrinchia cavignarii | brachiopod | Middle Bathonian | A17 | EG-2 | 30.677° . 33.355° | 3.3 | 1.36 | -3.01 | 24.9 | -0.16 | 28.9 |
| A17-2 | Burmimyrinchia cavignarii | brachiopod | Middle Bathonian | A17 | EG-2 | 30.677° . 33.355° | 3.3 | 0.33 | -3.43 | 26.8 | -0.16 | 30.9 |
| A17-3 | Burmimyrinchia cavignarii | brachiopod | Middle Bathonian | A17 | EG-2 | 30.677° . 33.355° | 3.3 | 1.28 | -3.28 | 26.1 | -0.16 | 30.2 |
| E7-1-ab | Afriogryphaea costellata | bivalve | Middle Bathonian | E7 | EG-3 | 30.698° . 33.387° | 3.3 | 1.33 | -3.72 | 28.2 | -0.16 | 32.4 |
| E7-2 | Afriogryphaea costellata | bivalve | Middle Bathonian | E7 | EG-3 | 30.698° . 33.387° | 3.3 | 1.41 | -3.81 | 28.6 | -0.16 | 32.8 |
| E7-3 | Afriogryphaea costellata | bivalve | Middle Bathonian | E7 | EG-3 | 30.698° . 33.387° | 3.3 | 1.67 | -3.54 | 27.3 | -0.16 | 31.5 |
| M11-1 | Afriogryphaea costellata | bivalve | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 1.34 | -3.22 | 25.9 | -0.16 | 29.9 |
| M11-2 | Afriogryphaea costellata | bivalve | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 1.36 | -3.19 | 25.7 | -0.16 | 29.7 |
| M11-3 | Afriogryphaea costellata | bivalve | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 0.48 | -2.91 | 24.4 | -0.16 | 28.4 |
| M11-4-ab | Daghanirynchia daghamiensis | brachiopod | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 0.56 | -2.95 | 24.6 | -0.16 | 28.6 |
| M11-5 | Daghanirynchia daghamiensis | brachiopod | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 1.02 | -2.85 | 24.1 | -0.16 | 28.1 |
| M11-6 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | 0.50 | -3.11 | 25.3 | -0.16 | 29.3 |
| E9-1 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | E9 | EG-3 | 30.698° . 33.387° | 3.3 | 1.27 | -3.14 | 25.5 | -0.16 | 29.5 |
| E9-2 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | E9 | EG-3 | 30.698° . 33.387° | 3.3 | 1.43 | -2.93 | 24.5 | -0.16 | 28.5 |
| E9-3 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | E9 | EG-3 | 30.698° . 33.387° | 3.3 | 1.37 | -2.99 | 24.8 | -0.16 | 28.8 |
| E9-4 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | E9 | EG-3 | 30.698° . 33.387° | 3.3 | 1.15 | -3.01 | 24.8 | -0.16 | 28.8 |
| E9-5 | Afriogryphaea costellata | bivalve | Middle Bathonian | E9 | EG-1 | 30.647° . 33.328° | 3.3 | 1.43 | -3.10 | 25.3 | -0.16 | 29.3 |
| H9-1 | Afriogryphaea costellata | bivalve | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 1.91 | -2.90 | 24.3 | -0.16 | 28.3 |
| H9-2 | Afriogryphaea costellata | bivalve | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 1.91 | -3.15 | 25.5 | -0.16 | 29.6 |
| H9-3 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 1.34 | -3.31 | 26.3 | -0.16 | 30.4 |
| H9-4 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 1.50 | -4.01 | 29.6 | -0.16 | 33.9 |
| H9-5 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 2.02 | -3.08 | 25.2 | -0.16 | 29.2 |
| H9-6 | Afriogryphaea costellata | bivalve | Middle Bathonian | H9 | EG-1 | 30.647° . 33.328° | 3.3 | 2.08 | -2.86 | 24.2 | -0.16 | 28.2 |
| H10-2 | Daghanirynchia angulocostata | brachiopod | Middle Bathonian | H10 | EG-1 | 30.647° . 33.328° | 3.3 | 2.26 | -2.85 | 24.1 | -0.16 | 28.1 |
| H11-1 | Daghanirynchia angulocostata | bivalve | Middle Bathonian | H11 | EG-1 | 30.647° . 33.328° | 3.3 | 1.86 | -2.89 | 24.3 | -0.16 | 28.3 |
| H11-2 | Afriogryphaea costellata | bivalve | Middle Bathonian | H11 | EG-1 | 30.647° . 33.328° | 3.3 | 2.32 | -3.06 | 25.1 | -0.16 | 29.1 |
| H11-3 | Afriogryphaea costellata | bivalve | Middle Bathonian | H11 | EG-1 | 30.647° . 33.328° | 3.3 | 1.53 | -2.85 | 24.1 | -0.16 | 28.1 |
| H11-4 | Daghanirynchia cavignarii | brachiopod | Middle Bathonian | H11 | EG-1 | 30.647° . 33.328° | 3.3 | 0.85 | -3.63 | 27.8 | -0.16 | 32.0 |
| H11-5 | Daghanirynchia cavignarii | brachiopod | Middle Bathonian | H11 | EG-1 | 30.647° . 33.328° | 3.3 | 2.61 | -3.39 | 26.6 | -0.16 | 30.7 |
| H11-7 | Daghanirynchia daghamiensis | brachiopod | Late Bathonian | E13 | EG-3 | 30.698° . 33.387° | 3.3 | 1.13 | -2.79 | 23.8 | -0.16 | 27.8 |
| E13-1 | Burmimyrinchia cavignarii | brachiopod | Late Bathonian | E13 | EG-3 | 30.698° . 33.387° | 3.3 | 1.40 | -3.26 | 26.0 | -0.16 | 30.1 |
| E13-2 | Burmimyrinchia cavignarii | brachiopod | Late Bathonian | E13 | EG-3 | 30.698° . 33.387° | 3.3 | 1.41 | -3.29 | 26.2 | -0.16 | 30.2 |
| E13-3 | Afriogryphaea costellata | bivalve | Late Bathonian | E14 | EG-3 | 30.698° . 33.387° | 3.3 | 1.79 | -3.36 | 26.5 | -0.16 | 30.6 |
| E14-1 | Daghanirynchia daghamiensis | brachiopod | Early Callovian | M15 | EG-4 | 30.682° . 33.422° | 3.4 | 2.21 | -3.47 | 27.0 | -0.15 | 31.2 |
| M15-1 | Afriogryphaea costellata | bivalve | Early Callovian | M15 | EG-4 | 30.682° . 33.422° | 3.4 | 1.87 | -2.75 | 23.6 | -0.15 | 27.6 |
| M15-2 | Daghanirynchia angulocostata | brachiopod | Early Callovian | M16 | EG-4 | 30.682° . 33.422° | 3.4 | 1.57 | -3.13 | 25.4 | -0.15 | 29.5 |
| M16-2-ab | Daghanirynchia angulocostata | brachiopod | Middle Callovian | E16 | EG-3 | 30.698° . 33.387° | 3.4 | 1.56 | -3.26 | 26.0 | -0.15 | 30.1 |

Table S2 (continued) Results of the stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of samples from northern Africa and western Asia ($^1\text{H}_2\text{O}$ water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰ , $^2\delta^{18}\text{O}_{\text{sea}}$ value calculated for the palaeolatitude with equation (4), $^3\delta^{18}\text{O}_{\text{sea}}$ value calculated with equation (4) for the palaeolatitude).

| sample no. | taxonomy | group | age | bed | locality | coordinates | palaeolatitude | $\delta^{18}\text{O}$ | $\delta^{18}\text{O}_{\text{sea}}^1$ | $\delta^{18}\text{O}_{\text{sea}}^2$ | T (°C) ³ |
|---------------------------------|--------------------------------------|---------------------------------|------------------|--------|-------------------|-------------------|----------------|-----------------------|--------------------------------------|--------------------------------------|---------------------|
| M20-1 | <i>Africogryphaea costellata</i> | bivalve | Middle Callovian | M20 | EG-4 | 30.682° . 33.422° | 3.4 | 1.78 | -3.35 | 26.5 | -0.15 |
| E18-1 | <i>Burmithynchia cavagnarii</i> | brachiopod | Late Callovian | E18 | EG-3 | 30.698° . 33.387° | 3.4 | 1.32 | -3.32 | 26.3 | -0.15 |
| E18-2 | <i>Burmithynchia cavagnarii</i> | brachiopod | Late Callovian | E18 | EG-3 | 30.698° . 33.387° | 3.4 | 1.47 | -3.66 | 27.9 | -0.15 |
| E18-3 | <i>Burmithynchia cavagnarii</i> | brachiopod | Late Callovian | E18 | EG-3 | 30.698° . 33.387° | 3.4 | 1.39 | -3.31 | 26.3 | -0.15 |
| H16-1 | <i>Daghanithynchia angulocostata</i> | brachiopod | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 0.80 | -3.17 | 25.6 | -0.15 |
| H16-2 | <i>Daghanithynchia angulocostata</i> | brachiopod | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 1.66 | -3.47 | 27.0 | -0.15 |
| H16-3 | <i>Daghanithynchia angulocostata</i> | brachiopod | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 1.29 | -3.48 | 27.0 | -0.15 |
| H16-5 | <i>Nanogyra nana</i> | bivalve | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 1.97 | -3.24 | 25.9 | -0.15 |
| H16-6 | <i>Nanogyra nana</i> | bivalve | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 2.32 | -2.90 | 24.3 | -0.15 |
| E22-1 | <i>Daghanithynchia angulocostata</i> | brachiopod | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 1.48 | -3.17 | 25.6 | -0.18 |
| E22-2 | <i>Daghanithynchia angulocostata</i> | brachiopod | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 1.22 | -2.96 | 24.6 | -0.18 |
| E22-3 | <i>Daghanithynchia angulocostata</i> | brachiopod | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 1.49 | -3.12 | 25.4 | -0.18 |
| E22-4 | <i>Nanogyra nana</i> | bivalve | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 1.43 | -2.16 | 21.0 | -0.18 |
| E22-5 | <i>Gryphaeligmus jabbokensis</i> | bivalve | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 1.74 | -3.04 | 25.0 | -0.18 |
| <i>sediment & cement</i> | | | | | | | | | | | |
| A7-2S | - | | Late Bajocian | A7 | EG-2 | 30.677° . 33.355° | 3.3 | -0.90 | -5.82 | - | - |
| H6-3S | - | | Middle Bathonian | H6 | EG-1 | 30.647° . 33.328° | 3.3 | 0.12 | -2.76 | - | - |
| M14-3C | - | | Middle Bathonian | A14 | EG-2 | 30.677° . 33.355° | 3.3 | 1.23 | -8.22 | - | - |
| M11-1S | - | | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | -0.53 | -6.38 | - | - |
| M11-4S | - | | Middle Bathonian | M11 | EG-4 | 30.682° . 33.422° | 3.3 | -1.12 | -6.51 | - | - |
| E9-2C | - | | Middle Bathonian | E19 | EG-3 | 30.698° . 33.387° | 3.3 | 0.61 | -6.19 | - | - |
| E13-1C | - | | Late Bathonian | E13 | EG-3 | 30.698° . 33.387° | 3.3 | 0.48 | -6.72 | - | - |
| H16-1C | - | | Late Callovian | H16 | EG-1 | 30.647° . 33.328° | 3.4 | 1.15 | -9.80 | - | - |
| E22-3C | - | | E. Kimmeridgian | E22 | EG-3 | 30.698° . 33.387° | 2.8 | 0.82 | -8.09 | - | - |
| <i>poorly preserved fossils</i> | | | | | | | | | | | |
| H10-1D | <i>Africogryphaea costellata</i> | bivalve | Middle Bathonian | H10 | EG-1 | 30.647° . 33.328° | 3.3 | 0.08 | -5.58 | - | - |
| Israel | | | | | | | | | | | |
| IS17-060 | well-preserved fossils | rhynchonellid | Late Bajocian | IS-8 | 30.627° . 34.842° | 2.7 | 2.88 | -4.03 | 29.8 | -0.18 | 33.9 |
| IS17-061 | | oyster | Late Bajocian | IS-9 | 30.617° . 34.836° | 2.7 | 1.14 | -4.09 | 30.0 | -0.18 | 34.1 |
| IS17-035 | | <i>Somalithynchia africana</i> | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 1.01 | -3.84 | 28.8 | -0.17 | 33.0 |
| IS17-036 | | <i>Eligmus</i> sp. | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.92 | -2.81 | 23.9 | -0.17 | 27.8 |
| IS17-037 | | oyster | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.17 | -3.46 | 27.0 | -0.17 | 31.0 |
| IS17-038 | | oyster | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.73 | -3.39 | 26.6 | -0.17 | 30.7 |
| IS17-039 | | <i>Nanogyra</i> sp. | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 3.10 | -3.34 | 26.4 | -0.17 | 30.4 |
| IS17-040 | | <i>Nanogyra</i> sp. | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.96 | -3.20 | 25.7 | -0.17 | 29.7 |
| IS17-041-2A | | <i>Nanogyra</i> sp. | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 3.09 | -3.53 | 27.3 | -0.17 | 31.4 |
| IS17-042 | | <i>Ostrea dubiensis</i> | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 3.10 | -3.26 | 26.0 | -0.17 | 30.0 |
| IS17-044 | | rhynchonellid | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.18 | -3.47 | 27.0 | -0.17 | 31.1 |
| IS17-045 | | oyster | Middle Callovian | gad-11 | 30.948° . 35.000° | 3.0 | 2.27 | -3.40 | 26.7 | -0.17 | 30.7 |
| IS17-001-AB | | <i>Burmithynchia jirbaensis</i> | Late Callovian | IS-2 | 30.941° . 34.983° | 3.0 | 2.38 | -3.91 | 29.2 | -0.17 | 33.3 |

Table S2 (continued) Results of the stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of samples from northern Africa and western Asia (¹Water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰, ² $\delta^{18}\text{O}_{\text{sea}}$ value calculated for the palaeolatitude with equation (4), ³Water temperatures based on the equation (4) for the palaeolatitude).

| sample no. | taxonomy | group | age | bed | locality | coordinates | palaearc latitude | T (°C) ¹ | $\delta^{18}\text{O}_{\text{sea}}$ | $\delta^{18}\text{O}_{\text{sea}}^{\text{2}}$ T (°C) ³ | |
|-------------------------------|--------------------------------------|-----------------------|------------------|-----------|----------|-------------------|-------------------|---------------------|------------------------------------|---|------------|
| IS17-004 | <i>Gryphaea</i> sp. rhynchonellid | bivalve brachiopod | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -3.58 | 27.5 | -0.17 31.6 | |
| IS17-005 | rhynchonellid | brachiopod | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -3.27 | 26.1 | -0.17 30.1 | |
| IS17-006 | | brachiopod | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -4.34 | 31.3 | -0.17 35.5 | |
| IS17-008 | <i>Nanogyra</i> sp. | bivalve | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -3.67 | 28.0 | -0.17 32.1 | |
| IS17-009 | <i>Somalirhynchia africana</i> | brachiopod | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -4.20 | 30.6 | -0.17 34.8 | |
| IS17-012 | <i>Nanogyra</i> sp. | bivalve | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | -3.23 | 25.9 | -0.17 29.9 | |
| IS17-020 | <i>Actinostreon</i> sp. | bivalve | Late Callovian | gad-47 | IS-3 | 30.941° . 34.982° | 3.0 | -3.86 | 28.9 | -0.17 33.1 | |
| IS17-055 | <i>Ostrea</i> sp. | bivalve | Late Callovian | gad-48 | IS-4 | 30.941° . 34.981° | 3.0 | -2.97 | 24.7 | -0.17 28.6 | |
| IS17-057 | <i>Ostrea</i> sp. | bivalve | Late Callovian | gad-48 | IS-4 | 30.941° . 34.981° | 3.0 | -3.29 | 26.2 | -0.17 30.2 | |
| IS17-058 | <i>Nanogyra</i> sp. | bivalve | Late Callovian | gad-48 | IS-4 | 30.941° . 34.981° | 3.0 | -2.64 | 23.2 | -0.17 27.0 | |
| IS17-046 | <i>Burmirhynchia jirbaensis</i> | brachiopod | Late Callovian | gad-60/61 | IS-5 | 30.942° . 34.980° | 3.0 | -3.07 | 25.1 | -0.17 29.1 | |
| IS17-059 | oyster | bivalve | Late Callovian | gad-62 | IS-6 | 30.943° . 34.980° | 3.0 | 2.68 | -3.58 | 27.5 | -0.17 31.6 |
| IS17-023-AB | | | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 3.49 | -3.30 | 26.2 | -0.17 30.2 |
| IS17-025 | | | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 3.93 | -3.56 | 27.4 | -0.17 31.5 |
| IS17-026 | | | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 4.71 | -3.31 | 26.3 | -0.17 30.3 |
| IS17-028-AB | <i>Actinostreon</i> sp. | bivalve | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 4.28 | -3.27 | 26.0 | -0.17 30.1 |
| IS17-029 | <i>Nanogyra</i> sp. | bivalve | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 3.88 | -3.20 | 25.7 | -0.17 29.7 |
| IS17-032 | rhynchonellid | brachiopod | Late Callovian | gad-63 | IS-7 | 30.943° . 34.980° | 3.0 | 2.35 | -3.59 | 27.6 | -0.17 31.7 |
| IS17-066 | <i>Exogyra</i> sp. | bivalve | Early Cretaceous | ram-Cr | IS-10 | 30.624° . 34.842° | 2.0 | 2.81 | -3.14 | 25.5 | -0.22 29.2 |
| IS17-067 | <i>Exogyra</i> sp. | bivalve | Early Cretaceous | ram-Cr | IS-10 | 30.624° . 34.842° | 2.0 | 2.73 | -3.36 | 26.5 | -0.22 30.3 |
| IS17-068 | <i>Exogyra</i> sp. | bivalve | Early Cretaceous | ram-Cr | IS-10 | 30.624° . 34.842° | 2.0 | 3.27 | -3.91 | 29.2 | -0.22 33.1 |
| IS17-069 | <i>Exogyra</i> sp. | bivalve | Early Cretaceous | ram-Cr | IS-10 | 30.624° . 34.842° | 2.0 | 2.72 | -3.90 | 29.1 | -0.22 33.0 |
| <i>sediment & cement</i> | | | | | | | | | | | |
| IS17-035Sed | - | sediment fill | Middle Callovian | gad-11 | IS-1 | 30.948° . 35.000° | 3.0 | 2.11 | -4.51 | - | |
| IS17-003Sed | - | sediment fill | Late Callovian | gad-43 | IS-2 | 30.941° . 34.983° | 3.0 | 1.86 | -3.81 | - | |
| IS17-013Cem | - | cement fill | Late Callovian | gad-47 | IS-3 | 30.941° . 34.982° | 3.0 | 2.29 | -6.74 | - | |
| IS17-014Cem | - | cement fill | Late Callovian | gad-47 | IS-3 | 30.941° . 34.982° | 3.0 | 1.86 | -7.84 | - | |
| IS17-036Sed | - | sediment fill | Middle Callovian | gad-11 | IS-1 | 30.948° . 35.000° | 3.0 | 2.01 | -3.45 | - | |
| <i>Jordan</i> | | | | | | | | | | | |
| <i>well-preserved fossils</i> | | | | | | | | | | | |
| JO18-028 | <i>Africogryphaea costellata</i> | bivalve | | | JO-2 | 32.125° . 35.654° | 1.7 | 2.15 | -2.55 | 22.7 | |
| JO18-029 | <i>Africogryphaea costellata</i> | bivalve | | | JO-2 | 32.125° . 35.654° | 1.7 | 2.22 | -3.04 | 25.0 | |
| JO18-033 | <i>Africogryphaea costellata</i> | bivalve | | | JO-3 | 32.175° . 35.705° | 1.8 | 2.13 | -2.69 | 23.4 | |
| JO18-034 | <i>Africogryphaea costellata</i> | bivalve | | | JO-3 | 32.175° . 35.705° | 1.8 | 2.33 | -2.63 | 23.1 | |
| JO18-024 | <i>Africogryphaea costellata</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 1.80 | -2.44 | 22.2 | |
| JO18-025 | <i>Africogryphaea costellata</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 2.14 | -2.78 | 23.8 | |
| JO18-027 | <i>Africogryphaea costellata</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 1.65 | -3.54 | 27.4 | |
| JO18-030 | <i>Gryphaeligmus jabboekensis</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 2.08 | -3.67 | 28.0 | |
| JO18-031 | <i>Gryphaeligmus jabboekensis</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 1.58 | -3.07 | 25.1 | |
| JO18-032 | <i>Gryphaeligmus jabboekensis</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 2.22 | -3.48 | 27.1 | |
| JO18-035 | <i>Eligmus rollandi</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 1.95 | -3.47 | 27.0 | |
| JO18-036 | <i>Eligmus rollandi</i> | bivalve | | | JO-1 | 32.108° . 35.660° | 1.7 | 1.59 | -3.12 | 25.4 | |

| sample no. | taxonomy | group | age | bed | locality | coordinates | paleolatitude | $\delta^{18}\text{O}$ | $\delta^{13}\text{C}$ | T (°C) ¹ | $\delta^{18}\text{O}_{\text{sea}}^{\text{2}}$ | T (°C) ³ |
|---------------------------------|---------------------------------|---------|-----------|------|----------|-------------------|---------------|-----------------------|-----------------------|---------------------|---|---------------------|
| JO18-003 | <i>Gryphaeolina jabbokensis</i> | bivalve | Bathonian | s-2 | JO-4 | 32.189° . 35.713° | 1.8 | -2.88 | 24.3 | -0.23 | 27.9 | |
| JO18-004 | <i>Gryphaeolina jabbokensis</i> | bivalve | Bathonian | s-2 | JO-4 | 32.189° . 35.713° | 1.8 | -3.08 | 25.2 | -0.23 | 28.9 | |
| JO18-005 | <i>Gryphaeolina jabbokensis</i> | bivalve | Bathonian | s-2 | JO-4 | 32.189° . 35.713° | 1.8 | 2.17 | -3.50 | 27.2 | -0.23 | 31.0 |
| JO18-016 | <i>Afriogryphaea costellata</i> | bivalve | Bathonian | s-h | JO-4 | 32.189° . 35.713° | 1.8 | 1.69 | -3.06 | 25.1 | -0.23 | 28.8 |
| JO18-018 | <i>Afriogryphaea costellata</i> | bivalve | Bathonian | s-h | JO-4 | 32.189° . 35.713° | 1.8 | 2.06 | -2.98 | 24.7 | -0.23 | 28.4 |
| JO18-023 | <i>Afriogryphaea costellata</i> | bivalve | Bathonian | s-h | JO-4 | 32.189° . 35.713° | 1.8 | 2.10 | -2.45 | 22.3 | -0.23 | 25.8 |
| JO18-019 | <i>Gryphaeolina jabbokensis</i> | bivalve | Bathonian | s-4 | JO-4 | 32.189° . 35.713° | 1.8 | 1.99 | -4.34 | 31.3 | -0.23 | 35.2 |
| JO18-013 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-1 | JO-5 | 32.191° . 35.800° | 1.8 | 1.78 | -4.00 | 29.6 | -0.23 | 33.5 |
| JO18-007 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-5 | JO-5 | 32.191° . 35.800° | 1.8 | 1.83 | -4.10 | 30.1 | -0.23 | 34.0 |
| JO18-001 | <i>Eligmus asiaticus</i> | bivalve | Callovian | r-11 | JO-1 | 32.108° . 35.660° | 1.8 | 2.13 | -3.99 | 29.5 | -0.23 | 33.4 |
| <i>sediment & cement</i> | | | | | | | | | | | | |
| JO18-003Sed | - | | | | | | | | | | | |
| JO18-016Sed | - | | | | | | | | | | | |
| <i>poorly preserved fossils</i> | | | | | | | | | | | | |
| JO18-020 | <i>Eligmus rollandi</i> | bivalve | Bathonian | h-6 | JO-3 | 32.175° . 35.705° | 1.8 | 2.35 | -2.39 | - | - | - |
| JO18-006 | <i>Gryphaeolina jabbokensis</i> | bivalve | Bathonian | s-2 | JO-4 | 32.189° . 35.713° | 1.8 | 1.66 | -3.19 | - | - | - |
| JO18-015 | <i>Afriogryphaea costellata</i> | bivalve | Bathonian | s-h | JO-4 | 32.189° . 35.713° | 1.8 | 1.08 | -3.28 | - | - | - |
| JO18-021 | <i>Afriogryphaea costellata</i> | bivalve | Bathonian | s-h | JO-4 | 32.189° . 35.713° | 1.8 | 0.95 | -3.51 | - | - | - |
| JO18-012 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-1 | JO-5 | 32.191° . 35.800° | 1.8 | 1.67 | -3.58 | - | - | - |
| JO18-014 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-1 | JO-5 | 32.191° . 35.800° | 1.8 | 1.36 | -4.19 | - | - | - |
| JO18-008 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-5 | JO-5 | 32.191° . 35.800° | 1.8 | 1.32 | -3.38 | - | - | - |
| JO18-009 | <i>Gryphaeolina jabbokensis</i> | bivalve | Callovian | t-5 | JO-5 | 32.191° . 35.800° | 1.8 | 1.07 | -3.36 | - | - | - |

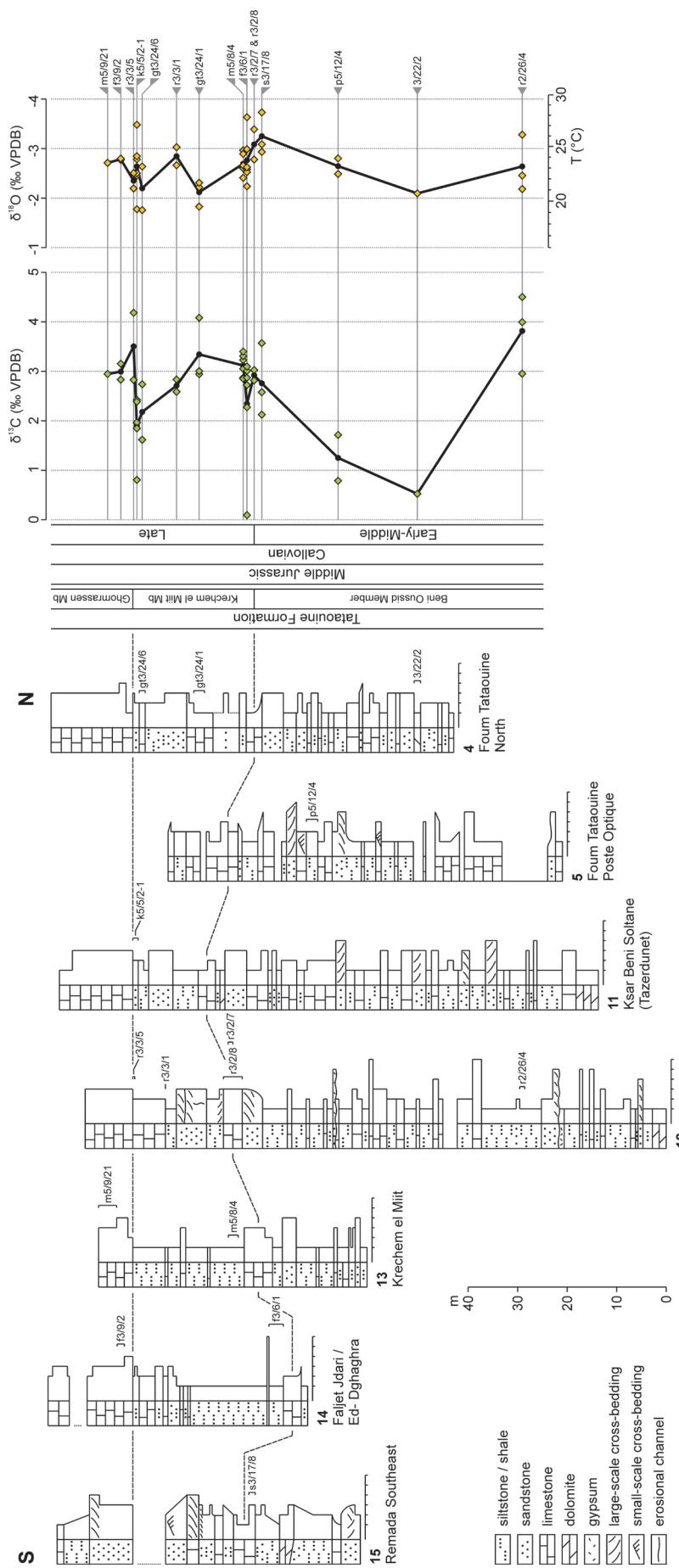


Fig. S3 Results of stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of Callovian brachiopod shells from Tunisia (sections modified after Holzapfel, 1998). Scale for water temperatures is based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰ .

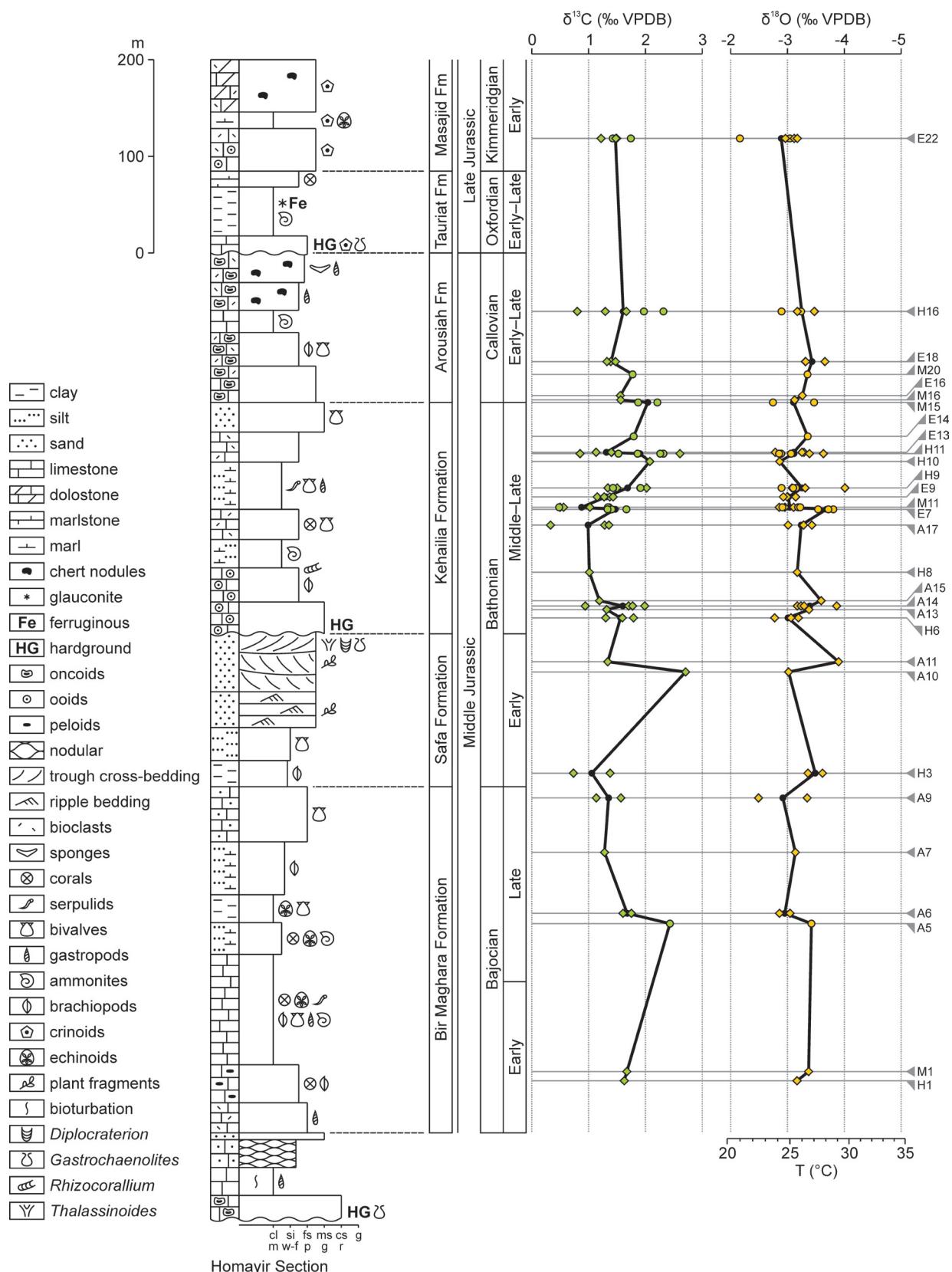


Fig. S4 Results of stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of Bajocian-Kimmeridgian bivalve and brachiopod shells from Egypt (modified after Abdelhady, 2014; Alberti et al., 2017). Scale for water temperatures is based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰.

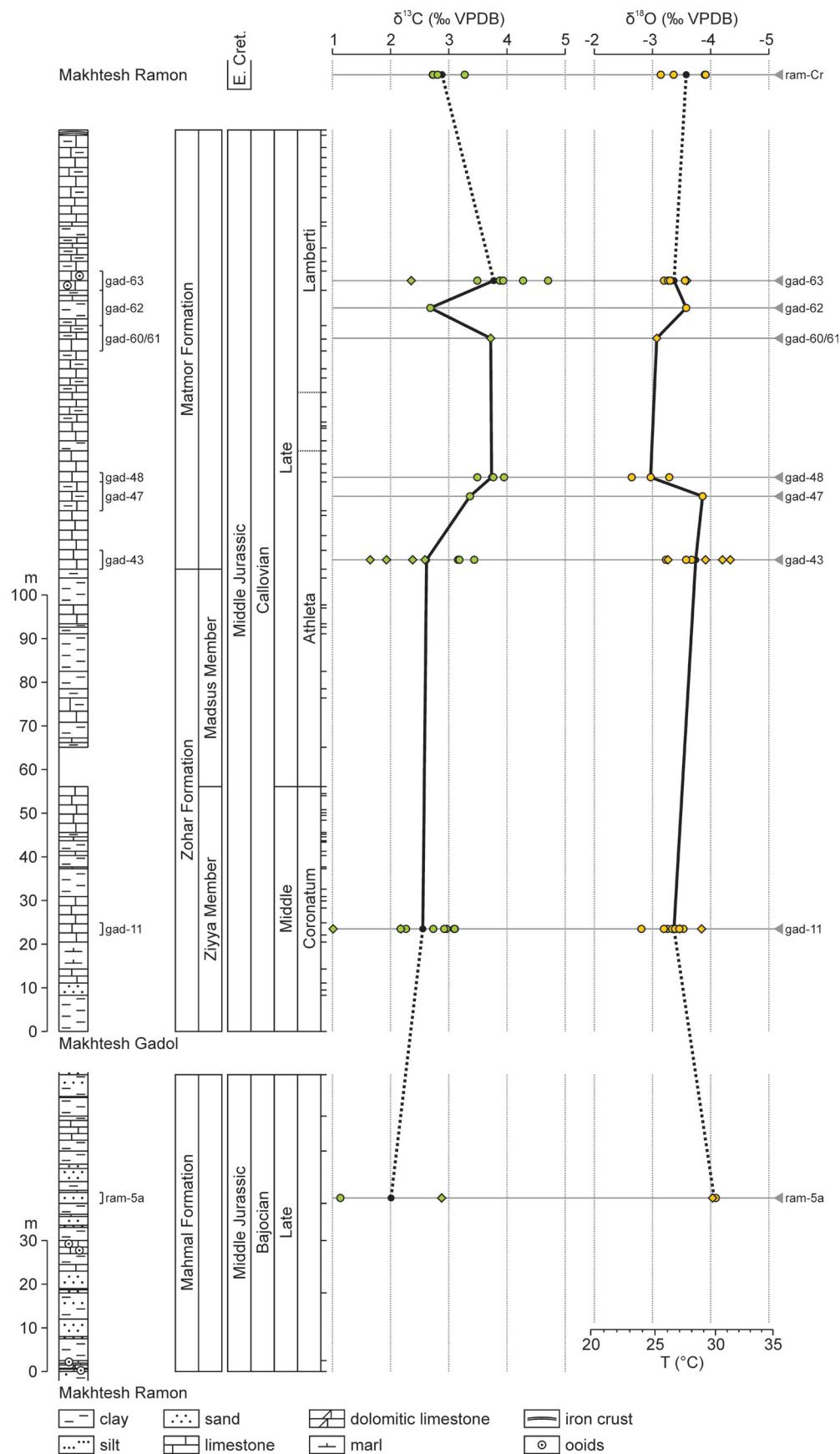


Fig. S5 Results of stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analyses of Bajocian-Early Cretaceous bivalve and brachiopod shells from Israel (sections modified after Goldberg, 1963; Parnes, 1981; Hirsch, 2005). Scale for water temperatures is based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰.

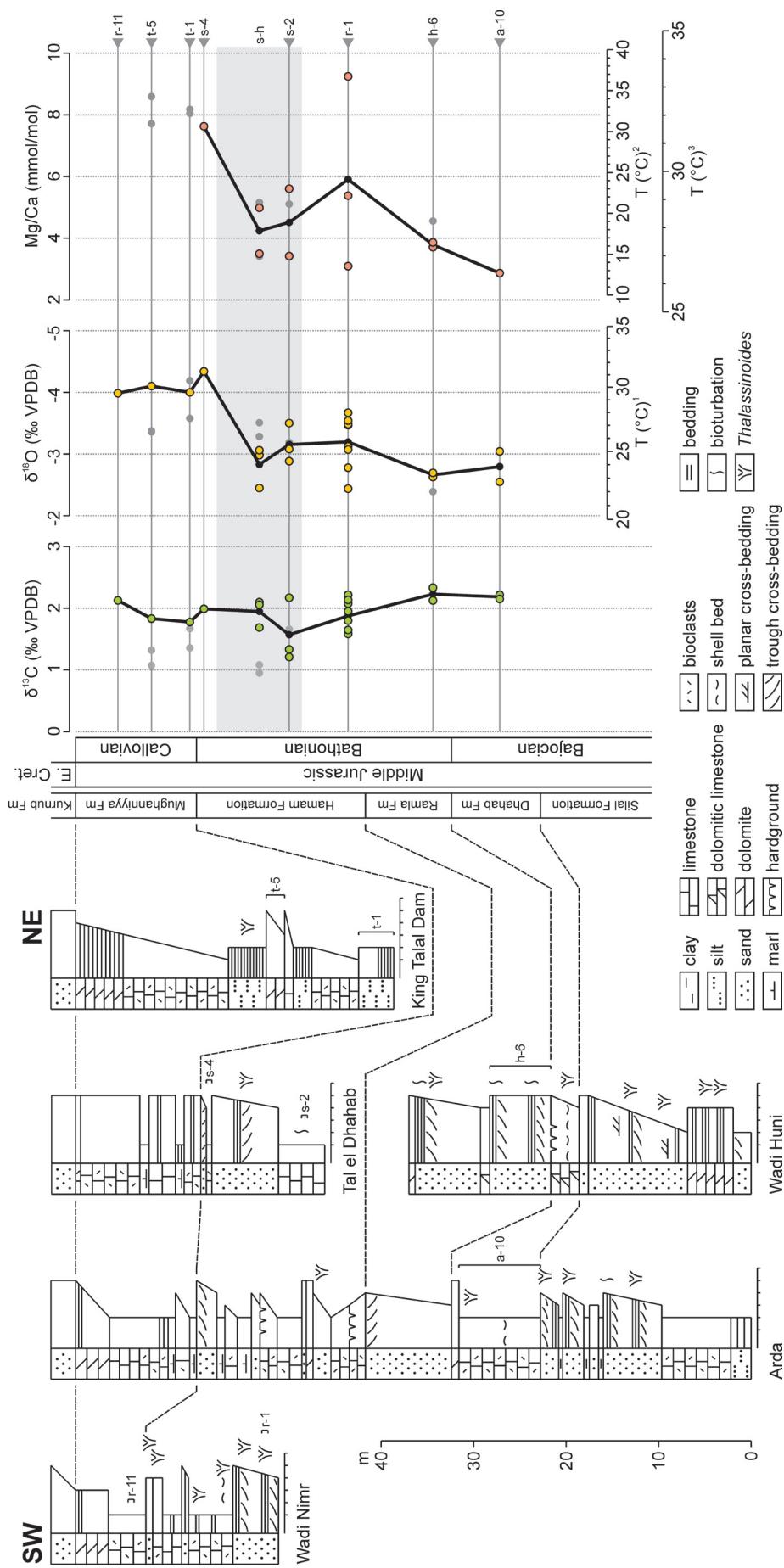


Fig. S6 Results of stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) and trace element (Mg/Ca) analyses of Bajocian-Callovian bivalve shells from Jordan (sections modified after Ahmad, 1999). Scales for water temperatures are based on (1) the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰, (2) the equation of Mouchi et al. (2013), or (3) equation (S2) as discussed in the text.

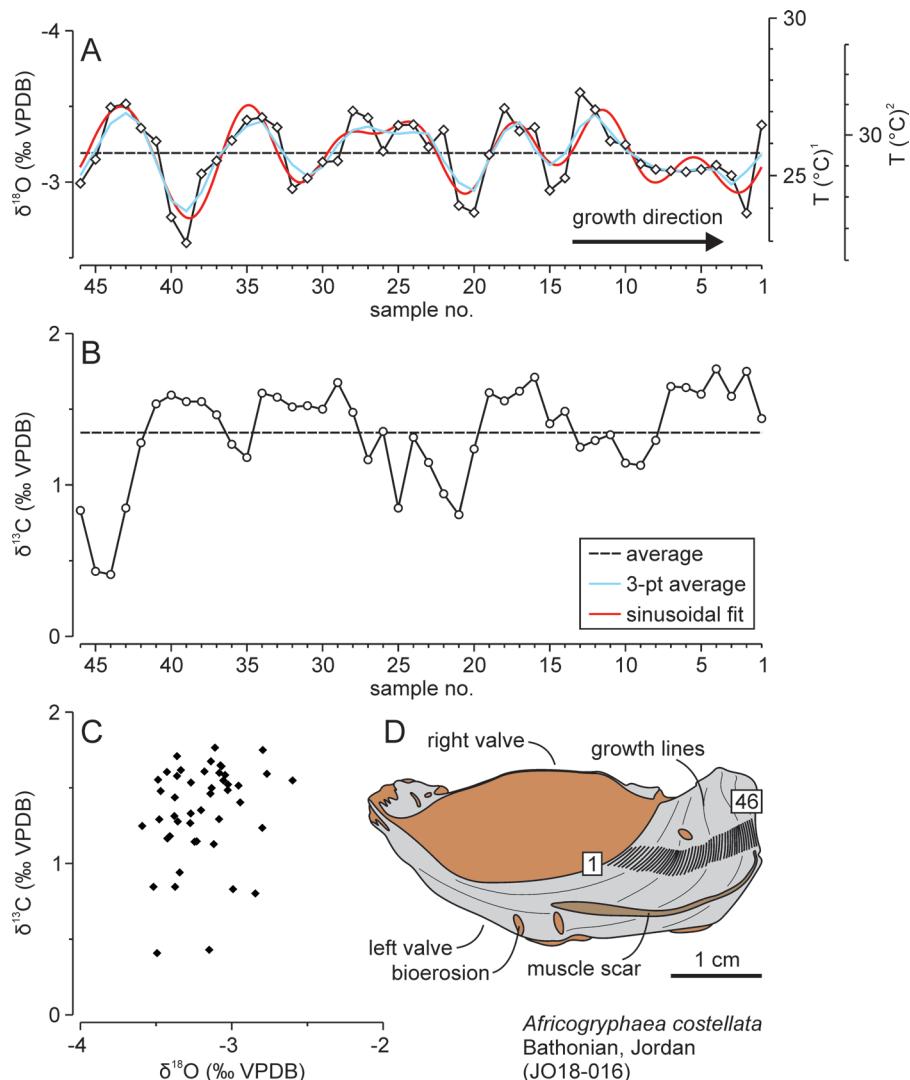


Fig. S7 Results of the high-resolution stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analysis of a specimen of *Africogryphaea costellata* from the Bathonian of Jordan. **A.** The measured $\delta^{18}\text{O}_{\text{shell}}$ values show a cyclic pattern. Scale for water temperatures is based on the equation of Anderson and Arthur (1983) and (1) a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰ or (2) a $\delta^{18}\text{O}_{\text{sea}}$ value of -0.23 ‰ corresponding to a palaeolatitude of 1.8 °N. **B.** High-resolution $\delta^{13}\text{C}$ data show fluctuating values not corresponding to the cycles in the $\delta^{18}\text{O}_{\text{shell}}$ values. **C.** Cross-plot of $\delta^{18}\text{O}$ versus $\delta^{13}\text{C}$ values showing no statistically significant correlation. **D.** Sketch of the used specimen of *Africogryphaea costellata* with the position of the individual samples.

ages (Fig. S6). Their $\delta^{18}\text{O}$ values show a more or less steady decrease through time with an average of -2.80 ‰ in the Bajocian and -3.13 ‰ in the Bathonian. Towards the Callovian, $\delta^{18}\text{O}$ values decrease again strongly to an average of -4.03 ‰. In contrast, the $\delta^{13}\text{C}$ record does not show very strong fluctuations from the Bajocian (average: 2.18 ‰) to the Bathonian (average: 1.88 ‰) and Callovian (average: 1.91 ‰). In addition to stable isotope analyses of individual shells, one oyster shell (*Africogryphaea costellata*; JO18-016) of the Bathonian was used for a high-resolution analysis with 46 samples taken perpendicular to the growth lines (Fig. S7; Table S3). Results from this study show a cyclic pattern in the $\delta^{18}\text{O}$ values around an average of -3.19 ‰ (minimum: -3.59 ‰; maximum: -2.60 ‰). A total of five cycles can be identified. The $\delta^{13}\text{C}$ values vary around an average of 1.35 ‰ (minimum: 0.41 ‰; maximum: 1.77 ‰), but their fluctuations do not correspond to those in the $\delta^{18}\text{O}$ values. In fact, $\delta^{18}\text{O}$ and $\delta^{13}\text{C}$ values show no correlation ($r_s = 0.27$; $p = 0.07$).

Table S3 Results of the high-resolution stable isotope ($\delta^{18}\text{O}$, $\delta^{13}\text{C}$) analysis of a specimen of *Africogryphaea costellata* from the Bathonian of Jordan.¹Water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰,²Water temperatures based on the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value of -0.23 ‰ corresponding to a palaeolatitude of 1.8 °N).

| sample no. | $\delta^{13}\text{C}$ | $\delta^{18}\text{O}$ | T (°C) ¹ | T (°C) ² |
|------------|-----------------------|-----------------------|---------------------|---------------------|
| JO18-16-1 | 1.44 | -3.38 | 26.6 | 30.3 |
| JO18-16-2 | 1.75 | -2.80 | 23.9 | 27.5 |
| JO18-16-3 | 1.59 | -3.04 | 25.0 | 28.7 |
| JO18-16-4 | 1.77 | -3.11 | 25.3 | 29.0 |
| JO18-16-5 | 1.60 | -3.08 | 25.2 | 28.9 |
| JO18-16-6 | 1.64 | -3.07 | 25.1 | 28.8 |
| JO18-16-7 | 1.65 | -3.08 | 25.2 | 28.8 |
| JO18-16-8 | 1.29 | -3.08 | 25.2 | 28.9 |
| JO18-16-9 | 1.13 | -3.12 | 25.4 | 29.1 |
| JO18-16-10 | 1.14 | -3.25 | 26.0 | 29.7 |
| JO18-16-11 | 1.33 | -3.27 | 26.1 | 29.8 |
| JO18-16-12 | 1.29 | -3.48 | 27.1 | 30.8 |
| JO18-16-13 | 1.25 | -3.59 | 27.6 | 31.4 |
| JO18-16-14 | 1.49 | -3.03 | 24.9 | 28.6 |
| JO18-16-15 | 1.40 | -2.95 | 24.5 | 28.2 |
| JO18-16-16 | 1.71 | -3.36 | 26.5 | 30.3 |
| JO18-16-17 | 1.62 | -3.34 | 26.4 | 30.1 |
| JO18-16-18 | 1.56 | -3.49 | 27.1 | 30.9 |
| JO18-16-19 | 1.61 | -3.18 | 25.6 | 29.4 |
| JO18-16-20 | 1.24 | -2.80 | 23.9 | 27.5 |
| JO18-16-21 | 0.80 | -2.85 | 24.1 | 27.7 |
| JO18-16-22 | 0.94 | -3.34 | 26.4 | 30.2 |
| JO18-16-23 | 1.15 | -3.23 | 25.9 | 29.6 |
| JO18-16-24 | 1.31 | -3.38 | 26.6 | 30.3 |
| JO18-16-25 | 0.85 | -3.38 | 26.6 | 30.3 |
| JO18-16-26 | 1.35 | -3.20 | 25.8 | 29.5 |
| JO18-16-27 | 1.17 | -3.43 | 26.8 | 30.6 |
| JO18-16-28 | 1.48 | -3.47 | 27.0 | 30.8 |
| JO18-16-29 | 1.68 | -3.14 | 25.5 | 29.2 |
| JO18-16-30 | 1.50 | -3.13 | 25.4 | 29.1 |
| JO18-16-31 | 1.52 | -3.03 | 24.9 | 28.6 |
| JO18-16-32 | 1.52 | -2.96 | 24.6 | 28.3 |
| JO18-16-33 | 1.58 | -3.36 | 26.5 | 30.3 |
| JO18-16-34 | 1.61 | -3.43 | 26.8 | 30.6 |
| JO18-16-35 | 1.18 | -3.41 | 26.7 | 30.5 |
| JO18-16-36 | 1.27 | -3.28 | 26.1 | 29.8 |
| JO18-16-37 | 1.46 | -3.14 | 25.5 | 29.2 |
| JO18-16-38 | 1.55 | -3.06 | 25.1 | 28.7 |
| JO18-16-39 | 1.55 | -2.60 | 22.9 | 26.5 |
| JO18-16-40 | 1.59 | -2.77 | 23.7 | 27.4 |
| JO18-16-41 | 1.54 | -3.27 | 26.1 | 29.8 |
| JO18-16-42 | 1.28 | -3.36 | 26.5 | 30.2 |
| JO18-16-43 | 0.85 | -3.52 | 27.2 | 31.0 |
| JO18-16-44 | 0.41 | -3.49 | 27.1 | 30.9 |
| JO18-16-45 | 0.43 | -3.15 | 25.5 | 29.2 |
| JO18-16-46 | 0.83 | -2.99 | 24.8 | 28.4 |

Table S4 Results of trace element (Mg/Ca, Sr/Ca) analyses of well-preserved bivalves from Jordan (¹Water temperatures based on the equation of Mouchi et al. (2013), ²Water temperatures based on equation (S2)).

| sample no. | taxonomy | Mg/Ca | T (°C) ¹ | T (°C) ² | Sr/Ca |
|------------|----------------------------------|-------|---------------------|---------------------|-------|
| JO18-003 | <i>Gryphaeligmus jabbokensis</i> | 3.42 | 14.8 | 27.0 | 0.73 |
| JO18-004 | <i>Gryphaeligmus jabbokensis</i> | 5.60 | 23.0 | 29.4 | 0.74 |
| JO18-018 | <i>Africogryphaea costellata</i> | 3.49 | 15.1 | 27.1 | 0.72 |
| JO18-019 | <i>Gryphaeligmus jabbokensis</i> | 7.63 | 30.6 | 31.6 | 0.85 |
| JO18-023 | <i>Africogryphaea costellata</i> | 4.98 | 20.7 | 28.7 | 0.58 |
| JO18-024 | <i>Africogryphaea costellata</i> | 3.09 | 13.5 | 26.6 | 0.70 |
| JO18-027 | <i>Africogryphaea costellata</i> | 9.25 | 36.7 | 33.4 | 2.67 |
| JO18-028 | <i>Africogryphaea costellata</i> | 2.87 | 12.7 | 26.4 | 0.82 |
| JO18-033 | <i>Africogryphaea costellata</i> | 3.86 | 16.4 | 27.5 | 0.71 |
| JO18-034 | <i>Africogryphaea costellata</i> | 3.71 | 15.9 | 27.3 | 0.75 |
| JO18-032 | <i>Gryphaeligmus jabbokensis</i> | 5.38 | 22.2 | 29.1 | 0.80 |

Trace element analyses

Eleven seemingly well-preserved shells of the bivalves *Africogryphaea costellata* and *Gryphaeligmus jabbokensis* from Jordan were analysed for their Mg/Ca- and Sr/Ca-ratios (Table S4). Measured Mg/Ca-ratios are between 2.87 and 9.25 mmol/mol and Sr/Ca-ratios vary between 0.58 and 0.85 mmol/mol with one outlier at 2.67 mmol/mol. Mg/Ca-ratios show a statistically significant negative correlation with $\delta^{18}\text{O}$ values ($r_s = -0.75$; $p = 0.007$), whether considered individually for each species or together (Fig. S2B). It should be noted, however, that in the case of *Africogryphaea costellata*, this correlation strongly depends on one sample with (suspiciously?) high Sr/Ca- and Mg/Ca-ratios. Similar to the $\delta^{18}\text{O}$ values, Mg/Ca-ratios exhibit a prominent trend through time. Their record starts with a minimum of 2.87 mmol/mol in the Bajocian, increasing to higher values over most of the Bathonian (average: 5.04 mmol/mol), before reaching a Mg/Ca-ratio of 7.63 mmol/mol just before the Bathonian-Callovian boundary (Fig. S6).

Water temperatures of the individual study areas

The reconstruction of absolute water temperatures based on $\delta^{18}\text{O}_{\text{shell}}$ values depends on a number of variables. Diagenetic alteration leading to a change in the stable isotope composition, vital effects of the used fossil taxa, and environmental factors are the most prominent. As explained above, steps have been taken to identify poorly preserved fossils in the collections. Furthermore, sedimentology and palaeoecology show that the study sites represent fully-marine settings above the thermocline, limiting the potential influence of salinity changes (e.g., via river discharge). Some climate models reconstructed seasonal rainfall patterns (and possibly upwelling) along the northern coast of Gondwana during the Jurassic which could influence local $\delta^{18}\text{O}_{\text{sea}}$ values (e.g., Price et al., 1995; Sellwood and Valdes, 2006). In contrast, high-resolution stable isotope analyses conducted on bivalve shells from Jordan and Egypt (compare Fig. S7 and Alberti et al., 2017) show very weak seasonal changes in stable isotope values ($\delta^{18}\text{O}$ and $\delta^{13}\text{C}$). These results disagree with the presence of strong seasonal rainfall or upwelling patterns which would be expected to result in much stronger fluctuations (compare Sadatzki et al., 2019). The location of the study sites in direct contact with the Tethys Ocean also suggest that conditions corresponded to an open-ocean setting. Finally, previous studies showed very similar stable isotope values for bivalves and rhynchonellid brachiopods sampled from the same horizons supporting the validity of combining data of both taxonomic groups (e.g., Alberti et al., 2012a).

The following water temperatures were reconstructed with the equation of Anderson and Arthur (1983) and a $\delta^{18}\text{O}_{\text{sea}}$ value determined with equation (4) for the individual palaeolatitudes (Table S2). There is a series of other equations available in literature (e.g., Epstein et al., 1953; Craig, 1965; Leng and Marshall, 2004; Takayanagi et al., 2013; Brand et al., 2013), but for the relatively low $\delta^{18}\text{O}_{\text{shell}}$ values recorded from the equatorial regions in this study, results are quite comparable (compare Fig. S8). Therefore, we decided to apply the equation of Anderson and Arthur (1983), which is very widely used in literature and thereby facilitates comparisons between articles.

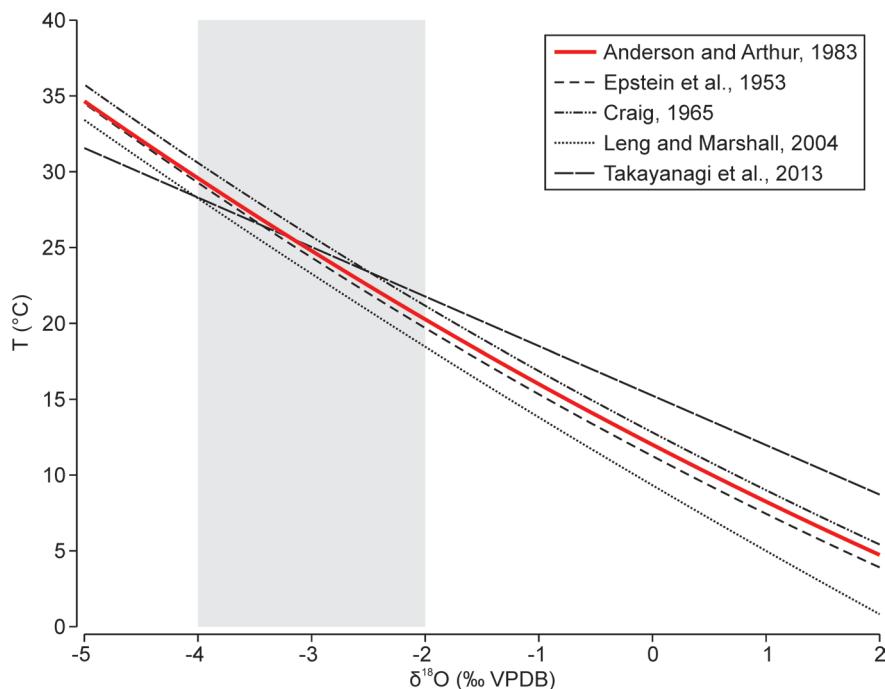


Fig. S8 A comparison of different equations used for temperature reconstructions based on stable oxygen isotope values shows that results are comparatively similar for lower $\delta^{18}\text{O}_{\text{shell}}$ values (=higher water temperatures), but differ more strongly at higher $\delta^{18}\text{O}_{\text{shell}}$ values (=lower water temperatures). Most samples from the equatorial study sites show $\delta^{18}\text{O}_{\text{shell}}$ values between -2 and -4 ‰. In this range, the widely used equation of Anderson and Arthur (1983) seems to be a reasonable fit. Temperatures in the graph were calculated for a $\delta^{18}\text{O}_{\text{sea}}$ value of -1 ‰.

Furthermore, temperatures reconstructed based on Anderson and Arthur (1983) fall well within the center compared to the other equations. Using a different equation would not change the conclusions of the study markedly.

Only four well-preserved shells were available from Morocco (palaeolatitude 26-28 °N) which point to water temperatures around 28.0 °C in the ?Toarcian-Aalenian, 23.9 °C in the Bajocian, 22.7 °C in the Callovian, and 22.7 °C in the Oxfordian. This seems to agree with reconstructions by Martin-Garin et al. (2010) for the Kimmeridgian of Morocco, who proposed water temperatures of around 24.6 °C based on stable oxygen isotope analyses of three brachiopods. However, Martin-Garin et al. (2010) used a $\delta^{18}\text{O}_{\text{sea}}$ value of -1.2 ‰ for their temperature calculations, which seems to be too low for a palaeolatitude of around 26 °N. If their original data is used with the method suggested within the present study, the Kimmeridgian water temperatures were much warmer (around an average of 30.6 °C).

Stable isotope analyses of 41 Callovian brachiopod shells of Tunisia (palaeolatitude 13-14 °N) point to water temperatures around an average of 28.3 °C (range: 24.0 to 33.6 °C). Water temperatures fluctuate considerably throughout the Callovian, but do not show a prominent long-term trend (Fig. 3A).

Results based on 73 well-preserved brachiopod and bivalve shells of Egypt (palaeolatitude 2-4 °N) indicate water temperatures around 29.3 °C in the Bajocian (range: 26.4 to 30.9 °C), 29.9 °C in the Bathonian (range: 27.7 to 33.9 °C), 30.2 °C in the Callovian (range: 27.6 to 32.1 °C), and 28.2 °C in the Kimmeridgian (range: 24.7 to 29.6 °C). Temperatures fluctuate slightly through time, but do not show prominent long-term trends (Fig. 3B; Alberti et al., 2017).

Thirty-five brachiopod and bivalve shells from Israel (palaeolatitude 2-3 °N) were analysed with their $\delta^{18}\text{O}_{\text{shell}}$ values translating into water temperatures around 34.0 °C in the Bajocian (range: 33.9 to 34.1 °C), 30.9 °C in the Callovian (range: 27.0 to 35.5 °C), and 31.4 °C in the Early Cretaceous (range: 29.2 to 33.1 °C). While the average temperatures seem to show a cooling from the Bajocian towards the Callovian (Fig. 3C), it should be noted that the Bajocian is represented only by two shells, which hinders the reliable recognition of long-term trends.

Stable isotope analyses of 22 bivalve shells from Jordan (palaeolatitude 1-2 °N) point to water temperatures around 27.5 °C in the Bajocian (range: 26.3 to 28.7 °C), 29.1 °C in the Bathonian (range: 25.8 to 35.2 °C), and 33.6 °C in the Callovian (range: 33.4 to 34.0 °C). Despite the limited number of samples, the results seem to indicate a warming in the study area starting in the latest Bathonian (Fig. 3D). This is supported by the measured Mg/Ca-ratios, which show a statistically significant, negative correlation to the $\delta^{18}\text{O}_{\text{shell}}$ values (Fig. S2B; Table S4). Mg/Ca-ratios of bivalve shells have been studied for their potential as temperature proxies, but their relationship to water temperatures seem to vary strongly between different species (e.g., Surge and Lohmann, 2008; Mouchi et al., 2013; Bougeois et al., 2016; Tynan et al., 2017). Mouchi et al. (2013) developed the following equation for the oyster *Crassostrea gigas* to calculate water temperatures based on Mg/Ca-ratios:

$$(S1) \quad T (\text{°C}) = 3.77 * \text{Mg/Ca} + 1.88$$

However, if this equation is used on the present dataset of *Africogryphaea costellata* and *Gryphaelimus jabbokensis*, the reconstructed water temperatures are mostly too low (range: 12.7 to 30.6 °C; Fig. S6). The correlation between Mg/Ca-ratios and $\delta^{18}\text{O}_{\text{shell}}$ values in the current dataset can also be used to establish a relationship with water temperatures reconstructed from the oxygen isotopes (Fig. S6):

$$(S2) \quad T (\text{°C}) = 1.0973 * \text{Mg/Ca} + 23.232$$

This relationship differs slightly, if equations are formed separately for both species, but the sample number is too low to come to a definite conclusion. Nevertheless, since both species are common in the Jurassic strata of northern Africa and western Asia, it might be useful to study the relationship between their Mg/Ca-ratios and $\delta^{18}\text{O}_{\text{shell}}$ values further in the future.

One specimen of *Africogryphaea costellata* from the Bathonian of Jordan was sampled at high resolution. Its shell recorded cyclic fluctuations around an average of 29.4 °C believed to be caused by seasonal weather changes (Fig. S7A). If explained only by water temperatures, the $\delta^{18}\text{O}_{\text{shell}}$ values translate into temperature between 26.5 and 31.4 °C. A seasonality of around 5 °C is slightly higher than that reconstructed for the Bathonian of Egypt by Alberti et al. (2017). Strong seasonal rainfall or upwelling patterns would have led to stronger changes in the $\delta^{18}\text{O}_{\text{shell}}$ values, but it cannot be excluded that minor seasonal changes in local $\delta^{18}\text{O}_{\text{sea}}$ values might have contributed to the signal (compare Moore et al., 1992).

Additional literature

Length-restrictions do not allow the citation of all studies connected with the conducted research within the article itself, therefore some additional literature will be mentioned in the following:

- Studies mentioning the Jurassic world being warmer than today with weak latitudinal temperature gradients, no polar glaciations, and/or a lack of major perturbations include Moore et al. (1992), Valdes and Sellwood (1992), Hallam (2001), and Sellwood and Valdes (2006). In contrast, some articles mentioning phases of polar glaciations in the Jurassic include Price (1999) and Dromart et al. (2003). In particular, the glaciation at the Callovian-Oxfordian boundary proposed by Dromart et al. (2003) has been debated in literature (e.g., Wierzbowski et al., 2009).

- There is a substantial amount of literature on stable isotope research connected to temperature reconstructions including for the Jurassic world. These studies concentrate on methodology, regional temperature developments, and/or global temperature changes. Some of these, which were considered, but ultimately not included in the discussion are Urey et al. (1951), Epstein et al. (1951), Podlaha et al. (1998), Wierzbowski (2002, 2004), Gröcke et al. (2003), Fürsich et al. (2005), Bragaud et al. (2008), Wierzbowski et al. (2009), Mutterlose et al. (2010), Alberti et al. (2012b, 2013, 2019), Martinez and Dera (2015), Ullmann et al. (2016), and Danise et al. (2020).

- Other methods applied for temperature reconstructions in the Jurassic include clumped isotope

analyses and the TEX_{86} palaeothermometer. Commonly, TEX_{86} -data result in temperatures warmer than those from the „classic“ stable oxygen isotopes. This has been interpreted in different ways, either supporting or discounting the applicability of the TEX_{86} proxy. In general, TEX_{86} -data might indicate warmer surface water temperatures compared to stable oxygen isotopes of bottom-dwelling or deep-swimming organisms, but the exact calibration method for TEX_{86} is also debated and might influence absolute temperature reconstructions (compare Jenkyns et al., 2012; Vickers et al., 2019; Price et al., 2020). Clumped isotope analyses are still very few for the Jurassic world and connected with a number of uncertainties. Results of Wierzbowski et al. (2018) for western Russia and Vickers et al. (2020) for Scotland point to fluctuating $\delta^{18}\text{O}_{\text{sea}}$ values due to the complex European palaeogeography (including changes in ocean currents and salinity fluctuations). Their reconstructed $\delta^{18}\text{O}_{\text{sea}}$ values span a wide spectrum, but do not necessarily correspond to open-ocean settings. Similarly, Vickers et al. (2019) used clumped isotope analyses on samples from the Falkland Plateau, but resulting $\delta^{18}\text{O}_{\text{sea}}$ values point to a restricted situation for their study area. Finally, Price et al. (2020) used clumped isotope analyses to reconstruct latitudinal gradients in the Early Cretaceous. While they reconstructed a latitudinal $\delta^{18}\text{O}_{\text{sea}}$ gradient almost parallel to the modern-day one, absolute $\delta^{18}\text{O}_{\text{sea}}$ values are much higher and therefore almost unrealistic as they would require enormous polar ice shields. The reasons for these high $\delta^{18}\text{O}_{\text{sea}}$ values reconstructed via clumped isotope analyses are still debated, but might be caused by the used calibration method and require additional research.

- The discussion on the ecological impact of the proposed elevated water temperatures along the equator during the Jurassic had to be kept very short, but literature focusing on this aspect is also relatively sparse yet. Some studies showing the importance of water temperatures on marine diversities include Roy et al. (1998), Figueira and Booth (2010), and Yamano et al. (2011). Additional articles including data on Jurassic faunas potentially pointing to generally elevated temperatures during the Jurassic or comparatively low diversities at low latitudes include Tintant (1987), Legarreta (1991), Fischer et al. (2001), and Leinfelder et al. (2005).

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