Ultra-shallow-marine anoxia in an Early Triassic shallow-marine clastic ramp (Spitsbergen) and the suppression of benthic radiation

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Abstract - Lower Triassic marine strata in Spitsbergen accumulated on a mid-to-high latitude ramp in which high-energy foreshore and shoreface facies passed offshore into sheet sandstones of probable hyperpycnite origin. More distal facies include siltstones, shales and dolomitic limestones. Carbon isotope chemostratigraphy comparison allows improved age dating of the Boreal sections and shows a significant hiatus in the upper Spathian. Two major deepening events, in earliest Griesbachian and late Smithian time, are separated by shallowing-upwards trends that culminated in the Dienerian and Spathian substages. The redox record, revealed by changes in bioturbation, palaeoecology, pyrite framboid content and trace metal concentrations, shows anoxic phases alternating with intervals of better ventilation. Only Dienerian-early Smithian time witnessed persistent oxygenation that was sufficient to support a diverse benthic community. The most intensely anoxic, usually euxinic, conditions are best developed in offshore settings, but at times euxinia also developed in upper offshore settings where it is even recorded in hyperpycnite and storm-origin sandstone beds: an extraordinary facet of Spitsbergen's record. The euxinic phases do not track relative water depth changes. For example, the continuous shallowing upwards from the Griesbachian to lower Dienerian was witness to several euxinic phases separated by intervals of more oxic, bioturbated sediments. It is likely that the euxinia was controlled by climatic oscillations rather than intra-basinal factors. It remains to be seen if all the anoxic phases found in Spitsbergen are seen elsewhere, although the wide spread of anoxic facies in the Smithian/Spathian boundary interval is clearly a global event.

Keywords: anoxia, Triassic, shallow marine, Spitsbergen, extinction.

1. Introduction

The development and intensification of marine anoxia during Permian–Triassic time is a well-known phenomenon widely regarded as a direct cause of the contemporary mass extinction (e.g. Wignall & Hallam, 1992; Isozaki, 1994; Wignall & Twitchett, 2002; Algeo *et al.* 2008; Bond & Wignall, 2010; Song *et al.* 2014). The subsequent Early Triassic interval saw widespread anoxia persist in shelf seas and oceanic waters (e.g. Isozaki, 1997; Wignall & Twitchett, 2002; Wignall *et al.* 2010). The consequent decrease in marine habitat area is thought to have been largely responsible for failure of benthic communities to recover and radiate in all but nearest-shore settings where a narrow belt of oxygenated conditions persisted (Wignall, Morante & Newton, 1998; Beatty, Zonneveld & Hender-

son, 2008; Knaust, 2010; Zonneveld, Gingras & Beatty, 2010).

The temporal development and intensity of Early Triassic marine anoxia varied considerably with the Smithian/Spathian (S/S) interval being especially noteworthy as an intense phase of oxygen deprivation (Galfetti et al. 2007; Wignall et al. 2010; Meyer et al. 2011; Song et al. 2012; Grasby et al. 2013; Sun et al. 2015). However, other than studies in South China (e.g. Song et al. 2012; Tian et al. 2014) and the Sverdrup Basin (Grasby et al. 2013), the spatial development of marine anoxia is unconstrained and the context of the events, such as the S/S episode, within the prolonged history of the Early Triassic 'superanoxic event' (sensu Isozaki, 1997), is poorly known. Here we aim to address this issue by establishing a depositional model for the Lower Triassic of Spitsbergen and then, using a combined ichnological/sedimentological/geochemical analysis, assess the temporal and spatial redox fluctuations in the region.

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Figure 1. Map of Spitsbergen within the Svalbard archipelago, showing the locations of studied sections.

2. Regional geology

The Lower Triassic succession in Spitsbergen forms a thick wedge of sediment peaking at \sim 500 m thickness in the westernmost outcrops at the entrance to Isfjorden and thinning to roughly half this thickness in more southerly and easterly outcrops on the island (Mørk, Knarud & Worsley, 1982). The thickest sections also display the most proximal facies and these are divided into the Vardebukta and Tvillingdodden formations that are roughly of Griesbachian–Dienerian and Smithian–Spathian ages, respectively. Shales and siltstone dominate both formations but a major sandbody is developed in the upper part of the Vardebukta Formation in western outcrops (Nakrem & Mørk, 1991) that records shoreface and foreshore deposition (Wignall, Morante & Newton, 1998). The more eastern outcrops lack major sandstone beds and are instead dominated by shale and siltstone strata of the Vikinghøgda Formation (Mørk *et al.* 1999).

The Lower Triassic successions have been examined in detail at four locations:

Festningen: This well-known coastal cliff section, at the western entrance to Isfjorden, provides a continuous exposure of the thickest-known development of Lower Triassic marine stratigraphy (Fig. 1). Only minor faulting in the lowest Tvillingdodden Formation and difficult-to-access sea cliffs in the uppermost part of the formation hindered sampling.

Forkastningsdalen: This valley section lies at the western end of Nathorst Land in southern Spitsbergen (Fig. 1). The Upper Permian Kapp Starostin Formation is well exposed but the basal (presumably shale-dominated) part of the Vardebukta Forma-

tion is obscured. The outcrop resumes in streamside sections c. 40 m stratigraphically above the highest Kapp Starostin outcrop, and the ensuing 95 m thick, sandstone-dominated succession is considered to belong to the upper Vardebukta Formation.

Tschermakfjellet: The basal metres of the Vikinghøgda Formation are exposed in a gully at Tschermakfjellet (Wignall, Morante & Newton, 1998; Dustira et al. 2013) but the overlying ~ 20 m is unexposed. The outcrop resumes a short distance to the south in coastal cliffs and reveals 56 m of section dominated by shales and siltstone with minor sandstone beds.

Vindodden: This steep hillside outcrop in eastern Sassenfjorden lies below Botneheia Mountain. This is the type locality for the Middle Triassic Botneheia Formation and the phosphatic black shales of this unit are well seen in the cliffs above the softer-weathering shales of the Vikinghøgda Formation. Only the topmost 110 m of the Vikinghøgda Formation are seen and it can be correlated with the eponymous type location of this unit, which lies ~ 15 km to the east of Vindodden. Comparison of sections between Vindodden and Vikinghøgda suggests a thinner development at the former locality. A distinctive, yellow-weathering, laminated dolostone with thin-shelled bivalves forms a distinct hillside ledge at both locations. At Vindodden this bed is 52 m below the Vikinghøgda/Botneheia boundary whereas it is recorded ~ 80 m below this contact at Vikinghøgda (Mørk et al. 1999).

3. Analytical protocol

The four study sections were logged in detail and facies types were identified with care given to identifying body and trace fossils and semi-quantifying the intensity of bioturbation in all beds using the six-point ichnofabric index (II) scheme of Droser & Bottjer (1986). Facies were determined in the field and further characterized by thin-section analysis. In addition polished blocks from a range of facies were examined using backscatter scanning electron microscopy to determine their pyrite petrography and (where present) the size range of pyrite framboids (cf. Wignall & Newton, 1998; Bond & Wignall, 2010).

Biostratigraphic control is available for the Vikinghøgda Formation at its type location and this allows the level of the S/S boundary to be estimated at the nearby Vindodden section. Samples were collected at this location and processed for conodonts in an attempt to further improve the biostratigraphic resolution. We employed three methods to disaggregate the samples. (1) samples were first treated with 10% acetic acid (buffered with tri-calcium phosphate) to remove carbonate contents; (2) samples were treated with 5 % H_2O_2 3– 5 times, and reaction between organic-rich materials and H_2O_2 further disintegrated the samples; (3) finally, samples were treated with the industrial surfactant Rewoquat for two weeks to disaggregate phyllosilicates (see Jarochowska et al. 2013 for details). The residues of these processes were wet sieved, washed with distilled water and dried at 55 °C. Sodium polytungstate heavy solution (2.81 g cm⁻³) was used for density separations. Conodonts were picked from the heavy fractions.

Early Triassic correlation can also be achieved using the δ^{13} C record, which shows substantial oscillations in this interval (Payne et al. 2004). Thus, detailed sampling was undertaken at Festningen and Vindodden and $\delta^{13}C_{org}$ values were generated. Stable isotope measurements were conducted at the Isotope Science Laboratory, University of Calgary and at GeoZentrum Nordbayern, University of Erlangen-Nuremberg. At Calgary, $\delta^{13}C$ was measured with a Finnigan DELTAplusXL mass spectrometer interfaced with a Costech 4010 elemental analyser. At Erlangen, δ^{13} C was determined with an elemental analyser (CE 1110) connected online to a ThermoFisher Delta V Plus mass spectrometer. All carbon isotope values are reported in the conventional δ -notation in per mil relative to the VPDB (Vienna Pee Dee Belemnite). Reproducibility of replicate standard analyses was ± 0.07 ‰ $(1\sigma, \text{Erlangen}) \text{ or } \pm 0.2 \% (1\sigma, \text{Calgary}).$

A total of 180 shale beds of the Festningen section were sampled (by SEG and BB) for trace metal assay. Determination of molybdenum, uranium and vanadium concentrations was performed at the Geological Survey of Canada. Elemental determinations were conducted on powdered samples digested in a 2:2:1:1 acid solution of H₂O–HF–HClO₄–HNO₃, and subsequently analysed using a PerkinElmer mass spectrometer, with $\pm 2\%$ analytical error.

4. Results

4.a. Chemo- and biostratigraphy

The $\delta^{13}C_{org}$ trends obtained at Festningen and Vindodden (Fig. 2) closely resemble the $\delta^{13}C_{carb}$ values from lower latitude Tethyan carbonate sections (Payne *et al.* 2004; Horacek *et al.* 2007) as well as the $\delta^{13}C_{org}$ record from the Smithian stratotype in the Sverdrup Basin (Grasby *et al.* 2013). This allows improved precision in age assignment for the Spitsbergen sections (Fig. 2).

At Festningen, the sharp negative excursion close to the base of the Vardebukta Formation is a marker for the Permian/Triassic boundary and this is followed by a rapid rise to heavier values (of -30 %) before a slowdown in the trend. The heaviest values are eventually attained around 200 m higher, in a sandstone unit. After this the $\delta^{13}C_{org}$ values begin a renewed fall that reaches a new low point 100 m higher in a shale-dominated section just above the Vardebukta/Tvillingdodden formational boundary (Fig. 2).

Comparison with Tethyan records suggests that, following the Permian/Triassic boundary low point, the slowdown in the rate of rise occurs within the Griesbachian Substage (Payne *et al.* 2004). A subsequent positive δ^{13} C peak occurs at the end of the Dienerian Substage in the Spitsbergen, Sverdrup Basin and China records (Grasby *et al.* 2013). The subsequent negative



Figure 2. Correlation of the Festningen and Vindodden sections based on their $\delta^{13}C_{org}$ trends. Stage boundaries are derived by comparing with the similar $\delta^{13}C_{carb}$ record from conodont-dated sections in South China (Payne *et al.* 2004; Horacek *et al.* 2007) and the Smith Creek $\delta^{13}C_{org}$ record of the Sverdrup Basin (Grasby *et al.* 2013). Aeg. – Aegean Substage; Bot. – Botneheia Formation.

low point is at the end of the Smithian Substage, and the remainder of the section (and nearly the entire Tvilling-dodden Formation) therefore belongs to the Spathian Substage (Fig. 2).

These age assignments differ slightly from those suggested previously by Mørk, Knarud & Worsley (1982) who placed the base of the Tvillingdodden Formation at the base of the Smithian, rather than in the mid Smithian as we suggest here (Fig. 2). The age of the eastern sections is also constrained by fossils, especially ammonoids, and is in much closer accord with the carbon isotope stratigraphy described above. Thus, the *Anawasatchites tardus* Zone straddles the S/S boundary (Mørk et al. 1999). No conodonts have been found from our Festningen samples but two thirds of samples from Vindodden yielded conodonts, of which neogondolellids were the most common and indicate a Spathian age. *Neogondolella* sp A. (Orchard, 2008) occurs in the upper part of the section, indicating a middle Spathian age (*Subrobustus* ammonoid zone). *N. regalis* was found 104 m above the base of the Vindodden section and is a long-lived form known to range from middle Spathian to early Middle Triassic time. Smithian conodonts from the lower part of the section are rare. The conodonts are similar to those described by Dagis (1984) from Siberia and contrast starkly with coeval, low-latitude faunas, which are dominated by *Neospathodus* species (e.g. Yan et al. 2013).

There is no conodont evidence for a latest Spathian age at Vindodden, and comparison of $\delta^{13}C_{org}$ trends suggest that this interval may be missing beneath the base of the Botneheia Formation (Fig. 2). Thus, the $\delta^{13}C_{org}$ values in the Spathian strata at Festningen show an initial positive trend followed by a gradual $\sim 2\%$ negative shift around 370 m followed by stable values for 100 m before a final minor, positive trend in the uppermost 30 m of the Tvillingdodden Formation. At Vindodden only the initial positive shift and the gradual negative shift (but not the prolonged stable phase) are seen above the S/S boundary (Fig. 2). This suggests that the upper part of the Spathian Substage has been removed by erosion (or the upper Spathian is highly condensed) at Vindodden. Further evidence for erosive removal comes from the basal bed of the overlying Botneheia Formation, which is an erosive-based sandstone with rounded phosphatic clasts. It is possible that a late Spathian hiatus occurs at Festningen too, albeit of briefer duration than that seen at Vindodden. The Sverdrup Basin and Tethyan records show a marked positive trend in the latest Spathian (of 4 ‰ amplitude in both organic and carbonate records); this trend is only weakly manifest at Festningen suggesting it has been truncated (Fig. 2).

4.b. Facies

The two westernmost sections (Forkastningsdalen and Festningen) show the presence of a major sandbody in the upper Vardebukta Formation whereas, further to the east, only thin sandstone beds (<50 cm thickness) are present at Tschermakfjellet, and at Vindodden they are very rare and <20 cm thick. This progressive eastward loss of sandstone supports a proximal-to-distal, west-to-east transition found by previous workers (Mørk, Knarud & Worsley, 1982; Nakrem & Mørk, 1991; Wig-nall, Morante & Newton, 1998). Nine facies types are present in the sections:

(1) Swash cross-stratified sandstone: Medium sandstone with planar laminations that show very lowangle truncation surfaces typical of swash crossstratification. Bioturbation intensity is low (II 2) and consists of vertical burrows (*Skolithos, Arenicolites* and *Diplocraterion*), which can penetrate downwards for ~50 cm. Examples of this facies type are only seen at two levels: near the top of the Forkastningsdalen section and, around the same stratigraphic level, ~200 m above the base of the Festningen section (cf. Wignall, Morante & Newton, 1998; Fig. 3). Desiccation cracks have been reported from *Skolithos*-bearing sandstone from around this level at Festningen (Mørk, Embry & Weitschat, 1989). This facies type is interpreted to record foreshore conditions and is the shallowest-water facies seen in this study.

(2) Sandy bioclastic grainstone: Limestone composed of thick-shelled bivalves and bryozoans arranged in tabular cross-sets ~ 1 m in height. Cross-beds record multimodal current flow directions. Sand-grade abraded bone material and phosphatic pellets are common. Where identifiable, the bivalves mostly belong to *Promyalina*, which have been recrystallized, and sometimes show slender borings (Fig. 4a). Prisms of calcite, present amongst the bioclastic debris, are probably from disintegrated valves of prismatic-shelled bivalves. Other fossils include echinoderms (probably ophiuroids) and bryozoans (*Paralioclema*). This facies is present in several beds at Forkastningsdalen (where they can be up to 2 m thick) and at one level at Festningen.

The sedimentology of this facies indicates persistent high-energy conditions (as testified by the abraded phosphatic debris) in a shoreface setting with dunescale bedforms recording variable flow directions.

(3) Cross-bedded sandstone: Medium-grained sandstone displaying trough cross-sets varying from 0.2– 0.6 m in height seen at Festningen and Forkastningsdalen. As with facies 2, palaeocurrents vary, although stacks of cross-sets with consistent dip are common (Wignall, Morante & Newton, 1998). Glauconite and phosphatic sand grains are present and the bottom sets can contain a lag of prismatic-shelled bivalve and brachiopod fragments together with small phosphatic concretions.

As with facies 2, cross-bedded sandstones are considered to be the product of fairweather processes on a shoreface, although the smaller dune height suggests a deeper setting.

(4) Hummocky cross-stratified (HCS) sandstone: Fine-grained sandstone showing the characteristic HCS-style thickening of laminae above erosive low points to produce hummocks. Generally found in thin sandstone beds (\sim 10 cm thick) the hummocks range from <20 cm in width to a few centimetres (Fig. 4b). Fine sand-grade pyrite grains are a common component. This very rare facies type is seen in a few beds at Festningen where it is interbedded with facies 7.

HCS bedforms are generally considered the product of storm-wave conditions (Dumas & Arnott, 2006). If this is the case with the Festningen examples then storms were rare in the Early Triassic of Spitsbergen.

(5) Sheet sandstone: This common facies type consists of thin sheets of fine- and medium-grained sandstone separated by siltstone beds (Fig. 5a). Individual beds range from 2–80 cm thick and can be stacked to produce composite beds up to 2 m thick that show little or no erosion at the bed boundaries. Internally, they commonly show fluctuating grain size with mud laminae intercalated with silt and fine sand laminae. Occasionally flute marks and small flame structures occur on the base of beds. Most beds are either planar laminated, current-ripple laminated or wave rippled (Fig. 4c). The



Figure 3. Correlation of Lower Triassic successions in Spitsbergen, thicknesses in metres. Stage-level age assignments are based on $\delta^{13}C_{org}$ chemostratigraphy supported by fossil evidence from conodonts and ammonoids. The grey-and-white 'barcodes' adjacent to the lithological columns distinguish between bioturbated and unbioturbated/laminated phases. These have been labelled (I–XI) at Festningen for ease of description in the text. Anoxic phases occur in the other sections but it is uncertain if these levels can be correlated. Aeg. – Aegean Substage; Bot. – Botneheia Formation.



Figure 4. (Colour online) Photomicrographs of (a) facies 2 at Forkastningsdalen showing a boring in a thick-shelled bivalve (labelled x), in which the shell is replaced with a blocky mosaic of calcite crystals, and bryozoans (*Paralioclema*) (labelled y). Field of view is 20 mm wide; (b) facies 4 at Festningen, upper Vardebukta Formation showing small-scale hummocky cross-stratification. Field of view is 32 mm wide; (c) facies 5, showing, in the lower half of the section, initial ripple cross-lamination, overlain by planar laminated sandstone and a mud lamina. The upper half of the section is planar laminated sandstone. Basal Tvillingdodden Formation, anoxic phase VII, Festningen. Field of view is 35 mm wide. (d) Graded silt laminae (facies 7) showing concentration of pyrite (opaque grains) in the lower part of the field of view. Anoxic phase IV, Vardebukta Formation, Festningen. Field of view is 1.3 mm wide.



Figure 5. (Colour online) Field photographs. (a) Sheet sandstone beds ~ 10 cm thick, interbedded with shale and siltstone from a level 265 m above the base of the Vardebukta Formation at Festningen. (b) Wrinkle structures, showing wrinkles present on numerous bedding planes. Forkastningsdalen. Occasional, short vertical burrows cut through the laminae (e.g. left of pencil tip). Pencil for scale is 12 cm long. (c) Laminated shale and dolomite found around the Smithian/Spathian boundary at Vindodden. Total height of field of view is ~ 2 m. (d) Bioturbated silty mudstone (facies 8) from the Smithian at Festningen (between anoxic phases V and VI). Hammer for scale is 28 cm long.

thicker beds often appear structureless but many are current rippled on careful inspection and the thickest individual beds commonly show climbing ripple lamination recording that flow was consistently offshore (eastwards). In contrast the wave-rippled beds rarely exceed 10 cm in thickness. Often beds show alternations of planar laminated and current-rippled parts and thicker beds may show a vertical succession from planar lamination to current-ripple lamination to waverippled top surfaces.

Quartz grains dominate the mineralogy but pyrite and glauconite grains can also be common (Fig. 4c). Fossils are locally common and dominated by examples of *Promyalina* and *Claraia*. Bioturbation varies in intensity (discussed in Section 4.c below) and typically consists of burrows that either penetrate the top surface (e.g. *Monocraterion*) or cut the entire bed (e.g. chevron escape traces).

Sheet sandstone records rapid deposition as indicated by the loading, escape traces and climbing ripple sets. Flows were only occasionally strong enough to erode the substrate prior to deposition (as shown by the presence of rare flutes) but the general lack of bed amalgamation indicates a deposition-dominated regime. In some regards the sheet sandstone facies resemble the product of decelerating turbidity currents but the frequent wave rippling does not fit this origin. Also, unlike typical turbidite systems there is no evidence for any organized stacking patterns typical of turbidite lobes (e.g. Macdonald *et al.* 2011) and neither are submarine channels apparent.

A storm deposition origin is possible for this facies, but typical storm bed attributes such as gutter fills and multimodal tool marks are absent whilst HCS (facies 4) is very rare. Arnott (1993) has described quasi-planar-laminated sheet sandstone beds often with current-rippled tops from lower shoreface/inner shelf environments. He attributed these beds to storms in which an initial high-velocity, oscillatory upper flow regime produces near-flat laminae and is followed by waning unidirectional flow recorded by ripples. This rather atypical storm facies fits some of the attributes of the sheet sandstone facies of Spitsbergen, although there is a much greater abundance of wave rippling than in Arnott's (1993) example.

River-fed hyperpycnites show many of the attributes of this sheet sandstone facies. These include sheet geometry and common planar and current ripples, although they lack an initial coarsening-upwards component that records surging discharge (Mulder *et al.* 2003; Plink-Björklund & Steel, 2004). However, they do show common clay laminae interbedded with fine sandstones suggesting fluctuating discharge during bed deposition (Fig. 4c).



Figure 6. (Colour online) Photomicrograph of a bryozoan-rich bed, Vardebukta Formation, Forkastningsdalen showing bryozoans (b), spirorbids (s), ophiuroid elements (o) and bivalves (c) in a siltstone matrix. Field of view is 1.5 mm wide.

(6) Wrinkle-laminated sandstone: This facies is only seen at Forkastningsdalen in laminated beds of sandstone 20–30 cm thick interbedded with facies 5 (Fig. 5b). Such structures are common in both the Precambrian and the Lower Triassic and show a range of linearity (e.g. Pruss, Fraiser & Bottjer, 2004). The Spitsbergen examples consist of flat-topped parallel ridges and grooves of millimetre heights that extend through many laminae. Bioturbation by *Planolites* and short, vertical burrows is common but not intense (II 2).

Wrinkle structures are thought to form from microbial binding of sediment in shallow-marine settings (Pruss, Fraiser & Bottjer, 2004).

(7) Siltstone: Beds of laminated siltstone are found in all locations and range from 1–20 cm thickness. The laminae are composed of weakly graded layers of quartz silt that often show concentrations of siltgrade pyrite grains at their base (Fig. 4d). Bioturbation intensity is highly variable and can reach II 6 (Fig. 5d). Only at Forkastningsdalen does the fossil content become significant, consisting of thin-valved, prismatic-shelled bivalves. At this location there is also a unique bed of highly bioturbated (II 6) siltstone that contains an abundant fauna dominated by bryozoans (*Paralioclema*), echinoderm grains (mostly ophiuroid material), *Promyalina* and spirorbids (Fig. 6).

The graded laminae of the siltstones suggest that this facies it is part of a continuum of event-bed-style deposition that includes the sheet sandstone facies. Benthic oxygenation clearly varied greatly from the well-oxygenated conditions of the thoroughly bioturbated bryozoan bed to the more common, unbioturbated beds of laminated, pyritic siltstone.

(8) Shale: Shale beds typically display fine lamination, consisting of alternations of silt-rich and clayrich laminae that are sometimes partially or completely destroyed by bioturbation. Carbonate concretions are locally common. The thickest shale unit at Festningen occurs between 275 and 310 m above the base of the Vardebukta Formation and it straddles the S/S boundary, whilst at Vindodden the majority of the section is composed of shale.

The shale beds record low-energy conditions in which the silt-rich laminae are probably deposited from weak traction currents. The variation from undisturbed lamination to intense bioturbation suggests oxygen levels varied considerably in this basinal facies.

(9) Laminated dolomite: Rare cementstone horizons, composed of silt-grade dolomite rhombs, occur interbedded amongst the shales at Vindodden. The thickest example is found 56 m above the base of the section where a 2 m thick bed of yellow, laminated dolostone with common '*Posidonia*' bivalves is found (Fig. 5c). Thin-sections show organic-rich laminae and abundant pyrite grains. Small microspheres are also present, which are possibly recrystallized radiolarians.

The laminated dolomite is an anoxic-dysoxic, hemipelagic facies possibly with a significant 'pelagic rain' of radiolarians, although substantial dolomitization has obliterated details of the fabric.

4.c. Bioturbation intensity

The intensity of bioturbation varies substantially in all sections (Figs 3, 5d, 7). Diversity of ichnogenera is modest and dominated by *Thalassinoides*, *Planolites* and *Diplocraterion* (Wignall, Morante & Newton, 1998). Burrow depths rarely exceed more than a few centimetres except in facies 1 where it can be more than 50 cm.

Fully bioturbated levels (II 6) occur at a level in the Dienerian of Festningen and in the bryozoan bed (probably also of Dienerian age) at Forkastningsdalen (Fig. 7). Both levels are also associated with peaks of benthic diversity in their respective sections. The fauna of the bryozoan bed fauna is noted above and the Festningen bioturbated horizon contains *Promyalina*, *Unionites*, *Claraia*, microgastropods, ophiuroids and brachiopods.

Perhaps surprisingly, there is little facies control to the laminated levels; only facies 1 and 6 are consistently bioturbated, whilst facies 2 and 3 are rarely bioturbated probably because a shifting substrate of dunes was difficult to colonize by mobile infauna. The other facies types occur as both burrowed and unburrowed varieties. Often the transition between burrowed/unburrowed strata is very sharp, occurring over a few centimetres, and is developed within a consistent facies type.

4.d. Framboid petrography

Pyrite grains and framboids are common in the Lower Triassic strata of Spitsbergen and occur both in finegrained facies and the sheet sandstone. At most levels the framboids are of a small diameter and show little size variation – a distribution typical of framboid populations from modern euxinic settings (Fig. 8). Larger, more variable framboid populations occur in some of the bioturbated intervals, although they still fall within the dysoxic field when compared to framboids



Figure 7. Detailed logs of the Festningen section (from 70-120 m) and the Forkastningsdalen section (from 40-90 m) showing fluctuations in ichnofabric index used to define the anoxic phases. Peak diversity occurs in the bioturbated strata between anoxic phases III and IV where bryozoans and bivalves are common. The laminated beds are generally unfossiliferous except for levels where *Promyalina* occurs.



Figure 8. Mean versus standard deviation plot of framboid populations (>100/sample) from Lower Triassic samples of Spitsbergen. The redox-related fields are derived from size-frequency distributions of framboids from modern euxinic, anoxic and dysoxic settings (cf. Wilkin, Barnes & Brantley, 1996 and Bond & Wignall, 2010) and reveals that many of the Spitsbergen samples formed in euxinic waters. No framboids occurred in the highly bioturbated (II \geq 4) samples we examined.

produced in modern environments (Wilkin, Barnes & Brantley, 1996). Framboids only become consistently rare/absent in the shoreface and foreshore sediments (facies 1–3) and in the fully bioturbated, high-diversity Dienerian strata noted above.

4.e. Trace metals

Analysis of the redox-sensitive trace metals uranium, molybdenum and vanadium at Festningen revealed two distinct trends. Uranium and vanadium show high concentrations (around 2 ppm and 50 ppm, respectively) and high U/Al and V/Al ratios through most of the Lower Triassic (Fig. 9). Somewhat higher uranium values (\leq 5 ppm) are also encountered in the shaledominated S/S boundary interval. In contrast, the behaviour of molybdenum is fundamentally different. Following an initial peak at the P/T boundary interval, Mo is not enriched (<1 ppm) and the Mo/Al ratio remains around 0 in the Vardebukta Formation. Mo concentrations then exhibit highly variable values (<1– 10 ppm) and Mo/Al ratios above the S/S boundary (Fig. 9).

5. Discussion

5.a. Depositional model

The proximal-distal trend in the Lower Triassic successions of Spitsbergen envisages nearest-shore conditions in the sandstone-rich westernmost outcrops passing distally to the east where the shale-dominated sections are found (Mørk, Knarud & Worsley, 1982; Wignall, Morante & Newton, 1998). This transition allows a facies model to be constructed in which high-energy foreshore and shoreface facies pass offshore into the hyperpycnite sand sheets of facies 5 (Fig. 10). These sheets were presumably river fed, although there is no record of fluvial conditions in the succession. Other potential origins for facies 5 include storm deposition (e.g. beds showing HCS), but the evidence is rare. A shoreface-attached turbidite system seems unlikely owing to the lack of organization of the sheets. It is note-worthy that the most distal development of the sheet sandstone beds, seen at Tschermakfjellet, are wave rippled indicating that water depths were still shallow enough for wave influence even though this location is >50 km offshore from the contemporaneous shoreface at Festningen. Shale and siltstone beds that are mostly finely laminated dominate the most offshore setting.

The vertical succession of facies consists of two long-term cycles. Following a major transgression at the start of the Griesbachian Substage, shale facies are developed throughout the region; this is followed by a shallowing/progradation trend that led to the establishment of shoreface/foreshore sandstones in westerly outcrops during Dienerian time. Gradual deepening during the following Smithian Substage culminated in widespread shale deposition once again around the S/S boundary. The succeeding Spathian succession records shallowing, although the trend only culminates in facies 5 at Festningen; the Dienerian littoral facies (facies 1–3) are not repeated. Base-level fall may also have eroded and removed upper Spathian strata at Vindodden.

5.b. Redox trends and shallow-marine anoxia

The lack of bioturbation and benthic fossils in multiple levels of the Lower Triassic succession of Spitsbergen suggests intensely anoxic/euxinic depositional conditions. This interpretation is supported by elevated trace metal concentrations (of U and V) and the size range of pyrite framboid populations that points to a sulphidic lower water column (Figs 8, 9). However, burrowed strata are also present indicating improved ventilation, and the associated framboid populations further indicate dysoxic bottom waters at these levels. Thus, the Early Triassic redox record can be divided into phases of euxinia (labelled I to XI in Fig. 3) separated by dysoxic/oxic phases. The best-oxygenated strata (II 6, diverse benthos, no framboids) occur in the Dienerian.

Unfortunately, it is not possible to confidently correlate the redox cycles between sections owing to insufficient stratigraphic resolution (further δ^{13} C study of all sections is needed). However, it seems likely that euxinic phases III and IV at Festningen are also seen at Forkastningsdalen because they occur around the same level (Fig. 3). Phases V to VIII at Festningen are separated by thin intervals of dysoxic strata, whilst further offshore at Vindodden a similar number of euxinic/dysoxic alternations is seen. If it is assumed that the upper part of the Spathian is not present at Vindodden (see discussion in Section 4a above) then it becomes possible to correlate the anoxic phases between



Figure 9. Carbon isotope and trace metal data from Festningen. Our two $\delta^{13}C_{org}$ records are in close accord (black circles, curve generated in Calgary; open triangles, curve generated in Erlangen). The concentrations of the redox-sensitive trace metals Mo, U and V have been normalized to Al. Aeg. – Aegean Substage; Bot. – Botneheia Formation; Chx – Changxingian Stage; KSF – Kapp Starostin Formation.



Figure 10. Depositional model for the Lower Triassic shallow-marine strata of Spitsbergen showing location of facies described in the text. FWWB – fairweather wave base; SWB – storm wave base.

locations. Thus, phase XI becomes the highest phase at Vindodden (Fig. 3).

The most extraordinary facet of the Early Triassic euxinic episodes is their development in exceptionally shallow-water settings. Many of the sheet sandstones, interpreted as hyperpycnites, formed in an upper offshore setting (Fig. 10), lack burrows and benthic fossils but possess framboids characteristic of euxinic deposition. The shale interbeds amongst the sandstone beds also show trace metal enrichment suggesting anoxic conditions. Typically, examples of nearshore-restricted euxinic black shales are considered to form in sheltered bottom waters beneath a pycnocline during base-level rise (e.g. Wignall & Newton, 2001). However, the Spitsbergen shallow-water euxinic facies are lithologically much more diverse than black shale facies and they differ significantly from other transgressive, anoxic facies because the Spitsbergen anoxic phases show little relationship to base-level changes. For example, the prolonged single episode of shallowing-up that spans the Griesbachian to late Dienerian substages is associated with four euxinic phases developed in successively shallower waters.

As a final observation, it is noteworthy that the $\delta^{13}C_{org}$ curve closely matches that found in Tethyan carbonates. This is despite the fact that it is obtained from a clastic and frequently nearshore environment. Such a setting, with high terrigenous run-off, might be expected to show more 'noise' caused by the mixing of variable amounts of marine and terrestrial organic carbon. That it does not suggests a uniform source perhaps dominated solely by marine organic matter. The Early Triassic period is noteworthy for its global 'coal gap' (Retallack, Veevers & Morante, 1996), and it may have been the case that there was little terrestrial organic productivity at this time. However, in a noteworthy counterpoint, it has been suggested that woody vegetation biomass recovered after the S/S boundary (Saito *et al.* 2013).

5.c. Global Early Triassic redox

The generally poor level of ventilation in Early Triassic shallow seas of Spitsbergen may have its origin in high oxygen demand, fostered by high productivity at a time of high run-off (e.g. Algeo & Twitchett, 2010). However, the phases of anoxia are not related to proximity to terrestrial nutrient supply nor to water depth suggesting another factor is more important. The anoxic phases correspond to the sea-surface temperature (SST) record of Sun et al. (2012). It is especially noteworthy that the two Early Triassic peaks of SST, in Griesbachian and late Smithian time, coincide with intervals of prolonged and extensive euxinia in Spitsbergen (Fig. 11). In contrast, the Dienerian Substage is the coolest interval of the Early Triassic SST curve and also the best-oxygenated period in Spitsbergen, and also globally (Wignall & Twitchett, 2002). There are three causal links between high temperatures and anoxia that may be at play here: (i) dissolved oxygen levels decline with temperature; (ii) organic matter remineralization rates increase with temperature thereby increasing oxygen demand within the water column; and (iii) a general link between warmer climates and increased humidity, thereby increasing nutrient run-off to the oceans.

If global temperature is a key factor in the Spitsbergen euxinic phases then it suggests such conditions should have been widespread. The Early Triassic period is indeed known as a 'superanoxic episode' characterized by widespread ocean anoxia (Isozaki, 1997), and recent work has demonstrated detailed fluctuations within this generally poorly ventilated oceanic interval (e.g. Wignall *et al.* 2010; Grasby *et al.* 2013; Tian *et al.* 2014). Anoxia was especially widespread during Griesbachian time, the S/S boundary interval and late Spathian time – all intervals of euxinia in Spitsbergen. Further work is needed to test whether these



in equatorial Tethys

Figure 11. Correlation of Spitsbergen anoxic phases with the sea-surface temperature (SST) record of Sun *et al.* (2012). The width of the SST curve denotes the minimum and maximum SSTs based on conodont apatite oxygen isotopes. A $\sim 2^{\circ}$ C warmer record was obtained around the Smithian/Spathian boundary from surface-dwelling conodonts. There is a correspondence between the warmest phases in the Griesbachian and Smithian and the frequent development of anoxic phases.

Spitsbergen euxinic phases are global cycles or regional ones.

The concept of widespread Early Triassic anoxia has, however, been challenged. Resurrecting an original idea of Erwin (1993), Hofmann et al. (2015, p. 9) suggested that the prevalence of laminated sediment was caused by 'extinction [at the end of the Permian], rather than the environmental exclusion, of bioturbators', producing an Early Triassic world with no animals capable of generating thoroughly bioturbated surficial sediments. This is suggested to have caused intense anoxia within the sediment producing a geochemical signal that reflects this. The Spitsbergen record clearly shows this scenario is untenable because it consists of alternations of laminated and intensely bioturbated strata. Intense burrowing is also known from other regions (e.g. Beatty, Zonneveld & Henderson, 2008; Zonneveld, Gingras & Beatty, 2010), and even Hofmann himself records intense bioturbation (II 4) from the Lower Triassic of Italy (Hofmann et al. 2011). Early Triassic bioturbators were readily capable of generating thoroughly mixed sediments but 'environmental exclusion' was common and the available evidence overwhelmingly favours anoxia as a cause.

6. Conclusions

A new $\delta^{13}C_{org}$ record from Spitsbergen, together with sporadic conodont finds, allows a major improvement in age dating of the Spitsbergen succession. The Lower Triassic record is seen to consist of hyperpycnite sandstone and shale-dominated ramp facies. Shoreward these pass into shoreface facies displaying shell-rich, cross-bedded strata and foreshore facies. Relative water depth changes saw deepest-water conditions in earliest Griesbachian time and around the S/S boundary with shallowest waters in late Dienerian time. Regression in later Spathian time may be responsible for the truncation of Spathian strata in the region.

Much of the Lower Triassic strata accumulated in oxygen-poor and frequently euxinic waters, including hyperpycnite sand sheets, and points to intervals of exceptionally shallow-water development of such conditions. However, the anoxic phases, of which at least a dozen are developed, do not correlate closely with water depth changes. Instead, there may have been a climatic control on the Early Triassic anoxic phases with the most intense anoxia developed at times of greatest warmth. The best-oxygenated Early Triassic interval occurred during the coolest phase, in earliest Dienerian time, and was witness to the development of a highly diverse benthic community found in thoroughly bioturbated sediments.

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References

- ALGEO, T. J., SHEN, Y., ZHANG, T., LYONS, T., BATES, S., ROWE, H. & NGUYEN, T. K. T. 2008. Association of ³⁴S-depleted pyrite layers with negative carbonate δ^{13} C excursions at the Permian–Triassic boundary: evidence for upwelling of sulfidic deep-ocean water masses. *Geochemistry, Geophysics, Geosystems* **9**, Q04025.
- ALGEO, T. J. & TWITCHETT, R. J. 2010. Anomalous Early Triassic sediment fluxes due to elevated weathering rates and their biological consequences. *Geology* 38, 1023–6.
- ARNOTT, R. W. C. 1993. Quasi-planar-laminated sandstone beds of the Lower Cretaceous Bootlegger Member, north-central Montana: evidence of combined flow sedimentation. *Journal of Sedimentary Petrology* 63, 488– 94.
- BEATTY, T. W., ZONNEVELD, J.-P. & HENDERSON, C. M. 2008. Anomalously diverse Early Triassic ichnofossil assemblages in northwest Pangea: a case for a shallowmarine habitable zone. *Geology* 36, 771–4.
- BOND, D. P. G. & WIGNALL, P. B. 2010. Pyrite framboid study of marine Permo-Triassic boundary sections: a complex anoxic event and its relationship to contemporaneous mass extinction. *Geological Society of America Bulletin* 122, 1265–79.
- DAGIS, A. A. 1984. Rannetriasovye konodonty severa Srednej Sibiri. (Early Triassic conodonts of northern Middle Siberia.) *Trudy Akademija SSSR, Sibirskoe otdelenie Instituta Geologii i Geofiziki* 554, 3–69.
- DROSER, M. L. & BOTTJER, D. J. 1986. A semiquantitative field classification of ichnofabric. *Journal of Sediment*ary Petrology 56, 558–9.
- DUMAS, S. & ARNOTT, R. W. C. 2006. Origin of hummocky and swaley cross-stratification – the controlling influence of unidirectional current and aggradation rate. *Geology* 34, 1073–6.
- DUSTIRA, A. M., WIGNALL, P. B., JOACHIMSKI, M., BLOMEIER, D., HARTKOPF-FRÖDER, C. & BOND, D. P. G. 2013. Gradual onset of anoxia across the Permian– Triassic boundary in Svalbard, Norway. *Palaeogeography, Palaeoclimatology, Palaeoecology* 374, 303–13.
- ERWIN, D. H. 1993. *The Great Paleozoic Crisis*. New York: Columbia University Press, 327 pp.
- GALFETTI, T., HOCHULI, P. A., BRAYARD, A., BUCHER, H., WEISSERT, H. & VIGRAN, J. O. 2007. Smithian–Spathian boundary event: evidence for global climatic change in the wake of the end-Permian biotic crisis. *Geology* 23, 291–4.
- GRASBY, S. E., BEAUCHAMP, B., EMBRY, A. & SANEI, H. 2013. Recurrent Early Triassic ocean anoxia. *Geology* 41, 175–8.
- HOFMANN, R., BUATOIS, L. A., MACNAUGHTON, R. B. & MANGANO, M. G. 2015. Loss of the sedimentary mixed layer as a result of the end-Permian extinction. *Palaeogeography, Palaeoclimatology, Palaeoecology* **428**, 1–11.
- HOFMANN, R., GOUDEMAND, N., WASMER, M., BUCHER, H. & HAUTMANN, M. 2011. New trace fossil evidence for an early recovery signal in the aftermath of the end-Permian mass extinction. *Palaeogeography, Palaeoclimatology, Palaeoecology* **310**, 216–26.

- HORACEK, M., RICHOZ, S., BRANDNER, R., KRYSTYN, L. & SPÖTL, C. 2007. Evidence for recurrent changes in Lower Triassic oceanic circulation in Tethys: the record from marine sections in Iran. *Palaeogeography, Palaeoclimatology, Palaeoecology* 252, 355–69.
- ISOZAKI, Y. 1994. Superanoxia across the Permo-Triassic boundary: recorded in accreted deep-sea pelagic chert in Japan. *Memoir of the Canadian Society of Petroleum Geologists* 17, 805–12.
- ISOZAKI, Y. 1997. Permo-Triassic boundary superanoxia and stratified superocean: records from lost deep sea. *Science* 276, 235–8.
- JAROCHOWSKA, E., TONAROVÁ, P., MUNNECKE, A., FERROVÁ, L., SKLENÁŘ, J. & VODRÁŽKOVÁ, A. S. 2013. An acid-free method of microfossil extraction from clayrich lithologies using the surfactant Rewoquat. *Palaeontologia Electronica* 16, 7T; 16 pp.
- KNAUST, D. 2010. The end-Permian mass extinction and its aftermath on an equatorial carbonate platform: insights from ichnology. *Terra Nova* **22**, 195–202.
- MACDONALD, H. A., PEAKALL, J., WIGNALL, P. B. & BEST, J. 2011. Sedimentation in deep-sea lobe elements: implications for the origin of thickening-upwards sequences. *Journal of the Geological Society, London* 168, 319–32.
- MEYER, K. M., YU, M., JOST, A. B., KELLEY, B. M. & PAYNE, J. L. 2011. δ^{13} C evidence that high primary productivity delayed recovery from end-Permian mass extinction. *Earth and Planetary Science Letters* **302**, 378–84.
- MØRK, A., ELVEBAKK, G., FORSBERG, A. W., HOUNSLOW, M. W., NAKREM, H. A., VIGRAN, J. O. & WEITSCHAT, W. 1999. The type section of the Vikinghøgda Formation: a new Lower Triassic unit in central and eastern Spitsbergen. *Polar Research* 18, 51–82.
- MØRK, A., EMBRY, A. F. & WEITSCHAT, W. 1989. Triassic transgressive-regressive cycles in the Sverdrup Basin, Svalbard and the Barents Shelf. In *Correlation in Hydrocarbon Exploration* (ed. J. D. Collinson), pp. 113–30. Norwegian Petroleum Society/Graham & Trotman.
- MØRK, A., KNARUD, R. & WORSLEY, D. 1982. Depositional and diagenetic environments of the Triassic and Lower Jurassic succession of Svalbard. In Arctic Geology and Geophysics (eds A. F. Embry & H. R. Balkwill), pp. 371–98. Canadian Society of Petroleum Geologists Memoir 8.
- MULDER, T., SYVITSKI, J. P. M., MIGEON, S., FAUGÈRES, J.-C. & SAVOYE, B. 2003. Marine hyperpycnal flows: initiation, behaviour and related deposits. *A review. Marine* and Petroleum Geology 20, 861–82.
- NAKREM, H. A. & MØRK, A. 1991. New early Triassic Bryozoa (Trepostomata) from Spitsbergen, with some remarks on the stratigraphy of the investigated horizons. *Geological Magazine* 128, 129–40.
- ORCHARD, M. J. 2008. Lower Triassic conodonts from the Canadian Arctic, their intercalibration with ammonoidbased stages and a comparison with other North American Olenekian faunas. *Polar Research* 27, 393–412.
- PAYNE, J. L., LEHRMANN, D. J., WEI, J., ORCHARD, M. J., SCHRAG, D. P. & KNOLL, A. H. 2004. Large perturbations of the carbon cycle during recovery from the end-Permian extinction. *Science* **305**, 506–9.
- PLINK-BJÖRKLUND, P. & STEEL, R. J. 2004. Initiation of turbidity currents: outcrop evidence of hyperpycnal flow turbidites. *Sedimentary Geology* 165, 29–52.
- PRUSS, S. B., FRAISER, M. & BOTTJER, D. J. 2004. Proliferation of Early Triassic wrinkle structures: implications for environmental stress following the end-Permian mass extinction. *Geology* 32, 461–4.

- RETALLACK, G. J., VEEVERS, J. J. & MORANTE, R. 1996. Global coal gap between Permian–Triassic extinction and Middle Triassic recovery of peat-forming plants. *Geological Society of America Bulletin* 108, 195–207.
- SAITO, R. KAIHO, K., OBA, M., TAKAHASHI, S., CHEN, Z.-Q. & TONG, J.-N. 2013. A terrestrial vegetation turnover in the middle of the Early Triassic. *Global and Planetary Change* 105, 152–9.
- SONG, H.-J., WIGNALL, P. B., CHU, D.-L., TONG, J.-N., SUN, Y.-D., SONG, H.-Y., HE, W.-H. & TIAN, L. 2014. Anoxia/high temperature double whammy during the Permian-Triassic marine crisis and its aftermath. *Scientific Reports* 4, 4132. doi: 10.1038/srep04132.
- SONG, H.-J., WIGNALL, P. B., TONG, J.-N., BOND, D. P. G., SONG, H.-Y., LAI, X.-L., ZHANG, K., WANG, H.-M. & CHEN, Y.-L. 2012. Geochemical evidence from bioapatite for multiple oceanic anoxic vents during end-Permian transition with end-Permian extinction and recovery. *Earth and Planetary Science Letters* 353–354, 12–21.
- SUN, Y.-D., JOACHIMSKI, M. M., WIGNALL, P. B., YAN, C.-B., CHEN,, Y.-L., JIANG, H.-S., WANG, L.-N. & LAI,, X.-L. 2012. Lethally hot temperatures during the Early Triassic greenhouse. *Science* 388, 366–70.
- SUN, Y.-D., WIGNALL, P. B., JOACHIMSKI, M. M., BOND, D. P. G., GRASBY, S. E., SUN, S., YAN, C. B., WANG, L. N., CHEN, Y. L. & LAI, X. L. 2015. High amplitude redox changes in the late Early Triassic of South China and the Smithian–Spathian extinction. *Palaeogeography, Palaeoclimatology, Palaeoecology* 427, 62– 78.
- TIAN, L., TONG, J.-N., ALGEO, T. J., SONG, H.-J., SONG, H.-Y., CHU, D.-L., SHI, L. & BOTTJER, D. J. 2014. Reconstruction of Early Triassic ocean redox conditions based on framboidal pyrite from the Nanpanjiang Basin, South China. *Palaeogeography, Palaeoclimatology, Palaeoecology* **412**, 68–79.
- WIGNALL, P. B., BOND, D. P. G., KUWAHARA, K., KAKUWA, K., NEWTON, R. J. & POULTON, S. W. 2010. An 80 mil-

lion year oceanic redox history from Permian to Jurassic pelagic sediments of the Mino-Tamba terrane, SW Japan, and the origin of four mass extinctions. *Global and Planetary Change* **71**, 109–23.

- WIGNALL, P. B. & HALLAM, A. 1992. Anoxia as a cause of the Permian/Triassic extinction: facies evidence from northern Italy and the western United States. *Palaeogeography, Palaeoclimatology, Palaeoecology* **93**, 21–46.
- WIGNALL, P. B., MORANTE, R. & NEWTON, R. 1998. The Permo-Triassic transition in Spitsbergen: $\delta^{13}C_{org.}$ chemostratigraphy, Fe and S geochemistry, facies, fauna and trace fossils. *Geological Magazine* **133**, 47–62.
- WIGNALL, P. B. & NEWTON, R. 1998. Pyrite framboid diameter as a measure of oxygen deficiency in ancient mudrocks. *American Journal of Science* 298, 537–52.
- WIGNALL, P. B. & NEWTON, R. 2001. Black shales on a basin margin: a model based on examples from the Upper Jurassic of the Boulonnais, northern France. *Sedimentary Geology* 144, 335–56.
- WIGNALL, P. B. & TWITCHETT, R. J. 2002. Extent, duration and nature of the Permian–Triassic superanoxic event. In *Catastrophic Events and Mass Extinctions: Impacts* and Beyond (eds C. Koeberl & K. C. MacLeod), pp. 395–413. Geological Society of America Special Paper no. 356.
- WILKIN, R. T., BARNES, H. L. & BRANTLEY, S. L. 1996. The size distribution of framboidal pyrite in modern sediments: an indicator of redox conditions. *Geochimica et Cosmochimica Acta* 60, 3897–912.
- YAN, C. B., WANG, L. N., JIANG, H. S, WIGNALL, P. B., SUN, Y. D., CHEN, Y. L. & LAI, X. L. 2013. Uppermost Permian to Lower Triassic conodonts at Bianyang Section, Guizhou Province, South China. *Palaios* 28, 509–22.
- ZONNEVELD, J.-P., GINGRAS, M. K. & BEATTY, T. W. 2010. Diverse ichnofossil assemblages following the P-T mass extinction, Lower Triassic, Alberta and British Columbia: evidence for shallow marine refugia on the northwestern coast of Pangea. *Palaios* 25, 368–92.