

# The history of exploration and stratigraphy of the Early to Middle Triassic vertebrate-bearing strata of Svalbard (Sassendalen Group, Spitsbergen)

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The palaeontology of the Lower to Middle Triassic succession in Spitsbergen has been studied for more than a century and a half. Our ability to properly interpret the evolutionary and ecological implications of the faunas requires precise stratigraphic control that has only recently become available. Within such a detailed stratigraphic framework, the Spitsbergen fossil material promises to contribute to our understanding of the faunal recovery after the end-Permian mass extinction.

**Keywords:** *Triassic stratigraphy, Boreal, Spitsbergen, Svalbard, Ichthyopterygia, Sedimentology, Mass extinction, Marmierfjellet, Vikinghøgda, Deltadalen, Botneheia, Olenekian, Anisian, Smithian, Spathian*

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## Introduction

The Permian-Triassic Mass Extinction (PTME) was by far the largest extinction in Earth's history, during which a staggering 90% of all species living in marine habitats are thought to have become extinct (Raup & Sepkoski, 1982; Erwin, 1994; Benton, 1995; Wignall et al., 1998; Scheyer et al., 2014; Roopnarine & Angielczyk, 2015). The mass extinction also affected life in terrestrial habitats, where approximately 70% of all vertebrates disappeared (Elewa, 2008; Metcalfe & Isozaki, 2009; Yin & Song, 2013). The extinction has been proposed to be caused by an increase in volcanic eruptions during the Late Permian, which reached its peak during the

waning stages of the Palaeozoic (Burgess et al., 2017). A direct result of this was the formation of the Siberian Traps (Yin & Song, 2013; Burgess et al., 2017). As a result of escalating volcanic activity, vast amounts of gas was released into the atmosphere (Burgess et al., 2017), creating a heating effect that contributed to a warmer global climate (Yin & Song, 2013). Intense heating during the final phases of the Permian may very well have been one of, if not the most dominant, causes for the mass extinction (Hallam & Wignall, 1997). The recovery stages in the wake of the end Permian extinctions resulted in the evolution of new Triassic marine communities, some of which still dominate today's oceans (Erwin, 1998).

During the Early Triassic, Svalbard formed part of a large

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embayment on the northern margin of Pangaea, known as the Boreal Ocean, although it was at approximately 45° palaeolatitude, or just north of the modern Mediterranean Sea if we compare to the present-day map (Nystuen et al., 2013; Lundschiem et al., 2014). Located in the northern regions of Panthalassa, it connected Eurasia/Baltica in the east with Laurentia/Gondwana in the west (Mørk, 2015). Recent excavations in what is today South China and Nevada have revealed extensive evidence of the biotic recovery after the PTME, with certain localities even preserving several ecosystems over millions of years (see Benton et al., 2013; Motani et al., 2014; Jiang et al., 2016; Kelley et al., 2016; Brayard et al., 2017). However, much less is known about this biotic recovery in other areas. The fossiliferous Early to Middle Triassic successions in Svalbard are one of the best, yet understudied localities to document and interpret the survival and recovery of marine ecosystems in the Boreal realm (Foster et al., 2016) with a crucial placement between the Chinese and Nevada localities.

Of special interest are several clades of tetrapods that evolved secondary aquatic lifestyles in the Triassic. Notably, Ichthyosauriformes is traditionally recognised as one of the clades originating and diversifying shortly after the PTME, during the Early Triassic (Smithian; e.g., Massare & Callaway, 1994; Sander, 2000; Motani, 2005; Motani et al., 2014; Jiang et al., 2016). Our current understanding of ecosystem recovery post-PTME is limited, but the new excavations from the Early and Middle Triassic of Svalbard, described in this volume, are increasing our knowledge of taxonomic diversity, niche partitioning and cladogenesis post-PTME.

### Early work (1864–1964)

Ichthyopterygian material has been collected and described from the Triassic strata of Svalbard since the late 19th century (Nordenskiöld, 1866; Hulke, 1873; Dames, 1895; Yakowlew, 1903). These collections were undertaken during the first scientific expeditions to map the geology of Svalbard; but the stratigraphic resolution was poor. As a result, none of the specimens collected from Spitsbergen in the nineteenth century can be placed in a more modern stratigraphic context with any certainty.

In 1908, the student Bertil Högbom participated in geologist De Geer's expedition to the Isfjorden area of central Spitsbergen on behalf of Swedish palaeontologist Carl Wiman. The expedition was a success, with a large amount of material collected. As a result, in 1909, Wiman instigated an expedition from Uppsala University, Sweden, led by Högbom, during which substantial amounts of ichthyopterygian material were collected from central Spitsbergen (Wiman, 1910a). In the following decades, Wiman published a series of papers on the ichthyopterygian remains from this area

(Wiman, 1916a, 1922, 1928, 1933). Smaller expeditions to Spitsbergen were also organised to collect Triassic vertebrates in the summers of 1912, 1913, 1915 and 1916–1918, but these expeditions mostly focused on Triassic fish and stegocephalians (Wiman, 1910b, 1914a, b, 1916b, 1917, 1918; Stensiö, 1918, 1921, 1925).

Wiman (1910a) described the Triassic strata around Isfjorden with three main vertebrate-bearing horizons placed according to the biostratigraphy as defined by Noetling (1905). A horizon rich in fossil fish, 'das Fischniveau' (= the Fish niveau = the Fish horizon of Wiman (1916a)) and two ichthyopterygian bone-bearing and stratigraphically restricted levels, 'das unteres Saurierniveau' (= Lower Saurian niveau = the Lower Saurian horizon of Wiman (1916a)) and 'das oberes Saurierniveau' (= the Upper Saurian niveau = the Upper Saurian horizon of Wiman (1916a)). The Fish niveau correlates to the Posidonomya shales, and the Upper Saurian niveau was correlated to the upper part of the 'Daonellen-niveau' as defined by Mojsisovics (1886). During the following decades, many workers adopted the biostratigraphic nomenclature of Wiman (1910a, 1916a) (e.g., Stolley, 1911; Böhm, 1912; Diener, 1916; Stensiö, 1918, 1921; Spath, 1921, 1934; Frebold, 1930). Stensiö (1921) located a bone-bearing level of 'rather local nature' 33 m above the Fish niveau (he used the term 'horizon'), containing chondrichthyans, parasuchians, unrecognisable tetrapod material and ichthyopterygians, and *Omphalosaurus*. The bone-bearing level also contained ammonoids and was assumed to be an endemic fauna (Frebold, 1930).

In 1928, Wiman described *Grippia longirostris* Wiman, 1928, based on material collected by the Hamburg Spitsbergen Expedition during fieldwork on the east side of Spitsbergen in 1927. Based on information from palaeontologist E. Stensiö and based on "the iron-rich matrix surrounding the specimen", a 'Grippia niveau' was interpreted to exist somewhere between the Fish niveau and the Lower Saurian niveau (Wiman, 1928). Further expeditions were carried out in 1929 and 1930, and a concretion-rich layer interpreted as the Grippia niveau was found in Agardhbukta and on Milne Edwardsfjellet on Spitsbergen (Wiman, 1933).

### Recent work (1965–present)

The Triassic stratigraphy of Spitsbergen as defined by Wiman remained the main framework for all subsequent workers until Buchan et al. (1965) revised the lithostratigraphy of central Spitsbergen, but they also commented on other areas of Svalbard. Using the scheme of Buchan et al. (1965), Tozer & Parker (1968) subsequently updated the Triassic biostratigraphy of Svalbard. Several revisions were later made by other workers studying the lithostratigraphy of Svalbard that also combined invertebrate, microfossil and palynological

data collected from additional localities (Birkenmajer, 1977 – Sørkapp Land; Lock et al., 1978 – Barentsøya and Edgeøya; Worsley & Mørk, 1978 – Sørkapp Land; Pčelina, 1980, 1983 – entire Svalbard; Mørk et al., 1982 – Svalbard, 1990 – Bjørnøya, 1999a – entire Svalbard, 1999b – Spitsbergen, Barentsøya and Edgeøya; Weitschat & Dagys, 1989 – Svalbard; Krajewski et al., 2007 – Western Svalbard; Krajewski, 2008 – Barentsøya and Edgeøya; Nakrem et al., 2008 – Spitsbergen; Vigran et al., 2014 – Svalbard). Few attempts were made to correlate the older biostratigraphic scheme of Wiman with the modern, revised, lithostratigraphic studies until recently (Maxwell & Kear, 2013). As a result, much of the vertebrate material that had been collected from the Lower to Middle Triassic of Svalbard has now been roughly placed in the stratigraphic context defined by Wiman (1910a). However, much of this vertebrate material has not been placed with any accuracy in detailed stratigraphic logs from the different localities (e.g., Mazin, 1981a, b, 1983, Maisch & Matzke, 2002b, 2003a, b). An example of difficulties arising from a lack of stratigraphic control is the supposedly articulated ichthyosaur specimen of *Merriamosaurus hulkei* (a junior synonym to *Pessopteryx nisseri*, see discussion in Engelschiøn et al., 2018). The material was collected in the 1960s by the Muséum National d'Histoire Naturelle (MNHN) in Paris, from various localities (e.g., Heimenfjellet and Ibsenfjella, approximately 4.5 km apart). Maisch & Matzke (2002b) assigned a significant amount of material to *M. hulkei* claiming it to represent one individual due to the similar size and type of preservation. Ultimately, more than 170 fragments, or two-thirds of the total material, was reassembled and assigned to a single specimen (Maisch & Matzke, 2002b).

Early Triassic ichthyopterygians from Svalbard have received sporadic attention in the latter half of the 20th century. Collections were made by the Cambridge Spitsbergen Expeditions during geological expeditions in 1948–1992 (= the Cambridge Arctic Shelf Programme (CASP) since 1975). Much of the material housed in the CASP collection in Cambridge, and material donated by, or bought from private collectors in the Natural History Museum in London remains undescribed. The Muséum National d'Histoire Naturelle (MNHN) in Paris conducted two expeditions to Spitsbergen; the 1964 Lehman-MNHN expedition and the 1969 CNRS-MNHN expedition (Centre National de la Recherche Scientifique). The focus of these expeditions was on Palaeozoic and Triassic fish, although ichthyopterygian remains were also collected. Mazin published a series of papers on the MNHN material (Mazin, 1981a, b, 1983, 1984), later revised by Maisch & Matzke (Maisch & Matzke, 2002a, b, 2003a, b). Maisch & Blomeier (2009) described a Middle Triassic ?toretocnemid collected in Dickson Land by geologists from the Norwegian Polar Institute in 2003. In 2008, scientists from the Norwegian Petroleum Directorate and the Natural History Museum in Oslo collected a Middle Triassic mixosaurid specimen

from Edgeøya, one of the eastern islands in the Svalbard archipelago (Hurum et al., 2014). In summary, the material collected by Wiman has formed the basis for our understanding of Triassic vertebrates from Svalbard until today, although much of the material has been repeatedly redescribed, and many specimens are now considered non-diagnostic (for a review see Sander & Faber, 1998; Motani, 1999; Maxwell & Kear, 2013).

## The work by the Spitsbergen Mesozoic Research Group at the Natural History Museum in Oslo (2014–now)

During three seasons of fieldwork in 2014–16, abundant vertebrate material has been recovered from the Lower and Middle Triassic Sassendalen Group (Vikingshøgda and Botneheia formations) in the Isfjorden area of central Spitsbergen. The newly collected material discussed below is from the Lower and Middle Triassic deposits of Marmierfjellet, central Spitsbergen, Svalbard (Fig. 1). The material is housed in the Natural History Museum, University of Oslo (collection prefix PMO). All localities are denominated by their approved name by the Norwegian Polar Institute, and all necessary permits were obtained from the Governor of Svalbard for the excavations; permits no. RiS 6725, 2013/1222–2; RiS 10227, 2015/00326–2; and RiS 10539, 2015/00326–5. The following papers (Bratvold et al., 2018; Ekeheien et al., 2018; Engelschiøn et al., 2018; Hansen et al., 2018; Økland et al., 2018) in this volume will highlight some of the early findings from the vast material still being processed at the museum in Oslo.

## Geological setting

During the Triassic, a large embayment of the Panthalassa Ocean was located on the northern rim of Pangaea, known as the Boreal Ocean (Nystuen et al., 2013; Lundschieen et al., 2014). At a palaeolatitude of approximately 45°N, the Boreal Ocean formed a low-relief, open-marine shelf, surrounded by Eurasia/Baltica and the Urals to the east and southeast, and the North American portion of Gondwana/Laurentia to the west, including Greenland to the southwest and the Canadian Arctic islands to the northwest (Mørk, 2015). During the Permian, Siberia moved towards Euramerica and eventually closed the Uralian Seaway that connected the Boreal Ocean to the equatorial Tethys, leading to the creation of the Uralian Mountains (Worsley, 2008; Lundschieen et al., 2014; Blomeier, 2015). The coastline in the Early Triassic was mainly located outside of present-day Svalbard, with sediment input originating from the southwest. Later, northeast-prograding clinoforms





is the Permian, chert-rich, Kapp Starostin Formation, and it is overlain by the Botneheia Formation, comprising mostly dark organic rich, calcareous shales of marine origin (Mørk et al., 1999a, b). The Vikinghøgda Formation is divided into three members; the Deltadalen Member consists of mudstones with sandstones and siltstones, the Lusitaniadalen Member is characterised by silty mudstones with limestone concretions, and the Vendomdalen Member comprises darker silty shales (Mørk et al., 1999b). The Vikinghøgda Formation spans the Upper Permian Changhsingian Stage to the boundary between the Lower Triassic Olenekian Stage and the Middle Triassic Anisian Stage. The basal few metres of the formation are composed of a sandstone or mudstone gradually grading into a mudstone with thin siltstone or sandstone beds, interpreted as transgressive units (Mørk et al., 1999b). With a total thickness is 250 m (Mørk et al., 1999b), the Vikinghøgda Formation is well exposed at the type section in Sassendalen, including exposures on the mountains of Vikinghøgda and Sticky Keep and the river valley Deltadalen between the. The uppermost part, the Vendomdalen Member, is also well exposed in the study area on the hillside of Marmierfjellet in Flowerdalen, immediately west of Sassendalen.

Ammonoids, bivalves, brachiopods and vertebrates occur in discrete horizons throughout the Vikinghøgda Formation, along with carbonate concretions (Mørk et al., 1999b). The biostratigraphy of the Vikinghøgda Formation has been defined by ammonoids, conodonts and palynology, with the ammonoid stratigraphy of W. Weitschat providing crucial age control (Weitschat & Lehmann, 1978; Weitschat & Dagys, 1989; Dagys & Weitschat, 1993; Mørk et al., 1999b). Tozer (1994), Dagys & Sobolev (1995) and Piazza et al. (2017) continued this work and Hounslow et al. (2008a) linked the magnetostratigraphy of the Vikinghøgda Formation to the ammonoid biostratigraphy. The palynological composite assemblage zones were defined by Mørk et al. (1999b) and Vigran et al. (2014).

#### Deltadalen Member

The Deltadalen Member extends from the upper Changhsingian through most of the Induan and is well constrained by a biostratigraphy of ammonoids and conodonts, and magnetostratigraphy (Mørk et al., 1999b; Hounslow et al., 2008a; Nakrem et al., 2008).

Nakrem et al. (2008) suggested that the Permian/Triassic boundary occurs between 5 and 11 m above the base of the Deltadalen Member. This was based on the conodont *Neogonodrella carinata*, indicative of a Griesbachian age. The disappearance of bioturbation in the same interval is a good indicator for the boundary in the field (Foster et al., 2016). The basal few metres are composed of bedded, glauconitic sandstones that are very similar to those in the underlying Kapp Starostin Formation, but the layers lack diagenetic chert nodules (Mørk et al., 1999a, b; Nabbefeld et al., 2010). These sandstones are described by

Nabbefeld et al. (2010) and Foster et al. (2016) to contain a diverse trace fossil assemblage. The sandstones mark the onset of a marine transgression (Mørk et al., 1999b; Nabbefeld et al., 2010). Above the glauconitic sandstone, silty mudstones indicate a significant deepening of the basin. The mudstones were deposited under anoxic and euxinic conditions, and Nabbefeld et al. (2010) found evidence of phytoplankton blooms in this part of the Deltadalen Member. Foster et al. (2016) described the oldest silicified invertebrate assemblage known from the Triassic from two carbonate concretions at 11.9 and 12.6 m above the base of the member. After acid treatment they were found to contain 14 species of bivalves and gastropods, as well as conodonts and ammonoids. Above the mudstone a greater proportion of siltstones and fine sandstones are interpreted as a progradation or a fall of sea level (Foster et al., 2016). The bioturbation shows a slow recovery with deposition under more oxygenated conditions.

#### Lusitaniadalen Member

The Lusitaniadalen Member of the Vikinghøgda Formation is situated above the Deltadalen Member and below the Vendomdalen Member (Fig. 2). The member consists mainly of dark-grey silty mudstones to sandstones, with carbonate concretions especially in the middle and upper parts of the unit. It is 88 m thick in the type section (Mørk et al., 1999b), but the lower part is generally scree covered. Of special interest regarding vertebrate fossils is the uppermost interval in the member, loosely referred to as the 'Fish niveau' following e.g., Wiman (1918) and Stensiö (1921, 1925). The Fish niveau contains abundant calcite concretions with well preserved ammonoids, bivalves, fishes and stegocephalian amphibians. Sedimentology and palynology (Mørk et al., 1999b) indicate a moderately deep shelf environment but closer to the sediment source than the overlying Vendomdalen Member. The base of the Fish niveau has never been formally designated, but the vertebrate remains are typically concentrated within the uppermost 10 m of the Lusitaniadalen Member.

The Fish niveau is well constrained biostratigraphically by the rich ammonoid fauna (Weitschat & Lehmann, 1978; Weitschat & Dagys, 1989; Dagys & Weitschat, 1993; Mørk et al., 1999b). The lower part belongs to the *Euflemingites romunderi* Zone of Smithian (early Olenekian) age, while the uppermost 3–4 m contain abundant ammonoids of the *Wasatchites tardus* Zone, late Smithian. The Smithian–Spathian boundary is marked by a positive carbon isotope excursion that is recognised globally, and provides an excellent chemostratigraphic marker in Spitsbergen (Galfetti et al., 2007). The peak of this excursion is found a few metres above the top of the Lusitaniadalen Member, placing the Fish niveau entirely within the Smithian.

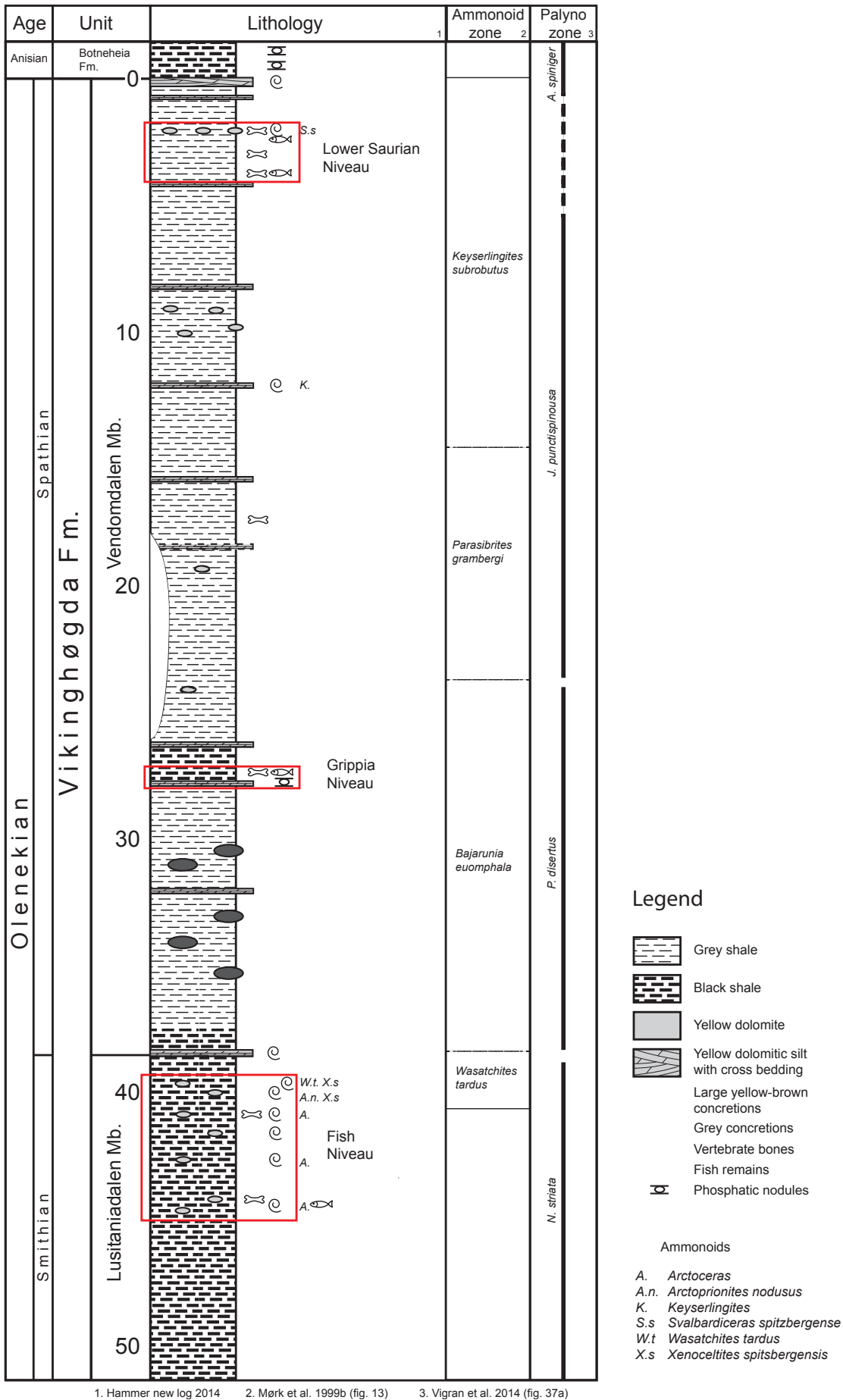


Figure 2. Litho- and biostratigraphy at Marmierfjellet. Parts modified from Mørk et al. (1999b) and Vigran et al. (2014).

### Vendomdalen Member

The Vendomdalen Member is the uppermost of three members in the Lower Triassic Vikinghøgda Formation (Fig. 2), which is overlain by the Botneheia Formation (Mørk et al., 1999b). The depositional environment is interpreted to be distal shelf, below wave base, with a high accumulation of organic material in a low oxic environment (Mørk et al., 1999b). The member represents a transgressive sequence, observed by a reduction in terrestrial palynodebris, increase in marine plankton, decreased sedimentation rate and current indicators (Mørk et al., 1999b). It is characteristically defined by its 94 m-thick transgression sequence at Vikinghøgda, consisting of dark grey, silty, laminated mudstone, together with yellow weathering dolomite nodules and beds (Mørk et al., 1999b).

At the mountain Vikinghøgda (Fig. 1), the Vendomdalen Member has been dated by ammonoids to be of Spathian age by Mørk et al. (1999b), where the lower part is identified as the *Bajarunia euomphala* Zone, above this the *Parasibirites grambergi* Zone, and the upper part as the *Keyserlingites subrobustus* Zone. The Spathian ammonoid zones are poorly resolved in the Boreal Ocean, but the *Bajarunia euomphala* Zone should be of earliest Spathian age while the *Parasibirites grambergi* Zone is of early to middle Spathian age. The *Keyserlingites subrobustus* Zone is late Spathian in age (Jenks et al., 2015). The section is also dated with palynomorphs by Vigran et al. (2014), where the lower part of the section is assigned to the *Pechorosporites disertus* Composite Assemblage Zone, early Spathian age, and the upper part to the *Jerseyiaspora punctispinosa* Composite Assemblage Zone of late Spathian age. Conodonts have been recovered throughout the Lower and Middle Triassic of Spitsbergen; however, they are sparse in the Spathian succession of both Svalbard and Arctic Canada. Nakrem et al. (2008) described four conodont collections from the upper part of the Vendomdalen Member at Milne Edwardsfjellet, sampled at 31, 48, 49.1 and 54.4 m above the base of the member. These levels concur with the range of the *Keyserlingites subrobustus* Zone, and an occurrence of *Keyserlingites subrobustus* was reported 36 m above the base of the member (Nakrem et al., 2008). So far, no conodonts older than the *Keyserlingites subrobustus* Zone are known from the Spathian Stage in Canada or Svalbard (Nakrem et al., 2008), and palynomorphs from the Vikinghøgda type sections are generally poorly preserved (Mørk et al., 1999b). The biostratigraphy of the Vendomdalen Member was correlated to magnetostratigraphy by Hounslow et al. (2008a). Hounslow et al. (2008b) further constructed a biomagnetostratigraphy for the Vikinghøgda Formation using the ammonoid biostratigraphy of the Arctic Boreal successions. Later, Hounslow & Muttoni (2010) correlated the Early Triassic strata of Svalbard with other Triassic sections globally. The Vendomdalen Member shows mostly normal polarity, with two polarity reversals that allowed for close correlation to other Spathian

sections of the Boreal Lower Triassic (Hounslow et al., 2008b).

The Vendomdalen Member at Marmierfjellet is approximately 39 metres thick and contains the vertebrate-bearing *Grippia* niveau and the Lower Saurian niveau (Fig. 2). The thickness at Marmierfjellet is comparable to nearby Ledalen (Vindodden), to one section at Stensiöfjellet in Sassendalen, and to Roslagenfjellet, East Spitsbergen, but is considerably thinner than at Vikinghøgda (94 m) and Milne Edwardsfjellet in Sassendalen (Vigran et al., 2014). This considerable thickness variation may be due to true lateral variation in sedimentation rate, and/or tectonic disturbance (especially thrust faults), and may complicate stratigraphic correlation. The carbonate beds consist either of calcite or dolomite, or a combination of these minerals. Some beds are formed mainly by flattened nodules or concretions, the latter of which are often septarian concretions. The beds are typically 5–20 cm thick and yellow-weathering; and these yellow-weathering horizons make the Vendomdalen Member distinct from the lower members of the Vikinghøgda Formation (Mørk et al., 1999b). The upper siltstone bed is rich in ammonoids, with *Keyserlingites subrobustus* and *Svalbardiceras spitsbergense* in limestone concretions (Hounslow et al., 2008a; Weitschat, 2008).

### Botneheia Formation

A possible transgressive event marks the boundary between the Vikinghøgda and Botneheia formations, and the upper sequence of the Botneheia Formation represents a moderately deep shelf environment, with distal deltaic lobes (Vigran et al., 2014). The formation is divided into the Anisian Muen Member and the Ladinian Blanknuten Member (Krajewski, 2008; Vigran et al., 2014). Dark, organic-rich shales with an abundance of phosphate nodules, infRACTED by alternating layers of bioturbation, characterise the formation (Lundschien et al., 2014; Vigran et al., 2014). As such, the depositional environment was mainly anoxic with short oxic periods (Vigran et al., 2014). An ammonoid zonation of the section has been produced (Weitschat & Lehmann, 1983), and correlated to the recent palynozonation (fig. 3a in Vigran et al., 2014) with the *Indigirites tozeri* zone as Upper Ladinian, the *Tsvetkovites varius* zone as Lower Ladinian and the *Frechites laqueatus* zone as Upper Anisian. The *Indigirites tozeri* ammonoid zone (Weitschat & Lehmann, 1983) was correlated to the Blanknuten Member by Hounslow et al. (2008b).

The Upper Saurian niveau is situated in the Blanknuten Member bordered by the Muen Member below and the Tschermarfjellet Formation above. The member is characterised by black shale and siltstone deposits with carbonate cementation, and an abundance of the bivalve *Daonella* (Weitschat & Lehmann, 1983; Vigran et al., 2014). Typically, the ichthyopterygians found in the Upper Saurian niveau are embedded in the

*Daonella* Shales. The preservation of ichthyopterygians is documented in at least the upper 10 metres of the formation.

## Conclusions

Although the palaeontology of the Lower to Middle Triassic succession in Spitsbergen has been studied for more than a century and a half, recent fieldwork has demonstrated that there is still much to discover. However, our ability to properly interpret the evolutionary and ecological implications of the faunas requires precise stratigraphic control that has only recently become available. Within such a detailed stratigraphic framework, the Spitsbergen material promises to contribute to our understanding of the faunal recovery after the end-Permian mass extinction, the early evolution of Ichthyopterygia, the dynamics of the Smithian–Spathian turnover event and the marine palaeobiogeography and palaeoecology of the Boreal Triassic.

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