

# The Devonian-Carboniferous boundary in the Graz Paleozoic (Eastern Alps, Austria) and its global significance

## El límite Devónico-Carbonífero en el Paleozoico de Graz (Alpes orientales, Austria) y su importancia global

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### ABSTRACT

Pelagic, outer-shelf sedimentary successions across the Devonian-Carboniferous boundary (DCB) in the western Graz Paleozoic at Trolp (Eastern Alps, Austria) display exceptional, continuous limestone facies of the Hangenberg Crisis interval. These enable the study of conodont successions, lithofacies, and geochemistry without noteworthy interruptions by siliciclastic intervals, which are equivalents of the Hangenberg Black Shale and Hangenberg Sandstone. The *praesulcata*, *kockeli*, and *sulcata/kuehni* conodont Zones are recognized by stratigraphically significant markers in the early siphonodellids and early protognathodids within thin-bedded limestones. The major conodont biofacies change and initial mass extinction phase (*costatus-kockeli* Interregnum) of the Hangenberg Crisis is succeeded by radiations among conodonts in the *kockeli* and *sulcata/kuehni* Zones. Carbon isotopes ( $\delta^{13}\text{C}_{\text{carb}}$ ) record positive excursions and enhanced  $\text{C}_{\text{org}}$  burial episodes; the characteristic signals can be used as chemo-indicators of the early, middle and late phases of the Hangenberg Crisis. The global common *Siphonodella sulcata* Morphotype 5 and *Protognathodus kuehni* have their first occurrence at the base of the Tournaisian at Trolp and elsewhere in Europe, Asia and North Africa, and thus can be used to define the base of the *sulcata/kuehni* Zone and the current DCB level. 'Protognathodus' faunas with a high morphological variability are recorded in the *praesulcata* and *kockeli* Zones, but spread mainly in the *sulcata/kuehni* Zone, or even later in the lower Tournaisian. Therefore, the extended *kockeli* Zone suggested recently, and even leaves the current DCB level undivided, as well as the use of *Protognathodus kockeli* as index fossil for the DCB, are not applicable as suggested herein.

A precise biozonation concept as well as multidisciplinary approaches used at Trolp are needed for understanding ultimate causes of the 1st order Hangenberg mass extinctions and major environmental changes at the DCB. Multidisciplinary approaches comprise taxonomically significant index conodonts, extinctions and radiations, as well as lithofacies and isotopic analyses. These markers at Trolp can be well correlated with those from many other settings worldwide, reflecting the global significance of the Graz Paleozoic.

**Keywords:** Hangenberg Crisis, conodont mass extinctions, stable isotopes, Trolp quarry, Protognathodus fauna, kockeli Zone, sulcata/kuehni Zone.

### RESUMEN

Las sucesiones sedimentarias pelágicas de la plataforma exterior a lo largo del límite Devónico-Carbonífero (DCB) en el Paleozoico occidental de Graz en Trolp (Alpes orientales, Austria) muestran facies de piedra caliza continuas y excepcionales en el intervalo de la crisis de Hangenberg. Estos permiten el estudio de sucesiones de conodontes, litofacies y geoquímica sin interrupciones notables por intervalos siliciclásticos, que son equivalentes de Hangenberg Black Shale y Hangenberg Sandstone. Las zonas de conodontes *praesulcata*, *kockeli* y *sulcata/kuehni* se reconocen mediante marcadores estratigráficamente significativos en los primeros sifonodélidos y los primeros protognatódidos dentro de calizas de estrato delgado. El cambio de biofacies de conodontes principales y la fase inicial de extinción masiva (interregno *costatus-kockeli*) de la crisis de Hangenberg es seguida por radiaciones entre conodontes en las zonas de *kockeli* y *sulcata/kuehni*. Los isótopos de carbono ( $\delta^{13}\text{C}_{\text{carb}}$ ) registran excursiones positivas y episodios de entierro de  $\text{C}_{\text{org}}$  mejorados; las señales características pueden utilizarse como quimioindicadores de las fases temprana, media y tardía de la crisis de Hangenberg. El morfotipo 5 de *Siphonodella sulcata*, común a nivel mundial y *Protognathodus kuehni* tienen su primera aparición en la base del Tournaisiano en Trolp y en otras partes de Europa, Asia y el norte de África y por lo tanto, pueden usarse para definir la base de la zona *sulcata/kuehni* y el nivel DCB actual. En las zonas *praesulcata* y *kockeli* se registran faunas de 'Protognathodus' con una alta variabilidad morfológica, pero se extendieron principalmente en la zona *sulcata/kuehni*, o incluso más tarde en el Tournaisiano inferior. Por lo tanto, la zona de *kockeli* extendida sugerida recientemente, e incluso deja indiviso el nivel actual de DCB, así como el uso de *Protognathodus kockeli* como fósil índice para el DCB, no son aplicables como se sugiere en este documento.

Se necesita un concepto de biozonificación precisa, así como enfoques multidisciplinarios utilizados en Trolp, para comprender las causas últimas de las extinciones masivas de primer orden del Hangenberg y los principales cambios ambientales en el DCB. Los enfoques multidisciplinarios comprenden conodontes índice taxonómicamente significativos, extinciones y radiaciones, así como litofacies y análisis isotópicos. Estos marcadores pueden correlacionarse bien con los de muchos otros entornos en todo el mundo, lo que refleja la importancia global del Paleozoico de Graz.

**Palabras clave:** Crisis de Hangenberg, extinciones masivas de conodontes, isótopos estables, cantera Trolp, fauna de Protognathodus, zona kockeli, zona sulcata/kuehni

## 1. Introduction

The Graz Paleozoic belongs to the Upper Austroalpine Nappe System (Schmid *et al.*, 2004) of the Eastern Alps and consists of Silurian to Carboniferous sedimentary rocks which crop out in the surroundings of Graz (Styria, southeastern Austria, Figure 1). In general terms, the internal tectonic architecture of the Graz Paleozoic has a structure consisting of a lower, tectonically, and metamorphically more stressed nappe system and an upper one comprising less metamorphic sequences (Gasser *et al.*, 2010; Figure 2). The slightly metamorphic and in places very fossiliferous sequence of the upper nappe system has been the subject of numerous paleontological and sedimentological studies (*e.g.*, Histon *et al.*, 2010; Ebner and Hubmann, 2012). In particular, biostratigraphic and geochemical studies of the stratigraphic younger parts (*e.g.*, Nössing, 1974a, 1974 b; Ebner, 1976b,

1980a; Kaiser, 2005; Bojar *et al.*, 2013; Kaiser *et al.*, 2020) have provided important new data on environmental dynamics during the 1st order mass extinctions of the multiphase Hangenberg Crisis at the Devonian-Carboniferous boundary (DCB; Kaiser *et al.*, 2016; Becker *et al.*, 2016a).

Worldwide the DCB is characterized by either transgressive Hangenberg black shales and regressive Hangenberg sandstone deposits, or by unconformities with stratigraphic gaps (Kaiser *et al.*, 2016). The latter phenomenon is also known from the Graz Paleozoic east of the Mur River (Figure 3). In contrast, stratigraphically comparable sections west of the Mur River show exceptionally continuous limestone sequences. The best outcrop is located in the abandoned Tropp quarry (Figures 2 and 3) and was examined with regard to high resolution biostratigraphy, lithostratigraphy and chemostratigraphy.

The Tropp section focused on herein, consists of pelagic cephalopod limestones of the upper



**Figure 1** Disconnected Paleozoic units in Austria. The Paleozoic units belong to the Eastern Alps (Graz Paleozoic, Greywacke Zone, Gurktal Nappe, South Burgenland) and the Southern Alps (Carnic Alps, South Karawanke Mts.) which are separated by the Periadriatic Fault, modified from Hubmann *et al.* (2003).

Famennian Steinberg Formation and Tournaisian Sanzenkogel Formation. The DCB is located between both units (Figures 4-8). A positive carbon isotope excursion ( $\delta^{13}\text{C}_{\text{carb}}$ ) measured previously in thin-bedded limestones coincides with the major change in conodont biofacies and the initial mass extinction episode of the Hangenberg Crisis (Figure 9-11). After Kaiser *et al.* (2020), recent high-resolution geochemical and biostratigraphic studies at Tropp, and the correlation with the Grüne Schneid section in the Carnic Alps (see reviews by Schönlaub, 2018; Spalletta *et al.*, 2021), reveal the utility of the joint *sulcata/kuehni* Zone, as well as uncertainties in the taxonomy of the *Protognathodus* fauna (Figures 12-14).

During the 1980s, the DCB working group at the time initially considered the base of the *kockeli* (upper *praesulcata*) Zone as a DCB position. However, this idea with the first occurrence (FO) of *Protognathodus kockeli* was rejected after intensive studies that included visits to relevant sections worldwide (*pers. comm.* C. Sandberg, 2019). The current DCB, based on the FO of *Siphonodella sulcata*, was established by the Heerlen Congress in 1935, confirmed by the DCB Working Group in 1988, and since has been applied worldwide. However, the current GSSP at La Serre E' (Montagne Noire, France) was reconsidered by Kaiser (2009) due to former uncertainties in the identification of early siphonodellids (Kaiser and Corradini, 2011; Spalletta *et al.*, 2017; see summary in Becker *et al.*, 2016a; Kaiser *et al.*, 2020; Hartenfels *et al.*, 2022). Also, according to Ziegler and Sandberg (1996), the current GSSP position at La Serre É - between Bed 88 and 89 - was chosen hastily and despite the objections of conodont specialists in the DCB working group. Kaiser's studies in 2005 and 2009 reveal an older FO of the first morphotype (M5) of the index fossil *Siphonodella sulcata* at La Serre É in Bed 84 (not in Bed 86 as shown in figure 1, Aretz and Corradini, 2021). However, from today's perspective (Aretz *et al.*, 2021; Hartenfels *et al.*, 2022), the profile at La Serre É would hardly be an option for a suitable GSSP.

Based on taxonomic studies during the last

decades, 1. uncertainties of the early siphonodellids (*Siphonodella praesulcata*, *Siphonodella sulcata*) resulted in morphotype groups established by Kaiser and Corradini (2011) and are used worldwide (see Kaiser *et al.*, 2020; Hartenfels *et al.*, 2022), 2. the Morphotyp 5 of *Siphonodella sulcata* is used as index fossil for the current DCB (*e.g.*, Kaiser, 2009; Kaiser *et al.*, 2020; Becker *et al.*, 2021; Hartenfels *et al.*, 2022), 3. the 'siphonodelloids' were discriminated from the early siphonodellids (Becker *et al.*, 2013), and 4. it is recently discussed to what extent *Protognathodus kockeli*, which was proposed as a supplementary index fossil for the DCB, would be applicable (Becker *et al.*, 2016a; Corradini *et al.*, 2011, 2016; Kaiser *et al.*, 2020; Aretz *et al.*, 2021; Hartenfels *et al.*, 2022).

In this review article we provide a brief overview of the stratigraphic architecture of the Graz Paleozoic Rannach Nappe and present the faunistic, geochemical and sedimentologically important markers at the DCB in the western Graz Paleozoic. We also highlight the potential for correlation with regions outside the study area. We discuss criteria currently proposed – 1. the extended *kockeli* Zone suggested by Spalletta *et al.* (2017), and 2. the FAD of *Protognathodus kockeli* suggested by Hartenfels *et al.* (2022) - for a revised DCB position and their application to the Graz Paleozoic as well as to other European, North African, and Asian areas. This discussion requires an evaluation of the applicability of the *kockeli* and the *sulcata/kuehni* Zones (= current DCB level). To this end, we provide a discussion of aspects of conodont biostratigraphy, biofacies, and taxonomy; stable isotope geochemistry ( $\delta^{13}\text{C}_{\text{carb}}$ ) and litho- and microfacies aspects are also included, as are earlier and more recent biozonation concepts (see Figure 12).

Abbreviations: DCB = Devonian-Carboniferous boundary, GSSP = Global Stratotype Section and Point, HBS = Hangenberg Black Shale, HBSE = Hangenberg Black Shale Event, FAD = first appearance datum, FO = first occurrence, *ckI* = *costatus-kockeli* Interregnum, HSS = Hangenberg Sandstone, HSSE = Hangenberg Sandstone Event.

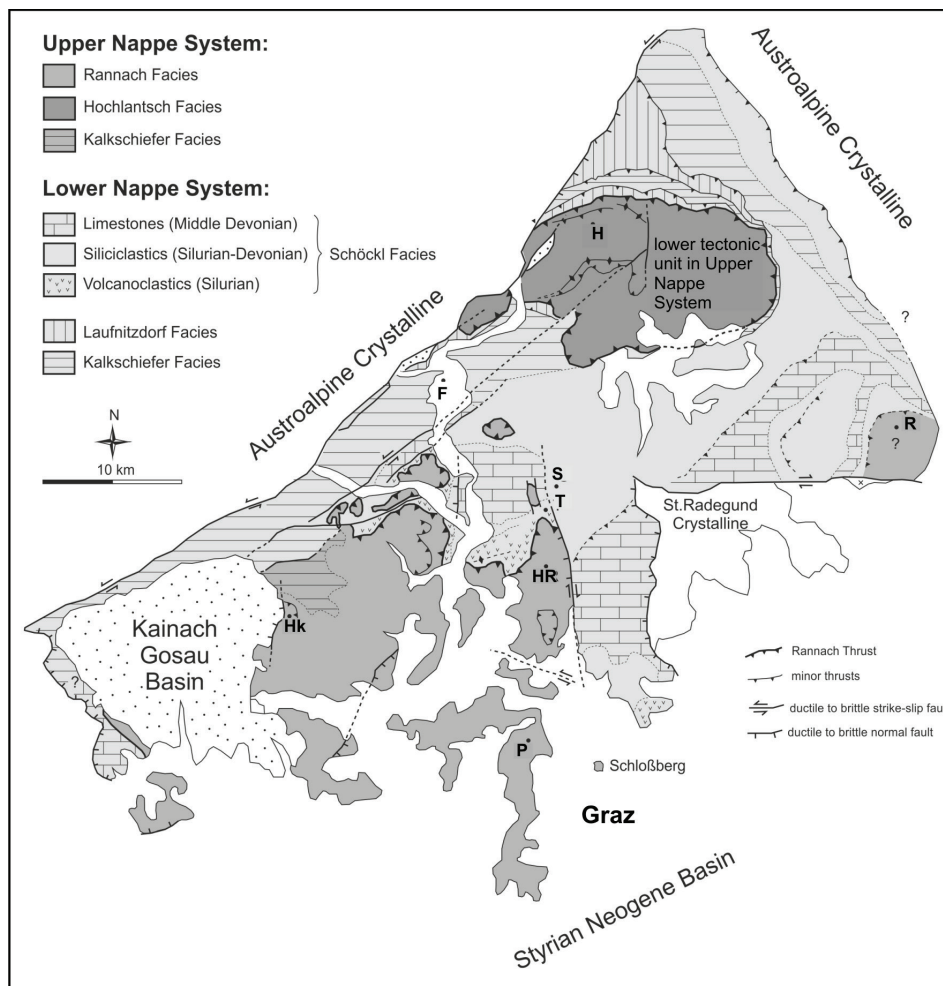
## 2. Overview of geology and facies development in the Graz Paleozoic

### 2.1. OVERVIEW OF GEOLOGY

Within Austria, weakly metamorphosed Paleozoic units are irregularly distributed throughout the Alpine orogen (Figure 1). Their original, pre-Alpine geographical location is uncertain, but it is generally accepted that the units were part of the northern Gondwana margin (e.g., Neubauer *et al.*, 2022). According to tectonic concepts of the Eastern Alps (e.g., Schmid *et al.*, 2004) – except for the Paleozoic remnants south of the Periadriatic fault – they are part of the uppermost tectonic nappes of

the Eastern Alps, the Upper Austroalpine Nappe System. The Upper Austroalpine Nappe System consists of highly varied crystalline units with a dominant pre-Alpine metamorphic overprint, and is tectonically overlain by the Greywacke Zone, which itself is largely unconformably overlain by thick Mesozoic sediments of the Calcareous Alps. The Graz Paleozoic and some isolated outcrops in southern Burgenland which both lack Permo-Mesozoic cover sequences, as well as the Gurktal nappe, are part of the Drauzug-Gurktal Nappe System, which is itself part of the Upper Austrian Basement Nappes.

The structure of the Upper Austroalpine Nappe System was triggered by the closure of the



**Figure 2** The Graz Paleozoic and its organisation in nappe groups (modified from Gasser *et al.*, 2010). H = Hochlantsch, Hk = Höllerkogel, HR = Hohe Rannach, P = Plabutsch, R = Raasberg, S = Semriach, T = Taschen.

Meliata Ocean (a marginal basin of the Tethys) during the Early Cretaceous. Therefore, the Upper Austroalpine Nappe System is sealed by Upper Cretaceous to Eocene sediments. Later, the Upper Austroalpine Nappe System was dismembered by lateral extrusion, *i.e.*, eastward displacement of the eastern parts of the Eastern Alps (Ratschbacher *et al.*, 1991; Neubauer *et al.*, 2000) and thus brought into its current position.

The internal architecture of the Graz Paleozoic (Silurian to Carboniferous) is composed of a high-grade metamorphosed lower nappe system and a low-grade metamorphosed upper nappe system (Gasser *et al.*, 2010; Figure 2). Each of them is differentiated into individual nappes. In short, the volcanosedimentary sequence contains basal pre-Devonian volcanoclastics, and a pronounced platform- to basinal-facies development during the Devonian. In the structurally higher units, especially in the Rannach Nappe, the depositional environment deepens steadily from the Upper Devonian to the Mississippian, and in the Pennsylvanian the depositional environment becomes abruptly shallow marine (Figure 3).

Resting on metamorphic basement, the nappes of the Graz Paleozoic are unconformably overlain by an Upper Cretaceous sequence, the “Kainach Gosau” (Ebner and Rantitsch, 2000) in the west and by Neogene sedimentary rocks of the Styrian Molasse Basin in the south (Gross *et al.*, 2007).

In contrast to the Lower Nappe System which experienced stronger deformation as well as higher metamorphism – up to lower amphibolite facies – sequences of the upper nappe system, among them the sequence of the Rannach Nappe, are quite fossiliferous.

## 2.2. RANNACH NAPPE OF THE GRAZ PALEOZOIC

The sequence of the Rannach Nappe (Figures 3-5) starts with predominantly alkaline metavolcanites (tuffs, lavas) which pass into slightly calcareous sediments and contain fossils from the late Silurian to early Devonian (Ludfordian - Lochkovian). This is followed by various Lower Devonian (Lochkovian)

to Upper Devonian (Frasnian) shallow-water platform carbonates. Formations of this depositional episode are comprised to the Rannach Group (Flügel, 2000). Approximately during the Frasnian to Famennian transition, the depositional environment changed to an open, pelagic marine area. This persisted until the late Mississippian (Serpukhovian). The formations of the pelagic facies are combined into the Forstkogel Group (Flügel, 2000). The end of the Rannach Nappe stratigraphic series is formed by lower Pennsylvanian (Bashkirian) shallow-marine birdseye limestones and phytoclast-rich shales of the Dult Group (Flügel, 2000). Detailed descriptions of formations of the Rannach, Forstkogel and Dult Groups, their lithologies, stratigraphic extents and the extensive literature can be found in the explanations for the “Austrian Stratigraphic Chart 2004” (Hubmann *et al.*, 2013).

The pelagic facies of the Forstkogel Group, which consists of micritic deep-marine cephalopod limestones of the Steinberg and Sanzenkogel Formations (Figures 4 and 5), reaches a maximum thickness of 100 m. In the Rannach Nappe east of the Mur (= Eastern Graz Paleozoic) it reduces down to about 30 m. The reasons for the reduced thickness are intraformational stratigraphic gaps in the area of the DCB, which were created by karstification (Ebner, 1978, 1980a, 1980b).

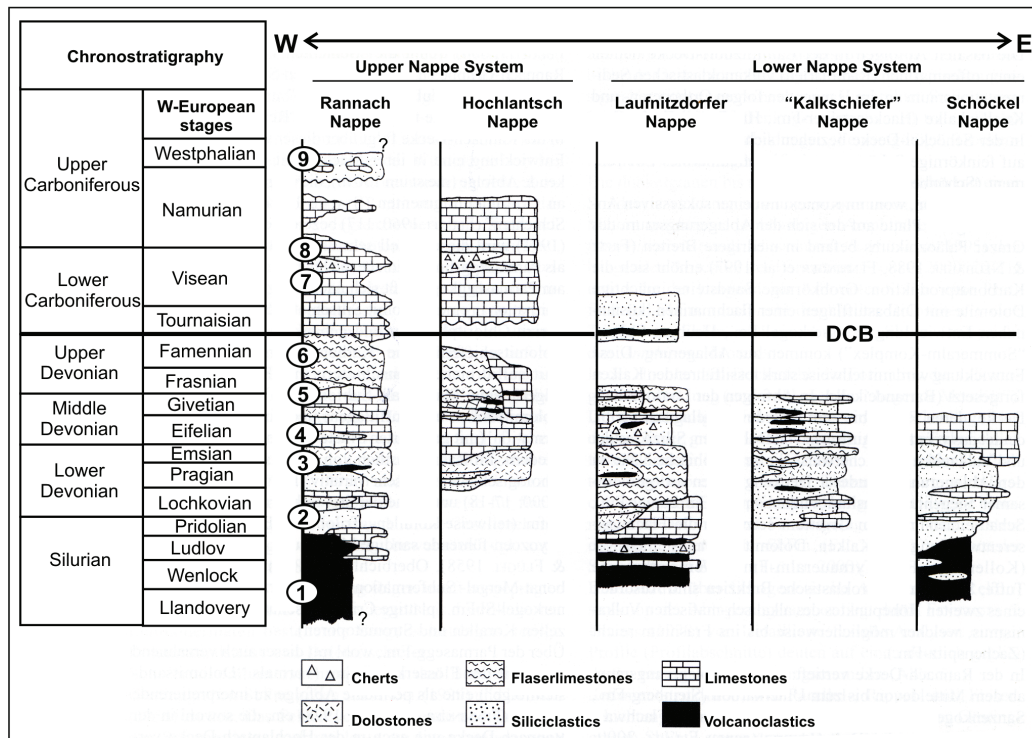
Elementary geochemistry used for interpretations of bathymetry shows clear geographical differences: While in the Graz Paleozoic west of the Mur (= western Graz Paleozoic) the depositional depth in the Mississippian period was around 200 m, in the eastern Graz Paleozoic the water depth was less than 50 m (Nössing, 1974b; Ebner and Prochaska, 1989; Buchroithner *et al.*, 1979). Continuous sequences covering the DCB interval only occur in the western Graz Paleozoic. In the eastern Graz Paleozoic, however, erosion gaps occur in this time interval, with stratigraphic gaps increasing in temporal extent towards the east (Figure 3).

### 3. The Devonian-Carboniferous boundary (DCB) in the western Graz Paleozoic

The Forstkogel Group is, according to conodonts, divided into the Upper Devonian Steinberg Formation and the Lower Carboniferous (Mississippian) Sanzenkogel Fm. (Figure 4). The strongly condensed and only 220 cm thick Lower Sanzenkogel Fm., comprising the Early Tournaisian *Siphonodella sulcata* Zone to the Late Tournaisian *Scaliognathus anchoralis* Zone, occurs only in the western Graz Paleozoic. Marker beds within the Upper Sanzenkogel Fm. are the Trolp Phosphorite Bed (shale and lydite layer with phosphorite nodules) in the western parts of the Rannach Facies (Ebner, 1978; Bosic, 1998, 1999; Flügel, 2000).

Numerous conodont studies from the Graz Paleozoic have been published since the 1950s

because the pelagic limestones of the Forstkogel Group/Rannach Facies yielded rich conodont faunas, which enable a fine-stratigraphic subdivision of the Paleozoic succession. Comprehensive and detailed conodont studies were conducted by Fritz Ebner (*e.g.*, Ebner, 1976a, 1977a, 1977b, 1978, 1980a, 1980b) in the western and eastern Graz Paleozoic. Also, conodont studies from the western Graz Paleozoic were published, for example, by Flügel and Ziegler (1957); Kodsí (1967); Khosrovi-Said (1962); Flajs (1966); Nössing *et al.* (1977); Surenian (1978), Buchroithner *et al.* (1979); Buchroithner and Ebner (1981), and succeeding conodont biostratigraphic and geochemical, high-resolution studies at the DCB at Trolp provide important comprehensive data towards a better understanding of the Hangenberg Crisis at the DCB (Kaiser, 2005; Kaiser *et al.*, 2008, 2009, 2020; Bojar *et al.*, 2013).



**Figure 3** Stratigraphic overview of nappes of the Graz Paleozoic, modified from Ebner *et al.* (2000), Hubmann and Hasenhüttl (1995), Kreutzer *et al.* (1997) and Ebner and Hubmann (2012). Lithostratigraphic column with main lithologies of the Rannach Nappe (Rannach Facies) in the western Graz Paleozoic near Graz: Reinerspitz Group: 1. Kehr Fm., Kötschberg Fm., Rannach Group: 2. Parmasegg Fm., 3. Flösserkogel Fm., Bameder Fm.; 4. Plabutsch Fm., 5. Kollerkogel Fm., Forstkogel Group: 6. Steinberg Fm., 7. Sanzenkogel Fm., Dult Group: 8. Höchkogel Fm., 9. Hahngraben Fm.

An overview of localities with DCB successions in the western Graz Paleozoic (Section 3.1), where successions of the Famennian Steinberg Fm. and Tournaisian Sanzenkogel Fm. crop out, is followed by a detailed description of the currently easily accessible Trolp Quarry (Section 3.2).

### 3.1. OVERVIEW OF LOCALITIES

Faunal and lithologic markers from different lithologic successions from the Graz Paleozoic, such as phosphorite nodules, lydites, fissure fillings, conodont index fossils and mixed conodont faunas, can be easily correlated (Figure 4) and resulted in the reconstruction of the depositional environments (see Section 2.2). The stratigraphic gap between the Steinberg Fm. and the Lower Sanzenkogel Fm. increases from west to east (Figure 3), and complete Famennian and Tournaisian sediments, including the DCB, are recorded at Forstkogel in the Steinberg area (Surenian, 1977, 1978; Nössing, 1974b, Buchroithner *et al.*, 1979; Ebner, 1980a), and at Eichkogel in the Rein area (Figures 4 and 5; Nössing, 1974a; Nössing *et al.*, 1977). Successions at Eichkogel are situated in a forest area and currently covered by forest soil. At Forstkogel, Buchroithner *et al.* (1979) described several quarries where DCB successions crop out, among them the famous Trolp quarry (Figure 6).

Complete successions of the Steinberg Fm. in the western Graz Paleozoic, reaching a thickness of 70 m, are reported at Forstkogel, Eichkogel and Weihermühle in the Gratberg-Au area (Figures 4 and 5; Ebner, 1980a, 1980b). Complete successions of the Sanzenkogel Fm., reaching a thickness of up to 35 m from the *Siphonodella sulcata* to the *Gnathodus bilineatus bollandensis* conodont Zones, are reported at Forstkogel, while a hiatus is already reported from Weihermühle from the lower Tournaisian (Figure 4). The Tournaisian successions at Forstkogel were selected as type locality for the Lower Sanzenkogel Fm., established by Nössing (1974b, 1975). Sedimentologic investigations of the Lower Carboniferous (Mississippian) at Forstkogel were summarized by Ebner and Prochaska

(1989).

In the western Graz Paleozoic, DCB successions at Trolp (see Section 3.2) and at Kanzelkogel (north of Graz, Kodosi, 1967) consist characteristically of condensed thin-bedded micritic limestones. At Kanzelkogel, the Upper Devonian and Lower Carboniferous (Mississippian) are separated by a several-decimetre-thick sequence of thin-bedded limestones with mixed Devonian and Carboniferous conodont faunas (Kodosi, 1967, Figure 12). However, the *Protognathodus* or *Siphonodella* index conodont faunas were not recorded at Kanzelkogel probably due to a stratigraphic gap, or condensation and an incomplete sampling. Unfortunately, these successions do no longer exist due to mining activity.

### 3.2. TROLP SECTION (FORSTKOGEL, STEINBERG AREA)

The Trolp section is an abandoned quarry (ÖK Sheet 163 Voitsberg) located at Forstkogel about 8 km from the city Graz, beside the street from Steinberg to Rohrbach (Figures 5 and 6). Trolp is the type locality of the Late Devonian conodont *Polygnathus styriacus* first described by Ziegler (1957). The Famennian-Tournaisian pelagic cephalopod limestones at Trolp have been intensively investigated due to rich conodont faunas, and sampling by Sandberg and Ebner in 1980 was followed by conodont studies of, for example, Ebner (1980a), Nössing (1975), Kaiser 2005, Kaiser *et al.* (2008, 2020). It was formally proposed as a candidate for the GSSP at the DCB, but the scarcity of macrofossils did not fulfill the criteria required (Ebner, 1980a; Ziegler and Sandberg, 1984a; Ji *et al.*, 1989).

The tectonically inverted (upside-down) successions at Trolp are dated as Late Famennian *expansa-praesulcata* Zone to Late Tournaisian *Gnathodus typicus* Zone (Ebner, 1980a; Kaiser, 2005). A detailed, bed-by-bed or cm-by-cm sampling of DCB beds is required at Trolp due to the condensation of successions (Figures 7 and 8) which may produce a bias of the conodont record; re-numbering of DCB beds was necessary due to weath-

ering and missing old numbers, as well as due to the high-resolution sampling of the successions. As shown in Figure 9, sample and bed numbers of Kaiser (2005) are correlated with bed numbers established by C. Sandberg and F. Ebner in 1980, and Ebner (1980a). The *praesulcata* and *kockeli* Zones, and the *sulcata/kuehni* Zone (DCB level) are readily recognized by their zonal index fossils (Ebner, 1980a, Kaiser, 2005; Kaiser *et al.*, 2009, 2020; Figures 9 and 10a). The Middle *praesulcata* Zone of the standard conodont zonation (Ziegler and Sandberg, 1984b), recognized at Trolp by Sandberg and Ebner in 1980, Ebner (1980a), Bojar *et al.* (2013), Kaiser (2005), and by Kaiser *et al.* (2008) was not considered recently due to the asynchronous last occurrence of *Palmatolepis gracilis gonioclymeniae* defining the base of the Middle *praesulcata* Zone (see discussions in Kaiser, 2005; Kaiser *et al.*, 2009; Corradini *et al.*, 2016; Spalletta *et al.*, 2017). The asynchronous last occurrence of this taxon resulted in different levels of the base of the Middle *praesulcata* Zone as previously recorded at Trolp by the above mentioned authors.

The eastern part of the quarry which is well accessible consists of well-exposed Lower Carboniferous beds, representing the lower member of the pelagic Sanzenkogel Formation (“Untere Sanzenkogel-Schichten”), and of bedded to massive upper Famennian beds, representing the Steinberg Formation. The succession consists of light-grey to brownish pelagic nodular cephalopod limestones and micritic Flaser-limestones of the Famennian Steinberg Fm. and Tournaisian Sanzenkogel Fm.; continuous limestone successions (Figures 7-10a and 10b) deposited during the Hangenberg Crisis interval are recorded at Trolp, which is globally characterized by black shales, sandstones or hiatuses. The stratigraphically lower part of the section consists of bedded to massive limestones of the *praesulcata* Zone, and a lithologic change is marked by a thin, ~1 cm thick shaly layer (basal Bed 10) at the base of thin-bedded limestones that disappears laterally (N 47°04'08", E 15°19'09"). Bojar *et al.* (2013, Figure 3d) recorded a continuous limestone succession lacking this layer in a lat-

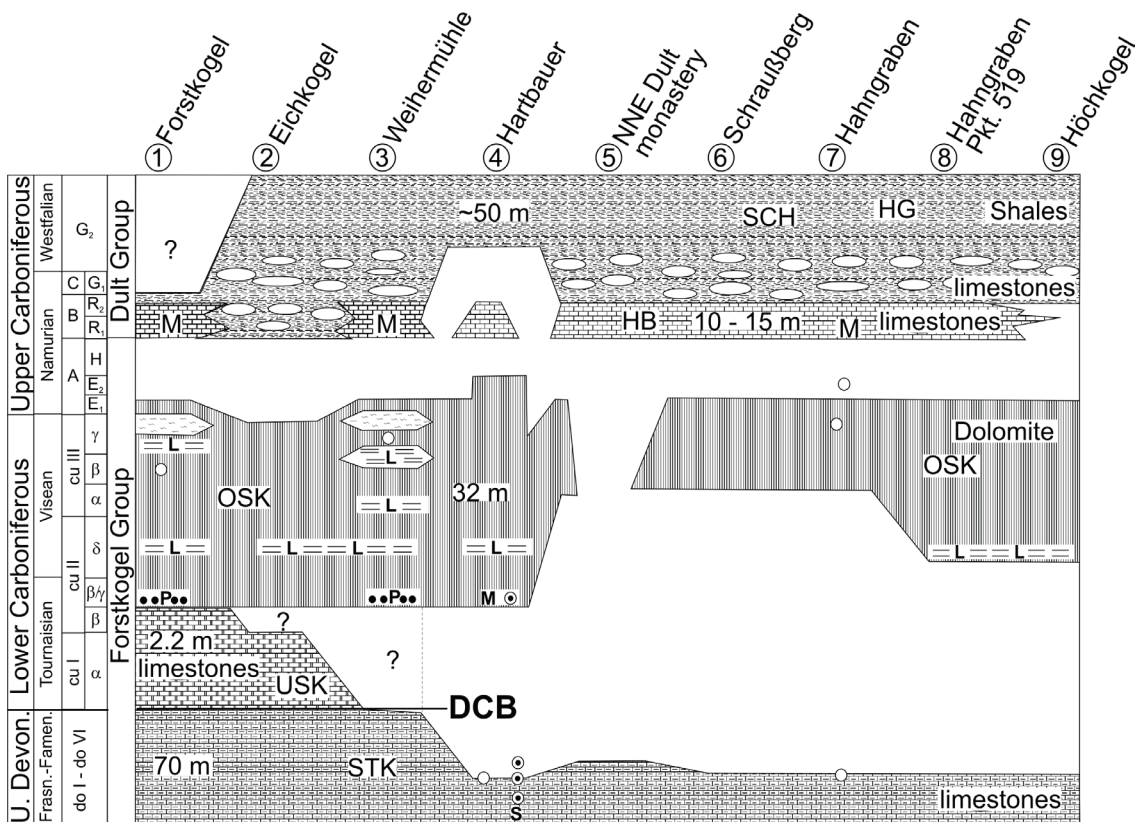
eral, different position next to previous mentioned sampling points (N 47°04'29", E 15°19'15"). This shaly or marly level was regarded by Flügel and Ziegler (1957) as a tectonic disturbance, which locally separated the Devonian and Carboniferous at Steinberg. Due to its stratigraphic position, Kaiser (2005) suggested an equivalent level of the Hangenberg Black Shale (HBS) since the lithologic change is accompanied by the major conodont biofacies change (Figure 11) indicating the main extinction phase of the Hangenberg Crisis at the *costatus-kockeli* Interregnum (*ckI*, after Kaiser *et al.*, 2009 = mass extinction-based level within the Middle *praesulcata* Zone, correlative with the HBSE and HSSE). This interpretation is supported by the onset of conodont radiations and the entry of *Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli* and *Protognathodus kockeli* immediately above in thin-bedded limestone successions conformably overlying the shaly layer (Kaiser *et al.*, 2020; Figures 9 and 10a). The occurrence of stratigraphically leaked younger Carboniferous faunas - mainly siphonodellids and *Gnathodus* - in Devonian samples were recognized about at the level of this shaly layer, or in younger levels, respectively.

The base of the Carboniferous has been fixed at Trolp in Bed 16 by the first occurrence of index conodonts at the same stratigraphic levels (Figures 8, 9 and 12): *Polygnathus purus subplanus*, an index conodont for the base of the Tournaisian (*e.g.*, Kaiser *et al.*, 2009; Spalletta *et al.*, 2017), as well as *Siphonodella sulcata* (M5) and *Protognathodus kuehni*, which were used to define the joint *sulcata/kuehni* Zone and the DCB (Kaiser *et al.*, 2020; Becker *et al.*, 2021). Conodont index fossils previously recorded by Ebner (1980a) and Nössing (1975) at Trolp, and their correlation with index fossils recorded by Kaiser (2005) and Kaiser *et al.* (2020) are shown in Figure 9; the biostratigraphic correlation of the Hangenberg Event and the DCB from the Carnic Alps and Rhenish Massif is shown in Figure 8. At stratigraphically younger levels, *Siphonodella bransoni* (*Siphonodella duplicata* M1), as well as *Pseudopolygnathus multistriatus* were recorded (Ebner

1980a). The condensed Tournaisian sequence is intercalated by the 20 cm thick Trolp Phosphorite Bed (Figures 4 and 14), marking the beginning of the Upper Sanzenkogel Fm. (base of the upper Tournaisian) after Nössing (1975), or the base of the middle Tournaisian as suggested by Kaiser (2005), respectively. It is correlative with the end of the erosional gap within parts of the Rannach Nappe, and interpreted as deepening and upwelling at the shelf margin at the base of the *Scaliognathus anchoralis* Zone (base of upper Tournaisian; Ebner, 1998; Ebner *et al.*, 2000).

Conodont biofacies at Trolp (Figure 11) indicates a dominance of *Bispathodus*, *Brammehla* and

*Palmatolepis* in pre-Hangenberg Crisis (pre-extinction) levels in the *praesulcata* Zone (Kaiser 2005). The conodont mass extinction in the *ckI* at Trolp (base of Bed 10), and major biofacies change to an impoverished *Polygnathus-Protognathodus* biofacies with stunted faunas, is connected to the extinction of Famennian taxa which predominated in the Famennian, such as *Brammehla suprema*, *Bispathodus costatus*, *Palmatolepis gracilis expansa*, and *Pseudopolygnathus marburgensis trigonicus*. The beginning of radiations in the basal *kockeli* Zone, recorded in the basal part of the thin-bedded limestones (Bed 10/11), is succeeded by faunal recovery and the 2nd radiation phase (Figure 10a) in the



**Figure 4** Outcrop areas (1 - 9, from west to east) within the Upper Devonian and Carboniferous of the Rannach Nappe (Rannach Facies) in the Graz Paleozoic (modified from Ebner, 1978, 1980b; Ebner *et al.*, 2000). 1. Forstkogel, Steinberg area (Trolp Quarry, locations 3, 4 in Figure 5), 2. Eichkogel, Rein area (for location see Figure 5), 3. Weihermühle, Gratwein/Au area (location 1 in Figure 5), 4. Hartbauer/Kanzelkogel (location 2 in Figure 5), 5. NNE Dult monastery (location 6 in Figure 5), 6. Schraußberg, 7. Hahngraben (location 5 in Figure 5), 8. Hahngraben Pkt. 519, 9. Höchkogel. Forstkogel Group: STK: Steinberg Formation, USK: Lower Sanzenkogel Fm., OSK: Upper Sanzenkogel Fm., P = phosphorites of Trolp Bed, L = lydites, the lowermost = Hart Bed. Dult Group: Höchkogel Fm.: HB Hartbauer Member, SCH Schrausbauer Mb., HG Hahngraben Fm., M = conodont mixed faunas (time of origin), S = fissure filling, black circle = components of fissure filling, white circle = mixed faunas (components).

basal Lower Carboniferous *sulcata/kuehni* Zone. It is connected to the return of normal-sized conodont faunas, and an increasing abundance of pseudopolygnathids and siphonodellids, but the *Polygnathus-Protognathodus* biofacies remains in the Lower Carboniferous.

Faunal evolution during the Hangenberg Crisis is precisely documented at Trolp (Kaiser *et al.*, 2020). A faunal assemblage of the Lower *Protognathodus* fauna, with *Protognathodus meischneri* and *Protognathodus collinsoni*, enters at a level with the initial litho- and biofacies change (Bed 10, Figures 9, 10a and 11), and the absence of *Bispathodus costatus* and *Protognathodus kockeli* at the same level in Bed 10 is interpreted as the event-based *ckI*. The succeeding entry of *Protognathodus semikockeli* recently established by Hartenfels *et al.* (2022) occurs immediately above, and the entry of *Protognathodus kockeli* s.str. slightly above marks the start of the *kockeli*

Zone (Figures 9 and 10a). The *Protognathodus* fauna at the DCB at Trolp even occurs in association with *Siphonodella praesulcata* and *Siphonodella sulcata* which was also recorded by Sandberg *et al.* (1983).

At Trolp, a phyletic lineage of the early protognathodids (*Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli*, *Protognathodus kockeli* s.str., advanced *Protognathodus kockeli*, *Protognathodus kuehni* (see figure 6 in Kaiser *et al.* 2020) as proposed by Ziegler (1973), Ziegler and Leuteritz (1970) and Hartenfels *et al.* (2022) is recognized on the basis of the shape and the ornamentation of the cup of typical morphotypes (Figures 12 and 13). Many specimens, however, can be designated as atypical morphotypes (Figure 13; see Kaiser *et al.*, 2019 *unpublished*) recorded by Nössing (1975), Ebner (1980a) and Kaiser *et al.* (2020).

Stable isotopes ( $\delta^{13}\text{C}$ ) at Trolp (Kaiser *et al.* 2008; Figure 10b), used for the reconstruction of

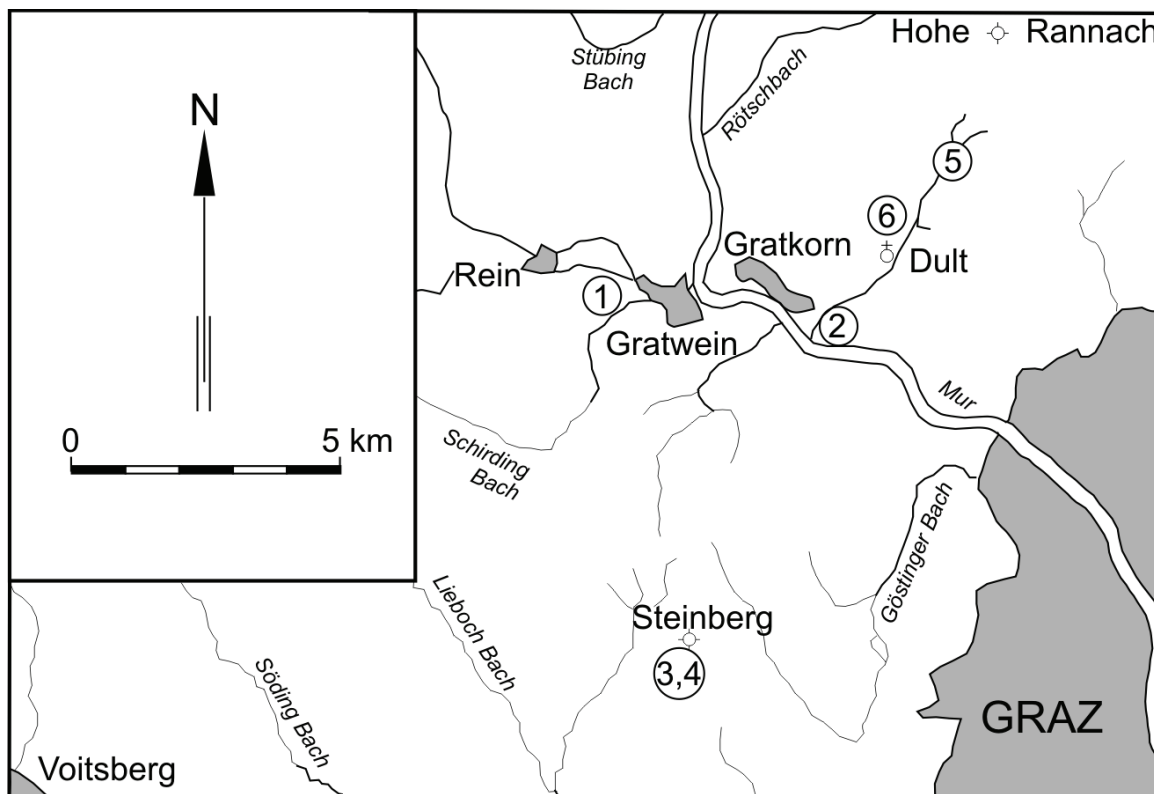


Figure 5 Geographic map and locations of outcrop areas near Graz in the western Graz Paleozoic, modified from Ebner (1980a). 1: Section Weihermühle, 2: Section Hartbauer, 3, 4: Steinberg area, 5: Section Hahngraben, 6: Section NNE Dult Monastery.

changes in the marine dissolved inorganic carbon reservoir, revealed a positive excursion in  $\delta^{13}\text{C}_{\text{carb}}$  in the *ckI* and *kockeli* Zone to values of up to 3‰ (V-PDB). Two minor positive  $\delta^{13}\text{C}_{\text{carb}}$  peaks were recorded in the *praesulcata* Zone, with values reaching 2.5‰ and 2.7‰, and one positive peak was measured in the Upper *expansa* Zone (*Bispathodus ultimis* Zone after Becker *et al.*, 2021). The onset of continuously decreasing carbon isotope values is at the base of the *sulcata/kuehni* Zone (DCB). A  $\delta^{13}\text{C}_{\text{org}}$  excursion was measured from the sedimentary organic matter at Trolp, and oxygen isotope values of conodont apatite at Trolp are remarkable high indicating low seawater temperatures (*unpublished data*). Element geochemistry conducted by Bojar *et al.* (2013) at Trolp indicates a decrease of detrital influence in the lower Tournaisian and was interpreted as transgression.

## 4. Discussion

### 4.1. LITHO-, BIO-, AND CHEMOSTRATIGRAPHY AT TROLP AND CORRELATIONS OUTSIDE THE REGION

Characteristic markers and abundant conodonts at Trolp allow a high time-resolution study of major environmental changes around the DCB. These can be well correlated with coeval markers from continuous limestone successions as well as siliciclastic-dominated successions in North-America, Asia, Europe, North-Africa from pelagic, hemipelagic and neritic realms as discussed as follows, indicating environmental changes on a global scale.

#### 4.1.1. LITHOSTRATIGRAPHY AND LITHOFACIES

At Trolp, a characteristic lithofacies change (Figure 7), from bedded or massive limestones to thin-bedded limestones (mud- to wackestones, Kaiser 2005), is correlative with the global change to siliciclastics (HBS, HSS) and the anoxic episode of the Hangenberg Crisis in the extinction-based *ckI* and *kockeli* Zone (Figure 8-10). In continuous, pelagic limestone successions in the Carnic Alps

(Grüne Schneid, Figure 8), and in hemipelagic to pelagic, more proximal successions in the Montagne Noire (La Serre É, *e.g.*, Flajs and Feist, 1988; Feist *et al.*, 2021) and in the Moravian Karst (Kumpan *et al.*, 2021), a lithofacies change from bedded to thin-bedded, condensed limestones can be correlated with equivalent changes at Trolp. This lithofacies change is globally recorded, and correlative for example with black shales and/or sandstones in pelagic but more deeper settings in the Carnic Alps (Kronhofgraben), Pyrenees (Milles), as well as in hemipelagic settings in the Rhenish Massif (Figure 8), for example at Drewer, or in neritic successions in the Meseta and Anti-Atlas, Morocco (Ain Jemaa, El Atrous, M'Karig, Lambidia), and in the Alborz Mts., Iran (Mighan, Chelcheli), as reported by several authors (*e.g.*, Cygan and Perret, 2002; Kaiser, 2005; Kaiser *et al.*, 2006, 2007, 2011, 2013, 2018; Becker *et al.*, 2016b; Bahrami *et al.*, 2021; Parvizi *et al.*, 2021; Königshof *et al.*, 2021).

The lithofacies change was caused by a global carbonate crisis at the end of the Devonian (Li *et al.*, 2022) due to ceased carbonate sedimentation, enhanced  $\text{C}_{\text{org}}$  burial and marine anoxia during the Hangenberg Black Shale Event (HBSE), and the succeeding “Hangenberg glaciation” during the Hangenberg Sandstone Event (HSSE; *e.g.*, Isaacson *et al.*, 2008; Kaiser *et al.*, 2006; Lakin *et al.*, 2016). The main regressive phase which led to the global deposition of Hangenberg Sandstone equivalents cannot be recognized by a clear sedimentological marker at Trolp, while the onset of the anoxic episode may be represented by the thin shaly layer (base of Bed 10), which is, however, only locally developed.

Tournaisian cherty sediments at Trolp (Trolp Phosphorite Bed, Figures 4 and 14), can be correlated with the basal middle Tournaisian as suggested by Kaiser (2005), and with the faunal turnover during the Lower Alum Shale Event which is globally characterized by a lithologic change from limestones to cherty rocks, the equivalents of the Rhenish Lower Alum Shale of the Middle *crenulata* Zone (Becker, 1993a, 1993b; Kaiser *et al.*,

2018). It is coeval with transgressive shales, partly associated with phosphorite nodules, for example at Kronhofgraben, Carnic Alps (Schönlaub, 1969; Kaiser, 2005), Puech de la Suque and La Serre É, Montagne Noire (Lethiers and Feist, 1990; Kaiser, 2005; Feist *et al.*, 2021), and at El Atrous, Jebel Ouououfilal, M'Fis, M'Karig, Anti-Atlas, Morocco (Kaiser *et al.*, 2011, 2013, 2018). The base of the upper Tournaisian anchoralis Zone is globally characterized by an end of an erosional gap, deepening and upwelling. It is correlative with cherty sediments all over the Graz Paleozoic and with the end of an erosional gap within the Rannach Nappe, interpreted as deepening and upwelling event by Ebner (1998) and Ebner *et al.* (2000).

#### 4.1.2. BIOSTRATIGRAPHY AND CONODONT BIOFACIES

A continuous record of biofacies change at the DCB can be well recognized at Trolp, as well as in continuous limestones in the Carnic Alps, Grüne Schneid (Figure 11), and there is no hiatus in each phases of the Hangenberg Crisis. The dominance of the *Protognathus* fauna and the *Polygnathus purus* group in the *kockeli* and *sulcata/kuehni* Zones at Trolp, Grüne Schneid and elsewhere characterizes the 1st (FO of *Protognathus kockeli*, = base of *kockeli* Zone) and 2nd (FO of *Protognathus kuehni*, FO of *Siphonodella sulcata* M5, = basal Tournaisian transgression) faunal recovery phases after the Hangenberg extinction (Figure 14). The 3rd radiation phase is in the middle *sulcata/kuehni* Zone (FO of *Siphonodella sulcata* M4, *Gattendorfia* level = Tournaisian transgression).

At Trolp, the Hangenberg extinction-based *ckI* is recognized by the absence of *Bispathodus costatus* and *Protognathus kockeli* at same level, and the extinction of almost all upper Famennian conodont taxa among the pseudopolygnathids, polygnathids, bispathodids, branmehlids and palmatolepids (see Kaiser *et al.*, 2009, 2020). It is recognized by a major biofacies change from a Famennian *Bispathodus-Branmehla-Palmatolepis* biofacies to a characteristic, impoverished latest Famennian and earliest Tournaisian *Polyg-*

*nathus-Protognathodus* biofacies. The *ckI* and the major conodont biofacies change are recorded at the base of the thin-bedded limestones at Trolp, as well as in continuous pelagic and hemipelagic limestone successions at the base of thin-bedded limestones in the Carnic Alps at Grüne Schneid (Kaiser, 2007; Figure 11), in the Moravian Karst at Krtny (Kumpan *et al.*, 2021), and in the Montagne Noire at La Serre É (Flajs and Feist, 1988; Feist *et al.*, 2021, Kaiser unpublished data) at correlative levels, and is caused by major environmental changes during the HBSE and HSSE.

Conodont faunal distributions and lithofacies at Trolp display distal, pelagic depositional environments, with sediment deposition in ca. 200 m water depth (*e.g.*, Nössing, 1975). For comparison, litho- and biofacies, and the depositional environments in the Carnic Alps at Grüne Schneid (Schönlaub *et al.*, 1988, 1991) are similar to that of Trolp, and a rich *Protognathodus* fauna occurs in both regions (Kaiser *et al.*, 2020). Differences only related to single occurrences of *Protognathodus* specimens in pre-Hangenberg extinction levels (*praesulcata* Zone) at Grüne Schneid, with minor differences in biofacies, only. Accordingly, a slightly higher abundance of *Bispathodus* than of *Palmatolepis* occurs in the Famennian in pre-crisis levels at Trolp (Figure 11) which is an indicator for more shallow depositional environments according to previous biofacies models (see Kaiser *et al.* 2017). For example, *Bispathodus* abundantly occurs in neritic settings in Iran (Alborz Mts., *e.g.*, Mighan, Chelcheli), Morocco (Tafilalt, *e.g.*, El Atrous), and in hemipelagic settings in the Rhenish Massif (*e.g.*, Hasselbachtal), Montagne Noire (*e.g.*, La Serre, Puech de la Suque), Moravian Karst (*e.g.*, Lesni Lom), as reported by several authors (*e.g.*, Kaiser, 2005; Hartenfels and Becker, 2016; Feist *et al.*, 2021; Kaiser *et al.*, 2011, 2018; Bahrami *et al.*, 2011, 2021; Königshof *et al.*, 2021; Kumpan *et al.*, 2021; Parvizi *et al.*, 2021).

In contrast, siphonodellids, as indicator for more deeper environments (see Kaiser *et al.*, 2017) are slightly more abundant at Trolp in pre-event levels when compared to Grüne Schneid, and *Sipho-*

*nodella* is not recorded in the *ckI* and *kockeli* Zone at Grüne Schneid and elsewhere in the Carnic Alps. Protognathodids, previously suggested as an indicator for more shallow environments, occur at Grüne Schneid in pre-event levels, while this faunal group is not recorded in pre-event levels at Trolp and elsewhere in the Graz Paleozoic. In summary, faunal distribution of *Protognathodus* and *Siphonodella* indicates more shallow depositional environments at Grüne Schneid than at Trolp in pre-event levels, which is, however, not in accordance with the *Bispathodus* biofacies at Trolp as mentioned above.

In this respect, previous biofacies concepts for example of Sandberg and Ziegler (1979), Dreesen

*et al.* (1986) and Ziegler and Weddige (1999), have to be re-considered (see Kaiser *et al.*, 2017, 2020). Paleogeographic provincialism and migration pattern, competition and feeding (Mossoni *et al.*, 2015), but also decreasing ecological niches around the DCB due to global shelf anoxia (Algeo *et al.*, 1995; Li *et al.*, 2022) and major eustatic sea-level fall with longterm and short-termed regressions (*e.g.* Johnson *et al.*, 1985; Kaiser *et al.*, 2011; Babek *et al.*, 2016), as well as climate cooling (Issacson *et al.*, 2008; Brezinski *et al.*, 2009; Lakin *et al.*, 2016), caused more likely characteristic faunal assemblages and biofacies changes. The occurrence of either *Protognathodus* or *Siphonodella* around the DCB is thus more likely related to major environ-

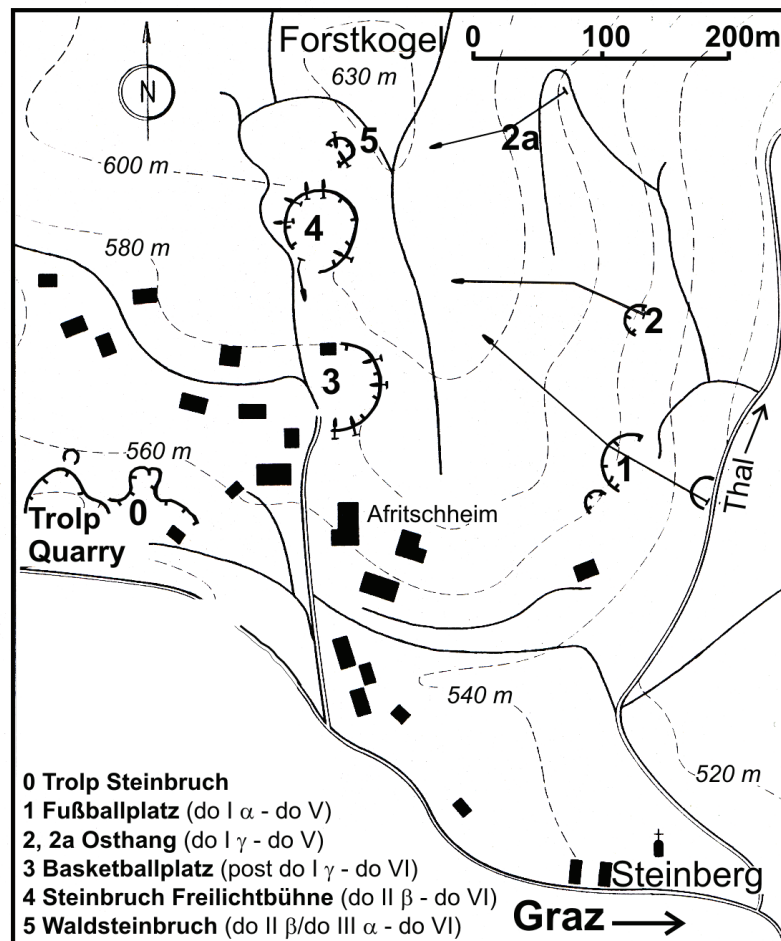


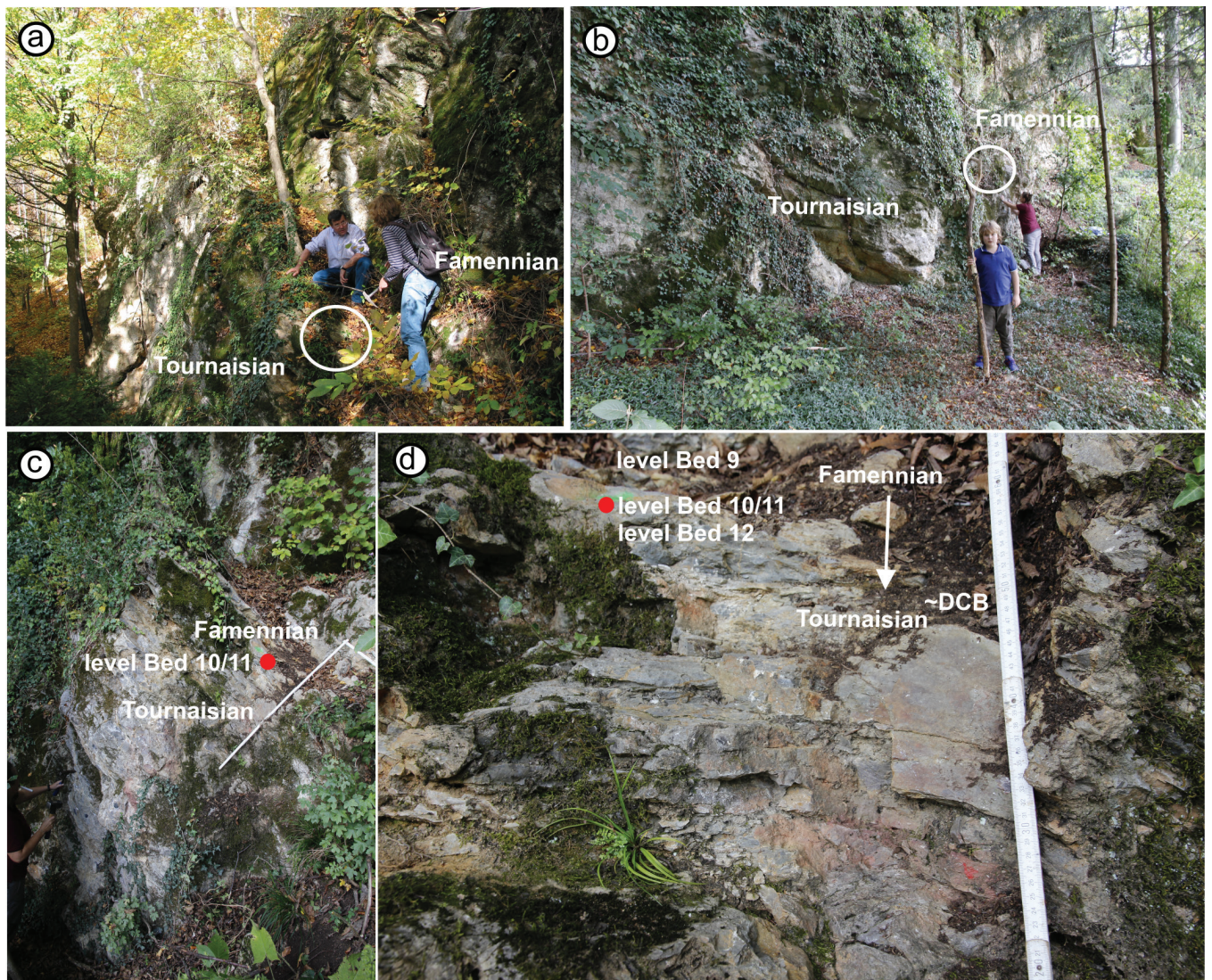
Figure 6 Locations of the Trolp Quarry (0) and other Upper Devonian outcrops (1-5) at Forstkogel in the Steinberg area, modified from Buchroithner *et al.* (1979).

mental changes during anoxic, transgressive and and regressive phases of the Hangenberg Event, and is regarded as biotic opportunism by several authors (e.g., Kaiser, 2005, Kaiser *et al.*, 2008, 2009; Corradini *et al.*, 2011; Spalletta *et al.*, 2017). Thus, disregarding taxonomic uncertainties as discussed below, the occurrence of *Protognathodus* in pre-Hangenberg Event levels in the Carnic Alps and other regions in Europe, Asia and North-Africa could thus be linked with an opportunistic

lifestyle during environmental changes probably related to the Drever Event (early Hangenberg phase) in the *praesulcata* Zone (see Section 4.1.3).

**4.1.3. STABLE ISOTOPES AND CHEMOSTRATIGRAPHY**

Chemostratigraphy as a formal stratigraphic method is a useful tool for global correlations of chronological boundaries (see Sial *et al.*, 2019), correlations of event-related successions (e.g.,



**Figure 7** Outcrop overview and field studies at the abandoned, tectonically disturbed Trolp Quarry, a) sampling in 2002, with Prof. Fritz Ebner (Leoben), b-d) new sampling in 2019, a-b) circles = thin-bedded limestones shown in c-d, and sampling locations. c-d) thin-bedded, continuous limestone successions at the DCB.

Algeo and Shen, 2024), and Hangenberg event-related successions at the DCB (Kaiser *et al.*, 2016). In this respect, positive carbon isotope excursions recorded at Trolp from different levels in the Upper *expansa* (*ultimus* Zone) and *praesulcata* Zones, as well as in the *ckI* and *kockeli* Zone (Figure 10b) can be well correlated outside the region (Figure 10c), and thus indicate enhanced marine  $C_{org}$  burial and global perturbations of the carbon cycle. Accordingly, a minor positive peak in  $\delta^{13}C_{carb}$  in the Upper *expansa* Zone at Trolp, with increasing values of more than 2‰, is widely known from the Middle (*costatus* Zone) and Upper *expansa* Zone as distinctive positive carbon isotope excursions of up to 3‰ in the Carnic Alps (Kaiser *et al.*, 2008), Franconia, southern Germany (*unpublished data*, Kaiser *et al.*, 2022), and the Moravian Karst (*unpublished data*, Kumpan *et al.*, 2018), and were correlated previously with the small-scaled “Epinette” (Middle *expansa*) and “Etreoungt” (Upper *expansa*) bioevents (Kaiser *et al.*, 2008). Increasing  $\delta^{13}C_{carb}$  values and positive carbon isotope excursions in shallow-water successions were also recorded from the Middle and Upper *expansa* Zone in South China (*costatus* and *ultimus* Zone, Zhang *et al.*, 2019), in northern Iran (Parvizi *et al.*, 2021), and in Belgium (Kumpan *et al.*, 2014b). In deeper pelagic settings (Kronhofgraben, Carnic Alps), a positive  $\delta^{13}C_{org}$  spike of 2‰ is recorded as well in the middle part of the Middle *expansa* Zone (Kaiser *et al.*, 2008). Worldwide, transgressive shale and black shale deposits (Algeo *et al.*, 1995; Li *et al.*, 2022; Sahoo *et al.*, 2023), and enhanced  $C_{org}$  burial episodes, as also indicated by an increasing  $\delta^{13}C_{carb}$  baseline on a global scale (Zhang *et al.*, 2019), are recorded in the *expansa* Zone probably connected with the Dasberg Crisis in the Lower and basal Middle *expansa* Zones (see Hartenfels, 2011), and with the Epinette and Etreoungt Events in the middle Middle and Upper *expansa* Zone (*e.g.*, Kaiser *et al.*, 2008; see also Di Pasquo *et al.*, 2019). Two minor positive  $\delta^{13}C_{carb}$  peaks in the *praesulcata* Zone at Trolp, reaching values of more than 2.5‰, can be correlated with coeval positive peaks and/or increasing carbon isotope values of up

to 3‰ recorded from the Carnic Alps at Grüne Schneid, Rio Boreado, Casera Malpasso, Großer Pal (Figure 10cA; Kaiser *et al.*, 2006, 2008), and in Franconia (Kaiser *et al.*, 2022, *unpublished data*), as well as by positive  $C_{org}$  peaks in the Rhenish Massif (Figure 10c B; Kaiser *et al.* 2006). These geochemical anomalies were interpreted as anoxic phases of the Drever Event during the early phase of the Hangenberg Crisis in the *praesulcata* Zone (Kaiser *et al.*, 2020). The Drever Event was previously correlated with the regressive Drever Sandstone and equivalents elsewhere (Streel, 1999; see also Kaiser *et al.*, 2011, 2016), and was previously regarded as a short-term glaciation episode during a humid event at the LL-LE miospore transition (*e.g.*, Streel *et al.*, 2000). The Drever level in the Rhenish Massif is characterized by at least two distinctive shale horizons (Becker *et al.*, 2021). Wet climate and enhanced detritus influx (= regressive Drever Sandstone Event) probably caused enhanced marine bioproductivity and succeeding global shelf anoxia already in the *praesulcata* Zone, well before the HBSE, and may expressed by enhanced carbon burial as indicated by the positive  $\delta^{13}C_{carb}$  peaks.

A significant increase of  $\delta^{13}C_{carb}$  values by almost 2‰ at Trolp is at the base of the thin-bedded limestones associated with the conodont mass extinctions and the major conodont biofacies change in Bed 10 as discussed in Section 4.1.2. The positive  $\delta^{13}C$  Hangenberg-excursion in the *ckI* and *kockeli* Zone of values up to 3‰ was globally recorded (Figure 10c, *e.g.*, Kaiser *et al.*, 2006, 2016; Cramer *et al.*, 2008; Kumpan *et al.*, 2014a; Matyja *et al.*, 2021; Qie *et al.*, 2021; Becker *et al.*, 2021). The onset of the positive excursion indicates global changes of the carbon cycle during the Hangenberg Crisis which was caused by enhanced  $C_{org}$  burial during the deposition of the HBS and equivalents (*e.g.*, Kaiser *et al.*, 2006). Enhanced bioproductivity, high rates of continental weathering and terrestrial nutrient influx due to erosion of organic matter during an upper Famennian sea-level lowstand (Babek *et al.*, 2016) contributed to enhanced organic carbon burial. This

finally resulted in a succeeding major, glacio-eustatic sea-level fall during wet and cold climate, time-equivalent to the deposition of the regressive Hangenberg Sandstone and global equivalents, and a glaciation pulse (Hangenberg glaciation) at the end of the Famennian (*e.g.*, Kaiser *et al.*, 2006; Isaacson *et al.*, 2008; Brezinski *et al.*, 2009; Lakin *et al.*, 2016).

A distinctive decrease of  $\delta^{13}\text{C}_{\text{carb}}$  values, starting in the lower Tournaisian *sulcata/kuehni* Zone at Troip, reflects an overall long-term trend in the lower Tournaisian at same levels in numerous settings in European, Asian and North-American regions (Figure 10c, Kaiser *et al.*, 2016, and references therein). This trend may be explained by a reduced sedimentation rate of marine organic matter, or by an enhanced delivery of light carbon into the oceans from continental weathering of organic-rich sediments during a short-term warming episode in the basal Tournaisian (Kaiser *et al.*, 2006; Marshall *et al.*, 2020). Decreasing carbon isotope values at Troip were reported as well by Bojar *et al.* (2013). However, their data and interpretations are related to uncorrect positions of the *kockeli* Zone (= Upper *praesulcata* Zone) and the DCB (see Bojar *et al.* 2013, Figure 3a). The DCB is fixed by the authors by the late FO of *Siphonodella sulcata* previously recorded by Ebner (1980a) from his Bed 8 (see Figure 9). The authors ignore that the DCB was correctly fixed by Ebner (1980a) in his Bed 4 with the FO of *Polygnathus purus subplanus* as an index fossil for the DCB (Kaiser *et al.*, 2009; Spalletta *et al.*, 2017). Bed 4 of Ebner is correlative with Bed 16 of Kaisers studies, according to thickness of the limestones and the FO's of *Polygnathus purus subplanus*, *Protognathodus kuehni* and *Siphonodella sulcata* (M5) in Bed 16. Thus, main parts of the *kockeli* Zone and the DCB of Bojar *et al.* (2013) can be regarded as younger, Tournaisian successions, and these biostratigraphic misinterpretations resulted in misinterpretations of geochemical proxies at Troip (see Zhang *et al.*, 2019, Figure 3). Further, the transgressive episode as interpreted from element geochemistry, and regarded as DCB-level by Bojar *et al.* (2013), is more likely the

expression of the lower Tournaisian transgression (*Gattendorfia* level) in the middle *sulcata/kuehni* Zone (*sulcata* Event after Kalvoda and Kukal, 1987).

## 4.2. SCENARIOS FOR A POTENTIAL GSSP POSITION AT THE DCB

### 4.2.1. AN EVENT-BASED SCENARIO

The event-based, early phase of the Hangenberg Crisis, recognized at Troip by bio-, litho- and chemostratigraphy (Section 4.1), is the initial main Hangenberg mass extinction Event among many marine organisms (*e.g.*, Kaiser *et al.*, 2016, Figure 3). It is correlative with the global deposition of the transgressive Rhenish HBS (= HBSE) and its equivalents elsewhere, and the main anoxic phase of the Hangenberg Crisis, indicated by the first significant positive peak in  $\delta^{13}\text{C}_{\text{carb}}$ ,  $\delta^{13}\text{C}_{\text{org}}$  (*e.g.*, Kaiser *et al.*, 2006). The global change from cephalopod limestones to black shales, or to high-condensed, thin-bedded limestones, has a worldwide correlation to numerous pelagic, hemipelagic and neritic settings in Asia, America and Europe (*e.g.*, Kaiser *et al.*, 2016; Becker *et al.*, 2016a, 2021; Aretz *et al.*, 2021, Kumpan *et al.*, 2021; Matyja *et al.*, 2021, Königshof *et al.*, 2021; Komatsu *et al.*, 2014; Qie *et al.*, 2021).

The HBSE (former middle part of the Middle *praesulcata* Zone) can be recognized biostratigraphically for example by miospores, and is dated as the LN miospore Zone (Sandberg *et al.*, 1972; Streel, 2009). The base of the LN miospore Zone provides correlation into the terrestrial realm and into macrofossil-free high-latitude settings. Furthermore, this level can be recognized by sequence stratigraphy and the drowning unconformity at the base of a maximum flooding interval (*e.g.*, Van Steenwinkel, 1993a, 1993b; Kaiser *et al.*, 2011). After summaries in Becker *et al.* (2016a), this episode of worldwide marine anoxia can be geochemically determined in various regions by organic geochemistry, inorganic carbon isotopes, magnetosusceptibility, element geochemistry, and by gamma ray spectroscopy (*e.g.*, Marynowski and Filipiak, 2007; Kaiser *et al.*, 2006; Cramer *et al.*,

2008; Kumpan *et al.*, 2014a, 2014b, 2015).

In siliciclastic-dominated successions, it is unclear which taxa range through the anoxic and regressive phase (HBSE, HSSE) due to fossil poor sediments, while in continuous pelagic limestone successions at Trolp and Grüne Schneid, the *ckI* (HBSE, HSSE) marks an important level. This level is characterized by main conodont extinctions, decreasing number of faunas, poor and stunted faunas, and major biofacies change. Alternative zonation concepts (Figure 12) do not consider the main Hangenberg extinction level instead, the level of the *ck* is the top part of the *ultimus* Zone (Spalletta *et al.*, 2021), or the base of the extended *kockeli* Zone (Corradini *et al.*, 2017).

The HBSE was previously proposed as a

natural DCB (Walliser, 1984). However, an event-based scenario is difficult to establish because 1. event-related successions are often accompanied by leakage, reworking, and impoverished faunas, and 2. hiatuses and  $C_{org}$  burial phases well before the HBSE, connected with the Dasberg, Etreoungt and Drewer Events as discussed in Section 4.1.3, resulted in the asynchronous onset of global black shale depositions and stratigraphic gaps (see figure 2a in Bojar *et al.*, 2013).

#### 4.2.2 THE BASE OF THE KOCKELI ZONE

The base of the *kockeli* Zone (= Upper *praesulcata* Zone), as well as the extended *kockeli* Zone (= *ckI*/*kockeli* Zone to base of the *bransoni* Zone, see Figure

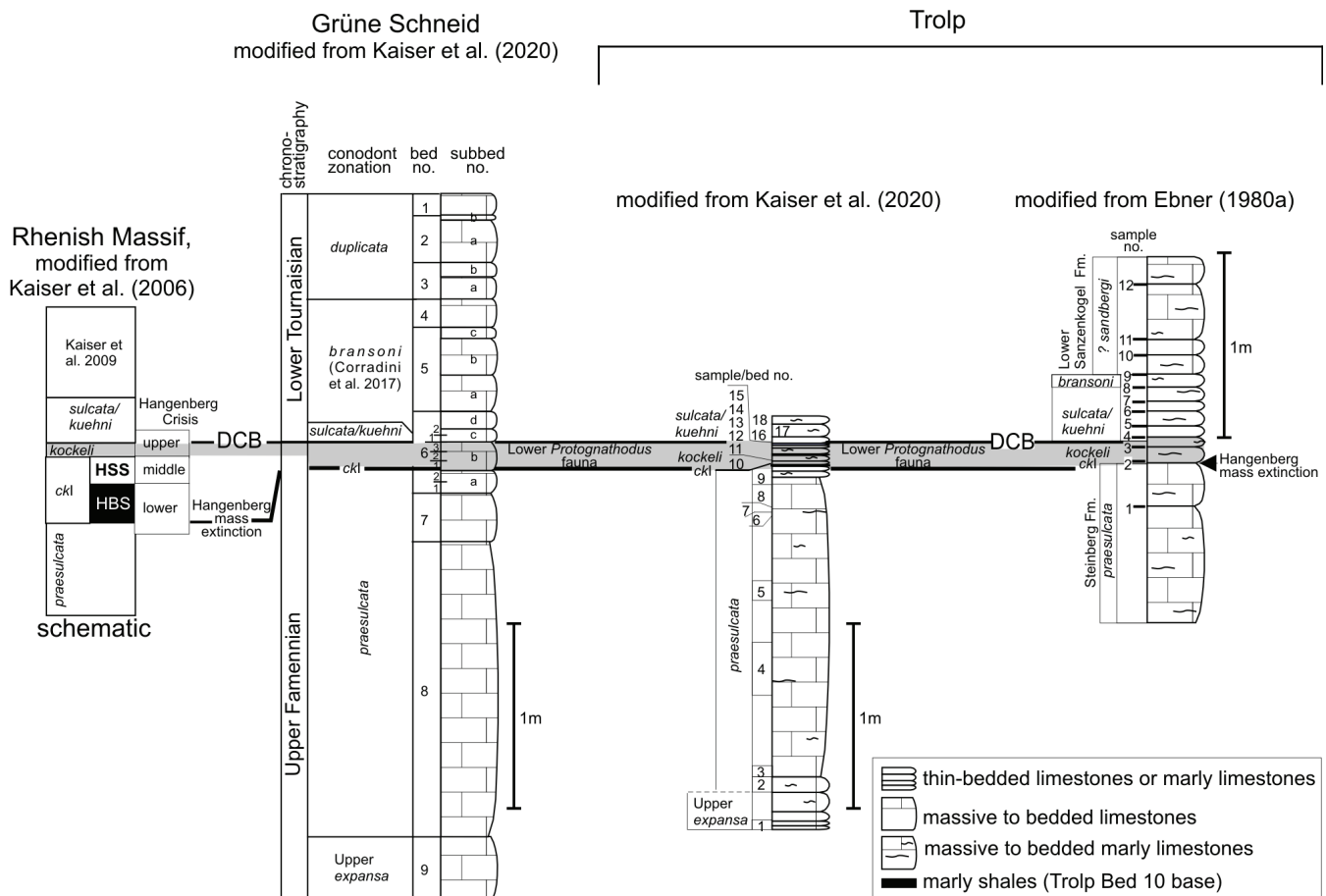


Figure 8 Stratigraphic correlation of Hangenberg Event successions at the DCB from Trolp (Graz Paleozoic), Grüne Schneid (Carnic Alps), and Rhenish Massif.

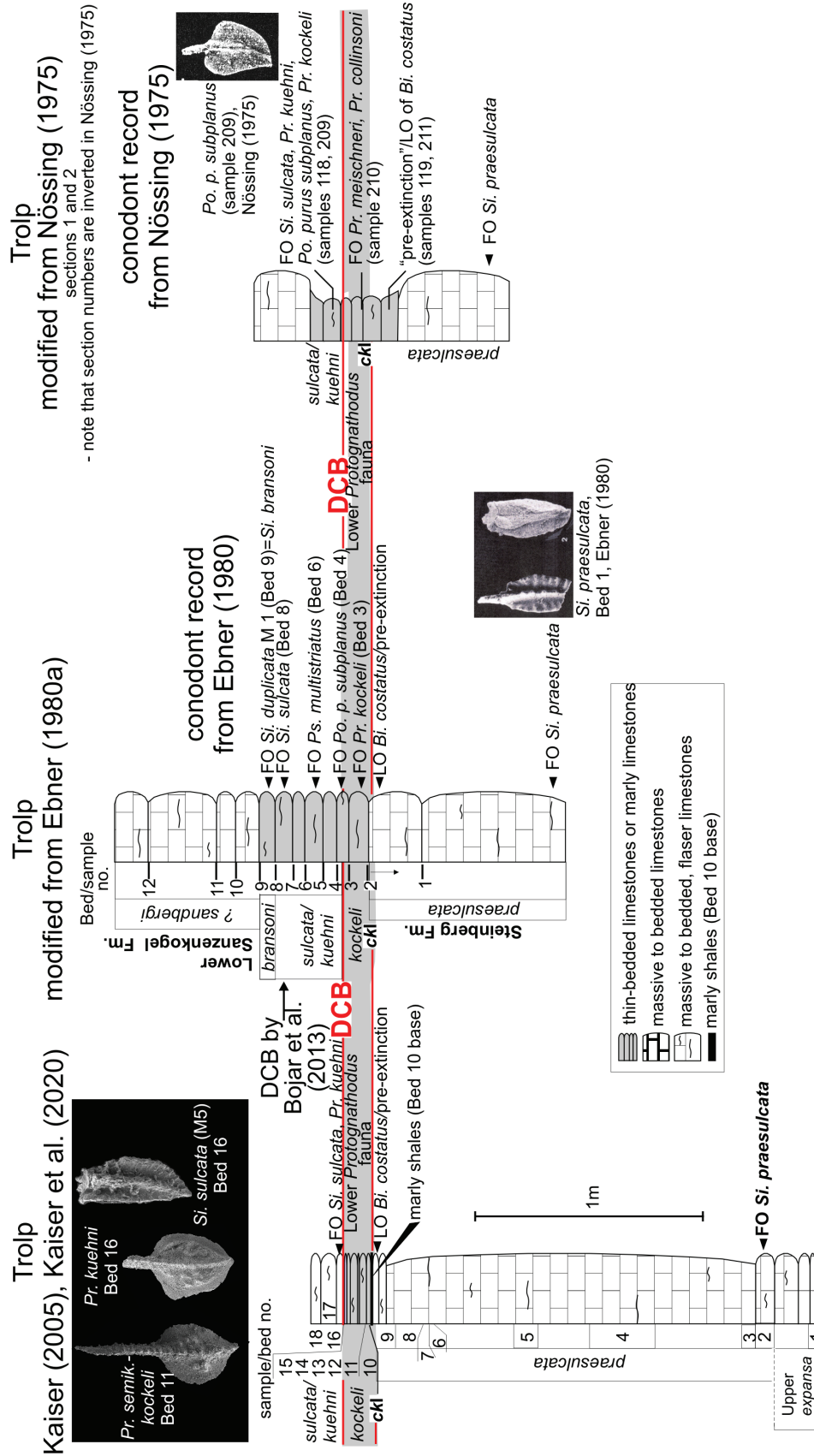


Figure 9 Conodont faunal correlation, and correlation of different but equivalent sample and bed numbers at Troip, after Nössing (1975), Ebner (1980), Kaiser (2005), Bojar et al. (2013) and Kaiser et al. (2020), and selected zonally diagnostic conodonts from Troip, ckl = *costatus-kockeli* Interregnum.

12) are not applicable as a potential DCB position as discussed as follows. The start of the *kockeli* Zone is probably within the upper LN miospore Zone, but the correlation into the terrestrial realm is difficult (e.g., Becker *et al.*, 2016a). The entry of *Protognathodus kockeli* in its former definition of Bischoff (1957) - related to diagnostic features as curvature (bending) and shape of the cup as well as platform ornamentation - is affected by the restricted distribution of *Protognathodus kockeli* in many regions and widespread facies-controlled, discontinuous conodont sampling across the base of the *kockeli* Zone due to carbonate-poor deposits and late entry of the index fossil with the first carbonates (see Kaiser *et al.*, 2020). The *kockeli* Zone is commonly not preserved due to the absence of *Protognathodus kockeli* at this level, as previously reported and additionally discovered recently in the Alborz Mts., Iran (e.g., Mighan, Chelcheli; Bahrami *et al.*, 2021; Königshof *et al.*, 2021; Parvizi *et al.*, 2021), Carnic Alps (e.g., Kronhofgraben, Plan di Zermula; Kaiser *et al.* 2009; Spalletta *et al.* 2021), Anti-Atlas (e.g., M'Karig, El Atrous; Kaiser *et al.*, 2011), Moravian Karst (Krtny; Kumpan *et al.*, 2021), or due to an impoverished *Polygnathus-Protognathodus* biofacies at the end of the Famennian (partly without *Protognathodus kockeli*), respectively, for example in the Rhenish Massif (e.g., Oese, Oberrödinghausen; Kaiser *et al.*, 2017; see Becker *et al.*, 2021), Holy Cross Mountains (Kowala; Matyja *et al.*, 2021), Illinois Basin, Indiana, USA (Over, 2021). Stratigraphic gaps are caused by the reduced formation of carbonates due to a global carbonate crisis during this time span, resulting from continuing Famennian sea-level fall (Babek *et al.*, 2016) as the longterm regression continued until the Carboniferous, and from continuing major global cooling (e.g., Isaacson *et al.*, 2008; Brezinski *et al.*, 2009; Kaiser *et al.*, 2022, *unpublished data*). The absence of the lower *Protognathodus* conodont assemblages (*Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus kockeli*) does not necessarily suggest stratigraphical gaps because erosional surfaces are missing (e.g., Rhenish Massif, Iran and elsewhere, see Becker *et al.*, 2021; Königshof *et al.*, 2021) but is

caused by biofacies changes and the deteriorating environment after the early and middle Hangenberg phase.

The entry of *Protognathodus kockeli* could be affected by local fossil reworking or leakage, for example in the Carnic Alps (Corradini *et al.*, 2017; see discussions in Kaiser *et al.*, 2020). Reworking and mixing of conodonts at this level is the result of the major eustatic fall associated with the HSSE and the Hangenberg glaciation in the middle Hangenberg phase, while leakage is the result of paleokarstic phenomena in the Graz Paleozoic and Carnic Alps (e.g., Ebner, 1980b; Schönlaub *et al.*, 1991).

Also, the uncertainties of the FAD of the “one-row” (*Protognathodus semikockeli* after Hartenfels *et al.*, 2022) and the “two-rows” morphotype of *Protognathodus kockeli* (*Protognathodus kockeli* s.str. after Hartenfels *et al.*, 2022), and the start of the *kockeli* Zone either during the middle (regressive episode of the HSSE, based on stable isotopes and the FO of *Protognathodus semikockeli* and *Protognathodus kockeli* at Tropl and Grüne Schneid, Kaiser *et al.*, 2022 unpublished data), or late phase of the Hangenberg Crisis (after the facies break related to the HSSE (Hartenfels *et al.*, 2022) have to be considered. The base of the *kockeli* Zone (former Upper *praesulcata* Zone) was previously established by the “two-rows” *Protognathodus kockeli* (Bischoff, 1957), while it was established by the “one-row” *Protognathodus kockeli* by Corradini *et al.* (2011). Based on conodont studies at Borkeweher (Rhenish Massif; Hartenfels *et al.*, 2022) from limestone successions overlying the regressive Hangenberg sandstone equivalent, the FO of *Protognathodus kockeli* s.str. (two-rows) within the evolutionary lineage of *Protognathodus semikockeli* (one-row) and *Protognathodus kockeli* s.str. in the post-regressive Hangenberg phase is proposed as future GSSP for the DCB.

#### 4.2.2.1. CONSIDERATIONS OF PROTOGNATHODUS

Taxonomic and biostratigraphic uncertainties within typical morphotypes of the early *Protognathodus* faunas are known for many years and were re-mentioned in Corradini *et al.* (2011), Kaiser *et*

*al.* (2019 unpublished, 2020) and Hartenfels *et al.* (2022). Originally, Bischoff (1957), Ziegler and Leuteritz (1970), and Ziegler (1969) only considered randomly the high spectrum of *Protognathodus* morphotypes. Generally, the high spectrum of intermediate morphotypes of *Protognathodus* species attests to the completeness of the stratigraphic record, as at Trolp, Grüne Schneid (Kaiser *et al.*, 2020), Puech de la Suque (France) and Borkewehr (Hartenfels *et al.*, 2022). However, taxonomic uncertainties are related to intraspecific variability concerning juvenile-adult and early-advanced forms, as well as intermediates between all species of the early *Protognathodus* fauna (Kaiser *et al.*, 2019, 2020; Hartenfels *et al.*, 2022). This was previously recognized in the early siphonodellids, too; the high spectrum of intermediate morphotypes of *Siphonodella praesulcata* and *Siphonodella sulcata* at La Serre É and elsewhere have been taxonomically re-evaluated (Kaiser and Corradini, 2011), and morphotype groups of both *Siphonodella praesulcata* and *Siphonodella sulcata* were established and widely used (Figure 12, see Section 4.2.3.1).

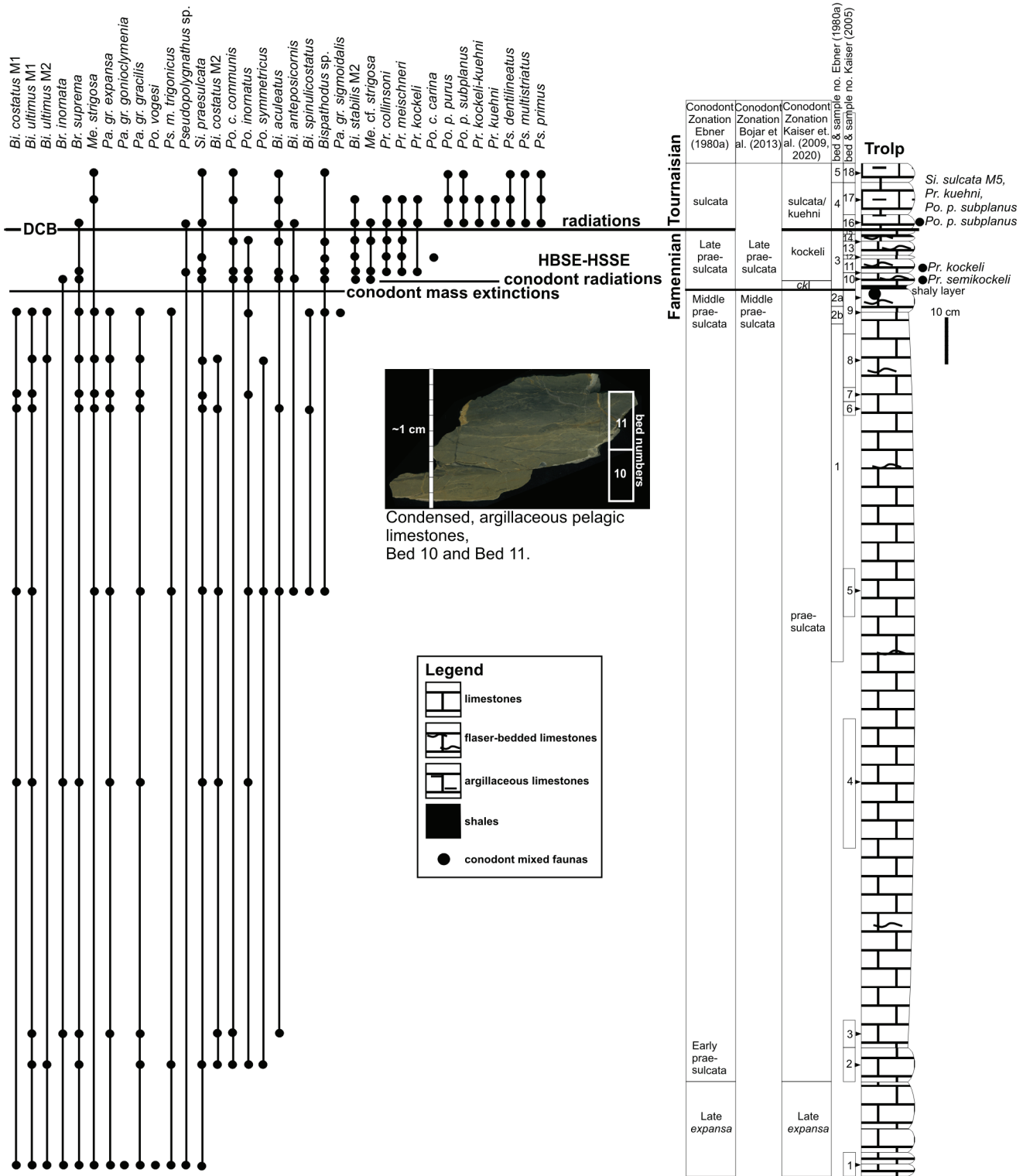
Diagnostic features of the *Protognathodus* fauna are related to the cup morphology (curvature and shape of the cup) and platform ornamentation (Bischoff, 1957; Ziegler and Leuteritz, 1970), and a phyletic lineage of the early protognathodids (Ziegler, 1973), based on the typical morphotypes and their transitional forms (intermediates), was suggested at Trolp and Grüne Schneid (Kaiser *et al.*, 2020), comparable to Puech de la Suque from Bed 10 overlying the HBS (Hartenfels *et al.*, 2022). Accordingly, a well-exposed succession of *Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli*, *Protognathodus kockeli* and *Protognathodus kuehni* could be recorded in these two regions. After Kaiser *et al.* (2020), the typical morphotype of *Protognathodus kuehni* at Trolp has at least on one side of the cup distinctive transverse ridges running radial from the platform edge to the carina, and the shape of the cup is wide and slightly asymmetric according to the original diagnosis of Ziegler and Leuteritz (1970).

Three new *Protognathodus* species are defined

based on the arrangement of the cup ornamentation which focused on the developing of row of nodes (Hartenfels *et al.*, 2022), without considering the shape and curvature of the cup, and the platform ornamentation 'scattered nodes' which are discussed by the authors, only (see also Corradini *et al.*, 2011, and Kaiser *et al.*, 2020). Transitional forms, transitional forms between *Protognathodus kockeli* and *Protognathodus kuehni*, but also morphotypes previously determined as *Protognathodus kuehni* are included in *Protognathodus kockeli* s.str. (Hartenfels *et al.*, 2022), and thus enlarged the spectrum of morphotypes to be determined as *Protognathodus kockeli*.

Atypical morphotypes of '*Protognathodus*' species with a high morphological variability, related to the shape of cup and the ornamentation of the platform, are recorded in the Graz Paleozoic (Figure 13), Carnic Alps, and other regions (see Kaiser *et al.*, 2020; Corradini *et al.*, 2011, Hartenfels *et al.*, 2022), and as stated in Hartenfels *et al.* (2022), "...atypical characteristics [within the genus *Protognathodus*],...result in a significant morphological complexity". At Trolp, within all species of *Protognathodus*, typical and also atypical morphotypes at Trolp are observed at the same stratigraphic levels in the uppermost Famennian and lower Tournaisian (Nössing, 1975; Kaiser *et al.*, 2020) which were partly included in the three new *Protognathodus* species (*Protognathodus semikockeli*, *Protognathodus kockeli* s.str., *Protognathodus kockeli*) recently established (Hartenfels *et al.*, 2022). For example, data at Trolp (Figure 13) indicate that beside specimens with a *Protognathodus*-like shape of cup, instead extremely narrow, atypical (instead of wide in comparison to holotypes) outlines of the shape of the cup in all species of the protognathodids are recorded from the *kockeli* to *bransonii* Zones (Kaiser *et al.*, 2020). Other morphologic characteristics are an extended posterior tip, a strong asymmetry due to an extended outer side of the cup, an enlarged outer margin which is commonly broken, or a strongly bent carina, in contrast to specimens with a straight carina.

Similar to the group of 'siphonodelloids',



**Figure 10a** Lithology and conodont biostratigraphy at Trolp, data from Kaiser (2005) and Kaiser *et al.* (2008, 2009). Updated detail records of *Protognathodus* at the DCB in Kaiser *et al.* (2020). *Si.* = *Siphonodella*, *Po.* = *Polygnathus*, *Ps.* = *Pseudopolygnathus*, *Bi.* = *Bispathodus*, *Br.* = *Branmehla*, *Pa.* = *Palmatolepis*, *Me.* = *Mehlina*, M = morphotype, *ckl* = *costatus-kockeli* Interregnum. Black circle = mixed conodont faunas. According to the recent taxonomic re-evaluation by Hartenfels *et al.* (2022), *Protognathodus semikockeli* and *Protognathodus kockeli* s.str. occurs at Trolp in the thin-bedded limestones.

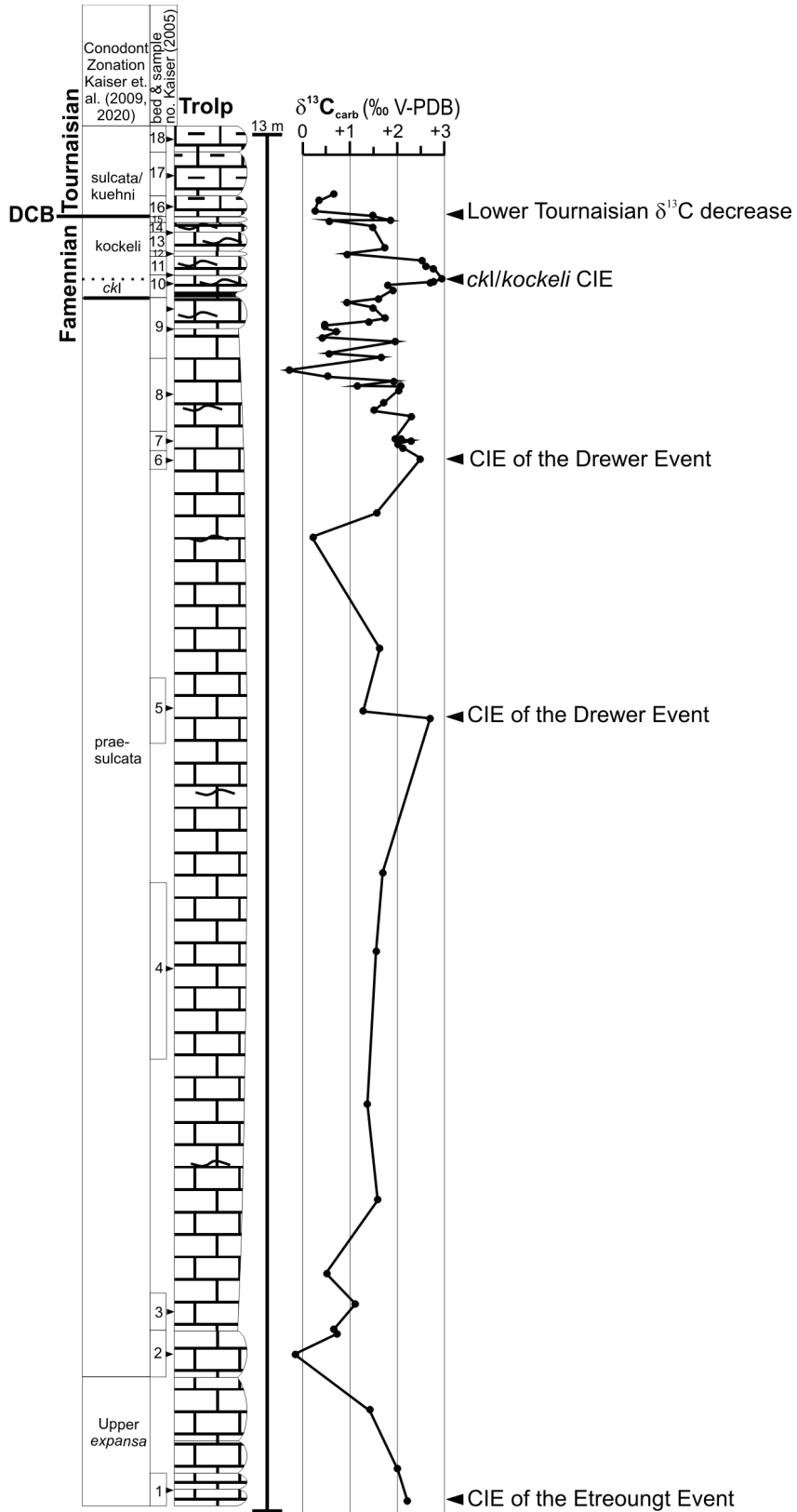
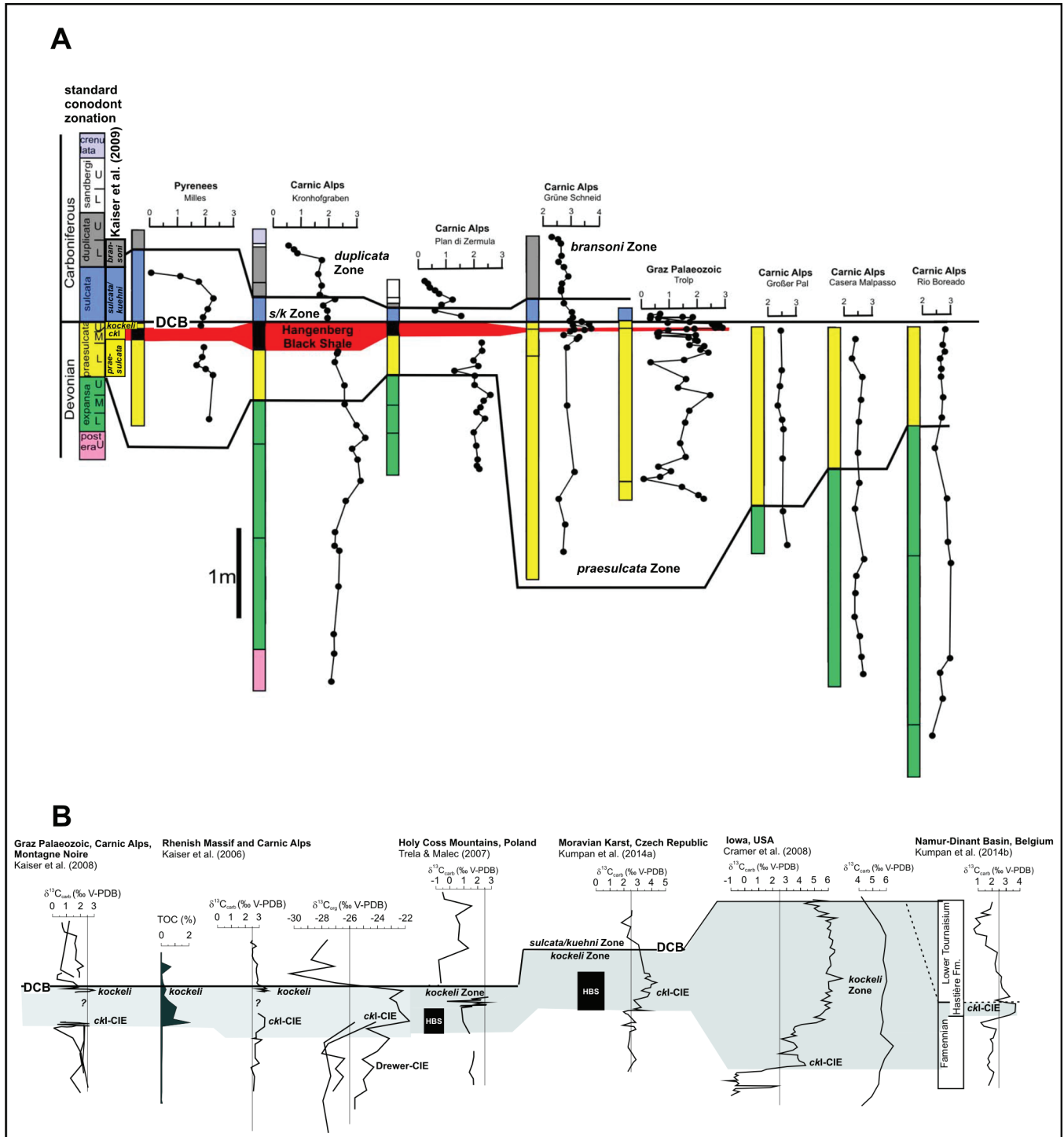


Figure 10b Geochemistry of carbon isotopes at Trolp, data from Kaiser (2005) and Kaiser *et al.* (2008), *ckl* = *costatus-kockeli* Interregnum, CIE=carbon isotope excursion.



**Figure 10c** Geochemical ( $\delta^{13}C_{carb}$ ) and biostratigraphic correlation of different DCB successions from southern and middle Europe and North America with the western Graz Paleozoic (Tropf), indicating same trends from the Famennian to Tournaisian in carbon isotopes. A. modified from Kaiser (2005), B. modified from Kaiser *et al.* (2016).

which has morphological features of *Siphonodella*, *Polygnathus* and *Pseudopolygnathus*, and were established previously (Becker *et al.*, 2013), the morphological complex group of *Protognathodus* faunas with atypical diagnostic features could be regarded as *Protognathodus*-like ‘protognathoids’, and can have morphological features of the late *Protognathodus*, of *Bispathodus* and of *Gnathodus* regarding platform ornamentation and shape of the cup as discussed below. Also, homeomorphic morphological features could be problematic because the early protognathodids can have overlapping ranges with the late *Protognathodus* (e.g., *Protognathodus* *praedelicatus*, see Habibi *et al.*, 2008; see discussion in Hartenfels *et al.*, 2022), with *Bispathodus* (e.g., *Bispathodus* *stabilis*, Ziegler 1973, 1974) and with *Gnathodus* (see below). These atypical faunas (‘protognathoids’) probably originated during the early phase of the Hangenberg Crisis (Drewer Event) in the *praesulcata* Zone, with ?*Protognathodus* *meischneri* and ?*Protognathodus* *collinsoni*, and evolved probably during the anoxic (HBSE) or the regressive phase (HSSE) of the Hangenberg Event, with ?*Protognathodus* *semikockeli* and ?*Protognathodus* *kockeli* (see Figure 13), and spreaded around the DCB and in the lower Tournaisian. Their occurrence could be related to an opportunistic lifestyle during major environmental changes related to the Hangenberg Event, which can be regarded as “bottleneck” for the evolution of the Idiognathodontidae to which *Protognathodus* belong. Because protognathoids spread in the *kockeli* Zone, but mainly in the *sulcata/kuehni* Zone, or probably even later (Figure 14; see Corradini *et al.*, 2011; Kaiser *et al.*, 2020), the extended *kockeli* Zone (base of the former *kockeli* Zone to the base of the *bransonii* Zone) is not applicable due to the spread of these still unknown faunas regarding their stratigraphic and distributional pattern, and also due to their unknown ancestor and descendent relationship.

At Trolp, a stratigraphically leaked younger fauna is recorded by single specimens of *Gnathodus* at the same stratigraphic level as the early *Protognathodus* fauna, and leaked *Gnathodus* were found in the *sulcata/kuehni* Zone, or in older (*praesulcata*

Zone) levels. Mixed faunas and homeomorphy (see below) within the Idiognathodontidae family (*Gnathodus* and *Protognathodus*) could produce a bias of the faunal record in the lower Tournaisian, especially in 1) condensed successions, 2) in stunted or juvenile-dominated assemblages, 3) in Lower Carboniferous, siliciclastic-dominated successions with fossil-poor limestones and scarce conodont faunas, without associated other diagnostic faunas and/or co-occurrence with ‘homeomorphic’ faunas (e.g., Kaiser *et al.*, 2020; see Königshof *et al.*, 2021 and Parvizi *et al.*, 2021). At Shahmirzad, (Alborz Mts., northern Iran), *Protognathodus* *kockeli* even occurs at same level as *Gnathodus* *cuneiformis* and *Protognathodus* *praedelicatus* in the upper Tournaisian Lower *typicus* Zone to *anchoralis-latus* Zone (Habibi *et al.*, 2008, see discussion in Hartenfels *et al.*, 2022). In contrast, leaked faunas among the late (*Siphonodella* *bransonii*, *Siphonodella* *duplicata*, *Siphonodella* *quadruplicata*) siphonodellids are generally easy to distinguish from the early (*Siphonodella* *praesulcata*, *Siphonodella* *sulcata*) siphonodellids, as at Trolp in mixed faunal assemblages around the thin-bedded limestone successions (Figure 10a).

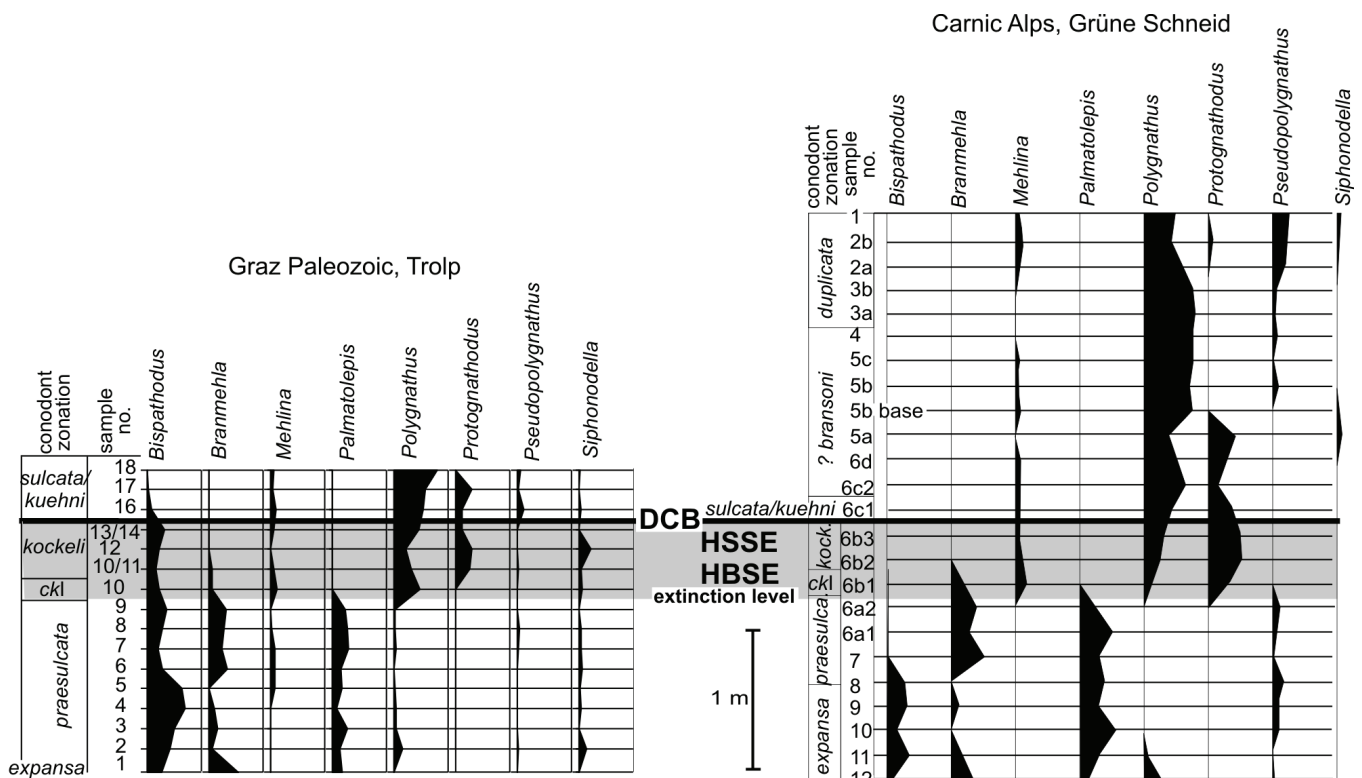
Lane *et al.* (1980) provided an extended discussion of the differences between *Gnathodus* and *Protognathodus* and the differences in cup shape and cup terminations between early and late *Protognathodus*, with diagnoses of both genera. However similar diagnostic features between the early *Protognathodus* fauna, the late *Protognathodus* fauna ([*Protognathodus* *cordiformis*], *Protognathodus* *praedelicatus*, see also discussion in Hartenfels *et al.*, 2022), and *Gnathodus* (Ziegler, 1973; Lane *et al.*, 1980; Dzik, 1997, 2006), and their overlapping ranges in the lower, middle or upper Tournaisian must be considered.

Examples of atypical morphotypes or homeomorphy as discussed as follows are partly described in synonymies for *Protognathodus* *kockeli* (Hartenfels *et al.*, 2022). Atypical ‘*Protognathodus*’ specimens can have a cup-morphology, which is similar to the late protognathodids regarding the termination of the margin and the wideness of the cup, and also confusion with *Gnathodus* may be related to the platform ornamentation of *Protognathodus* *kockeli*,

where, for example, nodes as ridge-forming row of “denticles”, partly with an adcarinal through, are developed (e.g., Corradini *et al.*, 2011, plate 1, figure 18; Kaiser *et al.*, 2020, plate 2, figure 1), or incipient parapet-like morphologies on one side of the platform and a strong asymmetry, e.g., in Sardinia, Corradini *et al.* (2011, pl. 1, Figure 15), at Troip, Kaiser *et al.* (2020; plate 3, figure 19, 22), in Morocco, Lalla Mimouna, Becker *et al.* (2013, plate 3, figure 9, with an unusual wide posterior carina, similar to specimen from Oklahoma figured by Over, 1992, see below), or in China, Wang and Yin (1988; Figure 15).

*Protognathodus* specimens from the DCB transition were previously determined as *Gnathodus* in the Rhenish Massif (e.g., Luppold *et al.*, 1984, pl. 6, Figure 2) or as *Protognathodus praedelicatus* in the Carnic Alps or Montagne Noire (Flajs and Feist, 1988; Schönlaub *et al.*, 1988). Subjective determinations

of *Protognathodus* species from North Africa (Korn *et al.*, 2004), and from China, are evident (Wang and Yin, 1988: pl. 22, Figure 1-19; and compare synonymy list in Corradini *et al.*, 2011, and discussions therein, p. 25). Over (1992) described *Protognathodus* n. sp. from the *kockeli* Zone (= Upper *praesulcata* Zone) from North-America, which can be distinguished after the author from *Gnathodus* by its relatively symmetrical platform and lack of the offset of the outer and inner anterior platform margins. Specimens of *Protognathodus kockeli* from this region (Oklahoma, Woodford Shale) have a wide posterior carina, similar to that figured specimen from Morocco (see above, Becker *et al.*, 2013). This species is regarded by Over (1992) as an intermediate between *Protognathodus kockeli* and *Protognathodus* sp. B. Over (1992) suggested that *Protognathodus* sp. B may represent the ancestor of *Gnathodus punctatus* (with FAD in the late Kinder-



**Figure 11** Conodont biofacies change and mass extinctions at the DCB from the Graz Paleozoic (Troip) and Carnic Alps (Grüne Schneid), data from Kaiser (2005, 2007), *ckl* = *costatus-kockeli* Interregnum.

hookian) which has a broadly expanded posterior carina. Similar specimens have been illustrated from La Serre, Montagne Noire (Flajs and Feist, 1988, pl. 9, Figure 9, 10), and fused nodes on the platform (plate 9, figure 10) could be transitional to *Protognathodus praedelicatus* but was determined as *Protognathodus kockeli* after Corradini *et al.* (2011).

Furthermore, juvenile specimens of *Protognathodus meischneri* and *Bispathodus stabilis* are difficult to distinguish (Ziegler, 1973). Both have overlapping ranges in the ?upper/uppermost Famennian and lower Tournaisian, and confusions with stunted or juvenile ?*Protognathodus meischneri* and *Bispathodus stabilis* occur at Trolp and elsewhere (Kaiser *et al.*, 2020; see also Corradini *et al.*, 2016). Ziegler *et al.* (1974) have demonstrated the ancestry of *Protognathodus*; *Bispathodus stabilis* Morphotype 2, which has an expanded basal cavity extending to the posterior tip of the blade (end of the platform), was shown to be the ancestor of the *Protognathodus* lineage by the authors, illustrated on plate 3, Figure 2. *Bispathodus stabilis* Morphotype 2 lacks only the wider expansion of the cup to distinguish it from *Protognathodus meischneri*, and *Protognathodus meischneri* has a distinctly flat basal cavity.

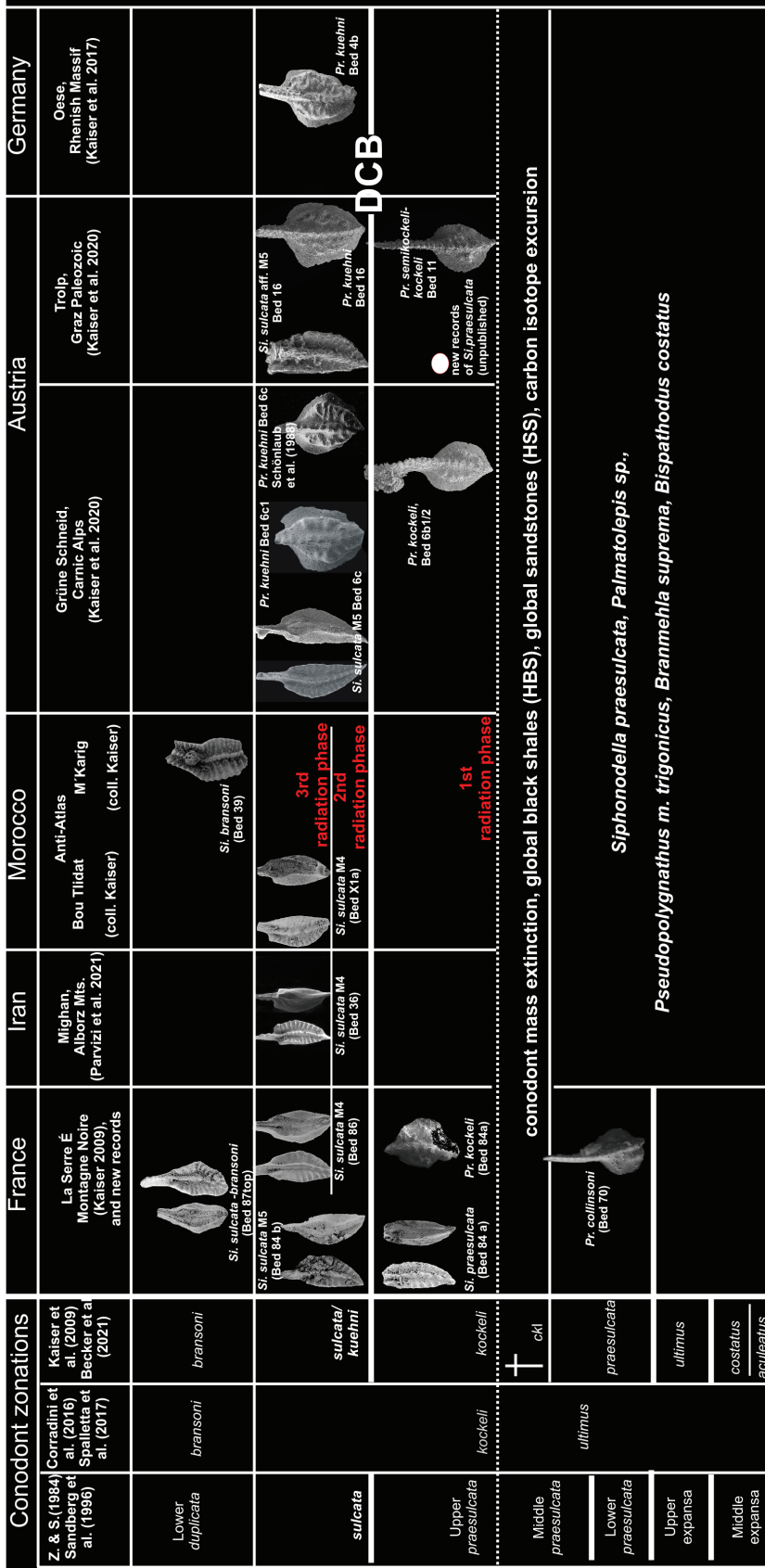
However, as stated in Kaiser *et al.*, (2020), the shape of the cup of adult specimens of ?*Protognathodus meischneri* at Trolp (Figure 13, narrow shape of cup) can be similar to adult specimens of *Bispathodus stabilis stabilis* (see Hartenfels, 2011), and thus confusions in the determinations of both species exist, also in regard to the ornamentation of the platform which can be similar between *Protognathodus collinsoni* and *Bispathodus stabilis bituberculatus* and *Bispathodus aculeatus*, or between *Bispathodus stabilis stabilis* and *Protognathodus meischneri* (Hartenfels, 2011; Hartenfels and Becker, 2016; see Hartenfels, 2011, plate 31), *e.g.*, Matyja *et al.*, (2021, Figure 5A.d: aff. *Protognathodus meischneri*); Königshof *et al.*, (2021); Becker *et al.*, (2013, plate 3, Figure 5: *Protognathodus meischneri* [?with initial nodes on the platform]); Matyja *et al.*, (2021, with a doubtful *Protognathodus*, Figure 7a-c [similar to her *Bispathodus aculeatus*]); Kumpan *et al.*, (2021, *Protognathodus cf. meischneri*, Figure 5.7 [with

an initial node on the platform]). The doubtful specimens of *Protognathodus meischneri* or *Protognathodus collinsoni*, partly discussed in Hartenfels *et al.*, (2022), occur in pre-extinction levels (*praesulcata* Zone) or in the Hangenberg Crisis level as single specimens in hemipelagic or neritic settings at La Serre É, Montagne Noire (Flajs and Feist, 1988; Kaiser, 2005, 2009; Feist *et al.*, 2021), Lesní Lom, Moravian Karst (Kumpan *et al.*, 2021), Mighan and Chelcheli, Alborz Mts., Iran (Parvizi *et al.*, 2021; Bahrami *et al.*, 2021; Königshof *et al.*, 2021), Lalla Mimouna, Anti-Atlas, Morocco (Becker *et al.*, 2013), Kowala, Holy Cross Mountains, Poland (Matyja *et al.* 2021), Grüne Schneid, Carnic Alps (Corradini *et al.*, 2017; Kaiser *et al.*, 2020). Also, the occurrence of questionable *Protognathodus meischneri* and *Protognathodus collinsoni* in pre-Hangenberg Event levels from different regions (*e.g.* at La Serre E) must be considered in respect to their FAD and ancestor relationship to the new species *Protognathodus semikockeli* recently established (see also discussion in Hartenfels *et al.*, 2022).

In summary, beside the diagnostic features of the shape and curvature of the cup of the *Protognathodus* fauna (see Figure 13), morphological complexity of the ornamentation of the platform may resulted in subjective determinations. Therefore, these morphological complexities in shape of the cup and ornamentation of the platform ('protognathoids') should be focused on in future studies, comparable to the 'siphonodellids' which are discriminated from the homeomorphic siphonodellids.

#### 4.2.3 THE DEVONIAN-CARBONIFEROUS BOUNDARY AND CURRENT INDICATORS IN THE SULCATA/KUEHNI ZONE

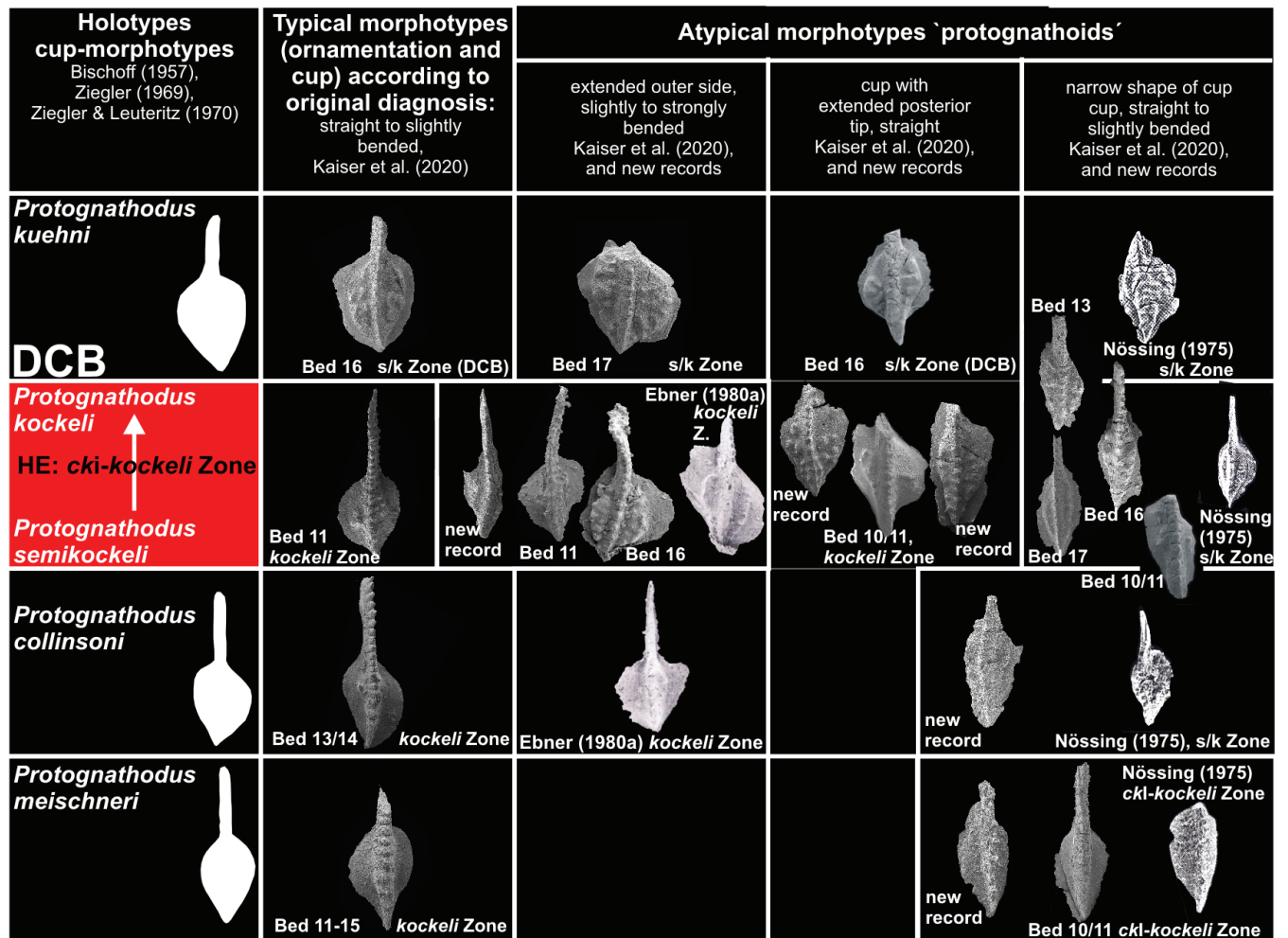
At the base of the *sulcata/kuehni* Zone, the 2nd faunal recovery among conodonts (Figure 14) is connected with an increasing abundance of pseudopolygnathids and siphonodellids (*e.g.*, Kaiser, 2005; Kaiser *et al.*, 2017), and with a change from poor, rare and stunted conodont faunas in the *ckI/kockeli* Zone (1st radiation phase) to more abundant and normal-size faunas indicating the post-extinction/



**Figure 12** Correlation of significant DCB index conodonts - *Siphonodella sulcata* Morphotype 5 and *Siphonodella sulcata* Morphotype 4, *Protognathodus kuehni* - recently discovered from different paleogeographical regions. s/k Zone = *sulcata/kuehni* Zone, *ckl* = *costatus-koekelli* Interregnum. The FAD of *Siphonodella sulcata* M5 is congruent to that of *Protognathodus kuehni*. Note that specimens from La Serre É shown are the earliest *Siphonodella sulcata*-morphotypes to occur at the base of the Tourmaisien in Bed 84, while specimens shown on Figure 1 in Aretz and Corradini (2021), are recorded by Kaiser (2009) from younger stratigraphic levels at La Serre É from Bed 86 (left specimen, advanced *Siphonodella sulcata*) and from Bed 85 (advanced *Siphonodella praesulcata*).

post-crisis level. The 1st and 2nd recovery phases (see Figure 12) each coincide with global post-extinction sea-level rises and deepening episodes, as interpreted, for example, by microfacies at Troip (Kaiser, 2005) and Grüne Schneid (Kaiser, 2007). At Troip and Grüne Schneid, it is characterized by more fine-grained micritic limestones – mud- and wackestones - when compared to under- and overlying beds which are pack- and floatstones. The

2nd faunal recovery phase are about time-equivalent to the return to full carbonate depositional conditions and continuing in the lower Tournaian (middle *sulcata/kuehni* Zone, *Gattendorfia* level, 3rd faunal recovery phase). The onset of a distinctive decrease of  $\delta^{13}C_{carb}$  values started in the *sulcata/kuehni* Zone at Troip with the onset of the 2nd radiation phase, and can be correlated to numerous other settings in European, Asian and



**Figure 13** Conodont faunas from the DCB at Troip. Typical and atypical ('protognathoids') morphotypes of *Protognathodus* related to the shape of the cup and platform ornamentation. A high morphological complexity within *Protognathodus semikockeli* and *Protognathodus kockeli* (for diagnosis see Hartenfels et al. 2022) is evident from the Hangenberg Event (cki-kockeli Zone). The 'three-rows' and 'four-rows' specimen related to the platform ornamentation are determined as *Protognathodus kockeli* (Hartenfels et al. 2022; Corradini et al. 2024), comparable to the record herein from Bed 16. Specimens from Beds 10, 11 (cki, base of kockeli Zone) belong to *Protognathodus semikockeli* ('one-row') or *Protognathodus kockeli* s.str. ('two-rows') after Hartenfels et al. (2022). The advanced *Protognathodus kockeli* (4-rows-specimen from Puech de la Suque, Bed 16, collection S.Kaiser) figured in Corradini et al. (2024) originated from the *bransoni* or even *duplicata* Zone (compare Corradini et al. 2024, p. 24, 25).

North-American regions (see Section 4.1.3).

Between the 1st (base of *kockeli* Zone) and 2nd recovery phases, a short regressive episode (DCB regression) corresponds to significant survivor extinctions in ammonoids, brachiopods, trilobites and foraminifers at the end of the *kockeli* Zone to a roughly contemporaneous terrestrial extinction, and to abruptly decreasing carbon isotope values globally recorded (Figure 14; e.g., Kaiser *et al.*, 2006, 2016). The level of the DCB regression enables a correlation of pelagic (conodonts, ammonoids) with shallow-water environments (brachiopods, corals, see review in Kaiser *et al.*, 2016), and can also be correlated into the terrestrial realm by the LN/VI miospore Zone boundary (e.g., Clausen *et al.*, 1994). The DCB regression is indicated at Trolp by microfacies change from micritic limestones to bioclastic pack- to floatstones reported by Kaiser (2005) and Bojar *et al.* (2013, Figure 3c) although the latter authors assigned this level to the Upper *praesulcata* Zone based on the incomplete conodont record as mentioned in Section 4.1.3.

The different morphotypes of *Siphonodella sulcata*, especially morphotypes 4 and 5, are globally widespread (see Section 4.2.3.1), and *Protognathodus kuehni* has a widespread geographic distribution which is reported in numerous studies from the *sulcata/kuehni* Zone (Corradini *et al.*, 2011; Kaiset *et al.*, 2020; see discussion in Hartenfels 2022). However, this taxon can have, as well as *Protognathodus kockeli*, a late entry, for example at the GSSP at La Serre É (e.g., Flajs and Feist, 1988; Feist *et al.*, 2021). Therefore, the *sulcata/kuehni* Zone has a high global correlation potential and the FO's of *Siphonodella sulcata* and *Protognathodus kuehni* can be correlated to many other regions globally (see Kaiser and Corradini, 2011; Corradini *et al.*, 2011; and extended references in Kaiser *et al.*, 2020; Königshof *et al.*, 2021; Qie *et al.*, 2021; Corradini *et al.*, 2021; Spalletta *et al.*, 2021; Kumpan *et al.*, 2021; Matyja *et al.*, 2021; Becker *et al.*, 2021; Feist *et al.*, 2021; Komatsu *et al.*, 2014). While the extended *kockeli* Zone leaves the event-related successions and the DCB undivided, as recently

used for example at Grüne Schneid, La Serre E, and Milles (Feist *et al.*, 2021, Spalletta *et al.*, 2021; Aretz *et al.*, 2021; Figure 12), a precise conodont biozonation at the DCB can be applied by the use of *Siphonodella sulcata* M5 and *Protognathodus kuehni*.

At Trolp, the typical morphotype of *Protognathodus kuehni* and *Siphonodella sulcata* M5 first occur at same stratigraphic levels (Figure 10a, 12). Also, *Siphonodella sulcata* ?M4 was figured in Nössing (1975) in the basal *sulcata/kuehni* Zone. The *sulcata/kuehni* Zone can thus be established in the Graz Paleozoic, and at correlative levels in the Carnic Alps and Moravian Karst, based on the co-occurrence of both *Siphonodella sulcata* M5 and *Protognathodus kuehni* (Figure 12; Kalvoda *et al.*, 2015; Kaiser *et al.*, 2020; Kumpan *et al.*, 2021). Sampling of continuous limestone successions there supports the coeval FAD of both taxa at the base of the Tournaisian. After Kaiser *et al.* (2020), this supports the use of a joint biozone in event-related successions, because their co-occurrences are evidenced from these three different regions; the utility of a joint biozone enhances the possibility of using either the biostratigraphic index *Siphonodella sulcata* or *Protognathodus kuehni*, if one or the other has a late entry or is as yet unrecorded in a region due to facies-controlled rare or diachronous occurrences (e.g., Kaiser *et al.*, 2009; Corradini *et al.*, 2011; Spalletta *et al.*, 2017).

#### 4.2.3.1. CONSIDERATIONS OF SIPHONODELLA

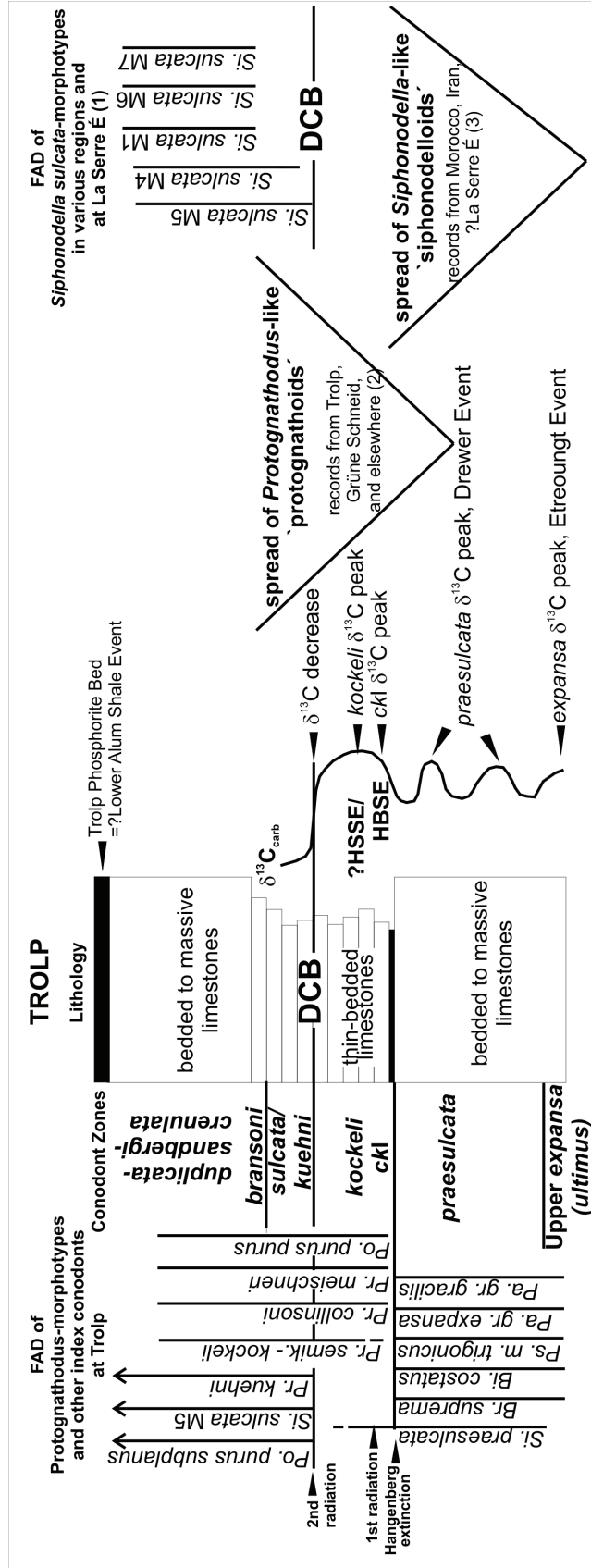
Within siphonodellids, the abundance of intermediate forms attests to the completeness of the stratigraphic record as at La Serre É (Figure 12; between *Siphonodella praesulcata*, *Siphonodella sulcata* M4 and M5, and advanced siphonodellids); the differences between *Siphonodella praesulcata* and *Siphonodella sulcata* are stated in Sandberg *et al.* (1972). However, the occurrence of intermediate forms may have resulted in subjective determination of the first occurrence of *Siphonodella sulcata* (e.g., Ziegler and Sandberg, 1996; Kaiser, 2009; Becker *et al.*, 2016a), but *Siphonodella*-morphotype groups can be clearly distinguished and recorded

the spectrum of morphotypes and intermediates, which spread in the lower Tournaisian (Kaiser and Corradini, 2011). These groups are used to distinguish between *Siphonodella praesulcata* (M2, M3) and *Siphonodella sulcata* (M1, M4-M7), and also between early (M4, M5) and advanced forms (M1, M6, M7) of *Siphonodella sulcata*, the latter are recorded especially in younger Tournaisian successions at La Serre É (Kaiser and Corradini, 2011). The morphotype groups are based mainly on the curvature of the carina, a feature which had been used for the current GSSP level at La Serre É (Flajs and Feist, 1988, see also Sandberg *et al.*, 1972), as well as shape of the platform, a pseudokeel which contains a pit (basal cavity), and the ornamentation. *Siphonodella sulcata* Morphotype 4 already has slight constrictions anteriorly but is otherwise similar to *Siphonodella sulcata* M5. The *Siphonodella sulcata* M1, M6 and M7 are representatives of advanced forms, and are intermediates between *Siphonodella sulcata* and *Siphonodella bransoni* (former *Siphonodella duplicata* M1) because they are strongly bended and have constrictions anteriorly.

*Siphonodella sulcata* M4 and M5 are easy to recognize, most widespread, and common in North America, Asia, and Europe (Figure 12, Kaiser and Corradini, 2011, figure 7; see Kaiser *et al.* 2020). The globally most widespread *Siphonodella sulcata* M5 is the earliest morphotype which occur at the base of the Tournaisian, for example at Troip, Grüne Schneid (Kaiser *et al.*, 2020), La Serre É (Kaiser, 2009; Kaiser and Corradini, 2011; Feist *et al.*, 2021), Lalla Mimouna (Anti-Atlas, Morocco, Becker *et al.*, 2013), Lesni Lom (Moravian Karst, Czech Republic; Kalvoda *et al.*, 2015), Kowala (together with *Protognathodus kuehni* and *Polygnathus purus subplanus*, Holy Cross Mountains, Poland; Malec, 2014; Matyja *et al.*, 2021), Borkeweher (Rhenish Massif, Hartenfels *et al.*, 2022). *Siphonodella sulcata* M4 was found in lower Tournaisian successions, for example at Bou Tlidat (Maidier, Anti-Atlas, Morocco; collection S.I. Kaiser; see Kaiser *et al.*, 2011; Becker and Kaiser, 2023), Mighan (Alborz Mts. northern Iran, Parvizi *et al.*, 2021) and at Shahmirzad (Alborz Mts.; Habibi

*et al.*, 2008). *Siphonodella sulcata* M4 evolved from *Siphonodella sulcata* M5 based on the record at La Serre É, and represents most probably the marker for the 3rd radiation phase in the *sulcata/kuehni* Zone at the *Gattendorfia* level (*sulcata* Event). *Siphonodella bransoni* is also recorded from successions at M'Karig, Tafilalt, Anti-Atlas (collection S.I. Kaiser; see Kaiser *et al.*, 2011; Becker and Kaiser, 2023). For the occurrence of *Siphonodella praesulcata* in neritic settings in Morocco, Iran, China, Vietnam, see Kaiser (2005), Kaiser *et al.* (2011), Becker *et al.* (2013), Komatsu *et al.* (2014), Parvizi *et al.* (2021), Bahrami *et al.* (2021), Qie *et al.* (2021). In shallow-water settings at Gedongguan, Muhua, Nanbiancun and Dapoushang (South China), *Siphonodella sulcata* occurs, also in association with *Protognathodus kuehni*, at the base of the Tournaisian (Qie *et al.*, 2021), and *Siphonodella sulcata* is also reported from shallow-water settings for example from northeastern Vietnam (Komatsu *et al.*, 2014). In summary, siphonodellids also occur in neritic successions, and the *sulcata/kuehni* Zone can be established in pelagic, hemipelagic and neritic settings based on *Siphonodella sulcata* and/or *Protognathodus kuehni*. However, an evaluation of various records from different paleogeographical settings of the earliest Morphotype 5 and Morphotype 4 of *Siphonodella sulcata* at the base of the Tournaisian is needed (*e.g.*, Becker *et al.*, 1984; Qie *et al.*, 2021; Komatsu *et al.*, 2014; Matyja *et al.*, 2021, Hartenfels *et al.*, 2022; see also Aretz *et al.*, 2021, and references therein).

*Siphonodella sulcata* M4 and M5 can be clearly distinguished from *Siphonodella praesulcata*, from the late siphonodellids (see Section 4.2.2.1), and from *Siphonodella*-like 'siphonodelloids'; the latter have overlapping ranges with *Siphonodella praesulcata* mainly in the *praesulcata* Zone, as documented *e.g.*, in Becker *et al.* (2013). The 'siphonodelloids' originated and evolved in pre-Hangenberg event levels in the *expansa* Zone probably connected with enhanced C<sub>org</sub> burial phases and/or shallowing phases during the Epinette and Etreoungt Events (for environmental changes during the events see Kaiser *et al.*, 2008), and spread in the *praesulcata*



**Figure 14** Schematic overview of bio- litho- and chemostratigraphic markers at Trolp in the western Graz Paleozoic. (1) Kaiser and Corradini (2011, and references), (2) Kaiser et al. (2020, and references), (3) Becker et al. (2013), Parvizi et al. (2021), S.I. Kaiser (2019, unpublished data). *Si.* = *Siphonodella*, *Po.* = *Polygnathus*, *Ps.* = *Pseudopolygnathus*, *Bi.* = *Bispathodus*, *Br.* = *Branmehla*, *Pa.* = *Palmatolepis*, *Me* = *Mehlina*, *M* = *morphotype*, *ckl* = *costatus-kockeli* Interregnum.

Zone; its occurrence is related to biotic opportunism comparable with the spread of 'protognathoids' during the Hangenberg Event (Figure 14). 'Siphonodelloids' are most widespread in hemipelagic or shallow-water settings, for example in Morocco (*e.g.*, Becker *et al.*, 2013), Iran (*e.g.*, Parvizi *et al.*, 2021), and probably at La Serre É (Figure 14).

## 5. Summary

Characteristic lithologic, chemo- and biostratigraphic markers at the DCB in the western Graz Paleozoic are well displayed at the famous Tropl section in the Steinberg area. Geochemical and lithologic characteristics, conodont extinctions, the end of mass extinctions and radiations, and the coeval FO of index conodonts are recognized at Tropl. These characteristic markers can be easily correlated globally, and afford the possibility for correlations into the pelagic, hemipelagic and neritic realms.

A hiatus is unlikely at Tropl. This is supported by the complete record of conodont zones during the Hangenberg Crisis, the abundance of intermediate forms which attests to the completeness of the conodont stratigraphic record, and by the absence of an erosional (sedimentary) gap. Conodonts are abundant and enable a high-resolution conodont stratigraphy, and radiation and extinction events are well recorded. The main Hangenberg extinction level in the former Middle *praesulcata* Zone (*costatus-kockeli* Interregnum = *ckI*) is easily recognized at the base of thin-bedded limestones, and is accompanied by a succeeding major conodont biofacies change and an impoverished and stunted *Polygnathus-Protognathodus* fauna. The index conodonts *Siphonodella praesulcata* and *Siphonodella sulcata* occur in the DCB beds at Tropl together with the early *Protognathodus* fauna (*Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli*, *Protognathodus kockeli*, *Protognathodus kuehni*) in the continuous, thinly bedded limestone succession.

The DCB at Tropl is recognized by the first

occurrence of both *Protognathodus kuehni* and *Siphonodella sulcata* M5 at same stratigraphic level and document the start of the *sulcata/kuehni* Zone. The supplementary diagnostic conodont fauna for the Lower Carboniferous, *Polygnathus purus subplanus* first appears at this level, too. *Siphonodella sulcata* Morphotype 5 and *Protognathodus kuehni* have a high correlation potential and a congruent FAD which can be used for defining the start of the joint *sulcata/kuehni* Zone; their phyletic lineages can be reconstructed in several regions. Thus, the use of the earliest morphotypes of *Siphonodella sulcata* M5 - and probably of *Siphonodella sulcata* M4 - and of *Protognathodus kuehni*, contribute to global correlations and biozonations. Morphotype 5 and Morphotype 4 are the earliest *Siphonodella sulcata*-morphotype to occur widely in Europe, North-Africa, Asia and North America at the base of the Tournaisian, without noteworthy overlapping ranges with 'siphonodelloids' (Figure 14).

A well-exposed succession of the early *Protognathodus* fauna, and the earliest morphotypes of *Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli*, *Protognathodus kockeli*, and *Protognathodus kuehni* are recorded at Tropl. However, atypical 'Protognathodus' morphotypes - proposed herein as *Protognathodus*-like 'protognathoids' - regarding diagnostic features of the cup and ornamentation of the platform (Figure 13) and partly evaluated by Hartenfels *et al.* (2022) - occur at Tropl and elsewhere around the DCB. Comparable with the *Siphonodella*-like 'siphonodelloids' which evolve and spread during late Famennian environmental changes in the *expansa* and *praesulcata* Zones connecting with the Epinette/Etreoungt and Drewer Events, the atypical morphotypes evolved and spread somewhat later, during late/latest Famennian and earliest Tournaisian environmental changes connecting with the Drewer Event in the *praesulcata* Zone (early phase of the Hangenberg Event) and in the *ckI* and *kockeli* Zone connecting with the middle and late phase of the Hangenberg Event (Figure 14). The record of their precise stratigraphic and distributional pattern is important for future

evaluation of phylogenetics because ancestor and descendent relationships of this faunal group are still unclear. Also, homeomorphy between the early and late protognathodids, gnathodids, and bispathodids, and a high morphological spectrum within the genera, could produce a bias of the conodont record in event-related successions. This is especially important due to overlapping ranges between these faunal groups at the DCB as well as in the Tournaisian. Thus, the extended *kockeli* Zone and the use of *Protognathodus kockeli* proposed recently as index fossil for the DCB, raises more questions than answers. The extinction-based *ckI* (HBSE), and the base of the former *kockeli* Zone (= Upper *praesulcata* Zone) are, however, needed for a precise biozonation scheme.

Carbon isotopes ( $\delta^{13}\text{C}_{\text{carb}}$ ) at Trolp show characteristic positive excursions and trends in the early, middle and late phases of the Hangenberg Crisis that can be readily correlated to many other regions worldwide. Accordingly, two positive  $\delta^{13}\text{C}_{\text{carb}}$  peaks in the *praesulcata* Zone represents  $\text{C}_{\text{org}}$  burial episodes of the early Hangenberg phase (Drewer Event). The onset of a distinctive positive carbon isotope excursion in the early/middle Hangenberg phase correlates with the initial conodont mass extinction episode (*ckI*) of the Hangenberg Crisis during globally widespread deposition of black shales (HBS), which precedes the main regressive episode during globally widespread deposition of regressive sandstones (HSS). This main anoxic episode of the Hangenberg Crisis (HBSE) is recorded in the thinly bedded limestone succession, and its onset may recorded in a locally developed thin shaly layer at the base of this succession. Because the pelagic fossil record of the regressive phase (HSSE) is generally unknown due to globally widespread, conodont-poor siliciclastics, stratigraphic gaps, it is recently unclear whether the main regressive phase of the Hangenberg Crisis is recorded in the thinly bedded limestone succession and needs further studies. The positive carbon isotope excursion in the late Hangenberg phase (*kockeli* Zone) at Trolp is time-equivalent to the first post-extinction recov-

ery phase; the onset of a distinctive decrease of carbon isotope values in the *sulcata/kuehni* Zone is coeval with the second conodont radiation phase; it can be correlated with same trends in many other successions worldwide.

## 5. Conclusions

The western Graz Paleozoic is of global significance because the phases of the Hangenberg Crisis at the DCB can be well reconstructed in continuous limestone successions at Trolp, and significant markers in bio-, litho- and chemostratigraphy can be well correlated outside the region. Conodont extinctions (*costatus-kockeli* Interregnum), conodont radiations and end of mass extinctions can be recognized at Trolp. The earliest morphotypes of *Siphonodella sulcata* M5, as well as of *Protognathodus meischneri*, *Protognathodus collinsoni*, *Protognathodus semikockeli*, *Protognathodus kockeli* and *Protognathodus kuehni* are recorded at Trolp. The globally common *Siphonodella sulcata* Morphotype 5 and Morphotype 4, as well as *Protognathodus kuehni*, have a high correlation potential worldwide and can be used for the joint *sulcata/kuehni* Zone; their phyletic lineages can be reconstructed from many regions.

Due to the spread of atypical morphotypes of *Protognathodus* ('protognathoids') at the DCB and in the Tournaisian related to the Hangenberg Event can be regarded as 'bottleneck' for the evolution of this faunal group. The previously suggested extended *kockeli* Zone and the recently proposed use of *Protognathodus kockeli* as index fossil for the DCB are not applicable due to the spread of these still unknown atypical morphotypes, and due to their unknown stratigraphic and distributional pattern and overlapping ranges with other homeomorphic groups. Also, the use of the extended *kockeli* Zone and the exceptional increasing amounts of morphological complex faunas especially among *Protognathodus kockeli* biased the conodont record and leaves the DCB and the Hangenberg event-related successions undivided. A conodont biozonation scheme with a fine-strati-

graphic subdivision, however, is needed for multi-disciplinary, high-resolution studies at the DCB. A strict biozonation and taxonomic concept enables further insights into the first-order Hangenberg mass extinction, also in respect to the age of the enigmatic end-Devonian glaciation. The loss of biozones in Hangenberg Event-related successions would present a potential loss of information and would throwback the scientific affords and progresses for several decades back.

## Contributions of authors

(1) Conceptualization: SIK; (2) Analysis or data acquisition: SIK, BH; (3) Methodologic/technical development: SIK; (4) Writing of the original manuscript: SIK, BH; (5) Writing of the corrected and edited manuscript: SIK, BH; (6) Graphic design: SIK, BH; (7) Fieldwork: SIK, BH; (8) Interpretation: SIK, BH; (9) Financing: BH.

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## Conflict of interest

There is no conflict of interest

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