

# A new Late Ordovician bubble-headed trilobite species from South West Wales and its implications for biostratigraphy

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A new species of bubble-headed trilobite, *Staurocephalus oliveae* sp. nov., is described from 250 specimens collected from a recently discovered locality in the Shoeshook Limestone Formation (upper Katian, Upper Ordovician; Ashgill, Cautleyan Zone 1 equivalent) in the village of Llanddowror, Carmarthenshire. Although no fully articulated specimens were found, the material is the most abundant and well-preserved of any Ordovician *Staurocephalus* species known from Britain and Ireland, and enables the complicated 3D structure of its cephalon to be reconstructed. *Staurocephalus oliveae* sp. nov. is the earliest representative of the genus recorded in Avalonia, following its presumed migration into the area from Laurentia or Baltica. Recognition of this Cautleyan species resurrects the potential usefulness of the long established, widely distributed, *Staurocephalus clavifrons* as a latest Katian (= Rawtheyan) indicator.

Key words: Trilobita, Phacopida, Encrinuridae, *Staurocephalus*, Late Ordovician, biostratigraphy, Wales, Great Britain.

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## Introduction

*Staurocephalus* was a long-ranging genus of “bubble-headed” trilobite (Phacopida, Encrinuridae), with the two oldest known species recorded from the lower Katian (uppermost Caradoc; Upper Ordovician) of Maine, USA (Owen and Bruton 2012) and the Oslo Region of Norway (Owen and Bruton 1980). It was an Ashgill (late Katian) immigrant from Laurentia or Baltica into the Anglo-Welsh area and Ireland and was the only member of the *Staurocephalinae* to survive into the Silurian (Owen 2007; Adrain 2013: 308). One species, *S. clavifrons* Angelin, 1854, has long been used as a marker for the uppermost Katian and possibly Hirnantian (both Upper Ordovician) in Sweden (Jaanusson 1963, Nielsen et al. 2023), Denmark and Poland (Kielan 1960) but its application has been questioned on the basis of occurrences of

*Staurocephalus* in Britain (Price 1973b). The material described herein provides some resolution to this matter.

A locality discovered by one of us (PDMcD) around 20 years ago at Talfan Farm near the village of Llanddowror, Carmarthenshire, South West Wales (Fig. 1; loc. 6 of Cocks 2014) has yielded a large number of specimens of a new species of *Staurocephalus* from the lower part of the Shoeshook Limestone Formation. The formation is largely correlated with the Cautleyan to lowest Rawtheyan stages of the Anglo-Welsh regional Ashgill Series (Fig. 2). The base of the Ashgill lies somewhere in the middle of the international Katian Stage (see Bergström et al. 2009) and thus the Shoeshook Limestone is upper Katian in global terms. The lowest part of the formation at Pengawse Hill near Whitland (see Fig. 1C) contains conodonts of the *Amorphognathus ordovicicus* Biozone (Ferretti et al. 2014) and chitinozoans from the upper part of the *Fungochitina spinifera* Biozone (Vandenbroucke et al. 2008). As noted by Vandenbroucke

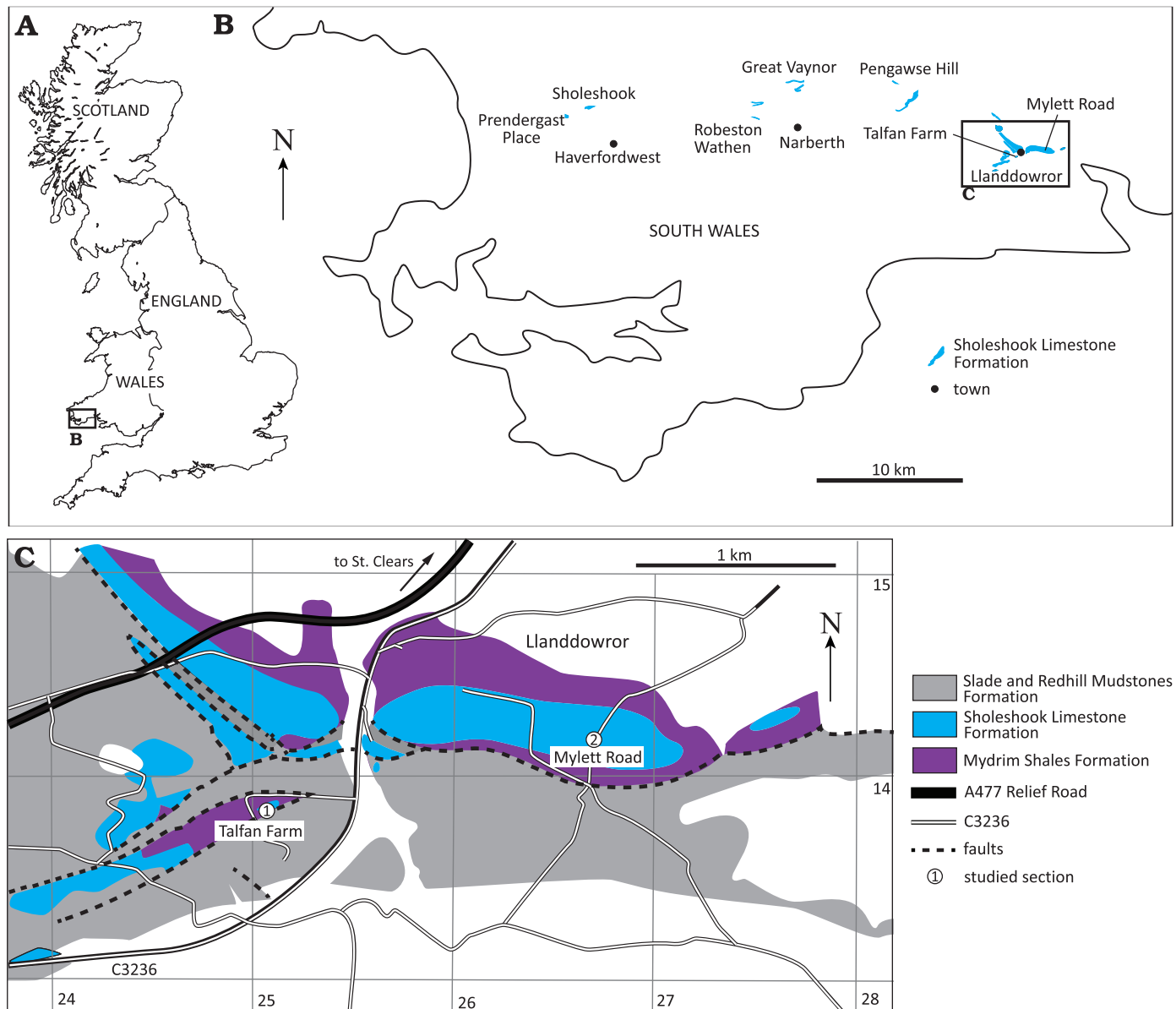


Fig. 1. Maps showing the location of the Talfan Farm fossil locality in Great Britain (A) and South West Wales (B). C. Geological map of the area around Llanddowror (greatly revised from BGS 1:50 000 Sheet 229 (Carmarthen), 1975 and Price (1973a: fig. 4) on the basis of recent field observations by PDMcD. Numbers on the x and y axis are co-ordinates (eastings and northings) on the UK National Grid. Talfan Farm is located at latitude  $51^{\circ}47'41''\text{N}$  and longitude  $4^{\circ}32'27''\text{W}$ .

et al. (2008: 209), the latter supports the correlation close to the Pusgillian/Cautleyan boundary (upper Katian) indicated by the trilobites (Zalasiewicz et al. 1995). The formation at Pengawse Hill overlies shales of the Mydrim Shales Formation with graptolites of the “*Normalograptus* proliferation interval”, the upper part of which might extend into the *Pleurograptus linearis* Biozone (Vandenbroucke et al. 2008: 209). As noted below, the trinucleid trilobite *Tretaspis moeldenensis moeldenensis* Cave, 1960, is present in the section at Talfan Farm indicating an earliest Cautleyan age for the occurrence of *Staurocephalus* at Talfan Farm, possibly slightly younger than the base of the formation at Pengawse Hill. It is the oldest recorded occurrence of the genus from Avalonia.

Although there are no complete exoskeletons and the very convex cephalon is subject to varying degrees of deformation, the Sholeshook Limestone specimens of *Staurocephalus oliveae* sp. nov. provide the most complete information on the morphology of any of the Ordovician occurrences of the genus in Britain and Ireland and, with the possible exception of *S. clavifrons*, the most complete of any Ordovician species of *Staurocephalus* so far described.

*Nomenclatural acts.*—This published work and the nomenclatural acts it contains have been registered in ZooBank: urn:lsid:zoobank.org:pub:BAF4499F-2ED3-4DB8-AE0C-3A3F58A5A280

*Institutional abbreviations.*—BIRUG, Lapworth Museum of

Geology, University of Birmingham, UK; GLAHM, Hunterian Museum, University of Glasgow, UK; NMW, National Museum Wales, Cardiff, UK; SM, Sedgwick Museum of Earth Sciences, University of Cambridge, UK.

*Other abbreviations.*—exsag., exsagittal; L, lateral glabellar lobe; max., maximum; S, lateral glabellar furrow; sag., sagittal; tr., transverse.

## Geological setting

The Sholeshook Limestone Formation is an upper Katian (Upper Ordovician) rock unit a little over 60 m thick in its most complete exposure near Sholeshook, comprising calcareous mudstones, siltstones and sandstones intercalated with some purer limestone horizons (Price 1973a; Owens 2000). It crops out in three main areas in South West Wales within a 30 km, roughly west-east trending band: at Sholeshook and Prendergast in the Haverfordwest area, at Robeston Wathen, and around the village of Llanddowror near St. Clears, south-west of Carmarthen (Fig. 1A–C). This study is based largely on fossils collected from the lower part of the formation at Talfan Farm near Llanddowror (Fig. 1D). In this area, the Sholeshook Limestone is represented by basal calcareous mudstones, which gradually coarsen upwards into coarse calcareous sandstone that makes up the bulk of the formation (Price 1973a). The majority of specimens at Talfan Farm were collected from a vertical section from three horizons in the basal Sholeshook Limestone at 1, 2, and 3 m above the boundary with the underlying Mydrim Shales Formation. A small number of additional specimens were from two spot samples, namely a deep 1 m diameter circular pit and a 1×2 m trench, both of which exposed the top of the Mydrim Shales Formation and the basal 1–3 m of the Sholeshook Limestone Formation. All five sampled horizons yielded very similar fossil faunas.

Marr and Roberts (1885) introduced the term “Sholeshook Limestone”, and Marr (1907) placed the formation at the base of his newly proposed Ashgill Series, though a subsequent revision (Ingham and Wright 1970) redefined the base of the series at a lower horizon. The Sholeshook Limestone has long been recognized for its diverse trilobite fauna, with Price (1974, 1977, 1980b, 1981) documenting in detail over fifty species. Price (1973a, 1980a) established the age of the Sholeshook Limestone Formation based on correlation of the trilobite species present with those in the shelly biozones in the type area for the Ashgill Series, the Cautley and Dent areas of northern England (Ingham 1966; Ingham and Wright 1970, 1972), with particular emphasis on the sequence of species of the trinucleid trilobite *Tretaspis*. The Upper Ordovician chitinozoan biozonal scheme established by Vandenbroucke (2008) for the largely siliciclastic successions of England and Wales incorporates the Ashgill zones documented in northern England by Vandenbroucke et al. (2005), which in turn, enhance the potential for international correlation of many

of the shelly zones (Fig. 2). Apart from the basal few metres at Pengawse Hill (Vandenbroucke et al. 2008), chitinozoans have not been sampled from the Sholeshook Limestone.

The basal Sholeshook Limestone was correlated with the upper part of Cautleyan Zone 1 of Ingham (1966) by Price (1973a) with the top of the formation having a low Rawtheyan (Zone 5, upper Katian) age (Price 1980a; Fig. 2).

Price (1973a: 243) recognized a stratigraphical gap between the Cautleyan (upper Zone 1) base of the Sholeshook Limestone and the underlying Mydrim Shales Formation at Mylet Road, Llanddowror (Fig. 1C), with the latter placed in the Caradoc Onnian Stage (now substage) (lower Katian) by Cave (1960) based on the graptolite fauna found in its highest beds. Hence, there appears to be a hiatus in the Mylet Road succession encompassing the Ashgill Pusgillian Stage and likely also parts of the Onnian and Cautleyan Zone 1 (Owens 2000: 159–161). However, further west at Pengawse Hill, Whitland (Fig. 1C), Zalasiewicz et al. (1995) used graptolite evidence to show a continuous succession through the top of the Mydrim Shales Formation and basal Sholeshook Limestone, with no apparent gap, and they suggested that the base of the Sholeshook Limestone may range down into the Pusgillian, a correlation subsequently supported by Vandenbroucke et al. (2008: 209). Hence, it appears that the base of the Sholeshook Limestone Formation is diachronous. The contact with the overlying Slade and Redhill Mudstone Formation (Fig. 2) is conformable, with some tri-

Global stratigraphy	British regional chrono-stratigraphy	Composite Anglo-Welsh chitinozoan biozonation (Vandenbroucke 2008)	Shelly zones (Ingham 1966; Ingham and Wright 1972)	Lithostratigraphy	
Upper Ordovician (part)	Hirnantian	Hirnantian	8	Haverford Mudstone Formation	
			Taugourdeau		
	Ashgill	Rawtheyan	Umbilicata	7	Portfield Formation
			Fossensis	6	Slade and Redhill Mudstone Formation
			Rugata	5	
		Cautleyan	?Bergstroemi	4	Sholeshook Limestone Formation
				3	
				2	
				1	
		Pusgillian	Spinifera (part)		*

Fig. 2. Stratigraphical column showing the age of the Sholeshook Limestone Formation (in blue) and the underlying and overlying formations together with the shelly and chitinozoan biozonations of the Anglo-Welsh succession. Asterisk shows approximate age of the Talfan Farm locality, in the lower part of the formation (equivalent to Cautleyan Zone 1).

lobite species ranging through the boundary. Price (1973a) argued that the base of the Slade and Redhill Mudstone Formation also appears to be diachronous, and part of the formation may be contemporaneous with the Shoeshook Limestone in its western outcrops although we consider that this may not prove to be the case.

Based on sedimentological evidence and the trilobite genera present, Price (1980b) interpreted the depositional setting of the Shoeshook Limestone as a relatively high energy environment, somewhere on the middle to upper part of the slope between the platform edge and basin. A diverse non-trilobite fauna has also been reported from the formation (e.g., Paul 1973, 1982, 1997; Donovan and Paul 1985; Barrs 2013; Ferretti et al. 2014; Cocks 2014) and the Talfan Farm locality has yielded abundant echinoderms (crinoids, cystoids, and coronates; Lanc et al. 2015; McDermott and Paul 2015) and brachiopods (Cocks 2014), along with rarer corals (Peel and McDermott 2016), bryozoans, gastropods, cephalopods, and ostracods. The presence of the trinucleid trilobite *Tretaspis moeldenensis moeldenensis* indicates an early Cautleyan (Zone 1) age.

Price (1973a, 1980b) recorded rare occurrences of *Stau-rocephalus clavifrons* (later *S. cf. clavifrons*) from various levels throughout the Shoeshook Limestone Formation, and from the basal Slade and Redhill Mudstone Formation (see below).

## Material and methods

*Stau-rocephalus oliveae* sp. nov. is described based on 250 specimens recently collected, with permission, from Talfan Farm and now housed in the Palaeontology collections of the National Museum of Wales, Cardiff. The collection encompasses all sclerites apart from the hypostome. No complete articulated exoskeletons were recovered and most of the highly convex cranidial specimens have undergone some degree of deformation or crushing, but the large number of specimens available collectively provides a comprehensive view of the original morphology of the species. The majority of specimens are preserved as internal moulds, with small numbers of external moulds and internal/external mould part/counterpart pairs.

Specimens from the National Museum of Wales collections were whitened with ammonium chloride before photographing using a Canon 80D camera attached to a Leica Z6 microscope; those from other museum collections were photographed unwhitened in accordance with the loan conditions of those institutions. Multiple images were taken of each specimen at different depths of field, which were then stacked to produce composite images using Helicon Focus software. Photographs were edited and photographic figures compiled using Adobe Photoshop Elements 2022. Some external moulds were cast in latex to reveal details of the external sculpture and were consolidated prior to casting using a 5% w/w solution of Paraloid B-72 adhesive in Acetone.

## Systematic palaeontology

Class Trilobita Walch, 1771

Order Phacopida Salter, 1864

Suborder Cheirurina Harrington & Leanza, 1957

Family Encrinuridae Angelin, 1854

Subfamily Stau-rocephalinae Prantl & Přibyl, 1947

*Remarks.*—Although given full family rank by many earlier workers (e.g., Kielan 1957; Evitt and Tripp 1977; Holloway 1980; Owen and Bruton 1980; Tomczykowa 1987), the concept of Stau-rocephalidae was largely based on the distinctive “bubble headed” morphology of derived members of the group. We concur with Adrain (2013: 307–308) in regarding Stau-rocephalinae as a subfamily of Encrinuridae, on the basis of shared morphological traits of protaspid and early meraspid stages of less derived stau-rocephalines such as *Libertella*, encrinurines, and “advanced” cybelines (see Edgecombe et al. 1988).

Genus *Stau-rocephalus* Barrande, 1846

*Type species:* *Stau-rocephalus purchisoni* Barrande, 1846, from the Liteň Formation (Wenlock) of Bohemia, Czech Republic.

*Emended diagnosis.*—Glabella with subparallel-sided posterior portion bearing short S1 and S2; S3 extends across glabella; frontal lobe greatly inflated and overhanging anterior margin of cephalon. Anterior part of rostral plate expanded and continuous with frontal glabellar lobe; short fixigenal spine present in most species. Thorax of 10 segments. Pygidium with 3 pairs of spines. Emended from Curtis and Lane (1997: 50)

*Stau-rocephalus oliveae* sp. nov.

Figs. 3, 4, 5, ?6A.

non1885 *Stau-rocephalus globiceps* Portlock; Marr and Roberts 1885: 481 [= *Stau-rocephalus cf. clavifrons*].

?1938 *Stau-rocephalus cf. purchisoni* Barrande; Whittington 1938: 451.

?1973 *Stau-rocephalus clavifrons* Angelin; Price 1973a: tables 1–3, non 7 [= *Stau-rocephalus cf. clavifrons*].

?1980 *Stau-rocephalus cf. clavifrons* Angelin; Price 1980b: 866 (Shoeshook Limestone specimens only), pl. 110: 18.

*ZooBank LSID:* urn:lsid:zoobank.org:act:99371F20-B988-498A-81E0-55729B248609

*Etymology:* After Olive Potts (St. Clears, Carmarthenshire, UK), who discovered the first specimen of this species.

*Type material:* Holotype: almost complete cephalon, NMW 2013.11G.1 (Fig. 3A). Paratypes: almost complete cephalon with eyes preserved, NMW 2013.11G.2 (Fig. 3B); one further cephalon, NMW 2013.11G.8 (Fig. 3E); three cranidia, NMW 2013.11G.6, 7, 9 (Fig. 3C, D, F); three librigenae, NMW 2013.11G.3, 10, 12 (Fig. 4B–D); three pygidia, NMW 2013.11G.4, 11, 15 (Fig. 4F, G, H); one rostral plate, NMW 2013.11G.5 (Fig. 4A); one thoracic segment, NMW 2013.11G.13 (Fig. 4E). All from the type locality and horizon.

*Type locality:* Talfan farm, Llanddowror, Carmarthenshire, Wales, UK.

*Type horizon:* Shoeshook Limestone Formation (Cautleyan, upper Katian), Upper Ordovician.



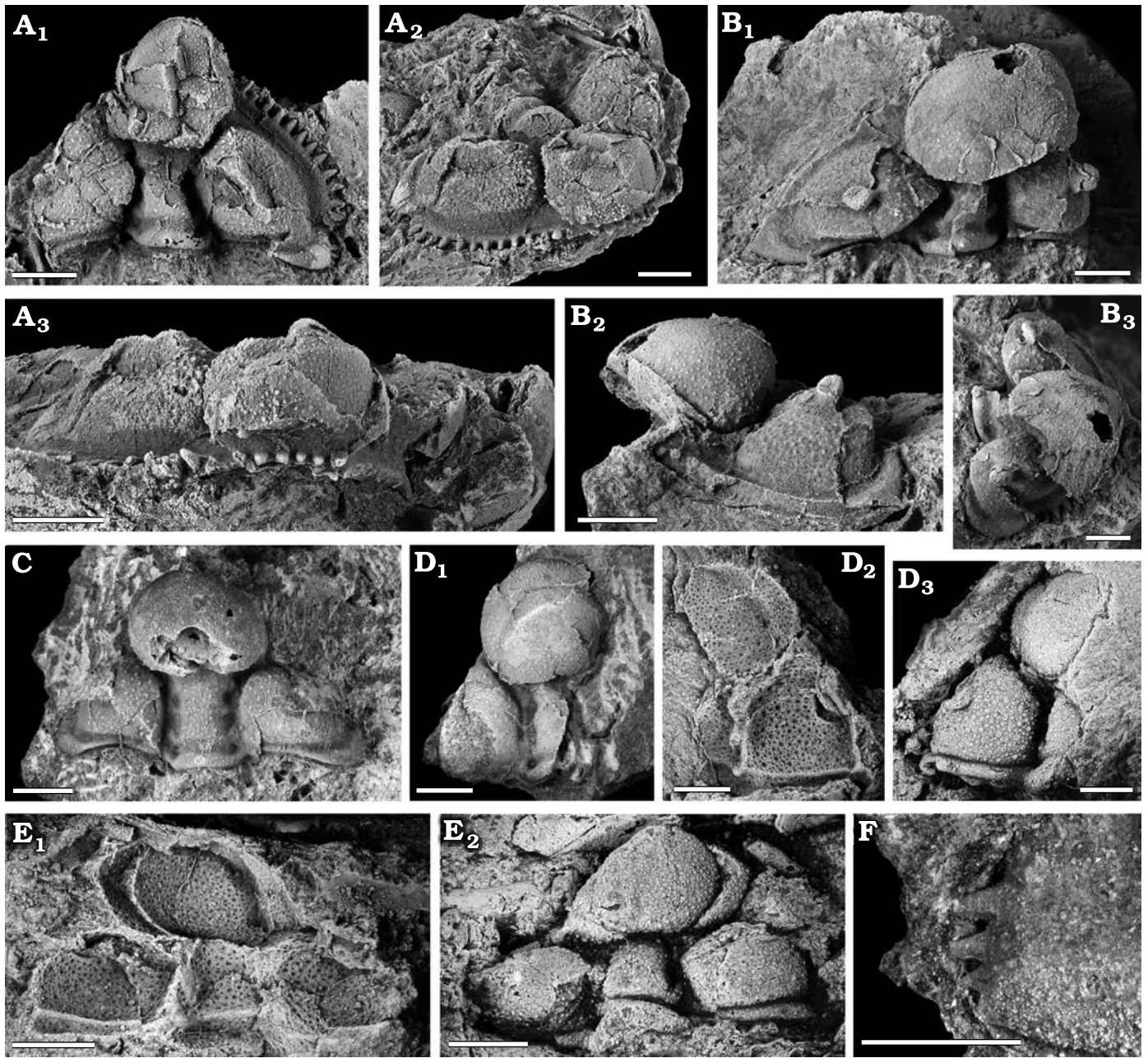


Fig. 3. Encrinurid trilobite *Staurocephalus oliveae* sp. nov. from the upper Katian (Upper Ordovician) Sholeshook Limestone Formation at Talfan Farm, Llanddowror, Carmarthenshire, Wales, UK. **A.** Holotype, NMW 2013.11G.1, cephalon in dorsal (A<sub>1</sub>), oblique anterolateral (A<sub>2</sub>), and frontal (A<sub>3</sub>) views. **B.** Paratype, NMW 2013.11G.2, cephalon in dorsal (B<sub>1</sub>), oblique anterolateral (B<sub>2</sub>), and dorsolateral (B<sub>3</sub>) views. **C.** Paratype, NMW 2013.11G.6, cranidium in dorsal view. **D.** Paratype, NMW 2013.11G.7, partial cranidium, dorsal view of internal mould (D<sub>1</sub>), ventral view of external mould (D<sub>2</sub>), and latex cast of external mould (D<sub>3</sub>). **E.** Paratype, NMW 2013.11G.8, cephalon, external mould in ventral view (E<sub>1</sub>) and latex cast of external mould (E<sub>2</sub>). **F.** Paratype, NMW 2013.11G.9, genal angle of cranidium, showing marginal denticles decreasing in size posteriorly. Scale bars 2 mm, except for F 1 mm.

**Material.**—Type material and 236 specimens (88 cephalo/cranidia, 33 pygidia, 80 librigenae, 33 rostral plates, one thoracic segment; NMW 2013.11G.13, 14, 16–249) from the type locality. One cranidium, (NMW 2013.11G.250) and one pygidium (NMW 2013.11G.251) from the Sholeshook Limestone Formation at Mylet Road near Llanddowror, Carmarthenshire. Price (1980b) reported 15 cranidia of *S.* cf. *clavifrons*, ranging throughout the Sholeshook Limestone Formation into the basal Slade and Redhill Mudstone

Formation, from the Sholeshook and Prendergast areas near Haverfordwest and from the Llanddowror area but not all of his specimens could be located in museum collections. The single cranidium (GLAHM: A 9751) figured by Price (1980b: pl. 110: 18), from low in the Sholeshook Limestone in Craig-y-deilo Quarry, Llanddowror, was collected by J. Keith Ingham (Glasgow, UK) and is very likely conspecific with our material (see Remarks, below). The specimen register at the Hunterian Museum, Glasgow, UK, notes those



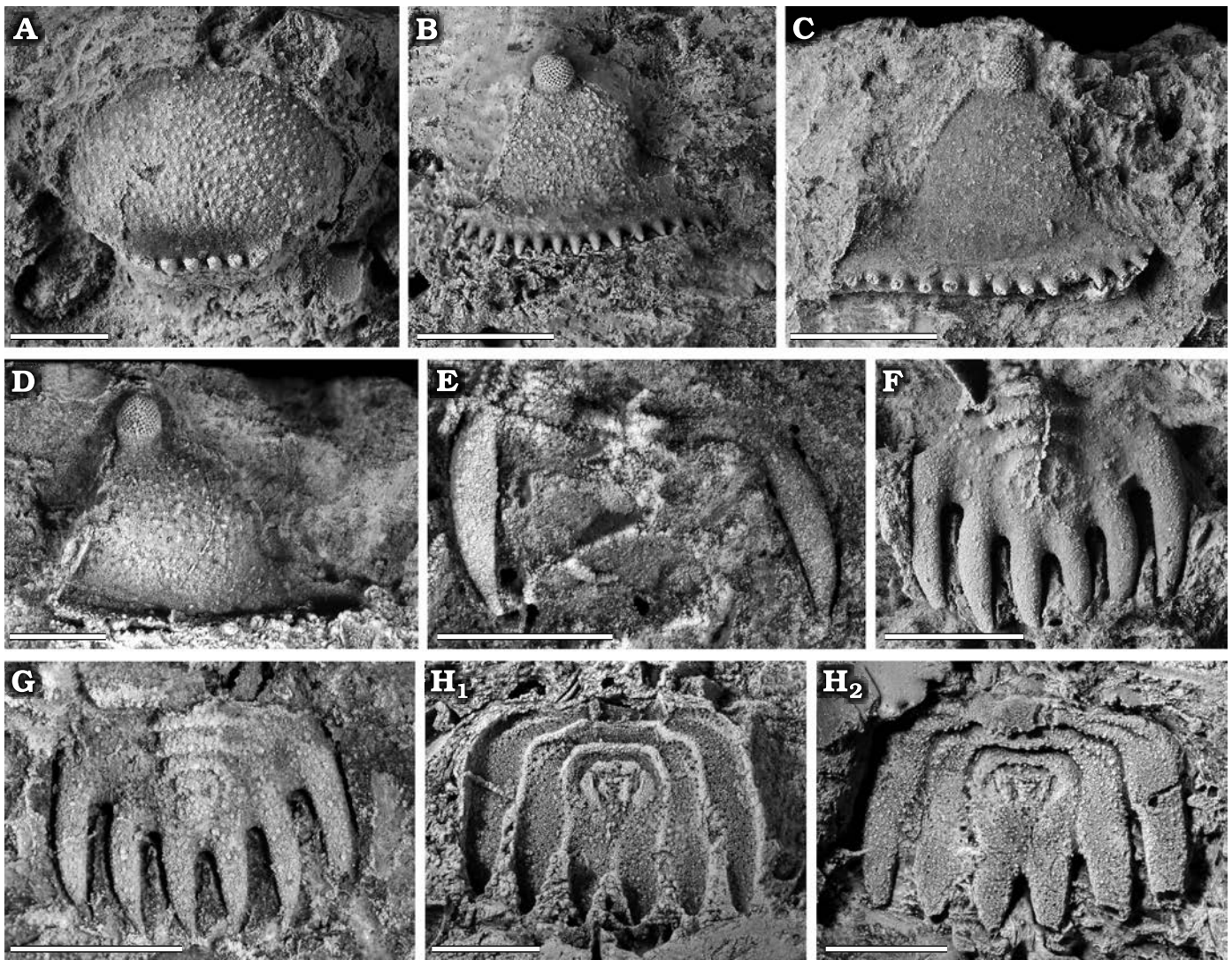


Fig. 4. Encrinurid trilobite *Staurocephalus oliveae* sp. nov. from the upper Katian (Upper Ordovician) Sholeshook Limestone Formation at Talfan Farm, Llanddowror, Carmarthenshire, Wales, UK. A. Paratype, NMW 2013.11G.5, rostral plate in ventral view. B. Paratype, NMW 2013.11G.3, right librigena in lateral view. C. Paratype, NMW 2013.11G.10, left librigena in lateral view. D. Paratype, NMW 2013.11G.12, right librigena in lateral view. E. Paratype, NMW 2013.11G.13, thoracic segment in dorsal view. F. Paratype, NMW 2013.11G.4, pygidium in dorsal view. G. Paratype, NMW 2013.11G.11, pygidium in dorsal view. H. Paratype, NMW 2013.11G.15, pygidium in ventral view of external mould (H<sub>1</sub>) and latex cast of external mould (H<sub>2</sub>). Scale bars 2 mm.

collected by J.K. Ingham in the 1960s from the Sholeshook Limestone that were studied by David Price as part of his Ph.D. research (1971), and these include the figured cranidium and six further *Staurocephalus* specimens (what became referred by Price 1973a: “one from Loc. 18c, four from Loc. 18d, and two from Loc. 25”). Unfortunately, these specimens cannot currently be located, but are presumably among the 15 *Staurocephalus* cf. *clavifrons* cranidia listed by Price (1980b). Two cranidia from the historic collections of the Sedgwick Museum, Cambridge, UK, collected from the Sholeshook Limestone Formation in the Sholeshook railway cutting, Haverfordwest area, were labelled as *Staurocephalus globiceps* (SM A31591) and *Staurocephalus* cf. *murchisoni* (SM A31592; Fig. 6A) but likely also belong in our new species. Three further fragmentary cranidia labelled “*Staurocephalus globiceps*” in the Sedgwick Museum col-

lections (SM A31587–9) were collected by Marr and Roberts (1885) from a locality near Pelcomb Cross, in strata now assigned to the basal Slade and Redhill Mudstone Formation (see Price 1973a). These specimens are generally too poorly preserved for specific identification, but one possesses a prominent genal spine (Fig. 6E) and is excluded from *S. oliveae* sp. nov. (see below).

**Diagnosis.**—Differs from other congeners by possessing the following combination of characters: *Staurocephalus* species with right-angled genal angle, lacking a genal spine or protrusion, with four lateral denticles anterior to the genal angle on the fixigena, decreasing in size posteriorly. Librigenae with 13–17 marginal denticles; mode and mean = 15. Inflated anterior glabellar lobe making up around two-thirds of preglabellar glabellar length (sag.). Pygidium around one

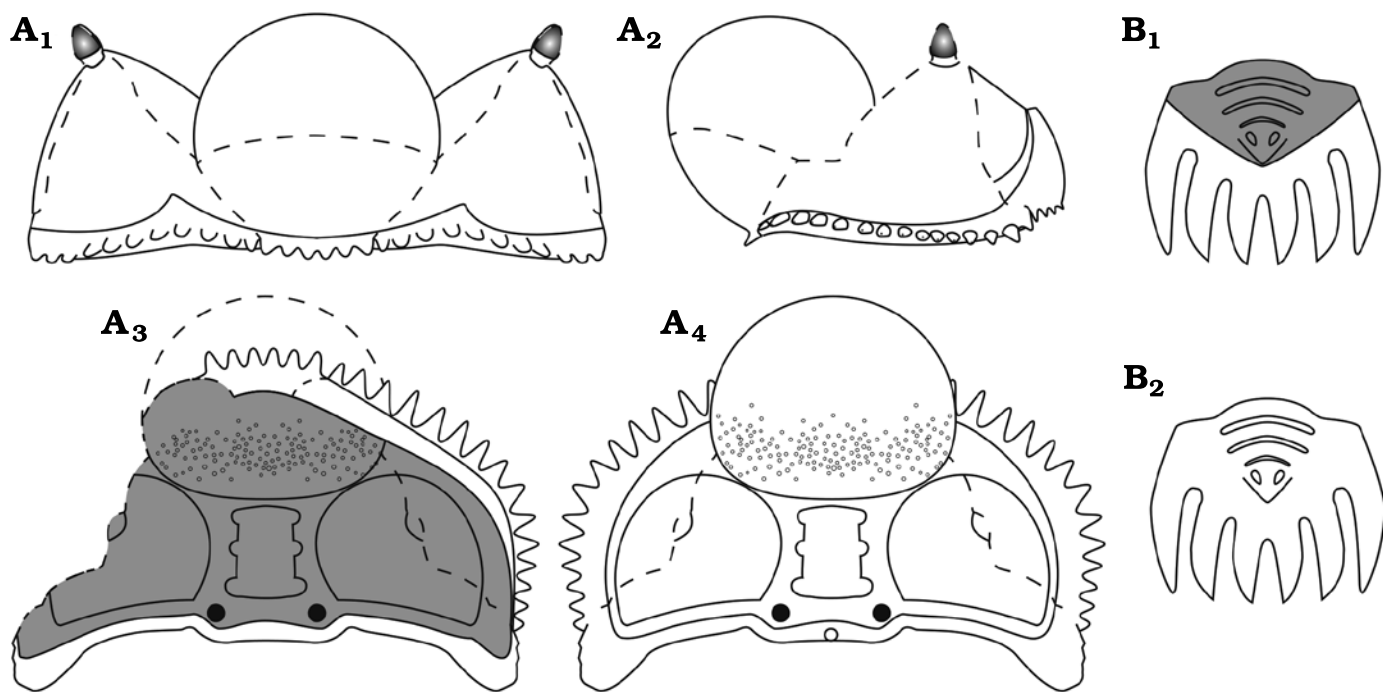


Fig. 5. Reconstruction drawings of the encrinurid trilobite *Staurocephalus oliveae* sp. nov. **A.** Cephalon, in anterior (A<sub>1</sub>), left lateral (A<sub>2</sub>), ventral (A<sub>3</sub>), and dorsal (A<sub>4</sub>) views. **B.** Pygidium, in ventral (B<sub>1</sub>) and dorsal (B<sub>2</sub>) views. Cephalon has a mean maximum width (tr.) of 10.15 mm; pygidium has a mean maximum width (tr.) of 3.96 mm.

fifth wider (tr.) than long. Inner of three pairs of spines making up just under two-fifths of total pygidial length. Cephalon (including rostral plate and librigenae) with dense covering of fine to medium granules, median occipital tubercle centrally located or on the posterior half of the occipital ring; pygidium with dense fine granules on spines with larger granules on axial rings and pleural ribs.

**Description.**—No complete articulated exoskeletons known. Cephalon approximately semicircular in outline; cranium subtriangular (Figs. 3, 5A). Cephalon max. length (sag.) mean = 6.86 mm, range = 2.39–10.08 mm (n = 62). Cephalon max. width (tr.) mean = 10.15 mm, range = 3.37–14.01 mm (n = 42). On those specimens (n = 42) that are complete enough to measure both dimensions, maximum width (tr.) (mean = 10.02 mm, range = 3.37–14.01 mm) of cephalon is about one and a half times (mean = 1.50, range = 0.75–2.18) the maximum length (sag.) (mean = 6.71 mm, range = 2.39–8.45 mm). Glabella strongly convex (tr.), consisting of a greatly inflated, subcircular anterior glabellar lobe in front of a rectangular glabellar stem. In some specimens, glabellar stem tapers gently forwards, however, in less crushed specimens (e.g., NMW 2013.11G.6; Fig. 3C), margins of stem are sub-parallel. Occipital ring accounts for about one fifth of transverse posterior width of cephalon and about a tenth of the total cephalic length (sag.) medially, tapering abaxially by about a third; separated from the posterior border by a very shallow furrow. Median tubercle present, centrally located or on the posterior half of the occipital ring (Fig. 3A<sub>1</sub>, B<sub>1</sub>, C). Occipital furrow shallow and follows the transverse convexity of the glabellar stem and occipital

ring, ending in deep pits where it meets the axial furrows. Anterior to the occipital furrow is the narrow (tr.), rectangular glabellar stem, indented by the lateral glabellar furrows S1 and S2; uniformly deep, semi-circular pits. S1 located approx. one fifth of length (sag.) of the glabellar stem in front of occipital furrow; L1 small, an elongate suboval abaxially, narrowing and subparallel-sided adaxially, weakly inflated, angled forwards at around 40° to the axis. S2 located approx. 70% along the length of the glabellar stem; L2 sub-rectangular, shorter, less inflated and less strongly angled forward than L1; L3 a very weakly inflated, transverse area behind S3. Transglabellar (S3) furrow marks the anterior end of the glabellar stem behind the inflated anterior lobe. S3 is consistently shallow, following the transverse convexity of the glabellar stem, and ends at the deep axial furrow where there is a suggestion of a pit in some specimens although this feature is not clearly preserved in any. Inflated anterior glabellar lobe (mean sag. length: 3.75 mm, n = 20) occupies about two thirds (mean = 0.59, range = 0.50–0.77) of total sagittal glabellar length (mean: 6.31 mm, n = 20), excluding occipital ring, while being approximately three times as wide (tr.) as the glabellar stem (excluding occipital ring), with about half of its length (sag.) protruding over the anterior cephalic margin. The axial furrows are consistently deep alongside the glabellar stem, then anterior of the transglabellar furrow they follow the curvature of the inflated anterior glabellar lobe, between the anterior glabellar lobe and inflated anterior fixigena, remaining consistently deep.

Fixigenae occupy about four fifths of the posterior (tr.) cephalic width. Palpebral lobe semicylindrical, bounded by



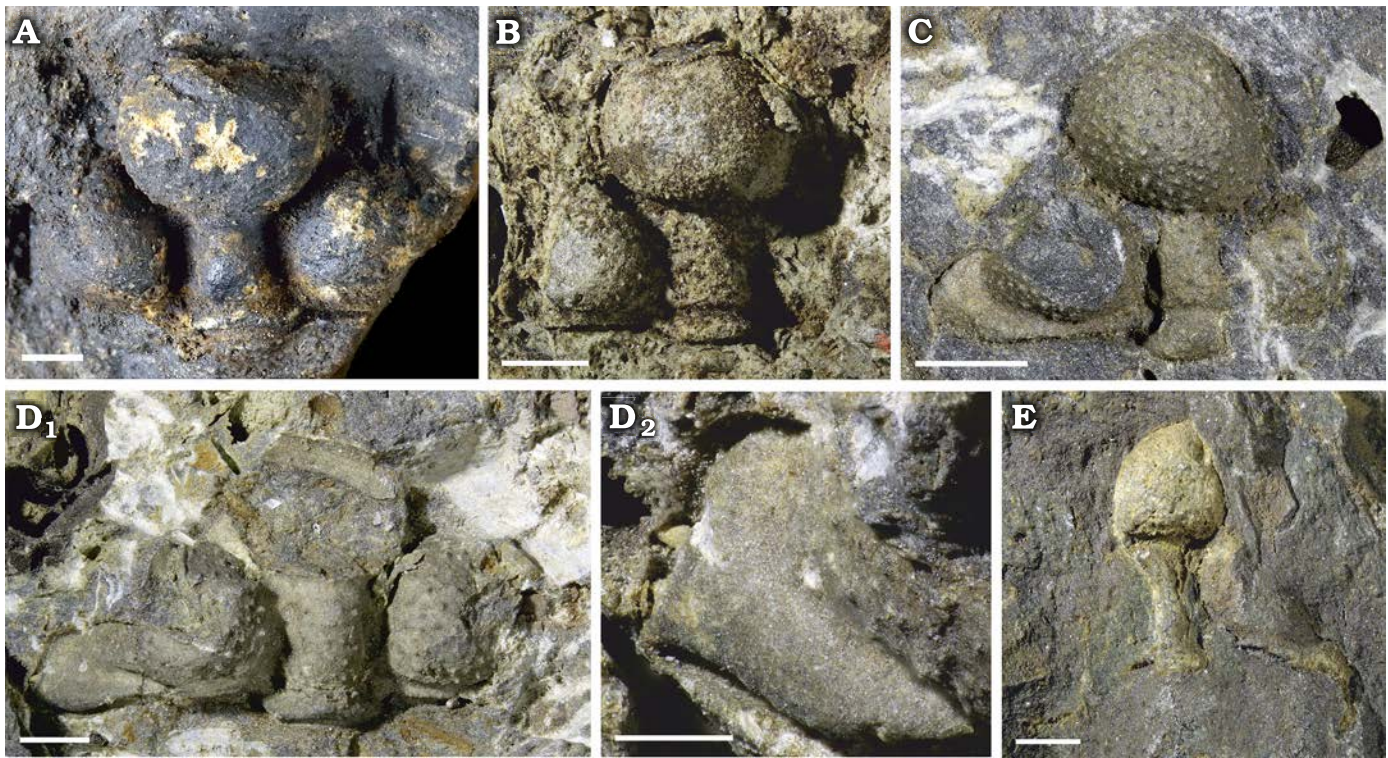


Fig. 6. Historic specimens of the encrinurid trilobite *Staurocephalus* from museum collections. **A.** SM A31592, cranium in dorsal view labelled “*Staurocephalus* cf. *murchisoni* Barrande” from the Sholeshook Limestone Formation (upper Katian, Upper Ordovician) in the Sholeshook railway cutting, Haverfordwest area, herein assigned to *S. oliveae* sp. nov.? **B.** BIRUG 48862, cranium in dorsal view labelled “*Staurocephalus* cf. *murchisoni*” from the lower Tre-wylan Beds (Cautleyan) (upper Katian, Upper Ordovician) in the Llansantffraid-ym-Mechain district of the south-eastern Berwyn Hills, Mid Wales (Whittington 1938: 451, locality 42), herein assigned to *S. oliveae* sp. nov.? **C, D.** Specimens from the King collection, Sedgwick Museum, from the (locally) basal Ashgill mudstones (lower Rawtheyan, Ashgill Zone 5) (upper Katian, Upper Ordovician) of Aber Marchnant, Berwyn Hills, Mid Wales, originally identified as *Staurocephalus* cf. *murchisoni* by King (1923) and assigned to *S. clavifrons* by Whittington (1968), herein assigned to *S. aff. oliveae*. **C.** SM A39813, incomplete cranium in dorsal view. **D.** SM A39814.A, incomplete cranium in dorsal view (**D<sub>1</sub>**) and close up of left genal area (**D<sub>2</sub>**) to show large lateral denticles (arrowed). **E.** SM A31589, dorsal view of incomplete cranium labelled “*Staurocephalus globiceps*” from the Marr collection, Sedgwick Museum, from the Slade and Redhill Mudstone Formation at Pelcomb Cross, Haverfordwest, herein assigned to *S. cf. clavifrons*. Note the prominent genal spine on the right side. Scale bars 1 mm, except **D<sub>2</sub>** 0.5 mm.

a shallow furrow; located at the approximate mid width (tr.) of the fixed cheek, slightly posterior to level of S2 furrow. Fixigenal field inflated, sloping down steeply anteriorly, highest parts positioned above the level of the glabellar stem and commonly reaching half the preserved height of the anterior glabellar lobe in posterior view, although in uncrushed specimens the anterior glabellar lobe was probably somewhat higher. The posterior border is weakly convex (exsag.), extending transversely (sag.) from the occipital ring to behind the palpebral lobe beyond which it widens (exsag.) markedly towards the genal angle where it joins the posterior end of the lateral cephalic border at a right angle. No genal spine present, however four small denticles present on lateral border, increasing in size anteriorly and continuous with the larger denticles of the librigena. Posterior border furrow shallow, broadening markedly abaxially alongside the broadening of the posterior border.

The posterior branch of the facial suture is directed transversely from the palpebral lobe curving very gently forwards, until it reaches the lateral border where it curves slightly posteriorly. The anterior branch of the facial suture extends

parallel to the sagittal line, from the palpebral lobe to opposite the middle (sag.) of the anterior glabellar lobe. Here the suture curves adaxially and is obscured by the anterior glabellar lobe in dorsal view, meeting the rostral and connective sutures on the flank of the anterior cephalic “bubble”.

Librigena triangular, maximum length subparallel to the lateral border approximately two fifths longer than width normal to this; librigenal field also has a maximum length subparallel to lateral border two fifths longer than width normal to the field (Fig. 4B–D). Librigenae are a little less convex in comparison to the fixigenae. Short (exsag.) eye lies at summit of inflated field, which descends relatively steeply abaxially, separated from it by a shallowing of the slope. Visual surface of eye sub-oval, convex, taller than wide, set on top of a socle of around half its height. Visual surface containing over 120 small hexagonal lenses, with over 16 around the base, (Fig. 4D; NMW 2013.11G.12, max. length 5.94 mm, max. width 4.31 mm), arranged in hexagonal packing. Librigenal border furrow uniformly broad and shallow. Librigenal border comprises a very narrow, flat-lying, very weakly swollen ridge, barely distinguished



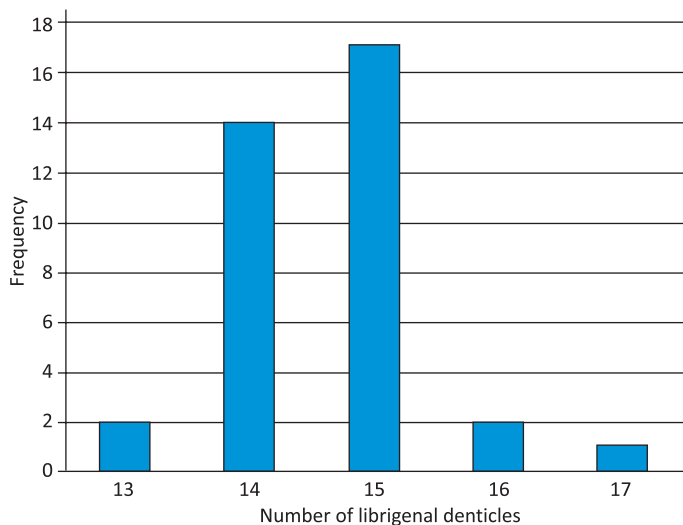


Fig. 7. Chart showing the frequency distribution of numbers of denticles on lateral margins of complete librigenae of *Staurocephalus oliveae* sp. nov. from the upper Katian (Upper Ordovician) Sholeshook Limestone Formation at Talfan Farm, Llanddowror, Carmarthenshire, Wales, UK.  $n = 36$ .

from the border furrow, and a row of denticles, each longer than the width of the ridge. On complete specimens ( $n = 36$ ), 13–17 denticles are present with a mean of 15 (rounded from 14.6) and mode of 15 (Fig. 7). The number of denticles is independent of the size of the librigena. Portion of librigena anterior of inflated field comprises a low-lying subtrapezoidal area, the anteriormost part of which is upturned to form a subtriangular flap, the librigenal precranial lobe (Figs. 4B, C, 5A<sub>2</sub>). Anterior edge of precranial lobe has a very gentle forwards curvature that matches the curve of the anterolateral margin of the rostral plate, the two librigenal precranial lobes fitting with the rostral plate along the connective sutures to form the underside of the inflated cephalic lobe (“bubble”).

Rostral plate sub-circular, split into two distinct parts (Fig. 4A). Anterior part of plate lies beneath the anterior glabellar lobe along the rostral suture and is inflated, matching the convexity of anterior glabellar lobe and forming the anteroventrally directed part of the anterior cephalic “bubble” (see Fig. 5). Separated by a transverse furrow, the posterior section consists of a set of six denticles. Behind the denticles there is short, flat ventral section, of similar length (sag.) to the denticles and continuing in the same plane as them.

Hypostome not known.

Internal mould of cranidium (including rostral plate) apart from the furrows and borders densely and evenly covered in relatively small granules. Latex casts of external moulds reveal the granules to be larger and more densely packed on external exoskeletal surfaces, although the number of granules remains consistent (Fig. 3D<sub>3</sub>, E<sub>2</sub>). Additionally, the larger granules shown on latex casts vary in size. Internal moulds and latex casts of external moulds of the librigenal field have similar granulation to those of the cranidium, with occasional larger granules situated on the marginal border, in line with the denticles.

Thorax known only from internal moulds of two isolated thoracic segments from the type locality that probably belong to this species (Fig. 4E) and are consistent with segments from the posterior part of the thorax (e.g., see specimen of *S. clavifrons* illustrated by Kielan 1957: pl. 3: 2). Axial ring occupies about a quarter of segment width (tr.). Pleura parallel-sided over most of its length (tr.) turning posteriorly through almost a right angle at the fulcrum and extended as a posteriorly directed pleural spine, almost parallel to the sagittal line, that tapers to a sharp point over the distal 30% of its length. The granulation of these spines is denser and finer than that of the cranidium. Prominent large granules present at the fulcrum. A small number of medium granules developed into short spines visible on a small part of the axial ring of one of the specimens, but preservation too poor to determine overall sculpture.

Pygidium subquadrate in outline, being about one fifth wider (tr.) than long, including spines (Figs. 4F–H). Axis occupies approximately 50% of pygidial length (sag., including spines), and approximately 30% of the anterior pygidial width. Axis convex (tr.) anteriorly, becoming progressively less so and tapering (tr.) posteriorly. Axis has three distinct axial rings, the first two being transverse with the third arching gently forward, separated by axial ring furrows, which become relatively shallower posteriorly following the decreasing convexity of the axis. Each furrow is shallow and narrow medially, widening and becoming deeper laterally. Axis ends with a subtriangular, flat terminal piece. Axial furrows very shallow, little more than a break in slope between the convex axis and flat-lying pleural areas. Axial rings confluent with pleural ribs, the first pair of which are directed transversely for a short distance before curving posteriorly through about 90°, the second and third pairs are essentially parallel to the sagittal line. On external surface (Fig. 4H<sub>2</sub>), the transverse, inner parts of the anterior two pleural ribs bear shallow furrows directed very slightly obliquely to the anterior edge of the ribs at about one third of the distance from that edge to the posterior edge. The pleural ribs each broaden and become robust, posteriorly directed, spines; the outermost being the longest. The inner edges of the outer and middle pair of spines remains parallel to the sagittal line, with their outer edges curving gently inward, tapering to form a point. On some specimens (e.g., Fig. 4F), both inner and outer edges of the inner pair of spines are gently curved to form a more symmetrical spine, although in others the inner edge is straighter (e.g., Fig. 4G). The spines are directed parallel to the sagittal line and terminate along an essentially transverse line. The innermost pair of spines (mean = 1.41 mm,  $n = 4$ ) account for around two fifths (mean = 0.42, range = 0.36–0.46,  $n = 4$ ) of total length (mean = 3.37 mm,  $n = 4$ ) of the pygidium.

The pygidium has a varied surface sculpture on the internal mould (Fig. 4F, G) and latex casts of external moulds (Fig. 4H<sub>2</sub>), with the spines having fine and dense granulation, relatively smaller and denser than on the cranidium. A large prominent tubercle is visible in the centre of each axial

ring, similar to the occipital ring, with a row of coarse granules either side (e.g., NMW 2013.11G.11, Fig. 4G). Larger granules are also present on the pleural field, of some specimens (e.g., NMW 2013.11G.4, Fig. 4F).

*Remarks.*—*Staurocephalus oliveae* sp. nov. is most similar to *Staurocephalus clavifrons* Angelin, 1854, which was first described from the *Staurocephalus* Beds (now Ulunda Mudstone, Rawtheyan, uppermost Katian) of Älleberg, Västergötland, Sweden (cranidium figured by Angelin 1854: pl. 34: 8; refigured by Kielan 1957: pl. 4: 1; see also Kielan-Jaworowska et al. 1991: 240). It was subsequently described in detail from an equivalent horizon in the Holy Cross Mountains, Poland, by Kielan (1957) and from the overlying *Mucronaspis mucronata* Zone (“*Dalmanitina mucronata* zone” then) there (probably Hirnantian, Upper Ordovician; but see Discussion section below) (Kielan 1960). *Staurocephalus oliveae* sp. nov. differs from *S. clavifrons* in lacking genal spines, in possessing four rather than three denticles on the lateral border of the fixigena, with these denticles increasing in size anteriorly whereas they are the same size in *S. clavifrons* (see e.g., Kielan 1957: fig. 2) and, in most specimens, a greater number of denticles (13–17, mean 15, cf. 13–14) on the lateral margin of the free cheeks. Some specimens also differ in having the medial occipital tubercle posteriorly (cf. centrally) placed. The cephalic and pygidial granulation of *S. oliveae* sp. nov. is denser and finer than that of *S. clavifrons* both on internal moulds and on the external surface. *Staurocephalus oliveae* sp. nov. may differ from *S. clavifrons* in the number of lenses in the eye (120 cf. 60 recorded by Kielan 1957: 165) although the size of the librigena on which this was counted by Kielan is not known.

*Staurocephalus clavifrons* was reported or described from Ashgill (Cautleyan and Rawtheyan, upper Katian) horizons in Wales (Whittington 1965; Price 1973a), northern England (Ingham 1974, McNamara 1979) and the Republic of Ireland (Dean 1971). However, Ingham (1977) later described some better-preserved specimens from the Rawtheyan of northern England, which showed that although they were morphologically very similar, there were differences in sculpture from the type and Polish specimens. He concluded that none of the occurrences from Britain and Ireland could confidently be placed in *S. clavifrons* and should instead be assigned to *S. cf. clavifrons* or *S. aff. clavifrons*. As a result, Price (1980b: 866, pl. 110: 18) described as *S. cf. clavifrons* the material that he earlier (1973a: 245) recorded as *S. clavifrons* from throughout the Sholeshook Limestone Formation and the base of the overlying Slade and Redhill Mudstone Formation. As is noted above, with the exception of the Slade and Redhill Mudstone Formation specimens, this material is here included in *S. oliveae* sp. nov., although questioningly so given the poor preservation of the available specimens. The posterolateral part of the left side of the cranidium from the Sholeshook Limestone at Craig-y-deilo quarry figured by Price (1980b: pl. 110: 18) appears to sweep rearwards slightly as if representing the innermost proximal part of a

genal spine. However, this area is only slightly beyond the level of the palpebral lobe and is where the width (exsag.) of the border begins to expand. Examination of the specimen in the Hunterian Museum showed that the outermost part of the fixigena curves steeply downwards into the matrix and the genal area is not exposed. Price did not mention a genal spine in his description and the cranidium is similarly proportioned to ours and has a similar sculpture and is assumed to be conspecific. An historic cranidium in the Sedgwick Museum labelled *Staurocephalus cf. purchisoni* from the Sholeshook Limestone in the Sholeshook railway cutting near Haverfordwest, is within the size range of *S. oliveae* sp. nov. has similar cranidial and glabellar proportions and has similar sculpture (Fig. 6A herein). Although the genal corner is not preserved, the posterior border width (exsag.) and posterior edge of the posterior border (which does not turn sharply rearward to indicate a spine) are consistent with *S. oliveae* sp. nov. suggesting that it may represent that species.

Hammann and Leone (2007: 76) discussed Ingham’s wider concerns (1977: 89) arising from the sculpture of the material of *Staurocephalus* from the highest Rawtheyan of the Cautley area in northern England in the light of their own specimens from the Ashgill of Sardinia and concluded that the differences that Ingham highlighted were insufficient to provide any separation from *S. clavifrons*. As a result, they included the material from North Wales described by Whittington (1965) and Ireland by Dean (1971) unequivocally in the synonymy of *S. clavifrons* and that described by Ingham (1974, 1977) from the uppermost Rawtheyan of the Cautley area of northern England questioningly so. They did, however, consider that Ingham’s (1977) *S. aff. clavifrons* from the lower Rawtheyan Swindale Limestone of the nearby Cross Fell Inlier differs from Angelin’s (1854) species in its dense, large, granulated cranidial tubercles of various sizes.

Whittington (1938: 451) reported *Staurocephalus cf. purchisoni* and *Staurocephalus* sp. from the Cautleyan lower Tre-wylan Beds (“*Phillipsinella parabola* zone”) at two localities in the Llansantffraid-ym-Mechain district of the south-eastern Berwyn Hills, Mid Wales. Two of his cranidia of *Staurocephalus cf. purchisoni* (Whittington locality 42; both BIRUG 48862, on same slab) were located in the Lapworth Museum, University of Birmingham. Although both specimens lack their genal areas, precluding definitive identification, their glabellar proportions and granulation are consistent with *S. oliveae* sp. nov. and they are tentatively assigned to the new species (Fig. 6B). Also from the Berwyn Hills, the historic King collection at the Sedgwick Museum includes three incomplete cranidia (SM A39813, 39814.a and 39814.b) and one librigena (SM A39812) from the (locally) basal Ashgill mudstones (lower Rawtheyan, Ashgill Zone 5, see Price 1980a, 1981, 1982: 61) at Aber Marchnant, originally assigned to *Staurocephalus cf. purchisoni* by King (1923: 497) and, presumably, are the material listed as *S. clavifrons* from there by Whittington (1968: table 4). The cranidia are



smaller than those described here and their granulation appears coarser and less dense, but they have similar glabellar proportions and posterior margin and fixigenal corner outline (Fig. 6C, D<sub>1</sub>), although the fixigenal denticles increase in size posteriorly rather than decrease as in *S. oliveae* sp. nov. (Fig. 6D<sub>2</sub>). These cranidia are much closer to *S. oliveae* sp. nov. than to *S. clavifrons* and are herein reassigned to *S. aff. oliveae*. The librigena from the same collection possesses 12–13 marginal denticles, which would place it at the lower end of the range for *S. oliveae* sp. nov.

The absence of genal spines distinguishes *S. oliveae* sp. nov. from the other material previously described as, compared with or considered to have affinity to *S. clavifrons* from North Wales (Whittington 1965) and northern England (Ingham 1974, 1977). Dean (1971) only found one cranidium of *Staurocephalus* from the Ashgill Kildare Limestone in eastern Ireland, with no associated librigenae. *Staurocephalus oliveae* sp. nov. has a similar internal mould sculpture to Dean's (1971: pl. 19: 1–4) cranidium and is similar in outline, glabellar proportions and overall size, although the Irish specimen is marginally narrower (tr.). However, the absence of a genal spine distinguishes *S. oliveae* sp. nov. from the Irish cranidium.

While details of the lateral cephalic margin are not known sufficiently in *S. cf. clavifrons* and *S. aff. clavifrons* of Ingham (1977) from the uppermost Rawtheyan and the lower Rawtheyan (Ashgill Zone 5) respectively of northern England, the librigena of material described by Whittington (1965: 53–54, pl. 15: 4–19) from the Rawtheyan Rhiwlas Limestone in North Wales as *S. clavifrons* has 14 denticles (cf. 13–17, mean and mode = 15, in *S. oliveae* sp. nov.). In addition to lacking a genal spine, *S. oliveae* sp. nov. also differs from the Rhiwlas Limestones specimens in having smaller, more closely spaced, granules on both the internal mould and the external surface of the cephalon. *Staurocephalus cf. clavifrons* of Owen and Romano (2011) from probable Cautleyan strata in eastern Ireland is too incomplete to determine whether there is a genal spine or for the numbers of denticles on the fixed or free cheeks to be determined. The specimens have the sparser but coarser external surface granulation of typical *S. clavifrons* but the librigena (Owen and Romano 2011: fig. 2m) also shows extremely fine pits between the granules (as in *S. cf. clavifrons* of Ingham 1977); the eye contains about 70 lenses. Specimens described as *S. aff. purchisoni* Barrande by Weir (1959) from the lower Ashgill of Slieve Bernagh in western Ireland are too poorly preserved for useful comparison with any named species.

The absence of genal spines and the surface sculpture also differentiate *S. oliveae* sp. nov. from material assigned to *S. clavifrons* from Sardinia from the upper part of the Rawtheyan Punta S'Agicola Member of the Domusnovas Formation underlying the Hirnatian Rio San Marco Formation (Loi et al. 2023) described by Hammann and Leone (2007). These differences also distinguish *S. oliveae* sp. nov. from material ascribed to *S. clavifrons* by Zhou

and Zhou (1982) from Inner Mongolia from a formation correlated by them with the Rawtheyan on the basis of the trilobite fauna. There is what appears to be a tubercle at mid-length (sag.) on the occipital ring of one of Zhou and Zhou's (1982: pl. 1: 6) figured cranidia but it is difficult to discern clearly and genal denticles are not visible on either cranidium. Details of the median occipital tubercle and genal denticles are not known from the Sardinian occurrence. Shaw (2000: 385) considered that material from the Králův Dvůr Formation (upper Rawtheyan) of Bohemia previously assigned to *S. clavifrons* is too poorly preserved for detailed species comparison. Although the genal area on the single incomplete cranidium known to Marek (1964: pl. 1: 3) from here is not fully exposed, it is possible that it does not possess a spine; *S. oliveae* sp. nov. differs from it in having markedly smaller and less dense granulation. The incomplete *S. clavifrons* cranidium featured by Štorch and Mergl (1989: pl. 5: 11) from the Králův Dvůr Formation has a surface sculpture like that illustrated by Marek (1964) and the details of the outermost parts of the fixigena are not preserved.

The cranidium described by Owen and Bruton (2012) from the uppermost Caradoc of Maine, USA, as *S. cf. clavifrons* is too incomplete to determine whether it is closest to *S. clavifrons* or *S. oliveae* sp. nov., though the granulation of the external surface is much closer to that of Angelin's (1854) species than the fine, dense coverage of *S. oliveae* sp. nov.

The other named Ordovician *Staurocephalus* species comprise: *Staurocephalus pilafrons* Owen and Bruton, 1980 (uppermost Caradoc, lower Katian of the Oslo Region, Norway), *Staurocephalus djarikensis* Apollonov, 1974 (Ashgill, Katian of Kazakhstan), and *Staurocephalus elongatus* von Freyberg, 1923 (redescribed by Knüpfner 1967 from the lower Ashgill, Katian of Thuringia, Germany). *Staurocephalus oliveae* sp. nov. differs from *S. pilafrons* in having a relatively broader and shorter anterior glabellar lobe in relation to the glabellar stem, smaller granules and in lacking a large median occipital spine (although it has a prominent tubercle). *Staurocephalus pilafrons* also has a distinct eye ridge; the details of its genal angle or numbers of genal denticles are not known. The cranidial border (and therefore the genal angle) is not preserved in the material of *S. djarikensis* described by Apollonov (1974). *Staurocephalus oliveae* sp. nov. differs from *S. djarikensis* in its glabellar proportions, having a relatively wider and longer (sag.) anterior glabellar lobe taking up two thirds of the glabellar length versus around half in *S. djarikensis*, and a narrower glabellar stem than the Kazakh species. *Staurocephalus oliveae* sp. nov. differs from *S. elongatus* by having much finer granules on the anterior glabellar lobe and by having a differently proportioned anterior glabellar lobe, which is about one fifth wider (tr.) than long (sag.), compared to that of *S. elongatus*, which is about one fifth longer (sag.) than wide (tr.). Knüpfner (1967: 15–16, pl. 1: 13, 14, pl. 17: 7, 8) stated that the posterior section of the genal angle is “directed backwards”, and although he considered that it cannot be determined if there

is a genal spine present similar to *S. clavifrons*, the figured holotype (Knüpfner 1967: pl. 17: 8) shows the left posterior margin extended strongly rearwards to form a narrow point. *Staurocephalus oliveae* sp. nov. clearly differs from this species in having a posterior margin that lacks any posteriorly directed portion like that seen on Knüpfner's cranidium. Ferretti and Barnes (1997) described the conodonts from the Kalkbank Limestone from which *S. elongatus* was recovered, placing the formation at a level somewhere near the base of the *A. ordovicicus* Zone of the Katian stratigraphy that was reiterated by Kraft et al. (2023) in their recent review of the Ordovician of the Bohemian Massif. In their paper on Katian conodonts from Wales, Ferretti et al. (2014: 812) noted that: "...the Kalkbank fauna is clearly not older than the *A. ordovicicus* Zone and it seems likely that it is at least broadly coeval with the Sholeshook Limestone", making it approximately contemporaneous with *S. oliveae* sp. nov.

In addition to revising *S. clavifrons*, Kielan (1957) described specimens that she termed "*Staurocephalus* sp. b" and "*Staurocephalus* sp. c" from the Ashgill of Sweden ("*Staurocephalus* sp. a" was assigned to *S. pilafrons* by Owen and Bruton 1980: 30). *Staurocephalus oliveae* sp. nov. differs from "*Staurocephalus* sp. b" described by Kielan (1957: 169, pl. 3: 1a–c) from the Upper Jonstorp Formation (Rawtheyan) in the Siljan region of Sweden, in lacking a genal spine and in commonly having more (13–17 cf. 14) marginal spines on the librigena and four rather than three on the fixigena. The single cranidium of "*Staurocephalus* sp. c" described by Kielan (1957: 170, pl. 4: 4) from the Jonstorp Formation in Västergötland, Sweden, is too incomplete for detailed comparison but *S. oliveae* sp. nov. appears to have a relatively narrower glabellar stem. Kielan (1957: pl. 4: 5) also assigned a single pygidium to the same species; *S. oliveae* sp. nov. has noticeably shorter spines, with its innermost pair of spines accounting for just under two fifths of maximum pygidial length (sag.), while those of "*Staurocephalus* sp. c" take up just under half of maximum pygidial length (around 10% longer). *Staurocephalus* sp. of Weber (1948), from the Ashgill of Mamontovo, Kazakhstan (see Kielan 1957: 160) is only known from a single, poorly preserved, cranidium. *Staurocephalus oliveae* sp. nov. has a distinctly larger anterior glabellar lobe, which occupies two thirds of maximum glabellar length (sag.) (excluding occipital ring), while that of *Staurocephalus* sp. occupies just over a half.

The absence of a genal spine distinguishes *S. oliveae* sp. nov. from most of the Silurian species of *Staurocephalus*, such as the type species *S. murchisoni* Barrande, 1846 (Beroun, Czech Republic, Wenlock), *S. susanae* Thomas, 1981 (Dudley and Malvern, UK, Wenlock), *S. clowni* Tomczykowa, 1987 (Poland, Sheinwoodian, Wenlock), *S. oarion* Holloway, 1980 (Arkansas, USA, Sheinwoodian, Wenlock), *S. aff. oarion* Holloway, 1980 (Arkansas, USA, Wenlock), *S. lagena* Holloway, 1980 (Arkansas, USA, Whitwellian, Wenlock), *S. aff. lagena* Holloway, 1980 (USA, same lo-

cality as *S. lagena*), *S. obsoletus* (Weller, 1907) (Illinois, USA, Ludlow), *S. struszi* Chatterton and Campbell, 1980 (Canberra, Australia, Whitwellian, Wenlock) and *S. mitchelli* Chatterton and Campbell, 1980 (New South Wales, Australia, Ludlow). Both *S. struszi* and *S. mitchelli* have pygidia with proportionally smaller and flatter spines than those of *S. oliveae* sp. nov., which also has more lateral denticles on its free cheeks, 13–17, compared to "11 or 12" for *S. struszi*, and 10 for *S. mitchelli*. *Staurocephalus mitchelli* is one of the youngest species of *Staurocephalus* known, occurring in the Yarwood Siltstone Member of the Black Bog Shale, which Bicknell and Smith (2021) placed within the Ludfordian. *Staurocephalus obsoletus* is potentially of similar age, but the stratigraphical horizon (Niagaran, from Chicago Drainage Canal, near Lemont, Illinois) provided by Weller (1907) for his single specimen is too imprecise to be certain.

Another Ludfordian species, *S. azuella* Šnajdr, 1980, was described from the *Cromus beaumonti* horizon in the Kopanina Formation (see Vaněk and Valíček 2002: 52; dated as Ludlow, Ludfordian, see Lehnert et al. 2007) in the Czech Republic. Although Šnajdr (1980) listed his single specimen as an "incomplete pygidium with corroded exoskeleton", his description and illustration focus only on the pygidium, which lacks the tips of the spines. Excluding the spines, *S. oliveae* sp. nov. differs in having a relatively longer pygidium (just over four fifths as wide [tr.] as long [sag.]) than that of *S. azuella* (twice as wide [tr.] as long [sag.]) The shape and proportions of the pygidial spines are also different, those of *S. oliveae* sp. nov. being slightly wider in the middle than the base and rapidly tapering to a point, with the abaxial edge of the first two pairs of spines curving gently abaxially, while the adaxial side of the spines remains straight giving these spines a conical, slightly outwardly inflated appearance. The preserved portions of the spines of *S. azuella* are distinctly narrower (tr.) and of more uniform width, a gently tapering rectangle in outline and appear to be directed straight rearwards.

*Staurocephalus* sp. of Whiteley et al. (2002) from Rochester Shale, New York, USA (Aeronian, Llandovery) is known only from a single librigena from which that of *S. oliveae* sp. nov. can be differentiated by having more lateral denticles, there being 12 in the American species. Like *S. oliveae* sp. nov., *S. trichochin* Kobayashi and Hamada, 1985 (Japan, Whitwellian, Wenlock) lacks genal spines but *S. oliveae* sp. nov. has a relatively smaller and less inflated anterior glabellar lobe as well as relatively larger fixigenae. *Staurocephalus* sp. of Curtis and Lane (1997, 1998) from the Aeronian, Llandovery, of northern England is too incompletely known for extensive comparisons to be made, although *S. oliveae* sp. nov. can be distinguished from it by having fewer denticles on the rostral plate.

*Stratigraphic and geographic range.*—Sholeshook Limestone Formation, upper Katian, Upper Ordovician. Llandowror, Carmarthenshire and Haverfordwest area, Pembrokeshire, Wales, UK.



## Discussion

**The “bubble-headed” morphology of *Staurocephalus*.**—With its inflated cephalic lobe, *Staurocephalus* is an example of what Fortey and Owens (1997) termed a “bubble-headed trilobite”. Fortey and Owens (1997) reviewed this morphology and noted that it was developed separately and in different ways in several trilobite families from the Cambrian to the Silurian. In most instances, they considered that the “bubble” was a development of all, or just the frontal part, of the glabella. These included what they termed the “*Staurocephalus* type” in which the anterior lobe of the glabella is inflated. This widely accepted interpretation of the condition in *Staurocephalus* was challenged by Tomczykowa (1987) who argued that the bulbous feature was a development of the anterior cranial border rather than of the glabella, effectively the bubble-head condition subsequently termed the “*Nepea* type” by Fortey and Owens (1997). However, Edgecombe et al. (1988: 792) provided a detailed and convincing critique of Tomczykowa’s (1987) interpretation and supported the long-held interpretation of the bulbous structure comprising the frontal lobe of the glabella together with the cranial anterior border, the pre-cranial lobe of each librigena and the rostral plate. This interpretation was endorsed by Kielan-Jaworowska et al. (1991: 240).

**Librigenal denticle numbers through time in *Staurocephalus*.**—The large sample size of *Staurocephalus oliveae* sp. nov. has shown that the number of librigenal denticles is highly variable; something that has not previously been reported in the genus. Taking those values that have been given for other species in the literature, there is some evidence for a consistent decrease in the number of denticles through time, ranging from the mean value of 15 (range 13–17) in *S. oliveae* sp. nov. to 10 in the late Silurian species *S. susanae* and *S. mitchelli* (Fig. 8). The number of species in which the number of denticles is known is limited but the decrease seems to be unidirectional and gradual and thus there are no markedly discrete groups of stratigraphically constrained species. At face value at least, there was a period of stasis in the number of denticles from the middle Llandovery to the early Wenlock but this may be an artefact of the lack of data from the relatively long Telychian (late Llandovery).

**Implications of the new species of *Staurocephalus* for the upper Katian *Staurocephalus clavifrons* Zone.**—The *Staurocephalus clavifrons* Zone has long been recognised in Sweden (Linnarsson 1869; Tullberg 1882; Jaanusson 1963: 112–114; Nielsen et al. 2023: 294), the Danish Island of Bornholm and Poland (Kielan 1960: 7–8, 13–16). As noted by Price (1973b: 538–539), the lower parts of the zone share many species with the underlying *Eodindymene pulchra* Zone, with some possibly extending lower in the succession (see also Jaanusson 1963: 133). In Sweden (the type region)

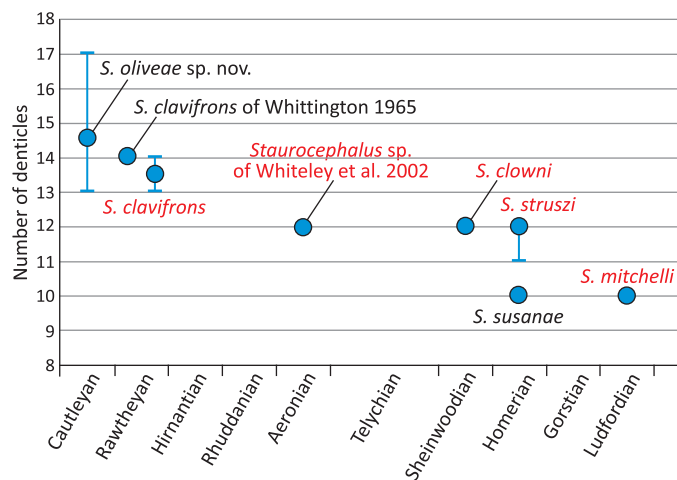


Fig. 8. Graph showing the number of lateral denticles on librigenae of various *Staurocephalus* species through time. Circle represents the mean, vertical lines the range where one was observed in that species. Chatterton and Campbell (1980) described *S. struszi* as having “twelve (possibly eleven on some specimens)” denticles. Species names in black font represent occurrences in England and Wales, those in red font occur outside Avalonia. Note that Cautleyan and Rawtheyan are regional stages from the Anglo-Welsh chronostratigraphy and belong in the upper part of the international Katian Stage. The remaining stages are international. The Telychian is approximately twice the length of the other stages (e.g., Ogg et al. 2008: fig. 1.1), hence the wider spacing allocated to it.

and on Bornholm, the *S. clavifrons* Zone is succeeded by rocks of the Regional Tommarpian (= Hirnantian) Stage and is equated with the uppermost Katian (Nielsen et al. 2023: fig. 3). It is thus essentially Rawtheyan in terms of the Anglo-Welsh regional chronostratigraphy. The same effectively applies in Poland although the zonal term is no longer applied, but what is now termed the Wólka Formation that contains the *S. clavifrons* and *E. pulchra* zone faunas of Kielan (1960) and is overlain by the Hirnantian Zalesie Formation (Trela 2021, 2023). *Staurocephalus clavifrons* in Scandinavia and Poland has not been recorded beyond its occurrence in rocks historically ascribed to the *S. clavifrons* Zone and, in Poland, the overlying Hirnantian beds containing “*Dalmanitina*” [now *Mucronaspis*] *mucronata* (see Kielan 1960: 32, 156–157; but see also Nilsson 1979 and Nielsen et al. 2023: 296, concerning the occurrence of *M. mucronata* in the *S. clavifrons* Zone in Skåne, SW Sweden).

The occurrences of *Staurocephalus* had an important role in the development of Upper Ordovician stratigraphy in Wales and northern England, including the initial formulation of the Ashgill Series by Marr (1907). This culminated in the proposal by Whittington (1968: 116–117) that the Rhiwlas Limestone of North Wales should be assigned to the *S. clavifrons* Zone. Whittington’s definition of the zone was based on an assemblage of ten trilobite species, including *S. clavifrons*, and he equated it with the shelly zones 4–7 identified by Ingham (1966) in northern England. These comprised the main part of what became the Rawtheyan Stage in the revision of the Ashgill proposed by Ingham and Wright (1970).

Price (1973b) provided a critique of the *S. clavifrons* Zone, especially its use in Wales and England together with that of the so-called *Phillipsinella parabola* fauna or zone (see Ingham 1966: fig. 2; Whittington 1968: 117). He noted that the first attempted correlation of units containing *Staurocephalus* in Britain was by Marr and Roberts (1885: 487–488) who linked its presence in the Shoeshook Limestone with the Rhiwlas Limestone and the *Staurocephalus* Zone in Sweden. Price (1973a: table 2) recorded *S. clavifrons*, from levels throughout most of the Shoeshook Limestone; a formation that he then correlated with the lower and middle parts of the Cautleyan Stage that precedes the Rawtheyan (he subsequently identified the very uppermost part as being lowest Rawtheyan [Price 1980a]). Price (1973b: 538) noted that in addition to *S. clavifrons*, the Shoeshook Limestone had yielded another four of the ten species previously listed by Whittington (1968) as comprising the assemblage characterizing the *S. clavifrons* Zone together with material probably belonging to three other species on the list and a different species from that assigned to another of the genera by Whittington (1968). Most significantly, Price (1973a: 538) stressed that all nine species in common with, or very closely allied to, Whittington's (1968) assemblage occur low in the Shoeshook Limestone in strata assigned to the lower Cautleyan (zones 1 & 2 of Ingham 1966). Price (1973b: 538) therefore considered that this discrepancy in age called into question the stratigraphical value of the *S. clavifrons* Zone.

Specimens in historical collections from all but the uppermost part of the Shoeshook Limestone are cautiously ascribed to *S. oliveae* sp. nov., thus extending its range through the Cautleyan. As noted in the Material and methods section (above) an incomplete cephalon (SM A31589) in the Marr collection from the overlying Slade and Redhill Mudstone Formation (probably lower Rawtheyan) at Pelcomb Cross, Haverfordwest, has a prominent genal spine (Fig. 6E) and therefore likely belongs in *S. clavifrons* or a closely similar species rather than *S. oliveae* sp. nov. The recognition herein that the species of *Staurocephalus* from at least the lower part of the Shoeshook Limestone (Ashgill Zone 1) and probably the entire Cautleyan, is a new species holds the prospect that *S. clavifrons* has more stratigraphical value than indicated by Price (1973a). As noted in the discussion of the new species above, the assignment of material to *S. clavifrons* from the Rhiwlas Limestone (lower Rawtheyan) by Whittington (1965) seems well founded. Specimens from northern England from the lower and uppermost Rawtheyan of the Cross Fell Inlier and Cautley areas, respectively, are closest to *S. clavifrons* although differing in their surface sculptures (see Ingham 1977; Hammann and Leone 2007 and above). Occurrences of the species in Sardinia (Hammann and Leone 2007), Inner Mongolia (Zhou and Zhou 1982) and possibly Ireland (Dean 1971) are in faunas of Rawtheyan (uppermost Katian) age as are those in Scandinavia and Poland noted above, with the possible extension of the species into the Hirnantian in

Poland. Thus, whilst many of the taxa that occur in association with *S. clavifrons* are known from lower horizons in the Ashgill, the species itself may well still have great value internationally as a Rawtheyan (i.e., uppermost Katian) and possibly lower Hirnantian indicator. If this is the case, the *S. clavifrons* Zone in Scandinavia could be defined on the basis of the first occurrence of *S. clavifrons* there and, at least, most of its range. Whether such a zone would have international application would require the precise levels of the occurrences of *S. clavifrons* elsewhere to be more tightly constrained.

As noted above, historic specimens from the lower Tre-wylan Beds collected by Whittington (1938) in the south-eastern Berwyn Hills are tentatively assigned to *S. oliveae* sp. nov. Price (1977: 764–766, pl. 98: 6–9) assigned trinucleid specimens from Whittington's (1938) locality to *Tretaspis moeldenensis* Cave, 1960 (sensu lato) Population B, (= *T. cf. kiaeri* Stømer, 1930 of Whittington 1938: 445, pl. 38: 1–3). This is consistent with the identification of *T. moeldenensis moeldenensis* from the Talfan Farm locality, confirming that the lower Tre-wylan Beds are of similar age (early Cautleyan, Zone 1, upper Katian). Thus *S. oliveae* sp. nov. may be a useful Cautleyan indicator in Wales and *S. clavifrons* may have very much wider stratigraphical value as a Rawtheyan indicator.

## Conclusions

The discovery of a new species of *Staurocephalus*, *S. oliveae* sp. nov., in the Shoeshook Limestone (equivalent to Ashgill early Cautleyan Zone 1, upper Katian, Upper Ordovician) of South West Wales represents the oldest known occurrence of the genus in Avalonia. Recognition of this Cautleyan species resurrects the potential usefulness of *Staurocephalus clavifrons* as an uppermost Katian (= Rawtheyan) and possibly Hirnantian indicator.

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## References

- Adrain, J.M. 2013. A synopsis of Ordovician trilobite distribution and diversity. In: D.A.T. Harper and T. Servais (eds.), *Early Palaeozoic Biogeography and Palaeogeography. Geological Society of London, Memoir 38*: 297–336.
- Angelin, N.P. 1854. *Palaeontologica Scandinavica I; Crustacea Formationis Transitionis, Fasc. 2* [in Russian], 21–92. Weigel, Lund.
- Apollonov, M.K. 1974. *Ashgillskie trilobity Kazakhstana*. 136 pp. Akademiya Nauk Kazakh SSR, Alma Ata.
- Barrande, J. 1846. *Notice préliminaire sur le système Silurien et les Trilobites de Bohême*. 97 pp. Hirschfeld, Leipzig.
- Barrs, C. 2013. Rugose corals from the Upper Ordovician Shoeshook Limestone of southwest Wales with an assessment of the coral affinities and biofacies. *Geological Journal* 48: 603–619.
- Bergström, S.M., Xu, C., Gutiérrez-Marco, J.C., and Dronov, A. 2009. The new chronostratigraphic classification of the Ordovician System and its relations to major regional series and stages and to the  $\delta^{13}\text{C}$  chemostratigraphy. *Lethaia* 42: 97–107.
- Bicknell, R.D.C. and Smith, P.M. 2021. Teratological trilobites from the Silurian (Wenlock and Ludlow) of Australia. *The Science of Nature* 108 (58): 1–11.
- Cave, R. 1960. A new species of *Tretaspis* from south Wales. *Geological Magazine* 97: 334–337.
- Chatterton, B.D.E. and Campbell, K.S.W. 1980. Silurian trilobites from near Canberra and some related forms from the Yass Basin. *Palaeontographica A* 167: 77–119.
- Cocks, L.R.M. 2014. The Late Ordovician brachiopods of southern Pembrokeshire and adjacent south-western Wales. *Special Papers in Palaeontology* 91: 5–89.
- Curtis, N.J. and Lane, P.D. 1997. The Llandovery trilobites of England and Wales. Part 1. *Monographs of the Palaeontographical Society* 151 (605): 1–50.
- Curtis, N.J. and Lane, P.D. 1998. The Llandovery trilobites of England and Wales. Part 2. *Monograph of the Palaeontographical Society* 152 (608): 51–101.
- Dean, W.T. 1971. The trilobites of the Chair of Kildare Limestone (Upper Ordovician) of eastern Ireland. Part 1. *Monograph of the Palaeontographical Society* 125 (531): 1–60.
- Donovan, S.K. and Paul, C.R.C. 1985. Coronate echinoderms from the Lower Palaeozoic of Britain. *Palaeontology* 28: 527–543.
- Edgecombe, G.D., Speyer, S.E., and Chatterton, B.D.E. 1988. Protaspid larvae and phylogenetics of encrinurid trilobites. *Journal of Paleontology* 62: 779–799.
- Evitt, W.R. and Tripp, R.P. 1977. Silicified Middle Ordovician trilobites from the families Encrinuridae and Staurocephalidae. *Palaeontographica A* 157: 109–174.
- Ferretti, A. and Barnes, C.R. 1997. Upper Ordovician conodonts from the Kalkbank limestone of Thuringia, Germany. *Paleontologia* 40: 15–42.
- Ferretti, A., Bergström, S.M., and Barnes, C.R. 2014. Katian (Upper Ordovician) conodonts from South Wales. *Palaeontology* 57: 801–831.
- Fortey, R.A. and Owens, R.M. 1997. Bubble-headed trilobites and a new olenid example. *Palaeontology* 40: 451–459.
- von Freyberg, B. 1923. Die fauna und gliederung der Thüringer Untersilurs. *Zeitschrift der Deutschen Geologischen Gesellschaft* 74 (for 1922): 237–276.
- Hammann, W. and Leone, F. 2007. Trilobites from the post-Sardic (Upper Ordovician) sequence of southern Sardinia. Part 2. *Beringeria* 38: 3–138.
- Harrington, H.J. and Leanza, A.F. 1957. Ordovician Trilobites of Argentina. *Special Publications, Department of Geology, University of Kansas* 1: 1–276.
- Holloway, D.J. 1980. Middle Silurian trilobites from Arkansas and Oklahoma, U.S.A. Part 1. *Palaeontographica A* 170: 1–85.
- Ingham, J.K. 1966. The Ordovician rocks in the Cautley and Dent districts of Westmorland and Yorkshire. *Proceedings of the Yorkshire Geological Society* 35: 455–505.
- Ingham, J.K. 1974. The Upper Ordovician trilobites from the Cautley and Dent districts of Westmorland and Yorkshire. Part 2. *Monograph of the Palaeontographical Society* 128 (538): 59–87.
- Ingham, J.K. 1977. The Upper Ordovician trilobites from the Cautley and Dent districts of Westmorland and Yorkshire. Part 3. *Monograph of the Palaeontographical Society* 130 (546): 89–121.
- Ingham, J.K. and Wright, A.D. 1970. A revised classification of the Ashgill Series. *Lethaia* 3: 233–242.
- Ingham, J.K. and Wright, A.D. 1972. The North of England. In: A. Williams, I. Strachan, D.A. Bassett, W.T. Dean, J.K. Ingham, A.D. Wright, and H.B. Whittington (eds.), *A Correlation of the Ordovician Rocks in the British Isles. Geological Society Special Report* 3: 43–49.
- Jaanuson, V. 1963. Classification of the Harjuan (Upper Ordovician) rocks of the mainland of Sweden. *Geologiska Föreningens i Stockholm Förhandlingar* 85: 110–144.
- Kielan, Z. 1957. On the trilobite family Staurocephalidae. *Acta Palaeontologica Polonica* 2: 155–180.
- Kielan, Z. 1960. Upper Ordovician trilobites from Poland and some related forms from Bohemia and Scandinavia. *Palaeontologica Polonica* 11: 1–198.
- Kielan-Jaworowska, Z., Bergström, J., and Ahlberg, P. 1991. Cheirurina (Trilobita) from the Upper Ordovician of Västergötland and other regions of Sweden. *Geologiska Föreningens i Stockholm Förhandlingar* 113: 219–244.
- King, W.B.R. 1923. The Upper Ordovician rocks of the south-western Berwyn Hills. *Quarterly Journal of the Geological Society* 79: 671–702.
- Knüpfel, J. 1967. Zur Fauna und Biostratigraphie der Ordoviciums (Gräfenhahler Schichten) in Thüringen. *Freiberger Forschungshefte C220*: 1–119.
- Kobayashi, T. and Hamada, T. 1985. Additional Silurian trilobites to the Yokokura-Yama fauna from Shikoku, Japan. *Transactions and Proceedings of the Paleontological Society of Japan* 139: 206–217.
- Kraft, P., Linnemann, U., Mergl, M., Bruthansová, J., Laibl, L., and Geyer, G. 2023. Ordovician of the Bohemian Massif. *Geological Society, London, Special Publications* 532: 433–464.
- Lanc, F.A., McDermott, P.D., and Paul, C.R.C. 2015. The identity of the British Ordovician cystoid “*Hemicosmites rugatus* Forbes”. *Geological Journal* 50: 1–16.
- Lehnert, O., Frýda, J., Buggisch, W., Munneki, A., Nützel, A., Kříž, J., and Manda, S. 2007.  $\delta^{13}\text{C}$  records across the Late Silurian Lau event: New data from middle palaeo-latitudes of northern peri-Gondwana (Prague Basin, Czech Republic). *Palaeogeography, Palaeoclimatology, Palaeoecology* 245: 227–244.
- Linnarsson, J.G.O. 1869. Om Vestergötlands cambriska och siluriska aflägringar. *Kongliga Vetenskaps-Akademiens Handlingar* 8: 1–89.
- Loi, A., Cocco, F., Oggiano, G., Funedda A., Vidal, M., Ferretti, A., Leone, F., Barca, S., Naitza, S., Ghienne, J.-F., and Pillola, G.L. 2023. The Ordovician of Sardinia (Italy): from the “Sardic Phase” to the end-Ordovician glaciation, palaeogeography and geodynamic context. *Geological Society, London, Special Publications* 532: 409–431.
- Marek, L. 1964. *Shumardia* Billings, 1862 a *Staurocephalus* Barrande, 1846 (Trilobita) v českém ordoviku (*Shumardia* Billings, 1862 and *Staurocephalus* Barrande, 1846 (Trilobita) from the Czech Ordovician). *Časopis Národního Muzea* 133: 153–154.
- Marr, J.E. 1907. On the Ashgillian Series. *Geological Magazine* 5 (4): 59–69.
- Marr, J.E. and Roberts, T. 1885. The Lower Palaeozoic rocks of the neighbourhood of Haverfordwest. *Quarterly Journal of the Geological Society* 41: 476–490.
- McDermott, P.D. and Paul, C.R.C. 2015. Coronate echinoderms from the Ordovician of the Llandowror area, South Wales. *Geological Journal* 50: 173–188.
- McNamara, K.J. 1979. Trilobites from the Coniston Limestone Group (Ashgill Series) of the Lake District, England. *Palaeontology* 22: 53–92.
- Nielsen, A.T., Ahlberg, P., Ebbestad, J.O., Hammer, Ø., Harper, D.A.T., Lindskog, A., Rasmussen, C.M. and Stouge, S. 2023. The Ordovician of Scandinavia: a revised regional stage classification. *Geological Society, London, Special Publications* 532: 267–315.

- Nilsson, R. 1979. A boring through the Ordovician–Silurian boundary in western Scania, South Sweden. *Sveriges Geologiska Undersökning C* 766: 1–18.
- Ogg, J.G., Ogg, G.M., and Gradstein, F.M. 2008. *A Concise Geologic Timescale*. vi+177 pp. Cambridge University Press, Cambridge.
- Owen, A.W. 2007. Trilobite diversity in Avalonia prior to the end Ordovician extinction—the peak before the trough. *Palaeogeography, Palaeoclimatology, Palaeoecology* 245: 264–271.
- Owen, A.W. and Bruton, D.L. 1980. Late Caradoc–early Ashgill trilobites of the central Oslo Region, Norway. *Palaeontological Contributions from the University of Oslo* 245: 1–63.
- Owen, A.W. and Bruton, D.L. 2012. The only known cyclopygid–“athelopic” trilobite fauna from North America: the upper Ordovician fauna of the Pyle Mountain Argillite and its palaeoenvironmental significance. *Geological Magazine* 149: 964–988.
- Owen, A.W. and Romano, M. 2011. Deep shelf trilobite biofacies from the upper Katian (Upper Ordovician) of the Grangegeeth Terrane, eastern Ireland. *Geological Journal* 46: 416–426.
- Owens, R.M. 2000. Arenig to Ashgill in South Wales. *Geological Conservation Review Series* 18: 125–180.
- Paul, C.R.C. 1973. British Ordovician Cystoids. Part 1. *Monograph of the Palaeontographical Society* 127 (536): 1–64.
- Paul, C.R.C. 1982. British Ordovician Cystoids. Part 2. *Monograph of the Palaeontographical Society* 136 (563): 65–152.
- Paul, C.R.C. 1997. British Ordovician Cystoids. Part 3. *Monograph of the Palaeontographical Society* 151 (604): 153–213.
- Peel, J.S. and McDermott, P.D. 2016. An association of problematic corals, crinoids and parasites from the Sholeshook Limestone Formation (Ordovician) of Wales. *Geological Journal* 51: 212–222.
- Prantl, F. and Přibyl, A. 1947. Roztržení některých českých cheiruridu. *Sborník Národního Muzea v Praze. Rada B. Přírodní Vědy* 3 (1): 1–44.
- Price, D. 1971. *The Stratigraphy and Trilobite Fauna of the Sholeshook Limestone of South Wales*. 256 pp. Unpublished Ph.D. Thesis, University of London, London.
- Price, D. 1973a. The age and stratigraphy of the Sholeshook Limestone of South-West Wales. *Geological Journal* 8: 225–246.
- Price, D. 1973b. The *Phillipsinella parabola*–*Staurocephalus clavifrons* fauna and Upper Ordovician correlation. *Geological Magazine* 110: 535–541.
- Price, D. 1974. Trilobites from the Sholeshook Limestone (Ashgill) of South Wales. *Palaeontology* 17: 112–116.
- Price, D. 1977. Species of *Tretaspis* (Trilobita) from the Ashgill Series in Wales. *Palaeontology* 20: 763–792.
- Price, D. 1980a. A revised age and correlation for the topmost Sholeshook Limestone Formation (Ashgill) of South Wales. *Geological Magazine* 117: 485–489.
- Price, D. 1980b. The Ordovician trilobite fauna of the Sholeshook Limestone Formation of South Wales. *Palaeontology* 23: 839–887.
- Price, D. 1981. *Tretaspis radialis* Lamont and allied species. *Geological Magazine* 118: 289–295.
- Price, D. 1982. *Calymene quadrata* King, 1923 and allied species of trilobites from the Ashgill Series of North Wales. *Geological Magazine* 119: 57–66.
- Salter, J.W. 1864. A monograph of the British trilobites from Cambrian, Silurian and Devonian formations. Part 1. *Monograph of the Palaeontographical Society* 16 (67): 1–80.
- Shaw, F.C. 2000. Trilobites of the Králův Dvůr Formation (Ordovician) of the Prague Basin, Czech Republic: *Věstník Českého geologického ústavu* 75: 371–404.
- Šnajdr, M. 1980. New Silurian trilobites from the Barrandian area Czechoslovakia. *Věstník Ústředního ústavu geologického* 55: 105–110.
- Štorch, P. and Mergl, M. 1989. Královor/Kosov boundary and the late Ordovician environmental changes in the Prague Basin (Barrandian area, Bohemia). *Sborník Geologických Věd* 44: 117–153.
- Stormer, L. 1930. Scandinavian Trinucleidae with special reference to Norwegian species and varieties. *Skrifter utgitt av Det Norske Videnskaps-Akademi i Oslo I. Matematisk-Naturvidenskapelige Klasse* 1930 4: 1–111.
- Thomas, A.T. 1981. British Wenlock trilobites. Part 2. *Monograph of the Palaeontographical Society* 134 (559): 57–99.
- Tomczykowa, E. 1987. Taksonomia Staurocephalidae Prantl et Přibyl, 1947 w związku z morfologią wyrostka przedniego. *Biuletyn Instytutu Geologicznego* 354: 183–196.
- Trela, W. 2021. Eustatic and local tectonic impact on the Late Ordovician–early Silurian facies evolution at SW margin of peri-Baltica (the southern Holy Cross Mountains, Poland). *Geological Magazine* 158: 1472–1486.
- Trela, W. 2023. Stratigraphy and sedimentary record of the Ordovician system in Poland: a review. *Geological Society, London, Special Publications* 532: 345–373.
- Tullberg, S.A. 1882. Skånes graptoliter. I: Allman ofversigt over de siluriska bildningarna i Skåne och jemforelse med ofriga kanda samtida aflagringar. *Sveriges Geologiska Undersökning C* 50: 1–44.
- Vandenbroucke, T.R.A. 2008. An Upper Ordovician chitinozoan biozonation in British Avalonia (England and Wales). *Lethaia* 41: 275–294.
- Vandenbroucke, T.R.A., Rickards, B., and Verniers, J. 2005. Upper Ordovician chitinozoan biostratigraphy from the type Ashgill area (Cautley district) and the Pus Gill section (Dufton district, Cross Fell Inlier), Cumbria, northern England. *Geological Magazine* 142: 783–807.
- Vandenbroucke, T.R.A., Williams, M., Zalasiewicz, J.A., Davies, J.A., and Waters, R.A. 2008. Integrated Upper Ordovician graptolite–chitinozoan biostratigraphy of the Cardigan and Whitland areas, southwest Wales. *Geological Magazine* 145: 199–214.
- Vaněk, J. and Valíček, J. 2002. New index of the genera, subgenera, and species of Barrandian trilobites Part C–D (Silurian and Devonian). *Palaeontologia Bohemiae* 8: 1–74.
- Walch, J.E.I. 1771. *Die Naturgeschichte der versteinungen zur erläuterung der Knorrischen Sammlung von Merkwürdigkeiten der Natur*. 235 pp. Felbeck, Nürnberg.
- Weber, V.N. 1948. Trilobites of the Silurian deposits of the USSR. Lower Silurian trilobites [in Russian]. *Monografii po Paleontologii SSSR* 69: 1–111.
- Weir, J.A. 1959. Ashgillian trilobites from Co. Clare, Ireland. *Palaeontology* 1: 369–383.
- Weller, S. 1907. The palaeontology of the Niagran Limestone in the Chicago area. *Bulletin of the Natural History Survey of the Academy of Science* 4 (2): 163–281.
- Whiteley, T.E., Kloc, G.J., and Brett, C.E. 2002. *Trilobites of New York. An Illustrated Guide*. xix + 380 pp. Cornell University Press, Ithaca.
- Whittington, H.B. 1938. The geology of the district around Llansantffraid ym Mechain, Montgomeryshire. *Quarterly Journal of the Geological Society* 94: 423–457.
- Whittington, H.B. 1965. A monograph of the Ordovician trilobites of the Bala area, Merioneth. Part 2. *Monograph of the Palaeontographical Society* 118 (504): 33–62.
- Whittington, H.B. 1968. A monograph of the Ordovician trilobites of the Bala area, Merioneth. Part 4. *Monograph of the Palaeontographical Society* 118 (504): 98–138.
- Zalasiewicz, J., Rushton, A.W.A., and Owen, A.W. 1995. Late Caradoc graptolite faunal gradients across the Iapetus Ocean. *Geological Magazine* 132: 611–617.
- Zhou, Z. and Zhou, Z. 1982. An Ashgill (Rawtheyan) trilobite faunule from Ejin Qi, Nei Monggol (Inner Mongolia) [in Chinese with English summary]. *Acta Palaeontologica Sinica* 21: 659–671.