A new Lower Cretaceous ichthyosaur from Russia reveals skull shape conservatism within Ophthalmosauromorpha

VALENTIN FISCHER*, MAXIM S. ARKHANGELSKY ‡, GLEB N. USPENSKY †, ILYA M. STENSHIN∥ & PASCAL GODOFROID ‡

*Geology Department, University of Liège, B-18, Allée du 6 Août, 4000 Liège, Belgium
‡Palaeontology Department, Royal Belgian Institute of Natural Sciences, 29, Rue Vautier, 1000 Brussels, Belgium
§Ecole Primaire, Geology Faculty, Saratov State University, Ulitsa Astrakhanskaya, 83, 410012, Saratov, Russia
∥Natural Science Museum, Ulyanovsk State University, Ulitsa L. Tolstogo, 42, 424320 Ulyanovsk, Russia
‡†Ulyanovsk Regional Museum of Local Lore named after I. A. Goncharov, 3/4, Prospekt Novy Venetz, 432601, Ulyanovsk, Russia

1. Introduction

Ichthyosaurs are Mesozoic marine reptiles that evolved a diversified array of pelagic forms, although only a handful of them have features considered as permitting deep-diving habits. One of these taxa, the Callovian–Berriasian ophthalmosaurines, are often regarded as a hyperspecialized deep diver, very little is known about the anatomy, evolutionary history and ecology of Cretaceous ophthalmosaurines because of the scarcity of the fossils and the lack of well-preserved skull material. Here, we describe the skull of a new basal ophthalmosaurine ichthyosaur, _Leninia stellans_ gen. et sp. nov., from the lower Aptian of western Russia, and compare the ocular characteristics of ophthalmosaurines. _Leninia_ is recovered as a basal ophthalmosaurine; it possesses unique traits such as a star-shaped frontal–parietal suture as well as features previously thought to be unique to _Ophthalmosaurus_ such as a supratemporal–stapes contact. A large sclerotic aperture – significantly larger than in platypterygiine ophthalmosaurids and similar to that of the largest-eyed modern animals (giant and colossal squids) – and reduced dentition appear widespread within ophthalmosaurines. This conservatism suggests ophthalmosaurine ophthalmosaurids occupied similar ecological niche(s) throughout their long evolutionary history.

Keywords: Ophthalmosauridae, Aptian, Ulyanovsk, vision, ecology.
Figure 1. Location of YKM 65931’s discovery site, Kriushi area, Ulyanovsk Region, Russia. Geographic coordinates of the site: 54°5′33″N; 48°33′25″E.

Figure 2. Stratigraphic log at YKM 65931’s discovery site, Kriushi locality, Russia. As the specimen was found in a limestone nodule on the riverbank, its position within the stratigraphic column is tentative.

2.b. Geography and Stratigraphy

The described material was found on the right bank of the Volga River, 2 km southeast from Kriushi village, Sengiley district of Ulyanovsk Region, Russian Federation (Fig. 1). Only a small part of the lower Aptian is exposed in this locality and is contained within the Deshayesites volgensis Zone (Baraboshkin & Mikhailova, 2002) (Fig. 2). The section is described in the online Supplementary Material at http://journals.cambridge.org/geo.

The specimen YKM 65931 was not found in situ but embedded in a large limestone nodule lying on the shoreline. Field observations by I. M. Stenshin & G. N. Uspensky suggest this nodule is similar to those of the ‘upper nodule bed’ and would thus belong to the lower A. matheronianum Zone. This age, based on the biostratigraphy of heteromorph ammonites, is, however, tentative; therefore, we date this specimen with no further precision than Deshayesites volgensis Zone, because the whole section is contained within this zone in the Kriushi area.

2.c. Phylogeny

We coded Leninia in the phylogenetic matrix of Fischer et al. (2012) with one additional character (see online Supplementary Material at http://journals.cambridge.org/geo). However, only 23.07% of the characters (12/52) could be coded owing to the absence of teeth and postcranial skeleton in the holotype. We used exact parsimony searches of TNT v1.1 (Goloboff, Farris & Nixon, 2010) to analyse the character matrix (exact algorithm, 10,000 trees in memory) and calculate the Bremer support, Jacknife (removal probability of 36, 1000 replications) and bootstrap (standard, 1000 replications) values.

We generated the phylogenetic tree using Winclada v.0.9 (Nixon, 1999) (detailed character state distribution in all optimizations, Bremer support, bootstrap and Jacknife values are available in Figs S1–S2 in the online Supplementary Material at http://journals.cambridge.org/geo). Characters were not weighted and, except for characters 17, 39 and 45, were not ordered (see Fischer et al. 2012 for an explanation of these characters).

2.d. Optics

We attempted to apply the calculations and methods of Motani, Rothschild & Wahl (1999) and Fernández et al. (2005) to assess the visual acuity and ecology of Leninia. We calculated precise orbital, sclerotic and sclerotic aperture areas using a plugin for the freeware program Inkscape v0.48.2 (Albert et al. 2012) on high-quality photographs. Then, rather than simply making an average of the dorsoventral and anteroposterior lengths, we recalculated precise diameters from the measured areas, assuming that the measured areas were those of circles ($d = 2\sqrt{\text{area}/\pi}$), in order to cope with diagenetic deformations. We used a similar technique to calculate the eye parameters of Sveltonectes insolitus (IRSNB R269). Because the skull of Leninia is crushed...
transversely, the axial length of the eye (a parameter in the estimation of the minimum f-number; Motani, Rothschild & Wahl, 1999) could not be assessed. We refrained from estimating this variable, as we feel the method used by Motani, Rothschild & Wahl (1999) is too subjective and hardly replicable since there is no constraint on the placement of the eyeball in the coronal plane. The suggestion of Fernández et al. (2005) to use the bending of the sclerotic plates as a proxy for eye size is also problematic because it involves a spherical eye, which is probably not the case in ichthyosaurs (Motani, Rothschild & Wahl, 1999; Maisch & Matzke, 2003; Fernández et al. 2005). Therefore, we feel that the minimum f-number cannot be satisfactorily calculated in ichthyosaurs yet. Moreover, it should be noted that Motani, Rothschild & Wahl (1999) examined a single specimen of O. natans (CM 878) to calculate the eye parameters but incorporated several specimens of O. icenicus (from the NHMUK) to analyse avascular necrosis. This is questionable, given the fact that the most recent phylogenies of ophthalmosaurids (including this one) do not recover Ophthalmosaurus as monophyletic (Druckenmiller & Maxwell, 2010; Fischer et al. 2012). Accordingly, we simply compare the absolute sclerotic aperture of ophthalmosaurids as a proxy for dim light vision but we do not assess their deep-diving habits.

3. Systematic palaeontology

Order ICHTHYOSAURIA Blainville, 1835
Family OPHTHALMOSAURIDAE Baur, 1887
Subfamily OPHTHALMOSAURINAE Baur, 1887 sensu Fischer et al. 2012
Leninia stellans gen. et sp. nov.
Figures 3–5

Holotype. YKM 65931, an articulated but incomplete skull preserved in three-dimensions.

Type horizon and locality. Deshayesites volgensis Zone, lower Aptian, Lower Cretaceous of the Kriushi locality, Sengiley district, Ulyanovsk Region, Russia.

Etymology. The museum where YKM 65931 is housed is located within the Lenin Memorial and Lenin school complex in Ulyanovsk; accordingly, the generic name reflects the geohistorical location of the find. The specific name refers to the peculiar frontal–parietal suture, which is star-shaped in the holotype.

Diagnosis. Leninia stellans is characterized by the following potential autapomorphies within Ophthalmosauridae: posterior process of the maxilla extending as far as the middle orbit (reminiscent of Ichthyosaurus intermedius; Maisch, 1997); presence of an anterior process on the prefrontal covering the nasal; frontal–parietal suture formed by the means of elongate medial forked processes of the frontal and the parietal; presence of an anterolateral process of the supratemporal contacting the parietal. Leninia stellans is also characterized by a combination of features shared by some other ophthalmosaurids (see Description, Section 4).

Note. Additional remains resembling the holotype of Leninia have been found in other Aptian localities near Ulyanovsk and Saratov; these fossils (UISI 115, a partial braincase; SSU 104a/28, a partial skull; SSU 14/53, a partial skull) are, however, fragmentary and cannot be determined unambiguously at the present time.

4. Description

4.a. Skull roof

4.a.1. Maxilla

The posterior end of the maxilla is preserved, but the narial lamella and its possible contribution to the narial aperture cannot be described because of the poor preservation of that region. The anterior cross-section suggests that the maxillary dental groove is extremely shallow relative to the size of the skull (height: 16 mm). Posteriorly, the maxilla forms a long palatal plate ventral and medial to the jugal. Unusually, this plate extends at least up to the middle-orbit level, whereas it ends more anteriorly in other ophthalmosaurids (e.g. Ophthalmosaurus icenicus, Platypterygius australis; Kear, 2005; A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983). The medial part of that plate is broken and pushed dorsally into the orbit by the dorsal surface of the mandible (Fig. 3).

4.a.2. Lacrimal

The posteroventral process of the lacrimal lies on the dorsal surface of the palatal plate of the maxilla. The lacrimal extends dorsally above the level of the narial aperture and contacts both the prefrontal and the nasal. The nasal–lacrimal contact is elongated and the lacrimal forms the posterior edge of the naris; therefore, it excludes the prefrontal from the narial aperture, unlike in Aegirosaurus and Sveltonectes (Bardet & Fernández, 2000; Fischer et al. 2011b, respectively).

4.a.3. Jugal

The anterior process and the shaft of the jugal are extremely reduced in dorsoventral height (shaft is 13 mm high at middle orbit). Anteriorly, it forms a transversely compressed ramus covering the lateral surface of the maxilla. The shaft is rounded in cross-section. The posterior plate of the jugal is well developed and expands dorsally up to the middle-orbit level, even more than in O. icenicus, and as in Mollesaurus (Fernández, 1999).
4.a.4. Postorbital

The postorbital is of usual shape, with a prominent dorsal orbital rim and a posteroventral plate articulating with the quadratojugal and the jugal. As in *Ophthalmosaurus* (e.g. Andrews, 1910), the postorbital is reduced in anteroventral length (44 mm) with respect to the size of the orbit (anteroposterior length $\approx 265$ mm).

Figure 3. (Colour online) Photograph and interpretation of the holotype specimen of *Leninia* (YKM 65931) in lateral view.
4.a.5. Quadratojugal

The quadratojugal is covered by all elements of the postorbital region. Nevertheless, it appears more robust and longer anteroposteriorly than in *Ophthalmosaurus icenicus* (Andrews, 1910; A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983). It articulates dorsally and anterodorsally with the squamosal. The anterior surface of the quadratojugal is thin, forming a concave lateral articular surface beneath the postorbital, the squamosal and the jugal. An angle is present between the squamosal and the postorbital/jugal facet, as in *P. hercynicus* (Fischer, 2012), but it is more conspicuous in *Leninia*. Posteroventrally to these large facets, the quadratojugal thickens and becomes pillar-shaped. Posteroventrally, this pillar slightly extends posteriorly to form the processus quadratus. As described by Kirton in *O. icenicus* (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983), there is a concave area for ligamentous attachment dorsomedially to the processus quadratus.

4.a.6. Squamosal

The squamosal is present, unlike in *P. australis* (Kear, 2005) and *P. americanus* (Romer, 1968). It is, however, incomplete; only its posteroventral part is preserved, but its general shape can be assessed because it covers numerous bones of the postorbital region (postfrontal, postorbital, quadratojugal), leaving a shallow facet textured by a series of subtle dorsoventral ridges. However, we refrained from coding character 14 (squamosal shape) for YKM 65931, but instead put the polymorphism 0/1 (as state ‘2’ stands for the absence of a squamosal). The squamosal covers the postfrontal extensively and the posterodorsal lamella of the postorbital. The squamosal–quadratojugal contact is not conspicuous.

4.a.7. Nasal

The nasal participates in the dorsal edge of the naris and presumably contacts the lacrimal over a long distance since the prefrontal is excluded from the nasal aperture (see below). Dorsally to the naris, the nasal forms a subtle lateral wing as in *Acamptonectes* (Fischer et al. 2012) and some derived platypterygiines (*P. australis, P. bannovakensis*; Kear, 2005 and V. Fischer, pers. obs., respectively). As in *Acamptonectes* (Fischer et al. 2012), the prefrontal facet is set directly posteriorly to the lateral wing, but there is no clear evidence for a foramen in that region. The posterior part of the nasal is medially incomplete, but the nasal seems to be excluded from the internasal foramen by the frontal.

4.a.8. Prefrontal

The prefrontal forms a thick anterior process constricting the nasal, unlike in other ophthalmosaurids (e.g. Andrews, 1910; Kear, 2005) where the prefrontal is restricted to the anterodorsal orbital rim. The lacrimal–prefrontal suture is straight, unlike in *Simbridskaiaouras* (Ochev & Efimov, 1985; V. Fischer, pers. obs.), *O. icenicus* (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983) and possibly *Sveltonectes* (Fischer et al. 2011b); this suture is placed lower than in *Athabascasaurus* (Druckenmiller & Maxwell, 2010) and higher than in *Aegirosaurus* (Bardet & Fernández, 2000).

4.a.9. Frontal

The frontal is roughly triangular. It encloses the so-called internasal foramen. A long posterior process interdigitating with the forked process of the parietal forms the anterior margin of the parietal foramen (Fig. 4). The frontal is strongly reduced laterally and does not participate in the anterior margin of the supratemporal fenestra unlike in many platypterygine ophthalmosaurids (Fischer et al. 2011b; Fischer, 2012), but as in *Athabascasaurus* (Druckenmiller & Maxwell, 2010) and *Ophthalmosaurus* (Gilmore, 1906; A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983).

4.a.10. Postfrontal

The anterior part of the postfrontal contacts the nasal over a short distance whereas its anteromedial part contacts the frontal over a long distance. As in *Ophthalmosaurus* (Gilmore, 1906; A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983) and unlike in *Sveltonectes* (Fischer et al. 2011b), *P. australis* (Kear, 2005), *P. hercynicus* (Fischer, 2012) and *Athabascasaurus* (Druckenmiller & Maxwell, 2010), the postfrontal does not form a Y-shaped anterior process. The postfrontal is excluded from the lateral margin of the supratemporal fenestra by an elongated anteromedial process of the supratemporal, which constitutes an autapomorphy. However, the postfrontal seems to contact the parietal internally, as suggested by the broken supratemporal on the right side. Medially, the postfrontal forms a long process interdigitating with the supratemporal. The posterolateral part of the postfrontal is sheet-like and contacts the postorbital laterally and the squamosal posteriorly.

4.a.11. Parietal

The anterior margin of the parietal forms a forked process receiving the posterior process of the frontal (Fig. 4). The parietal contacts the anteromedial process of the supratemporal, a unique condition among ichthyosaurs. Both pariets are crushed against each other medially; there is, however, no evidence of a parietal crest. The posteroomedial notch, receiving the dorsal surface of the supraoccipital, is markedly concave, even more than in *Athabascasaurus* (Druckenmiller & Maxwell, 2010); however, the moderate lateral crushing of the skull does not permit satisfactory assessment of the shape of this notch in YKM.
6 V. FISCHER AND OTHERS

Figure 4. (Colour online) Photograph and interpretation of the holotype specimen of *Leninia* (YKM 65931) in dorsal view.

65931. The parietal is covered posterolaterally by the supratemporal.

4.a.12. Supratemporal

The supratemporal is extensive and forms the entire lateral and posterior edges of the supratemporal fenestra. Anteromedially, the supratemporal forms a long finger-like process contacting the anterolateral part of the parietal. A similar process is present in *O. icenicus* (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983), but it does not contact the parietal anteriorly in this taxon. Laterally to that process, the supratemporal receives the posteromedial
finger-like process of the postfrontal. Ventrolaterally to that zone, the supratemporal is sheet-like and covers the postfrontal. It is, however, separated from the postorbital by the postfrontal and the squamosal, unlike in Caypullisaurus, where the supratemporal contacts the postorbital (Fernández, 2007). A squamosal facet is present posterolaterally. Posteriorly, the supratemporal is expanded, closing the dorsolateral part of the basicranium. A few foramina are present. Posteroventrally, the supratemporal forms a process contacting the stapes (Fig. 5), a feature previously regarded as unique to O. icenicus (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983) and O. natans (Gilmore, 1906).

4.a.13. Pterygoid

Both pterygoids are present but can only be seen in cross-section slightly posterior to the dorsal lamella and are distorted by the lateral compression of the skull. The lateral lamella is thin and extensively covers the quadratojugal.

4.b. Basicranium

4.b.1. Basioccipital

The extracondylar area of the basioccipital is partly reduced, and forms a wide concave area external to the articulating condyle (Fig. 5), which is an ophthalmosaurusine synapomorphy (Fischer et al. 2012). As in adult Acamptonectes but unlike in Mollesaurus (Fernández, 1999) and Ophthalmosaurus (Fischer et al. 2012), this groove is continuous ventrally, although a ventral notch is present. There is a depression anterior to the peripheral edge of the concave area, as in Acamptonectes (Fischer et al. 2012) and Ophthalmosaurus (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983). The occipital condyle is bulbous, slightly deflected peripherally, and the notochordal pit is set in the dorsal half of the condyle. Unfortunately, the dorsal surface of the basioccipital is not available for study, so the shape of the floor of the foramen magnum, which may be diagnostic in ophthalmosaurids (Fischer et al. 2012), cannot be assessed on this specimen.

4.b.2. Stapes

As in Acamptonectes, the occipital head of the stapes is markedly expanded dorsoventrally (≈53 mm high) and the shaft is a slender rod (≈ 25 mm high) (Fig. 5). A large hyoid process is present, but this feature appears variable in ophthalmosaurids (Fischer et al. 2012). The quadrate head of the stapes is not preserved.

4.b.3. Quadratojugal

The quadrate is ear-shaped as in most ophthalmosaurids. An occipital lamella is present, unlike in
**P. hercynicus** (Kolb & Sander, 2009; Kuhn, 1946) and some specimens referable to *Platypterygius* sp. (V. Fischer, pers. obs.).

### 4.c. Lower jaw

The posterior part of the lower jaw is preserved but lacks distinctive or diagnostic features. A coronoid process is present, although the presence of this process varies intraspecifically in some ophthalmosaurids (Fischer et al. 2012). The fossa surangularis is present but extremely reduced. As in all ophthalmosaurids (Motani, 1999), the angular is markedly exposed laterally.

### 4.d. Sclerotic ring

The left sclerotic ring is complete and almost undistorted. It is composed of 14 intermeshing trapezoidal plates. Each individual plate bears subtle striations on its lateral surface, and the internal edge is crenulated. The external edge is bent medially. The sclerotic aperture is moderately small compared to the orbital area (11.03 %, see Section 5), suggesting that the specimen was mature (Fernández et al. 2005).

### 5. Results and discussion

#### 5.a. Phylogenetic position

The cladistic analysis (a single most parsimonious tree, 103 steps long, consistency index: 0.55, retention index: 0.67) recovers *Leninia* as a basal ophthalmosaurus ophthalmosaurid, forming a polytomy at the base of the subfamily with the Bajocian taxon *Mollesaurus perialus* (Fig. 6). This polytomy is actually an ‘irresolvable’ node, because there is no character state in the dataset that could decipher which taxon is more derived than the other. In other words, the branch separating *Leninia* from *Mollesaurus* is of length = 0. This uncertainty and the surprisingly basal position of *Leninia* among ophthalmosaurines are possibly due to the large amount of missing data for both *Mollesaurus* and *Leninia*. Moreover, the skull of the only other Cretaceous ophthalmosaurine, *Acamptonectes*, is poorly known, preventing detailed comparison with that of *Leninia*. The large amount of missing data of the basal ophthalmosaurus *Mollesaurus perialus* is also the main reason why the clade Ophthalmosaurinae is supported by a very small number of unambiguous synapomorphies in the analysis undertaken by Fischer et al. (2012). The inclusion of *Leninia* in the phylogenetic analysis does not modify this scheme as it adds a large number of uncertainties on postcranial anatomy as well. Additionally, while the exclusion of *Mollesaurus* from the data matrix provides additional unambiguous synapomorphies to diagnose the clade Ophthalmosaurinae, it does not improve the support of the tree, quite the contrary. Nevertheless, our analysis adds an unambiguous and non-homoplastic synapomorphy uniting ophthalmosaurines: the presence of a supratemporal–stapes contact (character state 52.1; see online Supplementary Material at http://journals.cambridge.org/geo for optimizations).

#### 5.b. Evolutionary history of ophthalmosaurines

Only two groups of marine animals evolved a very large eye size: the modern giant and colossal squids and ichthyosaurs (Nilsson et al. 2012). The size of the eye is related to ecology, where large eyes usually indicate adaptations for feeding and/or predator detection in low light setting, such as the deep marine realm (Motani, Rothschild & Wahl, 1999; Humphries & Ruxton, 2002; Fernández et al. 2005; Nilsson et al. 2012). Whereas relative eye size is linked to ontogeny and/or possible evolutionary mechanisms such as paedomorphosis (Fernández et al. 2005), absolute eye size determines visual acuity (Motani, Rothschild & Wahl, 1999; Humphries & Ruxton, 2002; Fernández et al. 2005; Nilsson et al. 2012). The absolute size of the sclerotic aperture is a good estimation of the size of the cornea (and thus the dilated pupil) and is likely to remain rather constant during ontogeny (Fernández et al. 2005). Because it approaches the size of the cornea, the sclerotic aperture can give clues on the ability to see in low light conditions. Using this variable, among others, Motani, Rothschild & Wahl (1999) postulated that *Ophthalmosaurus* could dive to depths in excess of 500 m and still rely on vision to hunt for prey or detect predators.
These extreme adaptations of *Ophthalmosaurus* are considered as exceptional among ichthyosaurs (Bakker, 1993; Fernández et al. 2005). However, numerous ichthyosaurs closely related to *Ophthalmosaurus*, ophthalmosaurine ophthalmosaurids, are now known in strata ranging from the Bajocian to the Albian (Fernández, 1999; Fischer et al. 2012), and it is possible that a large sclerotic aperture is characteristic for the entire ophthalmosaurine clade; the peculiar anatomy of *Ophthalmosaurus* would therefore not be an exception but characteristic of a long-lived clade of ichthyosaurs. The skull of *Leninia* appears similar to that of *Ophthalmosaurus* or *Mollesaurus* in overall architecture as well as in morphological details, suggesting that ophthalmosaurines kept a similar skull shape and visual acuity throughout their evolution. To test this, we looked at the eyes and dental groove of *Leninia*, and compared them to those of other ophthalmosaurids.

When the sclerotic aperture diameter is plotted against the sclerotic area diameter (both relative to orbital area as in Fernández et al. 2005; see Fig. 7), *Leninia* and *Sveltonectes* fall outside the convex polygon encompassing adult and supposedly non-deep-diving parvipelvian ichthyosaurs and is located in between this group and the convex polygon grouping juveniles and ‘deep-diving’ forms (*Ophthalmosaurus* and *Eurhinosaurus*, according to Fernández et al. 2005). The interpretation of this result is hazardous because two factors, ontogeny and ecomorphology, affect the position on the graph. Because the specimens of *Sveltonectes* and *Leninia* used here appear osteologically mature, this graph suggests that the relative eye size of these taxa was in between ‘deep-diving’ and ‘non-conclusively deep-diving’ forms. However, as mentioned above, the absolute size of the sclerotic aperture is far more important to assess dim light vision than is the relative size. This remark is also important for *Mollesaurus*, which is set on the bottom-left corner of the ‘adult parvipelvian’ group, yet possesses a sclerotic aperture similar to other ophthalmosaurines (see below).

The absolute size of the sclerotic ring of *Leninia* (area of 252.5 cm², diameter of 17.93 cm) is among the largest ever measured in ichthyosaurs, although much smaller than those of *Temnodontosaurus* (23.5 cm) and *O. natans* (22 cm) (Motani, Rothschild & Wahl, 1999). The absolute size of the sclerotic aperture of *Leninia* is 7.74 cm long in diameter with a total area of 47.07 cm²; this falls close to *Mollesaurus* (diameter: 7.06 cm; Fernández et al. 2005). Fernández et al.’s dataset of absolute sclerotic aperture diameters does not contain adult specimens of *Ophthalmosaurus*; similarly, Motani, Rothschild & Wahl (1999) do not provide the sclerotic aperture of the specimens they measured. Nevertheless, we recorded a sclerotic aperture diameter of slightly below 10 cm in photographs of *O. natans* (CM 878, from Fernández et al. 2005; UW 24816, from Wahl, 2009) and of 7.11 cm in Kirton’s (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983) detailed reconstruction of *Ophthalmosaurus icenicus*. Whereas these calculations are rough, they suggest that the sclerotic aperture of both *Mollesaurus* and *Leninia* fall within the range of those of the largest-eyed animals ever, *Ophthalmosaurus* and the modern giant squid *Architeuthis* (Nilsson et al. 2012), and appears significantly higher than in most ptycterygine ophthalmosaurids (Fig. 8): *Sveltonectes*: 3.44 cm; *Platypterygius australis*: 3.15 cm (calculated on a photograph of AM F116939 from Kear, 2005); *Caypullisaurus bonapartei*: 5.05 to 5.69 cm (adults; Fernández et al. 2005). Moreover, it should be noted that of platypterygines, regardless of total size, and is within the range of that of the modern giant squid (Nilsson et al. 2012).

---

**Figure 7.** (Colour online) Sclerotic ring and sclerotic aperture diameters of various parvipelvin ichthyosaurs relative to the orbit diameter, modified from Fernández et al. (2005). Grey areas represent the convex polygon encompassing the two groups in the Fernández et al. paper: juveniles + supposed deep divers (*Ophthalmosaurus*, *Eurhinosaurus*) versus other adult parvipelvians. Both *Sveltonectes* and *Leninia* fall outside these groups, despite great differences in terms of absolute sclerotic apertures. This suggests parvipelvians actually display a continuum of relative sclerotic dimensions that may evolve with age, as suggested by Fernández et al. (2005).

**Figure 8.** Sclerotic aperture diameter of ophthalmosaurids. Vertical bars on the left denote two groups: ophthalmosaurine (top) and platypterygine (bottom) ophthalmosaurids. The aperture of ophthalmosaurines is usually markedly higher than that of platypterygines, regardless of total size, and is within the range of that of the modern giant squid (Nilsson et al. 2012).
that the theoretical modelling undertaken by Nilsson et al. (2012) indicated that the range of vision is roughly similar with a 7 or 10 cm wide pupil, but that such a pupil size provides a vision range increased by 10–20 m (depending on depth and the object to detect) when compared to the already large pupils of platypterygiine ichthyosaurs.

No teeth are preserved within the holotype of Leninia. However, the minute depth of the maxillary dental groove (16 mm high) at the level of the naris suggests a markedly reduced dentition (in acipinbasal size) for this moderately large taxon, at least in this part of the snout. A similarly reduced dentition is present in other ophthalmosaurines such as in Ophthalmosaurus natans (Gilmore, 1902, 1905, 1906; V. Fischer, pers. obs. on CM material), Acamptonectes densus (Fischer et al. 2012) and Mollesaurus (Fernández, 1999). The dentition of Ophthalmosaurus icenicus is subject to debate. This taxon has widely been considered as edentulous in the past (e.g. McGowan, 1976), but this was due to the weak attachment of teeth within the groove, as numerous specimens of this taxon are preserved with teeth, even adults (A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983; Zammit, 2012; V. Fischer, pers. obs. on GLAHM and MJML material). Interestingly, teeth of O. icenicus do not appear as small and slender as in other ophthalmosaurines, which may indicate a slightly different diet for this taxon.

Ophthalmosaurines appear conservative in cranial anatomy; large eyes and reduced dentition probably appeared early in the evolution of the group and were apparently conserved throughout their long history (Middle Jurassic – Early Cretaceous). Similar sclerotic apertures and absence of profound modifications in tooth size and skull shape suggest that ophthalmosaurines conserved similar ecological niche(s), as colonization of a particular ecological niche is usually linked to numerous craniodental modifications, as recently demonstrated for marine crocodyliforms (e.g. Pierce, Angelczyk & Rayfield, 2009; Young et al. 2012). Moreover, the metabolic cost of growing and maintaining eyes of such size is so high that it is extremely unlikely to be a plesiomorphic feature devoid of eco/ethological significance (Nilsson et al. 2012). On the contrary, the other ophthalmosaurid subfamily, Platypterygiinae, evolved a much more disparate assemblage of skull shapes and feeding guilds within the same time interval, with small piscivorous/teuthophagous forms (Sveltonectes: Fischer et al. 2011b), large opportunistic predators (Platypterygius: Kear, Boles & Smith, 2003; Brachypterygius: A. Kirton, unpub. Ph.D. thesis, Univ. Newcastle upon Tyne, 1983; McGowan, 1976) as well as forms in between these extremes (Aegirosaurus: Fischer et al. 2011a), but seemingly lacked the deep-diving/low light vision adaption of the eye that is found in ophthalmosaurines. Additional and independent methods to evaluate deep-diving behaviour such as analyses of bone histology and avascular necrosis (Motani, Rothschild & Wahl, 1999; Rothschild, Xiaoting & Martin, 2012) could be useful to further investigate the evolutionary trends outlined here.

6. Conclusion

A new ichthyosaur, Leninia stellans, from lower Aptian deposits of Russia highlights a remarkable conservatism of skull shape and eye size among ophthalmosaurine ophthalmosaurids throughout their long history (Middle Jurassic – Early Cretaceous), suggesting these ichthyosaurs kept similar ecological niche(s) during this time interval. These similarities are not biased by phylogeny because the features like eye architecture and dentition are not linked to the characters used in the cladistic analysis. However, the fragmentary nature of the basal ophthalmosaurines Mollesaurus and Leninia does not permit clarification of the tempo of character acquisition at the base of this subfamily. Nevertheless, this contribution highlights distinct evolutionary histories for the ophthalmosaurid subfamilies, by indicating that the peculiar morphology of Ophthalmosaurus – often considered as a hyperspecialized exception – is actually found within an increasing number of closely related Jurassic and Cretaceous ichthyosaurs.

Acknowledgements. We thank the staff of the YKM for making our study as easy as possible, and Erin Maxwell and one anonymous reviewer for their constructive and insightful comments. We also thank Ben Kear for the invitation to contribute to the present volume. This work was funded by a Fond de la Recherche Scientifique doctoral grant (V.F., Aspirant du F.R.S.–FNRS).

References

**Conservatism within Ophthalmosaurinae**


Supplementary Material

1. Stratigraphy

In the Ulyanovsk region, heteromorph ammonites of the family Ancyloceratidae can also be used as a biostratigraphic tool. The section can be described as follows:

The lower part (first 4 meters) of this section is composed of dark shale of the *Volgoceratoïdes schilovkensis* Zone and contains large and closely spaced carbonate nodules. Large wood fragments, broken shells and small phosphate concretions are also present at the base of the section, as well as possible erosional surfaces. A bed of smaller nodules is present in the upper part of the *V. schilovkensis* Zone. Alternating light and dark layers (1 to 5 mm thick) textures the shale. A large number of flattened ammonites aptychi and fish scales are present along some bedding planes. Some surfaces are almost completely covered by embryonic shells of ammonites. The following monomorph ammonites are present in this part of the section: *Deshayesites gracilis* Casey, 1964, *D. volgensis* Sasonova, 1958, *D. forbesi* Casey, 1961,

The following heteromorph ammonites are present in this part of the section:
Volgoceratoides schilovkensis Michailova and Baraboshkin, 2002 and Koeneniceras tenuiplicatum (Koenen, 1902).

A 3 meters-thick unit of homogeneous grey shale of the Ancyloceras matheronianum Zone overlies the dark shale of the V. schilovkensis Zone.

The shale contains scattered shell detritus at the bottom, followed by a bed of large to very large (up to 3 meters in diameter) carbonate nodule. The following monomorph ammonites are present in this part of the section:
Deshaysites multicostatus Swinnerton, 1953, D. consubrinoides (Sinzow, 1898), D. sp., Paradeshaysites sengillyensis (Sasonova, 1958), P. similis (Bogdanova, 1991), and P. imitator (Glasunova, 1968), among others. The following heteromorph ammonites are present in this part of the section:
Ancyloceras matheronianum d’Orbigny, 1842, Lithancylus aff. grandis (Sowerby, 1829), and L. glebi Michailova & Baraboshkin, 2001.

2. Phylogeny

2.1. New character

We added the new character (n°52) at the end of the character list of Fischer et al. (2012) so that the numbering of characters remains unchanged.

52. *Supratemporal–stapes contact: absent (0); present (1).

2.2. Coding of each taxa for character 52
2.3. Character coding for *Leninia stellans*

???????? 10?01 10?0/11 ?1?0? ?1???? ???? ?? ???? ??? ???? ???? 
???? ?1

2.4. Results

The bootstrap (standard bootstrap, 1000 replications) and Jacknife (removal probability of 36, 1000 replications) values are low: only the clade...
Ophthalmosauridae is recovered with values above 50: bootstrap = 54; Jacknife = 53.

Figure S1. Single most parsimonious tree recovered from the cladistic analysis, in unambiguous optimization.
extensive ichthyosaur survival across the Jurassic–Cretaceous boundary. New ophthalmosaurids from the Early Cretaceous of Europe demonstrate extensive ichthyosaur survival across the Jurassic–Cretaceous boundary. *PLoS ONE* 7(1), e29234.

**References**