

The first gladius-bearing coleoid cephalopods from the Lower Toarcian “Schistes Cartons” Formation of the Causses basin (southeastern France) (#91410)

1

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The first gladius-bearing coleoid cephalopods from the Lower Toarcian “Schistes Cartons” Formation of the Causses basin (southeastern France)

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The fossil record of gladius-bearing coleoids is scarce and based only on a few localities with geological horizons particularly favourable to their preservation (the so-called Konservat-Lagerstätten), which naturally leads to strongly limited data on geographical distributions. This emphasizes the importance of every new locality providing gladius-bearing coleoids. Here, we assess for the first time the coleoid taxonomic diversity within the Lower Toarcian “Schistes Cartons” of the Causses basin (southeastern France). The material includes two fragmentary gladii, identified as *Paraplesioteuthis sagittata* and *?Loligosepia* sp. indet. Just with these two specimens, two (Prototeuthina and Loligosepiina) of the three (Prototeuthina, Loligosepiina and Teudopseina) suborders of Mesozoic gladius-bearing coleoids are represented. Thus, our results hint at an unexpected early Toarcian coleoid diversity in the Causses basin and point out the need for further field investigations in the Lower Toarcian “Schistes Cartons” in this area. This new record of *Paraplesioteuthis sagittata* is only the second one in Europe and the third in the world (western Canada, Germany and now France). Based on these occurrences, we suggest that *P. sagittata* originated in the Mediterranean domain and moved to the Arctic realm through the Viking Corridor to eventually move even farther to North America.

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18

19 Abstract

20 The fossil record of gladius-bearing coleoids is scarce and based only on a few localities with
21 geological horizons particularly favourable to their preservation (the so-called Konservat-
22 Lagerstätten), which naturally leads to strongly limited data on geographical distributions. This
23 emphasizes the importance of every new locality providing gladius-bearing coleoids. Here, we
24 assess for the first time the coleoid taxonomic diversity within the Lower Toarcian “Schistes
25 Cartons” of the Causses basin (southeastern France). The material includes two fragmentary
26 gladii, identified as *Paraplesioteuthis sagittata* and ?*Loligosepia* sp. indet. Just with these two
27 specimens, two (Prototeuthina and Loligosepiina) of the three (Prototeuthina, Loligosepiina and
28 Teudopseina) suborders of Mesozoic gladius-bearing coleoids are represented. Thus, our results
29 hint at an **unexpected** early Toarcian coleoid diversity in the Causses basin and point out the need
30 for further field investigations in the Lower Toarcian “Schistes Cartons” in this area. This new
31 record of *Paraplesioteuthis sagittata* is only the second one in Europe and the third in the world
32 (western Canada, Germany and now France). Based on these occurrences, we suggest that *P.*
33 *sagittata* originated in the Mediterranean domain and moved to the Arctic realm through the
34 Viking Corridor to eventually move even farther to North America.

35

36 Introduction

37 Within the cephalopod subclass Coleoidea Bather, 1888, the cohort Neocoeloidea Haas, 1997
38 includes present-day organisms (e.g., vampire squid, octopods, squids, cuttlefishes and their
39 relatives) that are mainly characterized by their internal shell. The latter is often called gladius

40 and consists of a sturdy but flexible chitinous structure within the dorsal mantle. The
41 evolutionary history, anatomy and paleobiology of Mesozoic gladius-bearing coleoids has been
42 extensively studied in the last four decades (Hauff & Hauff, 1981; Reitner & Engeser, 1981,
43 1982; Fischer & Riou, 1982, 2002; Riegraf et al., 1984, 1998; Engeser & Reitner, 1985, 1986;
44 Bandel & Leich, 1986; Engeser, 1988; Donovan & Toll, 1988; Hall & Neuman, 1989; Mehl,
45 1990; Doyle, 1990; Guérin-Franiatte & Gouspy, 1993; Doyle et al., 1994; Weis, 1998; Engeser
46 & Keupp, 1997, 1999; Haas, 2002; Košťák, 2002; Fuchs et al., 2003, 2007a, b, c, 2009, 2010,
47 2015, 2016; Bizikov, 2004, 2008; Fuchs & Weis, 2004, 2008, 2009, 2010; Wilby et al., 2004;
48 Riccardi, 2005; Fuchs, 2006a, b, c, 2007, 2009, 2015, 2016, 2019, 2020; Fuchs & Schultze,
49 2008; Larson, 2010; Donovan & Strugnell, 2010; Keupp et al., 2010; Klug et al., 2010, 2015,
50 2021a, b; Fuchs & Larson, 2011a, b; Breton et al., 2013; Donovan & Boletzky, 2014; Fuchs &
51 Iba, 2015; Jattiot et al., 2015a; Donovan & Fuchs, 2016; Kruta et al., 2016; Marroquín et al.,
52 2018; Košťák et al., 2021; Moreau et al., 2022; Rowe et al., 2022, 2023) thanks to a few
53 localities with geological horizons particularly favourable to their preservation, the so-called
54 Konservat-Lagerstätten. Thus, studies on exceptionally preserved Mesozoic gladius-bearing
55 coleoids are based on Konservat-Lagerstätten such as the Lower Jurassic Posidonia Shales of
56 Holzmaden, Germany (e.g., Hauff & Hauff, 1981; Riegraf et al., 1984; Klug et al., 2021a), the
57 Middle Jurassic of Christian Malford and Rixon Gate, England (e.g., Wilby et al., 2004), the
58 Middle Jurassic of La Voulte-sur-Rhône, France (e.g., Fischer & Riou, 1982, 2002; Charbonnier,
59 2009; Kruta et al., 2016; Rowe et al., 2022, 2023), the Upper Jurassic of Eichstätt, Solnhofen,
60 Painten and Nusplingen Plattenkalks, Germany (e.g., Fuchs, 2006b, 2015; Klug et al., 2010,
61 2015; Keupp et al. 2010), the Upper Jurassic of the Causse Méjean, France (Moreau et al., 2022)
62 and the Upper Cretaceous of Hâkel, Hâdjoula and Sâhel Aalma, Lebanon (e.g., Fuchs, 2006c;
63 Fuchs et al., 2009; Fuchs & Larson, 2011a, b; Jattiot et al., 2015a; Klug et al., 2021b).
64 However, as these Lagerstätten are not continuous in space, the record of Mesozoic gladius-
65 bearing coleoids is intermittent. As noted by Fuchs et al. (2016), our knowledge on geographical
66 distributions of species still needs to be greatly improved, which confers great importance upon
67 every new coleoid-bearing locality.

68 Here, we describe fossils of gladius-bearing coleoids from the Lower Toarcian “Schistes
69 Cartons” Formation (contemporaneous with the famous Posidonia Shales of Holzmaden; see
70 references above) of the Causses basin in southeastern France. The material described here
71 represents the first coleoid remains documented from this area. Thus, this work is the first
72 attempt to assess the taxonomic diversity of coleoids from the Lower Toarcian “Schistes
73 Cartons” of the Causses basin.

74

75 Materials & Methods

76 The material studied herein was collected by one of us (N.C.P.) in the northern part of the
77 Causses basin, Lozère, France (more precisely in the vicinity of Saint Bauzile village in the
78 Valdonnez Valley, at the base of the Balduc plateau, 5 km southern of Mende; Fig. 1). It consists
79 of two gladius-bearing coleoid specimens (M486_2024.1.1 and M486_2024.1.2) preserved as

80 compressions on slabs, which are housed in the palaeontological collections of the Musée du
81 Gévaudan (Mende, Lozère). We used UV-light to highlight soft tissues, using a UV wavelength
82 of 360 nm. The anatomical terminology of the gladius and systematic palaeontology follows
83 Fuchs & Weis (2010), Fuchs & Larson (2011a, b) and Fuchs (2016, 2020). Measured characters
84 (see Appendix A) are: preserved gladius length, maximum gladius width, median field width_{hypz},
85 hyperbolar zone length, maximum lateral fields width. These measurements are standard in most
86 published studies on coleoids (e.g., Fuchs & Larson, 2011a, b; Fuchs, 2016, 2020).

87

88 Geological settings and stratigraphy

89 The Causses Basin area is constituted by Jurassic limestone plateaus that are located south of the
90 Massif Central (southeastern France). The studied material was collected *in situ* near Saint
91 Bauzile (Fig. 1) from a ravine exhibiting a 27 m thick stratigraphic section showing Upper
92 Pliensbachian to Middle Toarcian deposits (Lower Jurassic, Fig. 2). The Upper Pliensbachian
93 corresponds with the Villeneuve Formation and consists of grey marls alternating with
94 concretioned, nodular and lenticular limestone beds (Fig. 2). At Saint Bauzile, this formation
95 mainly yields large belemnite rostra, bivalves (*Plicatula (Harpax) spinosa*), brachiopods (e.g.,
96 *Cirpa boscensis*) as well as some ammonites (e.g., *Juraphyllites*, *Pleuroceras*) and invertebrate
97 burrows (*Tisoa siphonalis*). The Toarcian is divided into two formations, the “Schistes Cartons”
98 Formation (Lower Toarcian) and the Marnes de Fontaneilles Formation (Lower to Upper
99 Toarcian). The “Schistes Cartons” Formation is about 7.5 m thick and consists of dark grey,
100 thinly laminated, organic-rich “shales” (Fig. 2). This formation is characterized by the abundance
101 of ammonite compressions (e.g., *Harpoceras falciferum*, *Dactylioceras*), aptychus, belemnite
102 rostra and large wood trunks. In other sites from the Causses Basin, the “Schistes Cartons” also
103 yielded rare vertebrate remains (e.g., crocodiles, ichthyosaurs; Bomou et al., 2021). At Saint
104 Bauzile, the first meter of the formation displays an alternation of centimetric, orange and
105 oxidized shale beds with centimetric black shale beds. The lower part of the formation shows
106 two hard and thinly laminated limestone beds. The first one, regionally called the “*Leptolepis*
107 bed” by several authors (e.g., Mattei, 1969; Trümpy, 1983), is very fossiliferous, displays a
108 characteristic bituminous smell and constitutes a benchmark bed observed all over the Causses
109 Basin. It is in this bed that the two coleoid specimens were retrieved. At Saint Bauzile, the
110 “*Leptolepis* bed” is quite isopach and 20 cm thick. It shows a strong concentration of *Leptolepis*
111 cf. *coryphaenoides* fishes (Coquel-Poussy et al., 2013). Near the Saint Bauzile locality, this bed
112 also yields rare lobster crustaceans (*Gabalerion*; Audo et al., 2017). In the Causses Basin, the
113 biozone associated with the “*Leptolepis* bed” is variable depending on the localities and the
114 authors. Based on ammonites, most authors stratigraphically locate these beds in the
115 Serpentinum Zone (e.g., Harazim et al., 2013; Pinard et al., 2014; Gatto et al., 2015), others in
116 the Tenuicostatum Zone (e.g., Fonseca et al., 2018). In the Balduc area, the detailed
117 biostratigraphic analysis conducted by Harazim et al. (2013) demonstrated that the “*Leptolepis*
118 bed” corresponds to the Serpentinum Zone. Regionally, the Marnes de Fontaneilles Formation
119 consists of grey to blue marls yielding a diversified, pyritized, marine, Middle to Upper Toarcian

120 fauna mainly including ammonites (e.g., Mattei, 1969, 1987; Jattiot et al., 2015b), belemnites
121 (e.g., Pinard et al., 2014), bivalves (Fürsich et al., 2002), gastropods (e.g., Gatto et al., 2015) and
122 rare vertebrates remains (e.g., Sciau et al., 1990; Bomou et al., 2021).

123 Toarcian deposits of the Causses Basin were deposited in a shallow epicontinental sea
124 located at a palaeolatitude of 25 to 30°N. At the base of the “Schistes Cartons” Formation,
125 Bomou et al. (2021) documented a negative carbon isotope excursion and higher mercury fluxes
126 that were linked with the Toarcian Oceanic Anoxic Event (T-OAE). This event, originating from
127 the intense volcanic activity of the Karoo Ferrar igneous province, is characterized by a
128 widespread deposition of organic-rich shales concomitant with the onset to an episode of global
129 warming. Bomou et al. (2021) showed that the deposition of the “Schistes Cartons” Formation
130 took place during a prolonged period of widespread oxygen-deficiency and elevated carbon
131 burial.

132

133 **Results**

134 Among Mesozoic gladius-bearing coleoids, three different morphotypes of gladius can be
135 recognized: prototeuthid, loligosepiid and teudopseid (Fuchs, 2009). Each is associated with the
136 suborders Prototeuthina Naef, 1921, Lolidosepiina Jeletzky, 1965 and Teudopseina
137 Starobogatov, 1983 (Fig. 3), respectively. The two individuals described in this study belong to
138 the Prototeuthina and Lolidosepiina, indicating that at least two of the three Mesozoic gladius-
139 bearing suborders were present in this locality.

140

141 Subclass Coleoidea Bather, 1888

142 Superorder Octobrachia Haeckel, 1866

143 Suborder Prototeuthina Naef, 1921

144 *Diagnosis* (after Fuchs, 2020). Octobrachiates with torpedo-shaped body; gladius length (=
145 median field length) equals mantle length; gladius with triangular median field and ventrally
146 closed (funnel-like) conus; gladius very slender to moderately wide, maximum gladius width
147 usually coincides with maximum median field width (by contrast to Lolidosepiina and
148 Teudopseina); median field slender (compared to most loligosepiids and teudopseids), with
149 median and lateral reinforcements; lateral reinforcements and central median field may be
150 projected; median field area large to very large compared to lateral fields (gladius is median field
151 dominated); hyperbolar zone indistinct or absent, hyperbolar zone length to median field length <
152 0.6; lateral fields very slender to moderately wide.

153

154 Family Plesioteuthidae Naef, 1921

155 *Type genus*. *Plesioteuthis* Wagner, 1859

156 *Diagnosis* (after Fuchs, 2020). Medium-sized prototeuthids; gladius very slender to moderately
157 wide (gladius width_{max} to gladius length 0.05–0.25), with triangular median field and ventrally
158 closed (funnel-like) conus; median field very slender to slender (median field width_{hypz} to
159 hyperbolar zone length < 0.35 = opening angle < 20°); median field area large to very large

160 (median field area to gladius area 0.70–1.0); lateral fields very slender to moderately wide;
161 hyperbolar zone very short to long (hyperbolar zone length to median field length < 0.6); median
162 and lateral reinforcements present on the median field; vestiges of septa and guard unknown,
163 eight arms equipped with uniserial circular suckers, sucker-rings absent; arm length variable;
164 funnel-and nuchal-locking cartilages absent; fins terminal; fin shape variable.

165 *Included genera* (after Fuchs, 2020). *Plesioteuthis* Wagner, 1859; *Boreopeltis* Engeser and
166 Reitner, 1985; *Dorateuthis* Woodward, 1883; *Eromangateuthis* Fuchs, 2019; *Nesisoteuthis*
167 Doguzhaeva, 2005; *Normanoteuthis* Breton, Strugnell and Donovan, 2013; *Paraplesioteuthis*
168 Naef, 1921; *Romaniteuthis* Fischer and Riou, 1982; *Rhombopteuthis* Fischer and Riou, 1982;
169 *Senefelderiteuthis* Engeser and Keupp, 1999.

170 *Stratigraphical and geographical range* (after Fuchs & Larson, 2011a). (?)Late Triassic
171 (Rhaetian), Early Jurassic (Toarcian)–Late Cretaceous (Maastrichtian); Europe, Central Russia,
172 Lebanon, North America and Australia.

173

174 Genus *Paraplesioteuthis* Naef, 1921

175 *Type species*. *Geoteuthis sagittata* Münster, 1843 by the subsequent designation of Naef (1922,
176 p. 111).

177 *Diagnosis* (after Fuchs, 2020). Gladius medium-sized, slender to moderately wide (gladius
178 width_{max} to gladius length 0.15–0.25) with a bipartite median ridge. Median field slender to
179 moderately wide (median field width_{hypz} to hyperbolar zone length 0.25–0.35 = opening angle
180 14°–20°), triangular and with lateral platelike reinforcements. Lateral reinforcements and central
181 median field anteriorly projected. Median field area large to very large (median field area to
182 gladius area 0.75–0.85). Lateral fields slender (lateral fields width_{max} to median field width_{max}
183 0.85–0.95). Hyperbolar zone moderately long to long (hyperbolar zone length to median field
184 length 0.45–0.55). Soft parts poorly known.

185 *Included species*. *Paraplesioteuthis sagittata* (Münster, 1843) only.

186 *Stratigraphical and geographical range*. Upper Pliensbachian–lower Toarcian; southern
187 Germany (Holzmaden region; Fuchs, 2006b), western Canada (Fernie Formation; Hall, 1985,
188 Marroquín et al., 2018), southeastern France (Causse basin; this study).

189

190 *Paraplesioteuthis sagittata* (Münster, 1843)

191 Figure 4

192

193 1843. *Geoteuthis sagittata* Münster, pp. 672–673, pl. 7, fig. 3, pl. 8, fig. 4.

194 p 1843. *Geoteuthis hastata* Münster, p. 73, pl. 14, fig. 4.

195 non 1843. *Geoteuthis hastata* Münster, p. 73, pl. 8, fig. 3.

196 1860. *Geoteuthis sagittata* Münster; Wagner, p. 807.

197 1922. *Paraplesioteuthis hastata* (Münster); Naef, p. 114, fig. 41a–c.

198 1978. *Paraplesioteuthis sagittata* (Münster); Reitner, p. 210, fig. 6.

199 1984. *Paraplesioteuthis sagittata* (Münster); Riegraf et al. p. 36.

- 200 1984. *Paraplesioteuthis hastata* (Münster); Riegraf et al. p. 36.
201 1985. *Paraplesioteuthis hastata* (Münster); Hall, p. 871, fig. 1.
202 1990. *Paraplesioteuthis sagittata* (Münster); Doyle, p. 205.
203 2006b. *Paraplesioteuthis hastata* (Münster); Fuchs, pl. 16a–c.
204 2009. *Paraplesioteuthis hastata* (Münster); Fuchs, fig. 1a–b.
205 2011a. *Paraplesioteuthis sagittata* (Münster); Fuchs & Larson, fig. 7.1.
206 ? 2018. *Paraplesioteuthis cf. sagittata* (Münster); Marroquín et al., figs. 6–10.
207 2020. *Paraplesioteuthis sagittata* (Münster); Fuchs, p. 10, fig. 4,1a–b
208
209 *Holotype*. The original specimen of Münster (1843, pl. 7, fig. 3) was lost during World War II.
210 *Lectotype (designated by Marroquín et al. 2018)*. Original of Münster (1843, pl. 8, fig. 4),
211 Geologisch-Paläontologisches Museum Tübingen, GPIT 1529-2 (original of Reitner 1978, fig.
212 6).
213 *Type locality*. Holzmaden region, southern Germany.
214 *Type horizon*. Posidonia Shales Formation, lower Toarcian (Lower Jurassic).
215 *Stratigraphical and geographical range*. As for genus.
216
217 *Material*. One incomplete specimen (M486_2024.1.1) from the Lower Toarcian “Schistes
218 Cartons” Formation of the Causses Basin, in the vicinity of Saint Bauzile village (Lozère,
219 France).
220
221 *Description*. Although the single specimen (M486_2024.1.1) is only partially preserved,
222 morphological description and taxonomic identification at the species level remain possible. The
223 specimen consists of a long and slender gladius (interpreted here as in dorsal view) with a
224 triangular, anteriorly diverging median field (Fig. 4A–C). The preserved gladius length (=
225 median field length) is 203 mm. Although the anterior end of the gladius is not preserved, we
226 hypothesize that the original gladius length did not exceed 220 mm (according to Klug et al.
227 2021a, gladii of *Paraplesioteuthis sagittata* rarely reach 200 mm). Of note, we suspect that the
228 anterior part of the gladius, which is poorly preserved, was affected by slight disruptions and
229 distortions (Fig. 4A, B). In our opinion, this impedes providing a reliable measurement of the
230 anteriormost gladius width. Partially preserved lateral reinforcements (most conspicuous in the
231 posterior part of the gladius, Fig. 4A–C) diverge from posterior to anterior extremities. Based on
232 the estimated median field width_{hypz} to hyperbolar zone length (see Fig. 3 and Appendix A), we
233 estimate that the lateral reinforcements form an opening angle of ~11° (see Appendix A). The
234 lateral fields are relatively slender. Although the outline of the hyperbolar zone is hardly
235 discernible, it appears relatively long (estimated hyperbolar zone length to median field length
236 ratio is 0.44). Finally, it cannot be determined whether the median ridge is bipartite as commonly
237 described for *Paraplesioteuthis* representatives.
238

239 *Remarks.* In our opinion, the gladius is too poorly preserved to provide an accurate measurement
240 of the original gladius width_{max}. Nevertheless, based on the general shape of the gladius, we
241 consider that the original gladius width_{max} to gladius length ratio likely fell within the range of
242 values mentioned by Fuchs (2020, p. 10) for *Paraplesioteuthis* (i.e., 0.15–0.25). Of note, the
243 opening angle of ~11° framed by the lateral reinforcements in the present specimen is lower than
244 the range of values given by Fuchs (2020, p. 10) for *Paraplesioteuthis* (i.e., 14°–20°). On the
245 other hand, it is comparable to the opening angle of 10° mentioned by Hall (1985) for the *P.*
246 *hastata* (Münster, 1843) specimen from western Canada (*P. hastata* is regarded as conspecific
247 with *P. sagittata* by Fuchs & Larson, 2011a). Thus, we suggest that values of opening angle for
248 *Paraplesioteuthis* should be redefined as ranging from about 10° to 20°.

249 According to Fuchs & Larson (2011b), the Middle Jurassic genus *Romaniteuthis* differs from
250 *Paraplesioteuthis* by having a reduced median field width (i.e., gladius width_{max} to gladius length
251 ratio 0.05–0.15; Fuchs, 2020) and lateral reinforcements as keels. Although the specimen
252 described herein is too poorly preserved to provide a reliable measurement of the preserved
253 gladius width_{max}, its original gladius width_{max} to gladius length ratio was probably not less than
254 0.15. Furthermore, it does not exhibit prominent lateral keels.

255 *Paraplesioteuthis* lateral fields are relatively short (Fuchs & Larson, 2011a; see Fig. 4D, E). In
256 this regard, the lateral fields of the present specimen seem more similar to that of *Romaniteuthis*,
257 since they appear slightly elongated, in oval shape (compare Fig. 4A–C with fig. 7.2 in Fuchs &
258 Larson 2011a). This may however be due to slight intraspecific variability. Finally, the estimated
259 hyperbolar zone length to median field length ratio for the present specimen is 0.44, which
260 nearly falls within the range of values given by Fuchs (2020) for *Paraplesioteuthis* (i.e., 0.45–
261 0.55).

262 In sum, despite a possible slight difference in shape of lateral fields, we consider that other
263 features of the present gladius support its attribution to the species *Paraplesioteuthis sagittata*.

264

265 Order Vampyromorpha Robson, 1929

266 Suborder Loligosepiina Jeletzky, 1965

267 *Diagnosis* (after Fuchs, 2020). Small- to large-sized octobrachiates with bullet-shaped body;
268 gladius length (= median field length) equals mantle length; gladius with triangular median field
269 and cup-shaped conus; gladius slender to wide, maximum gladius width always exceeds
270 maximum median field width; median field width very slender to moderately wide without
271 pronounced median reinforcements, anterior median field margin concave, straight or convex;
272 median field area small to large; hyperbolar zone mostly well-arcuated, rarely indistinct, long to
273 very long; lateral fields usually moderately wide.

274

275 Family Loligosepiidae Regteren Altena, 1949

276 *Type genus.* *Loligosepia* Quenstedt, 1839.

277 *Diagnosis* (after Fuchs, 2020). Medium-sized loligosepiids; gladius slender to wide (gladius
278 width_{max} to gladius length 0.10–0.60), with deeply concave (V-shaped) hyperbolar zone; median

279 field very slender to moderately wide (median field width_{hypz} to hyperbolar zone length 0.10–
280 0.40 = opening angle 7°–23°), anterior median field margin slightly convex; median field area
281 small to moderate (median field area to gladius area 0.35–0.45); hyperbolar zone very long
282 (hyperbolar zone length to median field length 0.85–0.95); lateral fields moderately wide (lateral
283 fields width_{max} to median field width_{max} 1.30–1.85), anterior limit of lateral fields clearly pointed
284 (spine-like); inner and outer asymptotes ridge-like.

285 *Included genera.* *Loligosepia* Quenstedt, 1839 and *Jeletzkyteuthis* Doyle, 1990.

286 *Stratigraphical and geographical range.* Lower Sinemurian–lower Toarcian; Germany,
287 Luxembourg, France, Switzerland, UK, Canada (Alberta).

288

289 Genus *Loligosepia* Quenstedt, 1839

290 *Type species.* *Loligo aalensis* Schübler in Zieten, 1832 (by subsequent designation of Regteren
291 Altena, 1949, p. 57) from the lower Toarcian Posidonia Shales of Holzmaden (Germany).

292 *Diagnosis (after Fuchs, 2020).* Medium-sized loligosepiids, gladius moderately wide to wide
293 (gladius width_{max} to median field length 0.30–0.60); median field slender to moderately wide
294 (median field width_{hypz} to hyperbolar zone length 0.20–0.40 = opening angle 12°–23°); anterior
295 median field margin convex; median field area small to moderate (median field area to gladius
296 area 0.35–0.45); hyperbolar zone very long (hyperbolar zone length to median field length 0.85–
297 0.95); lateral fields moderately wide (lateral fields width_{max} to median field width_{max} 1.35–1.80;
298 arms short to moderate (arm length to mantle length ~0.45).

299 *Included species.* *Loligosepia bucklandi* (Voltz, 1840) from the Lower Sinemurian of Dorset
300 (UK) and *L. aalensis* (Schübler in Zieten, 1832) from the Lower Toarcian of Holzmaden.

301 *Stratigraphical and geographical range.* Lower Sinemurian–lower Toarcian of Germany,
302 Luxembourg, France, UK and Canada (Alberta).

303

304 ?*Loligosepia* sp. indet.

305 Figure 5

306

307 *Material.* One incomplete specimen (M486_2024.1.2) from the Lower Toarcian “Schistes
308 Cartons” Formation of the Causses Basin, in the vicinity of Saint Bauzile village (Lozère,
309 France).

310

311 *Description.* The preserved gladius length (= median field length) of this specimen
312 (M486_2024.1.2) is 76 mm. Although a significant part of the original gladius is most certainly
313 missing, we tentatively hypothesize that the original gladius did not exceed 150 mm.
314 Unfortunately, few features can be described on this specimen. On the posterior part of the
315 specimen, inconspicuous lines are interpreted as a disrupted median line and inner asymptotes
316 (Fig. 5A–C). The black structure preserved anteriorly is interpreted as the ink sac (Fig. 5A–C).
317 The specimen outline appears weakly constricted posteriorly, although it cannot be determined

318 whether it is a genuine feature, or if it is due to slight taphonomic disruptions. The median field
319 width_{max} of the original gladius cannot be assessed.

320

321 *Remarks.* The overall shape of the gladius combined with the presence of lines interpreted as a
322 median line and inner asymptotes hint at the possibility that this specimen belongs to the
323 suborder Lolidosepiina. Mesozoic gladius-bearing coleoids belonging **of** the suborder
324 Teudopseina exhibit a characteristically constricted median field anteriorly, which does not seem
325 to be the case in specimen M486_2024.1.2. Within the suborder Lolidosepiina, the assignment of
326 this specimen to the genus *Lolidosepia* is **unsettled**. It is only based on a conjectural original size
327 of the specimen (estimated around 150 mm) that is comparable with that of representatives of the
328 *Lolidosepia* species *L. bucklandi* (Voltz, 1840) and *L. aaalensis* (Schübeler in Zieten, 1832). In
329 our opinion, *Jeletzkyteuthis* species differ mostly by their larger gladius size. For example, *J.*
330 *coriaceus* (Quenstedt, 1849) gladiuses regularly exceed 200 mm in length, according to Klug et
331 al. (2021a).

332

333 Discussion

334

335 *Palaeobiogeography of Paraplesioteuthis sagittata*

336

337 The present record of *Paraplesioteuthis sagittata* is only the second one in Europe and the third
338 in the world (western Canada, Germany and now France; Fig. 6). During the Early Jurassic, the
339 NW Tethyan and Arctic realms (which consisted of two epicontinental seas) were linked by a
340 narrow seaway named the Viking Corridor (Ziegler, 1988; Fig. 6). In this time interval, the
341 Serpentium Zone (from **which** the coleoids described herein come **from**) marks the onset of the
342 **disruption of a previous provincialism**, with a strong homogenization of all Tethyan and Arctic
343 ammonite species (Dera et al., 2011). This event is linked with the origination of numerous
344 cosmopolitan **taxa** in the Mediterranean domain (Macchioni and Cecca, 2002). In this context,
345 the Viking Corridor probably regulated the mixing between Arctic and Euro-Boreal ammonites
346 (Smith et al., 2001; Dera et al., 2011).

347 Based on this, it can be hypothesized that some Early Jurassic coleoids (such as *P. sagittata*),
348 similarly to ammonites, broadly originated in the NW Tethyan realm and moved to the Arctic
349 realm through the **Viking Corridor, to eventually move even farther to North America**. Other
350 routes cannot be excluded, such as the Hispanic Corridor (Fig. 6), a narrow epicontinental
351 seaway that was sporadically active since the Late Sinemurian–Early Pliensbachian time interval
352 (Aberhan, 2001, 2002; Venturi et al., 2006; Dera et al., 2009). However, according to Dera et al.
353 (2011, p. 100), the Hispanic Corridor “...was certainly too shallow for allowing massive
354 movements of hemipelagic organisms such as ammonites”. Since coleoids are also hemipelagic
355 organisms, we presume that coleoids, similarly to ammonites, **were not able to go through the**
356 **Hispanic corridor.**

357 In the hypothesis that coleoids moved from the Mediterranean domain to North America through
358 the Viking Corridor, we would expect to find *P. sagittata* specimens in localities from the Arctic
359 Realm, provided that the geological time interval is represented and that there are geological
360 horizons peculiarly favourable to the preservation of gladius-bearing coleoids.

361

362 **Conclusions**

363 Two of the three suborders of Mesozoic gladius-bearing coleoids are represented in the studied
364 material, which is constituted of only two specimens. This hints at an unexpected early Toarcian
365 coleoid diversity in the Causses basin and points out the need for further field investigations in
366 the Lower Toarcian black shales in this area. New findings from the Causses basin would indeed
367 most certainly improve our understanding of the Mesozoic gladius-bearing coleoid
368 palaeobiogeography, ecology and taxonomy. Finally, based on the known worldwide
369 occurrences of *P. sagittata*, we suggest that this species originated in the Mediterranean domain
370 and moved to the Arctic realm through the Viking Corridor to eventually move even farther to
371 North America.

372

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376

377 **References**

- 378 Aberhan, M. 2001. Bivalve palaeobiogeography and the Hispanic Corridor: time of opening and
379 effectiveness of a proto-Atlantic seaway. *Palaeogeography, Palaeoclimatology, Palaeoecology*,
380 165(3–4): 375–394.
- 381 Aberhan, M. 2002. Opening of the Hispanic Corridor and the Early Jurassic bivalve biodiversity.
382 In: Crame, J.A. and Owen, A.W. (Eds.), *Palaeobiogeography and Biodiversity Change: The
383 Ordovician and Mesozoic–Cenozoic Radiations*: Geological Society of London, Special
384 publications, pp. 127–139.
- 385 Audo, D., Williams, M., Charbonnier, S. and Schweigert, G. 2017. *Gabaleryon*, a new genus of
386 widespread early Toarcian polychelidan lobsters. *Journal of Systematic Palaeontology*,
387 15(3):205–222.
- 388 Bandel, K. and H. Leich. 1986. Jurassic Vampyromorpha (dibranchiate cephalopods). Neues
389 *Jahrbuch für Geologie und Paläontologie Monatshefte*:129–148.
- 390 Bather, F. A. 1888. Shell-growth in Cephalopoda (Siphonopoda). *Annals and Magazine of
391 Natural History*, 6:421–427.
- 392 Bomou, B., Suan, G., Schlögl, J., Grosjean, A. S., Suchéras-Marx, B. et al. 2021. The
393 palaeoenvironmental context of Toarcian vertebrate-yielding shales of southern France (Hérault).
394 *The Geological Society, London, Special Publications*, 2021, *Carbon Cycle and Ecosystem
395 Response to the Jenkyns Event in the Early Toarcian (Jurassic)*, 514 p.

- 396 Bizikov, V. A. 2004. The shell in Vampyropoda (Cephalopoda): Morphology, functional role
397 and evolution. *Ruthenica Supplement*, 3:1–88.
- 398 Bizikov, V. A. 2008. Evolution of the shell in Cephalopoda. VNIRO Publishing, Moscow, 445 p.
- 399 Breton, G., Strugnell, J.M. and Donovan D.T. 2013. A coleoid gladius (Mollusca, Cephalopoda)
400 from the Albian of Normandy (France): A new squid genus and species. *Annales de*
401 *Paléontologie*, 99(3):275–283.
- 402 Charbonnier, S. 2009. Le Lagerstätte de La Voulte : un environnement bathyal au Jurassique.
403 *Mémoires du Muséum national d'Histoire naturelle*, 199:1–272.
- 404 Coquel-Poussy, N. 2013. Recherches paléontologiques dans le Toarcien basal de la Lozère.
405 *Fossiles*, 15:5–15.
- 406 Dera, G., Pucéat, E., Pellenard, P., Neige, P., Delsate, D., Joachimski, M.M., Reisberg, L. and
407 Martinez, M. 2009. Water mass exchange and variations in seawater temperature in the NW
408 Tethys during the Early Jurassic: evidence from neodymium and oxygen isotopes of fish teeth
409 and belemnites. *Earth and Planetary Science Letters*, 286(1–2):198–207.
- 410 Dera, G., Neige, P., Dommergues, J-L. and Brayard, A. 2011. Ammonite paleobiogeography
411 during the Pliensbachian–Toarcian crisis (Early Jurassic) reflecting paleoclimate, eustasy, and
412 extinctions. *Global and Planetary Change*, 78:92–105.
- 413 Doguzhaeva, L.A. 2005. A gladius-bearing coleoid cephalopod from the Aptian of Central
414 Russia. *Mitteilungen des Geologisch-Paläontologischen Instituts, Universität Hamburg*, 89:41–
415 48.
- 416 Donovan, D. T. and R. B. Toll. 1988. The gladius in coleoid (Cephalopoda) evolution, p. 89–
417 101. In M. R. Clarke, and E. R. Trueman. (eds.), *The Mollusca. Palaeontology and Neontology*
418 of Cephalopods
- , Vol. 12. Academic Press, San Diego.
- 419 Donovan, D. T. and J. Strugnell. 2010. A redescription of the fossil coleoid cephalopod genus
420 *Palaeololigo* Naef, 1921 and its relationship to recent squids. *Journal of Natural History*,
421 44:1475–1492.
- 422 Donovan, D. T. and Boletzky, S. v. 2014. Lolidosepia (Cephalopoda: Coleoidea) from the Lower
423 Jurassic of the Dorset coast, England. *Neues Jahrbuch für Geologie und Paläontologie*
424 *Abhandlungen*, 273:45–63.
- 425 Donovan, D. T. and Fuchs, D. 2016. Part M, Chapter 13: Fossilized soft tissues in Coleoidea.
426 *Treatise Online*, 73:1–30.
- 427 Doyle P. 1990. Teuthid cephalopods from the Lower Jurassic of Yorkshire. *Palaeontology*,
428 33(1):193–207.
- 429 Doyle, P., Donovan, D. T. and Nixon, M. 1994. Phylogeny and systematics of the Coleoidea.
430 *The University of Kansas Paleontological Contributions (New Series)*, 5 :1–15.
- 431 Engeser, T. 1988. Vampyromorpha (“Fossile Teuthiden”), p. 1–167. In F. Westphal (ed.),
432 *Fossilium Catalogus. I: Animalia*, vol. 130. Kugler Publications, Amsterdam.
- 433 Engeser, T. and Reitner, J. 1985. Teuthiden aus dem Unterapt (“Töck”) von Helgoland
434 (Schleswig-Holstein, Norddeutschland). *Paläontologische Zeitschrift*, 59:245–260.

- 435 Engeser, T. and Reitner, J. 1986. Coleoidenreste aus der Oberkreide des Libanon im Staatlichen
436 Museum für Naturkunde in Stuttgart. *Stuttgarter Beiträge zur Naturkunde, Ser. B*, 124:1–17.
- 437 Engeser, T. and Keupp, H. 1997. Zwei neue Gattungen und eine neue Art von vampyromorphen
438 Tintenfischen (Coleoidea, Cephalopoda) aus dem Untertithonium von Eichstätt. *Archaeopteryx*,
439 15:47–58.
- 440 Engeser, T. and Keupp, H. 1999. Zwei neue vampyromorphe Tintenfische (Coleoidea,
441 Cephalopoda) aus dem oberjurassischen Solnhofener Plattenkalk von Eichstätt. *Archaeopteryx*,
442 17:21–32.
- 443 Fischer, J.-C. and B. Riou. 1982. Les Teuthoïdes (Cephalopoda, Dibranchiata) du Callovien
444 inférieur de La Voulte-sur-Rhône (Ardèche, France). *Annales de Paléontologie*, 68:295–325.
- 445 Fischer, J.-C. and Riou, B. 2002. *Vampyronassa rhodanica* nov. gen. nov sp., vampyromorphe
446 (Cephalopoda, Coleoidea) du Callovien inférieur de La Voulte-sur-Rhône (Ardèche, France).
447 *Annales de Paléontologie*, 88:1–17.
- 448 Fonseca, C., Mendonça Filho, J. G., Lézin, C., Duarte, L. V. and Fauré, P. 2018. Organic facies
449 variability during the Toarcian Oceanic Anoxic Event record of the Grands Causses and Quercy
450 basins (southern France). *International Journal of Coal Geology*, 190:218–235.
- 451 Fuchs, D. 2006a. Re-description of *Doryanthes munsterii* (D'ORBIGNY, 1845), a poorly known
452 vampyropod coleoid (Cephalopoda) from the Late Jurassic Solnhofen Plattenkalks.
453 *Archaeopteryx*, 24:79–88.
- 454 Fuchs, D. 2006b. Fossil erhaltungsfähige Merkmalskomplexe der Coleoidea (Cephalopoda) und
455 ihre phylogenetische Bedeutung. *Berliner Paläobiologische Abhandlungen*, 8:1–122.
- 456 Fuchs, D. 2006c. Diversity, taxonomy and morphology of vampyropod coleoids (Cephalopoda)
457 from the Upper Cretaceous of Lebanon. *Memorie della Società Italiana di Scienze Naturali et*
458 *del Museo Civico di Storia Naturale di Milano*, 34:1–28.
- 459 Fuchs, D. 2007. Coleoid cephalopods from the Plattenkalks of the Late Jurassic of Southern
460 Germany and the Late Cretaceous of Lebanon—A faunal comparison. *Neues Jahrbuch für*
461 *Geologie und Paläontologie Abhandlungen*, 245:59–69.
- 462 Fuchs, D. 2009. Octobrachia – a diphylectic taxon? *Berliner paläobiologische Abhandlungen*,
463 10:181–192.
- 464 Fuchs, D. 2015. Tintenfische (Coleoidea, Endocochleata, Dibranchiata). In: Arratia, G., Schultze,
465 H.-P., Tischlinger, H. and Viohl, G. (eds.). Solnhofen – Ein Fenster in die Jurazeit. Vol. 1+2 [in
466 German]. 229–238. Pfeil, Munich.
- 467 Fuchs, D. 2016. Part M, Chapter 9B: The gladius and gladius vestige in fossil Coleoidea.
468 *Treatise Online*, 83:1–23.
- 469 Fuchs, D. 2019. *Eromangateuthis* n. Gen., a new genus for a late Albian gladius-bearing giant
470 octobrachian (cephalopoda: coleoidea). *Paleontological Contributions*, 21:1–3.
- 471 Fuchs, D. 2020. Part M, Chapter 23G: Systematic Descriptions: Octobrachia. *Treatise Online*
472 138:1–52.

- 473 Fuchs, D. and Weis, R. 2004. The “Schistes Cartons” (Lower Toarcian) of Luxembourg and its
474 unnoticed coleoid diversity. In: Mapes, R. H. and Landman, N. H. (eds.): Abstract volume of the
475 6th International Symposium Cephalopods – Present and Past, 2004, Fayette-ville: 43–44.
- 476 Fuchs, D. and Schultze, H.-P. 2008. *Trachyteuthis covacevichi* n. sp., a Late Jurassic coleoid
477 cephalopod from the Paleopacific. *Fossil Record*, 11:39–49.
- 478 Fuchs, D. and Weis, R. 2008. Taxonomy, morphology and phylogeny of Lower Jurassic
479 loligosepiid coleoids (Cephalopoda). *Neues Jahrbuch für Geologie und Paläontologie*
480 *Abhandlungen*, 249:93–112.
- 481 Fuchs, D. and Weis, R. 2009. A new Cenomanian (Late Cretaceous) coleoid (Cephalopoda) from
482 Hâdjoula, Lebanon. *Fossil Record*, 12:175–181.
- 483 Fuchs, D. and Weis, R. 2010. Taxonomy, morphology and phylogeny of Lower Jurassic
484 teudopseid coleoids (Cephalopoda). *Neues Jahrbuch für Geologie und Paläontologie*
485 *Abhandlungen*, 257:351–366.
- 486 Fuchs, D. and Larson, N. 2011a. Diversity, morphology and phylogeny of coleoid cephalopods
487 from the Upper Cretaceous Plattenkalks of Lebanon–Part I: Prototeuthidina. *Journal of*
488 *Paleontology*, 85:234–249.
- 489 Fuchs, D. and Larson, N. 2011b. Diversity, morphology and phylogeny of coleoid cephalopods
490 from the Upper Cretaceous Plattenkalks of Lebanon–Part II: Teudopseina. *Journal of*
491 *Paleontology*, 85:815–834.
- 492 Fuchs, D. and Iba, Y. 2015. The gladiuses in coleoid cephalopods homology, parallelism, or
493 convergence? *Swiss Journal of Palaeontology*, 134:187–197.
- 494 Fuchs, D., Keupp, H. and Engeser, T. 2003. New records of soft parts of *Muensterella scutellaris*
495 Muenster, 1842 (Coleoidea) from the Late Jurassic Plattenkalks of Eichstätt and their
496 significance for octobrachian relationships. *Berliner Paläobiologische Abhandlungen*, 3:101–
497 111.
- 498 Fuchs, D., Engeser, T. and Keupp, H. 2007a. Gladius shape variation in coleoid cephalopod
499 *Trachyteuthis* from the Upper Jurassic Nusplingen and Solnhofen Plattenkalks. *Acta*
500 *Palaeontologica Polonica*, 52:575–589.
- 501 Fuchs, D., Klinghammer, A. and Keupp, H. 2007b. Taxonomy, morphology and phylogeny of
502 plesiotethidid coleoids from the Upper Jurassic (Tithonian) Plattenkalks of Solnhofen. *Neues*
503 *Jahrbuch für Geologie und Paläontologie Abhandlungen*, 245:239–252.
- 504 Fuchs, D., Beard, G., Tanabe, K. and Ross, R. 2007c. Coleoid cephalopods from the Late
505 Cretaceous North eastern Pacific. In: Abstracts volume of the 7th International Symposium
506 International Symposium Cephalopods – Present and Past, 2007, Sapporo.
- 507 Fuchs, D., Bracchi, G. and Weis, R. 2009. New records of octopods (Cephalopoda: Coleoidea)
508 from the Late Cretaceous (Upper Cenomanian) of Hakel and Hadjoula (Lebanon).
509 *Palaeontology*, 52:65–81.
- 510 Fuchs, D., Stinnesbeck, W., Ifrim, C., Giersch, S., Gutierrez, J. M. P. and Frey, E. 2010.
511 *Glyphiteuthis rhinophora* n. sp., a trachyteuthidid (Coleoidea, Cephalopoda) from the
512 Cenomanian (Late Cretaceous) of Mexico. *Paläontologische Zeitschrift*, 84:523–532.

- 513 Fuchs, D., Iba, Y., Tischlinger, H., Keupp, H. and Klug, C. 2015. The locomotion system of
514 Mesozoic Coleoidea (Cephalopoda) and its phylogenetic significance. *Lethaia*, 49:433–454.
- 515 Fuchs, D., Reitano, A., Insacco, G., and Iba, Y. 2016. The first coleoid cephalopods from the
516 Upper Cenomanian of Sicily (Italy) and their implications for the systematic-phylogenetic
517 position of the Palaeololiginidae (Teudopseina). *Journal of Systematic Palaeontology*,
518 15(6):499–512.
- 519 Fürsich, F. T., Berndt, R., Scheuer, T. and Gahr, M. 2001. Comparative ecological analysis of
520 Toarcian (Lower Jurassic) benthic faunas from southern France and east-central Spain. *Lethaia*,
521 34(3):169–199.
- 522 Gatto, R., Monari, S., Neige, P., Pinard, J. D., and Weis, R. 2015. Gastropods from upper
523 Pliensbachian–Toarcian (Lower Jurassic) sediments of Causses Basin, southern France and their
524 recovery after the early Toarcian anoxic event. *Geological Magazine*, 152(5) :871–901.
- 525 Guérin-Franiatte, S., and Gouspy, C. 1993. Découverte de Céphalopodes Teuthides (Coleoidea)
526 dans le Lias supérieur de Haute-Marne, France. *Geobios, M.S.*, 15 :181–189.
- 527 Haas, W. 1997. Der ablauf der entwicklungsgeschichte der Decabrachia (Cephalopoda,
528 Coleoidea). *Palaeontographica, Abteilung A*, 254:63–81.
- 529 Haas, W. 2002. The evolutionary history of the eight-armed Coleoidea, p. 341–351. In H.
530 Summesberger, K. Histon, and A. Daurer (eds.), *Cephalopods—Present & Past*. Abhandlungen
531 der Geologischen Bundesanstalt, 57, Wien.
- 532 Haeckel, E. 1866. Generelle Morphologie der Organismen. Georg Reiner, Berlin, 462 p.
- 533 Hall, R. L. 1985. *Paraplesioteuthis hastata* (Münster), the first teuthid squid recorded from the
534 Jurassic of North America. *Journal of Paleontology*, 59(4):870–874.
- 535 Hall, R. L. and Neuman, A. G. 1989. *Teudopsis cadominensis*, a new teuthid squid from the
536 Toarcian (lower Jurassic) of Alberta. *Journal of Paleontology*, 63:324–327.
- 537 Harazim, D., Van De Schootbrugge, B. A. S., Sorichter, K., Fiebig, J., Weug, A., Suan, G. and
538 Oschmann, W. 2013. Spatial variability of watermass conditions within the European
539 Epicontinental Seaway during the Early Jurassic (Pliensbachian–Toarcian). *Sedimentology*,
540 60(2):359–390.
- 541 Hauff, B. and Hauff, R. B. 1981. Das Holzmadenbuch, 3rd edition. 136 p., Holzmaden.
- 542 Jattiot, R., Brayard, A., Fara, E. & Charbonnier, S. 2015a. Gladius-bearing coleoids from the
543 Upper Cretaceous Lebanese Lagerstätten: diversity, morphology, and phylogenetic implications.
544 *Journal of Paleontology*, 89:148–167.
- 545 Jattiot, R., Trincal, V., Moreau, J. D. and Brocard, A. 2015b. Guide des ammonites pyriteuses,
546 Toarcien moyen et supérieur des Causses (Lozère-France). Les Editions du Piat, 144 p.
- 547 Jeletzky, J. A. 1965. Taxonomy and phylogeny of fossil Coleoidea (=Dibranchiata). *Geological
548 Survey of Canada, Papers*, 65(2):76–78.
- 549 Keupp, H., Engeser, T., Fuchs, D., and Haechel, W. 2010. Ein *Trachyteuthis hastiformis*
550 (Cephalopoda, Coleoidea) mit Spermatophoren aus dem Ober-Kimmeridgium von Painten
551 (Ostbayern). *Archaeopteryx*, 28:23–30.

- 552 Košťák, M. 2002. Teuthoidea from the Bohemian Cretaceous Basin (Czech Republik) – A
553 critical review. *Abhandlungen der geologischen Bundesanstalt*, 57:359–369.
- 554 Košťák, M., Schlägl, J., Fuchs, D., Holcová, K., Hudáčková, N., Culka, A., Fözy, I.,
555 Tomašových, A., Milovský, R., Šurka, J. and Mazuch, M. 2021. Fossil evidence for vampire
556 squid inhabiting oxygen-depleted ocean zones since at least the Oligocene. *Communications
557 Biology*, 4, 216.
- 558 Klug, C., Schweigert, G., and Dietl, G. 2010. A new *Plesioteuthis* with beak from the
559 Kimmeridgian of Nusplingen (Germany). In: Fuchs, D. (Ed.), Proceedings of the Third
560 International Coleoid Symposium. *Ferrantia*, 59:73–77.
- 561 Klug, C., Fuchs, D., Schweigert, G., Röper, M., and Tischlinger, H. 2015. New anatomical
562 information on arms and fins from exceptionally preserved *Plesioteuthis* (Coleoidea) from the
563 Late Jurassic of Germany. *Swiss Journal of Palaeontology*, 134:245–255.
- 564 Klug, C., Schweigert, G., Fuchs, D., and De Baets, K. 2021a. Distraction sinking and fossilized
565 coleoid predatory behaviour from the German Early Jurassic. *Swiss Journal of Palaeontology*,
566 140(1):1–12.
- 567 Klug, C., Di Silvestro, G., Hoffmann, R., Schweigert, G., Fuchs, D., Clements, T., and Guériaud,
568 P. 2021b. Diagenetic phosphatic Liesegang rings deceptively resemble chromatophores in
569 Mesozoic coleoids. *PeerJ*, 9:e10703.
- 570 Kruta, I., Rouget, I., Charbonnier, S., Bardin, J., Fernandez, V., Germain, D., Brayard, A., and
571 Landman, N. 2016. *Proteroctopus ribeti* in coleoid evolution. *Palaeontology*, 59(6):767–773.
- 572 Larson, N. L. 2010. Fossil coleoids from the Late Cretaceous (Campanian & Maastrichtian) of
573 the Western Interior. *Ferrantia*, 59, 78–113.
- 574 Macchioni, F. and Cecca, F. 2002. Biodiversity and biogeography of middle–late liassic
575 ammonoids: implications for the early Toarcian mass extinction. *Geobios*, 35(Supplement
576 1):165–175.
- 577 Marroquín, S. M., Martindale, R. C., and Fuchs, D. 2018. New records of the late Pliensbachian
578 to early Toarcian (Early Jurassic) gladius-bearing coleoid cephalopods from the Ya Ha Tinda
579 Lagerstätte, Canada. *Papers in Palaeontology*, 4(2):245–276.
- 580 Mattei, J. 1969. Définition et interprétation de *Pseudopolyplectus*, nov. gen. (Harpoceratinae,
581 Ammonoidea) du Toarcien d'après un matériel des Causses et du Bas-Languedoc. *Geobios*, 2:7–
582 79.
- 583 Mattei, J., Combémorel, R., and Enay, R. 1987. Sur la présence du genre *Atractites*
584 (Aulacoceratida) dans le Lias moyen des Causses du Sud du Massif central
585 Français. *Geobios*, 20(1), 133–139.
- 586 Mehl, J. 1990. Fossilerhaltung von Kiemen bei *Plesioteuthis prisca* (Rüppell 1829)
587 (Vampyromorpha, Cephalopoda) aus untertithonen Plattenkalken der Altmühlalb.
588 *Archaeopteryx*, 8:77–91.
- 589 Moreau, J.-D., Trincale, V., Deconinck J.-F., Philippe, M., Bourel, B. 2021. Lowermost Jurassic
590 dinosaur ecosystem from the Bleymard Strait (southern France): sedimentology, mineralogy,

- 591 palaeobotany and palaeoichnology of the Dolomitic Formation. *Geological Magazine*,
592 158:1830–1846.
- 593 Moreau, J-D, Vullo, R, Charbonnier, S, Jattiot R, Trincal, V, Néraudeau, D, Fara, E, Baret, L,
594 Garassino, A, Gand, G, and Lafaurie, G. Konservat- Lagerstätten from the Upper Jurassic
595 lithographic limestone of the Causse Méjean (Lozère, southern France): palaeontological and
596 palaeoenvironmental synthesis. *Geological Magazine*, <https://doi.org/10.1017/S0016756821001382>
- 598 Münster, G. Graf zu. 1843. Die schalenlosen Cephalopoden im unteren Jura, den Lias-Schiefern
599 von Franken und Schwaben. *Beiträge zur Petrefaktenkunde*, 6:57–77.
- 600 Naef, A. 1921. Die Cephalopoden: Fauna und Flora des Golfes von Neapel, Monografia 35. R.
601 Friedländer und Sohn, Berlin, 863 p.
- 602 Naef, A. 1922. Die fossilen Tintenfische: eine paläozoologische Monographie. Fischerverlag, G.
603 Fischer, Jena, 322 p.
- 604 Pinard, J. D., Weis, R., Neige, P., Mariotti, N. and Di Cencio, A. 2014. Belemnites from the
605 Upper Pliensbachian and the Toarcian (Lower Jurassic) of Tournadous (Causses, France). *Neues
606 Jahrbuch für Geologie und Paläontologie, Abhandlungen*, 273(2):155–177.
- 607 Quenstedt, F. A. 1839. *Loligo bollensis* ist kein Belemnitenorgan. *Neues Jahrbuch für
608 Mineralogie, Geognosie, Geologie und Petrefaktenkunde*, 156–167.
- 609 Quenstedt, F. A. 1849. Petrefactenkunde Deutschlands, 1. Abteilung, 1. Band, Cephalopoden.
610 Verlag Fues, Tübingen, 581 p.
- 611 Regteren Altena, C. O. v. 1949. Teyler's Museum systematic catalogue of the palaeontological
612 collection—sixth supplement (Teuthoidea). *Archives du Musée Teyler*, 3:53–62.
- 613 Reitner, J. 1978. Ein Teuthiden-Rest aus dem Obernor (Kössener-Schichten) der Lahnewies-
614 Neidernachmulde bei Garmisch-Partenkirchen (Bayern). *Paläontologische Zeitschrift*,
615 52(3/4):205–212.
- 616 Reitner, J. and Engeser, T. 1981. Eine neue Teuthiden-Art aus dem unteren Sinemurium (Lias
617 alpha 3, Ölschiefer) von Dusslingen bei Tübingen (Baden-Württemberg). *Neues Jahrbuch für
618 Geologie und Paläontologie, Monatshefte*, 1981(7):425–430.
- 619 Reitner, J. and T. Engeser. 1982. Teuthiden aus dem Barrême der Insel Maio (Kapverdische
620 Inseln). *Paläontologische Zeitschrift*, 56:209–216.
- 621 Riccardi, A. C. 2005. First teuthid cephalopod from the Lower Jurassic of South America
622 (Neuquén Basin, Argentina). *Geologica Acta*, 3:179–184.
- 623 Riegraf, W., Werner, G. and Lörcher, F. 1984. *Der Posidonienschifer –Biostratigraphie, Fauna
624 und Fazies des südwestdeutschen Untertoarciums (Lias ε)*. Ferdinand Enke, Stuttgart, 195 pp.
- 625 Riegraf, W., Janssen, N. and Schmitt-Riegraf, C. 1998. *Fossilium Catalogus. I: Animalia*. Leiden
626 (Backhuys), 519 pp.
- 627 Robson, G.C. 1929. On the rare abyssal octopod *Melanoteuthis beebei* (sp. n.): a contribution to
628 the phylogeny of the Octopoda. *Proceedings of the Zoological Society of London*, 99(3):469–
629 486.

- 630 Rowe, A. J., Kruta, I., Landman, N. H., Villier, L., Fernandez, V., and Rouget, I. 2022.
631 Exceptional soft-tissue preservation of Jurassic *Vampyronassa rhodanica* provides new insights
632 on the evolution and palaeoecology of vampyroteuthids. *Scientific Reports*, 12(1): 8292.
633 Rowe, A. J., Kruta, I., Villier, L., and Rouget, I. 2023. A new vampyromorph species from the
634 Middle Jurassic La Voulte-sur-Rhône Lagerstätte. *Papers in Palaeontology*, 9(3): e1511.
635 Sciau, J., Crochet, J. Y. and Mattei, J. 1990. Le premier squelette de Plesiosaure de France sur le
636 Causse du Larzac (Toarcien, Jurassique inférieur). *Geobios*, 23(1):111–116.
637 Smith, P.L., Tipper, H.W. and Ham, D.M. 2001. Lower Jurassic Amaltheidae (Ammonitina) in
638 North America: paleobiogeography and tectonic implications. *Canadian Journal of Earth
Sciences*, 38:1439–1449.
640 Starobogatov, Y. I. 1983. The System of the Cephalopoda, p. 4–7. In: Y. I. Starobogatov and K.
641 Nesis (eds.), Taxonomy and ecology of Cephalopoda. Zoological Institute, USSR Academy
642 of Sciences, Leningrad.
643 Trümpy, D.M. 1983. Le Lias Moyen et Supérieur des Grands Causses et de la région de Rodez :
644 contributions stratigraphiques, sédimentologiques et géochimiques à la connaissance d'un bassin
645 à sédimentation marneuse. *Cahiers de l'Université, Université de Pau et des Pays de l'Adour*,
646 19:1–363.
647 Venturi, F., Bilotta, M. and Ricci, C. 2006. Comparison between western Tethys and eastern
648 Pacific ammonites: further evidence for a possible late Sinemurian–early Pliensbachian trans-
649 Pangaea marine connection. *Geological Magazine*, 143 :699–711.
650 Voltz, P. L. 1840. Observations sur les Belopeltis ou lames dorsales de Bélemnites. *Mémoires de
la Société d'Histoire Naturelle de Strasbourg*, 1:1–38.
652 Wagner, A. 1859. Revision der bisherigen systematischen Bestimmungen der Überreste von
653 nackten Dintenfischen aus dem Süddeutschen Juragebirge. *Gelehrte Anzeichen der königlich
654 bayerische Akademie der Wissenschaften, München*, 34:273–278.
655 Wagner, A. 1860. Die fossilen Überreste von nackten Dintenfischen aus dem lithographischen
656 Schiefer und dem Lias des süddeutschen Juragebirges. *Abhandlungen der königlich bayerische
657 Akademie der Wissenschaften, München*, 8:700–821.
658 Weis, R. 1998. Luxemburg für Fossiliensammler. *Fossilien*, 1:13–18.
659 Wilby P. R., J. D. Hudson, R. G. Clements, and N. T. J. Hollingworth. 2004. Taphonomy and
660 origin of an accumulate of soft-bodied cephalopods in the Oxford Clay Formation (Jurassic,
661 England). *Palaeontology*, 45:1159–1180.
662 Woodward, H. 1883. On a new genus of fossil “calamary” from the Cretaceous formation of
663 Sahel Alma, near Beirût, Lebanon, Syria. *Geological Magazine, new series*, 10:1–5.
664 Ziegler, P.A. 1988. Evolution of the Arctic–North Atlantic and the Western Tethys. *AAPG
665 Memoir*, 43:1–198.
666 Zieten, C. H. von. 1830–1833. Die Versteinerungen Württembergs. Verlag & Lithographie der
667 Expedition des Werkes unserer Zeit, Stuttgart, 102 p.
668

669 **Figure captions**

- 670 Figure 1. Geographical location of the Causses basin. The black star indicates the Saint Bauzile
671 site, where the studied material was retrieved (modified after Moreau et al. 2021).
- 672
- 673 Figure 2. Stratigraphic section of the Saint Bauzile site showing the location of the coleoid-
674 bearing bed (black star = “*Leptolepis* bed”). Thi., thickness (m); Form., formations; Lith.,
675 lithology.
- 676
- 677 Figure 3. Morphology, terminology and measurements of the three different gladius
678 morphotypes among Mesozoic gladius-bearing coleoids. Modified after Marroquín et al. (2018).
- 679
- 680 Figure 4. A, B. *Paraplesioteuthis sagittata* (Münster, 1843), specimen M486_2024.1.1 (Lower
681 Toarcian, Saint Bauzile, Causses Basin) in dorsal view, under natural (A) and UV (B) light. C.
682 Interpretative drawing of specimen M486_2024.1.1. D. *Paraplesioteuthis sagittata* specimen
683 from the Lower Toarcian Posidonia Shale Formation, Germany (UMH collection, see also
684 Fuchs, 2020, fig. 4, 1a). E. Gladius reconstruction of *Paraplesioteuthis sagittata* (see also Fuchs,
685 2020, fig. 4, 1b). Scale bars: 10 mm.
- 686
- 687 Figure 5. A, B. ?*Loligosepia* sp. indet., specimen M486_2024.1.2 (Lower Toarcian, Saint
688 Bauzile, Causses Basin) in dorsal view, under natural (A) and UV (B) light. C. Interpretative
689 drawing of specimen M486_2024.1.2. Scale bars: 10 mm.
- 690
- 691 Figure 6. Worldwide occurrences of *Paraplesioteuthis sagittata* (black stars, 1 = western
692 Canada, 2 = southern Germany and 3 = southeastern France) in the paleogeographical context of
693 the Pliensbachian–Toarcian interval. (A) Global paleogeography. (B) Paleogeographical details
694 of the NW Tethyan realm (scale bar = 250 km). Maps are modified after Dera et al. (2011).

Figure 1

Geographical location of the Causses basin

The black star indicates the Saint Bauzile site, where the studied material was retrieved (modified after Moreau et al. 2021).

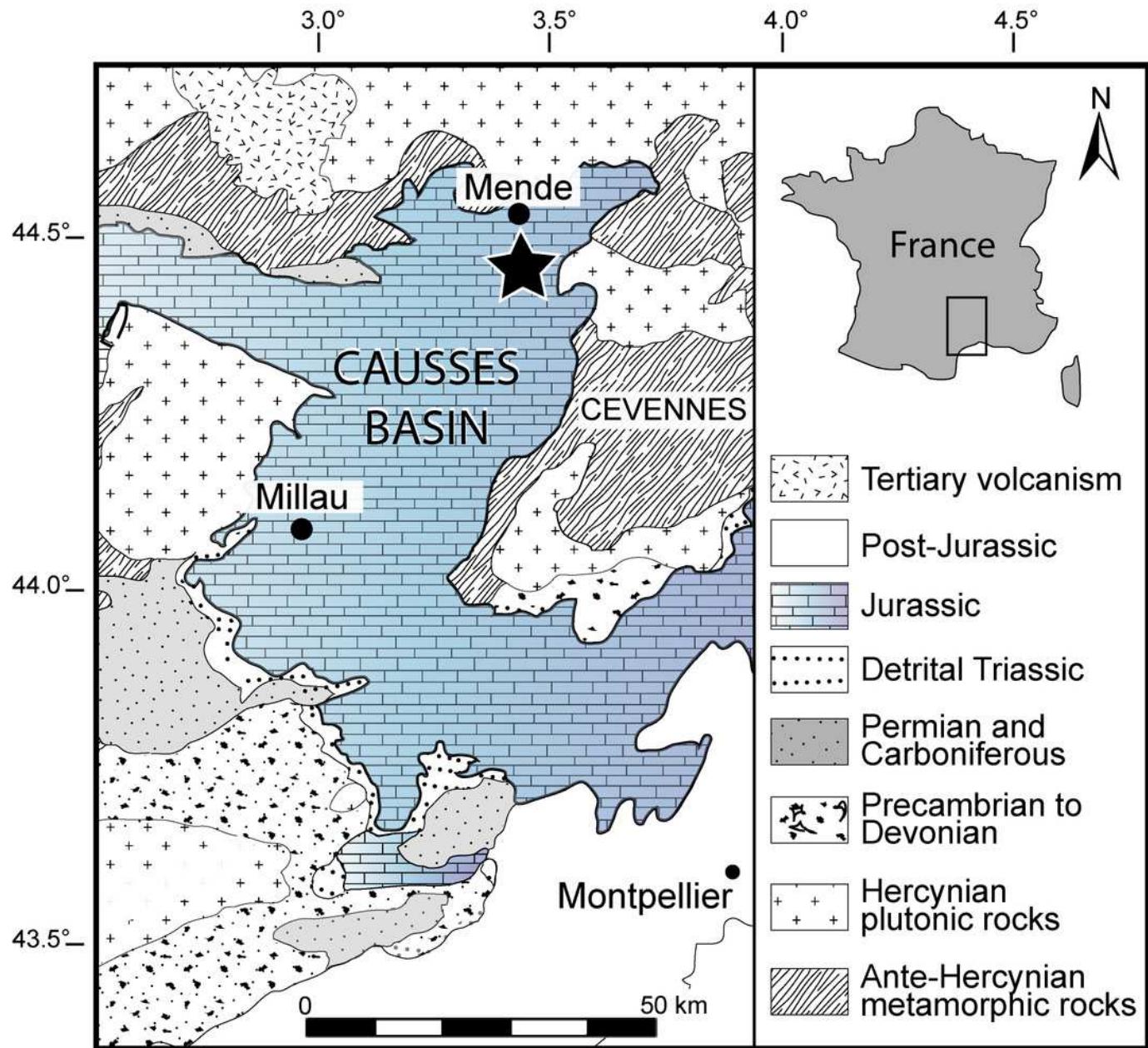


Figure 2

Stratigraphic section of the Saint Bauzile site

Stratigraphic section of the Saint Bauzile site showing the location of the coleoid-bearing bed (black star = “*Leptolepis* bed”). Thi., thickness (m); Form., formations; Lith., lithology.

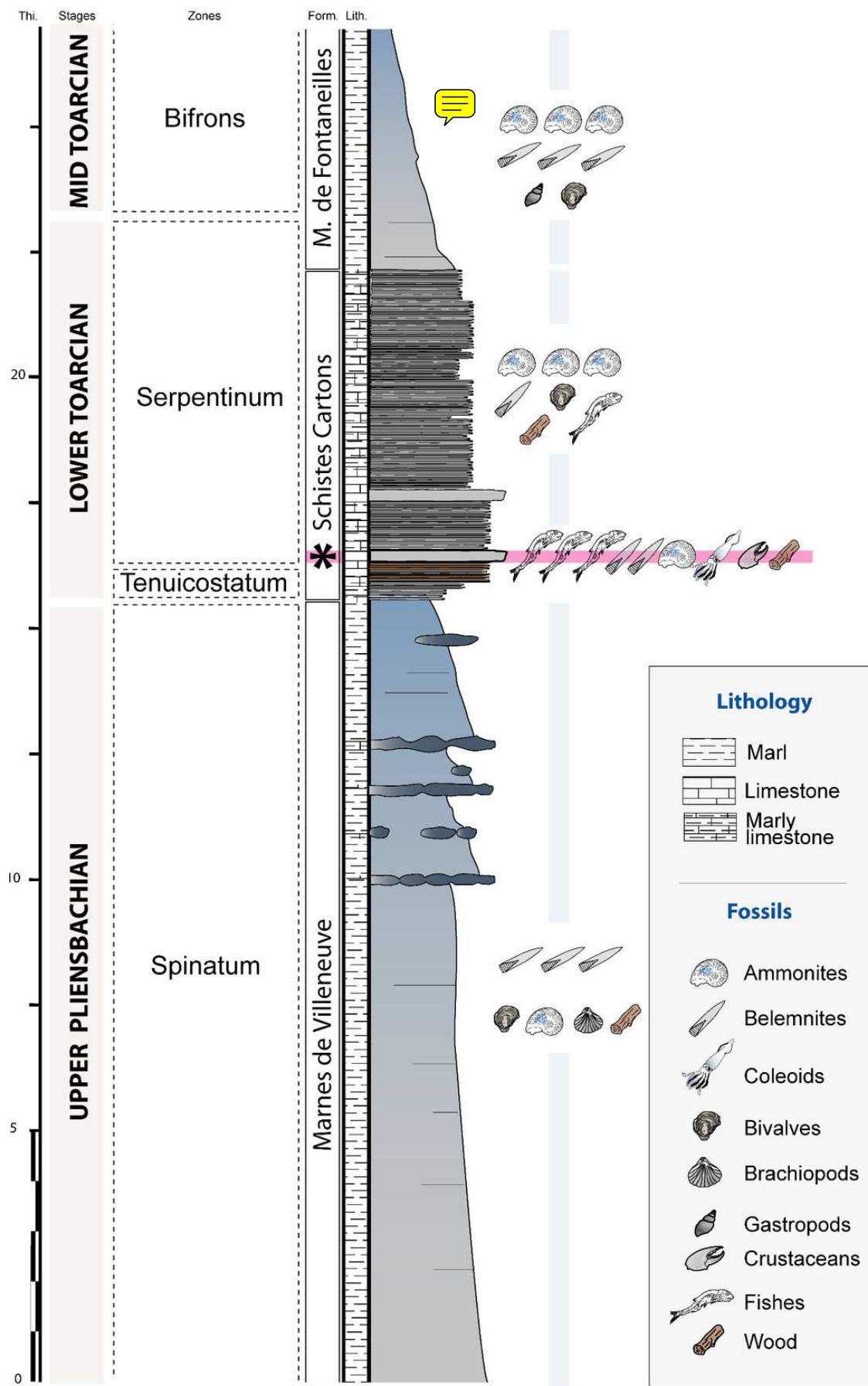


Figure 3

Morphology, terminology and measurements of the three different gladius morphotypes among Mesozoic gladius-bearing coleoids

Modified after Marroquín et al. (2018)

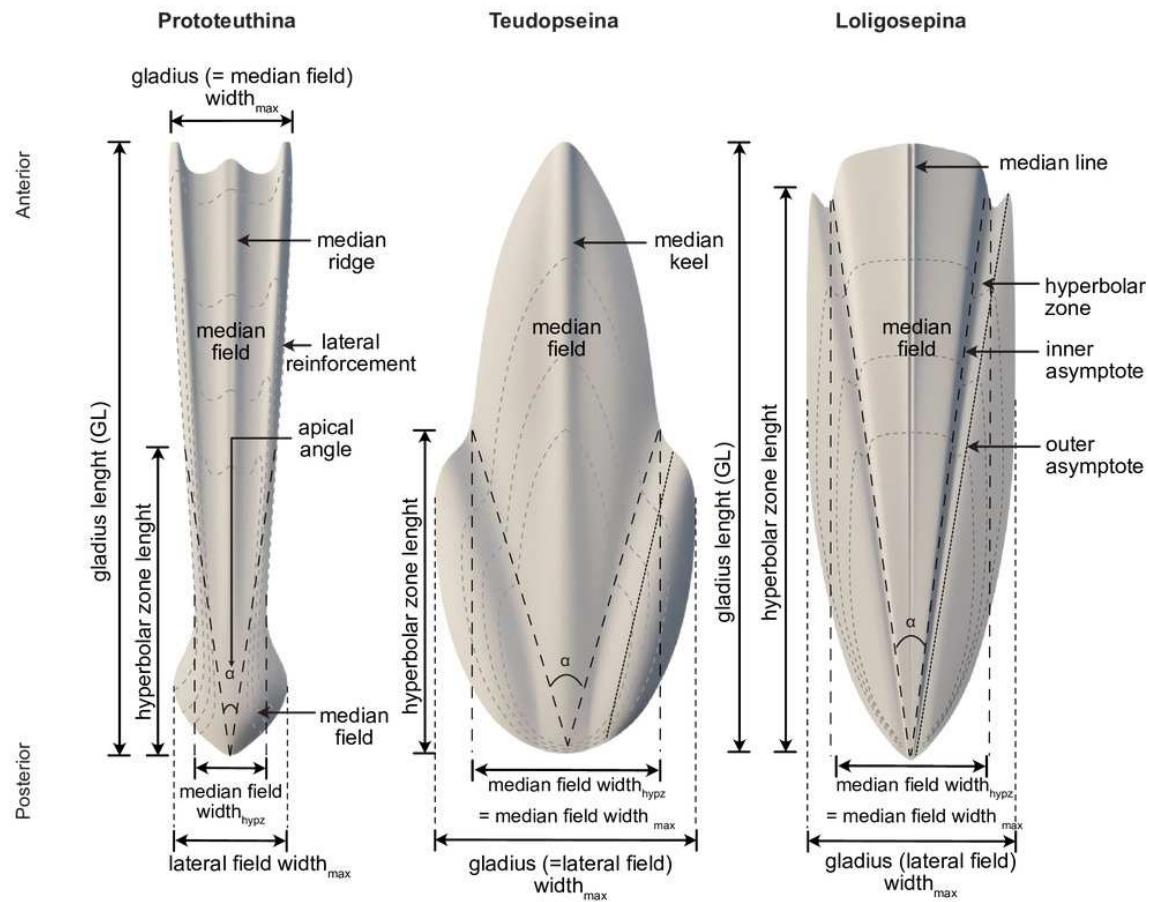


Figure 4

Paraplesioteuthis sagittata (Münster, 1843)

A, B. *Paraplesioteuthis sagittata* (Münster, 1843), specimen M486_2024.1.1 (Lower Toarcian, Saint Bauzile, Causses Basin) in dorsal view, under natural (A) and UV (B) light. C. Interpretative drawing of specimen M486_2024.1.1 . D. *Paraplesioteuthis sagittata* specimen from the Lower Toarcian Posidonia Shale Formation, Germany (UMH collection, see also Fuchs, 2020, fig. 4, 1a). E. Gladius reconstruction of *Paraplesioteuthis sagittata* (see also Fuchs, 2020, fig. 4, 1b). Scale bars: 10 mm.

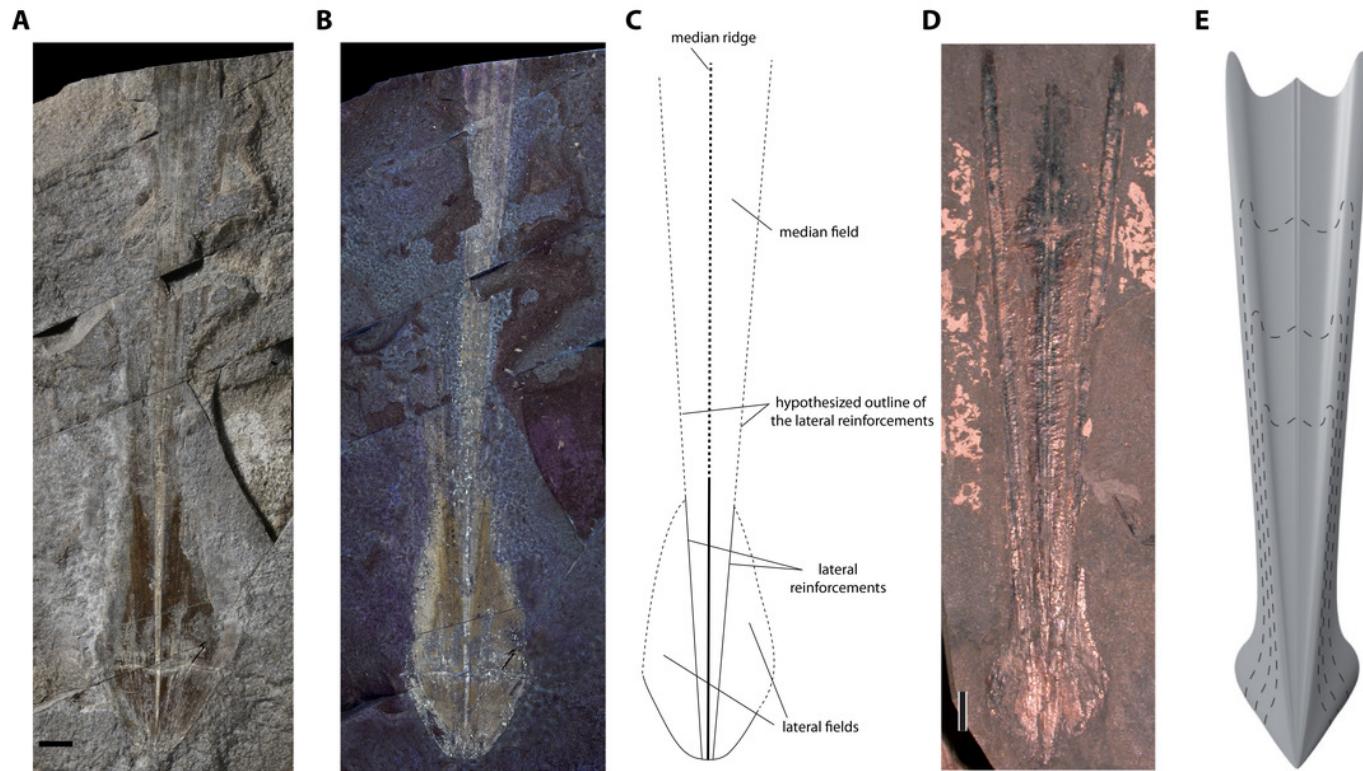


Figure 5

?*Loligosepia* sp. indet

A, B. ?*Loligosepia* sp. indet., specimen M486_2024.1.2 (Lower Toarcian, Saint Bauzile, Causses Basin) in dorsal view, under natural (A) and UV (B) light. C. Interpretative drawing of specimen M486_2024.1.2. Scale bars: 10 mm.

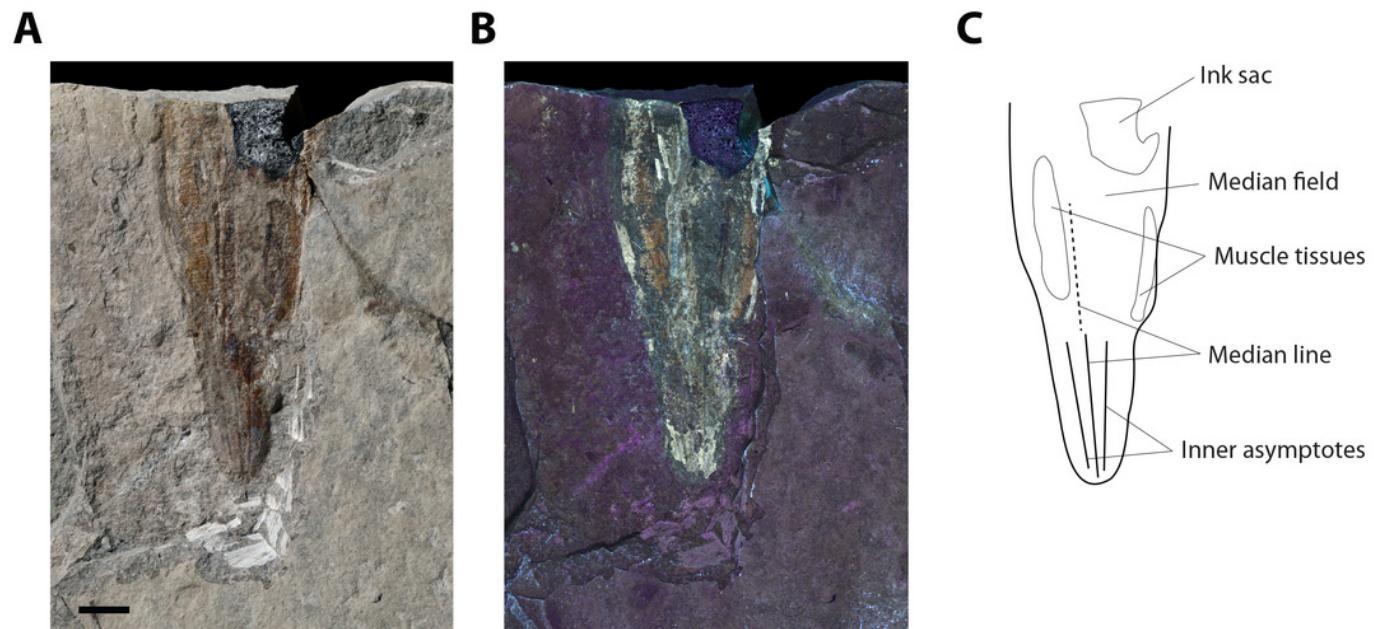


Figure 6

Worldwide occurrences of *Paraplesioteuthis sagittata*

Worldwide occurrences of *Paraplesioteuthis sagittata* (black stars, 1 = western Canada, 2 = southern Germany and 3 = southeastern France) in the paleogeographical context of the Pliensbachian–Toarcian interval. (A) Global paleogeography. (B) Paleogeographical details of the NW Tethyan realm (scale bar = 250 km). Maps are modified after Dera et al. (2011).

