The first Palaeodictyoptera (Insecta) from the Carboniferous-Permian basin of Graissessac (France)

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1 Historical biology

2
3 **The first Palaeodictyoptera (Insecta) from the Carboniferous-Permian basin**
4 **of Graissessac (France)**
5

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8

9 Running head

10 Palaeodictyopteran insect from Graissessac
11

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ABSTRACT

A new dictyoneurid insect, *Dictyoneura goujonorum* n. sp. from the Latest Ghzelian – Asselian basin of Graissessac (Hérault, France) is described in details. It is represented by a well-preserved specimen with wings of 32-35 mm long and 13-14 mm wide and other peculiar diagnostic characters such a MP with four branches and a CuP with three branches. As all the other *Dictyoneura* species are known from the Namurian and/or the Wesphalian, *Dictyoneura goujonorum* n. sp. is the youngest representative of the genus. It is also the first record of the order Palaeodictyoptera from the Graissessac basin. The Carboniferous-Permian palaeodictyopterans are well-known to have lived in rather humid swamp forests. The global warming and drying of the climate during the Permian and/or the rise of potential predators may be responsible of their extinction.

KEYWORDS

Insecta; Palaeoptera; Carboniferous-Permian; Capitanian extinction; climatic changes; France.

Introduction

The Palaeodictyoptera are major herbivorous sucking insects known from the Late Carboniferous to the Early Permian. They were well-diversified, with at least known 50 genera in about 16 families (see Fossilworks database at <http://fossilworks.org>). However the palaeodictyopteran species are generally represented by very few specimens in comparison to other clades such as the Dictyoptera or the Archaeorthoptera (A.N. pers. obs.). Thus, any new fossil of these strange six-winged insects is welcome to improve our knowledge on their diversity and temporal range. Palaeodictyoptera are known of the swamp forests of the Late

Carboniferous, with a maximum of diversity in the well-known outcrop of Commentry (northern part of the French Massif Central) (Kukalová 1969a-b, 1970). They are generally associated with the warm and humid evergreen coal forests, together with some of their emblematic predators, the giant odonatopterans Meganeuridae called griffenflies. Some Palaeodictyoptera are also very large insects (e.g. *Homoioptera gigantea* Agnus, 1902 with a wing span of ca. 40 cm), but most of them were relatively small, with wing span ca. 10 cm.

The Latest Carboniferous – Earliest Permian Graissessac basin (Hérault, France) is a quite favorable area to find fossils of plants. Numerous geological researches, mainly linked to the exploitation of the coal, have been undertaken since the 19th century. The collect of fossil animals is decreasing since these last decades with the closure of these coal mines. The insect record is especially scarce. Thanks to the efforts of Mr and Mrs Goujon Claude and Monique, more fossils have been found recently. Here we describe in details the first Palaeodictyoptera from the Graissessac basin. It is also the first fossil insect properly described for this outcrop. It is compared with all the other dictyoneurid genera. This new finding addresses our knowledge on the Carboniferous-Permian fauna of Graissessac. It is also the latest occurrence of this genus and therefore contribute to document the putative extinction of the clade before the Middle Permian.

Material and methods

The type specimen was collected in 2019 by Goujon and Monique Claude, in the dump issued of the open coal mine named ‘Carrière de Layrac’ (North East of the town of Graissessac). The Graissessac basin is situated in the southern part of the Massif Central, France. Its general orientation is west to east, forming a long synclinal of 30 km long and 2.5 km wide (Fig. 1) (Martín-Closas and Galtier 2005: fig. 1). Becq-Giraudon (1973) considered this basin as a channel of sedimentation flow and plant debris-flow alimenting a larger Carboniferous basin

located under the Permian sediments of Lodève. Seven to eight coal layers have been identified, one of them being up to six meters thick on average. They are interbedded between layers of sandstone and shale. The recent sedimentological study of these rocks allows to recognize a great diversity of depositional environments: torrential alluvial cones, fluvial environments of variable energy, flood plains and swamps in an intra-mountain sedimentary basin (Saint Martin 1993; Rilliart 2013). These environments may have developed in a relatively low basin region framed by very high mountains (Becq-Giraudon and Van den Driessche 1993), forming possibly what we called here a “lost valley”. Indeed sedimentological data (exhaust craters, solifluxion flows, ice crystal ghosts) suggest the presence of periglacial phenomena and a rather cool paleoclimate for the Graissessac area (Becq-Giraudon and Van Den Driessche 1994; Becq-Giraudon et al. 1996).

Concerning the age of the Graissessac basin, Bruguier et al. (2003) estimated it around 295.5 ± 5.1 Ma, that-is-to-say Asselian (earliest Permian) based on zircon geochronology. However its plant assemblage suggests a late Gzhelian (latest Carboniferous) age (Poschmann et al. 2016). This Graissessac flora includes about 70 taxa of lycopsids, sphenopsids, pteridosperms, and prephanerogams (Grand'Eury 1877; Becq-Giraudon 1973; Doubinger, 1983 Poschmann et al. 2016): these taxa are relatively well-known in the Carboniferous-Permian rocks of the other Massif Central and European basins (e.g. Doubinger et al. 1995). The presence of very large trunks of *Sigillaria* and *Calamites* suggests an environment of flood plains and swamps, also supported by Martín-Closas and Galtier (2005).

The animal fossil record is reduced. Mostly invertebrate have been recovered: Mollusca: *Carbonicola* sp. (Becq-Giraudon 1973); Arthropoda: xiphosuran *Euproops mariae* (Crônier and Courville, 2005), Chelicerata *Aenigmatarbus rasteli* Poschmann et al., 2016 and Scorpiones *Eoscorpius* sp., *Compsoscorpius* sp. (Poschmann et al. 2016); Insecta: Paoliida *Blattinopsis* sp., a wing of a Polyneoptera ‘Prothoptera’ (Becq-Giraudon 1972) and the endophytic

oviposition of an insect, possibly an Odonatoptera (Béthoux et al. 2004); Vertebrata: ‘fishes’ have been mentioned but not described (Bergeron 1889; Louis 1954) and a branchiosaurid recently excavated is currently under description by us. Trace fossils have also been found in the Graissessac basin with the presence of trackways of the millipede-like *Arthropleura* sp. (Moreau et al. 2019). Compared with the faunas of the other Massif Central basins, this fauna is reduced in specimen number but well-diversified, with representatives of several major groups of organisms. This suggests that the biota of the Graissessac “lost valley” already presents complex trophic chains.

The photographs were taken using Nikon D800 camera with 60 mm 2,8 Micro Nikkor lens in the 3D Lab (ISYEB, MNHN), and the reconstruction of the venation was done under a binocular microscope Nikon SMZ 1500, with a camera lucida.

We follow the wing venation terminology of Kukalová-Peck (1991).

Institutional abbreviations.—MHNE, Muséum d’Histoire Naturelle et d’Ethnologie of Colmar, France.

Other abbreviations.—A anal veins; C costal vein; CuA cubitus anterior; CuP cubitus posterior; MA median anterior; MP median posterior; RA radius anterior; RP radius posterior; ScP subcostal posterior.

Nomenclatural acts.—This published work and the nomenclatural acts it contains, have been registered in urn:lsid:zoobank.org:pub:9220BD12-40CC-4181-8F90-798E433CDE55

Systematic palaeontology

Order: Palaeodictyoptera Goldenberg, 1877

Superfamily: Dictyoneuroidea Handlirsch, 1906a

Family: Dictyoneuridae Handlirsch, 1906a

126 *Dictyoneura* Goldenberg, 1854

127 *Type species*

128 *Dictyoneura libelluloides* Goldenberg, 1854 (Westphalian D, Saar basin, Germany)

129

130 *Other species*

131 *Dictyoneura kemperi* Brauckmann and Koch, 1983 (Namurian, Hagen-Vorhalle, Germany),

132 *Dictyoneura nigra* Kliver, 1883 (Westphalian C, Saar basin), and *Dictyoneura goujonorum* n.

133 sp.

134

135 *Dictyoneura goujonorum* n. sp.

136 (Figure 2)

137 urn:lsid:zoobank.org:act:53F972EE-42A0-4F24-8A86-17D2A9073275

138 *Etymology*

139 Named after Goujon Claude and Monique who found the type specimen.

140

141 *Type material*

142 MHNE.2021.3.1 (imprint and counter-imprint of a fore- and a hind wing in life position with

143 fragments of abdomen and thorax), stored at the Muséum d'Histoire Naturelle et d'Ethnologie

144 of Colmar, France.

145

146 *Type locality*

147 Quarry of Layrac, Graissessac, Hérault, France (GPS coordinates available to qualified

148 researchers).

149

150 *Stratigraphic occurrence*

151 Latest Gzhelian to Asselian.

152

153 *Diagnosis*

154 Wings ca. 32.0-35.0 mm long, and 13.0-14.0 mm wide; ScP elongate; RP not pectinate; MA
155 simple; MP with four branches; CuA simple; CuP with three branches; bases of CuA, MA and
156 RP approximate.

157

158 *Description*

159 Wings without trace of coloration; forewing ca. 35.0 mm long, 13.0 mm wide; anterior margin
160 not curved basally; costal area 1.0 mm wide; ScP at least 29.0 mm long, terminating on C well
161 beyond midwing; RA simple; base of RP 11.0 mm from wing base; RP not pectinate, with two
162 main branches, anterior one forked and posterior one with two-three branches; MA unbranched,
163 weakly curved basally, separated from MP slightly basal to base of RP, 1.0 mm apart; MP with
164 two main branches, each of them being forked again; CuA unbranched, weakly curved basally,
165 separated from CuP slightly basal to base of MA, 2.0 mm apart; CuP with two-three branches;
166 anal area 4.0 mm wide, anal veins poorly preserved; archaedyction present but poorly visible.
167 Hind wing ca. 32.0 mm long, 14.0 mm wide; triangular-shaped and with a broader anal area
168 than in forewing; costal area as broad as that of forewing, 1.0 mm wide; venation identical to
169 that of forewing; cubito-anal area broader than that of forewing, 6.0 mm wide.

170

171 **Discussion**

172 This fossil belongs to the Palaeodictyoptera rather than to the other orders of the
173 Palaeodictyopterida because of the dense venation, with numerous branches of main veins, and
174 the simple pattern of branching of RP and MA (Carpenter 1992). Riek (1976) proposed a key
175 to palaeodictyopteran superfamilies, after which *Dictyoneura goujonorum* n. sp. would fall in

the Dictyoneuroidea, because of the following characters: CuA and MA simple; ScP separated from R; Archaedictyon present. But after Riek (1976), the representatives of this superfamily would also have MP simple or three-branched, which is not the case for the type genus *Dictyoneura* and *Dictyoneura goujonorum* n. sp.

The current classification of the Palaeodictyoptera is not satisfactory. Sinitshenkova (2002: fig. 138) first proposed a phylogenetic hypothesis, but the absence of real outgroup(s) to basically polarize the character states, and the basal-most dichotomy established on the character ['wings wide basally' vs. 'wing base narrow'], with the two states supposedly supporting the two branches, while one should be plesiomorphic, prevent a total confidence to these results. Sroka et al (2015: fig. 11) proposed another hypothesis, better supported by true outgroups and computer treatment of the data, in which the order Palaeodictyoptera falls as a grade, sister group of the (Megasecoptera + (Permothemistidae + Diaphanopteroidea)).

A comparison of this fossil to all the currently accepted families (as listed in the fossilworks database <http://fossilworks.org>) is necessary:

- the 'anterior margin of the wing not curved basally' excludes the Lithomanteidae Handlirsch, 1906;
- the posteriorly curved branches of M and Cu exclude the Megaptilidae Handlirsch 1906;
- the less elongated wings, with only three branches of MP instead of six, and the shorter abal area differs from *Archaemegaptilus kiefferi* Meunier, 1908 (Kukalová 1969: fig. 46) and exclude the Archaemegaptilidae Handlirsch, 1919;
- the 'RP and MA not coalescent nor strongly approximate' excludes the Eugereonidae Handlirsch, 1906a;
- the elongate ScP excludes the Calvertiellidae Martynov, 1931 and the Stobbsiidae Handlirsch, 1908 (Laurentiaux and Laurentiaux-Vieira, 1951: fig. 5);
- the 'branched MP' excludes the Tchirkovaeidae Sinitshenkova, 1979;

201 - the 'CuA simple' excludes the Homiopteridae Handlirsch, 1906a, Spilapteridae
 202 Handlirsch, 1906a, Fouqueidae Handlirsch, 1906a, Elmoboriidae Carpenter, 1976;
 203 - the 'MA simple' excludes the Homothetidae Scudder, 1885 based on a poorly known
 204 fossil;
 205 - the 'CuA simple' and the 'MA simple' exclude the Pteronidiidae Bolton, 1912,
 206 Mecynostomatidae Handlirsch, 1904, Straeleniellidae Laurentiaux-Vieira and
 207 Laurentiaux, 1986, and Eubleptidae Handlirsch, 1906a;
 208 - the 'CuA simple', and the broader and shorter wings and shorter cubito-anal area
 209 exclude the Peromapteridae Handlirsch, 1906a;
 210 - the 'branched CuP and RP with more than three branches' excludes the Psychroptilidae
 211 Riek, 1976 (originally considered as a Megasecoptera, but put in the Palaeodictyoptera
 212 in the fossilworks database <http://fossilworks.org>);
 213 - the 'MA not arising in a distal position, and not on first branch of MP' exclude' the
 214 Lycocercidae Handlirsch, 1906a;
 215 - the 'RP posteriorly pectinate' excludes the Saarlandiidae Guthörl, 1930 and the
 216 Mongolianidae Özdikmen 2008 (replacement name for Mongolodictyidae
 217 Sinitshenkova, 1992), two poorly known families;
 218 - the 'base of MA basal to that of RP' excludes the Heolidae Handlirsch, 1906a and the
 219 Graphiptilidae Handlirsch, 1906a;
 220 Finally, The Jongmansiidae Laurentiaux, 1950 have a reduced RP. The Namuroningxiidae
 221 Prokop and Ren, 2007 have a richer venation, with bases of RP and MA at the same level, and
 222 much more branches of RP, MP, and CuP. The Polycraegridae Handlirsch, 1906b have much
 223 more branches of RP and MP and a forked CuA (Handlirsch 1906b). The Synarmogidae
 224 Handlirsch, 1910 have the median vein appressed to the radial one.

Affinities with the Breyeriidae Handlirsch, 1906a are more complicate to exclude. They generally have long crossveins between the main veins and their branches (viz. in *Breyeria* de Borre, 1875, *Jugobreyeria* Brauckmann et al., 1985, *Hasala* Brauckmann, 1995, *Megaptiloides* Handlirsch, 1906, and *Vermooija* Prokop et al., 2018), not present in *Dictyoneura goujonorum* n. sp. Thus this character would be sufficient to exclude affinities of *Dictyoneura goujonorum* n. sp. with the Breyeriidae. But in *Aviobreyeria* Prokop et al., 2013, these crossveins are mainly present in the distal half of the wing while there is an archedictyon in basal part of wing. The Breyeriidae can have a short ScP, ending at the level of first branch of anterior branch of RP (in *Breyeria*, *Jugobreyeria*) or an elongate one (in *Hasala*, *Aviobreyeria*, *Megaptiloides*, and *Vermooija*) (Laurentiaux and Laurentiaux-Vieira, 1951; Brauckmann et al., 1985; Brauckmann, 1995; Prokop et al., 2013, 2018). *Hasala* and *Aviobreyeria* have a pectinate RP, unlike *Dictyoneura goujonorum* n. sp.

Dictyoneura goujonorum n. sp. has all the diagnostic characters of the Dictyoneuridae (as proposed by Carpenter 1992: 28). They share the absence of well-defined crossveins all over the wings (Guthörl 1934). It fits with the genus *Dictyoneura* (see diagnoses in Waterlot 1934: 139 or Guthörl 1934), while it strongly differs from the other genera of this family in the combination of the following characters: RP not posteriorly pectinate, base of CuA nearly at the same level as those of RA/RP and MA/MP, base of MA slightly more basal than that of RP, CuP not simple, main veins posteriorly curved, ScP elongate, no trace of crossveins in costal area.

Brauckmann and Koch (1983), following Guthörl (1934), made a revision of *Dictyoneura*, with the three species *Dictyoneura libelluloides*, *Dictyoneura kemperi*, and *Dictyoneura nigra*. Waterlot (1934) included the two species *Dictyoneura rugosa* Handlirsch, 1906a and *Dictyoneura sinuosa* Kliver, 1883 in this genus. Lastly *Dictyoneura higginsii* Handlirsch 1906a is a basal fourth of a wing, impossible to compare to the other species in the

genus. *Dictyoneura sinuosa* probably does not belong to this genus because its base of CuA and of RP are well basal to that of MA (Waterlot 1934: pl. 17, fig. 1). Also, *Dictyoneura rugosa* (based on a fragment of mid part of wing) probably does not belong to this genus because the base of CuA is well basal to that of MA (Waterlot 1934: text-fig. 43). *Dictyoneura kemperi* has five branches of MP, unlike four in the other species and *Dictyoneura goujonorum* sp. nov. But the value of this character remains uncertain because of the lack of information on the variations in the number of branches of the main veins in these insects. *Dictyoneura kemperi* has forewings ca. 43 mm long, 16 mm wide, and the hind wings ca. 39 mm long and 16 mm wide. *Dictyoneura libelluloides* has wings more than 55 mm long and 22 mm wide, after Waterlot (1934: 141), and/or wings more than 57 mm long and 22 mm wide, after Guthörl (1934: 52). *Dictyoneura nigra* has wings more than 44 mm long and 18 mm wide after Waterlot (1934: 142), and/or more than 35 mm long and 18 mm wide, after Guthörl (1934: 53). *Dictyoneura goujonorum* n. sp. is clearly a smaller insect with forewing ca. 35 mm long, 13 mm wide, and hind wing ca. 32 mm long, 14 mm wide.

Conclusion

All the previously described *Dictyoneura* spp. are Pennsylvanian ('Namurian B' or 'Westphalian C or D'), while *Dictyoneura goujonorum* n. sp. is latest Gzhelian to Asselian, thus the most recent known representative of the genus. The Namurian *Dictyoneura kemperi* is supposed to have lived in a moist area with rich vegetation along a coastal area in a Variscan foreland basin (Brauckmann 1988). *Dictyoneura libelluloides* was found in rocks formed in the Westphalian D swamp forests of the intramontane Saar basin (Uhl and Cleal, 2010), similarly to *Dictyoneura goujonorum* n. sp. *Dictyoneura nigra* was also found in the Westphalian C of the same area. Thus all these insects probably lived in similar environments, humid but possibly not very warm, even cool. This possible cool temperature could be linked with the proximity

of relatively high mountains, as suggested by Becq-Giraudon and Van den Driessche (1993) for the Graissessac basin.

The fossil record of the family Dictyoneuridae ranges between the ‘Namurian’ and the Artinskian (in Central Siberia and Northern China) (Sharov and Sinitshenkova 1977; Hong, 1985). Thus they possibly disappeared because of the global warming which started during the Early Permian and continued after, during the whole Permian. The mean annual temperatures of the period during which they are currently recorded seem to have been under 18°C (Scotese et al. 2021: fig. 6). The floristic changes to a ‘seed plant-dominated world’, due to a supposed global drying, would have happened earlier, around the Permian-Carboniferous boundary (DiMichele et al. 2001). But it seems that the drying was not uniform during the Early Permian, much more serious in the Western part of Pangea than in North China (Yang et al. 2016). Also, the central Siberia was clearly less arid than the central Pangea during the Early to middle Permian (Fujimoto et al. 2012), possibly allowing the survival of the dictyoneurids in these regions at least till the end of the Early Permian. Nevertheless, it seems that the *Dictyoneura* spp. that were living in the humid (and possibly relatively cool) forests did not survive the earliest Permian. Another explanation for the apparent extinction of these insects could be the rise of some of their predators such as the reptiles, whose diversity increased during the Carboniferous-Permian. But, as some other palaeodictyopterans (e.g some Calvertiellidae; Béthoux et al. 2007) of the same range of body size survived till the end of the Middle Permian, this hypothesis is more unlikely.

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Figure 1. Geological map of Carboniferous Graissessac basin (modified from Martín-Closas and Galtier 2005: fig. 1; Becq-Giraudon 1973; Saint Martin 1973; Pfeifer et al. 2018).

Figure 2. *Dictyoneura goujonorum* n. sp., holotype: MHNE.2021.3.1. Imprint. (A) forewing; (B) hind wing; (C) forewing under alcohol; (D) hind wing under alcohol; (E) drawing of forewing. Scale bars 10 mm.