

1 The literature on Triassic, Jurassic and earliest Cretaceous dinoflagellate cysts: supplement 4

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## 10 **ABSTRACT**

11 Since the publication of four compilations issued between 2012 and 2019, 93 further  
12 published contributions on Triassic, Jurassic and earliest Cretaceous (Berriasian)  
13 dinoflagellate cysts from Africa, North America, South America, the Arctic, Australasia, East  
14 Europe, West Europe, the Middle East and Russia have been discovered in the literature, or  
15 were issued in the last 12 months (i.e. between February 2018 and January 2019). Of these,  
16 55 were published during 2018 and 2019, making this period a very productive one. These  
17 studies are mostly on the Late Triassic and Early Jurassic of Europe. All the 93 items are  
18 listed herein with digital object identifier (doi) numbers where available, as well as a  
19 description of each item as a string of keywords. Publications on West Europe comprise  
20 31.2% of the total, and items on Africa, the Arctic, Australasia, East Europe and Russia are  
21 also significant (15.1%, 6.5%, 7.5%, 9.7% and 14.0% respectively). The least well-  
22 represented regions are North America, South America and the Middle East (2.2%, 1.1% and  
23 1.1% respectively).

24

25 **KEYWORDS** dinoflagellate cysts; earliest Cretaceous (Berriasian); Jurassic; literature  
26 analysis and compilation; Triassic; worldwide

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## 29 **1. Introduction**

30 The literature on Triassic to earliest Cretaceous (Berriasian) dinoflagellate cysts is extensive,  
31 and was listed and reviewed by Riding (2012, 2013, 2014, 2019). These four papers cited  
32 1347, 94, 89 and 266 publications respectively, with each citation followed by keywords  
33 detailing the scope of each of the 1796 studies. The reviews provided by Riding (2014, 2019)  
34 were substantially more interpretive than those in Riding (2012, 2013); the former two papers

35 reviewed and summarised each of the major publications listed. During the 12 months since  
36 the completion of Riding (2019), i.e. between February 2018 and January 2019, the author  
37 has compiled a further 93 relevant articles. Of these, 55 are recently published papers; the  
38 other 38 were previously overlooked. Thirty of the 93 items are considered to be of  
39 substantial scientific significance. The total of 55 articles published between February 2018  
40 and January 2019 makes this one of the most productive periods on this topic in recent years.

41 The 93 articles are largely on the Late Triassic and Early Jurassic of Europe  
42 (Tables 1, 2), and are listed in Appendix 1 of the Supplementary data. Papers on West Europe  
43 are most numerous (29), and comprise 31.2% of the overall total (Table 1). By contrast,  
44 Riding (2012, 2013, 2014, 2019) noted a substantial bias towards the Late Jurassic of Europe.  
45 Publications on Africa, the Arctic, Australasia, East Europe and Russia are also numerous  
46 (15.1%, 6.5%, 7.5%, 9.7% and 14.0% respectively; Table 1). Finally, relatively low  
47 proportions of articles are on North America, South America and the Middle East (2.2%,  
48 1.1% and 1.1% respectively; Table 1). In this compilation, six formally unpublished PhD  
49 theses are listed (e.g. Ruckwied 2009, Baranyi 2018, Correia 2018); these are all freely  
50 available online and the respective web addresses are quoted.

51

## 52 **2. Regional review and synthesis**

53 In this section, brief commentaries/reviews of selected articles from the 93 publications listed  
54 in Appendix 1 of the Supplementary data are presented. These items are from nine of the 14  
55 geographical subdivisions in Riding (2019). In the present compilation, there are no relevant  
56 single-region publications from Central America, Antarctica, Southeast Asia, China and the  
57 Indian subcontinent. Each contribution in Appendix 1 of the Supplementary data is referred  
58 to at least one of these 14 regions; furthermore, ‘multi-region’ and ‘no geographical focus’  
59 are also options (Table 1).

60 The publication by Mangerud et al. (2019) is a good example of the latter two  
61 categories. This article is a global synthesis of the available literature on Triassic  
62 dinoflagellate cysts; it reviewed data from Arabia, the present Arctic region, Europe, Oceania  
63 and South America. It is clear that, with the exception of *Sahulidinium ottii* in one well in  
64 offshore Australia, dinoflagellate cyst body fossils first appeared during the Late Triassic.  
65 The Norian–Rhaetian genus *Rhaetogonyaulax* appears to be a cosmopolitan pioneer taxon.  
66 There was migration into many formerly land-locked regions during the Rhaetian, and most  
67 Triassic dinoflagellate cyst taxa became extinct at the end-Triassic event. Other examples of  
68 ‘multi-region’ and ‘no geographical focus’ papers include Boersma et al. (1987), Lindstrom

69 et al. (2017), Londeix (2018) and Penaud et al. (2018). All dinoflagellate cysts and other  
70 palynomorph taxa at and below species level mentioned in this paper are listed in Appendix 2  
71 of the Supplementary data with full author citations.

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## 74 **2.1. Africa**

75 This compilation includes 14 single-region contributions from East and North Africa,  
76 including five that are deemed especially significant (Appendix 1 of the Supplementary data).  
77 The highlights of this research are outlined in the next two subsections.

78

### 79 **2.1.1. East Africa**

80 In this synthesis, four single-region contributions on Ethiopia and Tanzania in East Africa are  
81 considered. Msaky (2008) is a thesis on the Bajocian to Cenomanian palynology of coastal  
82 Tanzania, and is available online. The thesis was published as Msaky (2011a, 2011b), and  
83 these major publications were reviewed by Riding (2019).

84 The palynofloras of the Pindiro Group (Triassic to Lower Jurassic) of southern  
85 Tanzania were studied by Hudson and Nicholas (2014). These authors reported the  
86 dinoflagellate cysts *Dapcodinium priscum*, *Sahulidinium ottii*, *Scriniocassis* sp. cf. *S. weberi*  
87 and *Sverdrupiella* sp. from the Mbuo Formation (Hudson and Nicholas 2014, p. 59). This  
88 assemblage was interpreted as being Late Triassic in age. The presence of *Dapcodinium*  
89 *priscum* and *Sverdrupiella* sp. is consistent with this age determination. However,  
90 *Sahulidinium ottii* and *Scriniocassis* sp. cf. *S. weberi* are indicative of the Middle Triassic and  
91 the late Pliensbachian to Aalenian respectively (Helby et al. 1987, Riding and Thomas 1992).  
92 If confirmed, this report would be the first record of *Sahulidinium ottii* since this species was  
93 first described by Stover & Helby (1987). *Nannoceratopsis pellucida* was recorded from the  
94 Mihambia Formation by Hudson and Nicholas (2014, p. 65). The Mihambia Formation was  
95 interpreted as being Toarcian to Aalenian in age. Either the interpreted age, or the  
96 identification of *Nannoceratopsis pellucida* appears to be erroneous because the range base of  
97 this species in both hemispheres is Bathonian (Riding et al. 1985, Riding et al. 2010). It  
98 should be noted that the ‘probable reworked dinoflagellate’ figured by Hudson and Nicholas  
99 (2014, fig. 3.5M) is an indeterminate palynomorph, and has no demonstrable dinoflagellate  
100 affinity.

101 Smelror et al. (2018) is a relatively short paper on the Upper Jurassic and Lower  
102 Cretaceous palynostratigraphy of the Kipatimu, Mitole, Nalwehe and Kihuluhulu formations

103 of the onshore Mandawa Basin in southeastern coastal Tanzania. The authors concluded that  
104 the four formations span the Oxfordian–Tithonian to Aptian–Albian interval. Jurassic and  
105 earliest Cretaceous dinoflagellate cysts were recorded only from the Mitole Formation, and  
106 these were interpreted as being of Oxfordian to Berriasian age. They include *Canningia*  
107 *reticulata*, *Circulodinium distinctum*, *Cribroperidinium* spp., *Dingodinium jurassicum*,  
108 *Kaiwaradinium scruttinum* and *Systematophora areolata*. This assemblage is significantly  
109 reminiscent of the Late Jurassic and Early Cretaceous of Gondwana (Helby et al. 1987,  
110 Riding et al. 2010). Sample WP232-5-14 from the Mitole Formation contains a marine  
111 palynoflora reminiscent to the *Dingodinium jurassicum*-*Kilwacysta* assemblage of Schrank  
112 (2005), and is indicative of a correlation with the *Trigonia smeei* Bed of Tendaguru Hill in  
113 southeastern Tanzania.

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### 116 **2.1.2. North Africa**

117 Ten single-region contributions on North Africa are included herein. Nine of the articles are  
118 on northern Egyptian material, which reflects the intense hydrocarbon exploration and  
119 production activity in this region. One contribution (Jaydawi et al. 2016) is a study of  
120 Moroccan material.

121 Aboul Ela and Tahoun (2010) documented the stratigraphical palynology of the  
122 Middle Jurassic to Lower Cretaceous (Bathonian–Callovian to Albian) of the Mango-1 and  
123 Til-1 offshore wells, northern Sinai, Egypt. Based on 174 samples of ditch cuttings, the  
124 authors established 11 informal dinoflagellate cyst zones which were correlated with other  
125 successions in Egypt and surrounding Tethyan areas. Five of these zones cover the  
126 Bathonian–Callovian to ?Berriasian interval. A major depositional hiatus between the late  
127 Kimmeridgian and the ?Berriasian was identified, and was attributed to a major sea-level fall  
128 associated with the Cimmerian orogenic event (Aboul Ela and Tahoun 2010, figs 2, 3). The  
129 samples yielded diverse and rich marine and terrestrial palynofloras. This paper focuses  
130 entirely on biostratigraphy, and the ranges of all the palynomorphs were given in non-  
131 quantitative range charts (Aboul Ela and Tahoun 2010, p. 90–98). The Jurassic dinoflagellate  
132 cyst associations appear to be substantially similar in content and distribution to their  
133 European counterparts; for example *Cribroperidinium? longicorne*, *Ctenidodinium*  
134 *continuum*, *Gonyaulacysta jurassica*, *Korystocysta pachyderma* and *Systematophora areolata*  
135 were recorded.

136 Ied and Ibrahim (2010) studied the Jurassic and Early Cretaceous palynology of the  
137 Almaz-1 well in northern Egypt. This contribution focused mostly on miospores, but some  
138 dinoflagellate cysts were recorded from the Bajocian–Callovian to the Barremian. These  
139 include *Ctenidodinium sellwoodii*, *Gonyaulacysta jurassica*, *Pareodinia ceratophora* and  
140 *Systematophora penicillata* (see Ied and Ibrahim 2010, p. 10). A very similar  
141 biostratigraphical paper on the Middle Jurassic to Early Cretaceous (Callovian–Albian) of the  
142 Kabrit-1 well drilled west of Cairo in northeastern Egypt was published by Ied and Lashin  
143 (2016). They recorded dinoflagellate cysts from the entire succession examined, including  
144 *Cribroperidinium* spp., *Dichadogonyaulax? pannea*, *Gonyaulacysta jurassica*, *Lithodinia*  
145 *jurassica*, *Pareodinia prolongata* and *Systematophora areolata* (see Ied and Lashin 2016, fig.  
146 2). This assemblage is similar to floras from eastern North America and Europe.

147 Tahoun et al. (2012) undertook a study of the Middle Jurassic to Upper Cretaceous  
148 succession of the Alamein-IX well in northern Egypt. In this study, zone 5, which comprises  
149 the Masajid Formation, was interpreted as being of Callovian to possibly Kimmeridgian age  
150 (Tahoun et al. 2012, p. 68, fig. 3). This interpretation was based on the presence of  
151 *Acanthaulax* sp. cf. *A. crispa*, *Amphorulacysta? dodekova*, *Epiplosphaera reticulospinosa*,  
152 *Lithodinia jurassica*, *Meiourogonyaulax reticulata* and *Sentusidinium* spp. This assemblage  
153 appears to be somewhat biostratigraphically ambiguous; however, the presence of  
154 *Amphorulacysta? dodekova* and *Epiplosphaera reticulospinosa* strongly suggests a late  
155 Oxfordian to early Kimmeridgian age (Feist-Burkhardt and Wille 1992, fig. 2).

156 Gentzis et al. (2018) published a study on the petroleum prospectivity of the  
157 Matruh Basin, North Western Desert, Egypt. The dinoflagellate cysts *Ctenidodinium*  
158 *sellwoodii*, *Korystocysta gochtii*, *Mancodinium semitabulatum*, *Nannoceratopsis gracilis*  
159 *Rhynchodiniopsis cladophora* and *Systematophora penicillata* were recorded from the Wadi  
160 Natrun, Khattatba and Masajid formations. The two discrete intervals represented by these  
161 formations were interpreted as being of Toarcian–Aalenian and ?late Bathonian–Oxfordian  
162 (Gentzis et al. 2018, fig. 4). Some photographs were presented, although the images of  
163 *Nannoceratopsis gracilis* (Gentzis et al. 2018, pl. 1/1, 2) are not clearly of that species.

164 The stratigraphical palynology of the Middle and Upper Jurassic (Bathonian–  
165 Oxfordian) strata of the South Sallum well, North Western Desert, Egypt was studied by  
166 Mostafa et al. (2018). This interval yielded relatively diverse dinoflagellate cyst  
167 associations, and these were comprehensively illustrated (Mostafa et al. 2018, pls 2–5). Two  
168 dinoflagellate cyst biozones were recognised. These are the *Dichadogonyaulax sellwoodii* –  
169 *Wanaea acollaris* – *Wanaea digitata* Assemblage Zone, interpreted as Bathonian–Callovian

170 in age, and the *Amphorula dodekova* Interval Zone which was deemed to be Callovian–  
171 Oxfordian. Note that the species *Amphorula dodekova* is now questionably accommodated  
172 in *Amphorulacysta?* (see Williams and Fensome 2016, p. 139). Included in the  
173 *Dichadogonyaulax sellwoodii* – *Wanaea acollaris* – *Wanaea digitata* Assemblage Zone were  
174 *Ctenidodinium ornatum*, *Impletosphaeridium varispinosum*, *Korystocysta* spp.,  
175 *Mendicodinium groenlandicum*, *Pareodinia prolongata* and *Wanaea digitata*. By comparison  
176 with Europe, this interval is most likely to be entirely Callovian in age (e.g. Poulsen and  
177 Riding 2003). The index taxon for the *Amphorula dodekova* Interval Zone was first  
178 described from the Kimmeridgian and its range was determined as late Oxfordian to early  
179 Kimmeridgian (Zotto et al. 1987; Feist-Burkhardt and Wille 1992, fig. 2). Mostafa et al.  
180 (2018) interpreted this biozone as being of Callovian–Oxfordian age. However the presence  
181 of *Amphorulacysta? dodekova*, *Compositosphaeridium? polonicum*, *Endoscrinium*  
182 *galeritum*, *Gonyaulacysta jurassica* and *Neuffenia willei* is strongly suggestive that it is  
183 entirely of Oxfordian age (Riding 1984a, Riding and Thomas 1992). The biostratigraphical  
184 significance of selected Berriasian dinoflagellate cysts from northern Egypt was discussed in  
185 a review paper by Tahoun and Ied (2018). Sparse and low diversity dinoflagellate cyst  
186 associations were recorded from the Tithonian to Albian strata penetrated by the Minqar-IX  
187 well, northern Egypt by Mahmoud et al. (2019).

188           The paper by Jaydawi et al. (2016) is a major and well-illustrated contribution on  
189 the Callovian to Kimmeridgian dinoflagellate cyst biostratigraphy of the petroliferous  
190 Essaouira Basin in the Marrakesh–Safi region of central-western Morocco. These authors  
191 examined three boreholes. An early Callovian assemblage which includes *Ctenidodinium*  
192 *combazii*, *Ctenidodinium cornigerum* and *Impletosphaeridium varispinosum* was encountered  
193 in the MKL-110 borehole. Further material was studied from the NDK-2 and NDK-3  
194 boreholes. A rich late Callovian flora containing *Ctenidodinium continuum*, *Ctenidodinium*  
195 *ornatum* and *Wanaea thysanota* was recovered in the latter borehole. The NDK-2 well  
196 yielded established marker species such as *Cribroperidinium? longicorne*, *Egmontodinium*  
197 *polyplacophorum*, *Gonyaulacysta centriconnata*, *Scriniodinium crystallinum*,  
198 *Systematophora areolata* and *Trichodinium scarburghense*, indicative of the interval from the  
199 Callovian–Oxfordian transition to the Kimmeridgian.

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202 **2.2.       Sub-Arctic North America**

203 The only relevant single-region publication on sub-Arctic North America issued between  
204 February 2018 and January 2019 is that by Dodsworth and Eldrett (2018). These authors  
205 recorded the reworking of low numbers of *Chytroeisphaeridia chytroeides*, ?*Gonyaulacysta*  
206 *jurassica*, ?*Rhynchodiniopsis cladophora* and *Scriniodinium* spp. into the Upper Cretaceous  
207 (Cenomanian–Turonian) Bridge Creek Member (Greenhorn Formation) near Pueblo,  
208 Colorado, USA. These Middle to Late Jurassic forms are part of an extensive suite of  
209 allochthonous palynomorphs of Carboniferous to middle Cretaceous age (Dodsworth and  
210 Eldrett 2018, p. 9, 10).

211 Additionally, one older single-region contribution was also discovered. This is by  
212 van Helden (1987), and comprises a short article in a newsletter designed to encourage  
213 research on the Jurassic palynology of Alberta and Saskatchewan in western Canada. This  
214 author reported that the Nordegg, Poker Chip and Rock Creek formations of Alberta, and the  
215 Shaunavon and Vanguard formations in Saskatchewan contain abundant and diverse Jurassic  
216 dinoflagellate cyst assemblages. It was noted that detailed study of these floras would help  
217 the understanding of both the biostratigraphy and palaeoecology of the region. Van Helden  
218 (1987) expressed surprise that the palynology of these lithostratigraphical units had not been  
219 studied by contemporary palynologists in Canada, and recommended that this open field of  
220 research be advanced.

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### 223 **2.3. South America**

224 Only one publication is included here on South America. This is Olivera et al. (2018), which  
225 is a report of the pollen grain *Shanbeipollenites proxireticulatus* from the Vaca Muerta  
226 Formation of the Neuquén Basin, Argentina and its associated palynomorphs, including  
227 dinoflagellate cysts. *Shanbeipollenites proxireticulatus* was previously reported from the  
228 Upper Jurassic of Tanzania (Schrank 2004). The material in this study was interpreted as  
229 being of ?Berriasian–Valanginian age based on the overall palynoflora which includes  
230 *Meiourogonyaulax bulloidea*, *Sentusidinium villersense* and *Systematophora penicillata*  
231 (Olivera et al. 2018, figs 4, 5).

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### 234 **2.4. The Arctic**

235 Six recent single-region contributions from the Arctic are considered in this subsection. Four  
236 of these are from Arctic Russia, one of which (Nikitenko et al. 2018a) is considered to be  
237 especially significant (Appendix 1 of the Supplementary data).

238 The Lower Jurassic through Upper Cretaceous (Hettangian–Turonian)  
239 biostratigraphy and lithostratigraphy of the New Siberian Islands and adjacent areas of  
240 continental Arctic Siberia was studied by Nikitenko et al. (2017), who defined three  
241 depositional series with important reference sections. These strata have been substantially  
242 deformed. Despite the substantial structural complications, however, Nikitenko et al. (2017)  
243 demonstrated the applicability of these successions for correlation in the continental shelf east  
244 of the Laptev Sea and in the west of the East Siberian Sea. These authors used ammonites,  
245 bivalves, foraminifera, miospores and ostracods, in addition to dinoflagellate cysts. Nikitenko  
246 et al. (2018b) is a closely related study and involves an investigation of the same sections that  
247 were studied by Nikitenko et al. (2017). Nikitenko et al. (2018b) comprises a detailed  
248 examination of the micropalaeontology (dinoflagellate cysts, foraminifera, miospores and  
249 ostracods) and Hettangian and Pliensbachian organic geochemistry of the Hettangian to  
250 Turonian reference sections of the New Siberian Islands. A scheme of Boreal standard  
251 biozones was erected that have regional applicability in northern Siberia. Kashirtsev et al.  
252 (2018) involves a study based on organic geochemistry on the Oxfordian to Valanginian  
253 succession of the Nordvik Peninsula, western Anabar Bay, Arctic Russia. A comprehensive  
254 biostratigraphy has been developed for this succession including seven dinoflagellate cyst  
255 zones (Kashirtsev et al. 2018, fig. 2).

256 By far the most significant publication on the Arctic region in this review is that by  
257 Nikitenko et al. (2018a). This work details the biostratigraphy, geochemistry ( $\delta^{13}\text{C}_{\text{TOC}}$ ),  
258 palaeoecology and sedimentology of the Middle Jurassic to Lower Cretaceous (Bathonian–  
259 Valanginian) succession of the Olenek section in the Anabar-Lena region of Arctic Russia.  
260 The emphasis is on the uppermost Jurassic and Lower Cretaceous (Tithonian [=Volgian] to  
261 Valanginian) Buolkalakh and Iaedaes formations (Nikitenko et al. 2018a, fig. 3). Detailed  
262 range data was gathered for ammonites, dinoflagellate cysts, foraminifera and miospores  
263 (Nikitenko et al. 2018a, figs 6–9). Five ‘dinocyst local zones’ were recognised: 1 – the  
264 *Cometodinium whitei*, *Epiplosphaera gochtii*, *Gonyaulacysta eisenackii* ‘dinocyst local  
265 zone’; 2 – the *Bourkidinium* sp. ‘dinocyst local zone’; 3 – the *Gochteodinia villosa* ‘dinocyst  
266 local zone’; 4 – the *Batioladinium varigranosum*, *Occisucysta tentorium* ‘dinocyst local  
267 zone’; and 5 – the *Cyclonephelium cuculliforme*, *Batioladinium reticulatum* ‘dinocyst local  
268 zone’ (Nikitenko et al. 2018a, fig. 7). Numbers 1 to 4 of these ‘zones’ cover the ?uppermost



269 Kimmeridgian–Lower Volgian to uppermost Tithonian/Volgian–lower Boreal Berriasian  
270 interval. The ages of the five ‘dinocyst local zones’ from the Olenek section were calibrated  
271 to the current geological time scale via correlations with 11 coeval studies throughout the  
272 northern and southern hemispheres (Nikitenko et al. 2018a, fig. 11).

273 The only other single-region publications on the Arctic reviewed herein are those by  
274 Felix and Burbridge (1977) and Rismyhr et al. (2018). Felix and Burbridge (1977)  
275 established a new species of pteridophytic spore, *Ricciisporites umbonatus*, from the Upper  
276 Triassic (Carnian–Norian) of the Sverdrup Basin, Arctic Canada. This comprehensive study  
277 and it included details of the associated palynomorphs, including common occurrences of the  
278 dinoflagellate cysts *Sverdrupiella baccata*, *Sverdrupiella manicata*, *Sverdrupiella*  
279 *ornaticingulata*, *Sverdrupiella septentrionalis* and *Sverdrupiella usitata* in the borehole  
280 successions that were examined. Dinoflagellate cysts proved absent in the outcrop samples  
281 that were studied (Felix and Burbridge 1977, table 1). *Sverdrupiella usitata* is the most  
282 prominent species throughout the boreholes sections studied. The genus *Sverdrupiella* is  
283 therefore highly characteristic of the Norian of boreholes drilled in the Canadian Arctic  
284 (Bujak and Fisher 1976).

285 Rismyhr et al. (2018) provided a major study of the palynology, sedimentology and  
286 sequence stratigraphy of the Carnian to Callovian strata of western central Spitsbergen,  
287 Svalbard. Ten composite assemblage zones (CAZs) were established, of which the six for the  
288 Norian to Callovian interval are based on dinoflagellate cysts (Rismyhr et al. 2018, fig. 3).  
289 The principal focus of this study was the Knorringsfjellet Formation (Wilhelmøya Subgroup),  
290 in which three sequences were identified. Sequence 1 is Norian, and is characterised by the  
291 *Rhaetogonyaulax arctica* and *Heibergella* spp. CAZs. *Nannoceratopsis senex* gives its name  
292 to a Toarcian CAZ which is equivalent to Sequence 2. The Brentskardhaugen Bed is a highly  
293 condensed deposit of late Toarcian–early Aalenian age. It is assigned to the *Phallocysta*  
294 *eumekes* CAZ, and was assigned to Sequence 2 by Rismyhr et al. (2018).

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## 297 **2.5. Australasia**

298 Seven single-region contributions from Australasia are listed in Appendix 1 of the  
299 Supplementary data in this review. Zhang Wangping & Grant-Mackie (2001) described the  
300 palynology of the Late Triassic and Early Jurassic (Norian–Sinemurian) of New Zealand.  
301 This paper is chiefly on miospores, although the authors also reported undifferentiated

302 acritarchs and dinoflagellate cysts from various lithostratigraphical units of the Hokonui Hills  
303 and southwestern Kawhia.

304         The remaining six single-region studies are from Australia. Jones & Nicoll (1984) and  
305 Dixon et al. (2012) are short papers on the Late Triassic (Carnian–Norian) of the North West  
306 Shelf of Australia. Both mention dinoflagellate cysts briefly. Dixon et al. (2012) worked on  
307 the Upper Mungaroo Formation from the offshore Carnarvon Basin. These authors recorded  
308 *Dapcodinium* spp. from marginal marine to tidally influenced facies, and *Hebecysta balmei*  
309 and *Rhaetogonyaulax* spp. from open-marine settings. The paper by Paumard et al. (2018) is  
310 also on material from the North West Shelf of Australia. In this multidisciplinary study the  
311 authors examined the sedimentary architecture and sediment partitioning in the Barrow  
312 Group (Tithonian–Valanginian) of the Northern Carnarvon Basin. Offshore well data were  
313 integrated to establish a seismic stratigraphy of this economically important unit. The authors  
314 related seven third-order seismic sequences to the eustatic and tectonic history of the  
315 depocentre, and calibrated these sequences using the *Pseudoceratium iehiense* to  
316 *Systematophora areolata* dinoflagellate cyst zones of Helby et al. (1987).

317         However, the most significant Australasian contribution listed herein is by Wainman  
318 et al. (2018a). This paper is on the latest Jurassic (Tithonian) palynology of the Indy 3 well  
319 in the western Surat Basin, Queensland, southeastern Australia. These authors discovered  
320 low-diversity dinoflagellate cysts and colonial algae in the Walloon Coal Measures (Injune  
321 Creek Group). The Walloon Coal Measures were previously believed to be entirely  
322 nonmarine. The study showed that either a brief marine transgression is represented by this  
323 unit, or that these planktonic palynomorphs were freshwater forms and thus represent a rare  
324 report of pre-Cretaceous nonmarine dinoflagellate cysts. These records are coincident with an  
325 interval of high global sea level hence the former scenario appears to be the best explanation.  
326 The new dinoflagellate cyst species described by Wainman et al. (2018a) are *Moorodinium*  
327 *crispa* and *Skuadinium fusum*, and they also described the colonial alga *Palambages pariunta*  
328 as new. *Moorodinium crispa* and *Skuadinium fusum* are small, thin-walled proximate cysts  
329 from a thin (ca. 2 m) unit interpreted as a possible upper estuarine deposit (Wainman et al.  
330 2018a, pls 1, 2). This interpretation, together with the high dominance and low species  
331 richness nature of the assemblage, is consistent with a freshwater setting. Wainman et al.  
332 (2018a) provided the palynological basis for a wider study of the Middle–Upper Jurassic  
333 Walloon Coal Measures of the Surat Basin (Wainman and McCabe 2018, Wainman et al.  
334 2018b).

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337 **2.6. East Europe**

338 None of the nations generally considered to comprise this region are in the Arctic Circle, so  
339 the prefix ‘sub-Arctic’ is superfluous in this case. Nine single-region items concerning  
340 Bulgaria, Hungary, Poland and Slovakia in East Europe are listed in Appendix 1 of the  
341 Supplementary data. Two of these publications are deemed to have substantial significance.

342 Three multi-authored papers written in Bulgarian detail the Jurassic (Sinemurian–  
343 Tithonian) lithostratigraphy of northeastern Bulgaria (Sapunov et al. 1985, 1986a, 1986b).  
344 Specifically, the Dobrič, Drinovo, Esenica, Ginci, Javorec, Kalojan, Ozirovo, Polaten,  
345 Provadija, Sultanci and Tiča formations were considered in this set of contributions, all in the  
346 same journal. The material studied is from numerous deep, continuously cored, boreholes  
347 drilled as part of a partially successful petroleum exploration campaign throughout northern  
348 Bulgaria. In Sapunov et al. (1985), the four formations considered are treated separately in  
349 ascending stratigraphical order. By contrast in Sapunov et al. (1986a, 1986b), the  
350 lithostratigraphy was described borehole-by-borehole. Throughout each of these three papers,  
351 integrated biostratigraphy based on brachiopods, calpionellids, molluscs and palynomorphs  
352 was included. Selected occurrences of dinoflagellate cysts, especially in Sapunov et al.  
353 (1986a), were provided by the late Lilia Dodekova, who was one of the co-authors. No  
354 photographs or range charts were provided.

355 The paper by Bóna (1995) is a major work on the Upper Triassic  
356 palynostratigraphy of a large coal-bearing basin around Pécs, in the Mecsek Mountains of  
357 southern Hungary. Only ‘dinoflagellate indet.’ was recovered from the lowermost Mecsek  
358 Coal formations (Rhaetian) (Bóna 1995, table 2; pl. 8/17). This specimen is very poorly-  
359 preserved, but appears to be referable to *Rhaetogonyaulax rhaetica*. This author also reported  
360 questionable specimens of *Hystrichosphaeridium magnum* from the Karolinavölgy Sandstone  
361 and the lowermost Mecsek Coal formations, which are of Norian and Rhaetian age  
362 respectively (Bóna 1995, table 2; pl. 8/16; pl. 9/2, 3, 6). The original authorship of the  
363 spinose species *Hystrichosphaeridium magnum* was not provided by Bóna (1995) and was  
364 not located by the present author. Only one of the two specimens illustrated of  
365 *Hystrichosphaeridium magnum* appears to have possible dinoflagellate cyst affinity (Bóna  
366 1995, pl. 8/16). This specimen is a chorate form with an apparent apical archaeopyle and may  
367 be referable to *Beaumontella*.

368 As part of an unpublished, PhD thesis mainly focussed on miospores from  
369 southwestern England, Hungary and the southwestern USA, Baranyi (2018) analysed

370 borehole material from the the Veszprém Marl Formation (Carnian) of the southern  
371 Transdanubian Range, western Hungary and recorded ‘dinocyst indet.’ and *Heibergella* sp.  
372 (see Baranyi 2018, pl. 12/10, 11). These records are within the Carnian Pluvial Episode, and  
373 are highly unusual in that most Triassic dinoflagellate cyst occurrences are Norian–Rhaetian  
374 (Mangerud et al. 2019, fig. 2).

375           The dissertation by Ruckwied (2009) on the palynology of the Rhaetian and  
376 Hettangian strata of the northwestern Tethyan Realm of Hungary and northern Slovakia  
377 aimed to investigate biostratigraphy, palaeoclimate and palaeoecology. The principal thrust of  
378 this contribution is on miospores. However Ruckwied (2009) reported *Dapcodinium priscum*  
379 and *Rhaetogonyaulax rhaetica* from the Rhaetian and Hettangian of the Furkaska section in  
380 the Tatra Mountains of northern Slovakia (see also Ruckwied and Götz 2009). Ruckwied  
381 (2009) did not record dinoflagellate cysts from the successions investigated in Hungary.

382           The prominent Hungarian palynologist Mária Sütőné Szentai produced a  
383 compilation of all the genera and species of Silurian to Holocene organic-walled  
384 microplankton reported from Hungary since 1957 (Sütőné Szentai 2018). This compendium  
385 was an alphabetical listing of all published post-Silurian records of palynomorphs excluding  
386 miospores. By far the largest section of this book is on dinoflagellate cysts (Sütőné Szentai  
387 2018, p. 11–111). Every species recorded from Hungary is included; the holotype and the  
388 stratigraphical range are also given. The main papers on Triassic and Early Jurassic  
389 dinoflagellate cysts from Hungary included in this compilation are Bóna (1995), Bucefalo  
390 Palliani et al. (1997) and Baranyi et al. (2016).

391

392

## 393 **2.7.       Sub-Arctic West Europe**

394 A total of 29 single-region contributions on the Triassic, Jurassic and earliest Cretaceous of  
395 sub-Arctic West Europe are covered in this review, and 12 of these are considered to be  
396 considerably impactful (Appendix 1 of the Supplementary data). This section is subdivided  
397 into three subsections based on the stratigraphical coverage of the items.

398

### 399 **2.7.1.     Triassic and Early Jurassic**

400 Seven articles summarised here are focused on the Triassic and Early Jurassic interval. The  
401 paper by Karle (1984) involves a detailed study of the palynology of the Triassic–Jurassic  
402 boundary at Fonsjoch, western Austria. This author recorded *Rhaetogonyaulax rhaetica* from  
403 the Rhaetian. This species is especially common in the middle and upper Rhaetian, and the

404 range top is immediately below a prominent limestone bed which underlies the Pre-Planorbis  
405 Beds at the Triassic–Jurassic transition (Karle 1984, fig. 3).

406 The published PhD dissertation by Holstein (2004) details the palynofacies,  
407 palynology and sequence stratigraphy of the Kössen Beds (Upper Triassic, Norian–Rhaetian)  
408 of the Eiberg and Mörtlbachgraben sections in the Northern Calcareous Alps of northern  
409 Austria. The palynofloras are dominated by miospores, but the dinoflagellate cysts  
410 *Dapcodinium priscum* and *Rhaetogonyaulax rhaetica* were recognised. This author asserted  
411 that *Dapcodinium priscum* preferred high-energy, shallow-water settings, and  
412 *Rhaetogonyaulax rhaetica* had a preference for deep-water, low-energy palaeoenvironments.

413 A wide-ranging and multidisciplinary study on the Triassic–Jurassic boundary by  
414 Lindström et al. (2017) includes descriptions of palynofloras from several localities in  
415 Austria, Denmark, England and Germany. The range top of the last, last common and last  
416 consistent occurrences (LO, LCO and LCON respectively) of *Rhaetogonyaulax rhaetica*  
417 were used as a reliable regional markers. The LCO and LCON of this prominent and  
418 cosmopolitan species are consistently within, or immediately below and above, the extinction  
419 phase in the late Rhaetian. However, *Rhaetogonyaulax rhaetica* apparently became extinct  
420 (LO) during the post-extinction phase in Austria and England (Lindström et al. 2017, fig. 12).

421 Juncal et al. (2018) involves a multidisciplinary study of the Permian and Triassic  
422 of the Paris Basin in central France. These authors reported *Dapcodinium priscum* and  
423 *Rhaetogonyaulax rhaetica* from the uppermost Rhaetian of the Sancerre-Couy 1 borehole,  
424 within their SC-4 assemblage. Schneebeili-Hermann et al. (2018) provided a very detailed  
425 study of the palynology of the Norian, Rhaetian and Sinemurian strata of northern  
426 Switzerland, distinguishing five informal palynomorph associations. The main emphasis was  
427 on miospores, but nine dinoflagellate cyst taxa were recognised from the Rhaetian and  
428 Sinemurian (Schneebeili-Hermann et al. 2018, figs 2, 5). These include *Beaumontella langii*,  
429 *Dapcodinium priscum*, *Rhaetogonyaulax rhaetica* and ?*Suessia swabiana*; the greatest  
430 dinoflagellate cyst diversity is in the Rhaetian (Schneebeili-Hermann et al. 2018, figs 2, 5).

431 As part of a major multidisciplinary paper on the cores recovered by the  
432 Schandelah Scientific Drilling Project in northern Germany, van de Schootbrugge et al.  
433 (2018) investigated the palynology of an important Rhaetian to Toarcian succession. Full  
434 details of the palynomorphs recovered were not given, but these authors illustrated major  
435 bioevents and figured significant dinoflagellate cyst, miospore taxa (van de Schootbrugge et  
436 al. 2018, fig. 3, pl. 1). The authors recognised the Early Jurassic *Dapcodinium priscum*,  
437 *Liasidium variable* and *Nannoceratopsis* dinoflagellate cyst zones.

438                Several contributions on the Early Jurassic palynology of the Lusitanian Basin in  
439 western Portugal were published recently by Vânia Correia and her co-authors (Appendix 1  
440 of the Supplementary data). These are all associated with the author's PhD thesis (Correia  
441 2018), and the most significant is Correia et al. (2018). This paper is on the  
442 palynostratigraphy of the Lower Jurassic strata of this important Iberian depocentre. Correia  
443 et al. (2018) documented the Sinemurian to Toarcian palynomorph biostratigraphy based on  
444 six localities, with the principal emphasis being on dinoflagellate cysts. The Sinemurian  
445 proved devoid of dinoflagellate cysts. By contrast the Pliensbachian and Toarcian are  
446 characterised by the presence of the genera *Luehndea*, *Mancodinium*, *Mendicodinium*,  
447 *Nannoceratopsis* and *Scriniocassis*. *Luehndea* was apparently made extinct by the Toarcian  
448 Oceanic Anoxic Event (T-OAE). This event proved substantially more intense in the  
449 Lusitanian Basin than elsewhere in southern Europe, and the recovery of phytoplankton was  
450 protracted in this basin. Correia et al. (2018) proposed a biozonation for the late  
451 Pliensbachian and Toarcian, comprising the *Luehndea spinosa* and *Mendicodinium*  
452 *microscabratum* dinoflagellate cyst zones. The zones were subdivided into subzones (Correia  
453 et al. 2018, fig. 15).

454

#### 455 **2.7.2. Middle Jurassic**

456 In this subsection, four items focused on the Middle Jurassic are documented. A major study  
457 on the palynology of the Middle Jurassic Ravenscar Group, from the Cleveland Basin,  
458 northeastern Yorkshire, northern England, was undertaken by Hogg (1993). This unpublished  
459 PhD thesis focussed on outcrops of the Cloughton, Scarborough, Scalby and Cornbrash  
460 formations (Bajocian–Bathonian) in the Scarborough area of North Yorkshire. The emphasis  
461 was on miospores, but diverse dinoflagellate cyst floras were also recovered and 30 genera  
462 were recognised (Hogg 1993, p. 121–137, fig. 4.6, pls 17–23). Three new species were  
463 informally introduced. Furthermore, *Ambonosphaera calloviana* and *Tabulodinium senarium*  
464 were reported from the UK for the first time. Hogg (1993) determined that much of the Long  
465 Nab Member of the Scalby Formation is of latest Bathonian (*Clydoniceras discus* ammonite  
466 zone) age. This work refined the biostratigraphical results of Riding and Wright (1989), who  
467 reported a Bathonian (undifferentiated) age. Hogg (1993, p. 179–190) discussed the  
468 dinoflagellate cyst biostratigraphy of the Scarborough and Scalby formations in some detail,  
469 comparing his results with previous studies such as those by Riding and Wright (1989) and  
470 Gowland and Riding (1991). The sequence stratigraphy of the successions investigated was  
471 also analysed.

472 Powell et al. (2018, appendix B) documented the palynology of two samples from  
473 the Kellaways Sand Member (Lower Callovian) of Burythorpe Sand Quarry, North  
474 Yorkshire, UK. Sample 2 was relatively rich in dinoflagellate cysts and five specimens were  
475 illustrated (Powell et al. 2018, fig. 12).

476 The early Mesozoic phytoplankton radiation was investigated by Wiggan et al.  
477 (2018). The coccolithophores and dinoflagellates radiated substantially during the Bajocian  
478 (~170–168 Ma). Wiggan et al. (2018) described and interpreted a dominance of the genus  
479 *Dissiliodinium* in the mid-latitudes, followed by the explosive evolutionary expansion of the  
480 dinoflagellate family Gonyaulacaceae. The latter phenomenon was viewed as being strongly  
481 influenced by increases in sea level and changes in ocean gateways, and possibly related to  
482 the Mesozoic Marine Revolution. The key dinoflagellate cyst data in Wiggan et al. (2018)  
483 were from an important borehole succession in southern Germany initially published by  
484 Wiggan et al. (2017).

485 Correia et al. (2019) presents part of the senior author's PhD study (Correia 2018),  
486 providing an account of the palynostratigraphy of the Cabo Mondego and Póvoa de Lomba  
487 formations (uppermost Toarcian–Bathonian) at Cabo Mondego and São Gião in the northern  
488 Lusitanian Basin. The succession at Cabo Mondego includes the Global Stratotype Section  
489 and Point (GSSP) for the Bajocian. The samples from Cabo Mondego were by far the most  
490 palynologically productive. Here the uppermost Toarcian to lowermost Bajocian succession  
491 produced low diversity dinoflagellate cyst associations dominated by *Nannoceratopsis*.  
492 Within the *Witchellia laeviuscula* ammonite zone, the assemblages become markedly more  
493 diverse, reflecting the intra-Bajocian global evolutionary explosion of dinoflagellates. This  
494 predominantly involved the family Gonyaulacaceae, and was apparently strongly linked to  
495 sea-level rise (Wiggan et al. 2017, 2018). The upper part of the Lower Bajocian and much of  
496 the Upper Bajocian were not sampled by Correia et al. (2019, fig. 2); however the trend of  
497 increasing dinoflagellate cyst diversity continued through the Bajocian–Bathonian transition.  
498 It is clear from Correia et al. (2019) that the Middle Jurassic dinoflagellate cyst species  
499 richnesses in the Arctic region and the Boreal Realm are substantially higher than in southern  
500 Europe. This may be because more northerly palaeolatitudes were a phytoplankton-diversity  
501 hotspot during the Mesozoic, that the the recovery from the Toarcian Oceanic Anoxic Event  
502 (T-OAE) was more protracted in the Iberian region, or that regional palaeogeographical  
503 factors controlled dinoflagellate diversity in the Lusitanian Basin.

504

505 **2.7.3. Middle Jurassic to Early Cretaceous inclusive**

506 In this subsection, the six remaining substantial papers exclusively on West Europe are  
507 discussed. The works involved range from the Middle Jurassic to Early Cretaceous. Heunisch  
508 and Luppold (2018) present an important study of the Middle Jurassic to earliest Cretaceous  
509 (Callovian–Berriasian) micropalaeontological biostratigraphy of two boreholes drilled in the  
510 Lower Saxony Basin of northern Germany. It is a technical report with the  
511 micropalaeontology of selected intervals in the Eulenflucht-1 and Wendhausen-6 boreholes  
512 described one-by-one. In the palynology subsections, age-diagnostic dinoflagellate cysts such  
513 as *Compositosphaeridium? polonicum*, *Dingodinium tuberosum*, *Gonyaulacysta jurassica*,  
514 *Hystrichosphaerina? orbifera*, *Muderongia simplex* subsp. *microperforata*, *Nannoceratopsis*  
515 *pellucida*, *Pareodinia brevicornuta*, *Systematophora areolata* and *Systematophora*  
516 *penicillata* are mentioned (Heunisch and Luppold 2018, pl. 3).

517 An overview of the Middle Jurassic to Early Cretaceous (Bathonian–Barremian)  
518 basin evolution in the Central Graben area of the North Sea (representing the Danish, Dutch  
519 and German sectors) was presented by Verreussel et al. (2018). This study was entirely based  
520 on the correlations of wells that have been analysed for palynology (Verreussel et al. 2018,  
521 fig. 2). These authors recognised four intervals that they termed tectonostratigraphical  
522 megasequences (TMS). Each TMS represents a distinct phase of basin evolution; for example  
523 TMS-1 reflects the onset of basin rifting and the rift climax occurred during this phase. It was  
524 characterised by thick mud deposition.

525 A study focussed on the Oxfordian/Kimmeridgian boundary beds in the Flodigarry  
526 Shale Member (Staffin Bay Formation) of the Isle of Skye, northwestern Scotland, was  
527 undertaken by Barski (2018). This unit is notable because it includes a proposed Global  
528 Stratotype Section and Point (GSSP) for the base of the Kimmeridgian. Barski (2018) studied  
529 seven samples from the *Ringsteadia pseudocordata* and *Pictonia baylei* ammonite zones, and  
530 presented quantitative data (Barski 2018, table 1). The author recognised a eutrophication  
531 event in the lowermost Kimmeridgian. Furthermore, Barski (2018) noted that the range bases  
532 of sparse *Emmetrocyta sarjeantii*, *Perisseiasphaeridium pannosum* and *Senoniasphaera*  
533 *jurassica* can be used as markers for the base of the Kimmeridgian.

534 Turner et al. (2018) is a major study by on the comprehensive stratigraphy of the  
535 Upper Jurassic and Lower Cretaceous of the Arctic and Europe, and it included palynological  
536 analysis of the Kimmeridgian to Berriasian of the Norwegian Continental Shelf. These  
537 authors integrated carbon isotope, cyclostratigraphical, dinoflagellate cyst and gamma ray  
538 data to effect interregional correlations throughout this important interval. Turner et al. (2018,  
539 fig. 2) calibrated the palynological data from the Barents and North seas to the the current



540 geological time scale using ammonite-dated dinoflagellate cyst studies such as Riding and  
541 Thomas (1992) and Poulsen and Riding (2003).

542 Schneider et al. (2018) undertook a detailed study of the micropalaeontology and  
543 palynology of the Jurassic–Cretaceous transition (Tithonian–Berriasian) in the Lower Saxony  
544 Basin of northern Germany. This study is based mainly on miospores, but the authors noted  
545 the regional correlative significance of the range bases of *Batioladinium pomum*,  
546 *Cantulodinium speciosum*, *Muderongia simplex* subsp. *microperforata*, *Muderongia simplex*  
547 and *Pseudoceratium pelliferum* in the Berriasian (Schneider et al. 2018, fig. 2).

548 Stanley Duxbury continued his extensive research into dinoflagellate cysts from  
549 the Lower Cretaceous of the North Sea and surrounding areas that began with Duxbury  
550 (1977). Duxbury (2018) is a major study of the Berriasian to lower Hauterivian marine  
551 palynostratigraphy of the Speeton Clay and Valhall formations of the Central North Sea and  
552 northeastern England based on 1131 samples. The biozonation of Duxbury (2001) is  
553 substantially refined. Duxbury (2018) is dominated by a systematic section with taxonomic  
554 novelties including one new genus and 21 new species.

555

556

## 557 2.8. *The Middle East*

558 The one single-region contribution on the Middle East herein is the article by Eshet (1990).  
559 This author studied the Permian and Triassic successions of 11 boreholes drilled throughout  
560 Israel. Eshet (1990) erected seven interval zones for the Early Permian to Late Triassic. Only  
561 one dinoflagellate cyst zone, the *Rhaetogonyaulax rhaetica* Zone (VII; Norian–Rhaetian),  
562 was established. The base of this zone was defined by the range bases of post-Carnian  
563 miospores, and the top by the range top of *Rhaetogonyaulax rhaetica* and other dinoflagellate  
564 cysts (i.e. *Heibergella asymmetrica*, *Noricysta fimbriata*, *Suessia swabiana* and *Sverdrupiella*  
565 spp.). Two distinct palynofacies are present in this zone, one with dinoflagellate cysts and the  
566 other devoid of these marine palynomorphs. The *Rhaetogonyaulax rhaetica* Zone is restricted  
567 to the coastal plain in the extreme west of Israel; elsewhere the Norian–Rhaetian has been cut  
568 out by a regional unconformity. Its reference section is within the Shefaiym Formation,  
569 between 4860 m and 4495 m in the Ga’ash 2 Borehole, northwestern Israel (Cousminer 1981;  
570 Eshet 1990, fig. 1). The age interpretation of Norian–Rhaetian is based on the ranges of the  
571 dinoflagellate cysts *Rhaetogonyaulax rhaetica* and *Suessia swabiana* (see Visscher &  
572 Brugman 1981) and foraminifera.

573

574

## 575 **2.9. Sub-Arctic Russia**

576 Twelve single-region articles on sub-Arctic Russia are compiled in this review, six of which  
577 are deemed to be highly significant (Appendix 1 of the Supplementary data). Three of these  
578 are on sub-Arctic West Russia, i.e. west of the Ural Mountains, and the remaining nine are on  
579 southwestern Russia. The most significant of these contributions are detailed below in two  
580 subsections.

581

### 582 **2.9.1. Sub-Arctic West Russia**

583 The unpublished PhD thesis by Smith (1999), available online, details the palynostratigraphy  
584 of the Tithonian (Volgian) to Valanginian strata of the important Volgian lectostratotype  
585 successions at Gorodische and Kashpir, near Ulyanovsk, southwestern Russia. The key  
586 findings were later incorporated into Harding et al. (2011). Another noteworthy article based  
587 on the Upper Kimmeridgian and Tithonian strata of Gorodische is Pestchevitskaya (2018).  
588 This author focussed on the distinctive camocavate dinoflagellate cyst genus *Dingodinium*.  
589 The genus was emended, with the archaeopyle interpreted as of combination type  
590 (apical/anterior intercalary), and the tabulation partiform (Pestchevitskaya 2018, fig. 2).  
591 Pestchevitskaya (2018) thus placed *Dingodinium* into the family Cladopyxiaceae. She  
592 compiled the Middle Jurassic to the latest Cretaceous stratigraphical ranges of the genus  
593 worldwide, and discussed the morphologies of 12 species of *Dingodinium* (Pestchevitskaya  
594 2018, figs 1, 3). Pestchevitskaya (2018) identified *Dingodinium albertii*, *Dingodinium*  
595 *jurassicum* and *Dingodinium tuberosum* from Gorodische, and established the new species  
596 *Dingodinium nequeas*. In another publication Dzyuba et al. (2018) discussed the ranges of  
597 dinoflagellate cysts, mainly identified only at the generic level, from the Tithonian–  
598 Berriasian (Volgian–Ryazanian) of a fossiliferous succession exposed on the banks of the  
599 Maurynya River in the Northern Ural Mountains of West Siberia.

600

### 601 **2.9.2. Southwestern Russia (the Caspian Sea, the Caucasus Mountains and Crimea)**

602 The discussion of articles on the Caspian Sea, the Caucasus Mountains and Crimea is placed  
603 in the subsection on sub-Arctic Russia herein for purely geographical and pragmatic reasons.  
604 This strategy has absolutely no political significance whatsoever.

605 The remaining nine items from sub-Arctic Russia, all recently published, are based  
606 on material from southwestern Russia. Arkadiev et al. (2018) is a biostratigraphical and  
607 magnetostratigraphical synthesis of the Jurassic–Cretaceous boundary beds of the Crimean

608 Mountains. Tithonian and Berriasian strata are well-developed and highly fossiliferous  
609 throughout this region. The authors summarised research on ammonites, calpionellids,  
610 dinoflagellate cysts, foraminifera and ostracods from Tithonian to lowermost Valanginian  
611 strata across Crimea, and correlated these fossil records with magnetostratigraphy. Two  
612 subdivisions based on dinoflagellate cysts were recognised. These are the ‘beds with  
613 *Amphorula expirata*’ (now *Amphorulacysta? expirata*) and the ‘beds with *Phoberocysta*  
614 *neocomica*’ of latest Tithonian–earliest Berriasian and earliest Berriasian–latest Berriasian  
615 age respectively. The range bases of *Ctenidodinium elegantulum*, *Phoberocysta neocomica*  
616 and *Spiniferites* spp. define the boundary between these two informal biozones; these  
617 bioevents are within the *Tirnovella occitanica* ammonite zone (Arkadiev et al. 2018, fig. 21).

618 The short paper by Goryacheva and Ruban (2018) is on the Pliensbachian and  
619 Toarcian palynology of the Sjuk River valley in the northwestern Caucasus, where the  
620 authors identified *Nannoceratopsis senex*. Another short article by Goryacheva et al. (2018)  
621 is on the palynology of a sandstone representing a reportedly deep marine setting in the upper  
622 part of the Bagovskaja Formation from the River Belaja, south of Guzeripl in the northern  
623 Arkhyz-Guzereplskaja area, Western Caucasus region. This unit is Toarcian in age, based on  
624 ammonites recovered from its lowermost beds. Goryacheva et al. (2018) reported a  
625 dinoflagellate cyst assemblage dominated by *Nannoceratopsis*. The most abundant species is  
626 *Nannoceratopsis spiculata*. Less common forms include *Nannoceratopsis plegas*,  
627 *Nannoceratopsis senex*, *Phallocysta eumekes* and *Susadinium faustum*. Goryacheva et al.  
628 (2018) interpreted this association as being of late Toarcian age.

629 Mitta et al. (2017) provided an integrated palaeontological study of the Middle  
630 Jurassic (Bajocian–Bathonian) of the Bolshoi Zelenchuk River Basin in the Northern  
631 Caucasus region. Samples were collected from the uppermost Bajocian and lowermost  
632 Bathonian (*Parkinsonia parkinsoni* and *Zigzagiceras zigzag* ammonite zones) part of the  
633 Djangura Formation from localities 8, 11, 12 and 25 of Mitta et al. (2017, fig. 2). The authors  
634 examined ammonites, dinoflagellate cysts, foraminifera, miospores and ostracods from this  
635 succession. The five productive samples yielded moderately diverse dinoflagellate cyst  
636 associations. Prominent species recorded throughout include *Aldorfia aldorfensis*,  
637 *Chytroeisphaeridia chytroeides*, *Ctenidodinium sellwoodii*, *Korystocysta gochtii*,  
638 *Meiourogonyaulax caytonensis*, *Nannoceratopsis gracilis*, *Nannoceratopsis spiculata* and  
639 *Valensiella ovulum*. Mitta et al. (2017) recognised two informal zones. The uppermost  
640 Bajocian was termed ‘beds with *Rynchodiniopsis? regalis*’, the nominate species being  
641 confined to this interval. The most prominent dinoflagellate cysts are *Ctenidodinium*

642 *sellwoodii* and *Dissiliodinium* spp. The presence of *Acanthaulax* aff. *crispa* (as  
643 *Cribroperidinium* aff. *crispum*) is indicative of the late Bajocian (Wiggan et al. 2017).  
644 However, Mitta et al. (2017) also recorded *Nannoceratopsis dictyambonis* from this interval.  
645 This species, characteristic of the latest Toarcian to early Bajocian interval (Riding 1984a,  
646 1984b; Wiggan et al. 2017), may thus be reworked. The overlying lowermost Bathonian was  
647 assigned to ‘beds with *Ctenidodinium sellwoodii*’, named after one of the dominant species.  
648 The dinoflagellate cysts of this interval are substantially similar to those from the underlying  
649 interval. Only six range bases were observed in the ‘beds with *Ctenidodinium sellwoodii*’,  
650 including that of *Ctenidodinium continuum*.

651 Mitta et al. (2018) is a companion paper to that of Mitta et al. (2017). The former is  
652 an important biostratigraphical study of the Upper Bajocian Djangura Formation from the  
653 banks of the Kyafar River, a tributary of the Bolshoi Zelenchuk River, Karachay-Cherkessia,  
654 Northern Caucasus, southwestern Russia. The material is all from the *Rarecostites subarietis*  
655 ammonite subzone of the *Parkinsonia parkinsoni* ammonite zone. Ten samples were  
656 examined, and all yielded substantial proportions of dinoflagellate cysts in relatively diverse  
657 associations (Mitta et al. 2018, fig. 6). The samples are dominated by *Dissiliodinium* spp.  
658 Furthermore, the following species were found throughout: *Aldorfia aldorfensis*;  
659 *Chytroeisphaeridia chytroeides*; *Ctenidodinium continuum*; *Ctenidodinium sellwoodii*;  
660 *Durotrigia daveyi*; *Korystocysta* spp.; *Meiourogonyaualax caytonensis*; *Meiourogonyaualax*  
661 *valensii*; *Nannoceratopsis gracilis*; *Nannoceratopsis senex*; *Nannoceratopsis spiculata*;  
662 *Pareodinia ceratophora*; *Pareodinia prolongata*; *Rhynchodiniopsis? regalis*; *Sentusidinium*  
663 spp.; *Tubotuberella* spp.; and *Valensiella ovulum*. The following are also present, but rather  
664 less consistently: *Endoscrinium galeritum*; *Kalyptea stegasta*; *Leptodinium* sp.;  
665 *Nannoceratopsis dictyambonis*; *Nannoceratopsis raunsgaardii*; *Pareodinia halosa*;  
666 *Phallocysta elongata*; and *Wanaea acollaris* (see Mitta et al. 2018, fig. 6, pl. V). The authors  
667 assigned the entire succession that they studied to the ‘beds with *Meiourogonyaualax valensii*,  
668 *Rhynchodiniopsis? regalis*’ (Mitta et al. 2018, fig. 7). Many of the dinoflagellate cyst taxa  
669 recovered by Mitta et al. (2018) are entirely consistent with the latest Bajocian age  
670 determined by ammonites and other fossils. These marker taxa include *Aldorfia aldorfensis*,  
671 *Ctenidodinium continuum*, *Ctenidodinium sellwoodii*, *Dissiliodinium* spp., *Durotrigia daveyi*,  
672 *Kalyptea stegasta*, *Korystocysta* spp., *Leptodinium* sp., *Meiourogonyaualax caytonensis*,  
673 *Meiourogonyaualax valensii*, *Rhynchodiniopsis? regalis*, *Tubotuberella* spp. and *Wanaea*  
674 *acollaris* (see for example Riding and Thomas 1992, Wiggan et al 2017).

675 As in Mitta et al. (2017), Mitta et al. (2018), identified apparent reworking of  
676 dinoflagellate cysts. The diversity of the species involved is considerable, comprising  
677 *Nannoceratopsis dictyambonis*, *Nannoceratopsis raunsgaardii*, *Nannoceratopsis gracilis*,  
678 *Nannoceratopsis senex* and *Phallocysta elongata*. The presence of these taxa is clearly  
679 stratigraphically incompatible with a latest Bajocian age and reflects stratigraphical recycling  
680 of Upper Pliensbachian to Early Bajocian strata. The range bases of *Nannoceratopsis gracilis*  
681 and *Nannoceratopsis senex* are late Pliensbachian, and these species are common throughout  
682 the Toarcian and earliest Bajocian interval (e.g. Morgenroth 1970, Poulsen 1996, Riding et al.  
683 1999, Correia et al. 2018, Correia et al. 2019). The allochthonous species with the shortest  
684 ranges are *Nannoceratopsis dictyambonis* and *Phallocysta elongata*, forms that are  
685 characteristic of the latest Toarcian to earliest Bajocian (Riding 1984b, 1994).

686 The earliest Cretaceous (Berriasian) dinoflagellate cysts from the Uruh section in  
687 the North Caucasus, southwestern Russia were reported in a short contribution in Russian by  
688 Shurekova (2018). The article is well-illustrated and a semiquantitative range chart was  
689 presented (Shurekova, 2018, p. 283–285). The author distinguished the *Phoberocysta*  
690 *neocomica* and *Systematophora* cf. *palmula* dinoflagellate cyst zones which are broadly  
691 equivalent to the *Tirnovella occitanica* and *Fauriella boissieri* ammonite zones. The species  
692 *Systematophora palmula* is now known as *Palaecysta palmula*.

693

### 694 3. Conclusions

695 In a literature search from February 2018 to January 2019, 55 new publications pertaining to  
696 Triassic to earliest Cretaceous dinoflagellate cysts were discovered, and are compiled herein  
697 together with 38 older items which were not covered by Riding (2012, 2013, 2014, 2019).  
698 These 93 papers are based on material from Africa, North America, South America, the  
699 Arctic, Australasia, East Europe, West Europe, the Middle East and Russia. Thirty of them  
700 are deemed herein to be significantly impactful and are asterisked in Appendix 1 of the  
701 Supplementary data. All 93 contributions are listed in Appendix 1 of the Supplementary data,  
702 and most are on the Late Triassic and Early Jurassic of Europe (Tables 1, 2). This may be due  
703 to substantial recent interest in the Triassic–Jurassic transition, and the situation differs from  
704 previous compilations which demonstrated greater focus on the Late Jurassic of Europe  
705 (Riding 2012, 2013, 2014, 2019). Papers based on West Europe comprise 31.2% of the total,  
706 and publications on Africa, the Arctic, Australasia, East Europe and Russia are also  
707 significant (15.1%, 6.5%, 7.5%, 9.7% and 14.0% respectively). The least well-represented

708 regions are North America, South America and the Middle East (2.2%, 1.1% and 1.1%  
709 respectively; Table 1).

710

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723

### 724 **Disclosure statement**

725 The author has no potential conflict of interest.

726

### 727 **Notes on contributor**

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735 Jim is a past President of AASP – The Palynological Society, and became Managing Editor  
736 in 2004.

737

738

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1147

1148

1149 **Display material captions:**

1150

1151 Table 1. A breakdown of the 93 publications on Triassic to earliest Cretaceous dinoflagellate  
1152 cysts compiled herein, based on the nine relevant geographical regions (i.e. Africa, North  
1153 America, South America, the Arctic, Australasia, East Europe, West Europe, the Middle East  
1154 and Russia) and the initial letter of the family name of the first author. The number in the  
1155 geographical region cell refers to the number of relevant published items on that area alone.  
1156 An ellipsis (...) indicates a zero return for that particular parameter.

1157

1158 Table 2. A breakdown of of the 93 publications on Triassic to earliest Cretaceous (Berriasian)  
1159 dinoflagellate cysts compiled herein, subdivided into Triassic, Early Jurassic, Middle  
1160 Jurassic, Late Jurassic, Jurassic–Cretaceous transition, investigations comprising three or  
1161 more of the previous intervals and studies with no stratigraphical focus, and reworking. Some  
1162 latitude and pragmatism is used in this compilation; for example, if a publication is on the  
1163 Berriasian and Valanginian it is classified as covering the Jurassic–Cretaceous transition. One  
1164 item may be counted twice (e.g. if it spans the Oxfordian to Berriasian), but not three times.  
1165 An ellipsis (...) indicates a zero return for that particular parameter.

1166

1167

## 1168 **SUPPLEMENTARY DATA**

1169

### 1170 **Appendix 1. List of Literature**

1171

1172 Ninety-three contributions on Triassic to earliest Cretaceous dinoflagellate cysts issued after  
1173 the publication of Riding (2012, 2013, 2014, 2019), together with papers encountered after  
1174 these compilations were made, are listed in alphabetical/chronological order below. The

1175 reference citation format used is much the same as in Riding (2013, 2014, 2019), which was  
1176 slightly modified from Riding (2012). Digital object identifier (doi) numbers are included  
1177 where these are available. The 30 papers which are deemed to be of major significance are  
1178 asterisked. The language in which a paper was written in is indicated if it is not in English. A  
1179 synthesis of the scope of each item is given as a string of keywords in parentheses after each  
1180 citation. These keywords attempt to comprehensively summarise the principal subject matter,  
1181 age range, major geographical region(s) and country/countries. A distinction is made between  
1182 publications that present new data ('primary data'), and those that compile, review or  
1183 summarise existing datasets ('compilation'). A significant number of abstracts are listed here;  
1184 these are denoted by the word 'summary' in the keyword string. If the author(s) have  
1185 included photographs, occurrence charts and a zonal breakdown, these are indicated  
1186 respectively in the keywords. For the purpose of this work, the world is subdivided into 14  
1187 major geographical regions. These are Africa, Central America, North America, South  
1188 America, Antarctica, the Arctic, Southeast Asia, Australasia, China, East Europe, West  
1189 Europe, the Indian subcontinent, the Middle East and Russia (Table 1). The regional  
1190 assignments of any disputed territories, for example of Crimea, are merely for internal  
1191 consistency and geographical pragmatism, and have no political significance whatsoever.

1192

1193

1194

## A

1195

1196 \*ABOUL ELA, N.M., and TAHOUN, S.S. 2010. Dinoflagellate cyst stratigraphy of the  
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1199 *Valley University, Quena, Luxor, Egypt*, 85–115.

1200 (acritarchs; biostratigraphy; biozonation; calcareous nannofossils; correlation; eustasy;  
1201 foraminifera; foraminiferal test linings; fungal spores; hiatus; lithostratigraphy; pollen-spores;  
1202 tectonics; primary data; occurrence charts; photographs; Middle Jurassic–Early Cretaceous  
1203 [Bathonian–Albian]; North Africa [Mango-1 and Til-1 wells, offshore northern Sinai Basin,  
1204 Egypt])

1205

1206 \*ARKADIEV, V., GUZHIKOV, A., BARABOSHKIN, E., SAVELIEVA, J.,  
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1211 foraminifera; magnetostratigraphy; ostracods; prasinophytes; stratigraphical synthesis;  
1212 Tethyan palaeogeography; primary data; photographs; latest Jurassic–Early Cretaceous  
1213 [Tithonian–Valanginian]; sub-Arctic Russia [Crimean Mountains from Feodosiya to  
1214 Sevastopol, Crimea, southwestern Russia])  
1215  
1216 ARKADIEV, V.V., BARABOSHKIN E.Yu., GUZHIKOV, A.Yu., BARABOSHKIN, E.E.,  
1217 SHUREKOVA, O.V., and SAVELIEVA, Yu.N. 2018. *Tirnovella occitanica* zone  
1218 (Berriasian) of the Eastern Crimea. In: Baraboshkin E.Yu., Lipnitskaya, T.A., and Guzhikov,  
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1223 (ammonites; biostratigraphy; biozonation; magnetostratigraphy; ostracods; pollen-spores;  
1224 *Tirnovella occitanica* zone; summary; earliest Cretaceous [Berriasian]; sub-Arctic Russia  
1225 [Zavodskaya Balka section, eastern Crimea, southwestern Russia])  
1226  
1227

## 1228 B

1229  
1230 BARABOSHKIN, E.Yu., SHTUN, S.Yu., GUZHIKOV, A.Yu., KUZNETZOV, A.B.,  
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1234 *Cretaceous system of Russia and near abroad: problems of stratigraphy and*  
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1236 Research University, Belgorod, September 17th–21st, 2018. Polyterra Publishing House,  
1237 Belgorod, ISBN 978-5-98242-250-7, 54–58 (in Russian with an English abstract).  
1238 (biostratigraphy; correlation; palaeoecology; palaeomagnetism; sedimentology; stable  
1239 isotopes; summary; latest Jurassic–earliest Cretaceous [Tithonian–Berriasian]; sub-Arctic  
1240 Russia [the Khvalynskaya-5 and Sarmatskaya-3 wells, northern Caspian Sea, offshore  
1241 southwestern Russia])

1242  
1243 BARANYI, V. 2018. *Vegetation dynamics during the Late Triassic (Carnian–Norian):*  
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1245 thesis, University of Oslo, Norway, 164 p., ISSN 1501-7710 (available online at:  
1246 <https://www.duo.uio.no/handle/10852/61940>).  
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1248 history; Carnian Pluvial Episode; Mid-Norian Climate Shift; lithostratigraphy [the Chinle and  
1249 Veszprém Marl formations and the Mercian Mudstone Group]; palaeoclimatology;  
1250 palaeoecology; Sub-Boreal and Tethyas realms; taphonomy; terrestrial ecosystems; Late  
1251 Triassic [Carnian–Norian]; multi-region: sub-Arctic North America [Petrified Forest National  
1252 Park, Arizona, southwestern USA]; East Europe [southern Transdanubian Range, western  
1253 Hungary]; sub-Arctic West Europe [Wessex Basin, southwestern England])  
1254  
1255 \*BARSKI, M. 2018. Dinoflagellate cyst assemblages across the Oxfordian/Kimmeridgian  
1256 boundary (Upper Jurassic) at Flodigarry, Staffin Bay, Isle of Skye, Scotland - a proposed  
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1259 (ammonites; biostratigraphy; bloom of zygnetacean chlorophycean alga [*Spirogyra*];  
1260 correlation; eutrophication event [“green tide”]; lithostratigraphy [Flodigarry Shale Member,  
1261 Staffin Bay Formation]; palynofacies; proposed Global Stratotype Section and Point (GSSP);  
1262 primary data; quantitative occurrence chart; photographs; Late Jurassic [Oxfordian–  
1263 Kimmeridgian]; sub-Arctic West Europe [foreshore sections at Flodigarry, Trotternish  
1264 Peninsula, Isle of Skye, northwestern Scotland])  
1265  
1266 BARTH, G., PIENKOWSKI, G., ZIMMERMANN, J., FRANZ, M., and KUHLMANN, G.  
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1272 (ammonites; ammonite biozones; basin evolution; biofacies; biostratigraphy; Central  
1273 European Basin; correlation; *Dapcodinium priscum*; *Liasidium variabile*; *Luehndea spinosa*;  
1274 palaeoecology; palaeogeography; palaeontology; pollen-spores; sedimentary architecture;  
1275 sequence stratigraphy; compilation; Late Triassic–Middle Jurassic [Norian–Callovian]; multi-

1276 region: East Europe [Poland]; sub-Arctic West Europe [Denmark; England, France,  
1277 Germany])  
1278  
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1281 and Palynology (LPP), State University of Utrecht, The Netherlands.  
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1283 pollen-spores; species; compilation; Triassic [undifferentiated]; no specific geographical  
1284 focus)

1285  
1286 BÓNA, J. 1995. Palynostratigraphy of the Upper Triassic formations in the Mecsek Mts  
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1288 (acritarchs; algae; biostratigraphy; coal exploration boreholes; foraminiferal test linings;  
1289 hiatuses; lithostratigraphy [Kantavár, Karolinavölgy Sandstone and lowermost Mecsek Coal  
1290 formations]; pollen-spores; primary data; occurrence charts; photographs; Late Triassic  
1291 [Carnian–Rhaetian]; East Europe [Pécs region, Mecsek Mountains, southern Hungary])  
1292

## 1293 C

1294  
1295  
1296 \*CORREIA, V.D.P.F. 2018. *Jurassic dinoflagellate cyst biostratigraphy of the Lusitanian*  
1297 *Basin, west-central Portugal, and its relevance to the opening of the North Atlantic and*  
1298 *petroleum geology*. Unpublished PhD thesis, Universidade do Algarve, Portugal, xxxii + 283  
1299 p.  
1300 (acritarchs; ammonite biozones; biostratigraphy; biozonation; correlation; diversity;  
1301 foraminiferal test linings; geochemistry; lithostratigraphy; palaeobiology; palaeoecology;  
1302 palaeogeography; palynomorph fluorescence; petroleum geology; pollen-spores;  
1303 prasinophytes; provincialism; regional geology; tectonic history; thermal alteration index  
1304 [TAI]; Toarcian Oceanic Anoxic Event [T-OAE]; primary data; occurrence charts;  
1305 photographs; Early–Middle Jurassic [Sinemurian–Callovian]; sub-Arctic West Europe  
1306 [Lusitanian Basin, western-central Portugal])  
1307

1308 CORREIA, V.F., RIDING, J.B., FERNANDES, P., DUARTE, L.V., and PEREIRA, Z. 2016.  
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1312 108.

1313 (abundance and diversity fluctuations; *Luehndea spinosa*; palaeoenvironmental change;  
1314 Toarcian Oceanic Anoxic Event (T-OAE); recovery; summary; Early Jurassic [Toarcian];  
1315 sub-Arctic West Europe [Maria Pares, Peniche, Vale das Fontes, Lusitanian Basin, western-  
1316 central Portugal])

1317

1318 CORREIA, V.F., RIDING, J.B., FERNANDES, P., DUARTE, L.V., and PEREIRA, Z. 2017.  
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1322 (abundance and diversity; *Luehndea spinosa*; palaeobiology; palaeoceanography; Toarcian  
1323 Oceanic Anoxic Event (T-OAE); recovery; summary; photographs; Early Jurassic [Toarcian];  
1324 sub-Arctic West Europe [Maria Pares, Peniche, Vale das Fontes, Lusitanian Basin, western-  
1325 central Portugal])

1326

1327 CORREIA, V.F., RIDING, J.B., HENRIQUES, M.H., FERNANDES, P., and PEREIRA, Z.  
1328 2017. The dinoflagellate cysts of the Bajocian GSSP (Middle Jurassic) at Cabo Mondego,  
1329 Lusitanian Basin, Portugal. *Eleventh International Conference on Modern and Fossil*  
1330 *Dinoflagellates (DINO 11), Bordeaux, France, 17th–21st July 2017*, p. 52.

1331 (ammonites; biostratigraphy; diversity; evolutionary radiation; Global Stratotype Section and  
1332 Point (GSSP); Mesozoic Marine Revolution; summary; Early–Middle Jurassic [Toarcian–  
1333 Bajocian]; sub-Arctic West Europe [Cabo Mondego, Lusitanian Basin, western-central  
1334 Portugal])

1335

1336 CORREIA, V.F., RIDING, J.B., DUARTE, L.V., FERNANDES, P., and PEREIRA, Z. 2018.  
1337 An overview of the Lower Jurassic dinoflagellate cyst biostratigraphy in the Lusitanian  
1338 Basin, Portugal. In: Vaz, N., and Sá, A.A. (editors). *Yacimientos paleontológicos*  
1339 *excepcionales en la península Ibérica. Cuadernos Del Museo Geominero*, No. 27: 335–342.  
1340 Instituto Geológico y Minero de España, Madrid. ISBN 978-84-9138-066-5.

1341 (ammonite biozones; biostratigraphy; diversity; summary; occurrence chart; photographs;  
1342 Early Jurassic [Sinemurian–Toarcian]; sub-Arctic West Europe [Brenha, Fonte Coberta,



1343 Maria Pares, Peniche, São Pedro de Moel and Val das Fontes, Lusitanian Basin, western  
1344 central Portugal])  
1345  
1346 CORREIA, V.F., RIDING, J.B., DUARTE, L.V., FERNANDES, P., and PEREIRA, Z. 2018.  
1347 Lower Jurassic dinoflagellate cysts throughout the Toarcian Oceanic Anoxic Event (T-OAE)  
1348 in the Lusitanian Basin, Portugal. *In*: Silva, R.L., Duarte, L.V., and Sêco, S. (editors). *Second*  
1349 *International Workshop on the Toarcian Oceanic Anoxic Event, Coimbra, Portugal,*  
1350 *September 6–9, 2018. Abstracts Volume, 33–34. ISBN: 978-989-98914-6-3.*  
1351 (abundances; diversity; palaeoecology; palaeogeography; recovery; Toarcian Oceanic Anoxic  
1352 Event (T-OAE); summary; photographs; Early Jurassic [Pliensbachian–Toarcian]; sub-Arctic  
1353 West Europe [Fonte Coberta, Maria Pares, Peniche, Lusitanian Basin, western central  
1354 Portugal])  
1355  
1356 \*CORREIA, V.F., RIDING, J.B., DUARTE, L.V., FERNANDES, P., and PEREIRA, Z.  
1357 2018. The Early Jurassic palynostratigraphy of the Lusitanian Basin, western Portugal.  
1358 *Geobios*, 51(6): 537–557 (doi: 10.1016/j.geobios.2018.03.001).  
1359 (acritarchs; ammonite biozones; biostratigraphy; biozonation; diversity; foraminiferal test  
1360 linings; pollen-spores; prasinophytes; regional geology; Toarcian Oceanic Anoxic Event (T-  
1361 OAE); primary data; occurrence charts; photographs; Early Jurassic [Sinemurian–Toarcian];  
1362 sub-Arctic West Europe [Brenha, Fonte Coberta, Maria Pares, Peniche, São Pedro de Moel  
1363 and Val das Fontes, Lusitanian Basin, western central Portugal])  
1364  
1365 \*CORREIA, V.F., RIDING, J.B., HENRIQUES, M.H., FERNANDES, P., PEREIRA, Z.,  
1366 and WIGGAN, N.J. 2019. The Middle Jurassic palynostratigraphy of the northern Lusitanian  
1367 Basin, Portugal. *Newsletters on Stratigraphy*, 52(1): 73–96 (doi: 10.1127/nos/2018/0471).  
1368 (acritarchs; ammonite biozones; archaeopyle types; biostratigraphy; diversity; eustasy;  
1369 evolutionary radiation of the family Gonyaulaceae; foraminiferal test linings; Global  
1370 Stratotype Section and Point (GSSP); lithostratigraphy (Cabo Mondego and Póvoa de Lomba  
1371 formations); palaeobiology; palaeogeography; regional geology; pollen-spores;  
1372 prasinophytes; Toarcian Oceanic Anoxic Event (T-OAE); primary data; quantitative  
1373 occurrence charts; photographs; Early–Middle Jurassic [Toarcian–Bathonian]; sub-Arctic  
1374 West Europe [Cabo Mondego and São Gião, northern Lusitanian Basin, western central  
1375 Portugal])  
1376

1377  
1378 **D**  
1379

1380 DARWISH, M., EL-ARABY, A., ABU KHADRAH, A.M., and HUSSEIN, H.M. 2004.  
1381 Sedimentary facies models and organic geochemical aspects of the Upper Jurassic–Lower  
1382 Cretaceous sequences in northern Sinai. *Sixth International Conference on Geochemistry,*  
1383 *Alexandria University, Egypt, 15<sup>th</sup> to 16<sup>th</sup> September 2004*, 183–208.

1384 (correlation; foraminifera; lithofacies analysis; lithostratigraphy; organic geochemistry;  
1385 palaeogeography; palynofacies; petroleum geology; compilation; Middle Jurassic–Early  
1386 Cretaceous [Callovian–Aptian]; North Africa [offshore and onshore northern Sinai, Egypt])

1387  
1388 DIXON, M., MORGAN, R., GOODALL, J., and VAN DEN BERG, M. 2012. Higher-  
1389 resolution palynostratigraphy of the Norian–Carnian (Triassic) Upper Mungaroo Formation,  
1390 offshore Carnarvon Basin (extended abstract). *APPEA 2012 Journal and Conference*  
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1423 palaeoclimate; palaeoecology; palaeotemperature; pollen-spores; shallow water deposition;  
1424 primary data; semiquantitative occurrence charts; photographs; latest Jurassic–earliest  
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1487 Russia [Sjuk River valley, north of the Dakh Crystalline Massif, Republic of Adygeja,  
1488 northwestern Caucasus])  
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1496 *Nannoceratopsis spiculata*; pollen-spores; regional geology; sandstone; X-ray diffraction  
1497 [XRD] of minerals; primary data; photographs; Early Jurassic [Toarcian]; sub-Arctic Russia  
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1511 *Nannoceratopsis tricerias*; taxonomy; *Wanaea acollaris*; primary data; occurrence charts;

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1537 palaeobotany; palaeoclimate; palaeoecology; plant macrofossils; prasinophytes; Ravenscar  
1538 Group; sequence stratigraphy; primary data; quantitative occurrence charts; photographs;  
1539 Middle Jurassic [Bajocian–Bathonian]; sub-Arctic West Europe [Cloughton Wyke, Crook  
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1548 spores; prasinophytes; sequence stratigraphy; statistics; systematics; primary data;  
1549 quantitative occurrence charts; photographs; Late Triassic [Norian–Rhaetian]; sub-Arctic  
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1559 formations; ostracods; palaeoecology; palaeogeography; Pindi Group; pollen-spores;  
1560 regional geology; reworking; sedimentology; tectonic rifting of Gondwana; compilation;  
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1572 occurrence chart; photographs; Early Jurassic–Early Cretaceous [undifferentiated–  
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1611 regional geology; *Rhaetogonyaulax rhaetica*; sedimentology; sequence stratigraphy; well-log  
1612 analysis; primary data; non-quantitative occurrence chart; photographs; Permian



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1626 data; semi-quantitative occurrence chart; photographs; latest Triassic–earliest Jurassic  
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1636 potential; trace fossils; primary data; Late Jurassic–Early Cretaceous [Oxfordian–  
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1651 geochronology; hiatuses; lithostratigraphy; methane release; pollen-spores; prasinophytes;  
1652 *Rhaetogonyaulax rhaetica*; sedimentology; seismic activity; Triassic–Jurassic boundary;  
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1675 primary data; quantitative occurrence chart; photographs; Late Jurassic–Early Cretaceous  
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1693 foraminifera; ostracods; pollen-spores; primary data; semi-quantitative occurrence chart;  
1694 photographs; Middle Jurassic [Bajocian–Bathonian]; sub-Arctic Russia [Bolshoi Zelenchuk  
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1705 linings; lithostratigraphy [Djangura Formation]; ostracods, pollen-spores; primary data;  
1706 occurrence chart; photographs; Middle Jurassic [Bajocian]; sub-Arctic Russia [banks of the  
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1722 province; informal taxonomy; Kipatimi Formation; Makonde and Mkindani beds;  
1723 palaeoecology; palaeogeography; palynofacies; petroleum geology; pollen-spores;  
1724 quantitative bioevents; regional geology; systematics; primary data; occurrence charts;  
1725 photographs; Middle Jurassic–earliest Late Cretaceous [Bajocian–Cenomanian]; East Africa  
1726 [coastal Tanzania])  
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1737 pollen-spores; primary data; Early Jurassic–Cretaceous [Hettangian–Turonian]; Arctic Russia  
1738 [New Siberian Islands Archipelago, East Siberian and Laptev seas])  
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- 1740 NIKITENKO, B.L., DEVYATOV, V.P., LEBEDEVA, N.K., BASOV, V.A., FURSENKO,  
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1746 ostracods; pollen-spores; molecular biomarkers; organic geochemistry; petroleum geology;

1747 primary data; Early Jurassic–Late Cretaceous [Hettangian–Turonian]; Arctic Russia [New  
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1755 foraminifera; geochemistry ( $\delta^{13}\text{C}_{\text{TOC}}$ ); lithostratigraphy [Buolkalakh, Chekurovka and  
1756 Iaedaes formations]; palaeoclimatology; palaeoecology; pollen-spores; prasinophytes;  
1757 sedimentology; shallow water deposition; Volgian and Boreal Berriasian; primary data; semi-  
1758 quantitative occurrence chart; photographs; Middle Jurassic–Early Cretaceous [Bathonian–  
1759 Valanginian]; Arctic Russia [Outcrop O14, Olenek section, Anabar-Lena region, eastern  
1760 Siberia, northeastern Russia])  
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1770 Muerta Formation]; palaeoclimate; palaeogeography; pollen-spores; prasinophytes;  
1771 *Shanbeipollenites proxireticulatus* [pollen]; volcanic activity; primary data; semiquantitative  
1772 occurrence chart; photographs; Early Cretaceous [?Berriasian–Valanginian]; South America  
1773 [Mallin Quemado, north of Zapala, Neuquén Basin, central western Argentina])  
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1784 regional geology; rift tectonics; sedimentary architecture; sediment thickness maps; seismic  
1785 stratigraphy; sequence stratigraphy; shelf margin depositional setting; latest Jurassic–Early  
1786 Cretaceous [Tithonian–Valanginian]; Australasia [Northern Carnarvon Basin, North West  
1787 Shelf, offshore Western Australia])  
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1794 geographical distributions; history of research; living dinoflagellates; palaeoecology;  
1795 palaeogeography; palaeoceanography; primary productivity; transfer functions; literature  
1796 review; occurrence chart; photographs; no geographical and stratigraphical focus)  
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1805 summary; latest Jurassic–earliest Cretaceous [Tithonian–Berriasian (=Volgian–Ryazanian);  
1806 sub-Arctic Russia [Maurynya River outcrop section, south of Tolya, Northern Ural  
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1813 (archaeopyle; biostratigraphy; *Dingodinium*; family Cladopyxiaceae [suborder  
1814 Cladopyxiineae]; morphology; partiform tabulation pattern; taxonomy; primary data and

1815 compilation; line drawings; photographs; occurrence chart; Late Jurassic [Kimmeridgian–  
1816 Tithonian]; sub-Arctic Russia [Gorodishche, Volga River, near Ulyanovsk, southwestern  
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1824 organic geochemistry; palaeoclimate; palaeoecology; palynofacies; peat-forming  
1825 environments; pollen-spores; *Rhaetogonyaulax rhaetica*; sedimentology; sequence  
1826 stratigraphy; vegetational analysis/dynamics; Triassic–Jurassic boundary; primary data;  
1827 occurrence charts; photographs; latest Triassic [Rhaetian]; sub-Arctic West Europe [Lunnom  
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1834 palynofacies; pollen-spores; sedimentology; sequence stratigraphy; primary data;  
1835 photographs; Early Jurassic [Hettangian–Toarcian]; East Europe [Poland])

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1842 borehole; lithostratigraphy; palaeoclimate; palaeogeography; pollen-spores; *Rhaetogonyaulax*  
1843 *rhaetica*; sedimentology; sequence stratigraphy; compilation; latest Triassic–earliest Jurassic  
1844 [Rhaetian–Hettangian]; East Europe [near Szczecin, Pomerania, northwestern Poland])

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1851 Formation; Kellaways Sand Member; macrofossils; palaeogeography; sedimentology;  
1852 thickness variations; trace fossils; palynofacies; pollen spores; primary data; occurrence data;  
1853 photographs; Middle Jurassic [Callovian]; sub-Arctic West Europe [Burythorpe Quarry, near  
1854 Malton, Howardian Hills, North Yorkshire, northern England])

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1864 correlation; facies analysis/associations; lithostratigraphy [Brentskardhaugen Bed;  
1865 Wilhelmøya Subgroup; De Geerdalen, Knorringsfjellet and Agardhfjellet formations; Kapp  
1866 Toscana Group]; pollen-spores; provenance; regional geology; reworking; sedimentology;  
1867 sequence stratigraphy; trace fossils; primary data; Late Triassic–Middle Jurassic [Carnian–  
1868 Callovian]; Arctic [western central Spitsbergen, Svalbard])

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1874 (biostratigraphy; biozonation; botanical affinities; Central Atlantic Magmatic Province  
1875 (CAMP); clay mineralogy; *Dapcodinium priscum*; diversity; end Triassic mass extinction;  
1876 foraminiferal test linings; lithofacies; palaeoclimate; palaeoecology; palaeogeography;  
1877 palynofacies; Pangaea; pollen-spores; prasinophytes; *Rhaetogonyaulax rhaetica*; spore spike;  
1878 sporomorph ecogroups; stable isotope geochemistry; statistics; Tethyan Realm; primary data;  
1879 occurrence charts; photographs; latest Triassic–earliest Jurassic [Rhaetian–Hettangian]; East  
1880 Europe [Csövár, northern Hungary, the Mecsek Mountains, southern Hungary and the Tatra  
1881 Mountains, northern Slovakia])



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1919 mass extinction; eustasy; foraminiferal test linings; lithostratigraphy [Klettgau and Staffelegg  
1920 formations]; palaeoecology; pollen-spores; regional geology; primary data; quantitative  
1921 occurrence chart; photographs; Late Triassic–Early Jurassic [Norian–Sinemurian]; sub-Arctic  
1922 West Europe [Adlerberg, Chilchzimmersattel and Weiach, northern Switzerland])  
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1932 (biostratigraphy; freshwater algae; Jurassic–Cretaceous boundary; lithostratigraphy [Münder  
1933 Formation and Bückeberg Group]; Lower Saxony Basin; organic geochemistry; ostracods;  
1934 palaeoclimatology; palaeoecology; palynofacies; petroleum prospectivity; pollen-spores;  
1935 regional geology; Wealden type facies; primary data; photographs; latest Jurassic–earliest  
1936 Cretaceous [Tithonian–Berriasian]; sub-Arctic West Europe [the Husen 1/08 borehole,  
1937 southwest of Hannover, northern Germany])  
1938  
1939 SCHNEIDER, A.C., HEIMHOFER, U., HEUNISCH, C., and MUTTERLOSE, J. 2018.  
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1943 lithostratigraphy [Münder Formation and Bückeberg Group]; palaeoclimatology;  
1944 palaeoecology; pollen-spores; vegetational dynamics; Wealden type facies; primary data;  
1945 photographs; quantitative occurrence charts; latest Jurassic–earliest Cretaceous [Tithonian–  
1946 Berriasian]; sub-Arctic West Europe [boreholes C-1 and Eulenflucht-1 (E-1), west of  
1947 Hannover, Lower Saxony Basin, northern Germany])

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 1953 Jurassic–Cretaceous boundary; lithostratigraphy [Münder Formation and Bückeberg Group];  
 1954 ostracods; palaeoclimatology; palaeoecology; pollen-spores; provincialism; Purbeck and  
 1955 Wealden type facies; regional geology; primary data; photographs; latest Jurassic–earliest  
 1956 Cretaceous [Tithonian–Berriasian]; sub-Arctic West Europe [boreholes 1/08 Husen (H-1),  
 1957 “core-1” (C-1), Eulenflucht-1 (E-1) and Isterberg 1001 (I-1), west of Hannover, Lower  
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 1968 Timan-Northern Ural Region, Komi Republic])
- 1969
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 1976 (acritarchs; biostratigraphy; biozonation; correlation; summary of primary data; semi-  
 1977 quantitative range chart; photographs; earliest Cretaceous [Berriasian]; sub-Arctic Russia  
 1978 [Uruh, Crimean Mountains, North Caucasus, southwestern Russia])
- 1979
- 1980 SMELROR, M., FOSSUM, K., DYPVIK, H., HUDSON, W., and MWENEINDA, A. 2018.  
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1985 assemblage; eustasy; freshwater algae; fungal palynomorphs; lithostratigraphy [the Kipatimu,  
1986 Mitole, Nalwehe and Kihuluhulu formations]; palaeoecology; pollen-spores; prasinophytes;  
1987 regional geology; primary data; photographs; Late Jurassic–Early Cretaceous [Oxfordian–  
1988 Tithonian to Aptian–Albian]; East Africa [Mandawa Basin, southeastern coastal Tanzania])  
1989

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1991 *Basin, Russia*. Unpublished PhD Thesis, University of Bristol, UK, 527 p. (available online  
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1995 palynofacies; pollen-spores; prasinophytes; regional stratigraphical synthesis; sedimentology;  
1996 total organic carbon [TOC]; Volgian lectostratotype; primary data; occurrence charts;  
1997 photographs; latest Jurassic–Early Cretaceous [Tithonian–Valanginian]; sub-Arctic Russia  
1998 [Gorodische and Kashpir, Ulyanovsk region, Volga River Basin, western Russia])  
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2005 (*Beaumontella langii*; coral bleaching; ecology; evolution; fossil and modern coral reefs;  
2006 modern corals; photosymbiosis; *Suessia swabiana*; *Symbiodinium*; review; photographs;  
2007 occurrence chart; Late Triassic–Early Jurassic [Carnian–Toarcian]; multi-region: North  
2008 Africa [Libya]; Australasia [northwestern Australia]; sub-Arctic West Europe [Austria,  
2009 southwestern Germany, southern Sweden])  
2010

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2016 foraminiferal test linings; inorganic and organic geochemistry; palaeoceanography;  
2017 palaeogeography; palynofacies; Pieniny Klippen Belt; pollen-spores; prasinophytes; pyrite  
2018 framboid size; regional geology; scolecodonts; Tethyan Realm; Toarcian Oceanic Anoxic  
2019 Event [T-OAE]; primary data; Early Jurassic [Toarcian]; East Europe [Zázrivá, northwestern  
2020 Slovakia])

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2027 foraminiferal test linings; graptolites; history of research; *incertae sedis*; organic-walled  
2028 microplankton; prasinophytes; scolecodonts; testate amoebae [formerly thecamoebians];  
2029 zooplankton; compilation of genera and species; Silurian [undifferentiated]–Holocene; East  
2030 Europe [Hungary])

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2039 palynomorphs; palaeoecology; pollen-spores; primary data; occurrence charts; photographs;  
2040 Middle Jurassic–Late Cretaceous [Callovian–Oxfordian and Barremian–Cenomanian]; North  
2041 Africa [Alamein-IX well, northern Western Desert, Egypt])

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2046 biozonation; Tethys; review/synthesis; occurrence chart; Late Jurassic–Late Cretaceous  
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2054 chronostratigraphy; correlation; cyclostratigraphy; gamma ray logs; geochemistry;  
2055 lithostratigraphy [Kimmeridge Clay Formation]; pollen-spores; wavelet analyses; primary  
2056 data; occurrence chart; Late Jurassic–Early Cretaceous [Oxfordian–Valanginian]; multi-  
2057 region: Arctic [Barents Sea; Arctic Russia; Svalbard]; East Europe [Brodno, Slovakia]; sub-  
2058 Arctic West Europe [Dorset, southern England; North Sea, offshore Norway])

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## V

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2064 WONIK, T., BLAU, J., MEISTER, C., VAN DER WEIJST, C.M.H., SUAN, G.,  
2065 FRAGUAS, A., FIEBIG, J., HERRLE, J.O., GUEX, J., LITTLE, C.T.S., WIGNALL, P.B.,  
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2070 shales; calcareous nannofossils; carbon cycle perturbations; chemostratigraphy; correlation;  
2071 end-Triassic extinction; eustasy; geochemistry [ $\delta^{13}\text{C}_{\text{org}}$ ;  $\delta^{18}\text{O}$ ]; geophysical downhole  
2072 logging; glendonites; hydrocarbon seepage; lithostratigraphy; marine anoxia; ostracods;  
2073 palaeoecology; pollen-spores; reworking; sedimentology; seismic activity; taxonomy;  
2074 primary data; photographs; latest Triassic–Early Jurassic [Rhaetian–Toarcian]; sub-Arctic  
2075 West Europe [Schandelah-1 borehole, Lehre, Landkreis Helmstedt, Lower Saxony, northern  
2076 Germany])

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2084 interactions; Lepidoptera (butterflies and moths); lepidopteran wing scales; *Lunnomidinium*  
2085 *scaniense*; mass extinction; pollen-spores; primary data; quantitative selective occurrence  
2086 chart; latest Triassic–Early Jurassic [Rhaetian–Sinemurian]; sub-Arctic West Europe  
2087 [Schandelah-1 well, near Braunschweig, Lower Saxony, northern Germany])  
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2093 petroleum geology; plant fossils; pollen-spores; scolecodonts; research proposal; Jurassic  
2094 [undifferentiated]; sub-Arctic North America [Alberta and Saskatchewan, western Canada])  
2095

2096 \*VERREUSSEL, R.M.C.H., BOUROULLEC, R., MUNSTERMAN, D.K., DYBKJÆR, K.,  
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2104 electric logs; lithostratigraphy; palaeoecology; palaeoceanography; petroleum geology;  
2105 pollen-spores; sedimentary architecture and facies; sedimentology; structural geology;  
2106 tectonic extension and rifting; tectonostratigraphical mega-sequences (TMS); compilation;  
2107 Middle Jurassic–Early Cretaceous [Bathonian–Barremian]; sub-Arctic West Europe [Danish-  
2108 Dutch-German Central Graben, North Sea])  
2109

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2114 *rhaetica*; *Suessia swabiana*; Tethyan Realm; compilation; occurrence charts; photographs;  
2115 Triassic–earliest Jurassic [Induan–Hettangian]; multi-region: East Europe and sub-Arctic

2116 West Europe [The Alpine region, i.e. Austria; Germany, Hungary, Italy and the former  
2117 Yugoslavia])

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2126 colonial alga; correlation; estuarine, freshwater-brackish dinoflagellate cysts; marine and

2127 terrestrial [fluviolacustrine] facies analysis; eustasy; isopach maps; lithostratigraphy;

2128 palaeoclimatology; palaeodrainage patterns; palaeogeography; regional geology;

2129 sedimentology; sequence stratigraphy; Walloon Coal Measures [Injune Creek Group];

2130 uranium–lead [U–Pb] dating; wireline log analysis; primary data; photographs; Late Jurassic

2131 [Oxfordian–Tithonian]; Australasia [Surat Basin, Queensland, eastern Australia])

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2133 WAINMAN, C.C., HANNAFORD, C., MANTLE, D., and McCABE, P.J. 2018. Utilizing  
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2138 geochronology; *Moorodinium*; pollen-spores; regional geology; *Skuadinium*; tuff horizons;

2139 uranium–lead chemical abrasion thermal ionization mass spectroscopy [U–Pb CA-TIMS]

2140 dating; Walloon Coal Measures [Injune Creek Group]; primary data; photographs; Middle–

2141 Late Jurassic [Bathonian–Tithonian]; Australasia [Surat Basin, Queensland, eastern

2142 Australia])

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2144 \*WAINMAN, C.C., MANTLE, D.J., HANNAFORD, C., and McCABE, P.J. 2018. Possible  
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2148 morphology; palaeoecology; pollen-spores; sea levels; sedimentology; taxonomy; uranium–

2149 lead [U–Pb] dating; Walloon Coal Measures [Injune Creek Group]; primary data; occurrence



2150 chart; photographs; Late Jurassic [Tithonian]; Australasia [Indy 3 well, western Surat Basin,  
2151 Queensland, southeastern Australia])  
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2157 (acritarchs; biostratigraphy; palaeoecology; regression-transgression; spore-pollen;  
2158 sporomorph eco-grouping; statistics; summary; latest Triassic–earliest Jurassic [Rhaetian–  
2159 Hettangian]; sub-Arctic West Europe [Carnduff-1 Borehole, near Larne, Northern Ireland])  
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2164 (ammonites; belemnites; biological productivity; bivalves; bryozoan; carbon isotope  
2165 geochemistry; coccolithophores; continental weathering; *Dissiliodinium*; diversity; evolution;  
2166 evolutionary radiation; fish; Gonyaulacaceae; Mesozoic Marine Revolution; nutrient flux;  
2167 palaeoclimate; palaeoecology; palaeogeography; palaeoceanography; phytoplankton;  
2168 planktonic foraminifera; sea level; sequence stratigraphy; *Watznaueria*; compilation;  
2169 occurrence charts; photographs; Middle Jurassic [Aalenian–Bathonian]; sub-Arctic West  
2170 Europe [no specific geographical focus])  
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2177 (aquifers; biostratigraphy; boreholes; correlation; downhole logging; fluvial sandstones;  
2178 geothermal groundwaters and heat; lateral thickness variability; Nieuwerkerk Formation;  
2179 pollen-spores; sedimentary architecture; seismic interpretation; sequence stratigraphy;  
2180 sporomorph eco-grouping; tectonic activity; primary data; Early Cretaceous  
2181 [Berriasian/Ryazanian–Barremian]; sub-Arctic West Europe [West Netherlands Basin, The  
2182 Netherlands])  
2183

2184 WILLIAMS, G., FENSOME, R., MILLER, M., and BUJAK, J. 2018. Microfossils:  
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2188 miscellaneous palynomorphs; morphology; palaeoclimatology; palaeoecology;  
2189 palaeogeography; palynofacies; palynology; pollen-spores; preparation techniques;  
2190 photographs; review article; no specific geographical or stratigraphical focus)

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2199 (assemblage zones; biostratigraphy; correlations with China; lithostratigraphy; Murihiku  
2200 Terrane; pollen-spores; systematics; Triassic–Jurassic boundary; primary data; occurrence  
2201 charts; photographs; Late Triassic–Early Jurassic [Norian–Sinemurian]; Australasia  
2202 [Awakino Gorge, southwestern Auckland, Hokonui Hills, Southland, and southwestern  
2203 Kawhia, New Zealand])

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## 2206 **Appendix 2. List of palynomorph species and subspecies**

2207

2208 This Appendix alphabetically lists all valid, formally-defined palynomorph taxa below  
2209 generic level which are mentioned in this contribution with full author citations. This listing  
2210 largely comprises dinoflagellate cysts, together with colonial alga and miospores; these are  
2211 given separately below. References to the majority of the author citations for the  
2212 dinoflagellate cysts can be found in Williams et al. (2017). Note that the proposals of Correia  
2213 et al. (2017, p. 93, appendix 2) regarding the taxonomy of *Nannoceratopsis gracilis* and  
2214 *Nannoceratopsis senex* are followed herein.

2215

### 2216 **Dinoflagellate cysts:**

2217 *Acanthaulax crispa* (Wetzel 1967) Woollam & Riding 1983

- 2218 *Aldorfia aldorfensis* (Gocht 1970) Stover & Evitt 1978
- 2219 *Ambonosphaera calloviana* Fensome 1979
- 2220 *Amphorulacysta? dodekovae* (Zotto et al. 1987) Williams & Fensome 2016
- 2221 *Amphorulacysta? expirata* (Davey 1982) Williams & Fensome 2016
- 2222 *Batioladinium pomum* Davey 1982
- 2223 *Batioladinium reticulatum* Stover & Helby 1987
- 2224 *Batioladinium varigranosum* (Duxbury 1977) Davey 1982
- 2225 *Beaumontella langii* (Wall 1965) Below 1987
- 2226 *Canningia reticulata* Cookson & Eisenack 1960
- 2227 *Cantulodinium speciosum* Alberti 1961
- 2228 *Chytroeisphaeridia chytroeides* (Sarjeant 1962) Downie & Sarjeant 1965
- 2229 *Circulodinium distinctum* (Deflandre & Cookson 1955) Jansonius 1986
- 2230 *Cometodinium whitei* (Deflandre & Courteville 1939) Stover & Evitt 1978
- 2231 *Compositosphaeridium? polonicum* (Górka 1965) Lentin & Williams 1981
- 2232 *Cribroperidinium? longicorne* (Downie 1957) Lentin & Williams 1985
- 2233 *Ctenidodinium combazii* Dupin 1968
- 2234 *Ctenidodinium continuum* Gocht 1970
- 2235 *Ctenidodinium cornigerum* (Valensi 1947) Jan du Chêne et al. 1985
- 2236 *Ctenidodinium elegantulum* Millioud 1969
- 2237 *Ctenidodinium ornatum* (Eisenack 1935) Deflandre 1938
- 2238 *Ctenidodinium sellwoodii* (Sarjeant 1975) Stover & Evitt 1978
- 2239 *Cyclonephelium cuculliforme* (Davies 1983) Århus 1992
- 2240 *Dapcodinium priscum* Evitt 1961
- 2241 *Dichadogonyaulax? pannea* (Norris 1965) Sarjeant 1969
- 2242 *Dingodinium albertii* Sarjeant 1966
- 2243 *Dingodinium jurassicum* Cookson & Eisenack 1958
- 2244 *Dingodinium tuberosum* (Gitmez 1970) Fisher & Riley 1980
- 2245 *Dingodinium nequeas* Pestchevitskaya 2018
- 2246 *Durotrigia daveyi* Bailey 1987
- 2247 *Endoscrinium galeritum* (Deflandre 1938) Vozzhennikova 1967
- 2248 *Egmontodinium polyplacophorum* Gitmez & Sarjeant 1972
- 2249 *Emmetrocyta sarjeantii* (Gitmez 1970) Stover & Evitt 1978
- 2250 *Epiplosphaera gochtii* (Fensome 1979) Brenner 1988
- 2251 *Epiplosphaera reticulospinosa* Klement 1960

- 2252 *Gochteodinia villosa* (Vozzhennikova 1967) Norris 1978
- 2253 *Gonyaulacysta centriconnata* Riding 1983
- 2254 *Gonyaulacysta eisenackii* (Deflandre 1938) Górká 1965
- 2255 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965
- 2256 *Hebecysta balmei* (Stover & Helby 1987) Below 1987
- 2257 *Heibergella asymmetrica* Bujak & Fisher 1976
- 2258 *Hystrichosphaerina? orbifera* (Klement 1960) Stover & Evitt 1978
- 2259 *Impletosphaeridium varispinosum* (Sarjeant 1959) Islam 1993
- 2260 *Kaiwaradinium scrutinum* Backhouse 1987
- 2261 *Kalyptea stegasta* (Sarjeant 1961) Wiggins 1975
- 2262 *Korystocysta gochtii* (Sarjeant 1976) Woollam 1983
- 2263 *Korystocysta pachyderma* (Deflandre 1938) Woollam 1983
- 2264 *Luehndea spinosa* Morgenroth 1970
- 2265 *Liasidium variabile* Drugg 1978
- 2266 *Lithodinia jurassica* Eisenack 1935
- 2267 *Mancodinium semitabulatum* Morgenroth 1970
- 2268 *Meiourogonyaulax bulloidea* (Cookson & Eisenack 1960) Sarjeant 1969
- 2269 *Meiourogonyaulax caytonensis* (Sarjeant 1959) Sarjeant 1969
- 2270 *Meiourogonyaulax reticulata* Dodekova 1975
- 2271 *Meiourogonyaulax valensii* Sarjeant 1966
- 2272 *Mendicodinium groenlandicum* (Pocock & Sarjeant 1972) Davey 1979
- 2273 *Mendicodinium microscabratum* Bucefalo Palliani et al. 1997
- 2274 *Moorodinium crista* Wainman et al. 2018
- 2275 *Muderongia simplex* Alberti 1961
- 2276 *Muderongia simplex* Alberti 1961 subsp. *microperforata* Davey 1982
- 2277 *Nannoceratopsis dictyambonis* Riding 1984
- 2278 *Nannoceratopsis gracilis* Alberti 1961
- 2279 *Nannoceratopsis plegas* Drugg 1978
- 2280 *Nannoceratopsis pellucida* Deflandre 1938
- 2281 *Nannoceratopsis raunsgaardii* Poulsen 1996
- 2282 *Nannoceratopsis senex* van Helden 1977
- 2283 *Nannoceratopsis spiculata* Stover 1966
- 2284 *Neuffenia willei* Brenner & Dürr 1986
- 2285 *Noricysta fimbriata* Bujak & Fisher 1976

- 2286 *Occisucysta tentorium* Duxbury 1977
- 2287 *Palaecysta palmula* (Davey 1982) Williams & Fensome 2016
- 2288 *Pareodinia brevicornuta* Kunz 1990
- 2289 *Pareodinia ceratophora* Deflandre 1947
- 2290 *Pareodinia prolongata* Sarjeant 1959
- 2291 *Pareodinia halosa* (Filatoff 1975) Prauss 1989
- 2292 *Perisseiasphaeridium pannosum* Davey & Williams 1966
- 2293 *Phallocysta elongata* (Beju 1971) Riding 1994
- 2294 *Phallocysta eumekes* Dörhöfer & Davies 1980
- 2295 *Phoberocysta neocomica* (Gocht 1957) Millioud 1969
- 2296 *Pseudoceratium iehiense* Helby & May in Helby 1987
- 2297 *Pseudoceratium pelliferum* Gocht 1957
- 2298 *Rhaetogonyaulax arctica* (Wiggins 1973) Stover & Evitt 1978
- 2299 *Rhaetogonyaulax rhaetica* (Sarjeant 1963) Loeblich Jr. & Loeblich III 1968
- 2300 *Rhynchodiniopsis cladophora* (Deflandre 1938) Below 1981
- 2301 *Rynchodiniopsis? regalis* (Gocht 1970) Jan du Chêne et al. 1985
- 2302 *Sahulidinium ottii* Stover & Helby 1987
- 2303 *Scriniocassis weberi* Gocht 1964
- 2304 *Scriniodinium crystallinum* (Deflandre 1938) Klement 1960
- 2305 *Senoniasphaera jurassica* (Gitmez & Sarjeant 1972) Lentin & Williams 1976
- 2306 *Sentusidinium villersense* (Sarjeant 1968) Sarjeant & Stover 1978
- 2307 *Skuadinium fusum* Wainman et al. 2018
- 2308 *Suessia swabiana* Morbey 1975
- 2309 *Susadinium faustum* (Bjaerke 1980) Lentin & Williams 1985
- 2310 *Sverdrupiella baccata* Bujak & Fisher 1976
- 2311 *Sverdrupiella manicata* Bujak & Fisher 1976
- 2312 *Sverdrupiella ornaticingulata* Bujak & Fisher 1976
- 2313 *Sverdrupiella septentrionalis* Bujak & Fisher 1976
- 2314 *Sverdrupiella usitata* Bujak & Fisher 1976
- 2315 *Systematophora areolata* Klement 1960
- 2316 *Systematophora penicillata* (Ehrenberg 1843 ex Ehrenberg 1865) Sarjeant 1980
- 2317 *Tabulodinium senarium* Dodekova 1990
- 2318 *Trichodinium scarburghense* (Sarjeant 1964) Williams et al. 1993
- 2319 *Valensiella ovulum* (Deflandre 1947) Eisenack 1963

- 2320 *Wanaea acollaris* Dodekova 1975  
2321 *Wanaea digitata* Cookson & Eisenack 1958  
2322 *Wanaea thysanota* Woollam 1982

2323

2324 **Colonial alga:**

- 2325 *Palambages pariunta* Wainman et al. 2018

2326

2327 **Miospores:**

- 2328 *Ricciisporites umbonatus* Felix & Burbridge 1977

- 2329 *Shanbeipollenites proxireticulatus* Schrank 2004

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2331

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