

1 **The literature on Triassic, Jurassic and earliest Cretaceous dinoflagellate cysts:**
2 **supplement five**

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10 **ABSTRACT**

11 Since the publication of five literature compilations issued between 2012 and 2020, 63 further
12 published contributions on Triassic, Jurassic and earliest Cretaceous (Berriasian)
13 dinoflagellate cysts have been discovered, or were issued in the last 14 months (i.e. between
14 February 2019 and March 2020). These studies are on North Africa, Southern Africa, East
15 Arctic, West Arctic, east and west sub-Arctic Canada, China and Japan, East Europe, West
16 Europe, the Middle East, and sub-Arctic Russia west of the Ural Mountains, plus multi-
17 region studies and items with no geographical focus. The single-region studies are mostly
18 focused on Africa, the Arctic, Europe and the Middle East. All the 63 publications are listed
19 herein with doi numbers where applicable, and a description of each item as a string of
20 keywords.

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

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26 **1. Introduction**

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28 The literature on Triassic to earliest Cretaceous (Berriasian) dinoflagellate cysts is relatively
29 extensive, and it has been compiled and reviewed by Riding (2012, 2013, 2014, 2019a,
30 2020). These five items listed 1347, 94, 89, 266 and 93 publications respectively on this
31 topic, with each citation followed by a string of keywords detailing the scope of these 1889
32 studies. Unfortunately 11 publications were mentioned twice, hence the true cumulative total
33 of published items is 1878 (Riding 2019b). The works of Riding (2014, 2019a, 2020) were
34 substantially more interpretive than Riding (2012, 2013); the former three papers reviewed

35 and summarised the major items of literature which were listed. During the 14 months since
36 the finalisation of Riding (2020), the author has compiled 63 relevant items which were either
37 previously inadvertently overlooked or have been recently published (i.e. issued between
38 February 2019 and March 2020). The 63 contributions listed herein makes the current
39 cumulative total 1941 (Table 1).

40 These 63 articles are largely on the Jurassic of Africa, the Arctic, Europe and the
41 Middle East (Table 2), and are listed in Appendix 1 of the Supplementary data. Papers on
42 West Europe are most numerous (17), and comprise 27% of the overall total (Table 2). This
43 continues the substantial Euro-centric bias noted by Riding (2012, 2013, 2014, 2019a, 2020).
44 No single stratigraphical interval is dominant, but 19 papers are focused on, or include data
45 from, the Early Jurassic (Table 3).

46 In this compilation, more selected catalogues, contributions on suprageneric
47 classification, indexes and major taxonomic reviews are included where considered
48 stratigraphically appropriate. Examples of these are Stover and Evitt (1978), Wilson and
49 Clowes (1981) and Fensome et al. (2019a, 2019b).

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52 **2. Regional review and synthesis**

53 In this section, brief commentaries/reviews of selected articles from the 63 publications listed
54 in Appendix 1 of the Supplemental data are presented. These items are deemed particularly
55 worthy of mention, and are from nine of the 11 geographical regions relevant to this
56 contribution (Table 2). These 11 territories are North Africa, Southern Africa, East Arctic,
57 West Arctic, east and west sub-Arctic Canada, China and Japan, East Europe, West Europe,
58 the Middle East, and sub-Arctic Russia west of the Ural Mountains. Forty-nine of the 63
59 contributions in Appendix 1 of the Supplemental data is referred to one of these 11 regions;
60 the remaining 14 are assigned as either ‘multi-region’ or ‘no geographical focus’ (Tables 1,
61 2). Van de Schootbrugge et al. (2019) and Stover and Evitt (1978) are good examples of
62 ‘multi-region’ and ‘no geographical focus’ respectively. In this compilation, there are no
63 relevant single-region publications from East Africa, Central America, South America,
64 Antarctica, Southeast Asia, Australasia, the Indian subcontinent, sub-Arctic Russia east of the
65 Ural Mountains, and the U.S.A. east and west of the Rocky Mountains (Tables 1, 2). All the
66 dinoflagellate cysts, at and below species level, mentioned throughout this paper are listed in
67 Appendix 2 of the Supplemental data with full author citations. All the biozones referred to

68 herein are deemed to be have chronostratigraphical significance and the terminology used
69 reflects this.

70

71 **2.1. North Africa**

72 Four contributions on material from the continent of Africa are included in this review. Three
73 of these are on Egypt and Morocco in North Africa, and are summarised below.

74 Omran et al. (1990) is largely focussed on the palynology of the Lower Cretaceous
75 (Hauterivian/Barremian to Albian) successions penetrated by three boreholes in the northern
76 Western Desert of Egypt. However, eight dinoflagellate cyst taxa were reported from the
77 Middle and Upper Jurassic, below a substantial hiatus in the Alam el Bueib Formation
78 (Omran et al. 1990, figs 2, 9). These include the genera *Ctenidodinium*, *Escharisphaeridia*
79 and *Sentusidinium*. The presence of *Apteodinium* spp., *Cribroperidinium* spp.,
80 *Hystrichosphaerina* spp. and *Systematophora areolata* means that the Jurassic samples
81 studied are all Late Jurassic in age (e.g. Klement 1960, Riding and Thomas 1992).

82 El Atfy et al. (2019) is a detailed study of the palynology of the Alam El Bueib and
83 Alamein members within the Burg El Arab Formation from the Obaiyed Oilfield in the
84 northwest Matruh Basin, northern Western Desert, northwest Egypt. Sixty-two cuttings
85 samples were examined from two boreholes. This succession is of Early Cretaceous age and
86 ranges from Berriasian to Aptian, and the entire palynomorph spectra were thoroughly
87 documented, and these include 24 dinoflagellate cyst species. These authors recognised a
88 single dinoflagellate cyst ‘phase’, of Berriasian to Barremian age. Dinoflagellate cysts proved
89 subordinate to pollen and spores, however dinoflagellate cyst phase DI of Berriasian to
90 Barremian age was identified. This was defined as the base of the two successions studied to
91 the range top of *Tenua anaphrissa* (as *Pseudoceratium anaphrissum*) (see Costa and Davey
92 1992, fig. 3.5). Also present were *Cribroperidinium* spp., *Coronifera oceanica*,
93 *Cyclonephelium* spp., *Oligosphaeridium* spp., *Sentusidinium* spp., *Subtilisphaera* sp. and
94 *Trichodinium castanea*. The age of phase DI was discussed by El Atfy et al. (2019, p. 114);
95 the Berriasian–Barremian age was determined largely due to a correlation with a nearby
96 successions studied by El Beialy (1994) and Mahmoud and Deaf (2007).

97 The geochemistry and palynofacies of the Pliensbachian–Toarcian Event from Ait
98 Moussa and Issouka, northeast of Boulemane, in the Fès-Meknès region, Middle Atlas Basin,
99 northeast Morocco was studied by Rodrigues et al. (2020) in order to investigate
100 palaeoclimate, sea level fluctuations, sedimentology and tectonic history. The succession
101 investigated exhibits strong terrestrial affinity, and was deposited in nearshore settings. The

102 dinoflagellate cysts *Luehndea spinosa* and *Nannoceratopsis gracilis* were recorded from the
103 Pliensbachian–Toarcian transition at Ait Moussa, and were attributed to post Late
104 Pliensbachian cooling by Rodrigues et al. (2020). *Luehndea spinosa* was also encountered at
105 Issouka.

106

107 **2.2. Southern Africa**

108 Steeman et al. (2020), is a study on material from on Angola and it represents the first
109 relevant record from Southern Africa. These authors undertook a study of Paleogene
110 dinoflagellate cysts of the Landana section on the coast of Cabinda Province in Angola,
111 western Southern Africa. They noted some reworking from the underlying Mesozoic,
112 including the characteristically Middle Jurassic species *Aldorfia aldorfensis* (see, for
113 example, Gocht 1970; Wiggan et al. 2017). As mentioned above, this represents the first
114 record of Jurassic dinoflagellate cysts from the region of Southern Africa.

115

116 **2.3. East Arctic**

117 Eight contributions listed in Appendix 1 of the Supplemental data are on the East Arctic
118 region; four of these are deemed to be substantially impactful. One of them, van de
119 Schootbrugge et al. (2019), is a multi-region study. Five of the eight items are on the
120 Svalbard Archipelago in the Arctic Ocean, and four of these are described in subsection 2.3.1.
121 below. The following subsection, 2.3.2, concerns two major investigations of important
122 Lower and Upper Jurassic successions from northern Russia.

123

124 **2.3.1. The Svalbard region**

125 Koevoets et al. (2018) is a major multidisciplinary study of the Agardhfjellet Formation of
126 borehole material from central Spitsbergen, Svalbard that was drilled for the Longyearbyen
127 carbon dioxide storage project. These authors obtained two dinoflagellate cyst associations,
128 one from the uppermost Bathonian Oppdalen Member and the other from the uppermost
129 Kimmeridgian to Ryazanian (Berriasian) Oppdalssåta and Slottsmøya members (Koevoets et
130 al. 2018, figs 8, 16). Samples from the Oppdalen Member yielded *Atopodinium haromense*,
131 *Chytroeisphaeridia hyalina*, *Gonyaulacysta jurassica*, *Sirmiodinium grossii*, *Valensiella*
132 *ovulum* and *Valvaeodinium spinosum* amongst others. This succession was assigned to the
133 *Cadoceras calyx* boreal ammonite zone of late Bathonian age and Koevoets et al. (2018)
134 concluded that the dinoflagellate cysts are consistent with this assessment based on the
135 presence of *Sirmiodinium grossii* and *Valvaeodinium spinosum* (see Woollam and Riding

136 1983, Riding et al. 1985). The identification of *Atopodinium haromense* may be questionable
137 as this species is typical of the Oxfordian–Kimmeridgian transition in the Late Jurassic
138 (Thomas and Cox 1988). The Oppdalssåta and Slottsmøya members produced apparently
139 more diverse associations including *Leptodinium subtile*, *Rhynchodiniopsis cladophora*,
140 *Senoniasphaera jurassica* and *Tubotuberella apatela*, and typical of the Upper Jurassic. The
141 upper, palynologically productive, part of the Oppdalssåta Member was interpreted as being
142 late Kimmeridgian to Tithonian (early Volgian) in age. The overlying Slottsmøya Member
143 spans the lower Volgian to Berriasian (Ryazanian), which is consistent with the boreal
144 ammonites recovered. However, some reworking of dinoflagellate cysts from the
145 Kimmeridgian was noted (Koevoets et al. 2018, p. 12).

146 The occurrence of the late Pliensbachian to early Toarcian marker dinoflagellate
147 cyst *Mancodinium semitabulatum* in the Mohnhøgda Member (Svenskøya Formation) was
148 mentioned in Olausen et al. (2018, p. 48). This occurrence is coeval with a global flooding
149 event (Smelror et al. 2018). Paterson and Mangerud (2019) produced an extensively
150 illustrated revised palynomorph zonation for the Middle and Upper Triassic (Anisian–
151 Rhaetian) of the Barents Sea between Svalbard in the north, and Arctic Norway in the south.
152 Most of the biozones considered are based on spores and pollen, but the *Rhaetogonyaulax*
153 *arctica* and *Rhaetogonyaulax rhaetica* dinoflagellate cyst zones of late Carnian-early Norian
154 and early Norian age respectively were also established (Paterson and Mangerud 2019, p. 18–
155 19, fig. 3).

156 Smelror et al. (2018) is a comprehensive illustrated account of the Upper Triassic
157 to Lower Cretaceous (Norian–Aptian) palynostratigraphy of Kong Karls Land in the eastern
158 part of the Svalbard Archipelago, north of the Barents Sea in the Arctic Ocean. The material
159 comprises samples collected from seven formations in the Kapp Toscana and Adventdalen
160 groups (Smelror et al. 2018, fig. 3). The oldest material are three samples from the Flatsalen
161 Formation of Kapp Koburg, Kongsøya. The two uppermost samples yielded
162 *Rhaetogonyaulax* sp., thereby placing the Flatsalen Formation in the *Rhaetogonyaulax*
163 *rhaetica* Assemblage Zone of Paterson and Mangerud (2015) of early Norian age. This is
164 consistent with several other studies in the Barents Sea area.

165 The Svenskøya Formation (Norian–?Rhaetian to Toarcian) comprises two
166 members and is thought to include several hiatuses. The lower unit, the Sjøgrenfjellet
167 Member, is devoid of dinoflagellate cysts and is of Norian–?Rhaetian to Hettangian–early
168 Pliensbachian age based on pollen and spores. The overlying Moenhøgda Member yielded
169 abundant palynomorphs including the dinoflagellate cysts *Luehndea spinosa*, *Mancodinium*

170 *semitabulatum*, *Nannoceratopsis gracilis*, *Pareodinia halosa* (as *Caddasphaera halosa*) and
171 *Phallocysta* sp. This association indicates a late Pliensbachian to early Toarcian age,
172 correlating with the DSJ6 and DSJ7 dinoflagellate cyst zones of Poulsen and Riding (2003).

173 The Kongsøya Formation was interpreted as being of late Toarcian to Aalenian in
174 age by Smelror et al. (2018). This unit produced *Nannoceratopsis gracilis*, *Nannoceratopsis*
175 spp., *Ovalicysta hiata*, *Parvocysta* spp., *Phallocysta eumekes*, *Scriniocassis priscus* (as
176 *Eyachia prisca*), *Scriniocassis weberi* and *Susadinium scrofoides*. The co-occurrences of
177 *Ovalicysta hiata*, *Phallocysta eumekes* and *Susadinium scrofoides* allows a correlation to the
178 latest Toarcian to earliest Aalenian DSJ10 dinoflagellate cyst zone of Poulsen and Riding
179 (2003). The Flatsalen, Svenskøya and Kongsøya formations are largely conformable.
180 However there is a major hiatus above the Kongsøya Formation and there are no Bajocian
181 strata preserved on Kong Karls Land (Smelror et al. 2018, fig. 17).

182 The lowermost unit of the Agardhfjellet Formation, the Oppdalen Member has a
183 transgressive base. The latter unit generally yielded common dinoflagellate cysts. These
184 include *Arkellea teichophera* (as *Heslertonia teichophera*), *Chytroeisphaeridia cerastes*,
185 *Chytroeisphaeridia hyalina*, *Ctenidodinium continuum*, *Ctenidodinium ornatum*,
186 *Endoscrinium galeritum*, *Nannoceratopsis pellucida*, the *Paragonyaulacysta* group,
187 *Rhynchodiniopsis cladophora* and *Sirmiodinium grossii*. On the basis of the aforementioned
188 taxa, together with key ammonites, the Oppdalen Member was assigned to the upper
189 Bathonian to middle Callovian (Smelror et al. 2018, fig. 13). The overlying Lardyfjellet
190 Member of the Agardhfjellet Formation produced relatively diverse dinoflagellate cyst
191 associations. These include the marker species *Evansia deflandrei* (as *Crussolia deflandrei*),
192 *Gonyaulacysta eisenackii*, *Gonyaulacysta jurassica* subsp. *adepta* var. *longicornis*, *Kalyptea*
193 *diceras*, *Scriniodinium crystallinum*, *Stephanelytron redcliffense*, *Trichodinium*
194 *scarburghense*, *Wanaea fimbriata* and *Wanaea thysanota*. This association is indicative of
195 the late Callovian to early Oxfordian interval (Poulsen and Riding 2003). However, evidence
196 from ammonites and foraminifera indicates that the upper part of the Lardyfjellet Member is
197 early Kimmeridgian in age. There is a substantial hiatus above the Agardhfjellet Formation
198 (Smelror et al. 2018, fig. 17).

199 The overlying strata, the Klippfisk, Kolje and Helvetiafjellet formations are Early
200 Cretaceous (Valanginian–Aptian) in age. This study has allowed reliable correlations to
201 Triassic to Cretaceous successions in Franz Josef Land, Arctic Russia (Smelror et al. 2018,
202 fig. 17).

203

204 2.3.2. Northern Russia

205 The foraminifera and palynomorphs from samples collected from two sections of the Upper
206 Jurassic (Oxfordian–Tithonian [Volgian]) on the banks of the Lopsiya River immediately east
207 of the northern Ural Mountains in north-central Russia were studied by Lebedeva et al.
208 (2019). This is an extremely important reference section largely due to its stratigraphical
209 completeness, and the presence throughout of zonal ammonites and other molluscs. Lebedeva
210 et al. (2019, figs 4, 5) reported relatively diverse marine and terrestrial palynomorphs. The
211 dinoflagellate cyst associations are dominated by non-tabulate, proximate forms with apical
212 archaeopyles referred to the *Sentusidinium-Batiacasphaera-Kallosphaeridium* group (Wood
213 et al. 2016), and chorate taxa are relatively rare. The latter phenomenon is typical of the
214 Boreal Realm (Wierzbowski et al. 2002). However, the assemblages also include typically
215 Late Jurassic dinoflagellate cysts including *Ambonosphaera? staffinensis*, *Cribroperidinium*
216 *globatum*, *Cribroperidinium? longicorne*, *Dingodinium* spp., *Endoscrinium luridum*,
217 *Glossodinium dimorphum*, *Leptodinium* spp., *Scriniodinium crystallinum*, *Senoniasphaera*
218 *jurassica*, *Systematophora areolata* and *Tubotuberella apatela*.

219 Two dinoflagellate cyst biozones were described by Lebedeva et al. (2019). These
220 are the *Gonyaulacysta jurassica* subsp. *jurassica* and *Corculodinium inaffectum* assemblage
221 zones of early Kimmeridgian, and latest early Kimmeridgian to earliest Tithonian (Volgian)
222 age respectively. The *Gonyaulacysta jurassica* subsp. *jurassica* assemblage zone is
223 equivalent to the early Kimmeridgian *Eurorasenia pseudoouralensis* ammonite subzone of
224 the *Rasenia evoluta* ammonite zone. Lebedeva et al. (2019, p. 9) stated that the Lopsiya River
225 material is closely comparable with the dinoflagellate cysts from around the Oxfordian–
226 Kimmeridgian transition of northwest Europe and adjacent areas and the central Russian
227 Platform (e.g. Riding and Thomas 1988, 1997; Riding et al. 1999). By contrast, it is markedly
228 different from coeval material from central northern Siberia (e.g. Ilyina et al. 2005).
229 Lebedeva et al. (2019) deemed the dinoflagellate cyst assemblages from Lopsiya River to be
230 intermediate in floral character between coeval material from Subboreal northwestern Europe
231 and the Boreal Realm. The overlying *Corculodinium inaffectum* assemblage zone ranges
232 from the *Zonovia ulalensis* ammonite subzone of the *Rasenia evoluta* ammonite zone to the
233 *Eosphinctoceras magnum* ammonite zone, and is latest early Kimmeridgian to earliest
234 Tithonian (Volgian) in age (Lebedeva et al. 2019, fig. 4). The base of this biozone was
235 formally identified as the range base of *Corculodinium inaffectum*.

236 The same succession of dinoflagellate cyst biozones were established in western
237 Russia by Riding et al. (1999). A major difference in Lebedeva et al. (2019) is that the

238 inception of *Corculodinium inaffectum* is substantially older than documented by Riding et
239 al. (1999), i.e. immediately below the *Aulacostephanus mutabilis* ammonite zone. Riding et
240 al. (1999) placed the range base of *Corculodinium inaffectum* (as *Subtilisphaera? inaffecta*) at
241 the base of the *Aulacostephanus autissiodorensis* ammonite zone. As Lebedeva et al. (2019)
242 pointed out, there is substantial congruence between their data with dinoflagellate cyst ranges
243 established in northern Europe and adjacent regions. For example, the range bases of
244 *Corculodinium inaffectum* and *Cribroperidinium? longicorne*, and the apparent extinctions of
245 *Endoscrinium luridum* and *Gonyaulacysta jurassica* subsp. *jurassica* are extremely similar in
246 both areas in terms of their calibration with the ammonite zonations.

247 Two successions spanning the Kyra and Kelimyar formations of late Pliensbachian
248 and Toarcian age from two exposures near the Kelimyar River in northern Siberia were
249 studied by van de Schootbrugge et al. (2019) as part of a major multi-region study comparing
250 the Arctic with sub-Arctic West Europe. These were sections S16 and S5-D1, and the former
251 comprises Upper Pliensbachian to Upper Toarcian strata. Most of the Upper Pliensbachian
252 Kyra Formation yielded sparse dinoflagellate cyst associations dominated by the genus
253 *Nannoceratopsis*. By contrast, the uppermost Kyra Formation, and the overlying Kelimyar
254 Formation of Toarcian age produced much more species-rich dinoflagellate cyst palynofloras.
255 *Batiacasphaera* sp., *Dissiliodinium* sp., *Maturodinium inornatum*, *Pareodinia?*
256 *pseudochytroeides* (as *Dodekovia pseudochytroeides*), *Parvocysta* spp., *Phallocysta eumekes*,
257 *Scriniocassis weberi*, *Susadinium scrofoides*, *Valvaeodinium koessenium* (as *Comparodinium*
258 *koessenium*), *Valvaeodinium* spp. and *Walloodinium cylindricum* were recorded from this
259 succession, together with the acritarch *Limbicysta bjaerkei* (see van de Schootbrugge et al.
260 2019, fig. 7). Biostratigraphically, the most notable aspect of this succession is that the
261 *Parvocysta-Phallocysta* suite has a substantially younger inception, i.e. earliest Toarcian,
262 than further south in western Europe (e.g. Riding 1984, Riding et al. 1991).

263 The transition between the Kyra and Kelimyar formations (uppermost
264 Pliensbachian–Lower Toarcian) was sampled in the Kelimyar River S5-D1 section by van de
265 Schootbrugge et al. (2019). At the onset of the Early Toarcian negative Carbon Isotope
266 Excursion, there is a dramatic increase in dinoflagellate cyst diversity. *Mancodinium*
267 *semitabulatum*, *Moesiodinium raileanui*, *Nannoceratopsis* spp., *Parvocysta* spp., *Phallocysta*
268 spp., *Susadinium scrofoides* and *Valvaeodinium* spp. all appeared at this time (van de
269 Schootbrugge et al. 2019, fig. 9). This succession proves that the range base of the
270 *Parvocysta-Phallocysta* suite is early Toarcian in age in the high palaeolatitudes.

271 The data from the two Kelimyar River sections examined by van de Schootbrugge
272 et al. (2019) disproves the contention of, for example Riding et al. (1999, fig. 11) that the
273 *Parvocysta-Phallocysta* suite emerged in the high northerly palaeolatitudes during the Late
274 Toarcian. By contrast, this group emerged during the early Toarcian, and thrived during the
275 Toarcian Oceanic Anoxic Event (T-OAE) in northern Siberia. As the *Parvocysta-Phallocysta*
276 suite co-occurred with early representatives of the Gonyaulacales during the Early Toarcian,
277 the high northerly latitudes appear to represent the cradle of dinoflagellate evolution at this
278 critical interval in plankton evolution (Wiggin et al. 2018). The abundance of dinoflagellate
279 cysts in northern Siberia during the T-OAE is believed to be as a result of only sporadic
280 benthic anoxia due to seasonally-driven marine mixing. Further south, there was a virtual
281 blackout of dinoflagellate cysts during the early Toarcian (Correia et al. 2017). Furthermore,
282 the *Parvocysta-Phallocysta* suite migrated into Europe in southerly-moving currents through
283 the Viking Corridor after oceanic deepening during the middle part of the early Toarcian (van
284 de Schootbrugge et al. 2019, fig. 11). These authors suggested that this enhanced Arctic-
285 Tethys marine connectivity, specifically the influx of cold, low-salinity, nutrient-rich waters
286 from the Arctic region helped to end the T-OAE. These conclusions are supported by the fact
287 that the late Pliensbachian and Toarcian ammonite zonal schemes are substantially different
288 in the Arctic, Suboreal and Tethyan regions, indicating intense provincialism at this time (van
289 de Schootbrugge et al. 2019, fig. 1).

290

291 **2.4. West Arctic**

292 In this review there are three items relevant to the West Arctic region. These are one on
293 Arctic Canada, one on northeast Greenland and there is one multi-region contribution
294 (Appendix 1 of the Supplemental data). An abstract on the dinoflagellate cysts from the
295 Upper Jurassic to Lower Cretaceous (Oxfordian–Valanginian) succession of the Rollrock
296 section on northern Ellesmere Island in the Sverdrup Basin of Arctic Canada was issued by
297 Ingrams (2019). This succession is an important high latitude reference section for the
298 Jurassic–Cretaceous transition. Seven biozones were distinguished, defined by the range bases
299 and tops of marker taxa such as *Muderongia simplex* and *Oligosphaeridium complex*.
300 Glacioeustasy is thought to influence spine-bearing dinoflagellate cyst morphology with
301 major fluctuations in proximochorate forms reflecting relative sea level fall.

302 A major paper on the Cretaceous palynostratigraphy of northeast Greenland
303 between Traill Ø in the south and Store Koldeway in the north was published by Nøhr-
304 Hansen et al. (2019). The interval considered was latest Jurassic to Late Cretaceous

305 (Tithonian–Maastrichtian) in age and the biozonation, which comprises 15 zones, was
306 calibrated to an updated ammonite zonation and based on three boreholes and over 100
307 outcrop sections. It is the first palynozonation for the entire Cretaceous of East Greenland,
308 and can be correlated to other areas in the Arctic region. The *Gochteodinia villosa villosa*
309 (NEG Cr 1) and *Oligosphaeridium complex* (NEG Cr 2) zones cover the late Tithonian to
310 earliest Hauterivian interval. The base of the former was defined as the inceptions of
311 *Gochteodinia villosa villosa* and *Isthmocystis distincta* in the upper Tithonian. Bioevents in
312 the *Gochteodinia villosa villosa* zone include the range base of *Scriniodinium pharo*, the
313 ranges of *Lagenorhytis delicatula* and *Rotosphaeropsis thule* and the range top of
314 *Paragonyaulacysta? borealis* in the Berriasian of the Rødryggen-1 core 517001. The base of
315 the succeeding *Oligosphaeridium complex* zone was drawn in the uppermost Berriasian at the
316 range base of the index species (Nøhr-Hansen et al. 2019, fig. 7).

317 Krencker et al. (2019) is a contribution based largely on geochemistry and
318 sedimentology which posited a temporally short, high amplitude global forced regression,
319 due to polar ice sheet dynamics, which immediately preceded the major marine transgression
320 associated with the Toarcian Oceanic Anoxic Event (T-OAE). It suggests that this, and other,
321 hyperthermal events may have had their origins in short-lived ‘cold snaps’. This study was
322 based on data and material from the Central High Atlas Basin in Morocco and Jameson Land,
323 East Greenland. The palynology data in Krencker et al. (2019) is entirely from the uppermost
324 Pliensbachian, Toarcian and lowermost Aalenian strata within the Neill Klintner Group of
325 Jameson Land Basin in East Greenland. The samples used were originally collected from the
326 Gule Horn to Sortehat formations for the study of Koppelhus and Dam (2003). The material
327 yielded the dinoflagellate cysts *Luehndea spinosa*, *Mancodinium semitabulatum*,
328 *Nannoceratopsis gracilis*, *Phallocysta elongata* (as *Parvocysta elongata*), *Parvocysta* sp.,
329 *Phallocysta eumekes* and *Valvaeodinium armatum* (see Krencker et al. 2019, p. 6, 7; fig. 5).

330

331

332 2.5. *China and Japan*

333 Two contributions on the marine palynology of the Jurassic of China and Japan were issued
334 during the period of this review. Lin and Li (2019, fig. 4E) illustrated ‘?Dinoflagellate cyst’
335 from the Lower Cretaceous Duoni Formation of Wadga coal mine, near Baxoi, Qinghai-
336 Xizang Plateau, western China. This highly thermally mature specimen has a substantial
337 opening that may be an archaeopyle. However the lack of other microplankton, the

338 resemblance to certain Mesozoic gymnospermous pollen such as *Perinopollenites* and the
339 poor preservation strongly suggests it is not of dinoflagellate affinity.

340 Kemp et al. (2019) is the first report of Jurassic dinoflagellate cysts from Japan.
341 This paper is an integrated study on the isotope geochemistry, palynofacies and palynology of
342 a highly expanded succession through the Toarcian Oceanic Anoxic Event (T-OAE) in
343 southwest Japan. Palynomorphs were extracted from 32 samples of the Nishinakayama
344 Formation collected from the Sakuraguchi-dani stream section near Toyota Town. The
345 palynoflora is of relatively low diversity and two samples apparently yielded the
346 dinoflagellate cyst *Luehndea spinosa* (see Kemp et al. 2019, fig. 4). The two samples precede
347 the T-OAE, and this scenario is consistent with the results of Correia et al. (2017, fig. 3). The
348 latter study found that *Luehndea spinosa* is highly characteristic of the pre T-OAE succession
349 in the Lusitanian Basin in Portugal. Due to the intense tectonism which has affected Japan,
350 the Nishinakayama Formation is highly thermally altered and substantially overmature. This
351 is confirmed by the extremely poor preservation of the palynomorphs extracted by Kemp et
352 al. (2019, fig. 3). They are intensely blackened and degraded such that identification to
353 species level is highly problematical. This includes the photograph of *Luehndea spinosa* (see
354 Kemp et al. 2019, fig. 3T). This specimen is a poorly-preserved subangular polygonal body
355 approximately 40 µm in diameter and bearing irregular spines. The epicystal archaeopyle,
356 gonal spines and gonyaulacacean tabulation characteristic of *Luehndea spinosa* are not
357 evident (Morgenroth 1970), and the validity of the identification of this specimen is therefore
358 not considered to be secure.

359

360 **2.6. East Europe**

361 In this compilation, eight items concerning East Europe were listed in Appendix 1 of the
362 Supplemental data; these are studies from the Czech Republic, Poland and Ukraine. Four of
363 these items, Birkenmajer and Gedl (2019), Skupien and Doupovcova (2019), Svobodová et
364 al. (2019) and Kowal-Kasprzyk et al. (2020), have substantial contributions on dinoflagellate
365 cysts.

366 The study of Birkenmajer and Gedl (2019) investigated the Jurassic to Paleogene
367 dinoflagellate cyst biostratigraphy of borehole PD-9 drilled at Szczawnica in central southern
368 Poland. This well was drilled in the intensely tectonised northern boundary fault zone of the
369 Pieniny Klippen Belt in the West Carpathians. Specifically, this borehole indicates that the
370 Grajcarek Main Dislocation is virtually vertical and separates the Magura Nappe of the Outer
371 Carpathians to the north, and the Pieniny Klippen Belt to the south. The authors reported

372 dinoflagellate cyst assemblages from the Lower–Middle Jurassic, Upper Cretaceous and
373 Eocene.

374 A steeply-dipping thrust sheet of the Szlachtowa Formation of Jurassic age was
375 identified. This is the oldest unit of the Grajcarek Unit and two samples were collected at
376 716.4–710.4 m and 710.4–707.1 m (Birkenmajer and Gedl 2019, fig. 4, table 1). The
377 lowermost sample at 716.4–710.4 m yielded a low diversity dinoflagellate cyst association. It
378 is dominated by *Nannoceratopsis gracilis* and *Phallocysta elongata*, and some Eocene
379 contaminants are also present. The occurrence of the latter species, together with the absence
380 of *Dissiliodinium*, is indicative of a latest Toarcian to Aalenian age (Feist-Burkhardt 1990,
381 Riding 1994). By contrast, the uppermost sample at 710.4–707.1 m produced a relatively
382 abundant assemblage, which lacks contamination, and is overwhelmingly dominated by
383 *Nannoceratopsis dictyambonis*. Also present, but in lower proportions, are *Batiacasphaera*
384 sp., *Dissiliodinium* sp., *Kallosphaeridium?* sp., *Nannoceratopsis gracilis*, *Nannoceratopsis*
385 *raunsgaardii*, *Nannoceratopsis spiculata*, *Nannoceratopsis* sp. and *Sentusidinium*
386 *explanatum* (as *Kallosphaeridium praussii*) (Birkenmajer and Gedl 2019, table 1). The
387 authors used *Dissiliodinium* and *Nannoceratopsis dictyambonis* to interpret a latest Aalenian
388 age for the sample at 710.4–707.1 m. The overlapping ranges of this species and genus is
389 indicative of the latest Aalenian interval (Birkenmajer and Gedl 2019, p. 247). Furthermore,
390 the absence of *Dissiliodinium giganteum* provides substantial negative evidence that this
391 sample is not Bajocian in age (e.g. Gedl 2008; Segit et al. 2015).

392 Skupien and Doupovcova (2019) is of substantial regional significance because the
393 succession examined is one of the few localities in the Tethyan Realm where the Jurassic–
394 Cretaceous transition is suitable for palynological study. These authors undertook
395 biostratigraphical research on the calcareous and organic dinoflagellate cysts, and
396 calpionellids of the Vendryně Formation and Těšín Limestone (Tithonian and Beriasian
397 respectively) at Bruzovice, Outer Western Carpathians in the eastern Czech Republic. These
398 lower Tithonian and Beriasian strata were sampled and several biostratigraphically significant
399 dinoflagellate cyst taxa recovered. These include *Amphorulacysta metaelliptica* (as
400 *Amphorula metaelliptica*), *Diacanthum hollisteri*, *Dichadogonyaulax bensonii*, *Glossodinium*
401 *dimorphum*, *Muderongia longicorna*, *Phoberocysta tabulata* (as *Muderongia tabulata*),
402 *Prolixosphaeridium anasillum*, *Spiculodinium neptuni* (as *Achomosphaera neptuni*) and
403 *Spiniferites* sp. S. cf. *ramosus* (see Skupien and Doupovcova 2019, fig. 6). The
404 biostratigraphy was discussed in detail, and a very extensive set of photographs was
405 presented (Skupien and Doupovcova 2019, p. 221, 226 and figs 7–13 respectively). The

406 Jurassic–Cretaceous transition was established to occur between samples Br 12 and Br 10
407 (Skupien and Doupovcova 2019, fig. 6). Some reworking from the Pliensbachian to Bajocian
408 was noted; *Nannoceratopsis gracilis* and *Nannoceratopsis raunsgaardii* were encountered in
409 the lowermost Cretaceous Těšín Limestone (Skupien and Doupovcova 2019, figs 11F, G).

410 Svobodová et al. (2019) examined the micropalaeontological biostratigraphy and
411 palaeocological analysis of the Kurovice Limestone from Kurovice Quarry in southeast
412 Czech Republic as part of a larger project to determine a Global Stratotype Section and
413 Point (GSSP) for the Berriasian. This study includes analysis of the entire palynoflora and
414 integrated all results with magnetostratigraphy. A total of 24 samples were examined for
415 palynomorphs, and seven of these produced relatively rare and often poorly-preserved
416 material due largely to the organic-lean nature of the succession (Svobodová et al. 2019, p.
417 166, 168, figs 13–16). Because of this, the majority of the biostratigraphical conclusions are
418 based on the calcareous microfossils. However, bioevents such as the range tops of
419 *Amphorulacysta? dodekovae* (as *Amphorula dodekovae*) and *Glossodinium dimorphum*, and
420 the range bases of, for example, *Amphorulacysta metaelliptica* (as *Amphorula metaelliptica*),
421 *Dichadogonyaulax bensonii*, *Dingodinium tuberosum* (as *Dingodinium 'tuberculosis'*),
422 *Scriniodinium campanula*, *Spiculodinium neptuni* (as *Achomosphaera neptuni*) and
423 *Tehamadinium evittii* proved to be stratigraphically useful. Some reworking was discerned.

424 Kowal-Kasprzyk et al. (2020) studied the dinoflagellate cysts (calcareous and
425 organic) and foraminifera of exotic clasts of Upper Jurassic (Oxfordian–Kimmeridgian) strata
426 from southern Poland which have been reworked into an extensive Lower Cretaceous to
427 Eocene succession. These allochthonous fragments of sedimentary rocks deposited in shelfal
428 settings are proxies for the understanding of the palaeogeography of this region prior to the
429 development of the Outer Carpathian flysch basins. The clasts are from three carbonate facies
430 types. Key marker organic dinoflagellate cyst species recognised include *Endoscrinium*
431 *luridum*, *Glossodinium dimorphum*, *Gonyaulacysta jurassica*, *Leptodinium subtile* and
432 *Rhynchodiniopsis cladophora* (see Kowal-Kasprzyk et al. 2020, figs 8, 9).

433

434 **2.7. Sub-Arctic West Europe**

435 Seventeen contributions solely on the Triassic, Jurassic and lowermost Cretaceous
436 successions of sub-Arctic West Europe are covered in this review, one of which is deemed
437 especially significant (Appendix 1 of the Supplementary data). Of these 17 single-region
438 items, seven are briefly outlined below, and one highly impactful multi-region publication is
439 described.

440 Adloff and Doubinger (1982) recorded *Dapcodinium priscum* and
441 *Rhaetogonyaulax rhaetica* from the Rhaetian and lowermost Hettangian strata of Mersch,
442 central Luxembourg. Similarly, Hillebrandt et al. (2013) recorded *Dapcodinium priscum* and
443 *Rhaetogonyaulax rhaetica* from the Rhaetian of the Kuhjoch Pass in the Karwendel
444 Mountains, western Austria, although abundance, sample and range data are lacking.

445 The palaeontology of the lowermost Cretaceous (Berriasian and Valanginian) strata
446 of central Austria was studied by Boorová et al. (2015). This study is centered on the
447 Schrambach Formation at its type locality and was multidisciplinary, encompassing
448 ammonites, calpionellids and calcareous dinoflagellate cysts. The Oberalm, Schrambach and
449 Rossfeld formations yielded organic-walled dinoflagellate cysts (Boorová et al. 2015, p. 106–
450 107, figs 3A–3F, 7–8, table 1). Biostratigraphically significant taxa recorded by these authors
451 include *Amphorulacysta metaelliptica* (as *Amphorula metaelliptica*), *Ctenidodinium*
452 *elegantulum*, *Cribroperidinium? edwardsii*, *Dichadogonyaulax bensonii*,
453 *Kleithriasphaeridium corrugatum*, *Kleithriasphaeridium fasciatum*, *Phoberocysta*
454 *neocomica*, *Pseudoceratium pelliferum*, *Scriniodinium campanula*, *Spiculodinium neptuni* (as
455 *Achomosphaera neptuni*), *Spiniferites ramosus* and *Stanfordella? cretacea*. Some reworking
456 of specimens of *Nannoceratopsis*, including *Nannoceratopsis gracilis*, from the underlying
457 Lower–Middle Jurassic (Pliensbachian–Bajocian) was observed in the Oberalm and
458 Schrambach formations (Boorová et al. 2015, figs 7P, 7Q).

459 The palynology and sedimentology of the Rannoch Formation (Brent Group) in the
460 northern North Sea was studied by Slater et al. (2017). These authors reported the presence of
461 the dinoflagellate cyst genera *Evansia*, *Kallosphaeridium*, *Mancodinium*, *Nannoceratopsis*,
462 *Pareodinia* and *Phallocysta* in bioturbated sandy facies of three wells in block 211/14. This
463 association, together with rare *Botryococcus*, is indicative of shallow marine conditions
464 within the Rannoch Formation which is late Aalenian–early Bajocian in age (Richards et al.
465 1993).

466 Schobben et al. (2019) undertook a multidisciplinary study of the uppermost
467 Triassic and lowermost Jurassic (Rhaetian–Hettangian) strata of central Europe in order to
468 better understand the end-Triassic mass extinction. These authors recorded *Dapcodinium*
469 *priscum*, *Rhaetogonyaulax rhaetica* and *Suessia swabiana* from the Rhaetian and Hettangian
470 succession at a quarry northwest of Bonenburg in central Germany. *Suessia swabiana* was
471 confined to the Rhaetian, but *Dapcodinium priscum* and *Rhaetogonyaulax rhaetica* were
472 recorded throughout (Schobben et al. 2019, fig. 2).

473 The vegetational response to the Toarcian Oceanic Anoxic Event (T-OAE) in
474 northern England was investigated by Slater et al. (2019). Despite the focus on the terrestrial
475 realm, these authors discussed the dynamics of marine phytoplankton and illustrated the
476 dominance of sphaeromorphs, together with abundant amorphous organic material with much
477 reduced numbers of dinoflagellate cysts during the T-OAE, which is characterised by a
478 marked negative carbon isotope excursion (Slater et al. 2019, fig. 2).

479 A major multi-region study on the Lower Jurassic (Pliensbachian and Toarcian) of
480 the East Arctic and West Europe was published recently by van de Schootbrugge et al.
481 (2019). These authors worked on the Cleveland Basin in northern England and the
482 Norwegian North Sea. In the Cleveland Basin, van de Schootbrugge et al. (2019, fig. 5)
483 examined productive samples from the Cleveland Ironstone and Whitby Mudstone
484 formations (Upper Pliensbachian to Upper Toarcian). In broad terms, the floras recorded by
485 van de Schootbrugge et al. (2019) are complementary to, and consistent with, the
486 assemblages documented by Riding (1984) and Bucefalo Palliani and Riding (2000) from this
487 depocentre. The oldest dinoflagellate cyst species recorded by van de Schootbrugge et al.
488 (2019) was *Luehndea spinosa* in the *Amaltheus margaritatus* ammonite zone of the Upper
489 Pliensbachian and this was followed by a substantial influx of taxa at the Pliensbachian–
490 Toarcian transition. These include the inceptions of *Mancodinium semitabulatum*,
491 *Maturodinium inornatum*, *Nannoceratopsis gracilis*, *Nannoceratopsis senex* and
492 *Scriniocassis weberi*. There is a marked decrease in dinoflagellate cysts, but not a total
493 blackout, during the Carbon Isotope Excursion (CIE) interval at the base of the *Harpoceras*
494 *falciferum* ammonite zone in the T-OAE. The *Parvocysta-Phallocysta* suite are first observed
495 in the *Harpoceras falciferum*–*Hildoceras bifrons* ammonite zone transition, after the T-OAE.
496 Of this major plexus of forms, *Parvocysta bullula*, *Phallocysta eumekes* and *Susadinium*
497 *scrofoides* were recorded by van de Schootbrugge et al. (2019, figs 5, 12). The subsequent
498 diversity only increased marginally up-section, with *Scriniocassis priscus* appearing in the
499 middle part of the *Hildoceras bifrons* ammonite zone (van de Schootbrugge et al. (2019, fig.
500 5). Taken together, Riding (1984), Bucefalo Palliani and Riding (2000) and van de
501 Schootbrugge et al. (2019) provide an excellent composite palynological reference section for
502 the Sinemurian to the Aalenian of the Cleveland Basin.

503 The uppermost Pliensbachian and Toarcian succession from the Norwegian North
504 Sea, specifically well 34/10-35 in the Gulfaks South oilfield, was studied by van de
505 Schootbrugge et al. (2019, fig. 6) as part of an investigation of the Arctic and Europe. This is
506 the only succession in this study that is not calibrated to the ammonite zonation. It appears

507 that, in general terms, this North Sea record is similar to coeval floras from northern England,
508 but nonetheless and intermediate between northern Siberia and northwest Europe. *Luehndea*
509 *spinosa* ranges slightly stratigraphically higher (to the end of the CIE in the T-OAE) than in
510 southern Europe and Tethys, i.e. into the *Harpoceras falciferum* ammonite zone equivalent
511 based on chemostratigraphy (van de Schootbrugge et al. 2019, fig. 12). There is no virtual
512 blackout of dinoflagellate cysts in the T-OAE, as is the case further south in Europe, in the
513 Norwegian North Sea (e.g. Correia et al. 2017). Early representatives of the Gonyaulacales
514 such as the genera *Batiacasphaera* and *Dissiliodinium* are present in the upper Pliensbachian,
515 and throughout the T-OAE of well 34/10-35. This is similar to the records from northern
516 Siberia (van de Schootbrugge et al. 2019, figs 7, 9). Potentially most significantly, in well
517 34/10-35 is the range bases of *Parvocysta bullula* and *Parvocysta nasuta* within the T-OAE,
518 i.e. within the negative CIE (lowermost Toarcian). These species are typical of the
519 *Parvocysta-Phalloecysta* suite, and this inception is similar to the situation in northern Siberia
520 (see section 2.2; van de Schootbrugge et al. 2019, figs 6, 7, 9, 12).

521 Hesselbo et al. (2020) is a follow-up paper to Riding et al. (2013). The latter is an
522 account of acmes of the dinoflagellate cyst *Liasidium variabile* and the pollen species
523 *Classopollis classoides*, together with a marked negative CIE of 2–3‰ in the upper
524 Sinemurian strata of Lincolnshire, eastern England. These phenomena were collectively
525 termed the S-CIE and interpreted as a hyperthermal event of global extent. Hesselbo et al.
526 (2020) sampled the shallow marine Sinemurian succession at Robin Hood's Bay in the
527 Cleveland Basin, North Yorkshire, northern England at a high resolution. These authors
528 confirmed the presence of the S-CIE (and renamed it the Liasidium Event), which
529 corresponds very closely to the *Oxynoticeras oxynotum* ammonite zone. The Liasidium Event
530 at Robin Hood's Bay also is coeval with a negative CIE that exhibits a distinctive double
531 spike in the middle part of the *Oxynoticeras oxynotum* ammonite zone (Hesselbo et al. 2020,
532 fig. 3). The peak occurrences of *Liasidium variabile* correspond to deep water and maximum
533 flooding. Analysis of parasequences in this succession allow an age assessment of at least one
534 million years for the Liasidium Event. The intensity of this relatively minor hyperthermal is
535 far less than the subsequent T-OAE, and no evidence of significant bottom water
536 deoxygenation was developed. This study established a chronostratigraphical range for
537 *Liasidium variabile* at Robin Hood's Bay as middle late Sinemurian. Specifically this is the
538 base of the *Eparietites denotatus* ammonite subzone of the *Asteroceras obtusum* ammonite
539 zone, to close to the top of the *Oxynoticeras oxynotum* ammonite subzone of the
540 *Oxynoticeras oxynotum* ammonite zone. However, *Liasidium variabile* is only consistent and

541 common (i.e. >5 %) in the *Oxynoticeras oxynotum* ammonite zone, from the middle of the
542 *Oxynoticeras simpsoni* ammonite subzone to the base of the *Oxynoticeras oxynotum*
543 ammonite subzone (Hesselbo et al. 2020, fig. 3).

544

545 2.8. *The Middle East*

546 In this review, there are six contributions which are focused exclusively on the Middle East,
547 two of which are considered especially impactful (Appendix 1 of the Supplementary data).
548 Five of these six items are on the Lower and Middle Jurassic (Pliensbachian to Callovian)
549 successions of northern Iran. The material documented in these five articles is dominated by
550 pollen and spores, and all the palynomorphs are blackened and poorly-preserved due to
551 substantial levels of thermal alteration.

552 Four of the items on Iran were authored or co-authored by Fatemeh Vaez-Javadi,
553 and three of these are centered on northeast Iran. The first of these was Vaez-Javadi et al.
554 (2003), a study of the marine palynomorphs in six samples collected from the Shemshak
555 Formation of Jajarm County, northeast Iran. The material is highly blackened, and includes
556 nine species of dinoflagellate cysts and two acritarchs. Two zones, the *Nannoceratopsis*
557 *spiculata* and *Valensiella ovulum* biozones were established, and are of Pliensbachian–
558 Toarcian and Bajocian age respectively (Vaez-Javadi et al. 2003, fig. 2, pl. 1, 2). The
559 *Nannoceratopsis spiculata* biozone yielded four taxa; these are *Kalyptea diceras*, *Liesbergia*
560 *liesbergensis*, *Nannoceratopsis spiculata* and *Scriniodium? dictyophorum* (as ‘*Aldorfia*
561 *dictyophora*’) (see Vaez-Javadi et al. 2003, fig. 2). *Nannoceratopsis spiculata* does not
562 normally occur in the Pliensbachian–Toarcian interval with younger gonyaulacacean taxa
563 such as *Liesbergia liesbergensis* (see Berger 1986). In the succeeding *Valensiella ovulum*
564 biozone (Bajocian), a more diverse flora was recorded. However, as in the *Nannoceratopsis*
565 *spiculata* biozone, some species such as *Gonyaulacysta centriconnata* appear to be
566 stratigraphically anomalous (Riding 1983).

567 Vaez-Javadi (2018, 2019) are both on the palynology of the Middle Jurassic
568 (Aalenian–Bajocian) Hojedk Formation of the Tabas Block in northeast Iran. In a substantial
569 paper, Vaez-Javadi (2018, fig. 2) reported a moderately diverse dinoflagellate cyst
570 association dominated by the genera *Kalyptea*, *Nannoceratopsis* and *Pareodinia*. This
571 assemblage was assigned to the *Nannoceratopsis triceras-Pareodinia ceratophora*
572 assemblage zone. The presence of species such as *Nannoceratopsis gracilis*, *Nannoceratopsis*
573 *symmetrica*, *Nannoceratopsis triceras* and *Pareodinia ceratophora* is consistent with an
574 Aalenian–Bajocian age (Bucefalo Palliani and Riding 2000, 2003). Vaez-Javadi (2019) is a

575 report on 38 samples from the Hojedk Formation of the Chahrehkneh borehole, southwest of
576 Tabas, in South Khorasan Province, northeast Iran. A less diverse dinoflagellate cyst
577 assemblage was recovered than in Vaez-Javadi (2018), but it also comprised the genera
578 *Kalyptea*, *Nannoceratopsis* and *Pareodinia*. The *Nannoceratopsis* sp. cf. *N. gracilis* interval
579 zone was established, and was assigned an Aalenian–Bajocian age. Miospore evidence also
580 contributed to this age assignment, which is consistent with other studies on marine
581 microplankton (Riding and Thomas 1992, Poulsen and Riding 2003).

582 An integrated study on the palaeobotany and palynology of the Dansirit Formation
583 (Middle Jurassic) from the Soltanieh Mountains of Zanjan Province, northwest Iran was
584 undertaken by Vaez-Javadi and Abbassi (2018). The dinoflagellate cysts recorded were
585 *Nannoceratopsis tricerias*, *Pareodinia ceratophora* and *Pareodinia* sp. cf. *P. prolongata*. The
586 specimens figured are not in an optimal preservational state (Vaez-Javadi and Abbassi 2018,
587 pl. 1/16–19). The *Pareodinia ceratophora*-*Nannoceratopsis tricerias* assemblage zone, of
588 Aalenian–Bajocian age, was erected on the basis of this material.

589 Badihagh et al. (2019) is a detailed study of the palynomorphs and plant
590 macrofossils from the Hojedk Formation of Well 233, southwest of Tabas city, northeast Iran.
591 This part of the Hojedk Formation of the Tabas Block is interpreted as being Middle Jurassic
592 (?Bajocian–Bathonian) in age based on the pollen and spores which dominate the 48 samples
593 studied. The entire succession studied was assigned to the *Klukisporites variegatus* acme
594 zone by Badihagh et al. (2019). This interpretation was based on the consistent and abundant
595 occurrence of the pteridophytic spore *Klukisporites variegatus*. However rare unidentified
596 dinoflagellate cysts were recorded in samples 42 and 41, in the uppermost part of the Hojedk
597 Formation, by Badihagh et al. (2019, fig. 2, tables 1, 2). It is clear that all the palynomorphs
598 recovered from the Hojedk Formation of the South Khorasan Province are very dark and
599 relatively poorly-preserved (Badihagh et al. 2019, fig. 4). This is indicative that this unit had
600 been subjected to high levels of thermal alteration. These authors illustrated one
601 indeterminate dinoflagellate cyst (Badihagh et al. 2019, fig. 4r). It is a poorly-preserved
602 subpentagonal specimen which is circumcavate/epicavate, and has a cingulum and a
603 precingular archaeopyle. The overall morphology, plus the relatively small hypocyst and the
604 apparently broken/damaged apical horn strongly suggests that this specimen is referable to
605 *Gonyaulacysta jurassica* subsp. *adepta*. The total range of this subspecies is Bathonian to
606 Oxfordian, but it is only common and consistent between the Callovian and middle Oxfordian
607 (Riding et al. 1985, Riding and Thomas 1992, 1997, Wiggan et al. 2017). If the specimen
608 illustrated by Badihagh et al. (2019, fig. 4r) is *Gonyaulacysta jurassica* subsp. *adepta*, this is

609 substantially more suggestive that the uppermost Hojedk Formation is Callovian as opposed
610 to Bathonian; it cannot be of Bajocian age.

611 The five contributions reviewed herein on the Lower and Middle Jurassic
612 (Pliensbachian–Callovian) successions of northern Iran indicate clearly that the entire region
613 has been subjected to significant levels of sub-metamorphic thermal alteration over a
614 substantial interval. This is because, following faulting during the early part of the
615 Cimmerian orogeny, Middle Jurassic siliciclastic successions were deposited in northern Iran
616 and these were affected by the Mid Cimmerian orogenic event throughout the Iran Plate
617 (Zanchi et al., 2009). Unsurprisingly, this intense tectonism has badly affected palynomorph
618 preservation. The four contributions authored or co-authored by Fatemeh Vaez-Javadi clearly
619 prove that there was a low diversity dinoflagellate cyst association, dominated by the genera
620 *Nannoceratopsis* and *Pareodinia*, in Aalenian and Bajocian successions throughout northern
621 Iran. The units examined were the Dansirit Formation of northwest Iran and the Hojedk and
622 Shemshak formations of northeast Iran. Badihagh et al. (2019) also studied the Hojedk
623 Formation of northeast Iran. These authors found evidence that part of this unit appears to be
624 somewhat younger, i.e. Callovian in age.

625 The remaining contribution on the Middle East is Issautier et al. (2019). This is a
626 major work on the depositional environments, palynostratigraphy, sedimentology and
627 sequence stratigraphy of the Minjur Formation in central Saudi Arabia. This unit was studied
628 in detail via examination of 112 cuttings and 12 core samples collected from five exploration
629 wells in central and eastern Saudi Arabia (Issautier et al. 2019, figs 1, 11, 12). The
630 palynology of this material was documented in detail, and six palynomorph zones
631 ('palynozones') established which span the late Carnian to Pliensbachian interval. These
632 authors reported the occurrences of the dinoflagellate cyst species *Dapcodinium priscum*,
633 *?Hebecysta* spp., *Nannoceratopsis gracilis*, *Rhaetogonyaulax dilatata*, *Rhaetogonyaulax*
634 *rhaetica* and *Rhaetogonyaulax wigginsii*, together with acritarchs, foraminiferal test linings,
635 freshwater algae, pollen and spores, and prasinophytes (Issautier et al. 2019, p. 155–158;
636 170–179). It is clear that the cuttings samples are badly affected by uphole contamination or
637 caving of substantially younger Jurassic dinoflagellate cysts such as *Ctenidodinium*
638 *sellwoodii*, *Korystocysta* spp. and *Systematophora penicillata* (see Issautier et al. 2019, fig.
639 16, enclosures 1–4). Significantly, one of these allochthonous forms is *Wanaea verrucosa*
640 which is a marker for the late Bajocian to early Bathonian interval of Australasia (Mantle and
641 Riding 2012). These occurrences indicate that *Wanaea verrucosa* has a wider
642 palaeogeographical extent than was initially envisaged.

643

644 **2.9. Sub-Arctic Russia west of the Ural Mountains**

645 Holm-Alwmark et al. (2019) is the only item solely on sub-Arctic western Russia that is
646 relevant to this review (Appendix 1 of the Supplementary data). These authors analysed
647 samples from a basal breccia and the overlying Kovernino Formation, both from above the
648 Puchezh-Katunki impact structure east of Moscow in western Russia. Abundant pollen and
649 spores, together with *Mendicodinium* spp. and unidentified dinoflagellate cysts were reported,
650 and interpreted to be Pliensbachian to early Toarcian in age.

651

652

653 **3. Conclusions**

654 From February 2019 to March 2020, 63 publications pertaining to Triassic to earliest
655 Cretaceous dinoflagellate cysts were discovered which are further to the 1878 already
656 compiled by Riding (2012, 2013, 2014, 2019a, 2020). This makes a cumulative total of 1941
657 relevant items (Table 1). These 63 contributions are listed in Appendix 1 of the Supplemental
658 data, and are mostly on the Jurassic of Africa, the Arctic, Europe and the Middle East (Table
659 2). Items on East and West Europe are most numerous (eight and 17 respectively), and
660 overall comprise 39.7% of the total (Table 2). This marked bias towards Europe was
661 previously recorded by by Riding (2012, 2013, 2014, 2019a, 2020). Nine and six items are on
662 the Arctic and the Middle East, and this represents 14.2% and 9.5% respectively. Africa is
663 also well-represented with 4 papers (6.4%). The other regions represented, sub-Arctic
664 Canada, China and Japan and sub-Arctic Russia, together make up 8% of the total. Multi-
665 region studies and publications with no geographical focus comprise 6.3% and 16%
666 respectively (Table 2). In terms of the stratigraphical intervals investigated, the spread is
667 relatively equable. The Early Jurassic has most studies with 19 papers either entirely focused
668 on, or including data from, this interval (Table 3).

669

670

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679

680

681 **Disclosure statement**

682 The author has no potential conflict of interest.

683

684 **Notes on contributor**

685

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688 Sheffield and, several years later, Jim was awarded a PhD by the same institution. During
689 2004, Jim gained a DSc from the University of Leicester, where he did his Bachelor's degree
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692 palynomorph preparation techniques, the history of palynology, and the morphology,
693 systematics and taxonomy of dinoflagellate cysts. Jim is a past Director-at-Large and
694 President of AASP – The Palynological Society, and became Managing Editor in 2004.

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696

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Display material captions:

Table 1. A breakdown of the 1941 publications on Triassic to earliest Cretaceous dinoflagellate cysts compiled by Riding (2012, 2013, 2014, 2019a, 2020) and herein based on the 23 relevant specified geographical region(s), plus multi-region studies and those with no

1014 geographical focus, and the initial letter of the family name of the first author. The number in
1015 the geographical region cell refers to the number of relevant published items on that area
1016 alone. An ellipsis (...) indicates a zero return for that particular parameter.

1017

1018 **Table 2.** A breakdown of the 63 publications on Triassic to earliest Cretaceous dinoflagellate
1019 cysts compiled herein, based on 11 specified relevant geographical region(s) plus multi-
1020 region studies and those with no geographical focus, and the initial letter of the family name
1021 of the first author. The number in the geographical region cell refers to the number of
1022 relevant published items on that area alone. An ellipsis (...) indicates a zero return for that
1023 particular parameter.

1024

1025 **Table 3.** A breakdown of the 63 publications on Triassic to earliest Cretaceous dinoflagellate
1026 cysts compiled herein, subdivided chronostratigraphically. The intervals are Triassic, Early
1027 Jurassic, Middle Jurassic, Late Jurassic, Jurassic-Cretaceous transition, investigations
1028 comprising three or more of the previous intervals and studies with no stratigraphical focus,
1029 and reworking. Some latitude and pragmatism are used in this compilation. For example if a
1030 publication is on the Berriasian and Valanginian it is classified as covering the Jurassic-
1031 Cretaceous transition. One item may be counted twice if, for example, it spans the Toarcian
1032 to Bathonian i.e. Early and Middle Jurassic) but not three times. An ellipsis (...) indicates a
1033 zero return for that particular parameter.

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SUPPLEMENTARY DATA I

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1039

Appendix 1. List of Literature

1040

1041 Sixty-three contributions on Triassic to earliest Cretaceous dinoflagellate cysts issued after
1042 the publication of Riding (2012, 2013, 2014, 2019a, 2020), and older papers discovered after
1043 these compilations were made, are listed in alphabetical/chronological order below. The
1044 reference format used is much the same as in Riding (2013), which was slightly modified
1045 from Riding (2012). Digital Object Identifier (doi) numbers are included where these are
1046 available. The nine papers which are deemed to be of major significance are asterisked. The
1047 language in which a paper was written in is indicated if it is not in English. A synthesis of the
1048 scope of each item is given as a string of keywords in parentheses after each citation. These
1049 keywords attempt to comprehensively summarise the principal subject matter, age range,
1050 major geographical region(s) and country/countries. A distinction is made between
1051 publications which present new data ('primary data'), and those which compile, review or
1052 summarise existing data ('compilation' etc.). Two abstracts are listed here, and these are
1053 denoted by the word 'summary' in the keyword string. If the author(s) have included
1054 photographs, occurrence charts and a zonal breakdown, these are indicated respectively in the
1055 keywords. For the purpose of this work, the world is subdivided into 23 major geographical
1056 regions. These are East Africa, North Africa, Southern Africa, Central America, northern
1057 South America, southern South America, Greater Antarctica, the Antarctic Peninsula, East
1058 Arctic, West Arctic, Southeast Asia, Australasia, sub-Arctic East Canada, sub-Arctic West
1059 Canada, China and Japan, East Europe, sub-Arctic West Europe, the Indian subcontinent, the
1060 Middle East, sub-Arctic Russia east of the Ural Mountains, sub-Arctic Russia west of the
1061 Ural Mountains, U.S.A. east of the Rocky Mountains and U.S.A. west of the Rocky
1062 Mountains (Table 1).

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1073 charts; photographs; latest Triassic–earliest Jurassic [Rhaetian–Hettangian]; sub-Arctic West
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1084 acme zone; lithostratigraphy [Hojedk Formation]; Mid Asian part of the Indo-European floral
1085 province; palaeobiogeography; palaeobotany; palaeoclimate; palaeoecology; palynofacies;
1086 plant macrofossils; pollen and spores; primary data; occurrence charts; photographs; Middle
1087 Jurassic [?Bajocian–Bathonian]; Middle East [Well 233, South Kuchak-Ali area, South
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1120 Member and the Hałuszowa, Jarmuta, Malinowa Shale, Szczawnica and Szlachtowa
1121 formations]; Magura Nappe; Pieniny Klippen Belt; structural geology; tectonic thrust sheet;
1122 tectonics; West Carpathian Mountains; primary data; quantitative occurrence chart;
1123 photographs; Early–Middle Jurassic to Eocene [Toarcian–Aalenian to Ypresian]; East Europe
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1132 formations], reworking; taxonomy; tectonic slices; primary data; non-quantitative occurrence
1133 charts; photographs; Early Cretaceous [Berriasian–Valanginian]; sub-Arctic West Europe
1134 [Schrambachgraben, Salzachtal, near Hallein and Kuchl, south of Salzburg, central Austria])
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1153 palynofacies; pollen and spores; prasinophytes; primary data; photographs; non-quantitative
1154 and quantitative occurrence charts; Early Cretaceous [Berriasian–Aptian]; North Africa
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1165 fungi; glossary; *incertae sedis*; mineral grains; pollen; prasinophytes; radiolaria; rules of
1166 nomenclature; schizosporous algae; silicoflagellates; sponges; spores; taxonomy;
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1174 (11 genera); evolution; generic definitions; history; morphology (acavate/cavate; horn and
1175 ornamentation distribution; intergradation; sulcus offset to the left); palaeoecology;
1176 palaeogeography; stratigraphical occurrences; tabulation; taxonomy; type material;
1177 taxonomic review; photographs; Late Jurassic–Cretaceous–Paleogene/Neogene
1178 [Kimmeridgian–Holocene]; no geographical focus)
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1191 palaeoecology; palaeogeography; palynofacies; pollen and spores; sedimentology; primary
1192 data; quantitative occurrence charts; photographs; earliest–Late Cretaceous [Berriasian–
1193 Cenomanian]; multi-region: sub-Arctic West Europe [Spain]; U.S.A. east of the Rocky
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1200 forcing; eustacy; flooding surfaces; gas hydrate (methane) release; glacioeustacy;
1201 lithostratigraphy [Komorowo, Drzewica and Ciechocinek formations]; *Luehndea spinosa*;
1202 megaspores; palaeoclimate; Polish Basin; sedimentology; sediment supply; sequence

1203 stratigraphy; stepwise carbon isotope excursions; terrestrial organic matter; Toarcian Oceanic
1204 Anoxic Event [T-OAE]; weathering; primary data; Early Jurassic [Pliensbachian–Toarcian];
1205 East Europe [Brody-Lubienia, Gorzów Wielkopolski, Kozłowice, Mechowo, Parkoszowice
1206 and Suliszowice, central Poland])

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1213 processes; biostratigraphy; *Botryococcus braunii*; carbonate minerals; Carboniferous
1214 reworking; chemostratigraphy; *Classopollis classoides*; clay mineralogy; Cleveland Basin;
1215 diagenesis; geochemistry [carbon isotope analysis; carbon:nitrogen ratios, elemental
1216 analyses; total nitrogen; total organic carbon]; hand-held X-ray fluorescence analyses;
1217 hyperthermal event; kerogen; Liasidium Event; *Liasidium variabile*; lithostratigraphy
1218 [Silicious Shale Member of the Redcar Mudstone Formation]; organic matter;
1219 palaeoclimatology; palaeoecology; paragenesis; petrography; pollen and spores;
1220 prasinophytes; scanning electron microscopy; sea level changes and sequence stratigraphy
1221 [lithological cycles/parasequences, maximum flooding, regressive-transgressive facies trends;
1222 short eccentricity cycles]; X-ray diffraction; primary data; quantitative occurrence chart;
1223 photographs; Early Jurassic [Sinemurian]; sub-Arctic West Europe [Boggle Hole, Robin
1224 Hood's Bay, North Yorkshire, northern England])

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1233 crinoids; crustaceans; *Dapcodinium priscum*; echinoids; foraminifera; gastropods;
1234 geochemistry; geological setting; Global Stratotype Section and Point (GSSP); kerogen
1235 analysis; lithostratigraphy (the Eiberg Member of the Kössen Formation and the
1236 Tiefengraben Member of the Kendlbach Formation); ostracods; palaeobiogeography;

1237 palaeomagnetism; pollen and spores; radiometric geochronology; *Rhaetogonyaulax rhaetica*;
1238 Triassic-Jurassic (T-J) boundary; scaphopods; primary data; latest Triassic and earliest
1239 Jurassic [Rhaetian–Hettangian]; sub-Arctic West Europe [the Kuhjoch Pass, Karwendel
1240 Mountains, Northern Calcareous Alps, Tyrol, western Austria])

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1248 dating); geological setting; history of study; impactites; lithostratigraphy (Kovernino
1249 Formation); *Mendicodinium* spp.; Permian reworking; petrography; pollen and spores; post-
1250 impact crater lake sediments; prasinophytes; radiometric dating; thin sections; Early Jurassic
1251 [Pliensbachian–Toarcian]; sub-Arctic Russia west of the Ural Mountains [Puchezh-Katunki
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1262 eustacy; glaciations; glendonites; morphology; palaeoclimate; palaeoecology; summary; Late
1263 Jurassic–Early Cretaceous [Oxfordian–Valanginian]; West Arctic [Rollrock section, northern
1264 Ellesmere Island, Sverdrup Basin, Arctic Canada])

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1272 environments; facies analysis; foraminiferal test linings; freshwater algae; geological
1273 background; isopach map; lithostratigraphy [Minjur Formation]; palaeoclimate;
1274 palaeoecology; palaeogeography; pollen and spores; prasinophytes; reworking;
1275 sedimentology; sequence stratigraphy; primary data; semi-quantitative occurrence charts;
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1295 change/global warming; fluvial flood events; global carbon release; hydrological cycling;
1296 lithostratigraphy [Nishinakayama Formation, Toyora Group]; *Luehndea spinosa*;
1297 palaeoecology; palynofacies; pollen and spores; Tabe Basin; thermal maturity; thin sections;
1298 Toarcian Oceanic Anoxic Event (T-OAE); primary data; semi-quantitative occurrence chart;
1299 photographs; Early Jurassic [Toarcian]; China and Japan [Sakuraguchi-dani stream section,
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1307 chemostratigraphy; correlation; downhole logging; facies analysis; fish teeth; frost
1308 weathering; geological setting; lithostratigraphy [Agardhfjellet Formation]; Longyearbyen
1309 carbon dioxide storage project; palaeoecology; marine reptiles; molluscs; reworking; sea
1310 floor oxygenation levels; sedimentology; sequence stratigraphy; structural geology; this
1311 sections; total organic carbon [TOC]; trace fossils; X-ray fluorescence geochemistry; primary
1312 data; semi-quantitative occurrence chart; Middle Jurassic–earliest Cretaceous [Bathonian–
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1321 palaeobathymetry; palaeoecology; palaeogeography; provenance analysis; reworking;
1322 primary data; occurrence chart; photographs; Late Jurassic [Oxfordian–Kimmeridgian]; East
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1330 sedimentology; sequence stratigraphy; Toarcian Oceanic Anoxic Event [T-OAE]; trace
1331 fossils; primary data; Early–Middle Jurassic [Pliensbachian–Aalenian]; multi-region: North
1332 Africa [Central High Atlas Basin, Morocco]; West Arctic (Jameson Land Basin, East
1333 Greenland)

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1343 freshwater algae; geological setting; molluscs; pollen and spores; prasinophytes;
1344 *Sentusidinium-Batiacasphaera-Kallosphaeridium* group; primary data; photographs; Late
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1352 formations]; palaeoclimates; palaeoecology; palaeogeography; palaeovegetation; pollen and
1353 spores; primary data; photographs; Late Jurassic–Early Cretaceous [undifferentiated–
1354 Barremian]; China and Japan [Doilongdegqin County, Lhasa Block, central Qinghai-Xizang
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1365 palaeosalinity; Paratethyan basins; size variation; tabulation; taxonomy; *Thalassiphora*
1366 *robusta*; *Thalassiphora* species complex; primary data and review article; occurrence chart;
1367 photographs; Late Jurassic–Late Miocene [Tithonian (Volgian)–undifferentiated]; multi-
1368 region: sub-Arctic East Canada [Shubenacadie H-100 well, Scotian margin, offshore Nova
1369 Scotia]; East Europe [Medvednica, Slavonija, Zagorje and Žumberak, Croatia and unknown
1370 locations in southwestern Romania])
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1378 pollen; compilation/primary data; photographs; bioevent charts; latest Jurassic–Late
1379 Cretaceous [Tithonian–Maastrichtian]; West Arctic [Traill Ø to Store Koldeway, northeast
1380 Greenland])

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1391 geological setting; lithostratigraphy [Kapp Toscana and Adventdalen groups]; macrofossils;
1392 *Mancodinium semitabulatum*; palaeogeography; plant fossils; sedimentology; seismic
1393 interpretation; sequence stratigraphy; structural geology; trace fossils; volcanism; primary
1394 data and compilation; Late Triassic–Early Cretaceous [Norian–Aptian]; East Arctic [Kong
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1401 palaeoecology; pollen and spores; primary data; non-quantitative occurrence charts;
1402 photographs; Middle Jurassic to Early Cretaceous [?Bajocian–Tithonian to Albian]; North
1403 Africa (northern Western Desert, northern Egypt)

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1412 background; lithostratigraphy; palaeoclimate; palaeoecology; pollen and spores;
1413 prasinophytes; *Rhaetogonyaulax arctica*; *Rhaetogonyaulax rhaetica*; sequence stratigraphy;
1414 taxonomy; primary data/compilation; bioevent charts; photographs; Middle–Late Triassic
1415 [Anisian–Rhaetian]; East Arctic [Arctic Norway, Barents Sea, Svalbard Archipelago])

1416

1417 PATERSON, N.W., MORRIS, P.H., and MANGERUD, G. 2019. Lycopoid megaspores from
1418 the Upper Triassic of Svalbard and their relationship to the floras and palaeoenvironments of
1419 northern Pangaea. *Papers in Palaeontology*, 5(4): 577–599 (doi: 10.1002/spp2.1251).

1420 (agglutinated foraminifera; biostratigraphy; biozonation; eustacy; foraminiferal test linings;
1421 freshwater algae; geological setting; kerogen; lithostratigraphy [Kapp Toscana Group];
1422 megaspores from heterosporous lycopoids; micro-biofacies; ostracods; palaeoecology;
1423 preservation potential; pollen and spores; radiolaria; *Rhaetogonyaulax rhaetica*;
1424 sedimentology; primary data; Late Triassic [Carnian–Rhaetian]; East Arctic [Hopen Island,
1425 southeast Svalbard archipelago, Arctic Ocean])

1426

1427 POCOCK, S.A.J. 1962. Jurassic palynology in the Western Canada Basin. *Oil in Canada*,
1428 February 8th, 1962: 36–40.

1429 (acritarchs; biostratigraphy; correlation [with Europe]; foraminiferal test linings;
1430 *Gonyaulacysta jurassica*; Jurassic–Cretaceous transition; lithostratigraphy [Ferne Shale and
1431 Manneville groups]; palaeoecology; pollen and spores; West Canada Basin; review article;
1432 histograms; photographs; Late Jurassic–Early Cretaceous [Kimmeridgian–Hauterivian]; sub-
1433 Arctic West Canada [British Columbia, Alberta and Saskatchewan])

1434

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1438 REOLID, M., DUARTE, L.V., and RITA, P. 2019. Changes in foraminiferal assemblages
1439 and environmental conditions during the T-OAE (Early Jurassic) in the northern Lusitanian

1440 Basin, Portugal. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 520: 30–43 (doi:
1441 10.1016/j.palaeo.2019.01.022).

1442 (ammonite zones; biotic crisis and recovery; brachiopods; calcareous nannofossils; diversity;
1443 echinoderms; foraminifera; geological setting; Iberian palaeomargin; lithostratigraphy [Sao
1444 Giao Formation]; mass extinction; opportunistic taxa; organic geochemistry; ostracods;
1445 oxygen depletion; palaeoecology; palaeoproductivity; phosphorus; redox-sensitive elements;
1446 tempestite-turbidite facies; Toarcian Oceanic Anoxic Event [T-OAE]; trace fossils; data
1447 compilation; Early Jurassic [Toarcian]; sub-Arctic West Europe [Maria Pares section,
1448 Rabaçal, northern Lusitanian Basin, western Portugal])

1449

1450 RIDING, J.B., LEBEDEVA, N.K., and GORYACHEVA, A.A. 2019. Obituary. Vera
1451 Ivanovna Ilyina (1930–2018). *Palynology*, 43(3): 349–354 (doi:
1452 10.1080/01916122.2019.1586090).

1453 (acritarchs; biography; biostratigraphy; biozonation; history; Institute of Geology and
1454 Geophysics, Academgorodok, Novosibirsk, Siberia; International Association for the
1455 Promotion of Co-operation with Scientists from the New Independent States of the former
1456 Soviet Union [INTAS]; obituary, pollen and spores; prasinophytes; Third International
1457 Conference on Palynology [1971]; Vera I. Ilyina; review article; Early Jurassic–Early
1458 Cretaceous [Hettangian–Valanginian]; multi-region: East Arctic [northern Russia]; sub-
1459 Arctic Russia east of the Ural Mountains [undifferentiated], sub-Arctic Russia west of the
1460 Ural Mountains [undifferentiated] including Kazakhstan)

1461

1462 RODRIGUES, B., SILVA, R.L., FILHO, J.G.M., SADKI, D., MENDONÇA, J.O., and
1463 DUARTE, L.V. 2020. Late Pliensbachian–Early Toarcian palaeoenvironmental dynamics and
1464 the Pliensbachian–Toarcian Event in the Middle Atlas Basin (Morocco). *International
1465 Journal of Coal Geology*, 217, 103339 (doi: 10.1016/j.coal.2019.103339).

1466 (acritarchs; ammonite zones; *Botryococcus*; carbon cycle; continental weathering; eustacy;
1467 foraminiferal test linings; fungal spores; geological background; *Luehndea spinosa*;
1468 *Nannoceratopsis gracilis*; organic geochemistry; palaeoclimate; palaeoecology;
1469 palaeoenvironments; palynofacies; Pliensbachian–Toarcian event; pollen and spores;
1470 prasinophytes; sedimentology; sequence stratigraphy; tectonics; thermal maturity; total
1471 organic carbon [TOC]; vitrinite reflectance; zygospores; primary data; photographs; Early
1472 Jurassic [Pliensbachian–Toarcian]; North Africa [Ait Moussa and Issouka sections, northeast
1473 of Boulemane, Fès-Meknès region, Middle Atlas Basin, northeast Morocco])

1474

1475 ROGALSKA, M. 1962. Analiza sporowo-pyłkowa osadów jurajskich północnej części
1476 Pasma Krakowsko-Wieluńskiego. [Spore and pollen grain analysis of Jurassic sediments in
1477 the northern part of the Cracow – Wieluń Cuesta] *Instytut Geologii Czyny Odbitka z Prac*,
1478 30(3): 495–524 (in Polish with English and Russian summaries).
1479 (biostratigraphy; correlation; lithostratigraphy; *Pareodinia*; pollen and spores; prasinophytes;
1480 taxonomy; primary data; quantitative occurrence chart and non-quantitative occurrence chart;
1481 photographs; Triassic–Middle Jurassic [Rhaetian–?Bathonian]; East Europe [northern
1482 Kraków–Wieluń Cuesta/Scarp, southern Poland])

1483

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1487 SARJEANT, W.A.S., and DOWNIE, C. 1966. The classification of dinoflagellate cysts
1488 above generic level. *Grana Palynologica*, 6(3): 503–527 (doi:
1489 10.1080/00173136609430038).

1490 (history of study; morphological basis of dinoflagellate ctst classification; problems of the
1491 previous classification; suprageneric classification; taxonomy; compilation/review; no
1492 geographical or stratigraphical focus)

1493

1494 SARJEANT, W.A.S., and DOWNIE, C. 1974. The classification of dinoflagellate cysts
1495 above generic level: a discussion and revisions. *Symposium on Stratigraphical Palynology*.
1496 *Birbal Sahni Institute of Palaeobotany Special Publication*, No. 3: 9–32.

1497 (familial groupings; living dinoflagellates and their cysts; principles of classification;
1498 suprageneric classification; taxonomy; compilation/review; no geographical or stratigraphical
1499 focus)

1500

1501 SCHOBEN, M., GRAVENDYCK, J., MANGELS, F., STRUCK, U., BUSSERT, R.,
1502 KÜRSCHNER, W.M., KORN, D., SANDER, P.M., and ABERHAN, M. 2019. A
1503 comparative study of total organic carbon- $\delta^{13}\text{C}$ signatures in the Triassic–Jurassic
1504 transitional beds of the Central European Basin and western Tethys shelf seas. *Newsletters on*
1505 *Stratigraphy*, 52(4): 461–486 (doi: 10.1127/nos/2019/0499).

1506 (ammonites; biostratigraphy; biozonation; bivalves; carbon cycle; carbon isotope analysis;
1507 chemostratigraphy; clay mineralogy; conchostracans; correlation; *Dapcodinium priscum*;

1508 end-Triassic mass extinction; geological setting; Global Stratotype Section and Point [GSSP];
1509 lithostratigraphy [Postera Beds to the Pilonotenton Formation]; palaeoclimate; pollen and
1510 spores; *Rhaetogonyaulax rhaetica*; *Suessia swabiana*; total nitrogen (TN); total organic
1511 carbon (TOC); weathering; primary data; semiquantitative occurrence chart; latest Triassic–
1512 earliest Jurassic [Rhaetian–Hettangian]; sub-Arctic West Europe [clay quarry northwest of
1513 Bonenburg village, near Warburg, North Rhine-Westphalia, west central Germany; Kuhjoch,
1514 near Hinteriss, central Austria])

1515

1516 SCHÖLLHORN, I., ADATTE, T., VAN DE SCHOOTBRUGGE, B., HOUBEN, A.,
1517 CHARBONNIER, G., JANSSEN, N., and FÖLLMI, K.B. 2020. Climate and environmental
1518 response to the break-up of Pangea during the Early Jurassic (Hettangian–Pliensbachian); the
1519 Dorset coast (UK) revisited. *Global and Planetary Change*, 185, 103096 (doi:
1520 10.1016/j.gloplacha.2019.103096).

1521 (anoxia; bioproductivity; carbon and oxygen isotopes; chemical index of alteration; clay
1522 mineralogy; continental breakup; correlation; eustacy; geochemistry; mineralogy;
1523 palaeoclimatology; palaeogeography; palaeoceanography; Pangea; sediment deposition rates;
1524 compilation; Early Jurassic (Hettangian–Pliensbachian); sub-Arctic West Europe [Pinhay
1525 Bay to Eype Mouth, Dorset, and St Audries Bay, Somerset, southern England])

1526

1527 SHEVCHUK, O.A. 2018. *Microfossils and biostratigraphy of the Middle Jurassic–*
1528 *Cretaceous of Ukraine*. Thesis for the degree of Doctor of Geological Sciences by specialty
1529 04.00.09 “Paleontology and Stratigraphy (103 – Earth Sciences). Institute of Geological
1530 Sciences of the National Academy of Sciences of Ukraine, Kiev, 42 p. (in Ukrainian with an
1531 English summary).

1532 (acritarchs; *Botryococcus*; biostratigraphy; biozonation; correlation; foraminiferal test linings;
1533 fungal spores; megaspores; palynofacies; *Pediastrum*; pollen and spores; prasinophytes;
1534 thesis summary; quantitative range charts; Middle Jurassic–Cretaceous [Aalenian–
1535 Maastrichtian]; East Europe [Teteic and Boreal-Atlantic belt, Ukraine])

1536

1537 SHEVCHUK, O., SLATER, S.M., and VAJDA, V. 2018. Palynology of Jurassic (Bathonian)
1538 sediments from Donbas, northeast Ukraine. *Palaeobiodiversity and Palaeoenvironments*,
1539 98(1): 153–164 (doi: 10.1007/s12549-017-0310-3).

1540 (biostratigraphy; *Botryococcus*; Dnieper–Donets Basin; Donbas fold belt; insect remains;
1541 lithostratigraphy [Kamyanska suite]; parent plants; petroleum geology; pollen and spores;

1542 provincialism; regional geology; sedimentology; thermal alteration index [TAI]; vegetation
1543 dynamics; primary data; quantitative occurrence charts; photographs; Middle Jurassic
1544 [Bathonian]; East Europe [Kamyanka village, Kharkiv region, northeast Ukraine])
1545

1546 *SKUPIEN, P., and DOUPOVCOVÁ, P. 2019. Dinoflagellates and calpionellids of the
1547 Jurassic–Cretaceous boundary, Outer Western Carpathians (Czech Republic). *Cretaceous*
1548 *Research*, 99: 209–228 (doi: 10.1016/j.cretres.2019.02.017).
1549 (biostratigraphy; calcareous dinoflagellate cysts; calpionellids; Jurassic–Cretaceous
1550 boundary; lithostratigraphy [Vendryně Formation and Těšín Limestone]; *Nannoceratopsis*;
1551 reworking; primary data; non-quantitative occurrence chart; photographs; latest Jurassic–
1552 earliest Cretaceous [Tithonian–Berriasian]; East Europe [Bruzovice River locality,
1553 Bruzovice, Outer Western Carpathians, eastern Czech Republic])
1554

1555 SLATER, S.M., McKIE, T., VIEIRA, M., WELLMAN, C.H., and VAJDA, V. 2017.
1556 Episodic river flooding events revealed by palynological assemblages in Jurassic deposits of
1557 the Brent Group, North Sea. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 485:
1558 389–400 (doi: 10.1016/j.palaeo.2017.06.028).
1559 (acritarchs; *Botryococcus*; correlation; facies analysis; foraminiferal test linings; geological
1560 setting; hyperpycnites; lithostratigraphy [Rannoch Formation of the Brent Group];
1561 megaspores; non-metric multidimensional scaling; palaeoecology; palaeogeography;
1562 palynofacies; pollen and spores; prasinophytes; sedimentology; vegetational dynamics;
1563 primary data; photographs; Middle Jurassic [Aalenian–Bajocian]; sub-Arctic West Europe
1564 [Don North East and Penguins Cluster oilfields, Viking Graben, northern North Sea, UK
1565 sector])
1566

1567 SLATER, S.M., TWITCHETT, R.J., DANISE, S., and VAJDA, V. 2019. Substantial
1568 vegetation response to Early Jurassic global warming with impacts on oceanic anoxia. *Nature*
1569 *Geoscience*, 12: 462–467 (doi: 10.1038/s41561-019-0349-z).
1570 (acritarchs; global warming; lithostratigraphy [Cleveland Ironstone and Whitby Mudstone
1571 formations]; palynofacies; pollen and spores; prasinophytes; Toarcian Oceanic Anoxic Event;
1572 vegetation dynamics; primary data; photographs; Early Jurassic [Pliensbachian–Toarcian];
1573 sub-Arctic West Europe [Kettlethness, Port Mulgrave, Runswick Bay, Saltwick Bay and
1574 Staithes, North Yorkshire, northern England])
1575

1576 *SMELROR, M., LARSEN, G.B., OLAUSSEN, S., RØMULD, A., and WILLIAMS, R.
1577 2018. Late Triassic to Early Cretaceous palynostratigraphy of Kong Karls Land, Svalbard,
1578 Arctic Norway, with correlations to Franz Josef Land, Arctic Russia. *Norwegian Journal of*
1579 *Geology*, 98(4): 1–31 (doi: 10.17850/njg004).
1580 (acritarchs; ammonites; biostratigraphy; biozonation; correlation; freshwater algae; hiatuses;
1581 lithostratigraphy [Kapp Toscana and Adventdalen groups]; pollen and spores; prasinophytes;
1582 reworking; sedimentology; sequence stratigraphy; primary data; non-quantitative occurrence
1583 charts; photographs; Late Triassic–Early Cretaceous [Norian–Aptian]; East Arctic [Kong
1584 Karls Land, eastern Svalbard Archipelago, Arctic Ocean])
1585
1586 SMITH, D.G. 1982. Stratigraphic significance of a palynoflora from ammonoid-bearing
1587 Early Norian strata in Svalbard. *Newsletters on Stratigraphy*, 11(3): 154–161 (doi:
1588 10.1127/nos/11/1982/154).
1589 (ammonoids; ammonoid zones; biostratigraphy; biozonation; chronostratigraphy; correlation;
1590 lithostratigraphy [Flatsalen Formation]; pollen and spores; *Rhaetipollis germanicus*
1591 assemblage; *Rhaetogonyaulax rhaetica*; primary data and review; Late Triassic [Norian];
1592 East Arctic [Hopen Island, Svalbard archipelago])
1593
1594 STEEMAN, T., DE WEIRDT, J., SMITH, T., DE PUTTER, T., MEES, F., and LOUWYER,
1595 S. 2020. Dinoflagellate cyst biostratigraphy and palaeoecology of the early Paleogene
1596 Landana reference section, Cabinda Province, Angola. *Palynology*, 44(2), 280–309 (doi:
1597 10.1080/01916122.2019.1575091).
1598 (*Aldorfia aldorfensis*; biostratigraphy; biozonation; correlation; Darteville collection;
1599 foraminifera; organic geochemistry [total organic carbon - TOC]; palaeoecology; reworking;
1600 primary data; quantitative and semi-quantitative occurrence charts; photographs; Middle
1601 Jurassic reworking into Palaeocene–Eocene/Oligocene [Bathonian reworking into
1602 Danian/Selandian–?Priabonian/Rupelian]; Southern Africa [Landana coastal section, Cabinda
1603 Province, Congo Basin, Angola])
1604
1605 STORM, M.S., HESSELBO, S.P., JENKYN, H.C., RUHL, M., ULLMANN, C.V., XU, W.,
1606 LENG, M.J., RIDING, J.B., and GORBANENKO, O. 2020. Orbital pacing and secular
1607 evolution of the Early Jurassic carbon cycle. *PNAS (Proceedings of the National Academy of*
1608 *Sciences of the United States of America)*, 117(8): 3974–3982 (doi:
1609 10.1073/pnas.1912094117).

1610 (ammonite zones and subzones; astrochronology; carbon isotopes [$\delta^{13}\text{C}_{\text{TOC}}$] and their
1611 excursions; chemostratigraphy; chronostratigraphy; geochemistry; geological background;
1612 global carbon cycle; magmatic events; orbital forcing; organic geochemistry;
1613 palaeoenvironment; palaeogeography; palaeotemperature; sedimentary organic matter;
1614 Toarcian Oceanic Anoxic Event [T-OAE]; Triassic–Jurassic transition; compilation/data
1615 review; latest Triassic–Early Jurassic [Rhaetian–Toarcian]; sub-Arctic West Europe [Bristol
1616 Channel Basin, southwest England, UK; Mochras Borehole, Cardigan Bay Basin, West
1617 Wales, UK; Sancerre-Couy Borehole, Paris Basin, northern France])

1618

1619 STOVER, L.E., and EVITT, W.R. 1978. Analyses of pre-Pleistocene organic-walled
1620 dinoflagellates. *Stanford University Publications, Geological Sciences*, 15, 300 p.
1621 (appendices; archaeopyle types and variability; catalogue/index; classification;
1622 *Gonyaulacysta* complex [e.g. *Gonyaulacysta*, *Impagidinium*, *Leptodinium* and
1623 *Rhynchodiniopsis*]; *Herendeenia-Omatia*; *Kiokansium unituberculatum*;
1624 *Lanternosphaeridium* complex; line drawings; lists of species; morphology; opercula;
1625 peridiniacean genera; *Spiniferites* complex; synopsis of genera; taxonomy; compilation; no
1626 specific geographical and stratigraphical focus)

1627

1628 SVOBODOVÁ, A., ŠVÁBENICKÁ, L., REHÁKOVÁ, D., SVOBODOVÁ, M., SKUPIEN,
1629 P., ELBRA, T., and SCHNABL, P. 2019. The Jurassic/Cretaceous boundary and high
1630 resolution biostratigraphy of the pelagic sequences of the Kurovice section (Outer Western
1631 Carpathians, the northern Tethyan margin). *Geologica Carpathica*, 70(2): 153–182 (doi:
1632 10.2478/geoca-2019-0009).

1633 (acritarchs; biostratigraphy; calcareous dinoflagellate cysts; calcareous nannofossils;
1634 calpionellids; foraminiferal test linings; geological setting; Jurassic–Cretaceous boundary;
1635 Kurovice Limestone; limestones; magnetostratigraphy; microfacies; palaeobathymetry;
1636 palaeoecology; pollen and spores; prasinophytes; radiolarians; reworking; sponge spicules;
1637 Tethys; primary data; non-quantitative occurrence chart; photographs; latest Jurassic–earliest
1638 Cretaceous [Tithonian–Berriasian]); East Europe [Kurovice Quarry, near Zlín, Outer Western
1639 Carpathians, southeast Czech Republic])

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1644 TAUGOURDEAU LANTZ, J., and DONZE, P. 1971. Un aperçu de l'environnement végétal
1645 pendant l'épisode régressif du Berriasien terminal dans le Jura méridional (France). *Revue de*
1646 *Micropaléontologie*, 14(5): 102–120 (in French).
1647 (acritarchs; *Botryococcus*; eustacy; foraminiferal text linings; geological background;
1648 megaspores; palaeoecology; pollen and spores; prasinophytes; taxonomy; vegetational
1649 reconstructions; primary data; occurrence chart [percentages]; photographs; earliest
1650 Cretaceous [Berriasian]; sub-Arctic West Europe [France])

1651

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1654

1655 *VAEZ-JAVADI, F. 2018. Dinoflagellate palynostratigraphy of Middle Jurassic of the
1656 Hojedk Formation, Tabas, central-east Iran and its correlation to the other palynomorph zones
1657 in Iran and elsewhere. *Quarterly Journal of Geosciences*, 127: 265–276 (in Persian).
1658 (acritarchs; biostratigraphy; biozonation [*Nannoceratopsis gracilis* total range subzone and
1659 *Nannoceratopsis tricerias-Pareodinia ceratophora* assemblage zone]; correlation;
1660 lithostratigraphy [Hojedk Formation]; palaeoclimate; pollen and spores; prasinophytes; Tabas
1661 Block; primary data; occurrence chart; photographs; Middle Jurassic [Aalenian–Bajocian];
1662 Middle East [Tabas County, South Khorasan Province, northeast Iran])

1663

1664 VAEZ JAVADI, F. 2019. Middle Jurassic palynology of the southwest Tabas Block, Central-
1665 East Iran. *Palynology*, doi: 10.1080/01916122.2019.1637954.

1666 (acritarchs; biostratigraphy; biozonation [*Nannoceratopsis* sp. cf. *N. gracilis* interval zone];
1667 botanical affinity; Chahrekhneh borehole; correlation; Iran Plate; lithostratigraphy [Hojedk
1668 Formation]; palaeoclimate; palaeoecology; palaeogeography; pollen and spores; Tabas Block;
1669 Tethys Ocean; primary data; occurrence chart; photographs; Middle Jurassic [Aalenian–
1670 Bajocian]; Middle East [southwest of Tabas, Tabas County, South Khorasan Province,
1671 northeast Iran])

1672

1673 VAEZ-JAVADI, F., and ABBASSI, N. 2018. Middle Jurassic biostratigraphy of plant macro
1674 and microfossils in Soltanieh Mountains, south of Zanjan, NW Iran. *Geosciences*, 106: 91–
1675 102.

1676 (biostratigraphy; biozonation [*Pareodinia ceratophora-Nannoceratopsis tricerias* assemblage
1677 zone]; correlation; lithostratigraphy [Dansirit Formation, Shemshak Group]; palaeoecology;

1678 palaeogeography; plant macrofossils; pollen and spores; Tethys Ocean; primary data;
1679 occurrence chart; photographs; Middle Jurassic [Aalenian–Bajocian]; Middle East [Soltanieh
1680 Mountains, south of Zanjan city, Zanjan Province, northwest Iran])

1681
1682 VAEZ-JAVADI, F., GHAVIDEL-SYOOKI, M., and GHASEMI-NEJAD, I. 2003.
1683 Biostratigraphy of Shemshak Formation in Ozon Mountain, Jajarm based on dinoflagellata.
1684 *Journal of Science, University of Tehran*, 29(1): 141–160 (in Persian with an English
1685 abstract).
1686 (acritarchs; biostratigraphy; biozonation [*Nannoceratopsis spiculata* and *Valensiella ovulum*
1687 biozones]; lithostratigraphy [Shemshak Formation]; primary data; occurrence chart;
1688 photographs; Early–Middle Jurassic [Pliensbachian–Bajocian]; Middle East [Ozon Mountain,
1689 Jajarm County, North Khorasan Province, northeast Iran])

1690
1691 *VAN DE SCHOOTBRUGGE, B., HOUBEN, A.J.P., ERCAN, F.E.Z., VERREUSSEL, R.,
1692 KERSTHOLT, S., JANSSEN, N.M.M., NIKITENKO, B., and SUAN, G. 2019. Enhanced
1693 Arctic-Tethys connectivity ended the Toarcian Oceanic Anoxic Event in NW Europe.
1694 *Geological Magazine*, doi: 10.1017/S0016756819001262.
1695 (ammonite zones; anoxia; bioproductivity; biostratigraphy; black shale; carbon cycle; carbon
1696 isotopes; chemostratigraphy; correlation; diversity; eustacy; geochemistry; geological
1697 background; global warming; heterochroneity; lithostratigraphy; marine stratification;
1698 migrations; ocean circulation; palaeoenvironmental recovery; palaeoceanography;
1699 palaeosalinity; Toarcian Oceanic Anoxic Event [T-OAE]; total organic carbon; primary data;
1700 occurrence charts; photographs; Early Jurassic [Pliensbachian–Toarcian]; multi-region: East
1701 Arctic [Kelimyar River, Siberia, northeast Russia]; sub-Arctic West Europe [coastal outcrops
1702 between Staithes and Ravenscar, North Yorkshire, northern England and well 34/10-35,
1703 Tjalve Terrace, Gulfaks South oilfield, Norwegian sector of the northern North Sea])

1704

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1707

1708 WARRINGTON, G. 1976. British Triassic palaeontology. *Proceedings of the Ussher Society*,
1709 3(3): 341–353.

1710 (acritarchs; biostratigraphy; correlation; lithostratigraphy; macrofossils; microfossils; pollen
1711 and spores; prasinophytes; scolecodonts; compilation/review paper; Triassic [Induan–
1712 Rhaetian]; sub-Arctic West Europe [pan-United Kingdom])
1713

1714 WARRINGTON, G. 1980. Palynological studies of Triassic rocks in central Somerset
1715 (Abstract). *Proceedings of the Ussher Society*, 5(1): 90.
1716 (biostratigraphy; correlation; diversity; foraminiferal test linings; lithostratigraphy [Mercia
1717 Mudstone and Penarth groups]; pollen and spores; scolecodonts; summary; Late Triassic
1718 [Carnian and Rhaetian]; sub-Arctic West Europe [Burton Row and Puriton boreholes, near
1719 Bridgwater, central Somerset, southwest England])
1720

1721 WILLIAMS, G.L. 1965. Organic-walled microfossils aid oil search. *The Oil and Gas*
1722 *Journal*, November 22 1965: 108–112.
1723 (acritarchs; biostratigraphy; correlation; *Gonyaulacysta jurassica*; history of study;
1724 hystrichospheres; life cycle; modern dinoflagellates; morphology; oil/gas exploration;
1725 palaeoecology; review article; photographs; no geographical or stratigraphical focus)
1726

1727 WILLIAMS, G.L. 1974. 57. Biostratigraphy and paleoecology of the Mesozoic and Cenozoic
1728 rocks of the Atlantic Shelf. Project 710062. *Geological Survey of Canada Paper* 74–1, Part
1729 B: 150–152.
1730 (biostratigraphy; biozonation; correlation; lithostratigraphy [Western Bank, Nova Scotia and
1731 Gully groups]; offshore boreholes; oil/gas exploration; palaeoecology; pollen and spores;
1732 unconformity; review article; Middle Jurassic to Pliocene/Pleistocene [Bathonian/Callovia –
1733 undifferentiated]; sub-Arctic East Canada [Grand Banks and Scotian Shelf, offshore eastern
1734 Canada])
1735

1736 WILSON, G.J., and CLOWES, C.D. 1981. A concise catalogue of organic-walled fossil
1737 dinoflagellate genera. *New Zealand Geological Survey Report*, No. 92, 199 p.
1738 (archaeopyle type; catalogue; descriptions of genera; line drawings; morphology; range
1739 charts; compilation; Late Triassic [undifferentiated] to Holocene; no geographical or
1740 stratigraphical focus)
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SUPPLEMENTARY DATA II

1744

1745

Appendix 2. List of palynomorph species, subspecies and varieties

1746

1747 This Appendix alphabetically lists all valid palynomorph taxa below generic level which are
1748 mentioned in this contribution with full author citations. References to the author citations for
1749 the dinoflagellate cysts can be found in Williams et al. (2019 - *American Association of*
1750 *Stratigraphic Palynologists Contribution Series 50*, available at:

1751 [https://palynology.org/contribution-series-number-50-the-new-lentin-and-williams-index-](https://palynology.org/contribution-series-number-50-the-new-lentin-and-williams-index-2019/)

1752 [2019/](https://palynology.org/contribution-series-number-50-the-new-lentin-and-williams-index-2019/)). The recommendations of Williams et al. (2019) are followed with the following two

1753 exceptions. The proposals of Correia et al. (2017 - *Review of Palaeobotany and Palynology*

1754 237, p. 93) on the species *Nannoceratopsis senex* are followed herein. With regard to this

1755 species, Williams et al. (2019) adopted the taxonomic proposals of Ilyina et al. (1994 -

1756 *Russian Academy of Sciences, Siberian Branch, United Institute of Geology, Geophysics and*

1757 *Mineralogy, Transactions* 818), who proposed that *Nannoceratopsis senex* is a subspecies of

1758 *Nannoceratopsis deflandrei* Evitt 1961. Furthermore, the Linnaean binomial *Ctenidodinium*

1759 *sellwoodii* (Sarjeant 1975) Stover & Evitt 1978 is preferred herein to *Dichadogonyaulax*

1760 *sellwoodii* Sarjeant 1975. Most of the Jurassic tabulate gonyaulacoid species with epicystal

1761 archaeopyles are placed in *Ctenidodinium*. The species *sellwoodii* is clearly closely related to

1762 two contemporary species which are accommodated in *Ctenidodinium* according to Williams

1763 et al. (2019). These are *Ctenidodinium combazii* Dupin 1968 and *Ctenidodinium cornigerum*

1764 (Valensi 1953) Jan du Chêne et al. 1985. That said, there are substantial taxonomic issues

1765 with the two apparently very similar genera *Ctenidodinium* and *Dichadogonyaulax*. These

1766 genera require a thorough taxonomic review. It is eminently possible that *Dichadogonyaulax*

1767 is a junior synonym of *Ctenidodinium* as previously suggested by Lentin and Williams (1973

1768 - *Geological Survey of Canada Paper* 73-42, p. 46).

1769

1770 **Acritarch:**

1771 *Limbicysta bjaerkei* (Smelror, 1987) MacRae et al. 1996

1772

1773 **Dinoflagellate cysts:**

1774 *Aldorfia aldorfensis* (Gocht 1970) Stover & Evitt 1978

1775 *Ambonosphaera? staffinensis* (Gitmez 1970) Poulsen & Riding 1992

1776 *Amphorulacysta? dodekovae* (Zotto et al. 1987) Williams & Fensome 2016

1777 *Amphorulacysta metaelliptica* (Dodekova 1969) Williams & Fensome 2016

- 1778 *Arkellea teichophera* (Sarjeant 1961) Below 1990
- 1779 *Atopodinium haromense* Thomas & Cox 1988
- 1780 *Chytroeisphaeridia cerastes* Davey 1979
- 1781 *Chytroeisphaeridia hyalina* (Raynaud 1978) Lentin & Williams 1981
- 1782 *Corculodinium inaffectum* (Drugg 1978) Courtinat 2000
- 1783 *Coronifera oceanica* Cookson & Eisenack 1958
- 1784 *Cribroperidinium? edwardsii* (Cookson & Eisenack 1958) Davey 1969
- 1785 *Cribroperidinium globatum* (Gitmez & Sarjeant 1972) Helenes 1984
- 1786 *Cribroperidinium? longicorne* (Downie 1957) Lentin & Williams 1985
- 1787 *Ctenidodinium continuum* Gocht 1970
- 1788 *Ctenidodinium elegantulum* Millioud 1969
- 1789 *Ctenidodinium ornatum* (Eisenack 1935) Deflandre 1938
- 1790 *Ctenidodinium sellwoodii* (Sarjeant 1975) Stover & Evitt 1978
- 1791 *Dapcodinium priscum* Evitt 1961
- 1792 *Diacanthum hollisteri* Habib 1972
- 1793 *Dichadogonyaulax bensonii* Monteil 1992
- 1794 *Dingodinium tuberosum* (Gitmez 1970) Fisher & Riley 1980
- 1795 *Dissiliodinium giganteum* Feist-Burkhardt 1990
- 1796 *Endoscrinium galeritum* (Deflandre 1938) Vozzhennikova 1967
- 1797 *Endoscrinium luridum* (Deflandre 1938) Gocht 1970
- 1798 *Evansia deflandrei* (Wolfard & Van Erve 1981) Below 1990
- 1799 *Glossodinium dimorphum* Ioannides et al. 1977
- 1800 *Gochteodinia villosa* (Vozzhennikova 1967) Norris 1978 subsp. *villosa* autonym
- 1801 *Gonyaulacysta centriconnata* Riding 1983
- 1802 *Gonyaulacysta eisenackii* (Deflandre 1938) Górká 1965
- 1803 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965
- 1804 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965 subsp. *adecta* Sarjeant
- 1805 1982
- 1806 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965 subsp. *adecta* Sarjeant
- 1807 1982 var. *longicornis* (Deflandre 1938) Downie & Sarjeant 1965
- 1808 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965 subsp. *jurassica* autonym
- 1809 *Heibergella asymmetrica* Bujak & Fisher 1976
- 1810 *Isthmocystis distincta* Duxbury 1979
- 1811 *Kalyptea diceras* Cookson & Eisenack 1960

- 1812 *Kleithriasphaeridium corrugatum* Davey 1974
1813 *Kleithriasphaeridium fasciatum* (Davey & Williams 1966) Davey 1974
1814 *Lagenorhysis delicatula* (Duxbury 1977) Duxbury 1979
1815 *Leptodinium subtile* Klement 1960
1816 *Liasidium variabile* Drugg 1978
1817 *Liesbergia liesbergensis* Berger 1986
1818 *Luehndea spinosa* Morgenroth 1970
1819 *Mancodinium semitabulatum* Morgenroth 1970
1820 *Maturodinium inornatum* Morgenroth 1970
1821 *Moesiodinium raileanui* Antonescu 1974
1822 *Muderongia longicornis* Monteil 1991
1823 *Muderongia simplex* Alberti 1961
1824 *Nannoceratopsis dictyambonis* Riding 1984
1825 *Nannoceratopsis gracilis* Alberti 1961
1826 *Nannoceratopsis pellucida* Deflandre 1938
1827 *Nannoceratopsis plegas* Drugg 1978
1828 *Nannoceratopsis raunsgaardii* Poulsen 1996
1829 *Nannoceratopsis senex* van Helden 1977
1830 *Nannoceratopsis spiculata* Stover 1966
1831 *Nannoceratopsis symmetrica* Bucefalo Palliani & Riding 2000
1832 *Nannoceratopsis triceris* Drugg 1978
1833 *Noricysta fimbriata* Bujak & Fisher 1976
1834 *Oligosphaeridium complex* (White 1842) Davey & Williams 1966
1835 *Ovalicysta hiata* Bjaerke 1980
1836 *Paragonyaulacysta? borealis* (Brideaux & Fisher 1976) Stover & Evitt 1978
1837 *Pareodinia ceratophora* Deflandre 1947
1838 *Pareodinia halosa* (Filatoff 1975) Prauss 1989
1839 *Pareodinia prolongata* Sarjeant 1959
1840 *Pareodinia? pseudochytrioeides* (Below 1987) Lentin & Williams 1989
1841 *Parvocysta bullula* Bjaerke 1980
1842 *Parvocysta nasuta* Bjaerke 1980
1843 *Phallocysta elongata* (Beju 1971) Riding 1994
1844 *Phallocysta eumekes* Dörhöfer & Davies 1980
1845 *Phoberocysta neocomica* (Gocht 1957) Millioud 1969

- 1846 *Phoberocysta tabulata* Raynaud 1978
- 1847 *Prolixosphaeridium anasillum* Erkmen & Sarjeant 1980
- 1848 *Pseudoceratium pelliiferum* Gocht 1957
- 1849 *Rhaetogonyaulax arctica* (Wiggins 1973) Stover & Evitt 1978
- 1850 *Rhaetogonyaulax dilatata* (Wiggins 1973) Stover & Evitt 1978
- 1851 *Rhaetogonyaulax rhaetica* (Sarjeant 1963) Loeblich Jr. & Loeblich III 1968
- 1852 *Rhaetogonyaulax wigginsii* (Stover & Helby 1987) Lentin & Williams 1989
- 1853 *Rhynchodiniopsis cladophora* (Deflandre 1938) Below 1981
- 1854 *Rotosphaeropsis thule* (Davey 1982) Riding & Davey 1989
- 1855 *Sahulidinium ottii* Stover & Helby 1987
- 1856 *Scriniocassis priscus* (Gocht 1979) Below 1990
- 1857 *Scriniocassis weberi* Gocht 1964
- 1858 *Scriniodinium campanula* Gocht 1959
- 1859 *Scriniodinium crystallinum* (Deflandre 1938) Klement 1960
- 1860 *Scriniodinium? dictyophorum* (Deflandre 1938 ex Sarjeant 1967) Brenner 1988
- 1861 *Scriniodinium pharo* (Duxbury 1977) Davey 1982
- 1862 *Senoniasphaera jurassica* (Gitmez & Sarjeant 1972) Lentin & Williams 1976
- 1863 *Sentusidinium explanatum* (Bujak in Bujak et al. 1980) Wood et al. 2016
- 1864 *Sirmiodinium grossii* Alberti 1961
- 1865 *Spiculodinium neptuni* (Eisenack 1958) Duxbury 2018
- 1866 *Spiniferites ramosus* (Ehrenberg 1837) Mantell 1854
- 1867 *Stanfordella? cretacea* (Neale & Sarjeant 1962) Helenes & Lucas-Clark 1997
- 1868 *Stephanelytron redcliffense* Sarjeant 1961
- 1869 *Suessia swabiana* Morbey 1975
- 1870 *Susadinium faustum* (Bjaerke 1980) Lentin & Williams 1985
- 1871 *Susadinium scrofoides* Dörhöfer & Davies 1980
- 1872 *Systematophora areolata* Klement 1960
- 1873 *Systematophora penicillata* (Ehrenberg 1843 ex Ehrenberg 1854) Sarjeant 1980
- 1874 *Tehamadinium evittii* (Dodekova 1969) Jan du Chêne et al. 1986
- 1875 *Tenua anaphrissa* (Sarjeant 1966) Benedek 1972
- 1876 *Trichodinium castanea* Deflandre 1935 ex Clarke & Verdier 1967
- 1877 *Trichodinium scarburghense* (Sarjeant 1964) Williams et al. 1993
- 1878 *Tubotuberella apatela* (Cookson & Eisenack 1960) Ioannides et al. 1977
- 1879 *Valensiella ovulum* (Deflandre 1947) Eisenack 1963

- 1880 *Valvaeodinium armatum* Morgenroth 1970
- 1881 *Valvaeodinium koessenium* (Morbey 1975) Below 1987
- 1882 *Valvaeodinium spinosum* (Fenton et al. 1980) Below 1987
- 1883 *Wallodinium cylindricum* (Habib 1970) Duxbury 1983
- 1884 *Wanaea fimbriata* Sarjeant 1961
- 1885 *Wanaea thysanota* Woollam 1982
- 1886 *Wanaea verrucosa* Riding & Helby 2001
- 1887
- 1888 **Pollen and Spores:**
- 1889 *Classopollis classoides* Pflug 1953
- 1890 *Klukisporites variegatus* Couper 1958
- 1891