

1 Dinoflagellate cyst biostratigraphy of the Patasar Shale Member (Upper Jurassic) of the Wagad
2 Uplift, Kachchh, Gujarat, western India

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14 ABSTRACT

15 Two samples were analysed from the lowermost Patasar Shale Member of the Patasar Tank
16 section on the western margin of the Wagad Uplift in the Kachchh Basin of Gujarat, western
17 India. One of the samples produced an abundant, diverse and well-preserved palynobiota. The
18 other sample produced a significantly sparser association, but of similar character. The overall
19 assemblage is dominated by relatively long-ranging Jurassic gymnospermous pollen grains, but
20 also includes dinoflagellate cysts of definite Gondwanan affinity. The dinoflagellate cysts are
21 confidently correlated to the Australian *Dingodinium swanense* Interval Zone, which is of
22 Kimmeridgian (Late Jurassic) age. This indicates that the Australasian Jurassic dinoflagellate
23 cyst biostratigraphical scheme can be applied in western India, and probably throughout the
24 Indian subcontinent. Due to evidence from ammonites and calcareous nannofossils in the Patasar
25 Shale Member, allied with other biostratigraphical evidence from New Zealand and Papua New
26 Guinea, the age of the *Dingodinium swanense* Interval Zone is reinterpreted as being of Early
27 Kimmeridgian age. The *Dingodinium swanense* Interval Zone was previously assigned to the
28 Early to Late Kimmeridgian. Its refinement and revision to an older age is entirely consistent
29 with recent reassessments of these dinoflagellate cyst biozones.

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31 Keywords: biostratigraphy; dinoflagellate cysts; Late Jurassic; pollen and spores; provincialism;
32 Gujarat, western India

33

34 1. Introduction

35 In this study we develop an integrated biostratigraphy of the Patasar Shale Member in the
36 Wagad Uplift which is part of the Great Rann of Kachchh (or Kachchh Basin) of Gujarat,
37 western India (Fig. 1). The Patasar Shale Member is of Late Jurassic age, and is thought to be
38 close to the Oxfordian–Kimmeridgian transition based on its macrofaunas (Fig. 2; Pandey et al.
39 2012). The present work is based on a study of the palynology of two samples from the
40 lowermost Patasar Shale Member, with emphasis on marine palynomorphs. The resultant data
41 are integrated with existing interpretations based on ammonites and calcareous nannofossils in
42 order to achieve a comprehensive and refined biostratigraphical framework for these regionally
43 important strata. It was anticipated that the Patasar Shale Member would yield palynobiotas of
44 Gondwanan (Australasian) affinity. However little is known regarding the palaeobiogeographical
45 relationships of Jurassic dinoflagellate cysts from the Indian subcontinent. The north-central part
46 of Gondwana once occupied by India is a key area for understanding regional correlations and
47 shelfal marine connections within the Gondwanan supercontinent.

48

49 2. Regional geological background

50 The Wagad Uplift of Gujarat, western India, forms part of the Great Rann of Kachchh (or
51 Kachchh Basin). It is a pericratonic depocentre formed after continental rifting between Africa
52 and India during the Late Triassic (Biswas, 1982; 1991; 2005). It contains an extensive Mesozoic
53 succession comprising Upper Triassic terrestrially derived strata, Lower to Upper Jurassic
54 (Pliensbachian–Tithonian) marine sedimentary rocks and Lower Cretaceous (Berriasian–Albian)
55 marine to paralic beds. Most of the Jurassic strata, and some of the Lower Cretaceous intervals,
56 are richly fossiliferous, the faunas including key zonal ammonites. Bordering the Kachchh region
57 to the southwest is the Arabian Sea, and to the northeast the salt marshes of the Great Rann of
58 Kachchh (Fig. 1). The Deccan Traps occupy the southern part of the Kachchh region, and domal
59 structures with a west–east trending Mesozoic outcrop belt are present in the northern part. The
60 Wagad Uplift lies in the east of the Kachchh Basin; it is an elongate anticlinal dome with

61 Jurassic rocks of Callovian to Kimmeridgian age in its core (Fig. 1; Biswas, 1977). The various
62 uplifts and domes, including the area known as Kachchh Mainland, form 'islands' separated by
63 plains covered in blown sand (Fig. 1).

64

65 3. The geology of the Wagad Uplift

66 The Wagad Uplift is situated in the easternmost part of the Kachchh Basin (Fig. 1; Biswas
67 and Deshpande, 1970). A high angle fault with east-west strike separates the Wagad uplift in the
68 north from southern Kachchh. Extensive exposures of Middle Jurassic (Callovian) to Lower
69 Cretaceous strata are present in the Wagad Uplift (Biswas, 1971; Deshpande, 1972; Deshpande
70 and Merh, 1980). While the Oxfordian to Kimmeridgian succession is strongly condensed, with
71 stratigraphical gaps, in Kachchh (Fig. 2; Alberti et al., 2013), the Wagad Uplift preserves a
72 relatively continuous Oxfordian–Kimmeridgian succession of shallow-water sedimentary rocks.
73 It is therefore ideally suited for the present integrated biostratigraphical study (Fig. 2).

74 The first Mesozoic lithostratigraphical framework of the Wagad Uplift succession was by
75 Deshpande (1972). This reconnaissance research was followed by, for example, Biswas (1977),
76 Pandey et al. (2012) and Fürsich et al. (2013). Lithostratigraphically, the Callovian to
77 Kimmeridgian sedimentary rocks have been placed into the Washtawa, Kanthkot and Gamdau
78 formations (Fig. 2). The uppermost Washtawa Formation is the Kanthkot Ammonite Bed of the
79 Nara Shale Member. Overlying the Washtawa Formation is the Kanthkot Formation, which is
80 subdivided into three members; the Patasar Shale Member is the lowermost of these units. The
81 Gamdau Formation is undivided (Fig. 2; Fürsich et al., 2013).

82 The Patasar Shale Member, is the subject of this work. It comprises the lowermost part of the
83 Kanthkot Member of the lower Wagad Sandstone of Biswas (1977) and conformably overlies the
84 Kanthkot Ammonite Bed. The latter is richly fossiliferous and forms the uppermost part of the
85 Nara Shale Member of the Washtawa Formation. The Kanthkot Ammonite Bed is considered to
86 be coeval with the Dhosa Conglomerate Bed, an ammonite-rich condensed horizon which is
87 present throughout the Kachchh Mainland (Fig. 2).

88 The ammonite biostratigraphy of the Washtawa, Kanthkot and Gamdau formations has been
89 significantly refined in recent decades (e.g. Krishna et al., 1995; 1998; 2009a,b,c; Some and
90 Bardhan, 2005; Pandey et al., 2012; 2013a,b). By contrast, data on calcareous nannofossils and

91 dinoflagellate cysts from the Jurassic of the Kachchh Basin are relatively sparse (Krishna et al.,
92 1983; Jain et al., 1984; 1986; Kumar, 1986a,b,c; Rai, 2003; Saxena and Jafar, 2008; Rai et al.,
93 2015).

94 A single specimen of the ammonite genus *Erymnoceras* from an ooidal ferruginous marl
95 band, presumed to be within the Nara Shale Member of the Washtawa Formation, was assigned
96 to the Middle Callovian by Some and Bardhan (2005). This horizon is within the lower part of
97 the Nara Shale Member, underlying the Kanthkot Ammonite Bed (Fig. 2). Furthermore, a single,
98 poorly preserved fragment of an ammonite assigned to *Perisphinctes (Dichotomoceras)* cf.
99 *besairiei* was reported by Pandey et al. (2012) from the lowermost Patasar Shale Member. As a
100 result of the discovery of this biostratigraphically significant form, this part of the unit has been
101 tentatively dated as Late Oxfordian (Bifurcatus Zone, Grossouvrei Subzone).

102

103 4. Material and methods

104 The two samples from the Patasar Shale Member described in this study, PTS 1A and PTS 1,
105 were collected from the lowermost part of the scarp section exposed near Patasar Tank (i.e.
106 reservoir), c. 2 km east of Kanthkot village at N 23° 29' 48.4'', E 70° 28' 01.7'' (Fig. 1; Table 1).
107 This section is a relatively monotonous shale succession (Rai et al., 2015, fig. 3D). Of the 13
108 samples collected from this unit, only the lowermost two have yielded well-preserved
109 palynomorphs (Tables 1, 2). Sample PTS 1A is 15 cm stratigraphically below PTS 1. One of the
110 overlying eleven samples (PT6) yielded very rare and poorly preserved dinoflagellate cysts,
111 which are consistently indeterminate. Samples from the remaining ten horizons proved barren of
112 palynomorphs (Table 1). The generally poor recovery of palynomorphs throughout most of this
113 succession may be probably due to the effects of long-term weathering in a subtropical climate.

114 The samples were all processed using standard palynological techniques (Wood et al. 1996)
115 at the Birbal Sahni Institute of Palaeosciences (BSIP). After demineralisation using hydrochloric
116 acid and hydrofluoric acid, the neutralised residue was macerated for several minutes with dilute
117 nitric acid and then washed using distilled water onto a 15 µm sieve. The concentrated residue
118 was then mixed with polyvinyl alcohol and mounted onto microscope slides using Canada
119 Balsam as a mounting medium. Study and photography was carried out with a Olympus BH2
120 photomicroscope. The samples, prepared residues, microscope slides (BSIP numbers 15629–

121 15638) and figured specimens are all curated in the BSIP collections housed at 53 University
122 Road, Lucknow 226 007, India.

123

124 5. Palynology

125 The present study documents the dinoflagellate cyst assemblages from the lowermost part of
126 the Patasar Shale Member at its type section at Patasar Tank. This association is compared with
127 calcareous nannofossil assemblages in samples from the overlying beds (Table 1; Rai et al.,
128 2015, fig 3D). The ammonite, calcareous nannofossil and dinoflagellate cyst data from various
129 horizons in the Wagad Uplift are integrated to provide greater and more precise
130 biostratigraphical resolution. The dinoflagellate cysts are of Australasian affinity, and the present
131 study has provided a more refined age for the *Dingodinium swanense* Interval Zone of Helby et
132 al. (1987).

133 The two samples studied from the lower Patasar Shale Member of the Patasar Tank section
134 (PTS 1A and PTS 1) yielded well-preserved palynomorph assemblages. However, sample PTS 1
135 proved markedly more abundant and diverse than PTS 1A (Table 2). Both samples are
136 dominated by terrestrially derived forms, dominantly gymnospermous pollen. These mainly
137 comprise *Araucariacites* spp., *Callialasporites dampieri*, *Callialasporites turbatus* and
138 undifferentiated bisaccate pollen, with lesser proportions of *Classopollis* spp., *Perinopollenites*
139 *elatoides* and indeterminate forms. Pteridophyte spores are extremely rare, but are represented by
140 occasional smooth forms assigned to *Cyathidites* (Table 2).

141 Dinoflagellate cysts are subordinate in proportions to pollen and spores, however, they are
142 much more diverse. In sample PTS 1, the assemblage is dominated by *Dingodinium jurassicum*,
143 together with significant proportions of *Gonyaulacysta jurassica* subsp. *jurassica*, indeterminate
144 forms, *Pareodinia* spp. (including *Pareodinia halosa*), *Scriniodinium dictyotum*, *Sentusidinium*
145 spp. and *Stiphrosphaeridium* spp. Also present are *Acanthaulax* sp., *Apteodinium* sp.,
146 *Chlamydophorella wallala*, *Chlamydophorella* spp., *Egmontodinium polyplacophorum*,
147 *Endoscrinium luridum*, *Glossodinium dimorphum*, *Hadriana cinctum*, indeterminate chorate
148 forms, *Indodinium khariense*, *Leptodinium eumorphum*, *Mendicodinium granulatum*,
149 *Perisseiasphaeridium?* sp., *Prolixosphaeridium parvispinum*, *Rhynchodiniopsis cladophora*,
150 *Rigaudella aemula*, *Scriniodinium crystallinum*, *Scriniodinium inritibile*, *Trichodinium* sp. and

151 *Tubotuberella apatela*. The miscellaneous microplankton consist largely of foraminiferal test
152 linings and very rare scolecodonts; no acritarchs were recorded (Table 2).

153 The kerogen macerals are dominated by palynomorphs (sample PTS 1) and plant fragments
154 (sample PTS 1A). Wood is consistently present in significant proportions, and amorphous
155 organic material is extremely sparse (Table 2). The dominance of gymnospermous pollen with
156 relatively diverse dinoflagellate cysts and abundant plant tissue and wood is consistent with
157 deposition in an open marine, offshore shelfal environment.

158 Despite the disparity in diversity, the overall similarity of the palynobiotas in the two
159 samples confirm the field evidence that they are from the same genetic sedimentary unit (Table
160 2; Rai et al., 2015, fig. 3D). The palynomorph associations recorded in the two samples are fully
161 documented in Table 2, and selected dinoflagellate cyst specimens are illustrated in Plates I and
162 II. The dinoflagellate cysts recorded in this study, at and below the species level and with their
163 author citations, are listed in Appendix 1.

164

165 6. Palynomorph biostratigraphy

166 Because the pollen assemblages are dominated by araucarian forms, including
167 *Callialasporites dampieri*, the material studied is referable to the *Callialasporites dampieri*
168 Superzone of Helby et al. (1987) of Hettangian to Kimmeridgian age. The relatively sparse
169 occurrences of *Classopollis* spp. and *Perinopollenites elatoides* support this assertion (Helby et
170 al., 1987, fig. 13). The low-diversity association, especially of pteridophyte spores, prevents a
171 more refined biostratigraphical breakdown based on terrestrially derived palynomorphs.

172 The two dinoflagellate cyst associations are typical of the Late Jurassic of Gondwana. Forms
173 such as *Dingodinium jurassicum*, *Leptodinium eumorphum*, *Rigaudella aemula*, *Scriniodinium*
174 *crystallinum* and *Stiphrosphaeridium anthophorum/dictyophorum* are characteristic of the
175 Oxfordian to Tithonian of Australasia (Helby et al., 1987; Davey, 1988). *Egmontodinium*
176 *polyplacophorum* and *Scriniodinium inritibile* have not been previously reported from the
177 Gondwanan Realm. In Europe these species are typical of the Tithonian and Middle Oxfordian to
178 Tithonian respectively (Woollam and Riding, 1983; Riding, 1984). Somewhat unusually, the
179 typically Late Jurassic dinoflagellate cysts *Gonyaulacysta ceratophora* and *Nannoceratopsis*
180 *pellucida* were not encountered.

181 The key marker dinoflagellate cysts are *Chlamydothorella wallala*, *Gonyaulacysta jurassica*
182 subsp. *jurassica*, *Hadriana cinctum*, *Indodinium khariense*, *Rhynchodiniopsis cladophora* and
183 *Stiphrosphaeridium dictyophorum*. These indicate a correlation to the *Dingodinium swanense*
184 Interval Zone of Helby et al. (1987), which was deemed to be of Early to Late Kimmeridgian age
185 by Riding et al. (2010). The range tops of *Gonyaulacysta jurassica* subsp. *jurassica* and
186 *Rhynchodiniopsis cladophora* are of Middle Kimmeridgian age in Australia (Riding et al., 2010).
187 However, Davey (1988) reported the youngest consistent occurrence of *Gonyaulacysta jurassica*
188 subsp. *jurassica* as Late Kimmeridgian in Papua New Guinea. The inceptions of *Hadriana*
189 *cinctum* and *Stiphrosphaeridium dictyophorum* in Australia are also within the Middle
190 Kimmeridgian (Riding and Helby, 2001a; Riding et al., 2010). The summary range chart of
191 Riding et al. (2010, fig. 12) indicates a minor intra-Kimmeridgian stratigraphical gap between
192 the apparent extinctions of *Gonyaulacysta jurassica* subsp. *jurassica* and *Rhynchodiniopsis*
193 *cladophora*, and the range bases of *Hadriana cinctum* and *Stiphrosphaeridium dictyophorum*,
194 but this hiatus is not considered to be biostratigraphically significant. To summarise, the virtually
195 overlapping ranges of *Gonyaulacysta jurassica* subsp. *jurassica* and *Rhynchodiniopsis*
196 *cladophora*, with *Hadriana cinctum* and *Stiphrosphaeridium dictyophorum* is indicative of a
197 Middle Kimmeridgian age by comparison with Australian assemblages.

198 The occurrences of *Chlamydothorella wallala* and *Indodinium khariense* are entirely
199 consistent with this age assessment. The presence of *Chlamydothorella wallala* means that the
200 samples are no younger than latest Kimmeridgian; the range top of this species coincides with
201 the Kimmeridgian–Tithonian transition in Australia (Riding et al. 2010). The range of
202 *Indodinium khariense* is Middle Oxfordian to earliest Tithonian (Helby et al., 1988; Riding and
203 Helby, 2001a; Riding et al., 2010). Furthermore, the absence of the distinctive genus *Wanaea*
204 indicates that the samples are younger than Early Kimmeridgian (Riding and Helby, 2001b;
205 Riding et al., 2010). The lack of species such as *Herendeenia pisciformis*, *Komewuia glabra* and
206 *Omatia montgomeryi* preclude a Tithonian age (Riding et al., 2010).

207 The occurrence of *Glossodinium dimorphum* in this material is interesting. In Australia and
208 Papua New Guinea, this distinctive species ranges from the Callovian to the Middle Oxfordian
209 (Helby et al., 1987; Davey, 1988; Riding et al., 2010). By contrast in Africa and Europe, it is
210 present from the Middle Oxfordian to the Tithonian, and is especially prominent in the
211 Kimmeridgian (Woollam and Riding, 1983; Msaky, 2011). *Dingodinium jurassicum* is the

212 dominant taxon, but the samples are not assigned to the Middle Tithonian *Dingodinium*
213 *jurassicum* Opper Zone of Helby et al. (1987) because of the presence of key markers such as
214 *Hadriana cinctum* and *Indodinium khariense*.

215

216 7. Integrated biostratigraphy

217 The biostratigraphy of the Wagad Uplift in the Kachchh Basin, summarised in Fig. 3, was
218 discussed in detail by Rai et al. (2015). The Washatwa Formation of Oxfordian age underlies the
219 Kanthkot Formation, which spans the latest Oxfordian to Kimmeridgian, according to ammonite
220 biostratigraphy (Fig. 2). The Nara Shale Member is the youngest unit of the Washatwa
221 Formation, and the uppermost unit is the Kanthkot Ammonite Bed. The two samples analysed in
222 the present study are from the Patasar Shale Member, the lowermost member of the Kanthkot
223 Formation (Fig. 2; Table 1).

224 The integrated biostratigraphy of the Nara Shale and Patasar Shale members is illustrated
225 in Figs. 3 and 4. Calcareous nannofossils indicative of the Early and Middle Oxfordian have
226 been recorded from the Nara Shale Member, below the Kanthkot Ammonite Bed (Rai et al.,
227 2015). The Kanthkot Ammonite Bed is comprised of ferruginous, sandy strata with occasional
228 shales that have yielded relatively abundant ammonites indicative of the Middle to Late
229 Oxfordian (Pandey et al., 2012) and calcareous nannofossils characteristic of the Middle
230 Oxfordian (Rai et al., 2015). Therefore it appears entirely probable that the entire Nara Shale
231 Member is Middle Oxfordian in age (Fig. 4).

232 In the overlying lowermost part of the Patasar Shale Member, a single, poorly-preserved
233 Late Oxfordian ammonite has been recovered (Pandey et al., 2012). Rai et al. (2015) reported a
234 mixed assemblage of Middle–Late Oxfordian and Early Kimmeridgian calcareous nannofossils
235 from the Patasar Shale Member immediately above the two samples studied here. These authors
236 commented that the Middle and Late Oxfordian markers were probably reworked, and thus the
237 calcareous nannofossil assemblages are Early Kimmeridgian in age. Samples PTS 1A and PTS 1
238 studied here have produced a Middle Kimmeridgian age based on correlations with Australian
239 assemblages (section 6). These unequivocally Kimmeridgian dinoflagellate cysts significantly
240 strengthen the case that the Oxfordian calcareous nannofossils in the Patasar Shale Member are
241 reworked.

242 Thus, the lowermost part of the Patasar Shale Member appears to be Late Oxfordian,
243 based on a single, fragmentary ammonite (Fig. 4). Above this, the part of the Patasar Shale
244 Member sampled for microfossils is of Early–Middle Kimmeridgian age. However, Middle
245 Kimmeridgian dinoflagellate cysts appear to underlie Early Kimmeridgian calcareous
246 nannofossils.

247 In order to rationalise the apparently incoherent calcareous nannofossil and dinoflagellate
248 cyst evidence in the Patasar Shale Member, the age of the *Dingodinium swanense* Interval Zone
249 of Helby et al. (1987) must be reexamined. This zone was assigned to the Kimmeridgian and the
250 Early to Late Kimmeridgian by Helby et al. (1987, p. 29; 2004) and Riding et al. (2010, p. 567)
251 respectively. However, the latter authors stated that the evidence for this correlation was
252 equivocal, and that the zone could possibly be as old as Late Oxfordian (Davey, 1988; 1999;
253 Francis and Westermann, 1993). The *Dingodinium swanense* Interval Zone dinoflagellate cyst
254 associations described herein are clearly intercalated between a Late Oxfordian ammonite and
255 Early Kimmeridgian calcareous nannofossils. This is, thus far, the best and most direct control
256 on the age of the *Dingodinium swanense* Interval Zone. Due to the ammonite in the lowermost
257 Patasar Shale Member, the *Dingodinium swanense* Interval Zone cannot be older than Late
258 Oxfordian (Fig. 4). Likewise, it apparently cannot be younger than Early Kimmeridgian due to
259 the calcareous nannofossils stratigraphically above the two samples examined herein. Hence, due
260 to the other independent evidence of an Early Kimmeridgian age for this zone based on
261 radiolaria (Aita and Grant-Mackie, 1992), molluscs (Francis and Westermann, 1993) and
262 calcareous nannofossils (Rai et al., 2015), the *Dingodinium swanense* Interval Zone is herein
263 interpreted to be of Early Kimmeridgian age (Fig. 4). This reinterpretation of the age of an
264 Australasian Jurassic dinoflagellate cyst zone such that it is older than it was first envisaged is a
265 consistent theme. It appears that many of these dinoflagellate cyst zones originally established by
266 Helby et al. (1987) are significantly older than was originally thought (Riding, 2003; Riding et
267 al., 2010; Mantle and Riding, 2012). This biostratigraphical synthesis is summarised in Figs. 3
268 and 4.

269

270 8. The palaeogeographical affinity of the dinoflagellate cyst assemblage

271 The pollen and spore associations encountered comprise cosmopolitan forms, but the
272 dinoflagellate cyst assemblage recovered is distinctly Gondwanan in affinity. The dominant

273 species, *Dingodinium jurassicum*, was endemic to Gondwana. Furthermore, species such as
274 *Chlamydomphorella wallala*, *Hadriana cinctum*, *Indodinium khariense* and *Leptodinium*
275 *eumorphum* were also confined to Australasia. Many of the species observed, such as
276 *Glossodinium dimorphum*, *Rigaudella aemula*, *Scriniodinium crystallinum* and *Tubotuberella*
277 *apatela*, are cosmopolitan (Woollam and Riding, 1983; Helby et al., 1987). Due to the Austral
278 nature of the associations reported here from the Kachchh Basin, it seems that dinoflagellate cyst
279 zonation schemes established in Australia and Papua New Guinea such as those by Helby et al.
280 (1987), Davey (1988; 1999) and Riding et al. (2010) are applicable in western India and
281 probably the entire Indian subcontinent. These schemes appear to have utility throughout
282 Gondwana with the exception of South America. Research on the Jurassic marine palynology of
283 the Neuquén Basin, Argentina, has indicated that the extreme west of Gondwana is characterised
284 by associations with far more affinity to the northern hemisphere (e.g. Hedlund and Beju 1976;
285 Harris, 1977; Quattrocchio and Volkheimer, 1990; Quattrocchio and Sarjeant, 1992;
286 Quattrocchio et al., 2007; Riding et al. 2011).

287

288 9. Conclusions

289 One of two samples (PTS 1) from the lowermost Patasar Shale Member of the Patasar Tank
290 section in the Kachchh Basin, western India yielded an abundant, moderately diverse and well-
291 preserved palynobiota including a dinoflagellate cyst association of Gondwanan affinity. By
292 contrast, the other sample examined (PTS 1A) produced a relatively poorly preserved and
293 markedly less diverse assemblage. However, both the samples studied are clearly referable to the
294 *Dingodinium swanense* Interval Zone of Australia, which is of Kimmeridgian age. This study
295 indicates that Australasian dinoflagellate cyst biozonations can be applied in western India and,
296 by extension, probably throughout the Indian subcontinent. Due to evidence from ammonites and
297 calcareous nannofossils in the Patasar Shale Member outlined by Rai et al. (2015), together with
298 other key evidence from Gondwana, the age of the *Dingodinium swanense* Interval Zone of
299 Australia is reinterpreted herein as being Early Kimmeridgian in age (Fig. 4). This biozone was
300 previously assigned to the Early to Late Kimmeridgian and this reassessment to an older age
301 follows a familiar recent trend (e.g. Riding et al., 2010).

302

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313

314 Appendix 1

315 The palynomorphs recorded and mentioned herein, at and below the species level, are listed in
316 this appendix with their respective author citations. They are listed alphabetically in two
317 groupings, dinoflagellate cysts, and pollen and spores. The Plate and photograph numbers of the
318 taxa which are figured herein are provided where appropriate. The references pertaining to the
319 dinoflagellate cysts can be found in Fensome and Williams (2004).

320

321 Dinoflagellate cysts:

322 *Chlamydomorphella wallala* Cookson & Eisenack 1960

323 *Dingodinium jurassicum* Cookson & Eisenack 1958

324 *Dingodinium swanense* Stover & Helby 1987

325 *Egmontodinium polyplacophorum* Gitmez & Sarjeant 1972

326 *Endoscrinium luridum* (Deflandre 1938) Gocht 1970

327 *Glossodinium dimorphum* Ioannides et al. 1977 (Plate II/1)

328 *Gonyaulacysta ceratophora* (Cookson & Eisenack 1960) Riding 2005

329 *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965 subsp. *jurassica* (autonym)
330 (Plate I/1, 2)

331 *Hadriana cinctum* Riding & Helby 2001

332 *Herendeenia pisciformis* (Cookson & Eisenack 1958) Wiggins 1969

333 *Indodinium khariense* Kumar 1986

334 *Komewuia glabra* Cookson & Eisenack 1960

- 335 *Leptodinium eumorphum* (Cookson & Eisenack 1960) Sarjeant in Davey et al. 1969
336 *Mendicodinium granulatum* Kumar 1986 (Plate I/11, 12)
337 *Nannoceratopsis pellucida* Deflandre 1938
338 *Omatia montgomeryi* Cookson & Eisenack 1958
339 *Pareodinia halosa* (Filatoff 1975) Prauss 1989
340 *Prolixosphaeridium parvispinum* (Deflandre 1937) Davey et al. 1969
341 *Rhynchodiniopsis cladophora* (Deflandre 1938) Below 1981
342 *Rigaudella aemula* (Deflandre 1938) Below 1982
343 *Scriniodinium crystallinum* (Deflandre 1938) Klement 1960
344 *Scriniodinium dictyotum* Cookson & Eisenack 1960
345 *Scriniodinium inritibile* Riley in Fisher & Riley 1980
346 *Stiphrosphaeridium anthophorum* (Cookson & Eisenack 1958) Lentin & Williams 1985
347 *Stiphrosphaeridium dictyophorum* (Cookson & Eisenack 1958) Lentin & Williams 1985
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541 Fig. 1. A geological map of the Kachchh Basin of western India (b - top) with an inset map of
542 India in the top left (a), and a schematic map of the Wagad Uplift, adapted from Deshpande and
543 Mehr (1980) (c – below).
544
545 Fig. 2. Lithostratigraphy and ammonite biostratigraphy of the Upper Jurassic (Oxfordian and
546 Kimmeridgian) strata of the Kachchh Basin, western India (adapted from Pandey et al., 2012).
547

548 Fig. 3. The integrated biostratigraphy of the uppermost Washtawa Formation and the lowermost
549 Kanthkot Formation in the western part of the Wagad Anticline, Kachchh Basin, western India.
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571
572 Fig. 4. The stratigraphical occurrences of the key marker ammonites, calcareous nannofossils
573 and dinoflagellate cysts in the Nara Shale Member, Kanthkot Ammonite Bed and Patasar Shale
574 Member near Kanthkot, Wagad Anticline, Kachchh Basin, western India.

575
576 Table 1. The 13 micropalaeontology (calcareous nannofossil and palynomorph) samples
577 collected from the Patasar Shale Member at the Patasar Tank section near Kanthkot, in the
578 central west part of the Wagad Uplift, western India (N 23° 29' 48.4'', E 70° 28' 01.7''; see Fig.

579 1; Rai et al., 2015, fig 3D). The distances from the base of the exposure (in metres) are given,
580 and the productivity of the samples indicated by an X. An ellipsis (...) indicates that the
581 respective microfossil group was not recorded in that sample.

582

583 Table 2. The quantitative occurrences of palynomorphs and kerogen macerals in samples PTS
584 1A and PTS 1 from the Patasar Shale Member of the Patasar Tank section of the Wagad
585 Anticline. The data are arranged into three major palynomorph groups, and the four principal
586 kerogen macerals are listed as approximate percentages. The data are presented as actual counted
587 specimens and percentages of the overall palynobiota for each sample. An 'X' indicates where a
588 palynomorph or a kerogen maceral is present in the respective sample, but was not encountered
589 in the count. An ellipsis (...) indicates that the respective palynomorph was not recorded in that
590 sample.

591

592 Plate I. Selected dinoflagellate cysts from sample PTS 1 in the lower part of the Patasar Shale
593 Member (Kanthkot Formation) of the Patasar Tank section in the western part of the Wagad
594 Anticline, near Kanthkot, western India. The scale bars all indicate 20 µm, and the locations of
595 the specimens on the BSIP slides are indicated by England Finder coordinates.

596 1, 2. *Gonyaulacysta jurassica* (Deflandre 1938) Norris & Sarjeant 1965 subsp. *jurassica*
597 (autonym). 1 - BSIP slide number 15633, Q37/1; 2 - BSIP slide number 15635, O32/3.

598 3. *Tubotuberella apatela* (Cookson & Eisenack 1960) Ioannides et al. 1977, BSIP slide number
599 15633, J70/2.

600 4. *Egmontodinium polyplacophorum* Gitmez & Sarjeant 1972, BSIP slide number 15629, X60/1.

601 5. *Endoscrinium luridum* (Deflandre 1938) Gocht 1970, BSIP slide number 15636, Q51.

602 6. *Pareodinia halosa* (Filatoff 1975) Prauss 1989, BSIP slide number 15629, Q61/3.

603 7, 8. *Scriniodinium dictyotum* Cookson & Eisenack 1960. 7 - BSIP slide number 15636, T52; 8 -
604 BSIP slide number 15636, V36.

605 9. *Indodinium khariense* Kumar 1986, BSIP slide number 15635, Q55.

606 10. *Chlamydophorella wallala* Cookson & Eisenack 1960, BSIP slide number 15630, E53/2.

607 11, 12. *Mendicodinium granulatum* Kumar 1986. 11 - BSIP slide number 15630, H58; 12 - BSIP
608 slide number 15635, E54.

609

610 Plate II. Selected dinoflagellate cysts from sample PTS 1 in the lower part of the Patasar Shale
611 Member (Kanthkot Formation) of the Patasar Tank section in the western part of the Wagad
612 Anticline, near Kanthkot, western India. The scale bars all indicate 20 μm , and the locations of
613 the specimens on the BSIP slides are indicated by England Finder coordinates.

- 614 1. *Glossodinium dimorphum* Ioannides et al. 1977, BSIP slide number 15635, J67/1.
- 615 2, 3. *Leptodinium eumorphum* (Cookson & Eisenack 1960) Sarjeant in Davey et al. 1969. 2 -
616 BSIP slide number 15630, T68; 3 - BSIP slide number 15633, S55/2.
- 617 4. *Rhynchodiniopsis cladophora* (Deflandre 1938) Below 1981, BSIP slide number 15633,
618 L47/4.
- 619 5, 6. *Dingodinium jurassicum* Cookson & Eisenack 1958. 5 - BSIP slide number 15634, M53; 6 -
620 BSIP slide number 15636, Z39.
- 621 7. *Prolixosphaeridium parvispinum* (Deflandre 1937) Davey et al. 1969, BSIP slide number
622 15632, O57/3.
- 623 8, 9. *Stiphrosphaeridium anthophorum* (Cookson & Eisenack 1958) Lentin & Williams 1985. 8 -
624 BSIP slide number 15638, W53/1; 9 - BSIP slide number 15634, Y61/1.
- 625 10, 11. *Hadriana cinctum* Riding & Helby 2001. 10 - BSIP slide number 15637, R58; 11 - BSIP
626 slide number 15638, S44.
- 627 12. *Rigaudella aemula* (Deflandre 1938) Below 1982, BSIP slide number 15631, N40/4.
- 628