

1 **Two deep marine oxygenation events during the Permian-Triassic boundary**  
2 **interval in South China: relationship with ocean circulation and marine**  
3 **primary productivity**

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15 *Keywords:* Permian-Triassic; South China; Oxygenation events; Ocean circulation; Marine  
16 productivity

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18 **1. Introduction**

19 The Permian-Triassic (P-T) mass extinction (*ca.* 251.9 Ma) is the largest of the “Big Five”  
20 extinction crises in both marine and terrestrial settings (Stevens, 1977; Kerr, 2001; Zhang et al.,  
21 2014; Zhou et al., 2021). Biotic species suffered a severe loss of about 90% of marine  
22 invertebrate and 80% of terrestrial vertebrate taxa (Smith and Ward, 2001; Lehrmann et al., 2003;

23 Retallack et al., 2003; Ward et al., 2005; Zhou et al., 2021). In the marine environment,  
24 extinctions displayed a preferential selectivity for biota with heavy calcification and limited  
25 elaboration of respiratory organs (Knoll et al., 1996; Erwin, 2006; Payne et al., 2007). On land,  
26 the terrestrial ecosystem collapse and devegetation led to an earliest Triassic “coal gap”  
27 accompanied by a dramatic turnover from typical Permian floras to those dominated by  
28 lycopsids and ferns or conifers (Smith and Ward, 2001; Retallack et al., 2003; Ward et al., 2005;  
29 Yin et al., 2007; Cui et al., 2017; Song et al., 2018; Zhou et al., 2021). There documented  
30 evidence that the extinctions occurred earlier on land than in the marine environments (Shen et  
31 al., 2011; Kaiho et al., 2016; Liu et al., 2017; Zhou et al., 2017; Xiao et al., 2018).

32 A number of causes have been proposed for the P-T marine mass extinction events,  
33 including global warming, sea-level fall, marine anoxia, salinity change, ocean acidification, or a  
34 combination of these (e.g., Stevens, 1977; Isozaki, 1997; Grice et al., 2005a; Hinojosa et al.,  
35 2012; Joachimski et al., 2012; Kershaw et al., 2012a; Yin et al., 2014; Baresel et al., 2017;  
36 Zhang et al., 2018a; Li et al., 2021; Zheng et al., 2021; Zhou et al., 2021; Wu et al., 2022). Many  
37 of these potential causes appear to be linked to the massive volcanism in the Siberian Traps  
38 Large Igneous Province (e.g., Song et al., 2013; Chen et al., 2014; Zhou et al., 2021).

39 The excellent exposures of continuous uppermost Permian through lowermost Triassic  
40 marine strata on the South China Block have long been a focus for detailed studies of the P-T  
41 mass extinction episode (e.g., Wignall et al., 1995; Isozaki et al., 2007; Yin et al., 2014; Saitoh  
42 and Isozaki, 2021). A detailed conodont biostratigraphy established in these sections provides a  
43 high-resolution biostratigraphic and correlation framework for these P-T studies in South China  
44 (e.g., Jiang et al., 2007; Sun et al., 2012; Yin et al., 2014; Zhang et al., 2014). These studies led

45 to the Global Stratotype Section and Point (GSSP) for the P-T boundary being placed at the  
46 Meishan section in the northeastern part of South China (e.g., Yin et al., 2014; Shen et al., 2015;  
47 Wu et al., 2022). Paleogeographical reconstructions place the South China Block in low latitudes  
48 ( $< 30^{\circ}\text{N}$ ) during this time interval, with its northern margin facing the Paleotethys ocean and the  
49 southern margin facing the Panthalassa Ocean (Muttoni et al., 2009; Yin et al., 2014; Xiao et al.,  
50 2018).

51 Episodes of marine anoxia are an important mechanism for killing marine organisms.  
52 Therefore, among the multitude of P-T studies in South China, investigations of marine redox  
53 conditions have been conducted by applying multiple methods to sedimentary sections that were  
54 deposited at different water depths and under various paleogeographic conditions (**see literature**  
55 **in Table. 1**). The array of intensive studies has resulted in important controversies on how to  
56 interpret the marine redox history through the P-T boundary interval (Bond and Wignall, 2010;  
57 Loope et al., 2013; Proemse et al., 2013; Xiao et al., 2018; Li et al., 2021; Xiang et al., 2021).  
58 Studies of P-T sections from different depositional settings and/or applying various redox  
59 proxies to the same P-T section can yield conflicting marine redox results and interpretations  
60 (e.g., Kershaw et al., 2012b; Lehrmann et al., 2015; Li et al., 2016; Huang et al., 2019; Xiang et  
61 al., 2020). The incongruent interpretations of marine redox variations heavily impede an overall  
62 understanding of environmental scenarios for the P-T mass extinction episodes.

63 In order to better understand the marine redox history through the P-T boundary interval and  
64 the end-Permian mass extinction events in South China, this study summarizes and reassesses a  
65 large body of marine redox studies (**Table 1**) spanning the conodont zones mainly from *Clarkina*  
66 *changxingensis* to *Isarcicella isarcica* (**Fig. S1**) in South China (Paleotethys). The Paleotethys

67 redox history is then compared with more extensive Panthalassa Ocean. The synthesis suggests  
68 there were two deep marine oxygenation events during the P-T boundary interval, and their  
69 occurrences are argued to be possibly related to changes in ocean circulation and marine primary  
70 productivity.

## 71

## 72 **2. Permian-Triassic stratigraphic records of marine redox variation in South China**

73 During the P-T transition, South China was an isolated continental block lying to the east of  
74 the supercontinent Pangea (Isozaki et al., 2007; Muttoni et al., 2009; Saitoh et al., 2014) (**Fig. 1**).  
75 Within the South China Block, a central, shallow-marine Yangtze Platform was bounded by the  
76 Kangdian Oldland on the east, by the Cathaysian Oldland and Yunkai Oldland on the west, by  
77 deep-water Northern Marginal Basin on the north and by the deep-water  
78 Hunan-Guizhou-Guangxi Basin on the south (Yin et al., 2014; Lei et al., 2017; Zhang et al.,  
79 2018b; Wu et al., 2022) (**Fig. 1**). The Northern Marginal Basin includes the northern slope of the  
80 Yangtze Platform and the deep-sea Qinling seaway of the Paleotethys (Feng et al., 1997; Yin et  
81 al., 2014). The Hunan-Guizhou-Guangxi Basin, also known as the Nanpanjiang Basin, contained  
82 many small and isolated shallow-water platforms surrounded by deep water that faced toward  
83 the Panthalassa Ocean (Feng et al., 1997; Yin et al., 2014). The suite of well-studied P-T marine  
84 sections are mainly concentrated into two realms: the northern part of the Yangtze Platform and  
85 adjacent Northern Marginal Basin; and the small isolated carbonate platforms and adjacent deep  
86 water troughs of the Hunan-Guizhou-Guangxi Basin (**Fig. 1**) (Lehrmann et al., 2003; Yin et al.,  
87 2014; Shen et al., 2015; Tian et al., 2015; Zhang et al., 2020; Wu et al., 2022). Previous P-T  
88 marine redox studies from these two realms provide a good opportunity to investigate P-T

89 marine redox variations at different water depths (shallow vs. deep) and different  
90 paleogeographic locations (**Fig. 1; Table 1**). As shown below, the inferred redox conditions vary  
91 from oxic to euxinic conditions, and the definition and characterization of oxic, upper dysoxic,  
92 low dysoxic, anoxic and euxinic conditions and facies follow those proposed by Bond and  
93 Wignall (2010) (**Table 2**).

94

## 95 **2.1. Northern Yangtze Platform Margin and the adjacent Northern Marginal Basin facing** 96 **the Paleotethys ocean**

97

### 98 *2.1.1. Basinal to slope P-T sections in the Northern Marginal Basin (Fig. 2)*

99 2.1.1.1. Xiaojiaba section, Guangyuan City, northern Sichuan Province (Wei et al., 2015; **Fig.**  
100 **S2**)

101 The Xiaojiaba section records a lower slope to basinal P-T deposition. The Xiaojiaba  
102 section is near the Chaotian section (detailed in 2.1.1.2), but was deposited at a slightly deeper  
103 water depth than the latter. The upper Permian Dalong Formation consists of lime mud-rich  
104 chert in its lower and middle parts, and of lime mudstone and siliceous or nodular limestone in  
105 its upper part. Volcanic tuff layers are abundant in the upper part, but only a few occur in the  
106 lower and middle parts. The Lower Triassic Feixianguan Formation consists of interbedded  
107 shale and lime mudstone. Many volcanic tuff layers are present near the base of the Feixianguan  
108 Formation, and no fossils are recorded in its lower part.

109 Based on detailed lithological correlations with the nearby Chaotian section where conodont  
110 and ammonoid biostratigraphy are well established, the mass extinction horizon in the Xiaojiaba

111 section is placed at the contact between the Dalong Formation and the Feixianguan Formation,  
112 and the P-T boundary is assigned to a shale layer about 1 meter higher.

113 Multiple redox proxies including  $\delta^{34}\text{S}_{\text{py}}$ ,  $\text{C}_{\text{org}}/\text{P}$ ,  $\text{Mo}_{\text{auth}}$ ,  $\text{U}_{\text{auth}}$  and the size distribution of  
114 framboidal pyrite, indicate that marine redox conditions changed from anoxic (occasionally  
115 euxinic) in the lower Dalong Formation to dysoxic/oxic conditions in the upper Dalong  
116 Formation through lower Feixianguan Formation, followed by a return to anoxic conditions. An  
117 oxygenation event (i.e., an increase in water column oxygen levels) occurred during the latest  
118 Permian (*C. taylorae* - *C. zhejiangensis* - *C. changxingensis* - *C. yini* conodont zones) on the  
119 basis of an increase in framboidal pyrite sizes and  $\delta^{34}\text{S}_{\text{py}}$ , and a decrease in  $\text{C}_{\text{org}}/\text{P}$ ,  $\text{Mo}_{\text{auth}}$  and  
120  $\text{U}_{\text{auth}}$  (Fig. S2). This oxygenation event co-occurred with a fall in primary productivity (indicated  
121 by a reduction in TOC, biogenic  $\text{SiO}_2$  and  $\text{Ba}_{\text{xs}}$  (excess Ba)), a sea-level fall (based on subaerial  
122 exposure), increased influx of fine-grained terrigenous clastics (based on increase in REE and  
123  $\text{Al}_2\text{O}_3$ , and lower  $\text{Eu}/\text{Eu}^*$ ) and elevated volcanic activity (inferred from the increased abundance  
124 of tuff layers) (Fig. S2).

125

126 2.1.1.2. Chaotian section, Guangyuan City, northern Sichuan Province (Isozaki et al., 2007; Ji et  
127 al., 2007; Saitoh et al., 2014, 2021; Saitoh and Isozaki, 2021; **Fig. S3**)

128 The Chaotian section was located in a lower slope to basinal setting during the P-T  
129 boundary interval. The upper Dalong Formation of latest Permian (late Changhsingian) age is  
130 frequently bioturbated and consists mainly of calcareous/cherty (lime)mudstone with micritic  
131 limestone at the top, and contains abundant ammonoids, conodonts and radiolarians (Figs 6 A,  
132 B). The uppermost Dalong Formation spans the *C. taylorae* -*C. zhejiangensis*

133 -*C. changxingensis* –*C. yini* conodont zones (Isozaki et al., 2007; Ji et al., 2007). The lower  
134 Triassic Feixianguan Formation is largely unbioturbated and consists of micrite limestone with a  
135 basal marl bed that is barren of fossils but contains abundant pyrites (Figs 6C, D) (Saitoh and  
136 Isozaki, 2021). The P-T mass extinction is placed at the contact between the Dalong Formation  
137 and the Feixianguan Formation. The conodont *Hindeodus parvus* is known from the base of  
138 micrite limestone in the lowermost Feixianguan Formation at the base of the Unit F (Isozaki et  
139 al., 2007). The overlying *Isarcicella* Zone was assigned based on the co-occurrence of the  
140 conodonts *H. parvus*, *I. isarcica* and *I. staeschei* (Isozaki et al., 2007; Ji et al., 2007).

141 In the lower-middle portion of the Dalong Formation, the low degree of bioturbation,  
142 decreased burrow size, absence of trace fossils, high TOC content and abundance of framboidal  
143 pyrite are suggestive of anoxic water conditions (Saitoh et al., 2014, 2021). In contrast, in the  
144 uppermost Dalong Formation (mainly the *C. taylorae* -*C. zhejiangensis* -*C. changxingensis yini*  
145 conodont zones), the common occurrence of burrows (Fig. 6A), increase in ichnofabric index,  
146 low TOC and relative rarity of framboidal pyrite indicate oxic to dysoxic water conditions  
147 immediately before and during the end-Permian mass extinction (Saitoh et al., 2014, 2021) (Fig.  
148 S3), suggesting an oxygenation event. This enhanced oxygenation co-occurred with a rapid  
149 sea-level fall. Above the level of the end-Permian mass extinction in the lowermost Feixianguan  
150 Formation (*H. parvus* and *Isarcicella* conodont zones), the absence of bioturbation and other  
151 trace fossils and the abundance of framboidal pyrites (Fig. 6C, D) suggest that anoxic water  
152 returned in the aftermath of the extinction (Saitoh and Isozaki, 2021).

153  
154 2.1.1.3. Shangsi section, Guangyuan City, northern Sichuan Province (Wignall et al., 1995;

155 Xiang et al., 2016; Shen et al., 2021; **Fig. S4**)

156 During the P-T boundary interval, the Shangsi section was deposited in a lower slope to  
157 basinal setting in shallower water depths than nearby Chaotian section that is described above  
158 (Wignall et al., 1995; Shen et al., 2021). The Dalong Formation of latest Permian  
159 (Changhsingian) age consists of micrite limestone that is organic-rich in the lower part and  
160 chert-rich in the upper part. The Feixianguan Formation of earliest Triassic (Induan) age consists  
161 mainly of thin-bedded alternations of micrite and argillaceous limestones. The P-T mass  
162 extinction horizon is placed within the uppermost part of the Dalong Formation (Wignall et al.,  
163 1995). The P-T boundary was assigned within the Bed 28 at the lowest occurrence of the  
164 conodont *H. parvus* (Wignall et al., 1995; Xiang et al., 2016; Shen et al., 2021).

165 Based on iron speciation proxies and redox-sensitive elements (Mo, V and U) (Xiang et al.,  
166 2016), the uppermost *C. wangi* through middle *C. subcarinata* conodont zones have been  
167 interpreted as times of euxinic bottom-water conditions. The upper *C. subcarinata* conodont  
168 zone, the *C. changxingensis* - *C. yini* - *C. meishanensis* Assemblage Zone and the *H.*  
169 *changxingensis* Zone were mainly periods of anoxic conditions. However, an oxygenation event  
170 was indicated by oxic water conditions occurring immediately below and during the  
171 end-Permian mass extinction interval (*C. meishanensis* and *H. changxingensis* conodont zones),  
172 which is supported by lower values of  $Fe_{HR}/Fe_T$ ,  $Fe_{Py}/Fe_{HR}$ ,  $Mo_{xs}$ ,  $U_{xs}$  and  $V_{xs}$  (Fig. S4). The  
173 redox proxies in the overlying *H. changxingensis*, *H. parvus* and *I. isarcica* conodont zones of  
174 the earliest Triassic indicate anoxic conditions with sporadic euxinic conditions. These redox  
175 variations are consistent with other sedimentary features (fossil diversity, bioturbation degree)  
176 and size distributions of framboidal pyrites (Wignall et al., 1995; Bond and Wignall, 2010). The



177 oxygenation event immediately below and during the end-Permian mass extinction horizon has  
178 been proposed to be related to a decrease in primary productivity at that time (Xiang et al.,  
179 2016).

180

181 2.1.1.4. Changtanhe section, Cili County, Hunan Province (He et al., 2012, 2013; **Fig. S5**)

182 The Changtanhe section was deposited in a marine slope setting during the P-T boundary  
183 interval. The Dalong Formation of latest Permian age consists mainly of laminated wackestone,  
184 with thin mudstone and shale beds becoming more common towards the top. Abundant and  
185 diverse fossils (including radiolarians, foraminifers, sponge spicules and echinoderms) are found  
186 in the Dalong Formation. The overlying Daye Formation of earliest Triassic age consists mainly  
187 of lime mudstone, marlstone and micrite limestone.

188 The end-Permian mass extinction horizon is placed at the top of the Dalong Formation  
189 where radiolarians, sponges and most of the foraminifers disappear. Marine redox conditions  
190 based on the size distribution of framboidal pyrite suggest lower dysoxic water conditions  
191 developed during the latest Permian, which shifted to be partly anoxic just below the P-T  
192 extinction horizon. The decreased oxygen level is accompanied with a decrease of the diversity  
193 of benthic organisms. Immediately after the extinction horizon, framboidal pyrites start to  
194 increase in size and even disappear in the interval of 0.4 to 0.48 m above the extinction boundary,  
195 thereby representing upper dysoxic to oxic water conditions and indicating a post-extinction  
196 oxygenation event (Fig. S5). Meanwhile, the TOC contents show a sharp decrease at this  
197 oxygenation event from lower dysoxic to upper dysoxic-oxic conditions across the mass  
198 extinction horizon (Fig. S5). This oxygenation event is also supported by the simultaneous

199 increase in abundance of fossils (ammonites, brachiopods, bivalves and microconchids). After  
200 the oxygenation event, the water redox conditions returned again to lower dysoxic conditions,  
201 and fossils became rare.

202

203 2.1.1.5. Ganxi section, Puyang County, Sichuan Province (Lei et al., 2017; Wu et al., 2022; **Fig.**  
204 **S6**)

205 The Ganxi section was deposited in an upper slope setting during the P-T boundary interval  
206 (Wu et al., 2022). The Dalong Formation of latest Permian mainly consists of lime mudstone in  
207 the lower part and of marly limestone with interbedded lime mudstone in the upper part. The  
208 Daye Formation of earliest Triassic consists mainly of lime mudstone. A few volcanic ash beds  
209 occur within the uppermost Dalong Formation and the lowermost Daye Formation.

210 Based on conodont biostratigraphy and correlation with the Meishan reference section, the  
211 mass extinction horizon is assigned to be at the base of the Daye Formation, and the P-T  
212 boundary is placed within Bed 237 in the lower Daye Formation (Lei et al., 2017). Marine redox  
213 conditions indicated by iron speciation and the Mo concentration data suggest mainly anoxic to  
214 euxinic conditions within the *C. wangi*, *C. changxingensis*, *C. yini* and *C. meishanensis*  
215 conodont zones (Fig. S6); and anoxic to oxic conditions prevailed within the *H. parvus* and *I.*  
216 *isarcica* conodont zones (Fig. S6). The  $\delta^{34}\text{S}_{\text{py}}$  shows a positive excursion at the increasing  
217 oxygen levels (the oxygenation event) during the *H. parvus* and *I. isarcica* conodont zones, and  
218 the oxygenation event is consistent with lower values of TOC, MOEF,  $\text{Fe}_{\text{HR}}/\text{Fe}_{\text{T}}$ , and  $\text{Fe}_{\text{Py}}/\text{Fe}_{\text{HR}}$   
219 (Fig. S6).

220

221 2.1.1.6. Meishan section, Changxing County, Zhejiang Province (Grice et al., 2005b; Isozaki et  
222 al., 2007; Huang et al., 2017; Li et al., 2016; **Fig. S7**)

223 The Meishan section was deposited in a marine upper slope setting during P-T boundary  
224 interval. The section, despite being highly condensed, was selected as the GSSP for the P-T  
225 boundary (Isozaki et al., 2007). The Changxing Formation of latest Permian consists of  
226 thin-bedded limestone, and the overlying Yinkeng Formation of earliest Triassic consists mainly  
227 of thin interbedded marlstone and mudstone (Huang et al., 2017). A few volcanic ash layers  
228 occur in the boundary interval. The mass extinction horizon occurs between Beds 24e and 25 (Li  
229 et al., 2016). The first occurrence of the conodont *H. parvus* in the middle of Bed 27 (base of  
230 27c) defines the P-T boundary at the GSSP.

231 Different marine redox proxies applied to this section have led to different interpretations.  
232 Euxinic conditions of the photic zone during the P-T transition have been inferred from  
233 biomarkers (Grice et al., 2005b; Summons et al., 2006). In contrast, the size distributions of  
234 framboidal pyrites and changes in fossil content within the P-T boundary interval suggest  
235 unstable marine redox conditions (Li et al., 2016). Abundant small pyrite framboids (4 to 6  $\mu\text{m}$   
236 in diameter) and the absence of benthos within the *C. taylorae* -*C. zhejiangensis* –  
237 *C. changxingensis* –*C. yini* conodont zone interval indicate mainly anoxic conditions that were  
238 periodically disrupted by short-term oxic conditions with rare framboidal pyrites during  
239 intrusions of turbidity currents (Li et al., 2016). Upper dysoxic to oxic conditions during most of  
240 the *H. parvus* conodont zone suggest an oxygenation event and are indicated by the absence of  
241 framboidal pyrites, abundant trace fossils and diverse biota (Li et al., 2016) (Fig. S7). A return to  
242 anoxic water conditions in the *Isarcicella* Zone is suggested by the reappearance of small (5  $\mu\text{m}$

243 in diameter) framboidal pyrites and the disappearance of trace fossils (Li et al., 2016).

244 Based on iron specification proxies, Xiang et al. (2020) suggested oxic to dysoxic water  
245 conditions in Beds 21-24d (corresponding to the *C. taylorae*, *C. zhejiangensis*,  
246 *C. changxingensis* and *C. yini* zones); anoxic water conditions with intermittently euxinia in  
247 Beds 24e to 28 (corresponding to the *H. parvus* and *I. staeschei* zones); and anoxic and  
248 ferruginous conditions in Beds 29 to 34 (corresponding to *Isarcicella* conodont zone) (Yin et al.,  
249 2014). At the level of the oxygenation event indicated by the absence of framboidal pyrites,  
250 abundant trace fossils and diverse biota, a decreasing trend of  $Fe_{HR}/Fe_T$  and  $Fe_{Py}/Fe_{HR}$  is  
251 observed and also indicates increasing oxygenation (Fig. S7).

252

### 253 2.1.2. Shallow-marine P-T sections in Northern Part of Yangtze Platform (**Fig. 3**)

254 2.1.2.1. Liangfengya section, Chongqing Municipality (Bond and Wignall, 2010; Li et al., 2021;  
255 **Fig. S8**)

256 The Liangfengya section records shallow-water, inter-reef platform carbonate deposition in the  
257 P-T boundary interval. The conodont biostratigraphy and carbon isotope correlations with the  
258 Meishan section suggest that the Liangfengya section is complete without an obvious hiatus that  
259 characterizes some shallow marine sections around the P-T boundary.

260 In the P-T boundary strata, the upper part of the Changxing Formation of latest Permian  
261 consists of fossiliferous limestone (Bed 1). The overlying lower part (Beds 2-8) of the  
262 Feixianguan Formation of earliest Triassic comprises alternating beds of shale and marlstone.  
263 The mass extinction horizon is recorded at Bed 8 by the disappearance of a fossil assemblage  
264 including mainly foraminifera, calcareous algae and Permian brachiopods. This bed is overlain

265 by fine-grained lithologies (micrite limestone and marlstone) in which the lower interval (Beds  
266 9-15) contains a low diversity of biota including microforaminifera, micrograstropods and  
267 ostracods, and the upper interval contains a few bivalves and abundant framboidal pyrites (Bed  
268 16-18). The lowest occurrence of conodont *H. parvus* in the middle of Bed 10 marks the P-T  
269 boundary.

270 No framboidal pyrite is found in the bioclastic limestone of the Changxing Formation (Bed  
271 1) or in the micrite limestones of the Feixianguan Formation (Beds 2 to 15) (Fig. S8). This  
272 suggests mainly oxic water conditions during the conodont zones of *C. changxingensis*, *C. yini*,  
273 *C. meishanensis*, *H. parvus* and *I. staeschei*. However, lower dysoxic to euxinic water conditions  
274 are inferred from abundant small framboidal pyrites (3 to 8  $\mu\text{m}$  in diameter) in the marlstone and  
275 lime mudstone of the higher Feixianguan Formation (Beds 16 to 17) of the *I. isarcica* Zone (Fig.  
276 S8). The redox variations are also supported by changes in the fossil diversity and in the degree  
277 of bioturbation at the same sequence.

278

279 2.1.2.2. Laolongdong section, Chongqing Municipality (Liao et al., 2010; Li et al., 2021; **Fig.**  
280 **S9**)

281 The Laolongdong section is a shallow-marine P-T section (Liao et al., 2010). At the P-T  
282 boundary succession, the bioclastic limestone of the upper Permian Changxing Formation is  
283 sharply overlain by microbialites with well-preserved microfabrics (Figs 6E–G) at the base of  
284 the Feixianguan Formation. The contact between the bioclastic limestone and the overlying  
285 microbialites represents the mass extinction horizon. Although this contact seems conformable,  
286 it should be noted that it is characterized by pressure solution and so the possibility of

287 depositional break or subaerial exposure exists at the contact (Liao et al., 2010). The  
288 microbialites are overlain by a laminated lime mudstone with few fossils. Based on  
289 carbon-isotope ( $\delta^{13}\text{C}_{\text{carb}}$ ) correlation with the Meishan section, the bioclastic limestone of the  
290 Changxing Formation belongs to *C. changxingensis* -*C. yini* conodont zones, the microbialite  
291 belongs to *H. parvus* -*I. staeschei* conodont zones, and the overlying lime mudstone  
292 (Feixianguan Formation) belongs to the *I. isarcica* conodont zone (Li et al., 2021).

293 In terms of marine redox proxies, the abundant and diverse fossils and the absence of  
294 framboidal pyrites indicate oxic conditions prevailed during the deposition of the bioclastic  
295 limestone of the Changxing Formation (Fig. S9). Abundant small framboidal pyrites (mean  
296 diameter of ca. 8  $\mu\text{m}$ ) and reduced biodiversity suggest lower dysoxic conditions for the  
297 microbialites (base of the Feixianguan Formation) (Fig. S9). Remarkably, shelly benthic fossils  
298 occur commonly within the microbialites, and are associated with small framboidal pyrites,  
299 raising the prospect that these pyrites might be forming within anoxic/dysoxic  
300 microenvironments within the microbialites. In the overlying mudstone (lower Feixianguan  
301 Formation), abundant small framboidal pyrites with a mean size of about 7.66  $\mu\text{m}$  suggest a  
302 continued oxygen depletion.

303  
304 2.1.2.3. Zaixia section, Chongyang County, Hubei Province (Liu et al., 2010; Chen et al., 2011;  
305 Adachi et al., 2017; **Fig. S10**)

306 The Zaixia section is a shallow-marine P-T section. The upper Permian (Changhsingian)  
307 bioclastic limestone is overlain by sparsely fossiliferous microbialites, then by lower Triassic  
308 oolitic and micrite limestone without fossils. The mass extinction horizon is placed at the contact

309 between the bioclastic limestone and overlying microbialites, but the contact records a  
310 depositional hiatus or erosive event of unknown duration. The appearance of the conodont *H.*  
311 *parvus* within the lower part of the microbialites suggests that the onset of microbialite  
312 formation may have been in the very latest Permian. Small (2.1 to 5.4  $\mu\text{m}$  in diameter)  
313 framboidal pyrites found within the microbialites indicate oxygen-deficient water conditions  
314 (Adachi et al., 2017). Alternatively, biomarker analyses (e.g., Pristane/Phytane,  
315 Trisnorneohopane/Trisnorhopane) suggest (Fig. S10): (i) no severe anoxic event in this  
316 shallow-marine section; (ii) a transient development of more oxidizing marine conditions just  
317 above the mass extinction horizon; and (iii) more reducing conditions or enhanced salinity at the  
318 upward transition from microbialites to oolitic and micrite limestone in the earliest Triassic  
319 (Chen et al., 2011).

320

321 2.1.2.4. Gaohua section, Cili County, Hunan Province (Forel et al., 2009; Loope et al., 2013;  
322 Wang et al., 2016; **Fig. S11**)

323 The Gaohua section is a P-T section deposited under shallow-marine environments near the  
324 edge of a carbonate platform. The uppermost Changxing Formation of latest Permian consists  
325 mainly of bioclastic limestone. The Daye Formation of early Triassic consists of a basal unit of  
326 microbialites followed by alternating beds of micrite and oolitic limestone. The mass extinction  
327 horizon is defined at the contact between the Changxing Formation and the Daye Formation.  
328 The P-T boundary, as defined by the first occurrence of conodont *H. parvus*, is within the  
329 microbialites at the base of the Daye Formation. In terms of biostratigraphy, the uppermost  
330 Changxing Formation belongs to *H. cf. inflatus* conodont zone, the microbialites in the basal

331 Daye Formation belong to *H. parvus* zone, and the beds above the microbialites belong to the *I.*  
332 *isarcica* Zone.

333 Several redox histories have been put forward for this P-T section. Loope et al. (2013)  
334 suggested that this section was deposited under fully oxic water conditions throughout the P-T  
335 boundary interval, as indicated by the negative Ce anomaly (Fig. S11). This is consistent with  
336 the interpretation by Forel et al. (2009), who suggested oxic water conditions that allowed  
337 persistent deposit-feeding guilds to flourish through the P-T boundary interval in the section.

338 However, this oxic water interpretation is inconsistent with a framboidal pyrite study by  
339 Wang et al. (2016). An absence of framboidal pyrite within the uppermost Changxing Formation  
340 and the basal part of the microbialites of the Daye Formation suggests oxic water conditions  
341 (Wang et al., 2016). However, framboidal pyrites are present in the overlying part of the  
342 microbialites, and their size distribution (mean size between 4 and 8  $\mu\text{m}$ ) suggests lower dysoxic  
343 depositional conditions (Fig. S11). Above the microbialites, the absence of small framboidal  
344 pyrites within the oolitic limestone suggesting oxic conditions, but their occurrence in the  
345 micritic limestone suggests lower dysoxic conditions (Wang et al., 2016).

346  
347 2.1.2.5. Panjiazhuang section, Changxing County, Zhejiang Province (Huang et al., 2019; **Fig.**  
348 **S12**)

349 The Panjiazhuang P-T section was deposited on a shallow-marine carbonate platform. The  
350 Changxing Formation of latest Permian consists of bioclastic limestone with diverse biota.  
351 Between the bioclastic limestone and overlying microbialites of the basal Daye Formation is a  
352 distinctive purple-red weathering crust, which is considered to be coeval with the P-T mass



353 extinction horizon. The P-T boundary is placed at the base of the microbialites where the  
354 conodont *H. parvus* firstly occurs. The microbialites are overlain by thin-bedded micrite  
355 limestones with oncoidal and shelly interlayers.

356 Oxidic water settings during the deposition of the upper Permian bioclastic limestone and the  
357 lower part of microbialites are indicated by the absence of framboidal pyrites (Fig. S12). In the  
358 upper part of the microbialites and in overlying micrite limestone, the occurrence of abundant  
359 framboidal pyrites (8 to 14  $\mu\text{m}$  in diameter) suggests dysoxic conditions (Fig. S12). Notably, no  
360 framboidal pyrites are found in the oncoidal and shelly interlayers within the micrite limestone.

361

## 362 **2.2. Hunan-Guizhou-Guangxi Basin facing Panthalassa Ocean**

363

### 364 *2.2.1. Deep-marine P-T sections in the basin (Fig. 4)*

365 2.2.1.1. Dongpan section, Nanning City, Guangxi Province (He et al., 2007; Zhang et al., 2018b;  
366 **Fig. S13**)

367 The Dongpan section was deposited in a deep-marine slope to basinal setting during the P-T  
368 boundary interval. The uppermost Talung Formation of latest Permian (late Changhsingian) is  
369 dominated by cherty (lime)mudstone in Beds 2 to 12. The overlying Bed 13 at the base of the  
370 Luolou Formation of earliest Triassic (early Induan) is dominated by silty (lime)mudstone.  
371 Based on sedimentological and geochemical correlations with the Meishan section, the P-T mass  
372 extinction horizon is assigned within Beds 7 and 8 in the uppermost Talung Formation and the  
373 P-T boundary is placed at the contact between Beds 12 and 13.

374 Redox-sensitive trace elements (Co, Cr, Cu and V) indicate oxygen restriction from Bed 2 to

375 the lower part of Bed 5 of the Talung Formation, followed by a change to a normal benthic  
376 oxygenation state that continues to increase towards Bed 12. This upward trend in increasing  
377 oxygenation during the latest Permian indicates an oxygenation event and is supported by: i)  
378 lower values of Cr/Al, Co/Al, Cu/Al, V/Al, V/Cr (Fig. S13); and ii) a decreasing ratio of  
379 calcareous foraminifera to agglutinated foraminifera, as, in the modern ocean, calcareous  
380 foraminifera dominate in the oxygen minimum zone and the ratio of calcareous foraminifera to  
381 agglutinated foraminifera decreases with increasing oxygen (Gooday et al., 2000; Levin et al.,  
382 2002). The increased oxygenation is also supported by a progressive increase in the intensity of  
383 bioturbation and abundance of trace fossils until extinction horizon in the Dongpan section.  
384 Moreover, this inferred rise in oxygen level is associated with sea-level fall (indicated by  
385 changes in radiolarian taxa), decreasing marine primary productivity (indicated by lower Ba/Al  
386 ratios and reduction in radiolarian diversity) and brachiopod miniaturization (He et al., 2007).

387

388 2.2.1.2. Penglaitan section, Laibin City, Guangxi Province (Shen et al., 2019; Xiang et al., 2021;  
389 **Fig. S14)**

390 The Penglaitan section was deposited in a deep-marine slope setting during the P-T  
391 boundary interval (Xiang et al., 2021). The Talung Formation of latest Permian consists mainly  
392 of mixed carbonate-siliciclastic-tuffaceous sediments, and the overlying Luolou Formation of  
393 earliest Triassic consists of interbedded lime mudstone and micrite limestone. The mass  
394 extinction horizon occurs at the top of Bed 141 within the Talung Formation. The P-T boundary  
395 is defined where the conodont *H. parvus* first occurs within Bed 142 at the contact between the  
396 Talung and the Luolou Formation.

397 Iron speciation proxies and absolute values of excess redox-sensitive elements (Mo, V and  
398 U) (Fig. S14) indicate there were mainly oxic water conditions with sporadic anoxia  
399 immediately prior to and during the end-Permian mass extinction horizon (up to bed 141;  
400 characterized by lower values of  $Fe_{HR}/Fe_T$ ,  $Fe_{Py}/Fe_{HR}$ , TOC and enrichment factors of  
401 redox-sensitive elements), which corresponds to the *C. meishanensis* -*H. preparvus* conodont  
402 zones, suggesting an oxygenation event. Environmental stresses (e.g., decreasing ocean  
403 temperatures shown by conodont  $\delta^{18}O_{\text{apatite}}$  and increasing volcanism suggested by tuffaceous  
404 layers) have been proposed to begin before the end-Permian mass extinction event in the section  
405 (Shen et al., 2019). Decreasing oxygenation at the level of bed 142 is consistent with sudden  
406 disappearance of diverse biota and an enrichment of TOC and U concentration (Fig. S14).

407

### 408 2.2.2. Shallow-marine P-T sections on the isolated carbonate platforms (**Fig. 5**)

409 2.2.2.1. Dajiang section, Luodian County, Guizhou Province (Song et al., 2014; Liao et al., 2017;

#### 410 **Fig. S15**)

411 The Dajiang section records the P-T transition on an isolated, shallow-marine carbonate  
412 platform. The uppermost Permian part consists mainly of bioclastic limestone of the Wuchiaping  
413 Formation. The Lower Triassic part consists of microbialites overlain by thin-bedded, poorly  
414 bioturbated micrite limestone of the Daye Formation. Within the microbialites are several shelly  
415 interlayers containing small thin-shelled bivalves, gastropods and ostracods. The contact  
416 between the Wuchiaping Formation and the Daye Formation is an irregular surface thought to  
417 result from erosion and karstification and/or submarine dissolution (Liao et al., 2017). Both the  
418 mass extinction horizon and the P-T boundary (marked by the lowest occurrence of *H. parvus*)

419 are defined at the base of the Daye Formation, thereby indicating a stratigraphic gap.

420 The lack of framboidal pyrite and the presence of diverse, abundant fossils within the  
421 Wuchiaping Formation suggest the development of oxic marine conditions prior to the mass  
422 extinction horizon (Fig. S15). Immediately above the mass extinction horizon, the sudden  
423 appearance of abundant small (mean diameter between 5 and 9  $\mu\text{m}$ ) framboidal pyrites in the  
424 Daye Formation suggests lower dysoxic to anoxic marine conditions during the deposition of the  
425 microbialites and overlying micrite limestone (Song et al., 2014; Liao et al., 2017). These  
426 earliest Triassic lower dysoxic to anoxic conditions are also reflected by: (i) small carapace size  
427 and high intraspecific variability of ostracods (Forel, 2013); (ii) low diversity and high  
428 abundance of small body-size “opportunistic” or “disaster” species; and (iii) increasing  
429  $S_{\text{pyrite}}/\text{TOC}$  ratios and degree of pyritization values (Song et al., 2014) (Fig. S15). Notably, the  
430 framboidal pyrites in this microbialite layer are smaller than those in the coeval microbialites in  
431 the Cili and Laolongdong sections in the northern part of the Yangtze Platform.

432

433 2.2.2.2. Zuodeng section, Baise City, Guangxi Province (Yang et al., 2015; Fang et al., 2017; **Fig.**  
434 **S16**)

435 The Zuodeng section is a shallow-marine P-T boundary interval section located within the  
436 Debao isolated carbonate platform. The uppermost Heshan Formation consists of bioclastic  
437 limestone, and the lower Majiaoling Formation is comprised of microbialites in the lower part  
438 and micrite limestone in the upper part (Fang et al., 2017). There is an irregular contact between  
439 the two formations (Yang et al., 2015). The uppermost Heshan Formation yields abundant  
440 foraminifera of *Colaniella* sp. and *Palaeofusulina sinsensis*, which correspond to the *C. yini*

441 conodont zone. The microbialites of the basal Majiaoling Formation yield conodonts belonging  
442 to the *H. parvus* and *I. staeschei* Zones. Within the microbialites, large-sized framboidal pyrites  
443 (mean size of ca. 15  $\mu\text{m}$ ) and abundant and diverse biota (including foraminifera, ostracods and  
444 microconchids) suggest lower to upper dysoxic conditions. This interpretation is consistent with  
445 the common occurrence, within the microbialites, of *Gakhumella* and *Renalcis*-like (unique to  
446 the post-EPME microbialites) calcimicrobes (possibly photosynthetic calcimicrobes) and of  
447 other taxa which require oxygen (Yang et al., 2015; Kershaw et al., 2021) (Fig. S16).

448

#### 449 2.2.2.3. Taiping section, Pingguo County, Guangxi Province (Xiao et al., 2018; **Fig. S17**)

450 During the P-T boundary interval, the Taiping section was located on the small Pingguo  
451 Platform and displays an upward environmental change from a deeper basin to shallow-marine  
452 settings. The Heshan Formation consists of bioclastic limestone with some interbedded cherty  
453 limestone and cherts in the lower part (Fig. 6H). Permian fusulinid taxa (*Nankinella* sp. and  
454 *Reichelina* sp.) suggest a latest Permian age for the uppermost Heshan Formation. The overlying  
455 Majiaoling Formation of earliest Triassic consists of microbialites in the basal part overlain by  
456 oolitic limestone/dolostone. The microbialites at this location have yielded five species of  
457 *Hindeodus* (including *H. parvus*, *praeparvus*, *eurypyge*, *changxingensis* and *inflatus*). The  
458 contact between the Heshan Formation and Majiaoling Formation is a flat surface without signs  
459 of carbonate dissolution, although an irregular, carbonate-dissolved surface occurs 2 cm above  
460 the formation contact. At the contact of the two formations, the mass extinction horizon also  
461 coincides with the P-T boundary as indicated by the first occurrence of *H. parvus*.

462 Abundant small framboidal pyrites with low  $\delta^{34}\text{S}_{\text{pyrite}}$  values occurring in both the

463 uppermost Heshan Formation and the lower Majiaoling Formation indicate lower dysoxic to  
464 anoxic water conditions (Fig. S17). Euxinic-anoxic conditions are recorded in the uppermost  
465 Permian just below the P-T boundary (indicated by lower  $\delta^{34}\text{S}_{\text{py}}$  and smaller framboidal pyrites),  
466 but there is an oxygenation event recorded in the microbialites during the earliest Triassic above  
467 the P-T boundary, as inferred from higher  $\delta^{34}\text{S}_{\text{py}}$  and larger framboidal pyrites (Fig. S17).

468

### 469 **3. Discussion on the P-T marine redox history recorded in South China**

470

#### 471 *3.1. Inconsistent interpretations from different redox proxies*

472 The large body of P-T marine redox studies in South China reveals that the interpretations  
473 from different redox proxies, even when applied into the same P-T section, can vary  
474 substantially (Bond and Wignall, 2010; Lehrmann et al., 2015; Xiao et al., 2018; Xiang et al.,  
475 2020). A good example is the Meishan GSSP section, for which the redox interpretation from  
476 iron speciation data (Xiang et al., 2020) is different to that deriving from other redox proxies  
477 (e.g., framboidal pyrites and trace fossils) (e.g., Li et al., 2016) (**Table 1**). A second example is  
478 the shallow-marine microbialites above the P-T boundary, for which anoxic (even sulfidic)  
479 conditions were mainly inferred from abundant small-size framboidal pyrites, biomarkers and U  
480 isotopes (Grice et al., 2005b; Liao et al., 2010; Nabbefeld et al., 2010; Luo et al., 2013; Lau et  
481 al., 2016), but dysoxic to oxic conditions were indicated by multiple lines including low total  
482 organic carbon (TOC) and total sulfur content, large-size framboidal pyrites in some cases, some  
483 biomarkers, Rare Earth Element (REE) and iodine chemistry, diverse bioclasts and  
484 upward-increasing bioclast size and abundance, and the presence of photosynthetic

485 cyanobacteria within the microbialites (Lehrmann et al., 2015; Wang et al., 2016; Kershaw et al.,  
486 2018) (**Table 1; Figs 3 and 5**).

487 There are many possible explanations for these divergences, including the sensitivity of  
488 different proxies to redox changes, and the effects of early marine diagenetic alterations  
489 (Kershaw et al., 2018; Eroglu et al., 2021). Some of the potential artifacts affecting redox  
490 proxies are summarized below.

491

### 492 3.1.1. Redox sensitivity

493 Different redox proxies have different sensitivities to changes in marine redox. For example,  
494 the residence time of U isotopes (*ca.* 400 kyr) may be too long to capture short-term redox  
495 changes, while the biomarkers may record more episodic or instantaneous seawater redox  
496 changes (Bond and Wignall, 2010; Chen et al., 2011; Li et al., 2021; Newby et al., 2021).  
497 Therefore, different marine redox history can be inferred from U isotopes and biomarkers, even  
498 they are measured in the same section. Multi-proxy approaches in the same rocks will provide  
499 the best picture of marine redox changes.

500

### 501 3.1.2. Artifacts affecting the iron speciation proxy

502 Early marine diagenesis within sediments can modify iron speciation data inherited from  
503 seawater properties, thereby leading to increases in  $Fe_{HR}/Fe_T$  and  $Fe_{py}/Fe_{HR}$  ratios while  
504 maintaining constant  $Fe_T/Al$  ratios (Eroglu et al., 2021). In the Meishan section, although higher  
505  $Fe_{HR}/Fe_T$  and  $Fe_{py}/Fe_{HR}$  ratios in Beds 24e to 34 were suggested to indicate anoxic and/or  
506 ferruginous marine conditions, the  $Fe_T/Al$  ratios are relatively constant (Xiang et al., 2020) (**Fig.**

507 7). If the higher  $Fe_{HR}/Fe_T$  and  $Fe_{py}/Fe_{HR}$  ratios are related to anoxic and ferruginous marine  
508 conditions, one would also expect to see increasing  $Fe_T/Al$  ratios (Eroglu et al., 2021). It is  
509 possible that the iron speciation proxy data of the Meishan section may be altered during early  
510 marine diagenesis and can therefore be problematic to reflect marine redox conditions. In  
511 contrast, in the Shangsi and Penglaitan sections the iron speciation analyses through the P-T  
512 boundary interval yielded higher  $Fe_{HR}/Fe_T$  and  $Fe_{py}/Fe_{HR}$  ratios that were typically associated  
513 with increasing  $Fe_T/Al$  ratios (**Fig. 7**), thereby supporting the interpretation of marine redox  
514 conditions (Lei et al., 2017; Xiang et al., 2016, 2021; Eroglu et al., 2021).

515

### 516 3.1.3. Artifacts affecting the uranium isotope proxy

517 Early marine diagenesis in situations where there is a shallow redox boundary (i.e., the  
518 boundary changing from oxic to anoxic conditions within seafloor sediment porewater is very  
519 near the sediment-water interface) and dolomitization can drive a negative excursion of U  
520 isotopes (Romaniello et al., 2013), which otherwise could be interpreted as a fingerprint of  
521 marine anoxia (Zhang et al., 2018a). Li et al. (2018) reported an early marine diagenetic  
522 dolomitization event around the Permian-Triassic boundary interval in some South China  
523 sections. Therefore, the influence of early marine diagenesis and dolomitization on the P-T U  
524 isotope signals across the P-T boundary interval need consideration for a better understanding of  
525 marine redox conditions indicated by U isotopes in the absence of independent proxies.

526

### 527 3.1.4. Artifacts affecting the framboidal pyrite proxy

528 Framboidal pyrites can form both within an anoxic seawater column and as early diagenetic



529 products in sediments (Gomes et al., 2018; Kershaw et al., 2018). For example, in microbial  
530 mats, Gomes et al. (2018) reported small pyrites (< 25  $\mu\text{m}$ ) with high  $\delta^{34}\text{S}$  values at the mat  
531 surface, and large pyrite grains (> 35 $\mu\text{m}$ ) with low  $\delta^{34}\text{S}$  values from deeper parts of the same  
532 mats. The  $\delta^{34}\text{S}$  trend of pyrites of different sizes and from different locations within the  
533 microbial mats has been suggested to be related to differential microbial metabolic activities, but  
534 the origin of varying  $\delta^{34}\text{S}$  in different pyrites can be different in non-microbialite marine settings  
535 (Gomes et al., 2018). Proposed explanations for the observed trend in the microbialites are: high  
536 rates of sulfate reduction, partial closed-system sulfate consumption in microenvironments,  
537 and/or sulfide oxidation near the mat surface, leading to small pyrites with high  $\delta^{34}\text{S}$  values; and  
538 slower sulfate reduction rates deeper parts in the mats, leading to large pyrites with low  $\delta^{34}\text{S}$   
539 values (Gomes et al., 2018).

540 An early diagenetic origin of some framboidal pyrites within the P-T microbialites of South  
541 China is supported by the coexistence of framboidal pyrites of widely varying sizes in sediments  
542 with low TOC content, and well-preserved microbial structures and microfossils (e.g., Figs 6F–  
543 G) (Luo et al., 2010; Wang et al., 2016; Gomes et al., 2018; Kershaw et al., 2018; Zhang et al.,  
544 2020). The widespread microbialites have been reported to have highly variable  $\delta^{34}\text{S}$  values and  
545 higher  $\delta^{34}\text{S}$  (both  $\delta^{34}\text{S}_{\text{CAS}}$  and  $\delta^{34}\text{S}_{\text{pyrite}}$ ) values than coeval sediments deposited in deeper marine,  
546 non-microbialite settings (Luo et al., 2010; Zhang et al., 2011; Xiao et al., 2018). The higher  
547  $\delta^{34}\text{S}$  values of the microbialites have been explained by low seawater sulfate concentrations  
548 (Luo et al., 2010) or early diagenetic influence (e.g., oxidation of framboidal pyrite) (Gomes et  
549 al., 2018; Johnson et al., 2021). Further, multiple other redox proxies applied into the  
550 microbialites suggest oxic conditions prevailed during their formation (Forel et al., 2009; Chen

551 et al., 2011; Loope et al., 2013; Yang et al., 2015; Kershaw et al., 2018). Besides, redox proxies  
552 (including framboidal pyrites) of coeval landward and seaward non-microbialite sediments also  
553 suggest oxic conditions at the time of microbialite formation (Fig. 7). For the distinction of  
554 framboidal pyrites associated with P-T microbialites between early diagenetic and water column  
555 origin, it is tentatively proposed: i) multiple redox proxies are applied to strata containing  
556 framboidal pyrites to ascertain whether the results/interpretations are consistent (water column  
557 origin) or not (diagenetic origin); ii) detailed petrological description of framboidal pyrites is  
558 carried out to evaluate whether the size and morphologies are approximated (water column  
559 origin) or not (diagenetic origin); iii) evaluate the relationship between framboidal pyrites and  
560 TOC,  $\delta^{13}\text{C}$  and early diagenetic features, in light of the observation that early diagenetic  
561 framboidal pyrites tend to be associated with a decrease in TOC, the release of light carbon and  
562 the promotion of early cementation and preservation of primary sedimentary fabrics, while  
563 framboidal pyrites of water column are not; iv) an investigation of landward and seaward water  
564 redox condition adjacent to the localities developing microbialites containing framboidal pyrites  
565 to see if the redox conditions are consistent (water column origin) or not (diagenetic origin); v)  
566 advanced technologies, such as fine-scale and *in situ* analyses (e.g., secondary ion mass  
567 spectrometry) of  $\delta^{34}\text{S}$  signals of pyrites of different sizes, are suggested to help better distinguish  
568 between origins from seawater and those rooted in early diagenesis in sediments. These could  
569 combine to understand the environmental implications of the framboidal pyrites observed in the  
570 P-T microbialites (Gomes et al., 2018; Johnson et al., 2021).

571

572 3.1.5. Artifacts affecting the biomarker proxy

573 Upwellings can influence the validity of redox proxies, like biomarkers and framboidal  
574 pyrites. Biomarkers in different P-T microbialites indicate low oxygen marine conditions in  
575 some places but normal oxic marine conditions in other places (Luo et al., 2013; Zhou et al.,  
576 2017). One potential explanation for the phenomena is that there was an influx of biomarkers  
577 transported from deeper depths via ocean circulation, such as the upwelling of deep anoxic water  
578 (Kershaw et al., 2018). For the occurrence of small framboidal pyrites in the earliest Triassic  
579 microbialites, the influence of upwellings was proposed to reconcile the coexistence with oxic  
580 indicators (such as diverse biota and negative Ce/Ce\* anomaly) (Kershaw, 2015; Kershaw et al,  
581 2016).

582

### 583 3.1.6. Evaluating results from multiple proxies

584 The above observations and discussions indicate that one must evaluate redox proxies in  
585 terms of potential effects of early marine diagenesis, of sensitivity to redox changes and of  
586 influence of local upwellings before accepting their interpretations about marine conditions.  
587 Therefore, multiple redox proxies should be used and compared to determine the marine redox  
588 conditions. After evaluating these types of sensitivity, diagenetic and upwelling influences on  
589 the reported redox proxies, this study proposed that dysoxic-oxic marine conditions are preferred  
590 for the *H. parvus* conodont zone in the marine upper slope settings such as in the Meishan  
591 section, based on evidence including decreasing values of  $Fe_{HR}/Fe_T$ ,  $Fe_{Py}/Fe_{HR}$  and Mo, and  
592 increasing size or the absence of framboidal pyrites (Figs S6, S7). The dysoxic-oxic marine  
593 conditions are also proposed for the formation of the well-developed microbialites in the shallow  
594 marine settings, according to: i) in contrast with framboidal pyrites, multiple redox proxies

595 applied to the microbialites suggesting dysoxic-oxic conditions (Chen et al., 2011; Loope et al.,  
596 2013; Yang et al., 2015; Kershaw et al., 2018); ii) the large variation of sizes and morphologies  
597 of framboidal pyrites (e.g., Yang et al., 2015; ); iii) the co-occurrence of framboidal pyrites, low  
598 TOC, negative  $\delta^{13}\text{C}$  excursion and early cementation features, as early diagenetic framboidal  
599 pyrites tend to decrease TOC, release light carbon and promote early cementation and  
600 preservation of primary sedimentary fabrics (Luo et al., 2010; Wang et al., 2016, Kershaw et al.,  
601 2018); and iv) the development of oxic marine conditions in coeval adjacent landward and  
602 seaward settings (Li et al., 2016, 2021).

603

### 604 3.2. Redox variation with water depth and with paleogeography

605 The high-resolution conodont biostratigraphy of the P-T sections deposited at different  
606 paleo-water depth settings enables a spatial and temporal compilation of marine redox variation  
607 in South China (**Fig. 8**).

#### 608 3.2.1. Northern Margin of the Yangtze Platform and adjacent Northern Marginal Basin

609 During the latest Permian, upper dysoxic to oxic marine conditions characterize lower slope  
610 to basinal settings (Xiaojiaba, Chaotian and Shangsi sections) during the interval immediately  
611 before and during the P-T mass extinction horizon (mainly corresponding to *C. taylorae*,  
612 *C. zhejiangensis*, *C. changxingensis* and *C. yini* conodont zones). Before and after this interval  
613 of relatively oxygenated conditions, lower dysoxic to anoxic (even euxinic) conditions were  
614 prevalent (**Figs 2, 8**).

615 In contrast, lower dysoxic to anoxic (even euxinic) conditions were developed in upper  
616 slope settings (Changtanhe, Ganxi and Meishan sections) during the end-Permian extinction

617 interval, but with an oxygenation event immediately after the P-T boundary that was mainly  
618 during *H. parvus* -*I. isarcica* conodont zone (**Figs 2, 8**).

619 The shallow-marine settings display two types of redox history based on the depositional  
620 facies and environmental conditions. The first kind is typified by the Liangfengya section in a  
621 more nearshore location with no development of microbialites above the P-T boundary (**Fig. S8**).  
622 That section indicates continuously oxic marine conditions during the deposition of the  
623 uppermost Permian bioclastic limestone through lowermost Triassic micrite limestone  
624 (*C. changxingensis*, *C. yini*, *C. meishanensis*, *H. parvus* and *I. isarcica* conodont zones).

625 The second type is the more seaward locations where microbialites occurred above the P-T  
626 boundary (Laolongdong, Zaixia, Gaohua and Panjiazhuang sections) (**Figs S9 to S12**). In those  
627 locations, an oxic marine redox condition is interpreted for the deposition of the uppermost  
628 Permian bioclastic limestone (*C. changxingensis*, *C. yini*, *C. meishanensis* conodont zones). An  
629 oxic marine setting is also proposed herein for the microbialites that formed during the earliest  
630 Triassic (*H. parvus* Zone) based on: (i) multiple sedimentological, fossil and geochemical  
631 evidence favoring oxic marine settings; (ii) a reevaluation of apparent anoxic indicators from  
632 small framboidal pyrites, U isotopes and biomarkers as being possibly affected by early redox  
633 sensitivity, early diagenesis and influence from local upwelling; (iii) consistency with the redox  
634 history of the more nearshore section (Liangfengya) and more seaward section (Meishan) that  
635 display coeval oxic marine settings. Accordingly, different with the open and deep marine  
636 framboidal pyrites, the formation of framboidal pyrites within shallow marine microbialites may be  
637 heavily influenced by a closed system of early diagenesis characterized by anoxic porewater  
638 conditions (Gomes et al., 2018; Kershaw et al., 2018), as is supported by early diagenetic

639 petrological features and low TOC of associated carbonate components, uneven pyrite size and  
640 morphologies and variable and heavier  $\delta^{34}\text{S}_{\text{py}}$  values (Luo et al., 2010; Wang et al., 2016;  
641 Gomes et al., 2018; Kershaw et al., 2018; Zhang et al., 2020). In both types, the oxic condition  
642 of earliest Triassic is followed by a dysoxic-anoxic redox interval (corresponding to the  
643 conodont zone *I. staeschei*) associated with a marine transgression, although this interval is  
644 occasionally interrupted by oxic conditions expressed by oolitic and shelly sediments.

645

### 646 3.2.2. Hunan-Guizhou-Guangxi Basin facing Panthalassic Ocean

647 Deep marine slope to basinal P-T sections (Dongpan and Penglaitan) in the southern region  
648 record a similar redox history as those of the northern marginal basin (Xiaojiaba, Chaotian and  
649 Shangsi sections). An oxygenation event occurred immediately below and during the  
650 end-Permian mass extinction horizon. This event is sandwiched above and below by dysoxic to  
651 anoxic conditions (**Fig. 4**).

652 The shallow-marine sections on the isolated carbonate platforms display similar lithological  
653 features to the second type of shallow-marine sections of the northern margin of the Yangtz  
654 Platform. Mainly oxic-dysoxic marine conditions were present during the deposition of both the  
655 upper Permian bioclastic limestone and the basal Triassic microbialites.

656

### 657 3.3. Possible reasons for the P-T deep marine oxygenation events in South China

658 Based on the correlation of P-T marine redox conditions in South China (**Figs 2-5**), two  
659 deep marine oxygenation events are suggested to occur during the P-T boundary interval that  
660 affected different sedimentary environments. The first oxygenation event occurred immediately

661 before and during the end-Permian mass extinction horizon, corresponding to the *C. taylorae*,  
662 *C. zhejiangensis* and *C. changxingensis yini* conodont zones. The second oxygenation event  
663 occurred during the *H. parvus* -*I. isarcica* conodont zones immediately after the P-T boundary.  
664 The first oxygenation event is recorded mainly in basinal to lower slope settings, whereas the  
665 second oxygenation event is apparent mainly in upper slope settings.

666 Variations in ocean circulation and marine primary productivity are commonly invoked as  
667 causes for changes of marine redox conditions (Kuypers et al., 2002; Meyer et al., 2011;  
668 Georgiev et al., 2015; Xiang et al., 2016). The first oxygenation event in deeper water  
669 settings appeared around the end Permian mass extinction horizon and co-occurred with  
670 enhanced volcanic activities (e.g., Xie et al., 2007; Shen et al., 2021), climate cooling (Shen et  
671 al., 2010; Chen et al., 2016; Joachimski et al., 2020), decreased primary productivity (Zhang et  
672 al., 2007, 2018c; Algeo et al., 2010, 2011; Shen et al., 2014, 2015; Wei et al., 2015; Wu et al.,  
673 2022), a rapid (ca. 100 ky) sea-level fall (Kershaw et al., 2002; Yin et al., 2014), and increasing  
674 terrestrial influx (Korte et al., 2003; Xie et al., 2007). Likely, the oxygenation event may be  
675 caused by increased oxygen supply from shallow seawater, as is supported by enhanced ocean  
676 circulation driven by climate cooling related to volcanic activities releasing SO<sub>2</sub> (Shen et al.,  
677 2020; Newby et al., 2021) (**Fig. 9**). Moreover, given the time-equivalence with volcanic  
678 activities, it is feasible that there is a causal relationship between the rapid sea-level fall during  
679 the first oxygenation and climate cooling related to volcanism, and similar cases have been  
680 reported in the end-Triassic mass extinction events (Schoene et al., 2010).

681 While, during the oxygenation event in the basinal and lower slope settings, increased  
682 oxygen depletion in the upper slope settings are consistent with upwelling of deeper marine

683 anoxic water (Li et al., 2016; Lei et al., 2017; Xiao et al., 2018). Meanwhile, the decreased  
684 marine productivity doesn't favor expansion of oxygen minimum zone (OMZ) as the reason for  
685 oxygen depletion in upper slope settings (Wei et al., 2015; Xiang et al., 2016). On the other hand,  
686 the effects of upwelling may have been stronger in the southern part of South China facing  
687 Panthalassa, where shallow marine platforms also show oxygen depletion (e.g., Taiping section)  
688 (Xiao et al., 2018).

689 The second oxygenation event, which affected predominantly the upper slopes during the  
690 earliest Triassic, occurred during a time of greatly reduced marine primary productivity in the  
691 South China area (Shen et al., 2014, 2015; Zhang et al., 2018c; Wu et al., 2022). At this time, the  
692 very warm climate might have promoted ocean stratification and reduced upwelling of  
693 nutrient-rich deeper waters (Joachimski et al., 2012; Winguth and Winguth, 2012; Schobben et  
694 al., 2014; Chen et al., 2016) and, hence also contributed to changes of seawater chemistry  
695 (Heydari et al., 2013; Tavakoli, 2016; Li et al., 2018) and reduced marine primary productivity  
696 (**Fig. 9**). The dysoxic-anoxic conditions in lower slope-basinal settings (**Figs 2, 8**) are consistent  
697 with such a reduced ocean circulation. The second oxygenation event that mainly affected the  
698 upper slope settings in South China may be related to the low marine primary production and  
699 biomass extinction, which in turn led to less marine oxygen consumption by organic respiration  
700 and decomposition and consequently a recovery in oxygenation levels (Algeo et al., 2013; Xiao  
701 et al., 2018).

702 In contrast to marine redox fluctuations in slope and basinal settings, redox conditions in  
703 most shallow-marine settings are thought to have remained consistently oxic (and/or dysoxic)  
704 through the P-T boundary interval, including during the formation of the microbialites during the



705 earliest Triassic, while some local areas may be influenced by the upwellings of deeper and  
706 oxygen-poor waters.

707

### 708 *3.4. Correlation with carbon-isotope excursions in South China*

709 Two marine negative  $\delta^{13}\text{C}$  excursion events have been reported during the P-T boundary  
710 interval in South China (Krull et al., 2004; Song et al., 2013; Li et al., 2021; Wu et al., 2022);  
711 and both of these appear to be associated with the oxygenation events proposed in the study.

712 The first negative  $\delta^{13}\text{C}$  excursion in both  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  is present in both deep and  
713 shallow-marine settings during the end-Permian mass extinction interval (Newby et al., 2021).  
714 This is temporally consistent with the first oxygenation event recorded in basinal to lower slope  
715 settings. Possibly, enhanced oxidization and remineralization of organic matters that had  
716 previously accumulated under the recorded low oxygen conditions may serve as a possibility  
717 contributing to that negative  $\delta^{13}\text{C}$  excursion by releasing large amounts of isotopically light  
718 carbon. Other suggested drivers of this excursion include volcanic  $\text{CO}_2$ , methane release,  
719 biomass kill, erosion of soil carbon, reduced organic matter burial (Berner, 2002; Krull et al.,  
720 2004; Payne and Kump, 2007; Retallack and Jahren, 2008; Algeo et al., 2011).

721 The second negative  $\delta^{13}\text{C}$  excursion in  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  is mainly present in the  
722 shallow-marine microbialite interval in the earliest Triassic above the P-T boundary (Krull et al.,  
723 2004; Liao et al., 2010; Kershaw et al., 2012b; Wang et al., 2016; Xiao et al., 2018; Wu et al.,  
724 2022), whereas a positive  $\delta^{13}\text{C}$  excursion occurred in upper slope settings (e.g., Meishan,  
725 Shangsi, Chaotian) (Li et al., 2021; Zhou et al., 2021; Wu et al., 2022). The contrasting patterns  
726 of  $\delta^{13}\text{C}$  excursion along a proximal-to-distal direction are unexplained hitherto. The shallow

727 marine microbialites are benthic and well preserved, indicating that they formed in low energy  
728 and relatively restricted settings (Kershaw et al., 2012a). In relatively restricted, oxic shallow  
729 marine settings, isotopically light carbon released during organic decomposition may cause a  
730 local depletion in seawater  $^{13}\text{C}$  and a negative excursion  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  in the carbonate  
731 sediment. Similar patterns have been reported in modern shallow-marine carbonate settings at  
732 Florida, the Bahamas and Abu Dhabi (Lloyd, 1964; Patterson and Walter, 1994; Pederson et al.,  
733 2021). For the upper marine slope sections, reduced input of isotopically light carbon by  
734 decreased marine primary productivity (e.g., Shen et al., 2014) may have contributed to the  
735 positive  $\delta^{13}\text{C}_{\text{carb}}$  and  $\delta^{13}\text{C}_{\text{org}}$  excursion in the sediment. Previous studies also proposed a change  
736 in carbon source (e.g., influx of terrestrial carbon and changes in microbial composition) for the  
737 positive  $\delta^{13}\text{C}$  excursion in the upper slope settings (Wu et al., 2022).

738

### 739 *3.5. Implications for the P-T mass extinction events in South China*

740 The two oxygenation events around the P-T boundary interval may provide some  
741 implications for environmental scenarios of the P-T mass extinction scenario in South China.  
742 For the first oxygenation event immediately before and during the marine mass extinction  
743 horizon, enhanced ocean circulation and sea level fall have been attributed to short-term climate  
744 cooling superimposed on the overall warming trend as a consequence of massive  $\text{SO}_2$  release by  
745 the emplacement of the Siberian Traps large igneous province (Wu et al., 2014; Baresel et al.,  
746 2017; Newby et al., 2021). Therefore, volcanic activities seem to have already occurred  
747 immediately before the marine mass extinction events. Terrestrial P-T extinction and  
748 deforestation related to volcanism have been reported to occur before marine mass extinction,

749 which supports volcanic activities before the marine mass extinction events (Shen et al., 2011;  
750 Kaiho et al., 2016; Liu et al., 2017; Zhou et al., 2017; Xiao et al., 2018). This timing is also  
751 indicated by the wildfire biomarkers and black carbon (Xie et al., 2007; Zhou et al., 2021). On  
752 the other hand, although ocean anoxia has been argued as an important killing mechanism for  
753 end-Permian mass extinction events (e.g., Isozaki, 1997; Liao et al., 2010; Wu et al., 2022), this  
754 is inconsistent with the deep-water oxygenation event at the end-Permian mass extinction  
755 interval (Bond and Wignall, 2010; He et al., 2013), at least in some settings. However, combined  
756 with regression, the upwelling of deep and anoxic water may cause the biota crisis for shallow  
757 marine biota. The development of ocean anoxia has started considerably before the marine mass  
758 extinction events, and can't account for the biotic turnover and extinction events on land (e.g.,  
759 Bond and Wignall, 2010). This indicates clearly more complex scenarios than a simple  
760 death-by-anoxia model during the P-T mass extinction.

761 The second oxygenation event, which affected mainly the upper slopes during the earliest  
762 Triassic following the P-T boundary, occurred immediately prior to a second pulse of marine  
763 extinction from the *H. parvus* to the *I. isarcica* conodont zones (Song et al., 2013; Su et al.,  
764 2021). During this time, there was enhanced ocean stratification and a disturbance in the marine  
765 nutrient cycle and food web, as indicated by a negative excursion of  $\delta^{15}\text{N}_{\text{org}}$  in the P-T sections  
766 of South China (Luo et al., 2011; Wu et al., 2022). In the shallow-marine settings, oxic  
767 conditions and localized food webs associated with the benthic microbialites may have provided  
768 ecological refuges for diverse shallow-marine biota (e.g., gastropods, bivalves and ostracods)  
769 after the end-Permian extinction. Those shallow-marine biota, however, suffered a wave of  
770 extinction during the following anoxia event associated with a sea-level rise over the shallow

771 marine shelves.

772

### 773 *3.6. Correlation of marine redox conditions between South China and Panthalassa*

774 The first oxygenation event that affected South China seems to have also occurred in other  
775 localities in the Paleotethys and Panthalassa Ocean (**Fig. 10**). Based on the P-T section in  
776 northern Italy, Newton et al. (2004) has reported such a transient oxygenation event in deep  
777 water, which caused reoxidation of deep-water H<sub>2</sub>S and consequent negative excursion of  $\delta^{34}\text{S}$   
778 and  $\delta^{18}\text{O}$  of carbonate-associated sulphate. Newby et al. (2021) reported an ocean oxygenation  
779 event during the end-Permian mass extinction interval from shelf to abyssal settings in the  
780 low-latitude Panthalassa Ocean. The short-term oxygenation event was also supported by  
781 bioturbation and geochemical proxies (e.g.,  $\delta^{34}\text{S}$  values and positive Ce anomaly) in deep  
782 Panthalassan P-T chert sections (Kakuwa, 2008).

783 For the second oxygenation event, it has been mainly documented in South China showing  
784 decreased primary productivity, and not recognized in the Panthalassa, where, in contrast, there  
785 is evidence showing no-decreasing and even increasing primary productivity at that time (Algeo  
786 et al., 2010, 2011; Shen et al., 2015; Newby et al., 2021). Marine nitrogen nutrient has been  
787 recognized as an important control on the marine primary productivity (Falkowski, 1997; Moore  
788 et al., 2013; Buchanan et al., 2021). The P-T  $\delta^{15}\text{N}_{\text{org}}$  isotope records show: decreasing (and low)  
789  $\delta^{15}\text{N}_{\text{org}}$  values across the P-T boundary in South China, Paleotethys realm (Luo et al., 2011; Wu  
790 et al., 2022); while increasing (and high)  $\delta^{15}\text{N}_{\text{org}}$  values across the P-T boundary in the  
791 Panthalassa realm (Georgiev et al., 2015). A terrestrial influence on the  $\delta^{15}\text{N}_{\text{org}}$  records is thought  
792 to be negligible (Luo et al., 2011; Radice et al., 2019; Wu et al., 2022), and main reasons for the

793 nitrogen isotope variation are nitrogen fixation and denitrification related to upwellings  
794 (Robinson et al., 2012; Buchanan et al., 2021). When upwellings are frequent and strong,  
795 nutrient nitrogen from deep water would fertilize surface seawater and promote marine primary  
796 productivity and water column denitrification, leading to increasing and high  $\delta^{15}\text{N}_{\text{org}}$  values;  
797 while sluggish ocean circulation limits upwelling and nitrogen supply to surface seawater  
798 (possibly nitrogen limited), leading to nitrogen fixation and decreasing  $\delta^{15}\text{N}_{\text{org}}$  towards low  
799 values (Luo et al., 2011; Robinson et al., 2012; Buchanan et al., 2021; Wu et al., 2022). Thus, it  
800 appears that different ocean circulation and nutrient nitrogen cycle occurred between South  
801 China and Panthalassa around the P-T boundary (**Fig. 11**). In South China, sea-level fall and the  
802 paleogeographic configure of the time may lead to a restriction event (Hu et al., 2021; Zheng et  
803 al., 2021). In the relatively restricted setting, climate warming and increased precipitation related  
804 to intensified monsoon favor ocean stratification and decreasing upwellings. In the Panthalassa,  
805 due to much larger ocean size, no ocean restriction happened and enhanced monsoon activities  
806 may promote upwellings.

807

#### 808 **4. Conclusions**

809       Synthesis and reevaluation of paleogeographically diverse studies of P-T marine redox  
810 conditions at different paleo-depths in South China reveal that two deep marine oxygenation  
811 events have occurred during the P-T boundary interval. The first oxygenation event occurred just  
812 before and continued through the main end-Permian mass extinction interval, and mainly  
813 affected the lower slope and basinal settings. This oxygenation event is recorded in other  
814 locations in the Paleotethys and Panthalassa realm. The first oxygenation event potentially had

815 origins in enhanced deep-ocean circulation driven by short-lived climatic cooling associated  
816 with the release of large amounts of SO<sub>2</sub> during the eruptive phases of the Siberian Traps Large  
817 Igneous Province. The second oxygenation event recorded mainly in upper slope settings of  
818 South China may be related to climate warming, increased ocean stratification, reduced ocean  
819 circulation and a decline in marine primary productivity within a relatively restricted  
820 paleogeographic configuration. The complex P-T marine redox history at different paleo-depths  
821 in South China were driven by the combination of paleogeography, major climate changes,  
822 surface and deep-ocean circulation and changes in marine productivity.

823

#### 824 **Acknowledgements**

825 Prof. James Ogg and Prof. Paul Wignall are greatly acknowledged for useful comments and  
826 instruction during the preparation of the manuscript. Constructive reviews from Dr. S. Kershaw  
827 and an anonymous colleague as well as editorial comments by Dr. A. Negri were appreciated.  
828 This study is financially supported by National Natural Science Foundation of China (No.  
829 42202123).

830

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1234 **TABLES**

1235

1236 **Table 1.** Marine redox-studied sections around the P-T boundary interval in Paleotethys and  
 1237 Panthalassa

Section	Environment al setting	Marine redox proxy and interpretation	Reference
<i>Northern Marginal Basin sections in South China (Paleotethys)</i>			
Meishan	Upper slope	Biomarkers:  photic zone euxinic across the P-T boundary interval  Framboidal pyrites, biota diversity, trace fossils:  mainly anoxic in <i>C. taylorae</i> - <i>C. zhejiangensis</i>  - <i>C. changxingensis</i> - <i>C. yini</i> zones;  mainly oxic in <i>H. parvus</i> Zone;  mainly anoxic in <i>Isarcicella</i> Zone  Iron speciation:  mainly anoxic, even euxinic in <i>H. parvus</i> Zone  mainly anoxic, ferruginous in <i>Isarcicella</i> Zone	Grice et al.,  2005b;  Summons et al.,  2006; Li et al.,  2016; Xiang et  al., 2020
Ganxi	Upper slope	Iron speciation and Mo element:  anoxic, in <i>C. wangi</i> , <i>C. changxingensis</i> , <i>C. yini</i> and  <i>C. meishanensis</i> Zones;  anoxic to oxic, in <i>H. parvus</i> and <i>I. isarcica</i> zones	Lei et al., 2017

Changtanhe	Lower slope	Framboidal pyrites, fossil diversity and TOC content:  anoxic, in the mass extinction horizon;  oxic, immediately after the mass extinction event  anoxic after the oxygenation event	He et al., 2013
Shangsi	Lower slope to basin	Iron speciation, redox-sensitive elements, fossil diversity and bioturbation degree:  euxinic, in uppermost <i>C. wangi</i> to lower and middle <i>C. subcarinata</i> Zones;  mainly anoxic, <i>C. changxingensis</i> - <i>C. yini</i> - <i>C. meishanensis</i> Assemblage Zone, to <i>H. changxingensis</i> Zone  oxic in the mass extinction horizon (within <i>C. meishanensis</i> - <i>H. changxingensis</i> Zone)  anoxic, <i>H. changxingensis</i> - <i>H. parvus</i> - <i>I. isarcica</i> zones	Wignall et al., 1995; Bond and Wignall, 2010; Xiang et al., 2016
Chaotian	Lower slope to basin	Framboidal pyrites, bioturbation, trace fossils and TOC content:  oxic, in <i>C. taylorae</i> - <i>C. zhejiangensis</i> - <i>C. changxingensis</i> - <i>C. yini</i> Assemblage Zone;  oxic, immediately above the mass extinction horizon;  anoxic, in <i>H. parvus</i> and <i>Isarcicella</i> zones	Newton et al., 2009; Saitoh et al., 2021
Xiaojiaba	Lower slope to basin	Framboidal pyrite, $\delta^{34}\text{S}_{\text{py}}$ , $\text{C}_{\text{org}}/\text{P}$ , $\text{Mo}_{\text{auth}}$ and $\text{U}_{\text{auth}}$ :  an oxygenation event in the end-mass extinction	Wei et al., 2015

		horizon	
<b><i>Sections in Northern part of Yangtze Platform in South China (Paleotethys)</i></b>			
Liangfengya	Shallow marine	Framboidal pyrites and bioturbation:  oxic, in <i>C. changxingensis</i> , <i>C. yini</i> , <i>C. meishanensis</i> ,  <i>H. parvus</i> and <i>I isarcica</i> Zones;  dysoxic to euxinic, in <i>I. staeschei</i> Zone	Li et al., 2021
Laolongdong	Shallow marine	Framboidal pyrites and fossil diversity:  oxic, below the mass extinction horizon and P-T  boundary;  lower dysoxia, in <i>H parvus</i> Zone	Liao et al., 2010
Zaixia	Shallow marine	Biomarkers:  no severe anoxia;  an oxygenation event immediately above the mass  extinction horizon;	Chen et al., 2011
Gaohua	Shallow marine	Ce anomaly and guilds:  oxic across the P-T boundary interval  Framboidal pyrites:  an oxygenation event around the mass extinction  horizon and followed by dysoxic	Forel et al., 2009; Loope et al., 2013; Wang et al., 2016
Panjiashuang	Shallow marine	Framboidal pyrites:  oxic across the mass extinction horizon	Huang et al., 2019
<b><i>Deep-water sections in the Hunan-Guizhou-Guangxi Basin in South China (Paleotethys)</i></b>			

Penglaitan	Lower slope to basin	Iron speciation and redox-sensitive elements:  oxic across the mass extinction horizon (correspond to <i>C. meishanensis</i> - <i>H. preparvus</i> Zones)	Xiang et al., 2021
Dongpan	Lower slope to basin	Redox-sensitive elements:  oxic across the mass extinction horizon	He et al., 2007;  Zhang et al., 2018b
<b><i>Sections in isolated carbonate platforms in the Hunan-Guizhou-Guangxi Basin in South China (Paleotethys)</i></b>			
Dajiang	Shallow marine	Framboidal pyrites, fossil diversity, S <sub>pyrite</sub> /TOC, Uranium concentration:  oxic below mass extinction horizon and P-T boundary;  lower dysoxic above mass extinction horizon and P-T boundary	Forel, 2013;  Song et al., 2014; Lau et al., 2016; Liao et al., 2017
Zuodeng	Shallow marine	Framboidal pyrites and microorganism:  lower to upper dysoxic, in the <i>H. parvus</i> and <i>I. staeschei</i> Zones	Fang et al., 2017
Taiping	Shallow marine to upper slope	Framboidal pyrites and $\delta^{34}\text{S}_{\text{pyrite}}$ :  lower dysoxic to anoxic in the mass extinction horizon and P-T boundary;  an oxygenation event immediately above the P-T boundary	Xiao et al., 2018
<b><i>Other marine sections in Paleotethys</i></b>			
Demirtas	Shallow	REE, Ce/Ce* anomaly and bioclasts: oxic for latest	Loop et al.

(Turkey)	marine	bioclastic limestone and earliest Triassic microbialites; and anoxic in overlying early Triassic micrite	(2013)
<i>Marine sections in Panthalassa</i>			
Opal Creek (Canada); Gujo-Hachima n, Ubara (Japan)	Shelf to abyssal settings	Thallium isotope: a transient oxygenation at the end-Permian mass extinction	Newby et al. (2021)

1238

1239 **Table 2.** Characteristics of different redox conditions during deposition (Bond and Wignall,  
1240 2010)

Conditions	Framboid parameters	Sedimentary fabric
Euxinic (persistently sulfidic lower water column)	Small (mean 3-5 $\mu\text{m}$ ), narrow size range. Framboids dominate pyrite fraction	Finely laminated
Anoxic (no oxygen in bottom waters for long periods)	Small (mean 4-6 $\mu\text{m}$ ), abundant, with a few larger framboids. Framboids dominate pyrite fraction	Finely laminated
Lower dysoxic (weakly	Mean 6-10 $\mu\text{m}$ , moderately common, with a	Finely laminated

oxygenated waters)	bottom	few, larger framboids and some crystalline pyrite	with	rare bioturbation
Upper oxygen bottom waters)	dysoxic restriction (partial restriction in bottom waters)	Moderately common to rare, broad range of in sizes, only a small proportion <5 μm. Majority of pyrite as crystals		Microburrowed. Bioturbation may obscure laminated fabric
Oxic restriction)	(no oxygen restriction)	No framboids, rare pyrite crystals		Burrowed and/or massive

1241

1242

1243 **FIGURES**

1244 **Fig. 1.** Global and South China paleogeography during the P-T boundary interval. (A) Global  
1245 paleogeography. The South China Block is indicated by the red frame. Modified from Xiao et al.  
1246 (2018). (B) South China paleogeography. The sections in this study are indicated by red  
1247 triangles. Northern margin sections: XJB = Xiaojiaba, SS = Shangsi, CT = Chaotian, LFY  
1248 = Liangfengya, LLD = Laolongdong, GX = Ganxi, GH = Gaohua, CTH = Changtanhe, ZX =  
1249 Zaixia, MS = Meishan. Southern margin sections: PJZ = Panjiazhuang, DJ = Dajiang, ZD  
1250 = Zuodeng, PLT = Penglaitan, TP = Taiping, DP = Dongpan. Detailed descriptions of the  
1251 sections are provided in the text. Modified from Yin et al. (2014).

1252

1253 **Fig. 2.** Interpretation of marine redox changes in deep marine basinal to slope P-T sections of

1254 the Northern Marginal Basin, South China. PTB = Permian-Triassic boundary, EPME  
1255 = end-Permian mass extinction horizon. Blue color = oxic to upper dysoxic; Yellow color =  
1256 lower dysoxic to euxinic. Note: the blue-color interval with a red star represents the first  
1257 oxygenation event near the EPME; and the blue-color interval with a yellow start represents the  
1258 second oxygenation event in the earliest Triassic.

1259  
1260 **Fig. 3.** Interpreted marine redox changes in shallow-marine P-T sections of the northern margin  
1261 the Yangtze Platform. PTB= Permian-Triassic boundary, EPME= end-Permian mass extinction  
1262 horizon. Blue color = oxic to upper dysoxic; Yellow color = lower dysoxic to euxinic.

1263  
1264 **Fig. 4.** Marine redox changes in deep marine lower slope to basinal P-T sections in the  
1265 Hunan-Guizhou-Guangxi Basin, South China. PTB= Permian-Triassic boundary,  
1266 EPME= end-Permian mass extinction horizon. Blue color = oxic to upper dysoxic; Yellow color  
1267 = lower dysoxic to euxinic. Note: the blue-color interval with a red star means first oxygenation  
1268 event near the EPME.

1269  
1270 **Fig. 5.** Marine redox changes in shallow-marine P-T sections in the isolated carbonate platforms  
1271 within the Hunan-Guizhou-Guangxi Basin, South China. PTB= Permian-Triassic boundary,  
1272 EPME= end-Permian mass extinction horizon. Blue color = oxic to upper dysoxic; Yellow color  
1273 = lower dysoxic to euxinic.

1274  
1275 **Fig. 6.** Photographs of Permian-Triassic marine strata and sedimentary features from South

1276 China. (A) Burrows in the uppermost Dalong Formation (latest Permian), Chaotian section  
1277 (sourced from Saitoh et al., 2021); (B) Abundant radiolarians in the uppermost Dalong  
1278 Formation (latest Permian), Chaotian section (sourced from Saitoh et al., 2014); (C) Absence of  
1279 burrows in the lowermost Feixianguan Formation (earliest Triassic), Chaotian section (sourced  
1280 from Saitoh et al., 2021); (D) Gray marl with abundant pyrite particles (black dots), and a lack  
1281 of burrows in the lowermost Feixianguan Formation (earliest Triassic), Chaotian section  
1282 (sourced from Saitoh et al., 2014); (E) The transition from bioclastic limestone (latest Permian)  
1283 to microbialites (earliest Triassic), Lalongdong section (sourced from Liao et al., 2010); (F)  
1284 Well-preserved microfabrics of microbialites (earliest Triassic) showing upward-branching  
1285 architecture, Laolongdong section (sourced from Kershaw et al., 2018); (G) Calcimicrobes  
1286 within microbialites (earliest Triassic) showing well-preserved microfabrics (sourced from  
1287 Kershaw et al., 2018); (H) Bioclastic limestone (latest Permian) under thin section, Taiping  
1288 section (sourced from Xiao et al., 2018).

1289

1290 **Fig. 7.** Iron speciation data from the Meishan, Shangsi and Penglaitan sections. In the Meishan  
1291 section,  $Fe_T/Al$  decreases across the P-T boundary (PTB), and then remains constant through the  
1292 earliest Triassic. In the Shangsi and Penglaitan sections, no  $Fe_T/Al$  decrease occurs across the  
1293 PTB. Data sources: Meishan from Xiang et al. (2020); Shangsi from Xiang et al. (2016);  
1294 Penglaitan from Xiang et al. (2021).

1295

1296 **Fig. 8.** Selected P-T marine sections revealing spatial variations in marine redox conditions at  
1297 different paleo-water depths. Note: the blue-color interval with a red star means first



1298 oxygenation event near the EPME; and the blue-color interval with a yellow start means second  
1299 oxygenation event in the earliest Triassic.

1300

1301 **Fig. 9.** A sketch showing the P-T marine redox changes. Two deep marine oxygenation events  
1302 occurred during the end-Permian mass extinction interval (EPME) and during the *H parvus*  
1303 conodont zone of earliest Triassic. SL= sea level.

1304

1305 **Fig. 10.** Marine redox variations at the P-T boundary interval in other localities in Paleotethys  
1306 and Panthalassa. Data sources: Demirtas, Loope et al. (2013), Fig. S18; Opal Creek,  
1307 Gujo-Hachiman and Ubara, Newton et al. (2004), Fig. S19.

1308

1309

1310 **Fig. 11.** Comparison of P-T marine redox changes between South China and Panthalassa area.  
1311 The first marine oxygenation event was recorded in both areas, while the second marine  
1312 oxygenation event was recognized only in South China.

1313