



UNIVERSITY OF LEEDS

This is a repository copy of *Early Triassic disaster and opportunistic foraminifers in South China*.

White Rose Research Online URL for this paper:
<http://eprints.whiterose.ac.uk/90242/>

Version: Accepted Version

Article:

Song, H, Tong, J, Wignall, PB et al. (5 more authors) (2016) Early Triassic disaster and opportunistic foraminifers in South China. *Geological Magazine*, 153 (2). pp. 298-315. ISSN 0016-7568

<https://doi.org/10.1017/S0016756815000497>

Reuse

Unless indicated otherwise, fulltext items are protected by copyright with all rights reserved. The copyright exception in section 29 of the Copyright, Designs and Patents Act 1988 allows the making of a single copy solely for the purpose of non-commercial research or private study within the limits of fair dealing. The publisher or other rights-holder may allow further reproduction and re-use of this version - refer to the White Rose Research Online record for this item. Where records identify the publisher as the copyright holder, users can verify any specific terms of use on the publisher's website.

Takedown

If you consider content in White Rose Research Online to be in breach of UK law, please notify us by emailing eprints@whiterose.ac.uk including the URL of the record and the reason for the withdrawal request.



eprints@whiterose.ac.uk
<https://eprints.whiterose.ac.uk/>

Early Triassic disaster and opportunistic foraminifers in South China

Haijun Song*‡†, Jinnan Tong*†, Paul B. Wignall§,

Mao Luo||, Li Tian*, Huyue Song*, YunFei Huang¶, Daoliang Chu*

*State Key Laboratory of Biogeology and Environmental Geology, China University of Geosciences, Wuhan 430074, PR China

‡State Key Laboratory of Palaeobiology and Stratigraphy, Nanjing Institute of Geology and Palaeontology, Nanjing, 210008, PR China

§School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK

||School of Life and Environmental Sciences, Deakin University, Melbourne Burwood Campus, Burwood, Victoria 3125, Australia

¶School of Geoscience, Yangtze University, Wuhan 430100, PR China

†Author for correspondence: haijun.song@aliyun.com (H. Song), jntong@cug.edu.cn (J. Tong)

Abstract: Survival and recovery are important dynamic processes of biotic evolution during major geological transitions. Disaster and opportunistic taxa are two significant groups that dominate the ecosystem in the aftermath of mass extinction events. Disaster taxa appear immediately after such crises whilst opportunists predate the crisis but also bloom in the aftermath. This paper documents three disaster foraminiferal species and seven opportunistic foraminiferal species from Lower Triassic successions of South China. They are characterized by extreme high-abundance and low-diversity and occurred occasionally in the Griesbachian, Smithian, and Spathian. The characteristics (small size,

23 simple morphology) and stratigraphic ranges of these groups suggest that *r*-selection is a
24 commonly used strategy for survivors to cope with either harsh post-extinction conditions
25 and/or environments lacking incumbents.

26 Keywords: disaster, opportunist, foraminifers, Early Triassic, Permian-Triassic extinction

27

28 **1. Introduction**

29 For the survivors of mass extinctions, their fate can be highly variable but also to some
30 extent predictable. Groups with intrinsically high rates of extinction before the crisis often
31 radiate at high rates afterwards whilst evolutionary laggards often recover much more
32 slowly. This is exemplified by the ammonoids, a group characterized by exceptionally high
33 evolutionary rates throughout their history. Having suffered a severe extinction at the
34 Permian-Triassic (P-Tr) they radiated at typically high rates in the immediate aftermath
35 (Stanley, 2007). Similarly, the bivalves, the evolutionary carthorses of the marine
36 invertebrates, underwent little radiation in the aftermath of the P-Tr extinction with the
37 exception of the spectacular recovery of the claraiids, a family of “flat clams” belonging to
38 the Pterinopectinidae, that exhibit similarly high evolutionary rates before the extinction
39 (Yin, 1985).

40 One of the most interesting facets of the immediate post-extinction interval is the
41 presence of prolific abundances of opportunists, typically called disaster taxa (Harries,
42 Kauffman & Hansen, 1996; Kauffman & Harries, 1996). These are defined as species that
43 are adapted to the high stress conditions of an extinction crisis and its immediate aftermath
44 but are rare or absent at other times (Harries, Kauffman & Hansen, 1996; Kauffman &

45 Harries, 1996; Rodland & Bottjer, 2001). They thus differ from “normal” opportunists,
46 which have long species ranges, and often appear in high stress settings including (but not
47 restricted to) the aftermath of mass extinctions. Opportunists are, by definition, ecological
48 generalists that exhibit high fecundity – a facet of their lifestyle that is manifest by the
49 rapid attainment of sexual maturity and a small, simple morphology. Disaster taxa also
50 typically exhibit these features but their more restricted temporal distribution suggests that
51 they are suited to the specific and unusual conditions of the post-extinction interval.
52 Disaster taxa are evolutionary dead ends. However, they differ from a third category found
53 at this time – progenitor taxa which appear and radiate rapidly in the post-extinction
54 interval (Kauffman & Harries, 1996).

55 Several benthic biota have been considered as potential disaster forms in the aftermath
56 of the P-Tr extinction e.g. stromatolites (Schubert & Bottjer, 1992), lingulide brachiopods
57 (Rodland & Bottjer, 2001) and calcareous tubeworms (He *et al.*, 2012). Amongst the
58 foraminifers the small, tube-like *Earlandia* is regarded as a typical disaster genus found in
59 huge numbers immediately following both the Frasnian-Famennian (F-F) mass extinction
60 and the Permian-Triassic boundary (PTB) extinctions (Hallam & Wignall, 1997).
61 *Earlandia* is known in the aftermath of both P-Tr extinction pulses (Song *et al.*, 2013)
62 along with *Postcladella kahlori*. For example both these taxa are especially abundant in the
63 microbialite facies that developed following the latest Permian extinction in Turkey
64 (Altiner *et al.*, 1980; Altiner & Zaninetti, 1981; Groves, Altiner & Rettori, 2005), Italy
65 (Groves *et al.*, 2007) and South China (Song *et al.*, 2009). *Earlandia* also occurs in
66 wackestones the immediate aftermath of the earliest Triassic extinction at Meishan section

67 of South China (Wignall & Twitchett, 2002).

68 However, the ecological significance of disaster taxa is unclear. Traditionally,
69 opportunists should record high-stress environmental conditions and so, ostensibly, the
70 presence of post-extinction disaster forms could record the persistence of high-stress
71 conditions that caused the preceding mass extinction. Alternatively they may record the
72 expansion of hardy opportunists, capable of surviving the extinction episode, into vacated
73 environments once the environments had returned to normal. In this second alternative
74 disaster taxa fit a distinct ecological category (rather than a temporally-defined subset of
75 opportunist) – their success is due to their extinction-resistance but not to any specific
76 adaptation to the environments in which they find themselves in the post-extinction world.

77 The significance of disaster taxa is at the heart of a long-running debate on the delayed
78 recovery of benthic ecosystems in the Early Triassic in the aftermath of the P-Tr mass
79 extinction. Hallam (1991) was the first to note that the severity of the P-Tr mass extinction
80 and delayed recovery may be, in part, due to the prolongation of the harmful conditions that
81 triggered the extinction – specifically the extent and duration of global marine anoxia. In
82 contrast, Schubert & Bottjer (1992; 1995) noting the spread of stromatolites in Early
83 Triassic seas argued that they were filling an ecospace in which biotic factors (such as
84 gastropod grazing) were much reduced (and had yet to recover) but with normal, physical
85 environmental factors. Similarly, Rodland & Bottjer's (2001) work on lingulide
86 brachiopods in the Early Triassic of the western USA concluded that their proliferation
87 took place in well-oxygenated shelf seas. In contrast, Pruss & Bottjer (2004) and Fraiser &
88 Bottjer (2009) studied impoverished Early Triassic trace fossil assemblages in the same

89 strata and suggested repetition of stressful conditions. In support of this conclusion,
90 contemporaneous trace fossils from nearer shore strata in western Canada are much higher
91 diversity suggesting that there was indeed something stressful about offshore marine
92 settings in the Early Triassic (Zonneveld *et al.*, 2010).

93 In this study, we document the types, stratigraphic ranges and ecological behaviors of
94 disaster and opportunistic foraminifers during the biotic recovery from the P-Tr mass
95 extinction and address the issue of whether they were survivors living in a pleasant but
96 emptied nirvana or whether they were living in a harsh post-apocalyptic hell.

97

98 **2. Geological setting and studied sections**

99 We report on our analysis of the foraminifer content of the Lower Triassic successions
100 of South China and supplement our observations with literature records from elsewhere.
101 During the P-Tr transition, the South China block was located in the eastern Tethys near
102 the equator, consisting of islands, widespread shallow-water platforms and deep basins (Fig.
103 1). Numerous sections containing PTB strata and Early Triassic strata are known from
104 South China including the Global Stratotype Section and Point (GSSP) of the PTB —
105 Meishan (Yin *et al.*, 2001). Of these, Meishan, Huangzhishan, Yangou, Tieshikou,
106 Dongling, Cili, Wufeng, Shangsi, Liangfengya, Xiangkou, Dajiang, and Lekang sections
107 (Fig. 1) contain abundant disaster and opportunistic foraminifers and are selected herein to
108 study their stratigraphic ranges and palaeoenvironmental implications.

109

110 **2.a. Meishan section**

111 The Meishan section, the GSSP of the PTB, is situated 200 km west of Shanghai City,
112 eastern China (Fig. 1). The base of the Triassic is marked by the first occurrence of the
113 conodont *Hindeodus parvus* at the base of Bed 27c (Yin *et al.*, 2001). The PTB succession
114 immediately overlying the Changxing limestone, consists of two thin beds, a white clay
115 (Bed 25) and black shales (Bed 26), a wackestone (Bed 27), followed by a succession of
116 thinly interbedded succession of black shales, grey-green marls and pale grey micrites
117 interpreted to have accumulated in a generally dysoxic setting (Wignall & Hallam, 1993).

118 **2.b. Huangzhishan section**

119 The Huangzhishan section, located 40 km southeast of the classic Meishan section,
120 Zhejiang Province, eastern China (Fig. 1), records a similar P-Tr boundary succession. The
121 PTB strata (the Huangzhishan Formation), overlying the Changxing limestones, mainly
122 consists of marly limestones and marls. The first appearance datum (FAD) of *Hindeodus*
123 *parvus* at Huangzhishan is at the middle part of the Huangzhishan Formation, about 3.8 m
124 above the top of Changxing limestones (Chen, Henderson & Shen, 2008; Chen *et al.*, 2009).
125 The lowest Triassic strata include black shales and are thinly bedded and contain a low
126 diversity fauna with abundant *Claraia* and *Ophiceras* and small *Planolites* burrows
127 suggesting oxygen-restricted conditions once again (Chen *et al.* 2009).

128 **2.c. Yangou section**

129 The Yangou section, located in the northeast of the Yangou Coalmine, Leping County,
130 Jiangxi Province (Fig. 1), records a carbonate-dominated P-Tr boundary succession. The
131 top part of the Permian comprises a 12-m-thick massive packstone-grainstone, yielding
132 diverse fossil groups, e.g. calcareous algae, fusulinids, small foraminifers, and conodonts

133 (Song *et al.*, 2012a; Sun *et al.*, 2012a; Tian *et al.*, 2014b). The PTB succession is at the
134 lower part of the Daye Formation, mainly consisting of thin-bedded limestones containing
135 small foraminifers, ostracods, small gastropods, and conodonts (Zhu *et al.*, 1994; Sun *et al.*,
136 2012a). The base of the Triassic is marked by the FAD of *Hindeodus parvus* at the base of
137 Bed 21-4, about 21 cm above the base of the Daye Formation (Sun *et al.*, 2012a).

138 **2.d. Tieshikou section**

139 The Tieshikou section is located in the north of Zhaigao village, Xinfeng County,
140 Jiangxi Province (Fig. 1). The PTB succession immediately overlying the Changxing
141 limestones mainly comprises black shales and limestone lens with abundant conodonts and
142 brachiopods (Yang & Sun, 1990). The lowest Triassic strata include black shales with
143 limestone lens and are thinly bedded and contain a low diversity fauna with abundant
144 *Claraia* (Yang & Sun, 1990) suggesting oxygen-restricted conditions.

145 **2.e. Dongling section**

146 The Dongling section is situated in the northeast of Diaoyan village, Xiushui County,
147 Jiangxi Province (Fig. 1). The upper Changxing Formation is composed of massive
148 packstones and a 50 m-thick algae-sponge bindstone (reef), that contains diverse sponges,
149 corals, calcareous algae, fusulinids, small foraminifers, ostracods, and conodonts. The PTB
150 succession is at the lower part of the Daye Formation, mainly consisting of marly
151 limestones with conodonts, ostracods, gastropods, and small foraminifers (Zhu, 1999). The
152 FAD of *Hindeodus parvus* at Dongling section is at 25 cm above the top of Changxing
153 limestones (Zhu, 1999). The basal Triassic is a thinly interbedded succession of black
154 shales, grey-green marls and pale grey micrites.

155 **2.f. Cili section**

156 The Cili section, also called the Kangjiaping section, is situated near Kangjiaping
157 village of Cili County, Hunan Province (Fig. 1). It consists of a well-developed Upper
158 Permian coral-sponge reef sequence and the overlying PTB succession of calcimicrobialite
159 and oolite facies. The top of the coral-sponge reef succession is composed of skeletal
160 limestones yielding abundant fossils, e.g. calcareous algae, fusulinids, small foraminifers,
161 ostracods, and echinoderms (Wang *et al.*, 2009). The fusulinid *Palaeofusulina sinensis* and
162 many other species of this genus are found in the top of the latest Permian packstones
163 (Yang *et al.*, 2013). The PTB stratigraphic succession comprises calcimicrobialites, oolitic
164 grainstones, vermiculitic (bioturbated) limestones, thin-bedded intraclastic wackstones (Fig.
165 2), yielding ostracods, gastropods, small foraminifers, microconchids, and conodonts
166 (Wang *et al.*, 2009; Yang *et al.*, 2011). Compared to the other sections noted above, the
167 Early Triassic facies at Cili clearly record better oxygenation. The FAD of *Hindeodus*
168 *parvus* is in the upper part of the microbialite, about 4.5 m above the Changxing limestones
169 and the calcimicrobialites boundary (Wang *et al.*, 2009).

170 **2.g. Liangfengya section**

171 The Liangfengya section, also called the Beifengjing section, is located in the west of
172 Chongqing City, southwestern China (Fig. 1). The top part of the Permian is composed of a
173 60-m-thick massive bioclastic limestone, yielding abundant fossils such as foraminifers
174 (Tong & Kuang, 1990; Song, Tong & Chen, 2011), brachiopods (Shen & He, 1991),
175 calcareous algae, echinoids, and ostracods (Yang *et al.*, 1987; Wignall & Hallam, 1996).
176 The PTB is at the base of the Feixianguan Formation, which mainly comprises thin-bedded

177 limestones, marls, and claystones that are frequently pyritic (Fig. 2). Bivalves, brachiopods,
178 and small foraminifers are generally common. Tiny burrows are present but these have not
179 disrupted the cm-scale bedding in the unit and the overall depositional setting is considered
180 to be dysoxic (Wignall & Hallam, 1996; Wignall & Twitchett, 1999).

181 **2.h. Dajiang section**

182 The Dajiang section is situated in the middle part of an isolated carbonate platform
183 called the Great Bank of Guizhou in the Nanpanjiang basin of southwest China (Lehrmann,
184 Wei & Enos, 1998). A series of PTB sections are well exposed from the southeast to
185 northwest (from platform facies to basin facies), i.e. Dawen, Heping, Dajiang, Rongbo,
186 Langbai, Mingtang, Guandao, Bianzhonglu, and Bianyang sections. The Dajiang section
187 records a typical facies transition at the PTB: fossiliferous packstones of the Wuchiaping
188 Formation are succeeded by earliest Triassic microbialites of the Daye Formation which
189 contain a diverse ostracod fauna that indicates conditions were well oxygenated (Forel *et*
190 *al.*, 2009).

191 **2.i. Wufeng section**

192 The Wufeng section is situated in the Wufeng County of western Hubei Province (Fig.
193 1). During the P-Tr transition, Wufeng is located in the northern margin of the Yangtze
194 Platform. The latest Permian Dalong Formation consists of siliceous limestone and black
195 shales. The Lower Triassic sequence is composed of the Daye and Jialingjiang formations.
196 Of these, the Daye Formation consists of thinly laminated shales in its lower half and
197 medium- to thick-bedded limestones in its upper part. The Jialingjiang Formation
198 comprises interbeds of dolomite units and limestone units (Fig. 3).

199 **2.j. Shangsi section**

200 As one of the candidate GSSPs of the PTB, the Shangsi section contains one of the
201 most detailed records of events during the P-Tr mass extinction in a deep basinal setting (Li
202 *et al.*, 1989; Wignall *et al.*, 1995; Lai *et al.*, 1996). In the Early Triassic, Sichuan occupied
203 the northwestern margin of the Yangtze Platform (Fig. 1). The Shangsi section is located
204 30 km west of Guangyuan City, northern Sichuan Province (Fig. 1). Over 1200 m of strata,
205 spanning the entire Late Permian and Early Triassic, are continuously exposed. The latest
206 Permian Dalong Formation mainly consists of interbeds of limestones, cherts and dark
207 shales with pervasive bioturbation suggesting well oxygenated conditions (Wignall *et al.*,
208 1995). The Lower Triassic sequence is composed, in ascending order, of the Feixianguan,
209 Tongjiezi and Jialingjiang formations (Fig. 3). Of these, the Feixianguan Formation is
210 characterized by a 3.5-m-thick siliceous marly limestone at its base followed by a
211 95-m-thick unit of limestone and a 685-m-thick black shales. The siliceous marl is thinly
212 laminated, pyritic and interpreted to be a dysoxic-anoxic facies (Wignall *et al.*, 1995).
213 Higher levels in the Formation are dominated by chocolate-coloured marls and thin micrite
214 interbeds together with storm-generated flat-pebble conglomerates (Wignall & Twitchett,
215 1999).

216 **2.k. Xiangkou section**

217 The Xiangkou section is situated in the Xiangkou Town, Zunyi City, northern Guizhou
218 Province (Fig. 1). In the Early Triassic, Xiangkou occupied the southwestern margin of the
219 Yangtze Platform. Over 1200 m strata, spanning the latest Permian to Middle Triassic, are
220 continuously exposed. The latest Permian Changxing Formation consists of dark grey

221 cherty limestone. The Lower Triassic sequence is composed, in ascending order, of the
222 Yelang and Maocaopu formations (Fig. 3). Of these, the Yelang Formation is characterized
223 by a 15-m-thick marl at its base followed by a 175 m-thick unit of limestone and a 160
224 m-thick shales. The marl is thinly laminated and contains a low diversity fauna with
225 abundant *Claraia* and *Lingula* suggesting oxygen-restricted conditions. Higher levels in the
226 Formation are dominated by thin micrite interbeds together with storm-generated
227 flat-pebble conglomerates and chocolate-coloured marls. The Maochaopu Formation is
228 characterized by pale grey, medium- to thick- bedded micrite in its lower and middle part
229 and thick-dolomite in its top part.

230 **2.1. Lekang section**

231 The Lekang section is situated at the Lekang village of the Wangmo County, Guizhou
232 Province (Fig. 1). In the Early Triassic, Lekang section is located in the northern margin of
233 the Nanpanjiang Basin. The latest Permian Linghao Formation consists of interbeds of
234 limestones, cherts and dark shales with pervasive bioturbation, suggesting well oxygenated
235 conditions. The Lower Triassic sequence is composed of the Luolou Formation which, in
236 its lowest part, is dominated by unbioturbated laminated black shales and overlying thinly
237 bedded micrite interbeds.

238 **3. Disaster and opportunistic foraminifers**

239 A total of nine disaster and opportunistic foraminiferal species were identified from the
240 12 Lower Triassic sections in South China, i.e. *Postcladella kahlori*, *Earlandia* sp.,
241 *Globivalvulina lukachiensis*, *Hemigordiellina regularia*, *Hoyenella* spp., *Arenovidalina*
242 *chialingchiangensis*, *Aulotortus? bakonyensis*, *Triadodiscus eomesozoicus*, *Meandrospira*

243 *pusilla*. These are typical disaster and opportunistic forms that are prolifically common in
244 some beds after the P-Tr crisis (Figs. 2, 3).

245 **3.a. *Postcladella kahlori***

246 *Postcladella kahlori* (Brönnimann, Zaninetti & Bozorgnia, 1972) is the almost unique
247 taxon of foraminifera in the earliest Triassic. It has usually been identified as
248 "*Rectocornuspira kahlori*" (e.g. Groves *et al.*, 2005; 2007; Song *et al.*, 2009). This taxon
249 has an initial planispiral coiling part and an uncoiled last whorl (Fig. 4). Krainer & Vachard
250 (2011) designated this taxon as *Postcladella kahlori*. *P. kahlori*, as one of most common
251 disaster foraminifer in the Early Triassic, has been found in the base of microbialite at
252 Taskent section of Turkey (Altiner *et al.*, 1980; Altiner & Zaninetti, 1981; Groves, Altiner
253 & Rettori, 2005), in the lower Werfen Formation of northern Italy (Groves *et al.*, 2007) and
254 southern Austria (Krainer & Vachard, 2011), and the base of Lower Triassic at Lukač
255 section of western Slovenia (Nestell *et al.*, 2011). In South China, *P. kahlori* was found in
256 the earliest Triassic microbialite of the Dajiang section (Song *et al.*, 2009; Yang *et al.*, 2011)
257 and in the Langpai section (Ezaki *et al.*, 2008) in Guizhou Province, Cili section in Hunan
258 Province (Fig. 2), and Dongwan section in Sichuan Province (Ezaki, Liu & Adachi, 2003).
259 *P. kahlori* was also found in other shallow-water facies in the earliest Triassic, e.g. the
260 lowest Daye Formation at Dongling section of Jiangxi Province and the lower Yelang
261 Formation at Xiangkou section of Guizhou Province (Figs. 3, 4). Therefore, *P. kahlori* is a
262 typical and widespread disaster form that bloomed instantaneously in the Palaeotethys after
263 the P-Tr extinction. It is found in a range of environments spanning oxygenated,
264 shallow-water facies (e.g. Dajiang and Cili sections) and deeper, dysoxic facies (e.g.

265 Werfen Formation).

266 **3.b. *Earlandia* sp.**

267 *Earlandia* sp. is a tube-like foraminifer with a globular proloculus followed by a long,
268 straight, undivided tubular chamber. It is a common disaster taxon that bloomed
269 immediately in the aftermath of the latest Permian extinction and earliest Triassic
270 extinction (Table 1), as first identified by Hallam & Wignall (1997). *Earlandia* has been
271 found in the Permian-Triassic boundary interval of Demirtas and Taskent sections of
272 Turkey (Altiner *et al.*, 1980; Altiner, Groves & Özkan-Altiner, 2005; Groves & Altiner,
273 2005; Groves, Altiner & Rettori, 2005), Bulla and Tesero sections of northern Italy (Groves
274 *et al.*, 2007), and Andreasstrasse and Suchagraben sections in southern Austria (Krainer &
275 Vachard, 2011). In South China, *Earlandia* sp. bloomed during the conodont *Hindeodus*
276 *parvus* Zone in shallow-water sections, e.g. the microbialite of Cili and Dajiang sections
277 and other shallow-water facies such as Yangou, Dongling, and Tieshikou sections, and in
278 the *Isarcicella isarcica* Zone in platform margin and slope facies such as Liangfengya and
279 Meishan sections (Table 1). In this study, we also found that *Earlandia* sp. was very
280 abundant in one bed of the Maochaopu Formation of Xiangkou section in Guizhou
281 Province, South China (Fig. 5d). This is the first report of this opportunistic form from
282 Spathian strata.

283 **3.c. *Globivalvulina lukachiensis***

284 *Globivalvulina lukachiensis*, a new species for the *Globivalvulina* genus was
285 established by Nestell *et al.* (2011). *Globivalvulina lukachiensis*, rather small, planispirally
286 coiled with a biserial chamber arrangement, is a common foraminiferal species in the Late

287 Permian, and has been found in South China (see "*Globivalvulina bulloides*" in Song *et al.*,
288 2007; Song *et al.*, 2009), western Slovenia – where it occurs in Late Permian facies but not
289 after the mass extinction (Nestell *et al.*, 2011), northwestern Caucasus (see "*Globivalvulina*
290 *araxensis*" in Pronina-Nestell & Nestell, 2001). It survived the latest Permian extinction
291 and is found in the microbialite of Dajiang section (Song *et al.*, 2009) and in the *Hindeodus*
292 *parvus* Zone of Meishan section (Song *et al.*, 2007; Song, Tong & Chen, 2009), and in the
293 earliest Induan of Turkey as failed survivor (see "*Globivalvulina aff. cyprica*" in Altiner,
294 Groves & Özkan-Altiner, 2005). However, their abundance is very low at these two
295 sections and does not show any characteristics of a disaster taxon. In this study, we found
296 *Globivalvulina lukachiensis* with a high abundance at the base of microbialite from Cili
297 section (Fig. 6), indicating a typical disaster taxon's characteristics.

298 **3.d. *Hemigordiellina regularia***

299 *Hemigordiellina*, small glomospiroid porcelaneous test with a proloculus followed by
300 undivided tubular second chamber that is streptospirally coiled in a somewhat irregular
301 manner, is a controversial taxon (p. 85 in Gaillot & Vachard, 2007). For its glomospiroid
302 test, lots of species with calcareous tests have been attributed to *Glomospira*, e.g.
303 *Glomospira* sp. and *Glomospira regularis* from Meishan section (Song *et al.*, 2007),
304 *Glomospira* spp. from Nanpanjiang Basin (Song *et al.*, 2009; Song *et al.*, 2011),
305 *Glomospira* sp. from Japan (Kobayashi, 2004; Kobayashi, 2012). But *Glomospira* is an
306 agglutinated foraminifer (Loeblich & Tappan, 1988) and so this name is inappropriate. In
307 this study, glomospiroid porcelaneous species are attributed to *Hemigordiellina* Marie in
308 Deleau & Marie, 1961.

309 *Hemigordiellina regularia* is one of the most common foraminiferal taxa in the Early
310 Triassic strata (Song *et al.*, 2011). *Hemigordiellina regularia* has a long geological range,
311 from Early Permian to latest Triassic (Gaillot & Vachard, 2007). It has an extensive
312 distribution in the Late Permian with a low abundance, e.g. South China (Song *et al.*, 2009),
313 Tibet (Song's unpublished data), Middle East (Gaillot & Vachard, 2007), and Japan
314 (Kobayashi, 2012). However, a large number of *Hemigordiellina regularia* specimens
315 appear suddenly in some Early Triassic beds from South China, e.g. upper Maochaopu
316 Formation of Xiangkou section and lower Jialingjiang Formation of Wufeng section (Fig.
317 7), showing that *Hemigordiellina regularia* is an opportunistic form that appeared in the
318 late Early Triassic.

319 **3.e. *Hoyenella* spp.**

320 *Hoyenella* with its small porcelaneous test is homeomorphic with the agglutinating
321 *Glomospirella*. A lots of species with glomospirellid-like calcareous tests have been
322 attributed to *Glomospirella*, e.g. *Glomospirella irregularis*, *Glomospirella spirillinoides*,
323 *Glomospirella ammodiscoidea*, *Glomospirella shengi*, *Glomospirella vulgaris*, and
324 *Glomospirella facilis* from Jialingjiang Limestone of Sichuan Province (Ho, 1959),
325 *Glomospirella lampangensis* from Lampang Group of Northern Thailand (Kobayashi *et al.*,
326 2006), and *Glomospirella* spp. from Pakistan (Zaninetti & Brönnimann, 1975). In this
327 study, glomospirelloid porcelaneous species are attributed to *Hoyenella* Rettori, 1994.
328 *Hoyenella* is one of most common foraminiferal taxa in the Early Triassic strata of South
329 China (Song *et al.*, 2011). In this study, we found that a large number of *Hoyenella* spp.
330 specimens occurred suddenly in some Early Triassic beds in South China, mostly in

331 dysoxic settings, e.g. Wufeng, Xiangkou, and Shangsi sections (Figs. 3, 8; Table 1).

332 **3.f. *Arenovidalina chialingchiangensis***

333 *Arenovidalina chialingchiangensis* was firstly found in the Lower Triassic Jialingjiang
334 Limestone of South China (Ho, 1959). Subsequently, this species was reported world wide,
335 e.g. the Albarracín Formation (Anisian) of Spain (Horwitz & Pidgeon, 1993), the
336 Olenekian and Anisian strata of Karst Dinarides (Velić, 2007), and the Lower Triassic
337 Tütünlüktepe Formation of northwest Turkey (Okuyucu *et al.*, 2014). In this study, a large
338 number of *Arenovidalina chialingchiangensis* specimens appeared suddenly in some Early
339 Triassic beds from South China, i.e. upper Daye Formation of Wufeng section and the
340 basal Tongjiezi Formation of Shangsi section (Fig. 9), showing that *Arenovidalina*
341 *chialingchiangensis* is an opportunistic form that appeared in the Olenekian (Fig. 3).

342 **3.g. *Aulotortus? bakonyensis***

343 *Aulotortus? bakonyensis* was first reported in the Jurassic strata of the Dogger of
344 Hungary (Blau, 1989). Here, we found abundant *Aulotortus? bakonyensis* in two thin-beds
345 of upper Tongjiezi Formation of Shangsi section (Fig. 3). These two thin-beds contain
346 hundreds of specimens of *Aulotortus? bakonyensis*, showing that *Aulotortus? bakonyensis*
347 is an opportunistic taxon that occurred occasionally in the late Early Triassic. The thickness
348 of each thin-bed is only several millimeters (Fig. 10a, b), suggesting that this opportunistic
349 taxon bloomed each time for only a very short period.

350 **3.h. *Triadodiscus eomesozoicus***

351 *Triadodiscus eomesozoicus*, an involutinid-like form with , was originally established
352 by Oberhauser (1957) from the Carnian of the eastern Alps. It is a common foraminiferal

353 species in the Triassic oceans, and has been found in Egypt (Kuss, 1988), Tunisia (Kamoun
354 *et al.*, 2001), southern Spain (Pérez-López, Márquez & Pérez-Valera, 2005), Japan
355 (Kobayashi, Martini & Zaninetti, 2005), northern Thailand (Kobayashi *et al.*, 2006), and
356 Timor (Haig & McCartain, 2012). Although most specimens of *Triadodiscus eomesozoicus*
357 have been found in the Middle and Late Triassic, it firstly appeared in the late part of the
358 Early Triassic (Márquez, 2005). In this study, one *Triadodiscus eomesozoicus* bed was
359 found in the Early Triassic Tongjiezi Formation at Shangsi section (Fig. 3). In this bed,
360 *Triadodiscus eomesozoicus* is abundant but poorly preserved (Fig. 10c, d).

361 **3.i. *Meandrospira pusilla***

362 *Meandrospira pusilla* is one of most common foraminiferal species in the Early and
363 Middle Triassic. It has been found in Greece (Rettori, Angiolini & Muttoni, 1994), Italy
364 (Zaninetti, Rettori & Martini, 1994), Austria (Krainer & Vachard, 2011), Eastern
365 Carpathians (Popescu & Popescu, 2005), Tunisia (Kilani-Mazraoui, Razgallah-Gargouri &
366 Mannai-Tayech, 1990), Rumania (Bucur, Strutinski & Paica, 1997), Northern United Arab
367 Emirates (Maurer, Rettori & Martini, 2008), Western Caucasus and Eastern Precaucasus
368 (Vuks, 2007), Iran (Baud, Bronnimann & Zaninetti, 1974), Japan (Kobayashi, Martini &
369 Zaninetti, 2005), and South China (Ho, 1959; He, 1988; 1993; Song *et al.*, 2011). In this
370 study, we found that a large number of *Meandrospira pusilla* specimens appeared
371 instantaneously in some Lower Triassic beds in South China, e.g. lower Jialingjiang
372 Formation of Wufeng section, upper Tongjiezi Formation of Shangsi section, Lekang
373 Formation of Lekang section, and upper Maochaopu Formation of Xiangkou section (Fig.
374 11), showing that *Meandrospira pusilla* is an opportunistic form that appeared in the late

375 Early Triassic.

376 **4. Temporal distribution of disasters and opportunists**

377 **4.a. Temporal distribution of disasters**

378 In this study, we found that disaster foraminifers occurred in the immediate aftermath
379 of the P-Tr extinction (Fig. 12). Group *Postcladella kahlori-Earlandia* sp., usually
380 dominated by abundant *Postcladella kahlori*, *Earlandia* sp., *Globivalvulina lukachiensis*,
381 and rare *Nodosaria expolita*, occurred in the microbialites at Dajiang and Cili sections that
382 followed the latest Permian mass extinction. This foraminiferal group has also been
383 reported in the basal Triassic microbialites at Dongwan section of South China (Ezaki, Liu
384 & Adachi, 2003), Taskent and Taurides sections of Turkey (Altiner *et al.*, 1980; Altiner &
385 Zaninetti, 1981; Ünal *et al.*, 2003; Groves, Altiner & Rettori, 2005), and Bulla section of
386 Italy (Groves *et al.*, 2007). These foraminifers co-occurred with other disaster taxa such as
387 cyanobacteria (Ezaki, Liu & Adachi, 2003; Wang *et al.*, 2005), worm tubes (polychaete
388 *Spirorbis*), and microgastropods (Yang *et al.*, 2011). Another disaster group dominated by
389 *Earlandia* sp. has been identified above the earliest Triassic extinction horizon, e.g. Bed 29
390 at Meishan section and Beds 21c and 23 at Liangfengya section. Several peaks in the
391 abundance of *Earlandia* sp. have been recorded coinciding with an abrupt extinction of
392 foraminifers during the earliest Triassic crisis (Song, Tong & Chen, 2009; Song *et al.*,
393 2013b).

394 **4.b. Temporal distribution of opportunists**

395 In the Dienerian, we did not find any opportunistic or disaster foraminifers in South
396 China. The opportunistic fauna dominates within the Smithian and Spathian and is

397 characterized by the extremely prosperous *Hemigordiellina* and *Hoyenella* (Fig. 12). The
398 number of *Hemigordiellina regularia* and *Hoyenella* spp. specimens exceeds 200 in a
399 2.2×2.2 cm² thin-section in some levels at Shangsi section (Fig. 13). The lower boundary of
400 the opportunistic fauna interval is defined by the horizon where the *Arenovidalina*
401 *chialingchiangensis* first bloomed. The upper boundary of the opportunistic fauna interval
402 is defined by the horizon where the relative abundance of the opportunistic group decreases
403 to less than 50%.

404 Smithian opportunists are divided into three groups based on the stratigraphic ranges.
405 The first group dominated by *Arenovidalina chialingchiangensis* occurred in the lower
406 Tongjiezi Formation (Fig. 3). The second group dominated by *Hemigordiellina regularia*,
407 *Hoyenella* spp., and *Meandrospira pusilla* occurred in the middle Tongjiezi Formation (Fig.
408 3). The third group dominated by *Aulotortus? bakonyensis*, *Triadodiscus eomesozoicus*
409 occurred in the upper Tongjiezi Formation (Fig. 3). Opportunistic fauna in the Spathian
410 consists of *Earlandia* sp., *Hemigordiellina regularia*, *Hoyenella* spp., and *Meandrospira*
411 *pusilla* (Fig. 12).

412 **5. Survival strategy response to stressed environments**

413 Opportunistic taxa usually take advantage of high-stress, strongly fluctuating
414 environments as a result of dramatic changes in oceanic ecosystems. As such they are
415 capable of prolific population expansion and rapid biogeographical dispersal into stressed
416 environments (Harries, Kauffman & Hansen, 1996; Kauffman & Harries, 1996). The
417 bloom of opportunistic foraminifers coincided with the Early Triassic stressed
418 environments that have been frequently reported in recent years, e.g. widespread and

419 long-term anoxia (Wignall & Twitchett, 2002; Song *et al.*, 2012b), high sea surface
420 temperature (Joachimski *et al.*, 2012; Sun *et al.*, 2012b), intensified water-column
421 stratification (Song *et al.*, 2012a; Song *et al.*, 2013a), and expansion of oceanic oxygen
422 minimum zone (Algeo *et al.*, 2011; Song *et al.*, 2014; Tian *et al.*, 2014a).

423 In this study, we found that disaster foraminifers develop relatively large populations in
424 the early survival interval. They are replaced by opportunistic foraminifers and other
425 survivors early in the following repopulation period. Both disaster and opportunistic
426 foraminifers had a very short time span, and occurred repeatedly in the Early Triassic.
427 These beds containing disaster and opportunistic foraminifers usually have a low diversity
428 but a high abundance (Fig. 13). The 'normal' species (including these Late Permian and
429 Triassic foraminifers that have been reported in Song *et al.*, 2007; 2009a, b; 2011a, b; 2015
430 and some unpublished data at Dongling, Tieshikou, Cili, and Bianyang sections) beds
431 usually have a moderate diversity with a moderate abundance (see Fig.13). These disaster
432 and opportunistic taxa are very small compared to pre-extinction forms (Payne *et al.*, 2011;
433 Song, Tong & Chen, 2011; Rego *et al.*, 2012). All of these traits characterize *r*-selection
434 strategy, i.e. high fecundity, small body size, short generation time, and wide offspring
435 dispersion. When the environmental conditions tended to get better in the middle to late
436 Spathian, larger, more diverse K-selection foraminifers began to dominate the benthic
437 ecosystem.

438 **6. Conclusion**

439 Three disaster foraminiferal species were identified in the immediate aftermath of the
440 P-Tr mass extinction, i.e. *Postcladella kahlori*, *Earlandia* sp., and *Globivalvulina*

441 *lukachiensis*. Among them, *Postcladella kahlori* and *Earlandia* sp. have also been found as
442 disaster species in many other regions around the world. As such, the bloom (rather than
443 the occurrence) of these disaster forms could be used as evidence of post-extinction strata
444 in the case of lacking conodonts and ammonoids.

445 Disaster fauna were replaced by opportunistic fauna in the Smithian and Spathian.
446 Opportunistic fauna is composed of *Earlandia* sp., *Hemigordiellina regularia*, *Hoyenella*
447 spp., *Arenovidalina chialingchiangensis*, *Aulotortus? bakonyensis*, *Triadodiscus*
448 *eomesozoicus*, and *Meandrospira pusilla*. These opportunistic fauna are the main
449 component of the recovery fauna (see Song *et al.*, 2011b) and the relative abundance
450 decreases to less than 50% of foraminifers in the middle-late Spathian, in accord with the
451 improvement of marine environments.

452 Disaster and opportunistic foraminifers have often been found in the aftermath of many
453 extinction events in the Phanerozoic. This phenomenon shows *r*-selection is a commonly
454 used strategy for survivors to cope with the catastrophe events. After the crisis, many
455 opportunists live in limited area while others are likely to choose K-selection strategy and
456 become the dominant groups during the recovery interval.

457

458 **Acknowledgements**

459 This study was supported by the 973 Program (2011CB808800), the National Natural
460 Science Foundation of China (41302271, 41272372), State Key Laboratory of
461 Palaeobiology and Stratigraphy, Nanjing Institute of Geology and Palaeontology (133111),
462 the 111 Project (B08030), BGEG (GBL11202, GBL11302), and the Fundamental Research

463 Funds for the Central Universities (CUG130407).

464

465 **References**

466 ALGEO, T. J., CHEN, Z. Q., FRAISER, M. L. & TWITCHETT, R. J. 2011. Terrestrial-marine teleconnections in the

467 collapse and rebuilding of Early Triassic marine ecosystems. *Palaeogeography, Palaeoclimatology,*

468 *Palaeoecology* 308(1–2), 1–11.

469 ALTINER, D., BAUD, A., GUEX, J. & STAMPFLI, G. 1980. La limite Permien-Trias dans quelques

470 localités du Moyen-Orient: recherches stratigraphiques et micropaléontologiques. *Rivista Italiana di*

471 *Paleontologia* 85, 3–4, 683–714.

472 ALTINER, D. 1981. *Recherches stratigraphiques et micropaléontologiques dans le Taurus Oriental au NW de*

473 *Pinarbasi (Turquie)*. Thèse de l'Université de Genève.

474 ALTINER, D. & ZANINETTI, L. 1981. Le Trias dans la région de Pinarbasi, Taurus oriental, Turquie: unités

475 lithologiques, micropaléontologie, milieux de dépôt. *Rivista Italiana di Paleontologia* 86, 4, 705–760.

476 ALTINER, D., GROVES, J.R. & ÖZKAN-ALTINER, S., 2005. Calcareous foraminiferal recovery from the

477 end-Permian mass extinction, southern Turkey. *Albertiana* 33, 14–17.

478 BAUD, A., BRÖNNIMANN, P. & ZANINETTI, L. 1974. Sur la présence de *Meandrospira pusilla* (Ho)

479 (Foraminifère), dans le Trias inférieur de Kuh-e-Ali Bashi, Julfa, NW Iran. *Palaeontologische Zeitschrift*

480 48(3–4), 205–13.

481 BLAU, J. 1989. *Aulotortus* (?) *bakonyensis* n. sp. (Involutinina, Foraminifera) from the Dogger of Hungary.

482 *Neues Jahrbuch für Geologie und Paläontologie, Monatshefte* 1989(8), 459–66.

483 BRÖNNIMANN, P., ZANINETTI, L. & BOZORGNIA, F. 1972. Triassic (Skythian) smaller foraminifera from the

484 Elika Formation of the central Alborz, northern Iran, and from the Siusi Formation of the Dolomites,

- 485 northern Italy. *Mitteilung Gesellschaft der Geologie und Bergbaustudenten, Innsbruck* 21, 861–84.
- 486 BUCUR, I. I., STRUTINSKI, C. & PAICA, M. 1997. A new occurrence of Triassic deposits NE of Oravita
487 (Southern Carpathians, Rumania) and its paleotectonic significance. *Geologica Carpathica* 48, 39–48.
- 488 CHEN, J., HENDERSON, C. M. & SHEN, S. 2008. Conodont succession around the Permian-Triassic boundary at
489 the Huangzhishan section, Zhejiang and its stratigraphic correlation. *Acta Palaeontologica Sinica* 47(1),
490 91–114.
- 491 CHEN, Z. Q., TONG, J., ZHANG, K., YANG, H., LIAO, Z., SONG, H. & CHEN, J. 2009. Environmental and biotic
492 turnover across the Permian-Triassic boundary on a shallow carbonate platform in western Zhejiang,
493 South China. *Australian Journal of Earth Sciences* 56(6), 775–97.
- 494 DELEAU, P. & MARIE, P. 1961. Les fusulinides du Westphalien C du Bassin d'Abadla et quelques autres
495 foraminifères du Carbonifère algérien (région de Colomb-Bechar). *Bull. Service Carte Geol. Algeria, ns*
496 25, 43–160.
- 497 EZAKI, Y., LIU, J. & ADACHI, N. 2003. Earliest Triassic Microbialite Micro- to Megastructures in the Huaying
498 Area of Sichuan Province, South China: Implications for the Nature of Oceanic Conditions after the
499 end-Permian Extinction. *Palaios* 18(4–5), 388–402.
- 500 EZAKI, Y., LIU, J., NAGANO, T. & ADACHI, N. 2008. Geobiological aspects of the earliest Triassic
501 microbialites along the southern periphery of the tropical Yangtze platform: Initiation and cessation of a
502 microbial regime. *Palaios* 23, 356–69.
- 503 FENG, Z., BAO, Z. & LIU, S. 1997. *Lithofacies Palaeogeography of Early and Middle Triassic of South China*.
504 Beijing: Petroleum Industry Press.
- 505 FOREL, M. B., CRASQUIN, S., KERSHAW, S., FENG, Q. L. & COLLIN, P. Y. 2009. Ostracods (Crustacea) and
506 water oxygenation in the earliest Triassic of South China: implications for oceanic events at the

- 507 end-Permian mass extinction. *Australian Journal of Earth Sciences* 56(6), 815–23.
- 508 FRAISER, M. L. & BOTTJER, D. J. 2009. Opportunistic behaviour of invertebrate marine tracemakers during the
509 Early Triassic aftermath of the end-Permian mass extinction. *Australian Journal of Earth Sciences* 56(6),
510 841–857.
- 511 GAILLOT, J. & VACHARD, D. 2007. The Khuff Formation (Middle East) and time-equivalents in Turkey and
512 South China: biostratigraphy from Capitanian to Changhsingian times (Permian), new foraminiferal taxa,
513 and palaeogeographical implications. *Coloquios de Paleontología* 57, 37–223.
- 514 GROVES, J. R. & ALTINER, D. 2005. Survival and recovery of calcareous foraminifera pursuant to the
515 end-Permian mass extinction. *Comptes Rendus Palevol* 4(6–7), 487–500.
- 516 GROVES, J. R., ALTINER, D. & RETTORI, R. 2005. Extinction, survival, and recovery of lagenide foraminifers
517 in the Permian-Triassic boundary interval, Central Taurides, Turkey. *Journal of Paleontology* 79 suppl.,
518 1–38.
- 519 GROVES, J. R., RETTORI, R., PAYNE, J. L., BOYCE, M. D. & ALTINER, D. 2007. End-Permian mass extinction of
520 Lagenide foraminifers in the Southern Alps (Northern Italy). *Journal of Paleontology* 81(3), 415–34.
- 521 HAIG, D. W. & MCCARTAIN, E. 2012. Intraspecific variation in Triassic Ophthalmidiid Foraminifera from
522 Timor. *Revue de Micropaléontologie* 55(2), 39–52.
- 523 HALLAM, A. & WIGNALL, P. B. 1997. *Mass Extinctions and their Aftermath*. Oxford: Oxford University
524 Press.
- 525 HARRIES, P. J., KAUFFMAN, E. G. & HANSEN, T. A. 1996. Models for biotic survival following mass
526 extinction. In *Biotic Recovery from Mass Extinction Events* (ed M. B. Hart). pp. 41–60. Geological
527 Society Special Publication.
- 528 HE, L., WANG, Y., WOODS, A., LI, G., YANG, H. & LIAO, W. 2012. Calcareous tubeworms as disaster forms

- 529 after the end-Permian mass extinction in South China. *Palaios* 27(12), 878–86.
- 530 HE, Y. 1988. Early and Middle Triassic foraminifera from Jiangsu and Anhui Provinces, China. *Acta*
531 *Micropalaeontologica Sinica* 5(1), 85–92 (in Chinese with English abstract).
- 532 HE, Y. 1993. Triassic foraminifera from Northeast Sichuan and South Shanxi, China. *Acta Palaeontologica*
533 *Sinica* 32(2), 170–87 (in Chinese with English abstract).
- 534 HO, Y. 1959. Triassic foraminifera from the Chialingkiang limestone of south Szechuan. *Acta*
535 *Palaeontologica Sinica* 7(5), 387–418.
- 536 HORWITZ, R. & PIDGEON, R. 1993. 3.1 Ga tuff from the Sholl Belt in the West Pilbara: further evidence for d
537 diachronous volcanism in the Pilbara Craton of Western Australia. *Precambrian research* 60(1), 175–83.
- 538 JOACHIMSKI, M. M., LAI, X., SHEN, S., JIANG, H., LUO, G., CHEN, B., CHEN, J. & SUN, Y. 2012. Climate
539 warming in the latest Permian and the Permian–Triassic mass extinction. *Geology* 40(3), 195–98.
- 540 KAMOUN, F., PEYBERNÈS, B., CISZAK, R. & CALZADA, S. 2001. Triassic palaeogeography of Tunisia.
541 *Palaeogeography, Palaeoclimatology, Palaeoecology* 172(3–4), 223–42.
- 542 KAUFFMAN, E. G. & HARRIES, P. J. 1996. The importance of crisis progenitors in recovery from mass
543 extinction. In *Biotic Recovery from Mass Extinction Events* (ed M. B. Hart). pp. 15–39. Geological
544 Society Special Publication.
- 545 KILANI-MAZRAOUI, F., RAZGALLAH-GARGOURI, S. & MANNAI-TAYECH, B. 1990. The Permo-Triassic of
546 Southern Tunisia - biostratigraphy and palaeoenvironment. *Review of Palaeobotany and Palynology*
547 66(3–4), 273–91.
- 548 KOBAYASHI, F. 2004. Late Permian foraminifers from the limestone block in the southern Chichibu terrane of
549 West Shikoku, SW Japan. *Journal of Paleontology* 78(1), 62–70.
- 550 KOBAYASHI, F. 2012. Middle and Late Permian Foraminifers from the Chichibu Belt, Takachiho Area,

- 551 Kyushu, Japan: Implications For Faunal Events. *Journal of Paleontology* 86(4), 669–87.
- 552 KOBAYASHI, F., MARTINI, R., RETTORI, R., ZANINETTI, L., RATANASTHIEN, B., SAEGUSA, H. & NAKAYA, H.
553 2006. Triassic foraminifers of the Lampang Group (Northern Thailand). *Journal of Asian Earth Sciences*.
554 KOBAYASHI, F., MARTINI, R. & ZANINETTI, L. 2005. Anisian foraminifers from allochthonous limestones of
555 the Tanoura formation (Kurosegawa Terrane, West Kyushu, Japan). *Geobios* 38(6), 751–63.
- 556 KRAINER, K. & VACHARD, D. 2011. The Lower Triassic Werfen Formation of the Karawanken Mountains
557 (Southern Austria) and its disaster survivor microfossils, with emphasis on *Postcladella* n. gen.
558 (Foraminifera, Miliolata, Cornuspirida). *Revue de Micropaléontologie* 54(2), 59–85.
- 559 KUSS, J. 1988. Microfacies and foraminifera of Middle Triassic limestones (Anisian-Carnian?) from Gebel
560 Araif el Naqa (Sinai, Egypt). *Facies* 19(1), 61–75.
- 561 LAI, X. L., YANG, F., HALLAM, A. & WIGNALL, P. B. 1996. The Shangsi section candidate of the Global
562 Stratotype section and point of the Permian-Triassic boundary. In *The Paleozoic-Mesozoic Boundary*
563 *Candidates of Global Stratotype Section and Point of the Permian-Triassic Boundary* (ed H. Yin). pp.
564 113–24. Wuhan: China University of Geosciences Press.
- 565 LEHRMANN, D. J., WEI, J. & ENOS, P. 1998. Controls on facies architecture of a large Triassic carbonate
566 platform: the Great Bank of Guizhou, Nanpanjiang Basin, South China. *Journal of Sedimentary Research*
567 68(2), 311–26.
- 568 LI, Z., ZHAN, L., DAI, J., JIN, R., ZHU, X., ZHANG, J., HUANG, H., XU, D., YAN, Z. & LI, H. 1989. *Study on the*
569 *Permian-Triassic biostratigraphy and event stratigraphy of Northern Sichuan and southern Shaanxi*.
570 Beijing: Geological Publishing House.
- 571 LOEBLICH, A. R., JR. & TAPPAN, H. 1988. *Foraminiferal Genera and Their Classification*. New York: Van
572 Nostrand Reinhold Co.

- 573 MÁRQUEZ, L. 2005. Foraminiferal fauna recovered after the Late Permian extinctions in Iberia and the
574 westernmost Tethys area. *Palaeogeography, Palaeoclimatology, Palaeoecology* 229, 137-157.
- 575 MAURER, F., RETTORI, R. & MARTINI, R. 2008. Triassic stratigraphy, facies and evolution of the Arabian
576 Shelf in the northern United Arab Emirates. *International Journal of Earth Sciences* 97(4), 765–84.
- 577 NESTELL, G. P., KOLAR-JURKOVSEK, T., JURKOVSEK, B. & ALJINOVIC, D. 2011. Foraminifera from the
578 Permian-Triassic transition in western Slovenia. *Micropaleontology* 57(3), 197–222.
- 579 OBERHAUSER, R. 1957. Ein Vorkommen von Trocholina and Paratrocholina in der Ostalpinen Trias.
580 *Jahrbuch der Geologischen Bundesanstalt* 100, 257–67.
- 581 OKUYUCU, C., IVANOVA, D., BEDI, Y. & ERGEN, A. 2014. Discovery of an earliest Triassic, post-extinction
582 foraminiferal assemblage above the Permian-Triassic boundary, Strandzha nappes, north-west Turkey.
583 *Geological Quarterly* 58(1), doi: 10.7306/gq. 1145.
- 584 PAYNE, J.L., SUMMERS, M., REGO, B.L., ALTINER, D., WEI, J., Yu, M., & LEHRMANN, D.J. 2011.
585 Early and Middle Triassic trends in diversity, evenness, and size of foraminifers on a carbonate platform
586 in south China: implications for tempo and mode of biotic recovery from the end-Permian mass
587 extinction. *Paleobiology* 37, 409-425.
- 588 PÉREZ-LÓPEZ, A., MÁRQUEZ, L. & PÉREZ-VALERA, F. 2005. A foraminiferal assemblage as a bioevent marker
589 of the main Ladinian transgressive stage in the Betic Cordillera, southern Spain. *Palaeogeography,*
590 *Palaeoclimatology, Palaeoecology* 224(1), 217–31.
- 591 POPESCU, D. A. & POPESCU, L. G. 2005. The Olenekian carbonates of the Bucovinian nappe (the central
592 sector of the Haghimas syncline, Eastern Carpathians): lithology and microfacies. *Studia Universitatis*
593 *Babeş-Bolyai, Geologia* 50(1–2), 53–62.
- 594 PRONINA-NESTELL, G. P. & NESTELL, M. K. 2001. Late Changhsingian Foraminifers of the Northwestern

- 595 Caucasus. *Micropaleontology* 47(3), 205–34.
- 596 PRUSS, S. B. & BOTTJER, D. J. 2004. Early Triassic trace fossils of the western United States and their
597 implications for prolonged environmental stress from the end-Permian mass extinction. *Palaios* 19(6),
598 551–64.
- 599 REGO, B.L., WANG, S.C., ALTINER, D., & PAYNE, J.L. 2012. Within- and among-genus components of
600 size evolution during mass extinction, recovery, and background intervals: a case study of Late Permian
601 through Late Triassic foraminifera. *Paleobiology* 38, 627–643.
- 602 RETTORI, R. 1994. Replacement name *Hoyenella*, gen. n.(Triassic Foraminiferida, Miliolina) for *Glomospira*
603 *sinensis* Ho, 1959. *Bolletino Società Paleontologica Italiana* 33(3), 341–43.
- 604 RETTORI, R., ANGIOLINI, L. & MUTTONI, G. 1994. Lower and Middle Triassic foraminifera from the Eros
605 limestone, Hydra Island, Greece. *Journal of Micropalaeontology* 13, 25–46.
- 606 RODLAND, D. L. & BOTTJER, D. J. 2001. Biotic Recovery from the end-Permian Mass Extinction: Behavior of
607 the Inarticulate Brachiopod *Lingula* as a Disaster Taxon. *PALAIOS* 16, 95–101.
- 608 SCHUBERT, J. K. & BOTTJER, D. J. 1992. Early Triassic stromatolites as post-mass extinction disaster forms.
609 *Geology* 20(10), 883–86.
- 610 SCHUBERT, J. K. & BOTTJER, D. J. 1995. Aftermath of the Permian-Triassic mass extinction event:
611 paleoecology of Lower Triassic carbonates in the western USA. *Palaeogeography, Palaeoclimatology,*
612 *Palaeoecology* 116(1–2), 1–39.
- 613 SHEN, S. & HE, X. 1991. Changhsingian brachiopod assemblage sequence in Zhongliang Hill, Chongqing.
614 *Journal of Stratigraphy* 15(3), 189–96.
- 615 SONG, H., TONG, J., ALGEO, T. J., HORACEK, M., QIU, H., SONG, H., TIAN, L. & CHEN, Z.-Q. 2013a. Large
616 vertical $\delta^{13}\text{C}_{\text{DIC}}$ gradients in Early Triassic seas of the South China craton: Implications for

617 oceanographic changes related to Siberian Traps volcanism. *Global and Planetary Change* 105(0), 7–20.

618 SONG, H., TONG, J. & CHEN, Z. Q. 2009. Two episodes of foraminiferal extinction near the Permian-Triassic
619 boundary at the Meishan section, South China. *Australian Journal of Earth Sciences* 56, 765–73.

620 SONG, H., TONG, J. & CHEN, Z. Q. 2011. Evolutionary dynamics of the Permian-Triassic foraminifer size:
621 Evidence for Lilliput effect in the end-Permian mass extinction and its aftermath. *Palaeogeography,*
622 *Palaeoclimatology, Palaeoecology* 308(1–2), 98–110.

623 SONG, H., TONG, J., CHEN, Z. Q., YANG, H. & WANG, Y. 2009. End-Permian mass extinction of foraminifers
624 in the Nanpanjiang Basin, South China. *Journal of Paleontology* 83(5), 718–38.

625 SONG, H., TONG, J., XIONG, Y., SUN, D., TIAN, L. & SONG, H. 2012a. The large increase of $\delta^{13}\text{C}_{\text{carb}}$ -depth
626 gradient and the end-Permian mass extinction. *Science China Earth Sciences* 55, 1101–09.

627 SONG, H., TONG, J., ZHANG, K., WANG, Q. & CHEN, Z. Q. 2007. Foraminifers surviving from the end-Permian
628 mass extinction at Meishan, Changxing, China. *Palaeoworld* 22(1–3), 105–19.

629 SONG, H., WIGNALL, P. B., CHEN, Z. Q., TONG, J., BOND, D. P. G., LAI, X., ZHAO, X., JIANG, H., YAN, C., NIU,
630 Z., CHEN, J., YANG, H. & WANG, Y. 2011. Recovery tempo and pattern of marine ecosystems after the
631 end-Permian mass extinction. *Geology* 39(8), 739–42.

632 SONG, H., WIGNALL, P. B., CHU, D., TONG, J., SUN, Y., SONG, H., HE, W. & TIAN, L. 2014. Anoxia/high
633 temperature double whammy during the Permian-Triassic marine crisis and its aftermath. *Scientific*
634 *reports* 4(4132), DOI:10.1038/srep04132.

635 SONG, H., WIGNALL, P. B., TONG, J., BOND, D. P. G., SONG, H., LAI, X., ZHANG, K., WANG, H. & CHEN, Y.
636 2012b. Geochemical evidence from bio-apatite for multiple oceanic anoxic events during
637 Permian–Triassic transition and the link with end-Permian extinction and recovery. *Earth and Planetary*
638 *Science Letters* 353–354, 12–21.

- 639 SONG, H., WIGNALL, P. B., TONG, J. & YIN, H. 2013b. Two pulses of extinction during the Permian-Triassic
640 crisis. *Nature Geoscience* 6(1), 52–56.
- 641 SONG, H., YANG, L., TONG, J., CHEN, J., TIAN, L., SONG, H. & CHU, D. 2015. Recovery dynamics of
642 foraminifers and algae following the Permian-Triassic extinction in Qingyan, South China. *Geobios* 48,
643 71-83.
- 644 STANLEY, S. M. 2007. An analysis of the history of marine animal diversity. *Paleobiology* 33(4), 1–55.
- 645 SUN, D., TONG, J., XIONG, Y., TIAN, L. & YIN, H. 2012a. Conodont biostratigraphy and evolution across
646 Permian-Triassic boundary at Yangou Section, Leping, Jiangxi Province, South China. *Journal of Earth*
647 *Science* 23(3), 311–25.
- 648 SUN, Y., JOACHIMSKI, M. M., WIGNALL, P. B., YAN, C., CHEN, Y., JIANG, H., WANG, L. & LAI, X. 2012b.
649 Lethally Hot Temperatures During the Early Triassic Greenhouse. *Science* 338(6105), 366–70.
- 650 TIAN, L., TONG, J., ALGEO, T. J., SONG, H., SONG, H., CHU, D., SHI, L. & BOTJER, D. J. 2014a. Reconstruction
651 of Early Triassic ocean redox conditions based on framboidal pyrite from the Nanpanjiang Basin, South
652 China. *Palaeogeography, Palaeoclimatology, Palaeoecology* 412, 68–79.
- 653 TIAN, L., TONG, J., SUN, D., XIONG, Y., WANG, C., SONG, H., SONG, H. & HUANG, Y. 2014b. The microfacies
654 and sedimentary responses to the mass extinction during the Permian-Triassic transition at Yangou
655 Section, Jiangxi Province, South China. *Science China Earth Sciences* 57(9), 1–13.
- 656 TONG, J. & KUANG, W. 1990. A study of the Changxingian foraminifera and microfacies in Liangfengya,
657 Chongqing, Sichuan Province. *Earth Science—Journal of China University of Geosciences* 15(3),
658 337–44 (in Chinese with English abstract).
- 659 ÜNAL, E., ALTINER, D., YILMAZ, I.O., & OZKAN-ALTINER, S. 2003. Cyclic sedimentation across the
660 Permian-Triassic boundary (Central Taurides, Turkey). *Rivista Italiana di Paleontologia e Stratigrafia*

- 661 109, 359-376.
- 662 VELIĆ, I. 2007. Stratigraphy and Palaeobiogeography of Mesozoic Benthic Foraminifera of the Karst
663 Dinarides. *Geologia Croatica* 60(1), 1–86.
- 664 VUKS, V. J. 2007. Olenekian (Early Triassic) foraminifers of the Gorny Mangyshlak, Eastern Precaucasus and
665 Western Caucasus. *Palaeogeography, Palaeoclimatology, Palaeoecology* 252(1–2), 82–92.
- 666 WANG, Q., JINNAN, T., SONG, H. & YANG, H. 2009. Ecological evolution across the Permian/Triassic
667 boundary at the Kangjiaping Section in Cili County, Hunan Province, China *Science in China Series D:
668 Earth Sciences* 52(6), 797–806.
- 669 WANG, Y., TONG, J., WANG, J. & ZHOU, X. 2005. Calcimicrobialite after end-Permian mass extinction in
670 South China and its palaeoenvironmental significance. *Chinese Science Bulletin* 50(7), 665–71.
- 671 WIGNALL, P. B. & HALLAM, A. 1993. Griesbachian (Earliest Triassic) palaeoenvironmental changes in the
672 Salt Range, Pakistan and southeast China and their bearing on the Permo-Triassic mass extinction.
673 *Palaeogeography, Palaeoclimatology, Palaeoecology* 102, 215–37.
- 674 WIGNALL, P. B. & HALLAM, A. 1996. Facies change and the end-Permian mass extinction in SE Sichuan,
675 China. *Palaios* 11(6), 587–96.
- 676 WIGNALL, P. B., HALLAM, A., LAI, X. L. & YANG, F. 1995. Palaeoenvironmental changes across the
677 Permian/Triassic boundary at Shangsi (N. Sichuan, China). *Historical Biology* 10(2), 175–89.
- 678 WIGNALL, P. B. & TWITCHETT, R. J. 1999. Unusual intraclastic limestones in Lower Triassic carbonates and
679 their bearing on the aftermath of the end-Permian mass extinction. *Sedimentology* 46(2), 303–16.
- 680 WIGNALL, P. B. & TWITCHETT, R. J. 2002. Extent, duration, and nature of the Permian-Triassic superanoxic
681 event. In *Catastrophic events and mass extinctions; impacts and beyond: Geological Society of America
682 Special Publication 356* eds C. Koeberl and K. G. MacLeod). pp. 395–413.

- 683 YANG, H., CHEN, Z. Q., WANG, Y., TONG, J., SONG, H. & CHEN, J. 2011. Composition and structure of
684 microbialite ecosystems following the end-Permian mass extinction in South China. *Palaeogeography,*
685 *Palaeoclimatology, Palaeoecology* 308(1–2), 111–28.
- 686 YANG, L., SONG, H., TONG, J., CHU, D. & TIAN, L. 2013. Latest Permian extinction of fusulinids at the
687 Kangjiaping section, Cili, Hunan. *Acta Micropalaeontologica Sinica* 30(4), 353–46.
- 688 YANG, S. & SUN, C. 1990. Discovery of Permian-Triassic conodont fauna in Tieshikou Area, Xinfeng, Jiangxi
689 and its geological significance. *Acta Scientiarum Naturalum Universitatis Pekinesis* 26(2), 243–56.
- 690 YANG, Z., YIN, H., WU, S., YANG, F., DING, M. & XU, G. 1987. *Permian-Triassic boundary stratigraphy and*
691 *fauna of South China*. Beijing: Geological Publishing House.
- 692 YIN, H. 1985. Bivalves near the Permian-Triassic boundary in South China. *Journal of Paleontology* 59(3),
693 572–600.
- 694 YIN, H., ZHANG, K., TONG, J., YANG, Z. & WU, S. 2001. The Global Stratotype Section and Point (GSSP) of
695 the Permian-Triassic boundary. *Episodes* 24(2), 102–14.
- 696 ZANINETTI, L. & BRÖNNIMANN, P. 1975. Triassic foraminifera from Pakistan. *Riv. Ital. Paleontol* 81(3),
697 257–80.
- 698 ZANINETTI, L., RETTORI, R. & MARTINI, R. 1994. Paulbronnimanninae Rettori and Zaninetti, 1993
699 (Foraminiferida, Ammodiscidae) and other Anisian foraminifers from the Piz da Peres section
700 (Valdaora-Olang, Pusteria Valley, Dolomites, NE Italy). *Rivista Italiana di Paleontologia e Stratigrafia*
701 100, 339–50.
- 702 ZHU, X. 1999. On a taking shape era of the reef s in Dongling Area of northwest Jiangxi. *Journal of Jiangxi*
703 *Normal University* 23(3), 252–58.
- 704 ZHU, X., WANG, C., LU, H., MU, X., ZHANG, L., QIN, Z., LUO, H., YANG, W. & DENG, Z. 1994.

705 Permian-Triassic boundary in Jiangxi, China. *Acta Micropalaeontologica Sinica* 11(4), 439–52.
706 ZONNEVELD, J.-P., MACNAUGHTON, R. B., UTTING, J., BEATTY, T. W., PEMBERTON, S. G. & HENDERSON, C.
707 M. 2010. Sedimentology and ichnology of the Lower Triassic Montney Formation in the
708 Pedigree-Ring/Border-Kahntah River area, northwestern Alberta and northeastern British Columbia.
709 *Bulletin of Canadian Petroleum Geology* 58(2), 115–40.

710

711 **Figure Captions**

712 **Figure 1.** Early Triassic paleogeographic map of South China modified from (Feng, Bao &
713 Liu, 1997; Lehrmann, Wei & Enos, 1998). Black triangles show Early Triassic sections
714 containing disaster and opportunistic foraminifers whereas grey triangles show PTB
715 sections containing disaster foraminifers.

716

717 **Figure 2.** The stratigraphical distributions of disaster foraminifers in four PTB sections:
718 Meishan, Liangfengya, Cili, and Dajiang.

719

720 **Figure 3.** The stratigraphical distributions of disaster and opportunistic foraminifers in
721 three Early Triassic sections: Wufeng, Xiangkou, and Shangsi.

722

723

724 **Figure 4.** Disaster foraminifer *Postcladella kahlori* Brönnimann, Zanninetti, & Bozorgnia,
725 1972 from the Permian-Triassic boundary strata of South China. (a), Lowest Daye
726 Formation of Dongling section, Jiangxi Province; (b), Lower Yelang Formation of

727 Xiangkou section, Guizhou Province; (c, d), Lowest Daye Formation of Cili section, Hunan
728 Province. Triangular arrows indicate blurry specimens whereas long arrow indicates broken
729 specimen.

730

731 **Figure 5.** Disaster and opportunistic foraminifer *Earlandia* sp. from the Permian-Triassic
732 boundary strata and Early Triassic of South China. (a), Lowest Tieshikou Formation of
733 Tieshikou section, Jiangxi Province; (b), Lowest Feixianguan Formation of Liangfengya
734 section, Chongqing; (c), Lowest Daye Formation of Dongling section, Jiangxi Province; (d),
735 Upper Maocaopu Formation of Xiangkou section, Guizhou Province.

736

737 **Figure 6.** Disaster foraminifer *Globivalvulina lukachiensis* Nestell *et al.*, 2011 from the
738 Permian-Triassic boundary strata of South China. (a-d), Lowest Daye Formation of Cili
739 section, Hunan Province.

740

741 **Figure 7.** Opportunistic foraminifer *Hemigordiellina regularia* (Lipina, 1949) from the
742 Early Triassic of South China. (a, b), Upper Maocaopu Formation of Xiangkou section,
743 Guizhou Province; (c, d), Lower Jialingjiang Formation of Wufeng section, Hubei
744 Province.

745

746 **Figure 8.** Opportunistic foraminifer *Hoyenella* spp. from the Early Triassic of South China.
747 (a, b), Upper Tongjiezi Formation of Shangsi section, Sichuan Province; (c), Upper
748 Maocaopu Formation of Xiangkou section, Guizhou Province; (d), Lower Jialingjiang

749 Formation of Wufeng section, Hubei Province.

750

751 **Figure 9.** Opportunistic foraminifer *Arenovidalina chialingchiangensis* Ho, 1959 from the
752 Early Triassic of South China. (a-c), Upper Daye Formation of Wufeng section, Hubei
753 Province; (d), Lowest Tongjiezi Formation of Shangsi section, Sichuan Province.

754

755 **Figure 10.** Opportunistic foraminifer *Aulotortus? bakonyensis* Blau, 1989 and *Triadodiscus*
756 *eomesozoicus* (Oberhauser, 1957) from the Early Triassic of South China. (a, b), *Aulotortus?*
757 *bakonyensis* Blau, 1989, Upper Tongjiezi Formation of Shangsi section, Sichuan Province;
758 (c, d), *Triadodiscus eomesozoicus* (Oberhauser, 1957) from Upper Tongjiezi Formation of
759 Shangsi section, Sichuan Province. Triangular arrows indicate blurry specimens.

760

761 **Figure 11.** Opportunistic foraminifer *Meandrospira pusilla* (Ho, 1959) from the Early
762 Triassic of South China. (a), Lower Jialingjiang Formation of Wufeng section, Hubei
763 Province; (b), Upper Tongjiezi Formation of Shangsi section, Sichuan Province; (c),
764 Lekang Formation of Lekang section, Guizhou Province; (d), Upper Maocaopu Formation
765 of Xiangkou section, Guizhou Province.

766

767 **Figure 12.** Stratigraphic ranges of disaster and opportunistic foraminifers in South China
768 during the end-Permian and Early Triassic.

769

770 **Figure 13.** Number of specimens versus number of genera in a 2.2×2.2 cm² thin-section for

771 disaster and opportunistic foraminifers from Early Triassic and normal taxa from Late

772 Permian and Middle Triassic.

773

774

775

776 **Table 1.** The distributions of Early Triassic disaster and opportunistic foraminifers in South

777 China.

Sections	Griesbachian <i>H. parvus</i> Zone	Griesbachian <i>I. sarcica</i> Zone	Dienerian	Smithian	Spathian
Meishan		<i>Earlandia</i>			
Huangzhishan	<i>Earlandia</i>				
Yangou	<i>Earlandia</i>				
Dongling	<i>Earlandia</i> , <i>Postcladella</i>				
Tieshikou	<i>Earlandia</i>				
Cili	<i>Earlandia</i> , <i>Globivalvulina</i> , <i>Postcladella</i>				
Liangfengya		<i>Earlandia</i>			
Dajiang	<i>Earlandia</i> , <i>Postcladella</i>				
Wufeng				<i>Arenovidalina</i>	<i>Hemigordiellina</i> , <i>Hoyenella</i> , <i>Meandrospira</i>
Shangsi				<i>Arenovidalina</i> , <i>Aulotortus?</i> , <i>Hemigordiellina</i> , <i>Hoyenella</i> , <i>Meandrospira</i> , <i>Triadodiscus</i>	
Xiangkou		<i>Postcladella</i>			<i>Earlandia</i> , <i>Hemigordiellina</i> , <i>Hoyenella</i> , <i>Meandrospira</i>
Lekang					<i>Meandrospira</i>