



UNIVERSITY OF LEEDS

This is a repository copy of *Modelling determinants of extinction across two Mesozoic hyperthermal events*.

White Rose Research Online URL for this paper:
<https://eprints.whiterose.ac.uk/137021/>

Version: Accepted Version

Article:

Dunhill, AM orcid.org/0000-0002-8680-9163, Foster, WJ, Azaele, S orcid.org/0000-0002-5153-4833 et al. (2 more authors) (2018) Modelling determinants of extinction across two Mesozoic hyperthermal events. *Proceedings of the Royal Society B: Biological Sciences*, 285 (1889). 20180404. ISSN 0962-8452

<https://doi.org/10.1098/rspb.2018.0404>

© 2018 The Author(s). Published by the Royal Society. All rights reserved. This is an author produced version of a paper published in *Proceedings of the Royal Society B: Biological Sciences*. Uploaded in accordance with the publisher's self-archiving policy.

Reuse

Items deposited in White Rose Research Online are protected by copyright, with all rights reserved unless indicated otherwise. They may be downloaded and/or printed for private study, or other acts as permitted by national copyright laws. The publisher or other rights holders may allow further reproduction and re-use of the full text version. This is indicated by the licence information on the White Rose Research Online record for the item.

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/>

1 Modelling determinants of extinction across two Mesozoic
2 hyperthermal events

3 Alexander M. Dunhill¹, William J. Foster², Sandro Azaele³, James Sciberras⁴ and
4 Richard J. Twitchett⁵

5 ¹School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK

6 ²Museum für Naturkunde, Leibniz-Institut für Evolutions- und Biodiversitätsforschung, Berlin,
7 Deutschland

8 ³School of Mathematics, University of Leeds, Leeds LS2 9JT, UK

9 ⁴OnCorps, Exeter House, Lower Station Approach, Temple Means, Bristol, BS1 6QS, UK

10 ⁵Department of Earth Sciences, Natural History Museum, London, SW7 5BD, UK

11

12 Author for correspondence: Alexander M. Dunhill

13 e-mail: a.dunhill@leeds.ac.uk

14

15 The Late Triassic and early Toarcian extinction events are both associated with
16 greenhouse warming events triggered by massive volcanism. These Mesozoic
17 hyperthermals were responsible for the mass extinction of marine organisms and resulted
18 in significant ecological upheaval. It has, however, been suggested that these events
19 merely involved intensification of background extinction rates rather than significant
20 shifts in the macroevolutionary regime and extinction selectivity. Here, we apply a
21 multivariate modelling approach to a vast global database of marine organisms to test
22 whether extinction selectivity varied through the Late Triassic and Early Jurassic. We
23 show that these hyperthermals do represent shifts in the macroevolutionary regime and
24 record different extinction selectivity compared to background intervals of the Late
25 Triassic and Early Jurassic. The Late Triassic mass extinction represents a more
26 profound change in selectivity than the early Toarcian extinction but both events show a
27 common pattern of selecting against pelagic predators and benthic photosymbiotic and
28 suspension-feeding organisms, suggesting that these groups of organisms may be
29 particularly vulnerable during episodes of global warming. In particular, the Late
30 Triassic extinction represents a macroevolutionary regime change that is characterised
31 by (i) the change in extinction selectivity between Triassic background intervals and the
32 extinction event itself; and (ii) the differences in extinction selectivity between the Late
33 Triassic and Early Jurassic as a whole.

34

35 **Key words:** mass extinction, palaeoecology, modelling, Mesozoic, hyperthermal

36

37 **1. Introduction**

38 The Late Triassic to Early Jurassic interval contains two major extinction events: the Late
39 Triassic mass extinction (LTE; c.201Ma) [1] and the early Toarcian extinction (EToE;
40 c.187Ma) [2]. The LTE is recognised as the second largest marine biodiversity loss [3] and
41 third biggest ecological crisis of the Phanerozoic [4], resulting in a global reef crisis [5], the
42 most severe extinction of scleractinian corals [6, 7], significant extinctions amongst
43 ammonoids [8], bivalves [9] and marine vertebrates [10], and the final demise of the conodonts
44 [11]. By comparison, the EToE was smaller in magnitude, but records a similar pattern of
45 selective losses, with an associated reef crisis [5], high levels of extinction amongst bivalves
46 [12] and ammonoids [13], and the collapse of both benthic and pelagic marine ecosystems [7,
47 8]. Both of these events are associated with, and likely caused by, elevated atmospheric CO₂
48 levels and global warming [14-21]; i.e. they are hyperthermals. In each case, eruptions of Large
49 Igneous Provinces (LIPs) probably caused the rise in CO₂ [18, 20-25], with proposed extinction
50 drivers including rapid warming [14-17, 20, 26], ocean anoxia [16, 17, 26], and ocean
51 acidification [27-29] as a direct result of the volcanic greenhouse gas emissions.

52 Several biological and ecological traits appear to have been selected against during the
53 LTE, such as possessing a heavily calcified skeleton [27], inhabiting reef and/or inshore
54 environments [30, 31], and residing at tropical latitudes [31, 32]. The greatest reduction in both
55 taxonomic and functional richness occurred amongst sessile suspension-feeding guilds,
56 particularly those dwelling in tropical reefs in the Panthalassa Ocean [31]. Despite this
57 evidence for apparent selectivity during the LTE hyperthermal, it has been claimed that there
58 was little change in “macroevolutionary regime” [30] compared to the rest of the Late Triassic
59 and Early Jurassic, and that the LTE simply reflects intensification of the high rates of
60 background extinction already experienced through the Late Triassic [30, 33, 34]. There has
61 been less research on selectivity during the EToE hyperthermal, although there is some

62 evidence for loss of reef taxa [5], selection against endemic taxa [12], the motile benthos [35],
63 infaunal organisms [12, 17, 35, 36], as well as higher levels of extinction in the restricted basins
64 of northwest Tethys, northeast Panthalassa [35], and the Boreal Ocean [2, 36] as well as in the
65 southern hemisphere [37]. Whether this represents a macroevolutionary regime shift compared
66 to Jurassic background extinction is unknown.

67 Here, we provide the first multivariate analysis of ecological selectivity during the Late
68 Triassic and Early Jurassic, in order to determine whether there are any substantial differences
69 between the LTE and EToE hyperthermal events and the periods of normal background
70 extinction, and hence whether a shift in macroevolutionary regime occurred. A
71 macroevolutionary regime shift is recognised when the suite of traits that promote extinction
72 or survivorship are different, and particularly when the direction of selectivity changes so that
73 traits that conferred survivorship during background times become an extinction risk [38]. We
74 apply a generalised linear modelling (GLM) methodology to the largest and most
75 comprehensive global database yet analysed in order to assess the relative importance of a
76 number of intrinsic and extrinsic ecological variables as determinants of extinction in marine
77 ecosystems. We aim to test the following hypotheses: (i) do certain ecological variables (e.g.
78 latitudinal distribution, habitat preference, feeding mode and calcification) correlate with
79 higher extinction risk during the LTE and EToE hyperthermal events; (ii) are similar trends
80 recorded in both past hyperthermals despite differences in starting conditions and magnitude;
81 and (iii) are similar trends recorded during background times, or do the LTE and EToE
82 hyperthermals represent significantly different extinction selectivity?

83

84 **2. Methods**

85 We utilised a database of fossil occurrences of Middle Triassic to Middle Jurassic (Ladinian-
86 Aalenian) marine animal genera collated from the Paleobiology Database (PaleoDB) [39, 40].

87 The total dataset comprises 55,428 occurrences of 2,621 genera, which is more than double the
88 number that was available for previous analyses, e.g. [30, 32]. Each genus was then classified
89 according to a number of extrinsic (i.e. abiotic) and intrinsic (i.e. biotic) ecological variables:
90 (Table 1; see [31] for detailed download, vetting, and classification information). Proportional
91 generic extinction rates were calculated and plotted at the stage level for guilds of fossil
92 organisms defined by each ecological variable (Table 1). In order to account for biases brought
93 about by uneven sampling across space and through time, we applied a subsampling protocol
94 to standardize proportional extinction on the basis of the number of fossil occurrences. All
95 variables were subsampled to $n = 250$ per stage, for 1000 iterations apart from feeding, which
96 was subsampled to $n = 75$, due to the increased number of variable arguments and thus reduced
97 sample sizes after splitting occurrences via feeding mode. Lightly calcified taxa, polar latitude,
98 boreal ocean and reef taxa all fall short of the subsampling requirement for at least one of the
99 time bins and are, therefore, not plotted in the univariate time series. However, when
100 amalgamated with the other variables for the multivariate analyses, they provide sample sizes
101 that are sufficient for the GLM analyses.

102 Multiple ecological variables are not independent of one another in terms of
103 proportional extinction through time, therefore it is essential to test their effects on extinction
104 within a multivariate framework. For example, pelagic taxa within the database are
105 predominantly predatory and fast-moving as the majority of pelagic taxa are vertebrates or
106 cephalopods. Therefore, it is impossible to determine which, if any, of these three variables is
107 influencing extinction rates in a univariate analysis. We applied GLMs with a binomial
108 distribution and a logit link function (i.e. multiple logistic regression models) to test the effects
109 of multiple ecological variables on proportional generic extinction through the study interval
110 [41]. The major extinction episodes of the LTE (Rhaetian/Hettangian) and EToE
111 (Pliensbachian/Toarcian) were analysed separately and compared to the other stage boundaries

112 which, together, are treated as representing the background intervals of the Triassic and
113 Jurassic. However, because the binomial models were strongly underdispersed, we then used
114 quasibinomial models and estimated the dispersion parameter from the data [42].
115 Underdispersal, where the variance is less than the nominal mean [43], can lead to over-
116 conservatism and thus can result in type II errors. We applied the GLMs to two datasets: (i)
117 including all the ecological variables, and (ii) a separate dataset compiled without the
118 depositional setting variable, because reliable depositional setting data only exist for around
119 50% of the fossil occurrences in the entire data set. A number of model runs were carried out
120 with different variable combinations for each of the four broad time intervals: Triassic
121 background (Ladinian-Carnian, Carnian-Norian, Norian-Rhaetian); LTE (Rhaetian-
122 Hettangian); Jurassic background (Hettangian-Sinemurian, Sinemurian-Pliensbachian,
123 Toarcian-Aalenian); and EToE (Pliensbachian-Toarcian). Model selection was carried out by
124 using the drop1() command in R, which drops one explanatory variable in turn and each time
125 applies an analysis of deviance test (F-test) [41]. The data for the GLMs were not subsampled,
126 but any ecological guilds with consistently low sample sizes (<10 occurrences per bin) were
127 omitted from the analyses. All analyses were carried in R v.3.4.3 [44].

128

129 **3. Results**

130 (a) Univariate time series analysis

131 The data show clear differences in extinction magnitude and selectivity between the
132 hyperthermals and background intervals (Figure 1). For many, but not all variables (e.g.
133 infaunal taxa at EToE, moderate calcifiers at LTE, mid-latitude taxa at LTE), extinction
134 magnitude is greater during the hyperthermals than during their respective background times,
135 as expected, and the pattern of relative selectivity remains the same (e.g. Figure 1A, B). During
136 the LTE, however, feeding, calcification, latitude and ocean basin all record different patterns

137 of selectivity compared to the Triassic background (Figures 1C-F). Photosymbiotic taxa
138 suffered a substantially greater extinction during the LTE than during Triassic background
139 periods, overtaking suspension- and deposit-feeders in relative extinction risk (Figure 1C).
140 Likewise, heavily calcified taxa and those that live in the tropics and Panthalassa record the
141 greatest extinction magnitudes during the LTE hyperthermal, which represents a marked shift
142 in selectivity compared to the Triassic background (Fig. 1D-F). In contrast, the only similar
143 shift in selectivity during the EToE, compared to the Jurassic background, occurs with latitude,
144 with mid-latitude taxa showing greater extinction risk than tropical taxa during the
145 hyperthermal (Figure 1E). With a few exceptions, remaining variables all increase during the
146 EToE event, suggesting the event mainly represents an intensification of Jurassic background
147 extinction rates.

148 Unexpectedly, there also appear to be clear differences in background extinction
149 magnitude and selectivity between the Triassic and Jurassic. Background extinction appears
150 much higher during the Triassic than the Jurassic, with extinction magnitude in some guilds
151 being higher during the Triassic background intervals than during the EToE (Figure 1).

152

153 (b) Generalised linear modelling

154 Multivariate analyses demonstrate clear differences between background and mass extinction
155 intervals (see Table 2 for GLM results). In general, the suite of ecological variables (see Table
156 1) analysed in this study explain far less of the recorded extinction during background times
157 compared to the two hyperthermal events. During the Triassic background interval,
158 depositional setting is the only significant predictor of extinction, and only in the single model
159 that considers just the set of extrinsic factors. This is due to reef taxa having significantly lower
160 extinction risk than taxa that live in other settings. During the Jurassic background interval,
161 when all ecological variables are considered, motility, palaeo-ocean basin, and depositional

162 setting are all significant predictors of extinction, with pelagic taxa and taxa residing in the
163 Tethys Ocean having higher extinction and reef dwellers having lower extinction. The best
164 fitting model identifies only palaeo-ocean (i.e. Tethys) and depositional setting (reefs) as being
165 significant predictors of extinction. The only other model run that identifies a significant
166 predictor of extinction is the one that considers just the extrinsic ecological variables. In that
167 case, palaeo-oceanic basin is again identified as having a significant bearing on extinction, due
168 to the higher extinction in Tethys. In contrast to the background times, during the LTE and
169 EToE events, many more model runs identify significant ecological predictors of extinction.
170 Furthermore, those variables that are identified as being significant are different to the ones
171 identified during the background times.

172 During the LTE, feeding or latitude are the only variables identified as being significant
173 predictors of extinction. Latitude alone significantly predicts extinction when (a) all ecological
174 variables are considered and (b) when only extrinsic factors are considered, using the smaller
175 dataset that includes depositional setting. In each case, taxa residing at tropical latitudes show
176 significantly higher extinction than those inhabiting higher latitudes. In contrast, feeding is
177 identified as a significant predictor of extinction in two other model runs, but only using the
178 expanded dataset that excludes depositional setting. In both cases, where (a) all variables or (b)
179 just the intrinsic ones are considered, excluding depositional setting, the best-fitting models
180 identify predatory, photosymbiotic and suspension-feeding habits as being significant positive
181 predictors of extinction.

182 For the EToE event, significant predictors of extinction are only identified in model
183 runs that utilise the expanded dataset that excludes depositional setting. When all variables,
184 apart from depositional setting, are considered, five factors (motility, feeding, latitude, palaeo-
185 ocean basin and calcification) all appear to significantly predict extinction. Model selection
186 reveals that the best-fitting model identifies just feeding and palaeo-ocean basin, with

187 photosymbiotic taxa and taxa residing in the Boreal Ocean predicting significantly higher
188 extinction than other categories within those variables. When considering just the extrinsic
189 ecological variables, no variable predicts extinction. However, when we use the expanded data
190 set with no depositional environment data, palaeo-ocean basin significantly predicts extinction
191 with Boreal taxa having higher extinction and, after model selection, Panthalassa taxa show
192 lower extinction than both Boreal and Tethys taxa. When considering only the intrinsic
193 ecological variables, no variable predicts extinction until we use the expanded data set with no
194 depositional environment variable, after which, feeding significantly predicts extinction, with
195 photosymbiotic taxa showing higher extinction than other feeding guilds.

196

197 **4. Discussion**

198 There are marked changes in extinction selectivity between periods of normal background and
199 the two hyperthermals (Figure 1; Table 2). Extinction magnitude is higher in most ecological
200 guilds during the LTE and, although the EToE generally displays higher levels of extinction
201 than Jurassic background intervals, for some guilds Triassic background extinction is higher
202 than it is during the EToE (Figure 1). Not only do the LTE and EToE events represent an
203 increase in extinction intensity above respective Triassic and Jurassic background rates but,
204 more importantly, and contrary to previous claims [30], there are stark differences in extinction
205 selectivity between times of both background and mass extinction, and between Triassic and
206 Jurassic intervals in general. It is also evident that background extinction was higher in the Late
207 Triassic, prior to the LTE hyperthermal, than it was during the Early Jurassic, in the aftermath
208 of the mass extinction [30, 33] (Figure 1).

209 A tropical extinction peak characterises the LTE, whereas mid-latitude taxa display
210 higher extinction during the EToE and during background times. Although the LTE data are
211 consistent with expectations that an episode of global warming should result in extinction being

212 concentrated in the tropics [32, 45], the EToE data suggest that high tropical extinction may
213 not be ubiquitous to all hyperthermal events. Additionally, higher tropical extinction across the
214 LTE is only significant in the models where the data are also partitioned by depositional setting,
215 suggesting that some of the tropical extinction signal is rooted in a particular depositional
216 setting, most probably tropical reefs. Although extinction is higher in the mid-latitudes through
217 the EToE [31], latitude does not significantly predict extinction, suggesting that the higher rates
218 of extinction in the mid-latitudes during the EToE may be governed by other factors such as
219 ocean basin or that the warming was not as intense as during the LTE [2].

220 The peak in extinction in Panthalassa during the LTE is not replicated in the Tethys
221 Ocean, despite the Tethys showing higher background extinction rates. However, modelling
222 does not show palaeo-ocean as a significant predictor of extinction across the LTE, suggesting
223 that this peak is a result of other factors, such as the high proportion of tropical data in Rhaetian-
224 Hettangian Panthalassa. In contrast, Panthalassa displays significantly lower extinction than
225 the Tethys and Boreal oceans during the EToE. The EToE appears to be characterised by raised
226 extinction rates in the Tethyan and Boreal Oceans. This might be expected given the higher
227 prevalence of restricted basins, particularly in north-western Tethys, when considering the
228 repeated dysoxic conditions in the Early Jurassic, of which the EToE is the most severe [2, 26].
229 However, this pattern persists in light of evidence for prolonged anoxia and extinction in some
230 Panthalassa basins [35]. It is also likely that the mid to high palaeolatitude of the Boreal and
231 north-western Tethys basins of Europe are driving the mid-latitudinal peak in extinction
232 intensity through the EToE.

233 Although difficult to show because of very small sample sizes, the reef crises at the
234 LTE and EToE are evident in the data by the crashes in reef taxa abundances [31] and diversity
235 [7, 31]. The reef crises are also highlighted by the high levels of extinction witnessed amongst
236 photosymbiotic taxa and suspension feeders across the LTE, and photosymbiotic taxa across

237 the EToE. In contrast to the extinction events, background extinction for reef taxa and
238 photosymbiotic feeders was lower than those taxa residing in other depositional settings and
239 feeding via different strategies. This highlights a major change in extinction selectivity during
240 both the LTE and EToE and permits the rejection of the idea that the LTE is merely an
241 intensification of background extinction seen during the Late Triassic [30].

242 Tiering does not appear to have an influence on extinction selectivity across the LTE,
243 despite there being an increase in extinction magnitude across all guilds. However, the impact
244 of the mass extinction on level-bottom communities was particularly short-lived with full
245 recovery occurring by the upper Hettangian [46-48]. Therefore, the temporary disappearance
246 of the deep infaunal and erect benthic tiers in the earliest Hettangian recorded by previous
247 studies [46, 48], would not be detected here because of the coarser nature of the stage-level
248 time bins. There is some evidence of increased extinction risk to pelagic taxa during intervals
249 of background and mass extinction, possibly related to high turnover of ammonoids and
250 vertebrates, which also drives the consistently high levels of extinction in predatory taxa [13,
251 49]. We see a similar pattern in terms of motility, with no apparent selectivity across the LTE
252 or during periods of background extinction. There is some weak evidence for selectivity against
253 non-motile taxa across the EToE, although this is not significant in the best-fitting model. The
254 lack of any selectivity against non-motile and epifaunal taxa across the LTE suggests that the
255 mass extinction did not result in an indirect intensification of the Mesozoic Marine Revolution
256 (MMR) as previously suggested [31] and these previously detected high levels of extinction
257 amongst non-motile epifauna are a result of elevated extinction amongst
258 photosymbiotic/suspension feeding guilds in reef environments, which are predominantly non-
259 motile and epifaunal. Our analyses do detect higher levels of background extinction amongst
260 non-motile and epifaunal taxa during the Late Triassic compared to the Early Jurassic.
261 Crucially, however, motility and tiering do not predict extinction in the Late Triassic

262 background interval. Therefore, we cannot find solid evidence of selectivity against non-motile
263 epifauna during a time period (Carnian-Norian) that has been identified as key to the MMR
264 [50]. The cause of higher Triassic background extinction versus Jurassic background extinction
265 is likely a result of the high faunal turnover associated with the Carnian Pluvial Event [51],
266 rather than the MMR.

267 There is a peak in extinction amongst heavily calcified taxa during the LTE whereas
268 during background periods and the EToE heavy calcifiers display lower extinction than
269 moderate calcifiers. Although this may support the hypotheses that hypercapnia [52] and/or
270 ocean acidification may have played a role in extinction during the LTE [5, 27], our modelling
271 results show no evidence that calcification was a significant predictor of extinction during
272 either the LTE or the EToE hyperthermal. The multivariate analyses show no evidence of
273 selectivity against heavy calcifiers during the LTE and only very weak evidence at the EToE,
274 although this result is non-significant in the best-fitting model. Our analyses support previous
275 studies that found no strong link between calcification grade and extinction selectivity [30].
276 Therefore, it seems unlikely that hypercapnia or ocean acidification were the main or sole
277 drivers of extinction during the LTE and EToE hyperthermals.

278 The LTE and, albeit to a lesser extent, the EToE are both characterised by ~~marked~~ shifts
279 in extinction selectivity away from the macroevolutionary regimes of the Late Triassic and
280 Early Jurassic background intervals. Background extinction rates in the Late Triassic were
281 higher than those of the Early Jurassic [30, 33], but the LTE was not merely an intensification
282 of those background rates as has been previously suggested [30]. Extinction selectivity changed
283 dramatically across the LTE with the initiation of strong selection against tropical taxa with
284 photosymbiotic, suspension or predatory feeding strategies. This pattern is consistent with a
285 warming-driven tropical reef crisis. We find little evidence to support previous ideas that
286 palaeo-ocean basin [31] or calcification [27] were important determinants of extinction at the

287 LTE. Despite differences in starting conditions, species involved, and magnitudes of global
288 warming and environmental change, the LTE and EToE show some common patterns of
289 selectivity. Both events record strong extinction selectivity against pelagic predatory guilds
290 and against benthic photosymbiotic and suspension feeding organisms, suggesting that these
291 groups of marine organisms may be particularly vulnerable during episodes of global warming.
292 The effects of the LTE were most severe in the tropics whilst the EToE was felt more severely
293 at higher latitudes, which may reflect differences in magnitude of environmental change or
294 starting conditions, such as palaeogeography. However, the EToE shares some common
295 selectivity patterns with periods of Jurassic background extinction, i.e. high extinction in the
296 Tethys Ocean, suggesting that the EToE may have represented an intensification of Jurassic
297 background extinction, albeit with a switch to selecting against reef inhabiting photosymbiotic
298 taxa. The LTE shows a clear change in extinction selectivity and thus macroevolutionary
299 regime which is characterised not only by a shift in extinction selectivity from Triassic
300 background intervals across the LTE but also by the difference in extinction selectivity between
301 the Late Triassic and Early Jurassic as a whole.

302

303 **Data accessibility.** The data set used in the statistical analyses and the statistical results are
304 available in the Dryad Digital Repository: doi:10.5061/dryad.6t461hq).

305 **Author's contributions.** A.M.D., W.J.F. and R.J.T. conceived the study. A.M.D. and W.J.F.
306 collected the data. S.A and J.S. wrote analytical code and advised on analytical methods.

307 A.M.D. analysed the data. A.M.D. led the writing of the manuscript and W.J.F., S.A. and

308 R.J.T. contributed to the writing and editing of the manuscript and preparation of figures.

309 **Competing interests.** We declare we have no competing interests.

310 **Funding.** Funding for this work was provided by a Leverhulme Early Career Fellowship
311 (ECF-2015-044) and Natural Environmental Research Council research grant
312 (NE/P013724/1) to AMD.

313 **Acknowledgements.** The authors thank Xiaoya Ma, Matthew Clapham and four anonymous
314 reviewers for helpful editorial and analytical reviews that have greatly improved this
315 manuscript. The authors also thank the numerous authors of the original studies that provide
316 the source data on which this study is based, and the many data enterers of the Paleobiology
317 Database for the provision of fossil occurrence data, particularly: Matthew Clapham,
318 Wolfgang Kiessling, Franz Fürsich, Martin Aberhan, Andy Rees, József Pálffy, Matthew
319 Carrano, David Bottjer, Alistair McGowan, Arnold Miller, Luc Villier, Roger Benson, John
320 Alroy, and Richard Butler. This is Paleobiology Database publication ###.

321
322 **References**

- 323 [1] Lindström, S., van de Schootbrugge, B., Hansen, K.H., Pedersen, G.K., Alsen, P.,
324 Thibault, N., Dybkjær, K., Bjerrum, C.J. & Nielsen, L.H. 2017 A new correlation of
325 Triassic–Jurassic boundary successions in NW Europe, Nevada and Peru, and the
326 Central Atlantic Magmatic Province: A time-line for the end-Triassic mass extinction.
327 *Palaeogeography, Palaeoclimatology, Palaeoecology* **478**, 80-102.
328 (doi:<https://doi.org/10.1016/j.palaeo.2016.12.025>).
- 329 [2] Little, C.T. & Benton, M.J. 1995 Early Jurassic mass extinction: A global long-term
330 event. *Geology* **23**, 495-498.
- 331 [3] Alroy, J. 2010 The shifting balance of diversity among major marine animal groups.
332 *Science* **329**, 1191-1194.

- 333 [4] McGhee, G.R., Sheehan, P.M., Bottjer, D.J. & Droser, M.L. 2004 Ecological ranking of
334 Phanerozoic biodiversity crises: ecological and taxonomic severities are decoupled.
335 *Palaeogeography, Palaeoclimatology, Palaeoecology* **211**, 289-297.
- 336 [5] Kiessling, W. & Simpson, C. 2011 On the potential for ocean acidification to be a general
337 cause of ancient reef crises. *Global Change Biology* **17**, 56-67. (doi:10.1111/j.1365-
338 2486.2010.02204.x).
- 339 [6] Flügel, E. 2002 Triassic reef patterns. *SEPM Special Publication* **72**, 391-463.
- 340 [7] Martindale, R.C., Berelson, W.M., Corsetti, F.A., Bottjer, D.J. & West, A.J. 2012
341 Constraining carbonate chemistry at a potential ocean acidification event (the
342 Triassic-Jurassic boundary) using the presence of corals and coral reefs in the fossil
343 record. *Palaeogeography Palaeoclimatology Palaeoecology* **350**, 114-123.
344 (doi:10.1016/j.palaeo.2012.06.020).
- 345 [8] House, M.R. 1989 Ammonoid extinction events. *Philosophical Transactions of the Royal*
346 *Society of London. B, Biological Sciences* **325**, 307-326.
347 (doi:10.1098/rstb.1989.0091).
- 348 [9] Ros, S. & Echevarría, J. 2012 Ecological signature of the end-Triassic biotic crisis: what
349 do bivalves have to say? *Historical Biology* **24**, 489-503.
350 (doi:10.1080/08912963.2011.625568).
- 351 [10] Benson, R.B.J., Butler, R.J., Lindgren, J. & Smith, A.S. 2010 Mesozoic marine tetrapod
352 diversity: mass extinctions and temporal heterogeneity in geological megabiases
353 affecting vertebrates. *Proceedings of the Royal Society B* **277**, 829-834.
- 354 [11] Bond, D.P.G. & Grasby, S.E. 2017 On the causes of mass extinctions. *Palaeogeography,*
355 *Palaeoclimatology, Palaeoecology* **478**, 3-29.
356 (doi:https://doi.org/10.1016/j.palaeo.2016.11.005).

- 357 [12] Aberhan, M. & Baumiller, T.K. 2003 Selective extinction among Early Jurassic
358 bivalves: A consequence of anoxia. *Geology* **31**, 1077-1080. (doi:10.1130/G19938.1).
- 359 [13] Dera, G., Neige, P., Dommergues, J.-L., Fara, E., Laffont, R. & Pellenard, P. 2010 High-
360 resolution dynamics of Early Jurassic marine extinctions: the case of Pliensbachian–
361 Toarcian ammonites (Cephalopoda). *Journal of the Geological Society* **167**, 21-33.
362 (doi:10.1144/0016-76492009-068).
- 363 [14] McElwain, J.C., Beerling, D.J. & Woodward, F.I. 1999 Fossil plants and global warming
364 at the Triassic-Jurassic Boundary. *Science* **285**, 1386-1390.
- 365 [15] McElwain, J.C., Wade-Murphy, J. & Hesselbo, S.P. 2005 Changes in carbon dioxide
366 during an oceanic anoxic event linked to intrusion into Gondwana coals. *Nature* **435**,
367 479-482.
368 (doi:http://www.nature.com/nature/journal/v435/n7041/supinfo/nature03618_S1.htm
369 l).
- 370 [16] Danise, S., Twitchett, R.J. & Little, C.T.S. 2015 Environmental controls on Jurassic
371 marine ecosystems during global warming. *Geology* **43**, 263-266.
372 (doi:10.1130/g36390.1).
- 373 [17] Danise, S., Twitchett, R.J., Little, C.T.S. & Clémence, M.-E. 2013 The Impact of Global
374 Warming and Anoxia on Marine Benthic Community Dynamics: an Example from
375 the Toarcian (Early Jurassic). *PLoS ONE* **8**, e56255.
376 (doi:10.1371/journal.pone.0056255).
- 377 [18] Schaller, M.F., Wright, J.D. & Kent, D.V. 2011 Atmospheric PCO_2 Perturbations
378 Associated with the Central Atlantic Magmatic Province. *Science* **331**, 1404-1409.
379 (doi:10.1126/science.1199011).

- 380 [19] Blackburn, T.J., Olsen, P.E., Bowring, S.A., McLean, N.M., Kent, D.V., Puffer, J.,
381 McHone, G., Rasbury, E.T. & Et-Touhami, M. 2013 Zircon U-Pb Geochronology
382 Links the End-Triassic Extinction with the Central Atlantic Magmatic Province.
383 *Science* **340**, 941-945. (doi:10.1126/science.1234204).
- 384 [20] Whiteside, J., Olsen, P.E., Eglinton, T., Brookfield, M.E. & Sambrotto, R.N. 2010
385 Compound-specific carbon isotopes from Earth's largest flood basalt eruptions
386 directly linked to the end-Triassic mass extinction. *PNAS* **107**, 6721-6725.
- 387 [21] Ruhl, M., Bonis, N.R., Reichart, G.-J., Damsté, J.S.S. & Kürschner, W.M. 2011
388 Atmospheric Carbon Injection Linked to End-Triassic Mass Extinction. *Science* **333**,
389 430-434. (doi:10.1126/science.1204255).
- 390 [22] Burgess, S.D., Bowring, S.A., Fleming, T.H. & Elliot, D.H. 2015 High-precision
391 geochronology links the Ferrar large igneous province with early-Jurassic ocean
392 anoxia and biotic crisis. *Earth and Planetary Science Letters* **415**, 90-99.
393 (doi:<https://doi.org/10.1016/j.epsl.2015.01.037>).
- 394 [23] Davies, J.H.F.L., Marzoli, A., Bertrand, H., Youbi, N., Ernesto, M. & Schaltegger, U.
395 2017 End-Triassic mass extinction started by intrusive CAMP activity. *Nature*
396 *Communications* **8**, 15596. (doi:10.1038/ncomms15596
397 <https://www.nature.com/articles/ncomms15596#supplementary-information>).
- 398 [24] Corso, J.D., Marzoli, A., Tateo, F., Jenkyns, H.C., Bertrand, H., Youbi, N., Mahmoudi,
399 A., Font, E., Buratti, N. & Cirilli, S. 2014 The dawn of CAMP volcanism and its
400 bearing on the end-Triassic carbon cycle disruption. *Journal of the Geological Society*
401 **171**, 153-164. (doi:10.1144/jgs2013-063).

- 402 [25] Heimdal, T.H., Svensen, H.H., Ramezani, J., Iyer, K., Pereira, E., Rodrigues, R., Jones,
403 M.T. & Callegaro, S. 2018 Large-scale sill emplacement in Brazil as a trigger for the
404 end-Triassic crisis. *Sci Rep* **8**, 141. (doi:10.1038/s41598-017-18629-8).
- 405 [26] Caruthers, A.H., Smith, P.L. & Gröcke, D.R. 2014 The Pliensbachian–Toarcian (Early
406 Jurassic) extinction: A North American perspective. *Geological Society of America*
407 *Special Papers* **505**, 225-243. (doi:10.1130/2014.2505(11)).
- 408 [27] Hautmann, M., Benton, M.J. & Tomasovych, A. 2008 Catastrophic ocean acidification
409 at the Triassic-Jurassic boundary. *N. Jb. Geol. Palaont. Abh.* **249**, 119-127.
- 410 [28] Trecalli, A., Spangenberg, J., Adatte, T., Föllmi, K.B. & Parente, M. 2012 Carbonate
411 platform evidence of ocean acidification at the onset of the early Toarcian oceanic
412 anoxic event. *Earth and Planetary Science Letters* **357**, 214-225.
413 (doi:https://doi.org/10.1016/j.epsl.2012.09.043).
- 414 [29] Hautmann, M. 2004 Effect of end-Triassic CO₂ maximum on carbonate sedimentation
415 and marine mass extinction. *Facies* **50**, 257-261. (doi:10.1007/s10347-004-0020-y).
- 416 [30] Kiessling, W., Aberhan, M., Brenneis, B. & Wagner, P.J. 2007 Extinction trajectories of
417 benthic organisms across the Triassic–Jurassic boundary. *Palaeogeography,*
418 *Palaeoclimatology, Palaeoecology* **244**, 201-222. (doi:10.1016/j.palaeo.2006.06.029).
- 419 [31] Dunhill, A.M., Foster, W.J., Sciberras, J. & Twitchett, R.J. 2018 Impact of the Late
420 Triassic mass extinction on functional diversity and composition of marine
421 ecosystems. *Palaeontology* **61**, 133-148. (doi:10.1111/pala.12332).
- 422 [32] Kiessling, W. & Aberhan, M. 2007 Environmental determinants of marine benthic
423 biodiversity dynamics through Triassic-Jurassic time. *Paleobiology* **33**, 414-434.
- 424 [33] Bambach, R.K. 2006 Phanerozoic biodiversity and mass extinctions. *Annual Review of*
425 *Earth and Planetary Sciences* **34**, 127-155.

- 426 [34] Bambach, R.K., Knoll, A.H. & Wang, S.C. 2004 Origination, extinction, and mass
427 depletions of marine diversity. *Paleobiology* **30**, 522-542. (doi:10.1666/0094-
428 8373(2004)030<0522:OEAMDO>2.0.CO;2).
- 429 [35] Martindale, R.C. & Aberhan, M. 2017 Response of macrobenthic communities to the
430 Toarcian Oceanic Anoxic Event in northeastern Panthalassa (Ya Ha Tinda, Alberta,
431 Canada). *Palaeogeography, Palaeoclimatology, Palaeoecology* **478**, 103-120.
432 (doi:http://dx.doi.org/10.1016/j.palaeo.2017.01.009).
- 433 [36] Harries, P.J. & Little, C.T.S. 1999 The early Toarcian (Early Jurassic) and the
434 Cenomanian–Turonian (Late Cretaceous) mass extinctions: similarities and contrasts.
435 *Palaeogeography, Palaeoclimatology, Palaeoecology* **154**, 39-66.
436 (doi:https://doi.org/10.1016/S0031-0182(99)00086-3).
- 437 [37] Al-Suwaidi, A.H., Angelozzi, G.N., Baudin, F., Damborenea, S.E., Hesselbo, S.P.,
438 Jenkyns, H.C., Manceñido, M.O. & Riccardi, A.C. 2010 First record of the Early
439 Toarcian Oceanic Anoxic Event from the Southern Hemisphere, Neuquén Basin,
440 Argentina. *Journal of the Geological Society* **167**, 633-636. (doi:10.1144/0016-
441 76492010-025).
- 442 [38] Jablonski, D. 1986 Background and Mass Extinctions: The Alternation of
443 Macroevolutionary Regimes. *Science* **231**, 129-133.
444 (doi:10.1126/science.231.4734.129).
- 445 [39] Clapham, M.E., Kiessling, W., Fürsich, F., Aberhan, M., Rees, A., Pálffy, J., Carrano,
446 M.T., Bottjer, D.B., McGowan, A.J., Miller, A.I., et al. 2016 Taxonomic occurrences
447 of Triassic to Jurassic marine animals. Palaeobiology Database, accessed 13 April
448 2016, <http://paleobiodb.org>

- 449 [40] Dunhill, A.M., Sciberras, J., Foster, W.J. & Twitchett, R.J. 2018 Data from: Functional
450 diversity of marine ecosystems across the Late Triassic mass extinction. Dryad Data
451 Repository. <https://doi.org/10.5061/dryad.bg30k>
- 452 [41] Zuur, A.F., Ieno, E.N., Walker, N., J., Saveliev, A.A. & Smith, G.M. 2009 *Mixed Effects*
453 *Models and Extensions in Ecology with R*. New York, Springer.
- 454 [42] McCullagh, P. & Nelder, J. 1989 Generalized linear models. (p. 511. New York,
455 Chapman and Hall.
- 456 [43] Hilbe, J.M. 2011 *Negative binomial regression*, Cambridge University Press.
- 457 [44] R Development Core Team. 2017. R: A language and environment for statistical
458 computing. v 3.4.3. The R Foundation for Statistical Computing, Vienna, Austria.
459 ISBN 3-900051-07-0, URL <http://www.R-project.org/>.
- 460 [45] Van de Schootbrugge, B. & Wignall, P.B. 2016 A tale of two extinctions: converging
461 end-Permian and end-Triassic scenarios. *Geological Magazine* **153**, 332-354.
462 (doi:10.1017/s0016756815000643).
- 463 [46] Barras, C.G. & Twitchett, R.J. 2007 Response of the marine infauna to Triassic-Jurassic
464 environmental change: Ichnological data from southern England. *Palaeogeography,*
465 *Palaeoclimatology, Palaeoecology* **244**, 223-241.
- 466 [47] Hautmann, M., Stiller, F., Huawei, C. & Jingeng, S. 2008 Extinction-Recovery Pattern
467 of Level-Bottom Faunas Across the Triassic-Jurassic Boundary in Tibet: Implications
468 for Potential Killing Mechanisms. *PALAIOS* **23**, 711-718.
469 (doi:10.2110/palo.2008.p08-005r).
- 470 [48] Mander, L., Twitchett, R.J. & Benton, M.J. 2008 Palaeoecology of the Late Triassic
471 extinction event in the SW UK. *Journal of the Geological Society, London* **165**, 319-
472 332.

- 473 [49] Thorne, P.M., Ruta, M. & Benton, M.J. 2011 Resetting the evolution of marine reptiles
474 at the Triassic-Jurassic boundary. *Proceedings of the National Academy of Sciences*
475 **108**, 8339-8344. (doi:10.1073/pnas.1018959108).
- 476 [50] Tackett, L.S. & Bottjer, D.J. 2016 Paleoeological succession of Norian (Late Triassic)
477 benthic fauna in eastern Panthalassa (Luning and Gabbs formations, west-central
478 Nevada). *PALAIOS*, 190-202. (doi:10.2110/palo.2015.070).
- 479 [51] Dal Corso, J., Gianolla, P., Newton, R.J., Franceschi, M., Roghi, G., Caggiati, M.,
480 Raucsik, B., Budai, T., Haas, J. & Preto, N. 2015 Carbon isotope records reveal
481 synchronicity between carbon cycle perturbation and the “Carnian Pluvial Event” in
482 the Tethys realm (Late Triassic). *Global and Planetary Change* **127**, 79-90.
483 (doi:http://dx.doi.org/10.1016/j.gloplacha.2015.01.013).
- 484 [52] Knoll, A.H., Bambach, R.K., Payne, J.L., Pruss, S. & Fischer, W.W. 2007
485 Paleophysiology and end-Permian mass extinction. *Earth and Planetary Science*
486 *Letters* **256**, 295-313. (doi:https://doi.org/10.1016/j.epsl.2007.02.018).
- 487 [53] Bambach, R.K., Bush, A.M. & Erwin, D.H. 2007 Autecology and the filling of
488 ecospace: Key metazoan radiations. *Palaeontology* **50**, 1-22. (doi:10.1111/j.1475-
489 4983.2006.00611.x).
- 490

Table 1. Summary of intrinsic and extrinsic ecological determinants of extinction.

| | Determinant | Levels | Ref. |
|-----------|--------------------|--|-------------|
| Intrinsic | Motility | motile non-motile | [31, 53] |
| | Tiering | pelagic epifaunal infaunal | [31, 53] |
| | Feeding | suspension deposit/mining grazing predatory photosymbiotic | [31, 53] |
| | Calcification | heavy moderate light | [31, 52] |
| Extrinsic | Latitude | polar (>60°) mid-latitude (30-60°) tropical (<30°) | [31, 39] |
| | Palaeo-ocean basin | Panthalassa Tethys Boreal | [31, 39] |
| | Environment | Onshore Offshore Reef | [31, 39] |

Table 2. Summary of full and best fitting GLMs for predicting extinction through hyperthermal mass extinctions and periods of background extinction. LTE = Late Triassic mass extinction; EToE = early Toarcian extinction. “Full model” includes all variables (see electronic supplementary material for model definitions). “Best model” is the best fitting model following model selection procedure described in methods section. “Significant variables” identifies variables identified as significantly determining extinction; “significant arguments” identifies which arguments of a particular variables show significantly higher (+) or significantly lower (-) extinction than other arguments of that variables. “Explained deviance” = (null deviance-residual deviance)/null deviance; provides estimate of goodness-of-fit of model to extinction variable. Where no variables are listed, no variables significantly determine extinction.

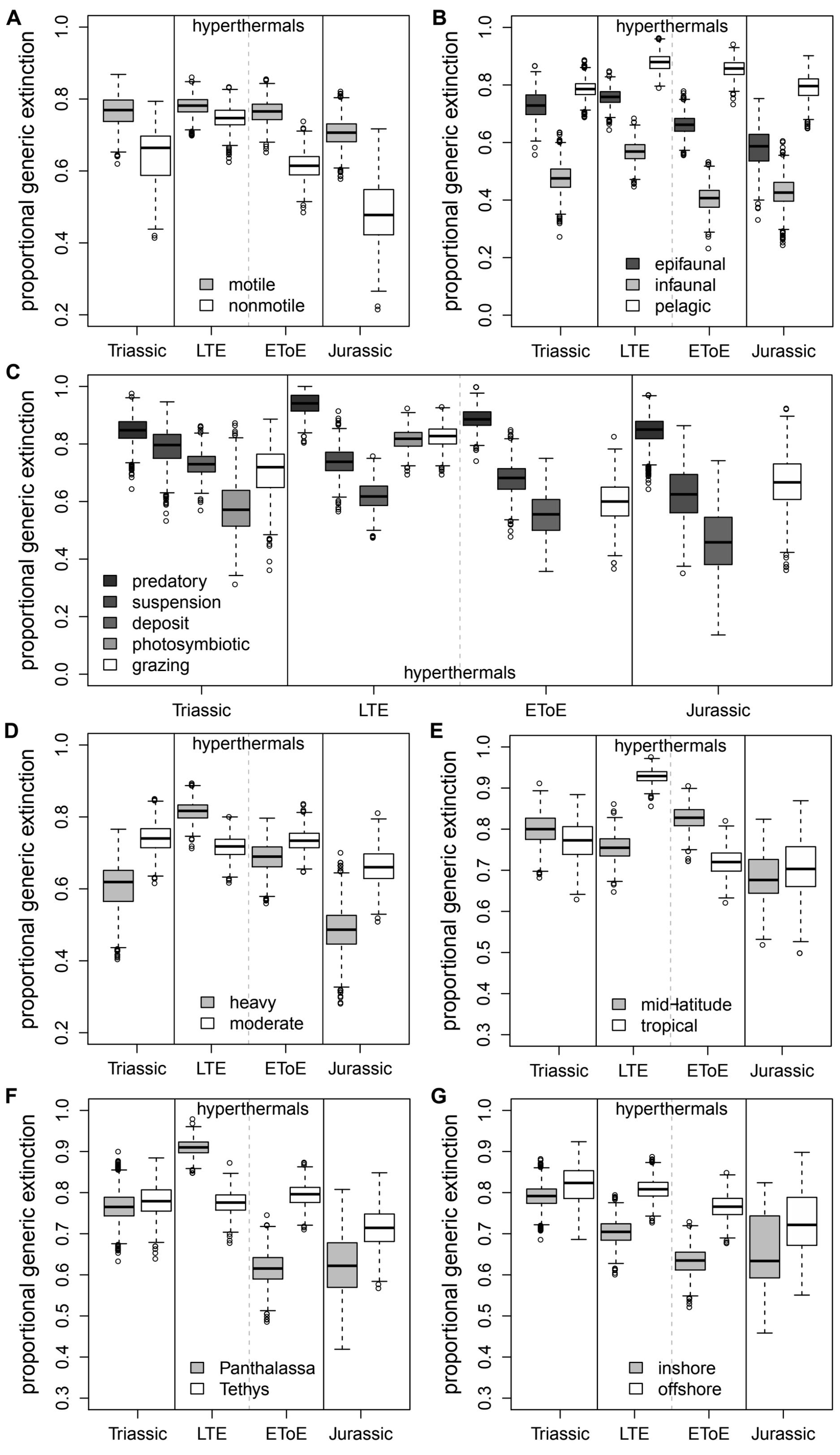
| Model | Full model | | | Best model | | | |
|---------------------|-----------------------|---|--|-----------------------|-----------------------|---|------|
| | significant variables | significant arguments | explained deviance | significant variables | significant arguments | explained deviance | |
| LTE | all | latitude | tropical (+) predatory (+) | 0.78 | latitude | tropical (+) predatory (+) | 0.32 |
| | all no env | feeding | photosymbiotic (+) suspension (+) | 0.60 | feeding | photosymbiotic (+) suspension (+) | 0.43 |
| | extrinsic | latitude | tropical (+) | 0.47 | latitude | tropical (+) | 0.32 |
| | extrinsic no env | - | - | 0.18 | - | - | - |
| | intrinsic | - | - | 0.40 | - | - | - |
| | intrinsic no env | feeding | predatory (+) photosymbiotic (+) | 0.48 | feeding | predatory (+) photosymbiotic (+) suspension (+) | 0.43 |
| EToE | all | - | - | 0.68 | - | - | - |
| | all no env | motility feeding latitude basin calcification | non-motile (+) photosymbiotic (+) polar (-) Boreal (+) light (-) | 0.71 | feeding basin | photosymbiotic (+) Boreal (+) | 0.5 |
| | extrinsic | - | - | 0.23 | - | - | - |
| | extrinsic no env | basin | Boreal (+) | 0.17 | basin | Panthalassa (-) | 0.14 |
| | intrinsic | - | - | 0.44 | - | - | - |
| | intrinsic no env | feeding | photosymbiotic (+) | 0.48 | feeding | photosymbiotic (+) | 0.33 |
| Triassic background | all | - | - | 0.34 | - | - | - |
| | all no env | - | - | 0.27 | - | - | - |
| | extrinsic | environment | reef (-) | 0.19 | environment | reef (-) | 0.19 |
| | extrinsic no env | - | - | 0.03 | - | - | - |
| | intrinsic | - | - | 0.31 | - | - | - |
| | intrinsic no env | - | - | 0.23 | - | - | - |
| Jurassic background | all | motility basin environment | pelagic (+) Tethys (+) reef (-) | 0.35 | basin environment | Tethys (+) reef (-) | 0.16 |
| | all no env | - | - | 0.28 | - | - | - |
| | extrinsic | basin | Tethys (+) | 0.16 | basin | Tethys (+) | 0.07 |
| | extrinsic no env | - | - | 0.03 | - | - | - |
| | intrinsic | - | - | 0.18 | - | - | - |
| | intrinsic no env | - | - | 0.25 | - | - | - |

493

494

495 **Figure legends**

496 **Figure 1.** Boxplots across all subsamples of proportional generic extinction per ecological
497 variable through Late Triassic background periods (Triassic), the Late Triassic mass extinction
498 (LTE), the early Toarcian extinction (EToE), and Early Jurassic background periods (Jurassic)
499 by (A) motility; (B) tiering; (C) feeding; (D) calcification; (E) latitude; (F) ocean basin; (G)
500 depositional setting. Proportional generic extinction is calculated from a subsample of $n=250$
501 across 1,000 iterations except for feeding which is calculated from a subsample of $n=75$ across
502 1,000 iterations. The solid black lines inside the boxes represent the medians, the top and
503 bottom edges of the boxes correspond to the first and third quartiles, and whiskers represent
504 the lowest and highest subsampled values within 1.5 times the interquartile range. Points
505 outside the whiskers are outliers.



Supplementary Material

Modelling determinants of extinction across two Mesozoic hyperthermal events

Alexander M. Dunhill¹, William J. Foster², Sandro Azaele³, James Sciberras⁴ and Richard J. Twitchett⁵

¹School of Earth and Environment, University of Leeds, Leeds LS2 9JT, UK

²Museum für Naturkunde, Leibniz-Institut für Evolutions- und Biodiversitätsforschung, Berlin, Deutschland³School of Mathematics, University of Leeds, Leeds LS2 9JT, UK

⁴Department of Biology and Biochemistry, University of Bath, Claverton Down, BA2 7AY, UK

⁵Department of Earth Sciences, Natural History Museum, London, SW7 5BD, UK

Author for correspondence: Alexander M. Dunhill

e-mail: a.dunhill@leeds.ac.uk

Supplementary methods

Table S1: List of variables in each GLM. “no env” refers to models run without depositional environmental setting data which is only present for around 50% of the data set. Therefore, the removal of the depositional environmental setting doubles the sample size.

| Model | Determinants |
|------------------|---|
| all | motility; tiering; feeding; calcification; latitude; palaeocean basin; depositional setting |
| all no env | motility; tiering; feeding; calcification; latitude; palaeocean basin |
| extrinsic | latitude; palaeocean basin; depositional setting |
| extrinsic no env | latitude; palaeocean basin |
| intrinsic | motility; tiering; feeding; calcification |
| intrinsic no env | motility; tiering; feeding; calcification |

Supplementary results

GLM outputs

LTE

Model = all

```
> GLM_LTE_Alii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fLATITUDE + fBASIN + fENVIRONMENT +
+ fCALCIFICATION,
+ family = quasibinomial, data = LTE)
> summary(GLM_LTE_Alii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fLATITUDE + fBASIN + fENVIRONMENT + fCALCIFICATION, family =
quasibinomial,
     data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.52039 | -0.07450 | 0.00003 | 0.15316 | 0.55019 |

Coefficients: (1 not defined because of singularities)

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 2.6340 | 1.5901 | 1.656 | 0.1320 |
| fMOTILITYnonmotile | -0.6121 | 0.7194 | -0.851 | 0.4170 |
| fTIERINGinfaunal | -1.0261 | 0.6461 | -1.588 | 0.1468 |
| fTIERINGpelagic | -2.2947 | 1.8030 | -1.273 | 0.2350 |
| fFEEDINGphotosymbiotic | 1.6102 | 2.0644 | 0.780 | 0.4554 |
| fFEEDINGpredatory | NA | NA | NA | NA |
| fFEEDINGSuspension | 0.7476 | 0.8971 | 0.833 | 0.4262 |
| fLATITUDEET | 2.9924 | 1.1182 | 2.676 | 0.0254 * |
| fBASINTethys | -0.9580 | 0.9703 | -0.987 | 0.3493 |
| fENVIRONMENTOffshore | 1.1036 | 0.7170 | 1.539 | 0.1581 |
| fENVIRONMENTReef | 16.1132 | 3723.9890 | 0.004 | 0.9966 |
| fCALCIFICATIONmoderate | -1.7223 | 1.0296 | -1.673 | 0.1287 |

Signif. codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1443989)

Null deviance: 6.1229 on 19 degrees of freedom
Residual deviance: 1.3575 on 9 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 19

```
> # find best model  
>  
> drop1(GLM_LTE_Alii, test = "F")
```

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
fENVIRONMENT + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|-------------|
| <none> | | 1.3576 | | |
| fMOTILITY | 1 | 1.4633 | 0.7010 | 0.424115 |
| fTIERING | 1 | 1.7343 | 2.4975 | 0.148485 |
| fFEEDING | 2 | 1.4873 | 0.4300 | 0.663196 |
| fLATITUDE | 1 | 3.1510 | 11.8898 | 0.007296 ** |
| fBASIN | 1 | 1.5097 | 1.0090 | 0.341381 |
| fENVIRONMENT | 2 | 1.7986 | 1.4619 | 0.281992 |
| fCALCIFICATION | 1 | 1.8601 | 3.3320 | 0.101236 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
>  
> # best model  
>  
> GLM_LTE_Aliiii <- glm(extinction ~ fLATITUDE,  
+ family = quasibinomial, data = LTE)  
>  
> summary(GLM_LTE_Aliiii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE, family = quasibinomial,  
data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.79806 | -0.18099 | 0.08947 | 0.30075 | 0.81341 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|-------------|----------|------------|---------|------------|
| (Intercept) | 0.9362 | 0.2806 | 3.336 | 0.00367 ** |
| fLATITUDE | 2.1372 | 0.8825 | 2.422 | 0.02623 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.207126)

Null deviance: 6.1229 on 19 degrees of freedom
Residual deviance: 4.1803 on 18 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 6

Model = all no env

```
> GLM_LTE_Blii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +  
+ fLATITUDE + fBASIN + fCALCIFICATION,  
+ family = quasibinomial, data = LTE)  
> summary(GLM_LTE_Blii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
fLATITUDE + fBASIN + fCALCIFICATION, family = quasibinomial,  
data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -0.9482 | -0.1619 | 0.1085 | 0.2678 | 0.7364 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 3.30683 | 1.90023 | 1.740 | 0.0965 . |
| fMOTILITYnonmotile | -0.16350 | 0.48100 | -0.340 | 0.7373 |
| fTIERINGinfaunal | 0.04299 | 0.51878 | 0.083 | 0.9347 |
| fTIERINGpelagic | -0.97680 | 1.26559 | -0.772 | 0.4488 |
| fFEEDINGgrazing | -1.00009 | 1.39971 | -0.715 | 0.4828 |
| fFEEDINGphotosymbiotic | 2.28581 | 1.50753 | 1.516 | 0.1444 |
| fFEEDINGpredatory | 3.24156 | 1.46390 | 2.214 | 0.0380 * |
| fFEEDINGSuspension | 0.80595 | 0.79223 | 1.017 | 0.3206 |

```

fLATITUDEP          -1.15779    1.14155   -1.014    0.3220
fLATITUDET          0.77683    0.48202    1.612    0.1220
fBASINPanthalassa  -2.38603    1.34792   -1.770    0.0912 .
fBASINTethys       -2.87926    1.46745   -1.962    0.0631 .
fCALCIFICATIONlight -1.79362    1.26434   -1.419    0.1707
fCALCIFICATIONmoderate -1.03692    0.65945   -1.572    0.1308

```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2016485)

```

Null deviance: 11.300  on 34  degrees of freedom
Residual deviance:  4.467  on 21  degrees of freedom
AIC: NA

```

Number of Fisher Scoring iterations: 6

>

> # find best model

>

> drop1(GLM_LTE_Blii, test = "F")

Single term deletions

Model:

```

extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
  fCALCIFICATION

```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|-----------|
| <none> | | 4.4670 | | |
| fMOTILITY | 1 | 4.4904 | 0.1099 | 0.74358 |
| fTIERING | 2 | 4.6018 | 0.3168 | 0.73189 |
| fFEEDING | 4 | 6.8103 | 2.7541 | 0.05507 . |
| fLATITUDE | 2 | 5.2620 | 1.8687 | 0.17909 |
| fBASIN | 2 | 5.2717 | 1.8916 | 0.17566 |
| fCALCIFICATION | 2 | 5.1245 | 1.5455 | 0.23650 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

>

> # best model

```

> GLM_LTE_Bliiii <- glm(extinction ~ fFEEDING,
+ family = quasibinomial, data = LTE)

```

```
>
> summary(GLM_LTE_Bliii)
```

Call:

```
glm(formula = extinction ~ fFEEDING, family = quasibinomial,
     data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.94003 | -0.24861 | 0.07654 | 0.30255 | 0.81753 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|------------|
| (Intercept) | -0.5878 | 0.6253 | -0.940 | 0.35471 |
| fFEEDINGgrazing | -1.0217 | 1.2975 | -0.787 | 0.43724 |
| fFEEDINGphotosymbiotic | 3.6779 | 1.2119 | 3.035 | 0.00494 ** |
| fFEEDINGpredatory | 2.5197 | 0.7893 | 3.192 | 0.00330 ** |
| fFEEDINGSuspension | 1.5121 | 0.6580 | 2.298 | 0.02871 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1795301)

Null deviance: 11.2997 on 34 degrees of freedom
Residual deviance: 6.4135 on 30 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 5

Model = extrinsic

```
> GLM_LTE_A3ii <- glm(extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
+ family = quasibinomial, data = LTE)
> summary(GLM_LTE_A3ii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
     family = quasibinomial, data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -0.7523 | -0.1184 | 0.0339 | 0.3520 | 0.8552 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|-----------|------------|---------|----------|
| (Intercept) | 7.396e-01 | 8.197e-01 | 0.902 | 0.381 |
| fLATITUDE | 1.933e+00 | 9.244e-01 | 2.090 | 0.054 . |
| fBASINTethys | 7.801e-02 | 8.717e-01 | 0.089 | 0.930 |
| fENVIRONMENTOffshore | 2.407e-02 | 5.685e-01 | 0.042 | 0.967 |
| fENVIRONMENTReef | 1.884e+01 | 4.216e+03 | 0.004 | 0.996 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1914137)

Null deviance: 6.1229 on 19 degrees of freedom
Residual deviance: 3.2701 on 15 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 19

```
> # find best model
>
> drop1(GLM_LTE_A3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN + fENVIRONMENT
          Df Deviance F value Pr(>F)
<none>          3.2701
fLATITUDE      1  4.3804  5.0929 0.03937 *
fBASIN         1  3.2716  0.0070 0.93446
fENVIRONMENT   2  4.0752  1.8465 0.19191
```

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
>
> # best model
>
> GLM_LTE_A3iii <- glm(extinction ~ fLATITUDE,
+ family = quasibinomial, data = LTE)
>
> summary(GLM_LTE_A3iii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE, family = quasibinomial,  
     data = LTE)
```

Deviance Residuals:

| | Min | 1Q | Median | 3Q | Max |
|--|----------|----------|---------|---------|---------|
| | -0.79806 | -0.18099 | 0.08947 | 0.30075 | 0.81341 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|-------------|----------|------------|---------|------------|
| (Intercept) | 0.9362 | 0.2806 | 3.336 | 0.00367 ** |
| fLATITUDET | 2.1372 | 0.8825 | 2.422 | 0.02623 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.207126)

Null deviance: 6.1229 on 19 degrees of freedom
Residual deviance: 4.1803 on 18 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 6

Model = extrinsic no env

```
> GLM_LTE_B3ii <- glm(extinction ~ fLATITUDE + fBASIN,  
+ family = quasibinomial, data = LTE)  
> summary(GLM_LTE_B3ii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN, family = quasibinomial,  
     data = LTE)
```

Deviance Residuals:

| | Min | 1Q | Median | 3Q | Max |
|--|----------|----------|---------|---------|---------|
| | -1.41178 | -0.28614 | 0.09095 | 0.42332 | 0.95988 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|-------------|----------|------------|---------|----------|
| (Intercept) | 2.8905 | 1.4932 | 1.936 | 0.0624 . |
| fLATITUDEP | -0.8110 | 1.1499 | -0.705 | 0.4861 |
| fLATITUDET | 0.8358 | 0.4875 | 1.714 | 0.0968 . |

```
fBASINPanthalassa -1.6740    1.4239 -1.176    0.2490
fBASINTethys      -2.3546    1.5076 -1.562    0.1288
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

(Dispersion parameter for quasibinomial family taken to be 0.2688368)

```
Null deviance: 11.2997 on 34 degrees of freedom
Residual deviance: 9.3068 on 30 degrees of freedom
AIC: NA
```

Number of Fisher Scoring iterations: 5

```
>
> # find best model
>
> drop1(GLM_LTE_B3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN
          Df Deviance F value Pr(>F)
<none>          9.3068
fLATITUDE  2  10.3885  1.7434 0.1922
fBASIN     2  10.2198  1.4715 0.2457
>
```

Model = intrinsic

```
> GLM_LTE_A4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fCALCIFICATION,
+ family = quasibinomial, data = LTE)
> summary(GLM_LTE_A4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fCALCIFICATION, family = quasibinomial, data = LTE)
```

Deviance Residuals:

```
      Min       1Q   Median       3Q      Max
-0.71433 -0.26270  0.09925  0.32029  0.81238
```

Coefficients: (1 not defined because of singularities)

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 2.7289 | 1.4630 | 1.865 | 0.0849 . |
| fMOTILITYnonmotile | -0.4503 | 0.8457 | -0.532 | 0.6034 |
| fTIERINGinfaunal | -0.4901 | 0.7625 | -0.643 | 0.5315 |
| fTIERINGpelagic | 0.8460 | 1.5784 | 0.536 | 0.6010 |
| fFEEDINGphotosymbiotic | 1.5392 | 2.4689 | 0.623 | 0.5438 |
| fFEEDINGpredatory | NA | NA | NA | NA |
| fFEEDINGSuspension | 0.6659 | 0.9222 | 0.722 | 0.4831 |
| fCALCIFICATIONmoderate | -1.9654 | 1.2570 | -1.564 | 0.1419 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2485554)

Null deviance: 6.1229 on 19 degrees of freedom
Residual deviance: 3.6698 on 13 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 6

```
>  
> # find best model  
>  
> drop1(GLM_LTE_A4ii, test = "F")
```

Single term deletions

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 3.6698 | | |
| fMOTILITY | 1 | 3.7409 | 0.2521 | 0.6240 |
| fTIERING | 1 | 3.7737 | 0.3683 | 0.5544 |
| fFEEDING | 2 | 3.8385 | 0.2988 | 0.7467 |
| fCALCIFICATION | 1 | 4.5139 | 2.9904 | 0.1074 |

```
>
```

Model = intrinsic no env

```
> GLM_LTE_B4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +  
+ fCALCIFICATION,  
+ family = quasibinomial, data = LTE)
```

```
> summary(GLM_LTE_B4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
     fCALCIFICATION, family = quasibinomial, data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -0.8854 | -0.2175 | 0.1099 | 0.2983 | 0.8137 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 0.05624 | 0.87501 | 0.064 | 0.9493 |
| fMOTILITYnonmotile | -0.12162 | 0.47360 | -0.257 | 0.7994 |
| fTIERINGinfaunal | -0.28702 | 0.49595 | -0.579 | 0.5680 |
| fTIERINGpelagic | -0.76296 | 1.31530 | -0.580 | 0.5671 |
| fFEEDINGgrazing | -1.16221 | 1.41697 | -0.820 | 0.4198 |
| fFEEDINGphotosymbiotic | 3.15546 | 1.43793 | 2.194 | 0.0377 * |
| fFEEDINGpredatory | 3.37268 | 1.53674 | 2.195 | 0.0377 * |
| fFEEDINGSuspension | 1.38256 | 0.75596 | 1.829 | 0.0794 . |
| fCALCIFICATIONlight | -1.68717 | 1.26360 | -1.335 | 0.1938 |
| fCALCIFICATIONmoderate | -0.50346 | 0.57528 | -0.875 | 0.3898 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2078436)

Null deviance: 11.2997 on 34 degrees of freedom
Residual deviance: 5.8432 on 25 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 5

```
>
```

```
> # find best model
```

```
>
```

```
> drop1(GLM_LTE_B4ii, test = "F")
```

Single term deletions

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|----------|
| <none> | | 5.8432 | | |
| fMOTILITY | 1 | 5.8569 | 0.0588 | 0.8104 |
| fTIERING | 2 | 5.9886 | 0.3111 | 0.7354 |
| fFEEDING | 4 | 9.0390 | 3.4183 | 0.0232 * |
| fCALCIFICATION | 2 | 6.2321 | 0.8320 | 0.4469 |

Signif. codes: 0 '****' 0.001 '***' 0.01 '**' 0.05 '.' 0.1 ' ' 1

>

> # best model

>

> GLM_LTE_B4iii <- glm(extinction ~ fFEEDING,

+ family = quasibinomial, data = LTE)

> summary(GLM_LTE_B4iii)

Call:

```
glm(formula = extinction ~ fFEEDING, family = quasibinomial,
     data = LTE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.94003 | -0.24861 | 0.07654 | 0.30255 | 0.81753 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|------------|
| (Intercept) | -0.5878 | 0.6253 | -0.940 | 0.35471 |
| fFEEDINGgrazing | -1.0217 | 1.2975 | -0.787 | 0.43724 |
| fFEEDINGphotosymbiotic | 3.6779 | 1.2119 | 3.035 | 0.00494 ** |
| fFEEDINGpredatory | 2.5197 | 0.7893 | 3.192 | 0.00330 ** |
| fFEEDINGSuspension | 1.5121 | 0.6580 | 2.298 | 0.02871 * |

Signif. codes: 0 '****' 0.001 '***' 0.01 '**' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1795301)

Null deviance: 11.2997 on 34 degrees of freedom

Residual deviance: 6.4135 on 30 degrees of freedom

AIC: NA

Number of Fisher Scoring iterations: 5

EToE

Model = all

```
> GLM_OAE_Alii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fLATITUDE + fBASIN + fENVIRONMENT +
+ fCALCIFICATION,
+ family = quasibinomial, data = OAE)
> summary(GLM_OAE_Alii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fLATITUDE + fBASIN + fENVIRONMENT + fCALCIFICATION, family =
quasibinomial,
     data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -0.4337 | -0.1391 | 0.0000 | 0.2102 | 0.4509 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 0.7777 | 1.1233 | 0.692 | 0.5031 |
| fMOTILITYnonmotile | 0.7891 | 0.5491 | 1.437 | 0.1785 |
| fTIERINGinfaunal | -0.1401 | 0.4658 | -0.301 | 0.7693 |
| fTIERINGpelagic | 1.1627 | 1.0537 | 1.103 | 0.2934 |
| fFEEDINGgrazing | 1.1905 | 0.7983 | 1.491 | 0.1640 |
| fFEEDINGphotosymbiotic | -1.7160 | 1.2277 | -1.398 | 0.1897 |
| fFEEDINGpredatory | 0.7080 | 0.9654 | 0.733 | 0.4787 |
| fFEEDINGSuspension | -0.3624 | 0.7043 | -0.515 | 0.6170 |
| fLATITUDET | -1.0360 | 0.5591 | -1.853 | 0.0909 . |
| fBASINTethys | 0.8423 | 0.5712 | 1.475 | 0.1683 |
| fENVIRONMENTOffshore | -1.3328 | 0.6372 | -2.092 | 0.0605 . |
| fENVIRONMENTReef | 3.3873 | 1.8921 | 1.790 | 0.1009 |
| fCALCIFICATIONmoderate | 0.1033 | 0.6401 | 0.161 | 0.8747 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1186548)

Null deviance: 4.1787 on 23 degrees of freedom
Residual deviance: 1.3477 on 11 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 5

```
>
> # find best model
>
> drop1(GLM_OAE_Alii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
  fENVIRONMENT + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|-----------|
| <none> | | 1.3477 | | |
| fMOTILITY | 1 | 1.5974 | 2.0381 | 0.18117 |
| fTIERING | 2 | 1.4956 | 0.6036 | 0.56398 |
| fFEEDING | 4 | 2.0222 | 1.3763 | 0.30423 |
| fLATITUDE | 1 | 1.7801 | 3.5289 | 0.08705 . |
| fBASIN | 1 | 1.6114 | 2.1526 | 0.17033 |
| fENVIRONMENT | 2 | 2.2438 | 3.6571 | 0.06058 . |
| fCALCIFICATION | 1 | 1.3508 | 0.0252 | 0.87681 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
>
> GLM_OAE_Aliiii <- glm(extinction ~ fENVIRONMENT,
+ family = quasibinomial, data = post)
>
> summary(GLM_OAE_Aliiii)
```

Call:

```
glm(formula = extinction ~ fENVIRONMENT, family = quasibinomial,
     data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|----------|---------|---------|
| -1.20688 | -0.35287 | -0.00746 | 0.35180 | 1.15096 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|-----------|------------|---------|----------|
| (Intercept) | 0.069080 | 0.205914 | 0.335 | 0.739 |
| fENVIRONMENTOffshore | -0.006514 | 0.308166 | -0.021 | 0.983 |

```
fENVIRONMENTreef      -1.860839   1.175805  -1.583   0.119
```

(Dispersion parameter for quasibinomial family taken to be 0.3282142)

```
Null deviance: 23.615  on 57  degrees of freedom  
Residual deviance: 22.425  on 55  degrees of freedom  
AIC: NA
```

Number of Fisher Scoring iterations: 4

Model = all no env

```
> GLM_OAE_Blii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +  
+ fLATITUDE + fBASIN + fCALCIFICATION,  
+ family = quasibinomial, data = OAE)  
> summary(GLM_OAE_Blii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
    fLATITUDE + fBASIN + fCALCIFICATION, family = quasibinomial,  
    data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.72266 | -0.16491 | 0.01584 | 0.13465 | 0.39736 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) | |
|------------------------|----------|------------|---------|----------|-----|
| (Intercept) | 1.5538 | 0.8060 | 1.928 | 0.065311 | . |
| fMOTILITYnonmotile | 0.6758 | 0.2645 | 2.554 | 0.017111 | * |
| fTIERINGinfaunal | -0.3910 | 0.2695 | -1.451 | 0.159156 | |
| fTIERINGpelagic | 1.3641 | 0.7013 | 1.945 | 0.063097 | . |
| fFEEDINGgrazing | 0.9163 | 0.6374 | 1.438 | 0.162935 | |
| fFEEDINGphotosymbiotic | 1.7089 | 0.7801 | 2.191 | 0.038023 | * |
| fFEEDINGpredatory | 0.2133 | 0.7339 | 0.291 | 0.773756 | |
| fFEEDINGSuspension | -0.1212 | 0.4878 | -0.248 | 0.805786 | |
| fLATITUDEP | -1.2973 | 0.5271 | -2.461 | 0.021108 | * |
| fLATITUDET | -0.2105 | 0.2368 | -0.889 | 0.382505 | |
| fBASINPanthalassa | -2.0205 | 0.5022 | -4.023 | 0.000467 | *** |
| fBASINTethys | -1.8369 | 0.5214 | -3.523 | 0.001667 | ** |
| fCALCIFICATIONlight | -1.5820 | 0.7751 | -2.041 | 0.051937 | . |
| fCALCIFICATIONmoderate | 0.2521 | 0.3502 | 0.720 | 0.478244 | |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.08575427)

Null deviance: 8.2940 on 38 degrees of freedom
Residual deviance: 2.3843 on 25 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

```
>  
> # find best model  
>  
> drop1(GLM_OAE_Blii, test = "F")
```

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) | |
|----------------|----|----------|---------|----------|----|
| <none> | | 2.3843 | | | |
| fMOTILITY | 1 | 2.9536 | 5.9686 | 0.021967 | * |
| fTIERING | 2 | 2.8716 | 2.5548 | 0.097824 | . |
| fFEEDING | 4 | 3.8265 | 3.7802 | 0.015446 | * |
| fLATITUDE | 2 | 2.9693 | 3.0667 | 0.064403 | . |
| fBASIN | 2 | 4.0320 | 8.6382 | 0.001406 | ** |
| fCALCIFICATION | 2 | 3.0072 | 3.2653 | 0.054965 | . |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
>  
> # motility, feeding, basin and calcification  
>  
> GLM_OAE_Bliii <- glm(extinction ~ fMOTILITY + fFEEDING +  
+ fBASIN + fCALCIFICATION,  
+ family = quasibinomial, data = OAE)  
> summary(GLM_OAE_Bliii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fFEEDING + fBASIN + fCALCIFICATION,  
family = quasibinomial, data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.05867 | -0.06024 | 0.06304 | 0.17024 | 0.47845 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|------------|
| (Intercept) | 0.7819 | 0.7250 | 1.078 | 0.28975 |
| fMOTILITYnonmotile | 0.5395 | 0.2818 | 1.914 | 0.06551 . |
| fFEEDINGgrazing | 1.0024 | 0.6641 | 1.509 | 0.14200 |
| fFEEDINGphotosymbiotic | 1.9016 | 0.8129 | 2.339 | 0.02641 * |
| fFEEDINGpredatory | 1.3548 | 0.5620 | 2.411 | 0.02249 * |
| fFEEDINGSuspension | -0.1287 | 0.5147 | -0.250 | 0.80428 |
| fBASINPanthalassa | -1.4785 | 0.4084 | -3.620 | 0.00111 ** |
| fBASINTethys | -1.2531 | 0.4027 | -3.112 | 0.00415 ** |
| fCALCIFICATIONlight | -1.1713 | 0.7957 | -1.472 | 0.15178 |
| fCALCIFICATIONmoderate | 0.2469 | 0.3424 | 0.721 | 0.47666 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1015277)

Null deviance: 8.294 on 38 degrees of freedom
Residual deviance: 3.348 on 29 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

>

> drop1(GLM_OAE_Bliii, test = "F")

Single term deletions

Model:

extinction ~ fMOTILITY + fFEEDING + fBASIN + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|---------------|
| <none> | | 3.3480 | | |
| fMOTILITY | 1 | 3.7264 | 3.2771 | 0.0806257 . |
| fFEEDING | 4 | 6.9142 | 7.7223 | 0.0002299 *** |
| fBASIN | 2 | 4.8737 | 6.6073 | 0.0043207 ** |
| fCALCIFICATION | 2 | 3.7888 | 1.9088 | 0.1664198 |

Signif. codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

>

> # best model

> GLM_OAE_Bliv <- glm(extinction ~ fFEEDING + fBASIN,

+ family = quasibinomial, data = OAE)

> summary(GLM_OAE_Bliv)

Call:

glm(formula = extinction ~ fFEEDING + fBASIN, family = quasibinomial,
data = OAE)

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.06388 | -0.16613 | 0.02477 | 0.21597 | 0.51343 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|------------|
| (Intercept) | 0.9920 | 0.6327 | 1.568 | 0.12674 |
| fFEEDINGgrazing | 1.0024 | 0.7081 | 1.416 | 0.16656 |
| fFEEDINGphotosymbiotic | 2.1544 | 0.7617 | 2.828 | 0.00801 ** |
| fFEEDINGpredatory | 1.0702 | 0.5717 | 1.872 | 0.07039 . |
| fFEEDINGSuspension | 0.0665 | 0.5207 | 0.128 | 0.89918 |
| fBASINPanthalassa | -1.3316 | 0.4153 | -3.206 | 0.00305 ** |
| fBASINTethys | -1.2164 | 0.4081 | -2.981 | 0.00546 ** |

Signif. codes: 0 '****' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1154421)

Null deviance: 8.2940 on 38 degrees of freedom

Residual deviance: 4.1213 on 32 degrees of freedom

AIC: NA

Number of Fisher Scoring iterations: 4

Model = extrinsic

> GLM_OAE_A3ii <- glm(extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,

+ family = quasibinomial, data = OAE)

> summary(GLM_OAE_A3ii)

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
     family = quasibinomial, data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -0.82785 | -0.19980 | 0.02194 | 0.30345 | 0.68123 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|----------|------------|---------|----------|
| (Intercept) | 0.4679 | 0.5296 | 0.884 | 0.388 |
| fLATITUDE | -0.3157 | 0.4468 | -0.707 | 0.488 |
| fBASINTethys | 0.8458 | 0.5224 | 1.619 | 0.122 |
| fENVIRONMENTOffshore | -0.4707 | 0.4937 | -0.953 | 0.352 |
| fENVIRONMENTReef | 2.0464 | 1.9653 | 1.041 | 0.311 |

(Dispersion parameter for quasibinomial family taken to be 0.1625729)

Null deviance: 4.1787 on 23 degrees of freedom
Residual deviance: 3.2273 on 19 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 5

```
>
> # find best model
>
> drop1(GLM_OAE_A3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN + fENVIRONMENT
```

| | Df | Deviance | F value | Pr(>F) |
|--------------|----|----------|---------|--------|
| <none> | | 3.2273 | | |
| fLATITUDE | 1 | 3.3089 | 0.4802 | 0.4967 |
| fBASIN | 1 | 3.6534 | 2.5082 | 0.1298 |
| fENVIRONMENT | 2 | 3.7659 | 1.5854 | 0.2308 |

Model = extrinsic no env

```
> GLM_OAE_B3ii <- glm(extinction ~ fLATITUDE + fBASIN,
+ family = quasibinomial, data = OAE)
> summary(GLM_OAE_B3ii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN, family = quasibinomial,  
     data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|----------|---------|---------|
| -0.85850 | -0.31552 | -0.02386 | 0.29006 | 0.88380 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|-------------------|----------|------------|---------|----------|
| (Intercept) | 1.76124 | 0.67687 | 2.602 | 0.0136 * |
| fLATITUDEP | -0.83310 | 0.70090 | -1.189 | 0.2428 |
| fLATITUDET | 0.08532 | 0.30341 | 0.281 | 0.7803 |
| fBASINPanthalassa | -1.73653 | 0.69143 | -2.512 | 0.0169 * |
| fBASINTethys | -1.53097 | 0.70095 | -2.184 | 0.0359 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1816417)

Null deviance: 8.2940 on 38 degrees of freedom
Residual deviance: 6.8489 on 34 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_OAE_B3ii, test = "F")

Single term deletions

Model:

extinction ~ fLATITUDE + fBASIN

| | Df | Deviance | F value | Pr(>F) |
|-----------|----|----------|---------|-----------|
| <none> | | 6.8489 | | |
| fLATITUDE | 2 | 7.1629 | 0.7795 | 0.46668 |
| fBASIN | 2 | 8.2454 | 3.4663 | 0.04265 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

>
> # best model
>
> GLM_OAE_B3iii <- glm(extinction ~ fBASIN,
+ family = quasibinomial, data = OAE)
> summary(GLM_OAE_B3iii)

```

Call:

```
glm(formula = extinction ~ fBASIN, family = quasibinomial, data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|----------|---------|---------|
| -1.18153 | -0.25543 | -0.03494 | 0.30631 | 0.86954 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) | |
|-------------------|----------|------------|---------|----------|----|
| (Intercept) | 1.2160 | 0.4466 | 2.723 | 0.00991 | ** |
| fBASINPanthalassa | -1.2064 | 0.5035 | -2.396 | 0.02190 | * |
| fBASINTethys | -0.9534 | 0.4832 | -1.973 | 0.05621 | . |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.175851)

Null deviance: 8.2940 on 38 degrees of freedom
Residual deviance: 7.1629 on 36 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

Model = intrinsic

```

> GLM_OAE_A4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fCALCIFICATION,
+ family = quasibinomial, data = OAE)
> summary(GLM_OAE_A4ii)

```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
    fCALCIFICATION, family = quasibinomial, data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|----------|---------|---------|
| -0.50167 | -0.26405 | -0.00598 | 0.31718 | 0.51395 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|-----------|------------|---------|----------|
| (Intercept) | 0.865721 | 0.835866 | 1.036 | 0.317 |
| fMOTILITYnonmotile | 0.586772 | 0.546392 | 1.074 | 0.300 |
| fTIERINGinfaunal | -0.408294 | 0.492615 | -0.829 | 0.420 |
| fTIERINGpelagic | 0.792790 | 1.106530 | 0.716 | 0.485 |
| fFEEDINGgrazing | 0.718156 | 0.810390 | 0.886 | 0.390 |
| fFEEDINGphotosymbiotic | 0.001459 | 1.107503 | 0.001 | 0.999 |
| fFEEDINGpredatory | 0.570980 | 1.078225 | 0.530 | 0.604 |
| fFEEDINGSuspension | -0.281368 | 0.685727 | -0.410 | 0.687 |
| fCALCIFICATIONmoderate | -0.338089 | 0.645988 | -0.523 | 0.608 |

(Dispersion parameter for quasibinomial family taken to be 0.1482437)

Null deviance: 4.1787 on 23 degrees of freedom
 Residual deviance: 2.3502 on 15 degrees of freedom
 AIC: NA

Number of Fisher Scoring iterations: 4

```
>
> # find best model
>
> drop1(GLM_OAE_A4ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 2.3502 | | |
| fMOTILITY | 1 | 2.5215 | 1.0938 | 0.3122 |
| fTIERING | 2 | 2.5248 | 0.5572 | 0.5842 |
| fFEEDING | 4 | 2.6545 | 0.4857 | 0.7462 |
| fCALCIFICATION | 1 | 2.3916 | 0.2642 | 0.6148 |

Model = intrinsic no env

```
> GLM_OAE_B4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
```

```
+ fCALCIFICATION,  
+ family = quasibinomial, data = OAE)  
> summary(GLM_OAE_B4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
     fCALCIFICATION, family = quasibinomial, data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.09670 | -0.21032 | 0.05265 | 0.18855 | 0.54955 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | -0.60947 | 0.64645 | -0.943 | 0.3536 |
| fMOTILITYnonmotile | 0.32053 | 0.30686 | 1.045 | 0.3049 |
| fTIERINGinfaunal | -0.45118 | 0.31438 | -1.435 | 0.1619 |
| fTIERINGpelagic | 1.28321 | 0.84016 | 1.527 | 0.1375 |
| fFEEDINGgrazing | 0.77966 | 0.77486 | 1.006 | 0.3226 |
| fFEEDINGphotosymbiotic | 2.17951 | 0.91122 | 2.392 | 0.0235 * |
| fFEEDINGpredatory | 0.18390 | 0.90933 | 0.202 | 0.8411 |
| fFEEDINGSuspension | 0.09612 | 0.56283 | 0.171 | 0.8656 |
| fCALCIFICATIONlight | -1.14531 | 0.93795 | -1.221 | 0.2319 |
| fCALCIFICATIONmoderate | 0.60790 | 0.40898 | 1.486 | 0.1480 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1320506)

Null deviance: 8.2940 on 38 degrees of freedom
Residual deviance: 4.3038 on 29 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

```
>  
> # find best model  
>  
> drop1(GLM_OAE_B4ii, test = "F")  
Single term deletions
```

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|-----------|
| <none> | | 4.3038 | | |
| fMOTILITY | 1 | 4.4486 | 0.9758 | 0.33142 |
| fTIERING | 2 | 4.8737 | 1.9200 | 0.16479 |
| fFEEDING | 4 | 5.9884 | 2.8378 | 0.04223 * |
| fCALCIFICATION | 2 | 5.1747 | 2.9342 | 0.06910 . |

Signif. codes: 0 '****' 0.001 '***' 0.01 '**' 0.05 '.' 0.1 ' ' 1

>

> # best model

>

```
> GLM_OAE_B4iv <- glm(extinction ~ fFEEDING,
```

```
+ family = quasibinomial, data = OAE)
```

```
> summary(GLM_OAE_B4iv)
```

Call:

```
glm(formula = extinction ~ fFEEDING, family = quasibinomial,  
     data = OAE)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.16779 | -0.21337 | 0.07018 | 0.28404 | 0.70741 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | -0.2243 | 0.5481 | -0.409 | 0.6849 |
| fFEEDINGgrazing | 1.0024 | 0.8026 | 1.249 | 0.2202 |
| fFEEDINGphotosymbiotic | 2.1149 | 0.8568 | 2.468 | 0.0188 * |
| fFEEDINGpredatory | 1.1659 | 0.6367 | 1.831 | 0.0758 . |
| fFEEDINGSuspension | 0.2016 | 0.5693 | 0.354 | 0.7254 |

Signif. codes: 0 '****' 0.001 '***' 0.01 '**' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.1483219)

Null deviance: 8.294 on 38 degrees of freedom

Residual deviance: 5.546 on 34 degrees of freedom

AIC: NA

Number of Fisher Scoring iterations: 4

Triassic background

Model = all

```
> GLM_pre_Alii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fLATITUDE + fBASIN + fENVIRONMENT +
+ fCALCIFICATION,
+ family = quasibinomial, data = pre)
> summary(GLM_pre_Alii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
    fLATITUDE + fBASIN + fENVIRONMENT + fCALCIFICATION, family =
quasibinomial,
    data = pre)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -1.0239 | -0.2825 | 0.0781 | 0.3665 | 0.7342 |

Coefficients: (1 not defined because of singularities)

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|-----------|------------|---------|----------|
| (Intercept) | 2.307822 | 1.126560 | 2.049 | 0.0479 * |
| fMOTILITYnonmotile | -0.001491 | 0.475700 | -0.003 | 0.9975 |
| fTIERINGinfaunal | -0.205156 | 0.553408 | -0.371 | 0.7130 |
| fTIERINGpelagic | 0.611297 | 0.891845 | 0.685 | 0.4975 |
| fFEEDINGphotosymbiotic | -1.734707 | 1.161870 | -1.493 | 0.1441 |
| fFEEDINGpredatory | NA | NA | NA | NA |
| fFEEDINGSuspension | -0.609533 | 0.809037 | -0.753 | 0.4561 |
| fLATITUDET | -0.369263 | 0.552370 | -0.669 | 0.5081 |
| fBASINTethys | -0.029463 | 0.632788 | -0.047 | 0.9631 |
| fENVIRONMENTOffshore | 0.146258 | 0.821322 | 0.178 | 0.8597 |
| fENVIRONMENTReef | -0.438626 | 0.491499 | -0.892 | 0.3781 |
| fCALCIFICATIONlight | -0.778322 | 0.724403 | -1.074 | 0.2898 |
| fCALCIFICATIONmoderate | -0.821088 | 0.571830 | -1.436 | 0.1597 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2401432)

Null deviance: 13.8327 on 47 degrees of freedom
Residual deviance: 9.1254 on 36 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_pre_Alii, test = "F")

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
fENVIRONMENT + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 9.1254 | | |
| fMOTILITY | 1 | 9.1254 | 0.0000 | 0.9976 |
| fTIERING | 1 | 9.1584 | 0.1302 | 0.7204 |
| fFEEDING | 2 | 9.7804 | 1.2920 | 0.2872 |
| fLATITUDE | 1 | 9.2353 | 0.4334 | 0.5145 |
| fBASIN | 1 | 9.1260 | 0.0021 | 0.9641 |
| fENVIRONMENT | 2 | 9.3238 | 0.3912 | 0.6791 |
| fCALCIFICATION | 2 | 9.6563 | 1.0471 | 0.3614 |

>

Model = all no env

```
> GLM_pre_Blii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +  
+ fLATITUDE + fBASIN + fCALCIFICATION,  
+ family = quasibinomial, data = pre)  
> summary(GLM_pre_Blii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
fLATITUDE + fBASIN + fCALCIFICATION, family = quasibinomial,  
data = pre)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.30649 | -0.24242 | 0.03913 | 0.31504 | 1.06498 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|--------------------|----------|------------|---------|----------|
| (Intercept) | 0.493783 | 0.609139 | 0.811 | 0.420 |
| fMOTILITYnonmotile | 0.132692 | 0.255165 | 0.520 | 0.604 |

| | | | | |
|------------------------|-----------|----------|--------|-------|
| fTIERINGinfaunal | -0.249179 | 0.288659 | -0.863 | 0.390 |
| fTIERINGpelagic | 0.422510 | 0.386715 | 1.093 | 0.278 |
| fFEEDINGgrazing | -0.221395 | 0.879548 | -0.252 | 0.802 |
| fFEEDINGother | -0.773554 | 1.111817 | -0.696 | 0.488 |
| fFEEDINGphotosymbiotic | -0.867489 | 0.713520 | -1.216 | 0.227 |
| fFEEDINGpredatory | 0.415223 | 0.689365 | 0.602 | 0.549 |
| fFEEDINGSuspension | -0.501679 | 0.610900 | -0.821 | 0.414 |
| fLATITUDEP | -0.002709 | 0.397414 | -0.007 | 0.995 |
| fLATITUDET | -0.245379 | 0.223035 | -1.100 | 0.274 |
| fBASINPanthalassa | 0.438921 | 0.406209 | 1.081 | 0.283 |
| fBASINTethys | 0.129194 | 0.440649 | 0.293 | 0.770 |
| fCALCIFICATIONlight | -0.275846 | 0.441214 | -0.625 | 0.533 |
| fCALCIFICATIONmoderate | -0.048012 | 0.292650 | -0.164 | 0.870 |

(Dispersion parameter for quasibinomial family taken to be 0.1870975)

Null deviance: 24.442 on 100 degrees of freedom
 Residual deviance: 17.897 on 86 degrees of freedom
 AIC: NA

Number of Fisher Scoring iterations: 4

```
> # find best model
>
> drop1(GLM_pre_Blii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
  fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 17.897 | | |
| fMOTILITY | 1 | 17.948 | 0.2431 | 0.6232 |
| fTIERING | 2 | 18.279 | 0.9181 | 0.4032 |
| fFEEDING | 5 | 19.132 | 1.1867 | 0.3224 |
| fLATITUDE | 2 | 18.141 | 0.5861 | 0.5587 |
| fBASIN | 2 | 18.465 | 1.3633 | 0.2613 |
| fCALCIFICATION | 2 | 17.979 | 0.1956 | 0.8227 |

>

Model = extrinsic

```
> GLM_pre_A3ii <- glm(extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
+ family = quasibinomial, data = pre)
> summary(GLM_pre_A3ii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
     family = quasibinomial, data = pre)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.21401 | -0.28266 | 0.08969 | 0.32510 | 0.83503 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|----------|------------|---------|----------|
| (Intercept) | 0.9568 | 0.5688 | 1.682 | 0.0998 . |
| fLATITUDE | 0.1052 | 0.5020 | 0.210 | 0.8350 |
| fBASINTethys | -0.1877 | 0.5423 | -0.346 | 0.7309 |
| fENVIRONMENTOffshore | 1.0662 | 0.6413 | 1.663 | 0.1036 |
| fENVIRONMENTReef | -0.7343 | 0.3295 | -2.229 | 0.0311 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2421453)

Null deviance: 13.833 on 47 degrees of freedom
Residual deviance: 11.197 on 43 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

```
>
> # find best model
>
> drop1(GLM_pre_A3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN + fENVIRONMENT
          Df Deviance F value Pr(>F)
<none>          11.197
fLATITUDE     1  11.207  0.0405 0.84141
```

```

fBASIN      1  11.226  0.1132 0.73814
fENVIRONMENT 2  13.831  5.0590 0.01064 *
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
>
> # best model
>
> GLM_pre_A3iii <- glm(extinction ~ fENVIRONMENT,
+ family = quasibinomial, data = pre)
> summary(GLM_pre_A3iii)

```

```

Call:
glm(formula = extinction ~ fENVIRONMENT, family = quasibinomial,
     data = pre)

```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -1.2179 | -0.2782 | 0.1173 | 0.3223 | 0.8322 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|----------|------------|---------|--------------|
| (Intercept) | 0.8824 | 0.2038 | 4.330 | 8.24e-05 *** |
| fENVIRONMENTOffshore | 1.0400 | 0.6238 | 1.667 | 0.102 |
| fENVIRONMENTReef | -0.7314 | 0.3222 | -2.270 | 0.028 * |

```

---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

(Dispersion parameter for quasibinomial family taken to be 0.232193)

```

Null deviance: 13.833 on 47 degrees of freedom
Residual deviance: 11.228 on 45 degrees of freedom
AIC: NA

```

Number of Fisher Scoring iterations: 4

Model = extrinsic no env

```

> GLM_pre_B3ii <- glm(extinction ~ fLATITUDE + fBASIN,
+ family = quasibinomial, data = pre)
> summary(GLM_pre_B3ii)

```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN, family = quasibinomial,
     data = pre)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.29562 | -0.30561 | 0.02562 | 0.39844 | 1.02334 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|-------------------|----------|------------|---------|----------|
| (Intercept) | 0.35325 | 0.39494 | 0.894 | 0.373 |
| fLATITUDEP | -0.07958 | 0.38308 | -0.208 | 0.836 |
| fLATITUDET | -0.24219 | 0.22525 | -1.075 | 0.285 |
| fBASINPanthalassa | 0.26273 | 0.39198 | 0.670 | 0.504 |
| fBASINTethys | 0.02883 | 0.42502 | 0.068 | 0.946 |

(Dispersion parameter for quasibinomial family taken to be 0.2223776)

Null deviance: 24.442 on 100 degrees of freedom
Residual deviance: 23.767 on 96 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 3

```
>
> # find best model
>
> drop1(GLM_pre_B3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN
      Df Deviance F value Pr(>F)
<none>      23.767
fLATITUDE  2   24.025  0.5220 0.5950
fBASIN     2   24.093  0.6592 0.5196
>
```

Model = intrinsic

```
> GLM_pre_A4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fCALCIFICATION,
+ family = quasibinomial, data = pre)
```

```
> summary(GLM_pre_A4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
     fCALCIFICATION, family = quasibinomial, data = pre)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|---------|---------|--------|--------|--------|
| -0.9961 | -0.2672 | 0.1008 | 0.3637 | 0.6452 |

Coefficients: (1 not defined because of singularities)

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 1.85156 | 0.84290 | 2.197 | 0.0339 * |
| fMOTILITYnonmotile | -0.06775 | 0.45827 | -0.148 | 0.8832 |
| fTIERINGinfaunal | -0.02356 | 0.47980 | -0.049 | 0.9611 |
| fTIERINGpelagic | 0.78405 | 0.78742 | 0.996 | 0.3254 |
| fFEEDINGphotosymbiotic | -1.88930 | 0.97094 | -1.946 | 0.0587 . |
| fFEEDINGpredatory | NA | NA | NA | NA |
| fFEEDINGSuspension | -0.71681 | 0.74704 | -0.960 | 0.3431 |
| fCALCIFICATIONlight | -0.88807 | 0.69239 | -1.283 | 0.2070 |
| fCALCIFICATIONmoderate | -0.76356 | 0.55221 | -1.383 | 0.1744 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2293279)

Null deviance: 13.8327 on 47 degrees of freedom
Residual deviance: 9.5384 on 40 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

```
>
```

```
> # find best model
```

```
>
```

```
> drop1(GLM_pre_A4ii, test = "F")
```

Single term deletions

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION  
Df Deviance F value Pr(>F)
```

```

<none>          9.5384
fMOTILITY       1   9.5434  0.0210 0.8854
fTIERING        1   9.5389  0.0023 0.9618
fFEEDING        2  10.6146  2.2568 0.1179
fCALCIFICATION  2  10.0601  1.0941 0.3447
>

```

Model = intrinsic no env

```

> GLM_pre_B4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fCALCIFICATION,
+ family = quasibinomial, data = pre)
> summary(GLM_pre_B4ii)

```

Call:

```

glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fCALCIFICATION, family = quasibinomial, data = pre)

```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.43602 | -0.23872 | 0.07058 | 0.28717 | 1.09648 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 0.55518 | 0.58195 | 0.954 | 0.343 |
| fMOTILITYnonmotile | 0.11194 | 0.25563 | 0.438 | 0.663 |
| fTIERINGinfaunal | -0.26538 | 0.27564 | -0.963 | 0.338 |
| fTIERINGpelagic | 0.44883 | 0.38590 | 1.163 | 0.248 |
| fFEEDINGgrazing | -0.38720 | 0.80757 | -0.479 | 0.633 |
| fFEEDINGother | -0.92316 | 1.07130 | -0.862 | 0.391 |
| fFEEDINGphotosymbiotic | -0.77628 | 0.66080 | -1.175 | 0.243 |
| fFEEDINGpredatory | 0.44291 | 0.63581 | 0.697 | 0.488 |
| fFEEDINGSuspension | -0.41397 | 0.54340 | -0.762 | 0.448 |
| fCALCIFICATIONlight | -0.42882 | 0.42755 | -1.003 | 0.319 |
| fCALCIFICATIONmoderate | -0.05979 | 0.28712 | -0.208 | 0.836 |

(Dispersion parameter for quasibinomial family taken to be 0.1891681)

```

Null deviance: 24.442 on 100 degrees of freedom
Residual deviance: 18.780 on 90 degrees of freedom
AIC: NA

```

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_pre_B4ii, test = "F")

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 18.780 | | |
| fMOTILITY | 1 | 18.816 | 0.1738 | 0.6777 |
| fTIERING | 2 | 19.212 | 1.0348 | 0.3595 |
| fFEEDING | 5 | 19.953 | 1.1242 | 0.3533 |
| fCALCIFICATION | 2 | 19.003 | 0.5347 | 0.5877 |

>

Jurassic background

Model = all

```
> GLM_post_Alii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fLATITUDE + fBASIN + fENVIRONMENT +
+ fCALCIFICATION,
+ family = quasibinomial, data = post)
> summary(GLM_post_Alii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fLATITUDE + fBASIN + fENVIRONMENT + fCALCIFICATION, family =
quasibinomial,
     data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.51234 | -0.21406 | 0.08584 | 0.31578 | 0.87603 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 0.04533 | 1.14542 | 0.040 | 0.9686 |
| fMOTILITYnonmotile | 0.44139 | 0.48417 | 0.912 | 0.3668 |
| fTIERINGinfaunal | 0.09458 | 0.44038 | 0.215 | 0.8309 |
| fTIERINGpelagic | 1.93865 | 0.90292 | 2.147 | 0.0372 * |
| fFEEDINGgrazing | -0.13636 | 0.94827 | -0.144 | 0.8863 |
| fFEEDINGphotosymbiotic | -1.08787 | 1.29808 | -0.838 | 0.4064 |
| fFEEDINGpredatory | -0.07292 | 0.99946 | -0.073 | 0.9422 |
| fFEEDINGSuspension | -0.81351 | 0.88483 | -0.919 | 0.3628 |
| fLATITUDET | -0.61547 | 0.42484 | -1.449 | 0.1544 |
| fBASINTethys | 1.08666 | 0.50557 | 2.149 | 0.0370 * |
| fENVIRONMENTOffshore | -1.20987 | 0.45935 | -2.634 | 0.0115 * |
| fENVIRONMENTReef | -1.66180 | 1.39474 | -1.191 | 0.2397 |
| fCALCIFICATIONmoderate | 0.15080 | 0.53518 | 0.282 | 0.7794 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2844998)

Null deviance: 23.615 on 57 degrees of freedom
Residual deviance: 15.391 on 45 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

```
> # find best model
>
> drop1(GLM_post_Alii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
  fENVIRONMENT + fCALCIFICATION
```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|-----------|
| <none> | | 15.391 | | |
| fMOTILITY | 1 | 15.629 | 0.6965 | 0.40836 |
| fTIERING | 2 | 16.803 | 2.0641 | 0.13878 |
| fFEEDING | 4 | 15.811 | 0.3069 | 0.87182 |
| fLATITUDE | 1 | 15.991 | 1.7543 | 0.19203 |
| fBASIN | 1 | 16.748 | 3.9680 | 0.05246 . |
| fENVIRONMENT | 2 | 18.316 | 4.2762 | 0.01994 * |
| fCALCIFICATION | 1 | 15.414 | 0.0661 | 0.79830 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```
>
> # best model
> GLM_post_Aliiii <- glm(extinction ~ fBASIN + fENVIRONMENT,
+ family = quasibinomial, data = post)
>
> summary(GLM_post_Aliiii)
```

Call:

```
glm(formula = extinction ~ fBASIN + fENVIRONMENT, family = quasibinomial,
     data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.39697 | -0.23517 | 0.01493 | 0.36667 | 1.15096 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|----------|------------|---------|------------|
| (Intercept) | -0.7316 | 0.3532 | -2.072 | 0.04309 * |
| fBASINTethys | 1.2343 | 0.4345 | 2.841 | 0.00633 ** |
| fENVIRONMENTOffshore | -0.4401 | 0.3350 | -1.314 | 0.19450 |

```
fENVIRONMENTreef      -2.2945      1.1370    -2.018    0.04856 *
```

```
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1
```

(Dispersion parameter for quasibinomial family taken to be 0.3009023)

Null deviance: 23.615 on 57 degrees of freedom

Residual deviance: 19.858 on 54 degrees of freedom

AIC: NA

Number of Fisher Scoring iterations: 4

Model = all no env

```
> GLM_post_Blii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fLATITUDE + fBASIN + fCALCIFICATION,
+ family = quasibinomial, data = post)
> summary(GLM_post_Blii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
     fLATITUDE + fBASIN + fCALCIFICATION, family = quasibinomial,
     data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.17245 | -0.30500 | 0.02441 | 0.33190 | 0.97939 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|-----------|------------|---------|----------|
| (Intercept) | -0.652952 | 0.840734 | -0.777 | 0.4392 |
| fMOTILITYnonmotile | 0.022546 | 0.265609 | 0.085 | 0.9325 |
| fTIERINGinfaunal | 0.013179 | 0.276256 | 0.048 | 0.9620 |
| fTIERINGpelagic | 0.953836 | 0.574465 | 1.660 | 0.1000 . |
| fFEEDINGgrazing | 1.153372 | 0.628432 | 1.835 | 0.0694 . |
| fFEEDINGother | 1.335053 | 1.178406 | 1.133 | 0.2600 |
| fFEEDINGphotosymbiotic | 0.625865 | 0.691337 | 0.905 | 0.3675 |
| fFEEDINGpredatory | 1.241876 | 0.690106 | 1.800 | 0.0749 . |
| fFEEDINGSuspension | 0.264316 | 0.497767 | 0.531 | 0.5966 |
| fLATITUDEP | 0.180565 | 0.518066 | 0.349 | 0.7282 |
| fLATITUDET | -0.004722 | 0.228922 | -0.021 | 0.9836 |
| fBASINPanthalassa | -0.462281 | 0.516853 | -0.894 | 0.3732 |

```

fBASINTethys          -0.002421   0.539806  -0.004   0.9964
fCALCIFICATIONlight  -0.577693   0.674157  -0.857   0.3935
fCALCIFICATIONmoderate 0.005010   0.354818   0.014   0.9888
---
Signif. codes:  0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

```

(Dispersion parameter for quasibinomial family taken to be 0.251139)

```

Null deviance: 42.295 on 114 degrees of freedom
Residual deviance: 30.418 on 100 degrees of freedom
AIC: NA

```

Number of Fisher Scoring iterations: 4

```

>
> # find best model
>
> drop1(GLM_post_Blii, test = "F")
Single term deletions

```

Model:

```

extinction ~ fMOTILITY + fTIERING + fFEEDING + fLATITUDE + fBASIN +
  fCALCIFICATION

```

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 30.418 | | |
| fMOTILITY | 1 | 30.420 | 0.0059 | 0.9387 |
| fTIERING | 2 | 31.094 | 1.1106 | 0.3334 |
| fFEEDING | 5 | 32.449 | 1.3355 | 0.2555 |
| fLATITUDE | 2 | 30.449 | 0.0512 | 0.9501 |
| fBASIN | 2 | 31.475 | 1.7371 | 0.1813 |
| fCALCIFICATION | 2 | 30.660 | 0.3979 | 0.6728 |

Model = extrinsic

```

> GLM_post_A3ii <- glm(extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
+ family = quasibinomial, data = post)
> summary(GLM_post_A3ii)

```

Call:

```

glm(formula = extinction ~ fLATITUDE + fBASIN + fENVIRONMENT,
    family = quasibinomial, data = post)

```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.42597 | -0.23507 | 0.02656 | 0.37338 | 1.14446 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|----------------------|----------|------------|---------|----------|
| (Intercept) | -0.6267 | 0.4674 | -1.341 | 0.1857 |
| fLATITUDE | -0.1288 | 0.3728 | -0.345 | 0.7312 |
| fBASINTethys | 1.1943 | 0.4527 | 2.638 | 0.0109 * |
| fENVIRONMENTOffshore | -0.4896 | 0.3676 | -1.332 | 0.1886 |
| fENVIRONMENTReef | -2.2306 | 1.1610 | -1.921 | 0.0601 . |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.3059362)

Null deviance: 23.615 on 57 degrees of freedom
Residual deviance: 19.822 on 53 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_post_A3ii, test = "F")

Single term deletions

Model:

extinction ~ fLATITUDE + fBASIN + fENVIRONMENT

| | Df | Deviance | F value | Pr(>F) |
|--------------|----|----------|---------|-----------|
| <none> | | 19.822 | | |
| fLATITUDE | 1 | 19.858 | 0.0975 | 0.75609 |
| fBASIN | 1 | 22.058 | 5.9786 | 0.01784 * |
| fENVIRONMENT | 2 | 21.794 | 2.6368 | 0.08096 . |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

>

> # best model

> GLM_post_A3iii <- glm(extinction ~ fBASIN,

```
+ family = quasibinomial, data = post)
>
> summary(GLM_post_A3iii)
```

Call:

```
glm(formula = extinction ~ fBASIN, family = quasibinomial, data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.25583 | -0.36205 | 0.04317 | 0.45311 | 1.10095 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|--------------|----------|------------|---------|----------|
| (Intercept) | -0.7316 | 0.3614 | -2.024 | 0.0477 * |
| fBASINTethys | 0.9141 | 0.3971 | 2.302 | 0.0251 * |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.3151467)

Null deviance: 23.615 on 57 degrees of freedom
Residual deviance: 21.850 on 56 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 3

Model = extrinsic no env

```
> GLM_post_B3ii <- glm(extinction ~ fLATITUDE + fBASIN,  
+ family = quasibinomial, data = post)  
> summary(GLM_post_B3ii)
```

Call:

```
glm(formula = extinction ~ fLATITUDE + fBASIN, family = quasibinomial,  
data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.18921 | -0.41443 | 0.07206 | 0.36821 | 1.23843 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|--|----------|------------|---------|----------|
|--|----------|------------|---------|----------|

| | | | | |
|-------------------|----------|---------|--------|-------|
| (Intercept) | -0.15073 | 0.50156 | -0.301 | 0.764 |
| fLATITUDEP | 0.09206 | 0.51178 | 0.180 | 0.858 |
| fLATITUDET | 0.04691 | 0.22530 | 0.208 | 0.835 |
| fBASINPanthalassa | -0.35305 | 0.50998 | -0.692 | 0.490 |
| fBASINTethys | 0.13156 | 0.51975 | 0.253 | 0.801 |

(Dispersion parameter for quasibinomial family taken to be 0.3064222)

Null deviance: 42.295 on 114 degrees of freedom
 Residual deviance: 40.896 on 110 degrees of freedom
 AIC: NA

Number of Fisher Scoring iterations: 3

```
>
> # find best model
>
> drop1(GLM_post_B3ii, test = "F")
Single term deletions
```

Model:

```
extinction ~ fLATITUDE + fBASIN
          Df Deviance F value Pr(>F)
<none>      40.896
fLATITUDE  2  40.917  0.0279 0.9725
fBASIN     2  42.294  1.8799 0.1575
>
```

Model = intrinsic

```
> GLM_post_A4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +
+ fCALCIFICATION,
+ family = quasibinomial, data = post)
> summary(GLM_post_A4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +
    fCALCIFICATION, family = quasibinomial, data = post)
```

Deviance Residuals:

| | Min | 1Q | Median | 3Q | Max |
|--|----------|----------|---------|---------|---------|
| | -1.23841 | -0.35885 | 0.07215 | 0.42268 | 1.11761 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|----------|------------|---------|----------|
| (Intercept) | 0.48986 | 0.93482 | 0.524 | 0.6026 |
| fMOTILITYnonmotile | 0.02496 | 0.46913 | 0.053 | 0.9578 |
| fTIERINGinfaunal | -0.16568 | 0.44754 | -0.370 | 0.7128 |
| fTIERINGpelagic | 1.77230 | 0.92260 | 1.921 | 0.0606 |
| fFEEDINGgrazing | -0.15181 | 0.97205 | -0.156 | 0.8765 |
| fFEEDINGphotosymbiotic | -1.00033 | 1.12559 | -0.889 | 0.3785 |
| fFEEDINGpredatory | -0.20319 | 1.05564 | -0.192 | 0.8482 |
| fFEEDINGSuspension | -0.37252 | 0.87741 | -0.425 | 0.6730 |
| fCALCIFICATIONmoderate | -0.28667 | 0.51093 | -0.561 | 0.5773 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.3181259)

Null deviance: 23.615 on 57 degrees of freedom

Residual deviance: 19.476 on 49 degrees of freedom

AIC: NA

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_post_A4ii, test = "F")

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 19.476 | | |
| fMOTILITY | 1 | 19.477 | 0.0023 | 0.9622 |
| fTIERING | 2 | 20.768 | 1.6259 | 0.2072 |
| fFEEDING | 4 | 19.804 | 0.2067 | 0.9335 |
| fCALCIFICATION | 1 | 19.576 | 0.2526 | 0.6175 |

>

Model = intrinsic no env

> GLM_post_B4ii <- glm(extinction ~ fMOTILITY + fTIERING + fFEEDING +

```
+ fCALCIFICATION,  
+ family = quasibinomial, data = post)  
> summary(GLM_post_B4ii)
```

Call:

```
glm(formula = extinction ~ fMOTILITY + fTIERING + fFEEDING +  
     fCALCIFICATION, family = quasibinomial, data = post)
```

Deviance Residuals:

| Min | 1Q | Median | 3Q | Max |
|----------|----------|---------|---------|---------|
| -1.07893 | -0.38812 | 0.07983 | 0.34490 | 1.05144 |

Coefficients:

| | Estimate | Std. Error | t value | Pr(> t) |
|------------------------|-----------|------------|---------|----------|
| (Intercept) | -0.617533 | 0.530221 | -1.165 | 0.2468 |
| fMOTILITYnonmotile | -0.029025 | 0.255465 | -0.114 | 0.9098 |
| fTIERINGinfaunal | -0.042630 | 0.264686 | -0.161 | 0.8724 |
| fTIERINGpelagic | 0.810725 | 0.557628 | 1.454 | 0.1490 |
| fFEEDINGgrazing | 1.123169 | 0.615087 | 1.826 | 0.0707 . |
| fFEEDINGother | 1.358299 | 1.150768 | 1.180 | 0.2406 |
| fFEEDINGphotosymbiotic | 0.410438 | 0.637146 | 0.644 | 0.5209 |
| fFEEDINGpredatory | 1.157417 | 0.682559 | 1.696 | 0.0929 . |
| fFEEDINGSuspension | 0.123304 | 0.467416 | 0.264 | 0.7925 |
| fCALCIFICATIONlight | -0.452490 | 0.659257 | -0.686 | 0.4940 |
| fCALCIFICATIONmoderate | -0.004989 | 0.332452 | -0.015 | 0.9881 |

Signif. codes: 0 '***' 0.001 '**' 0.01 '*' 0.05 '.' 0.1 ' ' 1

(Dispersion parameter for quasibinomial family taken to be 0.2494198)

Null deviance: 42.295 on 114 degrees of freedom
Residual deviance: 31.582 on 104 degrees of freedom
AIC: NA

Number of Fisher Scoring iterations: 4

>

> # find best model

>

> drop1(GLM_post_B4ii, test = "F")

Single term deletions

Model:

extinction ~ fMOTILITY + fTIERING + fFEEDING + fCALCIFICATION

| | Df | Deviance | F value | Pr(>F) |
|----------------|----|----------|---------|--------|
| <none> | | 31.582 | | |
| fMOTILITY | 1 | 31.585 | 0.0106 | 0.9182 |
| fTIERING | 2 | 32.103 | 0.8577 | 0.4271 |
| fFEEDING | 5 | 33.904 | 1.5293 | 0.1871 |
| fCALCIFICATION | 2 | 31.732 | 0.2482 | 0.7806 |

>