

# Evolutionary paleoecology and the biology of extinction

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# 1 Introduction

2 Evolutionary paleoecology is the study of the effects of ecological traits and factors on  
3 differential rate dynamics, particularly rates of faunal turnover and diversification [73].  
4 Ecological traits are expressed by a taxon which are involved in biotic–biotic or biotic–abiotic  
5 interactions while ecological factors are the environmental conditions in which a taxon  
6 exists (the set of all biotic and abiotic interactors). Diversification is the difference between  
7 origination and extinction and is the net pattern of macroevolution. The study of evolutionary  
8 paleoecology is therefore the link between environmental (biotic–biotic and biotic–abiotic)  
9 interactions and macroevolution. As a corollary to Kitchell [73]’s definition, Allmon [2] states  
10 that in order to correctly link ecological interactions to macroevolution, one must focus on  
11 the specific traits and factors that may affect the speciation process. Tacitly included in this  
12 is the study of how ecological traits are related to extinction [72].

13 It is expected that for the majority of geological time, extinction is non-random with respect  
14 to biology [54]. Determining how different traits, both alone or in concert, influence extinction  
15 risk is then extremely important for understanding the differential diversification of taxa over  
16 the Phanerozoic. Periods of background extinction also represent the majority of geologic  
17 time, remain relatively predictable and change slowly, thus providing a better opportunity to  
18 study how traits are related to survival than periods of mass extinction [54, 106]. The Law of  
19 Constant extinction [132] posits that extinction risk of taxa within a given adaptive zone is  
20 age independent (memoryless), however the generality of this statement is possibly suspect  
21 [29, 36, 104, 115]. By analyzing survival patterns within adaptive zones during periods of  
22 background extinction, it should be possible to determine if extinction is best modeled as age  
23 independent or dependent.

24 Organismal traits relating to environmental preference are good candidates for estimating  
25 differences in extinction risk. A variety of organismal traits have been shown to be related  
26 to differential extinction [11, 41, 77, 90], especially with regards to the relationship between  
27 adaptation to variable environments and increased species longevity. A simple expectation  
28 is that taxa with preference for rare environments will be more at risk than taxa which  
29 prefer abundant environments based on purely stochastic grounds. As environments change  
30 in availability, a taxon’s instantaneous risk of extinction would then be expected change in  
31 concert. Taxa are also expected to be adapting to their environment, possibly increasing or  
32 decreasing their environmental tolerance and thus changing their instantaneous extinction risk.  
33 Related to environmental preference is species-level geographic range size. Species with larger  
34 geographic ranges tend to have lower extinction rates than species with smaller geographic  
35 ranges [48, 54, 60, 90, 113]. However, how range size is “formed” is different between clades  
36 [55] and thus remains a black box for most taxa. Thus, the utility of focusing on organismal  
37 traits related to environmental preference is that the black box can be “opened.”  
38 In addition to understanding patterns of survival, how community composition changes  
39 over time is extremely important for understanding how trophic structure changes or is  
40 maintained over time. Additionally, community connectedness is important for understanding

the degree to which global, regional, or local scale processes are important for shaping the environment, or the set of all possible biotic and abiotic interactors. In addition to total community connectedness, the dynamics of taxa within various ecological categories are important for understanding whether different adaptive conditions are differently affected by global, regional, or local scale processes. The Law of Constant extinction is theorized to hold during periods of environmental stability and is thus considered extremely difficult/impossible to test [79]. However, if environmental shifts are incorporated into the analysis of survival distributions, it may be possible to actually test the relationship between taxon age and extinction risk in the context of their adaptive zone and environment. Additionally, this may allow for illumination of what actual processes underly extinction during the majority of geologic time.

It is under this framework that I propose to study how ecological traits associated with environmental preference have affected both differential survival and cosmopolitan-endemism dynamics. I will be studying two distantly related and biotically different groups: Permian brachiopods and Cenozoic mammals. Both of these groups are considered to have very good fossil records able to reflect massive long term evolutionary patterns [84]. These two time periods were chosen because they represent periods of approximately the same length (47 My and 65 My) and of climatic change, global warming and global cooling respectively. Also, these two groups are a marine and terrestrial system respectively and the traits associated with environmental preference and range size (described below) are fundamentally very different. Both patterns of survival (Section 1.1) and community connectedness (Section 1.2) will be measured for both of these groups. The differences between these two groups in terms of life-habit and environmental preference, along with global climatic context, provides a fantastic scenario to understand how long-term, large-scale processes away from mass extinctions proceed.

## 1.1 Survival analysis

Survival analysis is the analysis of time till event data. In the case of paleontological analysis this is the time from the origination (first appearance date; FAD) of a taxon till the time of extinction (last appearance date; LAD), also known as the duration of a taxon. Survival analysis has a long history in paleontology [122, 132], though these initial analyses differ from modern approaches [74]. Survival is described primarily by two functions,  $S(t)$  and  $h(t)$ , or probability of survival at time (age)  $t$  and instantaneous failure rate at time  $t$  respectively. The survival function, and corresponding hazard function, can be parameterized in a variety of different ways. Survival curves can also be estimated non-parametrically using, for example, the Kaplan-Meier (K-M) estimator. This approach provides a useful method for graphically representing the observed survival distribution. While other methods, such as semiparametric Cox models [74], can be used to compare patterns of survival, only fully non-parametric and fully parametric methods are used here. Parametric techniques are favored because the shape of the hazard function is of interest.

- 80 Survival analysis shares some similarities with linear and logistic regression. While these  
 82 approaches use continuous (duration) or dichotomous (extinct/not extinct) responses, re-  
 spectively, survival analysis combines these concepts by measuring the duration till event  
 84 or follow-up time. In addition to using both duration and death/extinction information,  
 86 censoring information can also be incorporated in analysis. Censoring is the term for when  
 there is uncertainty of exact survival time. In a paleobiological context this would be when a  
 taxon either originated or went extinct outside of a period of interest, or the exact duration  
 is known only as a range and not a single value.
- 88 The Law of Constant extinction [132] predicts that extinction risk is random with respect  
 90 to taxon age. In the language of survival analysis, this means that  $h(t) = \lambda$  where  $\lambda$  is a  
 constant. This only occurs when survival times are exponentially distributed, formulated as

$$S(t) = \exp^{-\lambda t} \quad (1)$$

. Importantly, this observation underlies the “validity” of birth-death models of population  
 92 and clade dynamics where death risk is memoryless.

There are many alternatives to constant extinction risk, however. For example, there is a  
 94 frequently observed inverse relationship between genus age and extinction risk [36], which  
 could be modeled using, among many others, a Weibull distribution. In comparison to the  
 96 exponential distribution which has only a single parameter  $\lambda$ , the Weibull distribution has  
 two parameters,  $\lambda$  and  $k$ . The  $\lambda$  of the Weibull distribution behaves as in the exponential  
 98 distribution while  $k$  is a shape parameter which describes how failure is proportional to a  
 power of time. If  $k < 1$  then failure rate monotonically decreases with age, and if  $k > 1$   
 100 then failure rate monotonically increases with age. When  $k = 1$  the Weibull distribution  
 reduces to the exponential distribution and failure rate is constant. Other potential survival  
 102 distributions, such as the log-logistic, extended Gamma, and log-normal distributions [74],  
 all of which describe different patterns of age-dependent failure/extinction (monotonic and  
 104 nonmonotonic).

### 1.1.1 Effect of heterogeneous preservation

- 106 While some amount of uncertainty is possible to incorporate in a survival analysis framework  
 via censoring, this may be complicated when dealing with the fossil record. Because the  
 108 observed duration of a taxon is virtually always shorter than the actual duration of that  
 taxon, it is important to understand how this affects models of survival. If preservation is  
 110 homogeneous across taxa, space, and time, then this is expected to have a minimal and  
 uniform effect on estimates of duration and survival [39, 42, 115].
- 112 However, a major concern is the systematic differential preservation of one group of organisms  
 compared to another, for example between taxa inhabiting open versus closed environments  
 114 (Section 3.2). Any systematic bias in estimation of survival times would affect estimating  
 model parameters of  $S(t)$  and potentially lead to the wrong conclusions. In order to investigate

the effect of systematic range truncation on one set of observations compared to another, I propose four different sets of simulations: (1) two groups with identical diversification and identical preservation, (2) two groups with identical diversification but different preservation, (3) two groups with different diversification but identical preservation, and (4) two groups with different diversification and different preservation. Both diversification and preservation will be simulated using very simple models in order to limit the complexity of interpreting results.

Diversification will be simulated as a time-homogeneous birth-death process. This model has only two parameters, birth ( $\lambda$ ) and death ( $\mu$ ), which are the expected number of events per unit time. Both events are memoryless and thus have no association with an observations age. When death is random with respect to age, the expected hazard function is constant and survival is expected to be exponentially distributed (Eq. 1). This process was chosen because it is extremely simple, well understood, and is a common model used to model phylogenies [69, 88, 89, 107].

The simplest model of fossil preservation is a Poisson process [39, 42, 126, 127]. Following an exponential distribution, preservation rate ( $\phi$ ) is defined as the number of expected observations per unit time. For a given observation (taxa), a series waiting times are randomly drawn until the sum of these random times is equal to or greater than the duration of the observation. Each of these waiting times represents a fossil occurrence. The time between the first occurrence and the penultimate occurrence is then the duration of the “fossilized” observation.

Varying  $\lambda$ ,  $\mu$ , and  $\phi$  across a wide range of values, I will measure the deviance in median survival time between both the “true” and the “fossilized” distributions. Any systematic bias away from the range of deviances in median survival times of the “true” distributions will represent the effect of differential preservation for the simplest possible diversification and fossilization models. Importantly, this approach may illuminate situations where survival may be biased away from being exponentially distributed.

## 1.2 Biogeographic networks

Community connectedness is the degree to which localities are composed of endemic versus cosmopolitan taxa, and how similar this relationship is across localities. If localities and taxa are defined as nodes in a bipartite network, different network measures can be used to measure the how nodes are linked and describe both how taxa are distributed among and between localities and how similar different localities are. A bipartite network is defined as a network where nodes can be divided into two disjoint sets such that connections are only between sets and not within sets [28], in this case meaning that taxa can only be linked to localities but not to taxa and *vice versa*.

Modified from Sidor et al. [118], community connectedness can be measured via four different summary statistics: average relative number of endemic taxa per locality ( $E$ , Eq. 2), average

<sup>154</sup> relative locality occupancy per taxon ( $Occ$ , Eq. 3), biogeographic connectedness ( $BC$ , Eq. 4),  
 and code length [111, 112, 118]. These summary statistics describe how unique each locality  
<sup>156</sup> is on average compared to all others during a time period, how relatively widespread taxa are  
 on average during a time period, how evenly distributed taxa are amongst localities during a  
<sup>158</sup> time period, and the degree of biogeographic provincially during a time period. Of course, the  
 first three of these measures can be represented as distributions instead of means allowing for  
<sup>160</sup> better understanding of the spread of taxa over the landscape.  $E$  is defined as

$$E = \frac{\sum_{i=1}^L \frac{u_i}{n_i}}{L} \quad (2)$$

<sup>162</sup> where  $L$  is as the number of localities,  $u$  is the number of taxa unique to a locality, and  $n$   
 is the number of taxa present at a locality.  $Occ$  is the number of localities a taxon is, on  
 average, found at and is defined

$$Occ = \frac{\sum_{i=1}^N \frac{l_i}{L}}{N} \quad (3)$$

<sup>164</sup> where  $N$  is as the number of taxa present in the biogeographic network and  $l$  is the number  
 of localities a taxon occurred in.  $BC$  is defined

$$BC = \frac{O - N}{LN - N} \quad (4)$$

<sup>166</sup> where  $O$  is the total number of taxonomic occurrences.  $BC$  ranges from 0 to 1, with 0 meaning  
 that each locality completely disconnected from all other localities and 1 indicating that all  
<sup>168</sup> taxa are presents at all localities.

<sup>170</sup> Code length is a measure of the information flow [117] of a graph as estimated from the  
 behavior of a random surfer [16] on a graph and how often it visits each node and it's behavior  
<sup>172</sup> with regards to moving into and out of different regions of the graph. Code length is the  
 minimum length of a binary code for describing the behavior the surfer and the community  
<sup>174</sup> structure of the nodes and is estimated via the map equation [111, 112]. The logic of the  
 map equation is that a good map compresses reality into as few symbols as possible thus we  
<sup>176</sup> want to compress as many nodes as possible into as few symbols as possible. The goal is to  
 compress a graph better than just assigning a unique Huffman code to each node [52, 111]. By  
<sup>178</sup> compressing multiple nodes into a single code block, we decrease the minimum coding length  
 of a network. A network with a low code length has more nodes compressed into distinct  
<sup>180</sup> subunits/provinces without losing the underlying information flow of the graph. In the case  
 of measuring community connectedness, a low code length means greater site distinctness  
<sup>182</sup> and provinciality than a graph with a high code length [118]. For further examples, visit  
<http://mapequation.org/>.

<sup>184</sup> Analysis of these measures of community connectedness both within and between different  
 regions across the globe allows for the expected relative importance of global versus regional  
<sup>186</sup> versus local scale processes, and how this might change over time, to be estimated. If global pro-  
 cesses are important to patterns of community connectedness and environmental interactions

than it is expected that these will be correlated with global climate measures. Additionally,  
188 if two or more regions have similar or correlated patterns of community connectedness, it  
is expected that global processes may play a role in shaping these environments. Regional  
189 processes are expected to dominate when  $E$  is low,  $Occ$  is high,  $BC$  is high, and code length  
is high. In contrast, local processes are expected to dominate when  $E$  is high,  $Occ$  is low,  $BC$   
190 is low and code length is low. The different scales are not mutually exclusive, however, and  
one or more scales might be involved in shaping patterns of community connectedness and  
191 environmental interactions. Importantly, which process scales are dominant may change over  
time.

<sup>196</sup> **2 Australian Permian Brachiopods**

## 2.1 Traits and environmental preference

<sup>198</sup> Brachiopods are suspension feeders that are not actively motile, thus the availability of optimal environmental conditions is extremely important for both establishment and survival.  
<sup>200</sup> Brachiopod occurrence has been found to be strongly linked to the type ocean floor on which they occur [108, 109]. Environmental preference is estimated here using three traits,  
<sup>202</sup> two environmental and one biological: substrate preference, habitat preference, and surface interface/affixing strategy. Each of these three traits relate to a different aspect of the environment and a taxon's specific adaptive zone. Importantly, the loss of suitable environments  
<sup>204</sup> may determine different trait associated extinction risk. While larval mode is considered  
<sup>206</sup> an important trait associated with geographic range and extinction risk [58, 59], it does not persevere in brachiopods and thus cannot be used to model survival [59]. Additionally,  
<sup>208</sup> taxa found on the east coast of Australia during the Permian would have been facing the Panthalassic Ocean and would have had few "good" areas to disperse to because of the  
<sup>210</sup> distance required to reach a different shore line and the high latitude of the region.

<sup>212</sup> Substrate preference is related to the chemical and physical processes present in a given environment. Substrate selection is mitigated via larval chemosensory abilities and thus may act as a weak proxy for larval dispersal ability [58, 59]. The three generally used states of  
<sup>214</sup> substrate affinity are carbonate, clastic, or mixed [7, 40, 70, 87, 90]. The Pharenozoic is characterized by an overall decline in carbonates relative to clastics [40, 87]. Additionally,  
<sup>216</sup> the Australian Permian is dominated by clastic beds [13, 32, 33, 95, 130]. It is expected then that the majority of brachiopod taxa will prefer clastic type substrates compared to the rarer  
<sup>218</sup> carbonate type substrates.

<sup>220</sup> Habitat preference is a description of the environment in which a taxon was found at the time of fossilization. Because brachiopods are sessil and frequently fixed to the environment, it is expected that the inferred environment is akin to that in which they lived. The range of environments are quite broad, representing many different marine settings. Because of the large range and difficulty of precisely inferring paleoenvironment a frequently used, albeit  
<sup>224</sup> coarse, classification is on-shore versus off-shore [15, 57, 61, 70, 116] along with the option of a taxon having no particular habitat preference. Importantly, habitat availability is broadly  
<sup>226</sup> related to sea-level which can change both dramatically and rapidly over time [91]. Because of this, on-shore type habitats are potentially very volatile and unstable for long periods of time. During the Permian of Australia there were four major glaciation events which covered most of the entire continent [13, 31–33], which most likely strongly impacted sea-level as well  
<sup>230</sup> as the availability and constancy of on-shore versus off-shore habitats. Additionally, habitat preference might capture other factors relating to the environment not captured in substrate preference.

Affixing strategy is the manner by with an individual interfaces with the ocean floor. Unlike

<sup>234</sup> bivalves which can burrow or snails which are motile, a brachiopod is sessil and has to maintain  
<sup>236</sup> their commissure at or above substrate level in order to expose their lophophore. Because of  
<sup>238</sup> this, brachiopods have evolved a variety of different methods to position themselves in various  
<sup>240</sup> different environmental conditions such as flow speed or mud depth [1, 75, 76, 108, 114].  
<sup>242</sup> Broadly, these methods can be classified as pedunculate (presence of a pedicle), reclining  
(absence of pedicle), and cementing. During the Permian, pedunculate taxa are associated  
with shallow on-shore environments while reclining taxa are associated with deep off-shore  
environments [19] however these associations are weak as most assemblages are composed of  
a heterogeneous mix of strategies.

Additionally, during the Permian there was a shift from an “ice house” to a “hot house” world  
<sup>244</sup> [13, 31, 33, 68, 98]. Australian taxa are of particular interest because of their proximity to the  
south pole during the Permian and the repeated glacial activity in the region [13, 31, 32, 68].  
<sup>246</sup> According to Olszewski and Erwin [91], however, sea-level and climate change do not wholly  
explain the brachiopod ecological dynamics experienced in the Permian of Texas. It is then  
<sup>248</sup> predicted that climate will not be the best sole predictor of brachiopod survival, and that  
some combination of one or more the above organismal traits will be necessary to best model  
survival.  
<sup>250</sup>

## 2.2 Environmental preference and extinction

### <sup>252</sup> 2.2.1 Questions

Which traits relating to environmental preference in brachiopods are predictors, either  
<sup>254</sup> separate or together, of differential survival? Do Permian glacial periods relate to differences  
in trait-correlated extinction? What is the distribution of brachiopod generic survival?

### <sup>256</sup> 2.2.2 Hypotheses and predictions

Because of both the long-term decline in carbonates versus clastics [97] and the dominance of  
<sup>258</sup> Permian-age clastic beds [13, 32, 33, 95, 130] described above (Section 2.1), taxa with clastic  
type affinities are expected to have longer durations than taxa with any other preference.  
<sup>260</sup> Additionally, this substrate dominance may have been a strong selection pressure for taxa  
to adapt to the common clastic types and/or away of the rarer carbonates. Because of this,  
<sup>262</sup> it is expected that taxa with clastic or mixed affinities will have greater survival than taxa  
associated with carbonate substrates. Additionally, it is predicted that substrate preference,  
<sup>264</sup> if it captures the same information as modern substrate type, will be a predictor in the best  
model(s) of survival [108, 109]. However, if substrate affinity is not found to be important  
<sup>266</sup> for modeling survival this may be due to one or more reasons. First, substrate affinity, as  
quantified here, may not be capturing the same information as modern substrate type and  
<sup>268</sup> thus may act as a poor predictor of survival. Second, it may mean that because clastic type

substrates were so dominate during the Permian of Australia that survival may be better  
270 explained by other factors, either measured or unmeasured.

While other environmental factors beyond substrate type, such as temperature or water depth,  
272 have not been found to limit the distribution of modern brachiopods [108, 109] it is unknown  
274 if these traits affect survival. Predictions of differential survival based on habitat preference  
and affixing strategy can be made on the basis of environmental preference and availability.

During the Permian of Australia there were four major glaciation events where most of the  
276 continent was covered [13, 31–33]. Because of this it is expected that off-shore adapted taxa  
278 will have greater durations than on-shore adapted taxa. This prediction is based on the  
expected constancy and availability of off-shore habitats and the expected high volatility of  
on-shore habitats. If habitat preference is not found to be a predictor for modeling survival,  
280 this may mean that sea-level mediated environmental availability does not determine long  
term survival. Specifically, while sea-levels may have fluctuated greatly due to high latitude  
282 glaciation [13, 32, 33] it may be that the long term continual availability of habitat over-  
shadows short term fluctuations. Also, it has been found in the case of Permian brachiopods  
284 from Texas that sea-level along with climate change do not wholly explain the observed  
ecological dynamics [91], which may mean that habitat availability may not be the singly  
286 dominate factor when modeling brachiopod survival.

Previous global level analysis of brachiopod durations showed that affixing strategy is  
288 correlated with longevity [1] and that among endemic taxa, reclining taxa had longer durations  
than other affixing strategies. Additionally, differential survival between affixing strategies has  
290 been observed at the Cretaceous/Paleogene mass extinction [66]. Among cosmopolitan taxa,  
however, pedunculate and cementing taxa had longer durations than all other taxa, both  
292 cosmopolitan and endemic. This global analysis mixed taxa from many different geological  
periods and geographic regions which may have led to unfair and biased comparisons. By  
294 restricting analysis to a single continuous region and geological time period, I hope to  
alleviate these concerns and instead focus on survival of an evolving taxonomic assemblage in  
296 a continuous environmental context. If affixing strategy is found to not be a predictor in the  
best model(s) of survival this would mean that, while it is correlated with differential survival  
298 [1], it may only be a minor factor. For example, this may indicate that the environmental  
energetics of Australia were rather uniform or constant with respect to time.

300 An important consideration is that taxonomic survival might not be linked to single environments *per se*, but the variability of environments [41, 50, 77] which has been found to relate  
302 strongly with survival past origination. Adaptation to variability of environments may be  
captured in taxa with mixed substrate preference and/or no habitat preference. Based on  
304 this observation, it is predicted that taxa with mixed substrate preference and/or no habitat  
preference will have longer durations than taxa with single preferences. However, this may  
306 also mean that taxa with mixed substrate and/or no habitat preferences will be of similar  
duration to clastic type and off-shore preferences, which are predicted to have the longest  
308 durations for their respective traits.

### 2.2.3 Proposed research

310 In order to investigate which traits best model survival and how, I propose a survival analysis  
 311 approach (Section 1.1). I choose to restrict this analysis to Australia because it represents a  
 312 relatively continually sampled and well worked area that preserves the majority of the entire  
 313 Permian [8, 13, 21, 22, 33, 136]. The traits described above (Section 2.1) will be used as  
 314 predictors of survival. The distribution of survival durations will be modeled using a variety  
 315 of different distributions which are tied to different hypotheses of extinction risk (Section  
 316 1.1).

Permian brachiopod occurrence information is available via the Paleobiology Database  
 318 (PBDB; <http://fossilworks.org>) and is primarily sourced from the work of Clapham  
 319 [19–23] and Waterhouse [136]. While lithological and paleoenvironmental information is  
 320 available for some occurrences through the PBDB, this information is frequently missing or  
 321 too coarse. Lithological information and paleoenvironmental reconstructions will be heavily  
 322 supplemented using the extensive geological unit information from Geosciences Australia  
 323 (<http://www.ga.gov.au/>) as well as the literature on the stratigraphy of Australian Permian  
 324 basins [13, 31–34, 44, 68, 95, 135, 136]. For example improvements to the initial PBDB  
 assignments, see Appendix A.

326 Duration will be measured as the difference between FAD and LAD. If a taxon originates  
 327 prior to the Permian or goes extinct within 5 million years of the Permo-Triassic (P/T)  
 328 boundary or after the P/T it will be censored. The possibility of accounting for the affect  
 329 of sampling on decreasing observed durations versus true durations may be done through  
 330 interval censoring (Section 1.1). For example, a range of LAD values between the observed  
 331 and that estimated via unbiased point estimation [6, 127] can be used. The unbiased point  
 332 estimation of true extinction time is calculated using

$$r = \frac{R}{H - 1} \quad (5)$$

where  $r$  is the average gap size between fossils,  $R$  is the stratigraphic range, and  $H$  is the  
 334 number of fossil horizons.  $r$  is then added to the LAD value for an estimate of the true  
 extinction time.

336 The most probable genus substrate and habitat preferences are estimated from the distribution  
 337 of occurrences. Preliminarily, the lithological setting of all occurrences will be classified into  
 338 one of three substrate affinity categories following Foote [40] while paleoenvironmental settings  
 339 will be classified following Kiessling et al. [71]. Both of these traits will be assigned to all  
 340 taxa following the Bayesian approach of Simpson and Harnik [121] where assignments are  
 341 determined as the posterior probability of a taxon’s occurrences in comparison to available  
 342 options during the duration of said taxon. The probability that a genus prefers, for example,  
 on-shore habitat ( $P(H_1|E)$ ) is calculated as

$$P(H_1|E) = \frac{P(E|H_1)P(H_1)}{P(E|H_1)P(H_1) + P(E|H_2)P(H_2)} \quad (6)$$

344 where the prior probability  $P(E)$  is the proportion of all occurrences that are on-shore.  
 The null hypotheses,  $P(H_1)$  and  $P(H_2)$ , differ for assignments of substrate and habitat. For  
 346 substrate,  $P(H_1) = P(H_2) = 0.5$ , meaning that the null is that a genus has no preference.  
 For habitat preference, probability of assignment is calculated three times with  $P(H_1) = \frac{1}{3}$   
 348 and  $P(H_2) = \frac{2}{3}$ , meaning that the null is that there is an equal chance that a genus prefers  
 on-shore, off-shore, or neither habitat. The conditional probabilities,  $P(E|H_1)$  and  $P(E|H_2)$ ,  
 350 are calculated using the binomial probability of observing the number of occurrences in, for  
 example, on-shore habitats,  $k$ , out of the total number of occurrences,  $n$ . The conditional  
 352 probability is calculated as

$$P(E|H_1) = \binom{n}{k} p^k (1-p)^{n-k} \quad (7)$$

354 where  $p$  is the proportion of collections in on-shore habitats observed during a taxon's  
 duration.

356 In the case of the coarse classification schemes of Foote [40] and Kiessling et al. [71], the  
 following rules are used assign preference. For substrate affinity, if  $P(H_1|E) > \frac{2}{3}$  then the taxon  
 358 was considered of carbonate affinity while if  $P(H_1|E) < \frac{1}{3}$  then the taxon was considered to  
 have a clastic affinity. Otherwise, the taxon was considered to have mixed affinity. For habitat  
 360 affinity, the posterior probability for each habitat (inshore, offshore, none) was calculated  
 using Eq. 6 and the preference with maximum of the three posterior probabilities was assigned.

362 Each of the three traits will be considered constant throughout the duration of a genus and  
 will be modeled as time-independent covariates of survival. If and how these traits may have  
 evolved will remain for future study.

364 Because there is no obvious single best model, multiple models will be compared in order to  
 determine which is the most likely model of survival. It is important, however, that each model  
 366 be well justified and be tied to a realistic biological hypothesis/prediction [18]. Below are a list  
 of possible models of brachiopod survival, based solely on time-independent covariates and not  
 368 time-dependent covariates (below) nor distribution of survival, and the associated hypotheses  
 (Table 1). This does not represent an exhaustive list of plausible models or hypotheses.

370 Because the four major periods of glacial activity during the Permian of Australia may have  
 had dramatic impacts on survival and environmental availability, it is necessary to model  
 372 glacial activity as a time-dependent covariate. In the simplest case, it is possible to model  
 glacial activity as a step-function with two states: ice or no ice. The ages of the onset and  
 374 retreat for all of the glacial period are fairly well constrained [32, 33]. Other options for  
 modeling climatic change are to use various Australian Permian isotope records [13] as more  
 376 fine grained estimates of environmental change.

378 Because survival models are fit in a maximum likelihood framework [74], model comparison  
 and selection can be done via AICc scores [18, 53].

## 2.2.4 Preliminary results

- 380 The preliminary results presented here are based entirely on the data present in the PBDB without modification (Appendix A). Observations were censored following the procedure  
382 described above (Section 2.2.3). Uncertainty of duration was not taken into account via interval censoring. Substrate and habitat preference were the only covariates of survival  
384 and were classified coarsely following Foote [40] and Kiessling et al. [71] respectively. Model formulations with each covariate alone, additively together, or interacting were used. Only  
386 two different survival distributions were considered in this initial analysis: exponential and Weibull. In total, 11 models were fit and compared (Table 2).
- 388 The best model of survival had substrate preference as the sole predictor of survival, which followed a Weibull distribution with increasing risk of failure with age (Table 2). This model  
390 was closely followed by the second best model of survival which had both substrate and habitat preference as additive predictors of survival, also following a Weibull distribution with  
392  $k > 1$ . The difference between the AICc best model and the second best model was small ( $\Delta$  AIC  $\approx 1.3$ ), meaning that both models can be considered almost equivalent.
- 394 The AICc best model is illustrated below (Fig. 1c) as well as the model with habitat as the sole predictor of survival (Fig. 1d)
- 396 The shape parameter ( $k$ ) of the AICc best model (Fig. 1c) is estimated to be approximately 1.85 (Table 2). As described above (Section 1.1), values of  $k$  greater than 1 indicate that  
398 failure (extinction) risk accelerates with taxon age, which may mean that the Law of Constant Extinction does not hold when modeling generic level extinction in brachiopods.
- 400 For brachiopod survival based on substrate affinity (Fig. 1c), survival was greater for both carbonate and clastic affinities and lowest for taxa with mixed affinity. Visual inspection  
402 of the estimated survival functions compared to the nonparametric Kaplan–Meier curves indicates that they are adequate fits to the data (Fig. 1a).
- 404 The model with habitat preference being the sole predictor of survival following a Weibull distribution was a poor estimate, with an approximate  $\Delta$ AICc of 22 between this model and  
406 the AICc best model. There is a great degree of deviance between the nonparametric Kaplan–Meier curves and model predictions (Fig. 1b). Additionally, this model is not significantly  
408 different from the model with only an intercept ( $\chi^2 = 1.14$ ,  $df = 2$ ,  $p = 0.57$ ). This means, preliminarily, that habitat preference alone makes no difference in generic level survival.
- 410 Further refinements to these models include modeling survival using other distributions of survival such as a log-normal distribution. Additionally the inclusion of affixing strategy and  
412 climate as predictors will increase the understanding of the biology underlying brachiopod generic survival.

414 **2.3 Brachiopod distribution and community connectedness**

2.3.1 Questions

- 416 Given the repeated major glacial activity during the Permian, how stable was community  
connectedness in Permian brachiopods? Are patterns of community connectedness different  
418 for taxa favoring different environments?

2.3.2 Hypotheses and predictions

- 420 During the Permian, the east coast of the Australian continent faced towards the massive  
Panthalassic Ocean. Because of this, the establishment of populations was most likely limited  
422 to within the local area because the amount of distance required to establish else was  
most likely too great. Additionally, individuals which settled across the ocean would have  
424 been almost instantly genetically isolated and not increase community connectedness, *per*  
*se*. Because of this, it is expected that community connectedness in Australian Permian  
426 brachiopods would be fairly high at any given time and that changes, specifically decreases  
in connectedness, would be expected during the four glacial periods [32, 33].
- 428 Dispersal ability of modern brachiopods appears to be most limited by availability and  
proximity of substrate types [108, 109]. The Permian of Australia is dominated by widespread  
430 clastic beds compared to relatively few carbonate beds. The expectation is that the distribution  
of taxa with a carbonate preference will be extremely patchy with a high  $E$  (Eq. 2), low  
432  $Occ$  (Eq. 3), low  $BC$  (Eq. 4), and low code length [111, 118] compared to the distribution of  
clastic preferring taxa. However, if community connectedness is approximately equal between  
434 carbonate and clastic preferring taxa this could be caused by approximately equal dispersal  
ability in both groups, either high or low.
- 436 Habitat would be expected to influence community structure if there is an uneven distribution  
of available habitats in space and time. Rarity of preferred habitat would be expected to lead  
438 to high  $E$ , low  $Occ$ , low  $BC$ , and low code length compared to an abundance of preferred  
habitat. Because of the four major glaciation events during the Permian of Australia, it is  
440 expected that the availability of on-shore habitats would be highly variable. It is then expected  
that during periods of glacial activity community connectedness of on-shore preferring taxa  
442 would be extremely low because of rarity of environments in comparison to both periods of  
non-glacial activity and off-shore habitats at all times. If habitat preference has no effect on  
444 community connectedness this may mean that the dispersal ability of on-shore taxa is very  
high and able to maintain gene flow between potentially isolated habitats.
- 446 It is expected that affixing strategy alone will have minimal effect on community connectedness  
unless affixing strategy is highly correlated with substrate and/or habitat preference. If  
448 community connectedness is found to be different between affixing strategies but affixing  
strategy is not highly correlated with substrate or habitat preference this may be because of

<sup>450</sup> spatial heterogeneity in energy levels which limits reclining versus fixed taxon distributions.  
This scenario is highly unlikely given knowledge of modern and fossil brachiopod distributions  
<sup>452</sup> [108, 109, 114].

### 2.3.3 Proposed research

<sup>454</sup> Using a biogeographic network approach (Section 1.2), I will construct networks between  
brachiopod genera and localities defined as 2x2 latitude–longitude grid cells from an equal-area  
<sup>456</sup> map projection. Biogeographic networks will be constructed for the entire Permian using 2 My  
bins. In addition to community wide networks, separate networks will be constructed for taxa  
<sup>458</sup> within ecological categories. This facilitates comparison of community connectedness patterns  
during the Permian both within and between categories as well as with the community wide  
<sup>460</sup> pattern. The data necessary to complete this study is the same as for the above analysis of  
brachiopod survival (Section 2.2). Importantly, sampling will be restricted to the east coast  
<sup>462</sup> of Australia because this represents a continuous coast line that faced the Panthalassic Ocean  
during the Permian.

<sup>464</sup> Trait assignment will follow the procedure outlined for analysis of brachiopod survival (Section  
2.2.3).

<sup>466</sup> The next step is to compare patterns of community connectedness both within and between  
regions in order to understand if global, regional, or local scale processes dominate. Addition-  
<sup>468</sup> ally, comparisons will be done between the different ecological traits both within and between  
regions to determine which scale processes may be dominate. The approach and methodology  
<sup>470</sup> to accomplish these analyses is currently under development. Additionally, the possibility  
of integrating locality–locality distance or some other measure of topology will be explored,  
<sup>472</sup> especially how this relates to code length and provinciality in general.

### 2.3.4 Preliminary results

<sup>474</sup> Preliminary results are based solely on the brachiopod occurrence information in the PBDB.  
Preliminary networks were constructed with taxa being defined as genera and localities  
<sup>476</sup> defined from a 2x2 latitude-longitude grid from an equal area map projection. All localities  
were restricted to those occurring in basins not present in the state of Western Australia.  
<sup>478</sup> Networks were also constructed for taxa divided by substrate and habitat preferences. No  
initial comparisons with the Permian glacial record have been made. These results are based  
<sup>480</sup> on the lithological and paleoenvironmental data present in the PBDB which will be improved  
as discussed above (Section 2.3.3).

<sup>482</sup> The summary statistics for community connectedness for all brachiopods show a qualitatively  
random pattern (Fig. 2) with no observable trends. Three of four summary statistics fluctuate  
<sup>484</sup> continually ( $E$ ,  $Occ$ , code length) while  $BC$  is qualitatively stationary throughout the Permian.  
Importantly, this pattern is effectively the same as that seen in clastic preferring taxa (Fig. 3a).

- <sup>486</sup> These preliminary results are also demonstrate the predicted rarity of carbonate preferring  
taxa (Fig. 3a)
- <sup>488</sup> Additionally, taxa with both in-shore and no habitat preference have approximately identical  
patterns that are also qualitatively random in contrast to the qualitatively stable off-shore  
<sup>490</sup> preferring taxa (Fig. 3b).

Because these results are based on only preliminary substrate and habitat assignments, there  
<sup>492</sup> is still major room for improvement. Additionally, patterns have not been explored for taxa  
based on affixing strategy, which may or may not follow the same pattern as substrate (Fig.  
<sup>494</sup> 3a). There are many further analyses to accomplish. Most importantly, comparisons both  
within and between the different ecological traits as well as with the timing of the four glacial  
<sup>496</sup> periods are necessary in order to better understand what environmental factors may affect  
measuring the four network summary statistics alternative methods for summarizing network  
<sup>498</sup> and taxon distributions will be explored.

formulation	hypothesis
~ 1	No differential survival based on measured ecological traits.
~ substrate	Substrate availability is the best predictor of survival as expected based on the distribution of modern taxa [108, 109].
~ habitat	Habitat stability is the best predictor as expected by models of Phanerozoic diversification.
~ affixing strategy	Environmental homogeneity/stability means that differentiation can only occur via differences in how a taxon interfaces with the ocean floor.
~ substrate + habitat	Substrate and habitat combine to best describe the environmental context of a taxon and the availability of its adaptive zone.
~ habitat + affixing strategy	By combining well adapted affixing strategy to the energetics of the habitat, survival increases.
~ substrate + affixing strategy	By combining well adapted affixing strategy to the state of the ocean floor increases survival.
~ substrate + habitat + affixing strategy	The adaptation of affixing strategy along with the environmental context represents the best approximation of the adaptive zone.

Table 1: Some example candidate models of brachiopod survival based on substrate affinity, habitat preference, and affixing strategy. Each model is presented with an associated hypothesis of biological meaning. A formulation of ~ 1 is a model with only an intercept and no covariates. Formulations are without reference to the distribution of survival.

formula	distribution	shape	df	AICc	weight
$\sim \text{aff}$	weibull	1.85	4	941.6757	0.65
$\sim \text{aff} + \text{hab}$	weibull	1.87	6	942.9977	0.34
$\sim \text{aff} * \text{hab}$	weibull	1.89	10	949.0816	0.02
$\sim 1$	weibull	1.74	2	960.2550	0.00
$\sim \text{hab}$	weibull	1.75	4	963.3091	0.00
$\sim \text{aff}$	exponential		3	993.1724	0.00
$\sim \text{aff} + \text{hab}$	exponential		5	996.4089	0.00
$\sim 1$	exponential		1	1000.2592	0.00
$\sim \text{aff} * \text{hab}$	exponential		9	1003.7639	0.00
$\sim \text{hab}$	exponential		3	1003.9227	0.00

Table 2: Model selection table for the preliminary models of brachiopod survival. As in Table 1, a formulation of  $\sim 1$  is a model with only an intercept and no covariates. The  $*$  symbol corresponds to covariate interaction.  $w$  are Akaike weights [18].

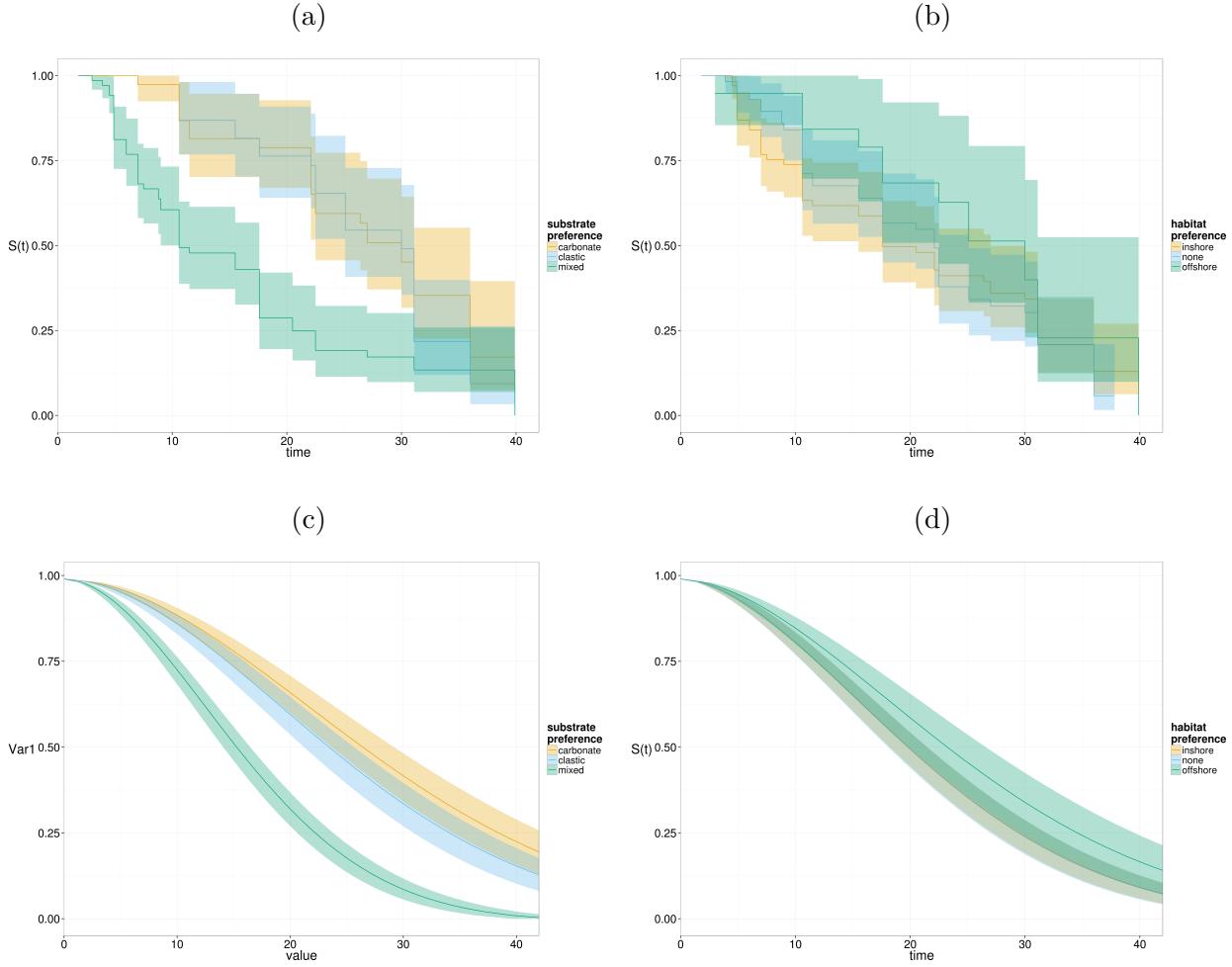


Figure 1: Survivorship curves of Australian Permian brachiopod genera based on substrate affinity and habitat preference. The stepwise functions are nonparametric Kaplan–Meier survival curves for both substrate affinity (a) and habitat preference (b). K–M curves are illustrated with 95% confidence intervals. The predicted survival curves based on the best parametric models with substrate (c) and habitat (d) as predictors. The parametric curves are illustrated with the standard errors of prediction.

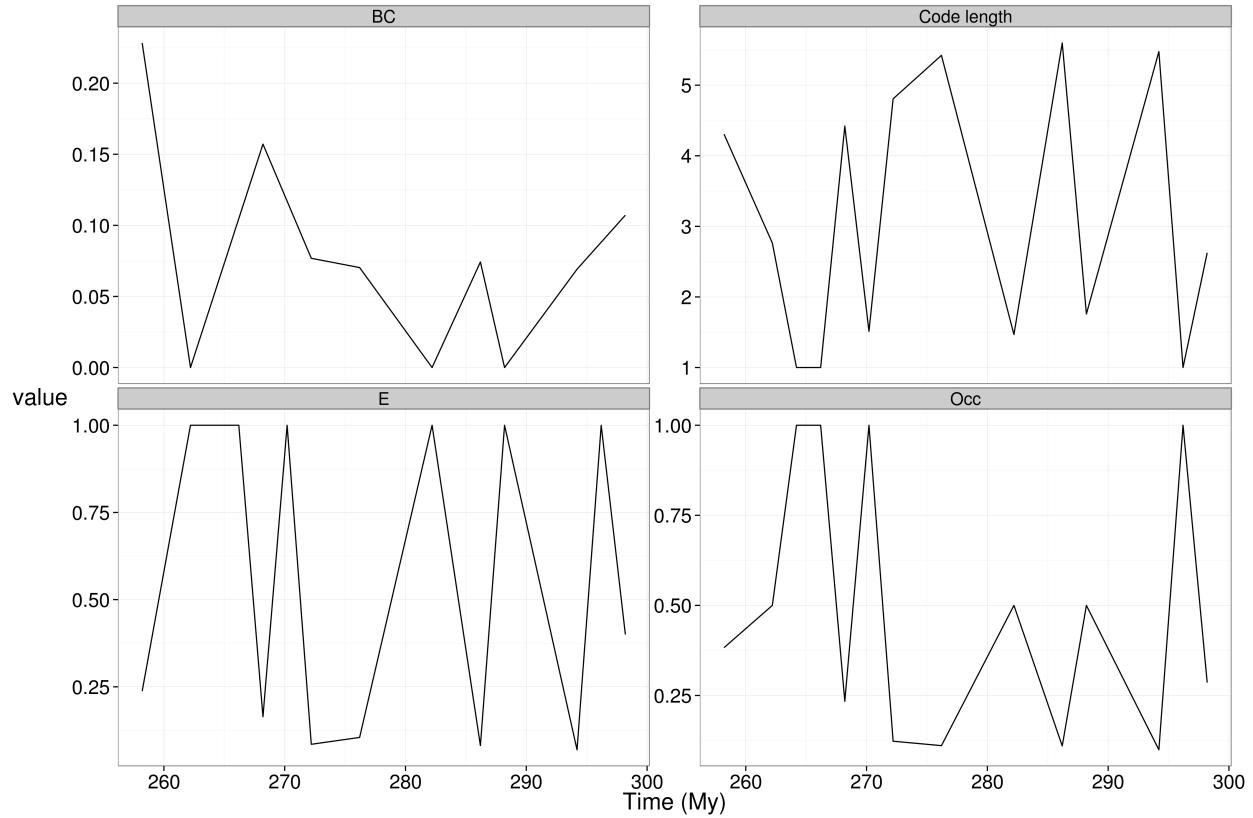


Figure 2: Summary statistics of community connectedness for all brachiopods occurring on the East coast of Australia during the Permian. The summary statistics are, clockwise from top left: biogeographic connectedness (BC), code length, average relative locality occupancy per taxon (Occ), and average relative number of endemic taxa per locality (E).

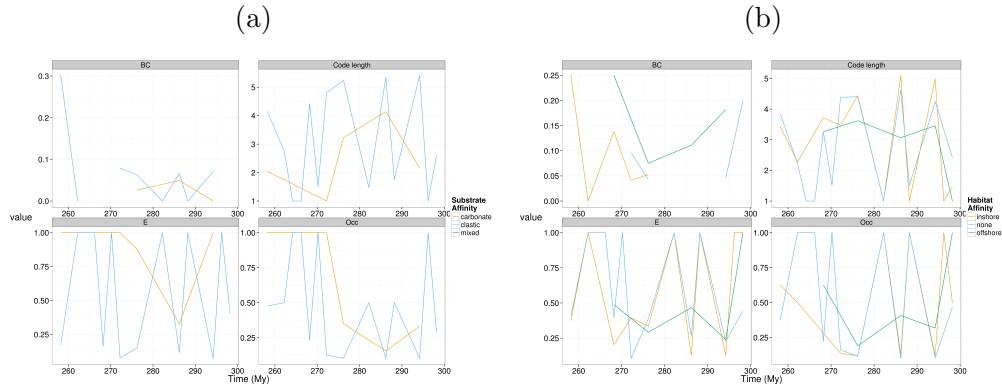


Figure 3: Community connectedness statistics for brachiopods separated by substrate (a) and habitat (b) preference. The summary statistics are, clockwise from top left: biogeographic connectedness (BC), code length, average relative locality occupancy per taxon (Occ), and average relative number of endemic taxa per locality (E).

500 **3 Cenozoic Mammals**

### 3.1 Traits and environmental context

502 Mammals are motile organisms which can track their preferred environmental context over  
time. However, if a taxon requires rare or fragile environmental conditions, or is a poor  
504 disperser, this would limit the availability of suitable environments or ability to track the  
preferred environment. Three important traits that describe the relationship between mammals  
506 and their environmental context are body size, dietary category, and locomotor category  
[25, 26, 65, 80, 81, 124, 125]. Each of these traits describe different aspects of a taxon’s  
508 adaptive zone such as energetic cost, population density, expected home range size, set of  
potential prey items, and dispersal ability among others. Additionally, these three traits are  
510 relatively easy to estimate from fossils.

Environmental availability, along with stability, is crucial for both the establishment and  
512 persistence of a species. During the Cenozoic, primarily between the Paleogene–Neogene, there  
was a shift from a predominately closed environment to a predominately open environment  
514 [14, 62, 110]. This environmental shift was differently timed between continents [128, 129].  
Because of the differential timing of environmental shift, along with the different biotic  
516 context, the community and survival patterns are expected to vary between continents.

Dietary categories are coarse groupings of similar dietary ecologies: carnivores, herbivores,  
518 omnivores, and insectivores. Each of these categories is composed of taxa with a variety of  
ecologies. For example, herbivores include both browsers and grazers which are known to  
520 have had different diversification dynamics during the Cenozoic [63]. Dietary categories are  
roughly linked with position in trophic hierarchy, with decreasing stability away from the  
522 “base.” Stability here meaning trophic “distance” from primary productivity, with herbivores  
having greater stability than carnivores because of the increased likelihood of prey item  
524 occurrence. Additionally, with increased likelihood of prey item occurrence, abundance can  
increase [17, 25, 63, 119, 134].

526 Locomotor categories describe the motility of a taxon, the plausibility of occurrence, and the  
dispersal ability. For example, an obligate arboreal taxon can only occur in locations with a  
528 minimum of tree cover and can most likely only disperse to other environments with suitable  
tree cover. Locomotor categories are similar to dietary categories as they represent coarse  
530 groupings of taxa with similar life habits. Here, the categories are arboreal, ground dwelling,  
and scansorial. Similar to dietary category, this trait is considered constant at the specific  
532 level. Dispersal ability is important for determining the extent of a taxon’s geographic range  
[12, 45, 58] and affects both the taxon’s extinction risk and regional community evenness.  
534 With the transition from primarily closed to open environments, there is an expected shift in  
stability associated with arboreal and ground dwelling taxa.

536 An organisms body size, here defined as (estimated) mass, has an associated energetic cost in  
order to maintain homeostasis which in turn necessitates a supply of prey items. Many life

538 history traits are associated with body size: reproductive rate, metabolic rate, home range  
size, among many others [17, 25, 96, 124]. While studies of body size dynamics are very  
540 common [5, 24, 26, 67, 78], the interactions or processes that are correlated with body size  
might be underlying the observed diversity pattern more than body size itself. By combining  
542 analysis of body size and both dietary and locomotor categories, it should be possible to  
better understand what processes underly patterns of survival and community connectedness.

## 544 3.2 Ecologically mediated survival

### 3.2.1 Questions

546 Which ecological traits relating to environmental selection in mammals are predictors, either  
separately or together, of differential survival? How does both regional and global environmen-  
548 tal shift relate to differential survival? Are the distributions of generic and specific survival  
different?

### 550 3.2.2 Hypotheses and predictions

Because dietary category describes, roughly, the trophic position of a taxon and its related  
552 stability, it is predicted that more stable categories will have longer durations than less stable  
categories. Stability here being “distance” from primary productivity, thus it is expected  
554 that herbivores will have greater duration than carnivores. Omnivorous taxa are expected to  
have average taxon durations compared to the other two categories. If dietary category is  
556 not found to be important for modeling survival it may mean that trophic category is not a  
major factor for determining species level survival and that other factors, such as body size,  
558 may dominate.

Mammalian herbivores and carnivores have been found to have a greater diversification rate  
560 than omnivores [99] which may indicate that these traits are better for survival. However  
diversification can be caused either by an increase in origination relative to extinction or a  
562 decrease in extinction relative to origination. Which scenario occurred, however, is (currently)  
impossible to determine from a phylogeny of only extant organisms [100] which means that  
564 analysis of the fossil record is required. If survival is found to be similar between all dietary  
categories, this may mean that the differential diversification patterns observed by Price et al.  
566 [99] are due to differences in speciation and not extinction.

It is expected that arboreal taxa during the Paleogene will have a greater expected duration  
568 than Neogene taxa, and the opposite will be true for ground dwelling taxa. In comparison,  
taxon duration of scansorial taxa is expected to remain relatively similar between the two  
570 time periods because it represents a mixed environmental preference that may be viable  
in either closed or open environments. If locomotor category is not included in the best  
572 model of survival this may mean that it is either a poor descriptor of dispersal ability which

may or may not affect mammalian survival. However, it may be the case that other factors, measured or unmeasured, may be of greater importance in determining differential survival. The difficulty of a Paleogene–Neogene comparison, which is potentially undermined by heterogeneous preservation, will be explored in simulation (Section 1.1.1).

Body size can possibly scale up to affect species level patterns because as body size increases, home range size increases [25]. If individual home range size scales up to reflect minimum total species geographic range, we would expect that taxa with larger body sizes would have lower extinction rates than species with smaller body sizes. This expectation, however, may not be right. As body size increases, reproductive rate decreases [67], populations get smaller [138], and generations get longer [86] all of which can increase extinction risk, as has been observed [27, 78]. However, the relationship between body size and extinction rate at the generic level has been found to vary between continents [78, 131]. By expanding to include a third continent, South America, and analyzing specific level data I hope to elucidate how differences in taxonomic diversity at a continental level might affect body size mediated extinction rate. If body size is found to be unimportant for modeling survival, as in the generic level analysis of Tomiya [131], this means that other biotic or abiotic factors may dominate. Also, this may mean that individual level home range size does not scale into increased species level range size, and there is therefore no correlated decrease in extinction rate. If increase in body size increases extinction risk, this may be due to traits correlated with body size and not necessarily body size itself [67].

The interaction of body size, locomotor category, and dietary category is also extremely important. For example, a small bodied arboreal taxon of any trophic category during the heavily forested and warm time of the Paleogene would be expected at once to have both a small body size determined range, a large potential geographic range determined by locomotion, as well as an increased availability of resources. Together this would mean that relative survival would be expected to be less than, greater than, and greater than average respectively. Determining which factors dominate during the Paleogene, as well as other parts of the Cenozoic, must be done empirically.

### 3.2.3 Proposed research

To analyze differential mammalian survival, I propose a survival analysis approach (Section 1.1) similar to that described above for Permian brachiopods (Section 2.2). Mammalian occurrence data will be collected primarily through a combination of the PBDB, Neogene Old World Database (NOW; <http://www.helsinki.fi/science/now/>), and museum collections. North American fossil mammal data are well represented in the PBDB because of the extensive work of Alroy [3–5]. European fossil mammal data is also well represented between the PBDB and NOW. South American fossil mammal data is available through the PBDB, but has poor overall coverage. Because of this, South American fossil mammal data will be gathered via various museums such as the Field Museum of Natural History and the American Museum of Natural History as well as published occurrence compilations. With

612 the South American taxa, taxonomy and sampling may not be as well resolved as for North  
613 and South America and it may be necessary to restrict analysis to the most taxonomically  
614 resolved and sampled groups such as Notoungulata, Marsupials, Carnivora, and Primates.

As described above (Section 2.2.3), duration will be measured as the difference between the  
616 observed FAD and LAD of every taxon. Taxa which originated prior to the Cenozoic and all  
617 taxa that are either extant or went extinct within 2 My of the present will be censored. This  
618 threshold is to limit the effect of the improved record of the Recent.

Dietary category, locomotor category, and body size will be considered constant throughout  
620 the duration of a taxon and will be modeled as time-independent covariates of survival. While  
621 body size is actually a distribution of values, it is quite common to use a single estimate of  
622 mean body size as an aggregate trait in studies of clade-wise dynamics [56]. While all three  
623 of these traits may have evolved over a taxon’s duration, this will not be considered as part  
624 of this study.

While many analyses of survivorship are done using generic data [36, 40, 48, 78, 131], there  
626 are potential biases in accurately modeling a specific level process using generic level data  
627 [104, 105, 115, 120, 133]. In order to assess some of the differences between generic and  
628 specific level survival, I will estimate specific and generic level survival models. Using an  
629 approach similar to previous work on estimating specific level origination and extinction  
630 rates from generic level survival curves [38], I will measure the deviance between extinction  
631 rate directly estimated from the specific survivorship and the specific level extinction rates  
632 estimated from the generic level survival data. In addition to empirical comparison between  
633 generic and specific level survival, simulations of diversification with varying levels of cryptic  
634 speciation (anagenesis). This may also act as a proxy for generic level diversification because  
635 a lineage having a long duration because it is not correctly broken up can be considered  
636 analogous to a genus persisting because it continues to speciate.

As with the brachiopods (Section 2.2.3), there is no obvious single best model of survival,  
638 so multiple models must be compared in order to determine which is the most likely. It is  
639 important, however, that each model be well justified and be tied to a realistic biological  
640 hypothesis/prediction [18].

In order to account for environmental shifts, two different time-dependent covariates will be  
642 used.  $\delta O^{18}$  isotope information for the whole Cenozoic [140] will be used as a global climate  
643 proxy. Additionally, the Paleogene–Neogene divide, which may reflect global environmental  
644 shift, will be modeled as a time-dependent step-function.

### 3.2.4 Preliminary results

646 Preliminary results are based solely on Cenozoic mammal occurrence data from North America  
647 and Europe from the PBDB. Nonparametric Kaplan-Meier survival curves were estimated  
648 for both dietary and locomotor categories (Fig. 4). These are shown on a log-linear scale for

more direct visual comparison with linearity [132, 133].

The North American species-level survival curves, both based on dietary (Fig. 4a) and locomotor categories (Fig. 4b), are semi-log linear as expected under the Law of Constant Extinction [132]. All dietary categories have approximately equivalent patterns of survival while ground dwelling taxa have a qualitatively higher probability of long duration. In comparison, the species-level survival curves for European mammals, both dietary (Fig. 4c) and locomotor categories (Fig. 4d), are qualitatively not semi-log linear which is not consistent with the Law of Constant Extinction. Diet qualitatively appears to have little effect on European mammal survival, while locomotor category appears to differentiate arboreal taxa from both ground dwelling and scansorial taxa.

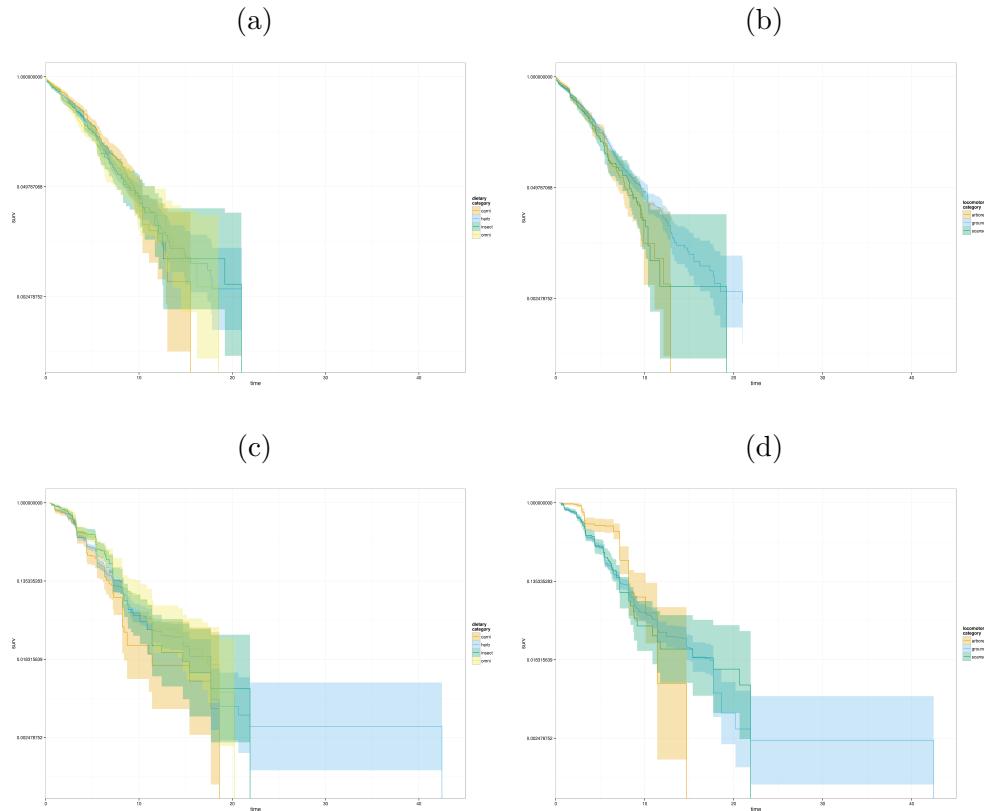


Figure 4: Nonparametric Kaplan-Meier species-level survival curves for North American and European mammals based on dietary category (a and c respectively, and locomotor category (b and d respectively). K-M curves are illustrated with 95% confidence intervals. The vertical axes are on a natural log scale.

These results are extremely preliminary and based solely on qualitative patterns present in the nonparametric K-M survival curves and without reference to estimated median survival times. Additionally, possible differences in survival based on body-size were not estimated. Also, no comparison has been made with the different climatic histories of both Continents.

664 By including all three time-independent covariates in a parametric modelling of survival  
framework it should be possible to better understand the underlying process behind survival.  
666 The inclusion of a third continent, South America, will also greatly improve the overall  
understanding of how extinction in mammals proceeds and how this may differ across  
environments.

### 668 **3.3 Community connectedness: global, regional, local**

#### **3.3.1 Questions**

670 How does the ratio between endemic and cosmopolitan taxa at a locality change over time? Is  
this pattern different between ecological categories? Does this pattern reflect global, regional,  
672 and/or local processes?

#### **3.3.2 Background and Predictions**

674 During the Cenozoic there was a global shift from a “hot house” environment to an “ice  
house” environment [139, 140]. This transition was accompanied by major shifts in global  
676 climatic envelopes and the reorganization of mammalian communities [5, 14, 35, 43, 62].  
For mammalian community connectedness there are two possible scenarios. First, while the  
678 environment was shifting, lineages may have adapted in place and overall trophic structure  
and community connectedness would have remained relatively constant through time, as  
680 observed during the Neogene of Europe [65]. Alternatively, species may have shifted ranges  
and changed the average set of taxa present at a locality which would be associated with  
682 non-stationary trophic structure and community connectedness.

Based on prior work, it is expected that the patterns of biogeographic community connected-  
684 ness for herbivorous taxa in a region would be most similar to that for all regional taxa and  
potentially “drive” the regional pattern, partially because on average this category represents  
686 the majority or plurality of taxa [64]. In contrast, community connectedness for carnivorous  
taxa is expected to remain constant over time or be correlated with herbivore patterns. Finally,  
688 omnivorous taxa are not expected to be correlated with the patterns of either herbivorous or  
carnivorous taxa and have either a relatively constant or random pattern of connectedness  
690 over time. These predictions are based on the differences in resilience and relationship to  
primary productivity, with herbivores being more resilient than carnivores and omnivores  
692 being random in their resilience [65]. Resilience is defined here as the ability for a taxon to  
increase in occupancy following a decline [65].

694 The Cenozoic global shift from closed, forested habitat in the Paleogene to open, savanna-like  
habitat during the Neogene would have greatly affected the possible distributions of both  
696 arboreal and ground dwelling taxa. Additionally, the timing of this environmental shift was  
different between continents [128, 129], so patterns of community connectedness may not be

698 globally uniform and instead reflect regional differences. Generally this transitions would cause  
699 forested environments to become increasingly patchier in distribution while transitioning  
700 from the Paleogene to the Neogene. The global prediction then is that there would have been  
701 a relative increase in  $E$  (Eq. 2) and code length accompanied by a decrease in  $BC$  (Eq. 4)  
702 and  $Occ$  (Eq. 3) in arboreal taxa over time. The opposite is expected for terrestrial taxa.

703 At a regional scale, North American community connectedness is expected to follow the  
704 global predictions described above because the vast amount of prior synthesis has focused  
705 on North America [3–5, 9, 10, 14, 35, 46, 47, 122, 123]. However, the effect of global climate  
706 change on North American diversity remains unresolved and controversial [5, 10, 14, 35],  
707 thus it is necessary to determine empirically when global versus regional versus local scale  
708 processes may have dominated and how that may have changed over time.

709 The European mammalian fossil record is also well studied, though research has primarily  
710 focused on the Neogene [64, 65, 78, 101–103]. An important aspect about the European  
711 record is that during the Neogene there was little shift in relative dietary category abundance  
712 [65] and that the patterns within herbivores (browse–graze transition) were mostly driven  
713 by abundant, cosmopolitan taxa [64]. It is predicted then that herbivores will demonstrate  
714 the same patterns of community connectedness as Europe as a whole, while omnivores and  
715 carnivores will be different from that of herbivores and may demonstrate random or constant  
716 patterns of community connectedness through time.

717 Patterns of community connectedness for South American mammalian fauna are comparatively  
718 less synthesized than those of North American and Europe. Instead, cross-continental  
719 dynamics between North and South America during the Neogene are much more studied [85].  
720 The South American mammalian faunal record reflects two distinct biotic provinces between  
721 the North and the South [37, 82, 83, 94]. Because of this, it is expected that South America  
722 will have a different pattern of community connectedness than either North America or Europe.  
723 Also, there is an expected dramatic increase occupancy in land-dwelling herbivores relative  
724 to arboreal and scansorial taxa related to the aridification of high-latitude South America.  
725 Additionally, because of this strong biome distinction, it is predicted that provinciality will  
726 be high but remain constant over time.

### 3.3.3 Proposed research

727 In order to estimate changes in community connectedness during the Cenozoic I will be using  
728 the network-based approach described above (Section 1.2). Biogeographic networks will be  
729 constructed for each region (North America, Europe, South American) between species and  
730 localities defined as 2x2 latitude–longitude grid cells from an equal-area map projection.  
731 Networks will be made for every 2 My span of the Cenozoic. This bin width was chosen to in  
732 order to maximize the chance that two localities are present at the same time. Networks will  
733 also be constructed for subsets of taxa defined by dietary and locomotor categories order to  
734 compare patterns both within and between categories, as well as to the combined regional and

736 global patterns. Because previous studies of mammalian occurrence patterns have restricted  
737 analysis to large bodied and well studied groups such as Primates and Artiodactyls in order to  
738 account for potential sampling and taxonomic biases, analysis will be done using all available  
739 taxa and with a restricted sample of just major groups in order to observe any differences in  
740 patterns of community connectedness. The data necessary to complete this study is the same  
741 as for the above analysis of mammalian survival (Section 3.2).

742 The degree of phylogenetic similarity between taxa at a locality may play an important  
743 role in community structuring [137]. For example, closely related taxa may be repulsed  
744 “repulsed” due to competitive exclusion or “clumped” because of environmental filtering.  
745 While it is infeasible to create an explicit phylogenetic hypothesis for all taxa sampled on all  
746 continents, almost all taxa have some hierarchical taxonomic information. Using taxonomy as  
747 the structure of an information phylogeny, it should be possible to estimate the distribution  
748 of phylogenetic similarity across localities.

749 For each locality, an informal phylogeny will be constructed based solely on available taxonomic  
750 information such as order, family, and genus assignments with each of these levels being a  
751 completely unresolved polytomy. Using this informal phylogeny, a number of measures of  
752 phylogenetic similarity can be estimated. For example the relative mean pairwise distance  
753 between all taxa at a locality [137] or the related phylogenetic species variability of a single  
754 locality [51]. These values calculated for all localities can then be used as a partial correlates  
755 or covariates when modeling changes in community connectedness.

756 As with the Permian brachiopods (Section 2.3), patterns of community connectedness will be  
757 compared both within and between ecological categories. Additionally, the correspondence  
758 of changes in environmental conditions and community connectedness will also be analysed.  
759 The approach and methodology to accomplish these analyses is currently under development.  
760 Additionally, the possibility of integrating locality–locality distance or some other measure  
761 of topology will be explored, especially how this relates to code length and provinciality in  
762 general.

### 3.3.4 Preliminary results

763 Preliminary analysis was done using only the occurrence information of both North American  
764 and European fossil mammals available in the PBDB. Both regions have qualitatively different  
765 patterns of community connectedness, primarily during the Paleogene (Fig. 5). Almost all  
766 four of the summary statistics are extremely volatile over the Cenozoic, especially for Europe.  
767 However, some interesting qualitative patterns are present. Importantly, the network summary  
768 statistics will be calculated only with reference to the localities at which arboreal taxa occur,  
769 and not all possible localities occurring at a specific time.

770 There is a qualitative decrease in *Occ* in Europe until approximately the start of the Neo-  
771 gene (approximately 23 My), indicating that the average taxon is becoming generally less  
772 cosmopolitan over time. In contrast, North American *Occ* is qualitatively stationary over the

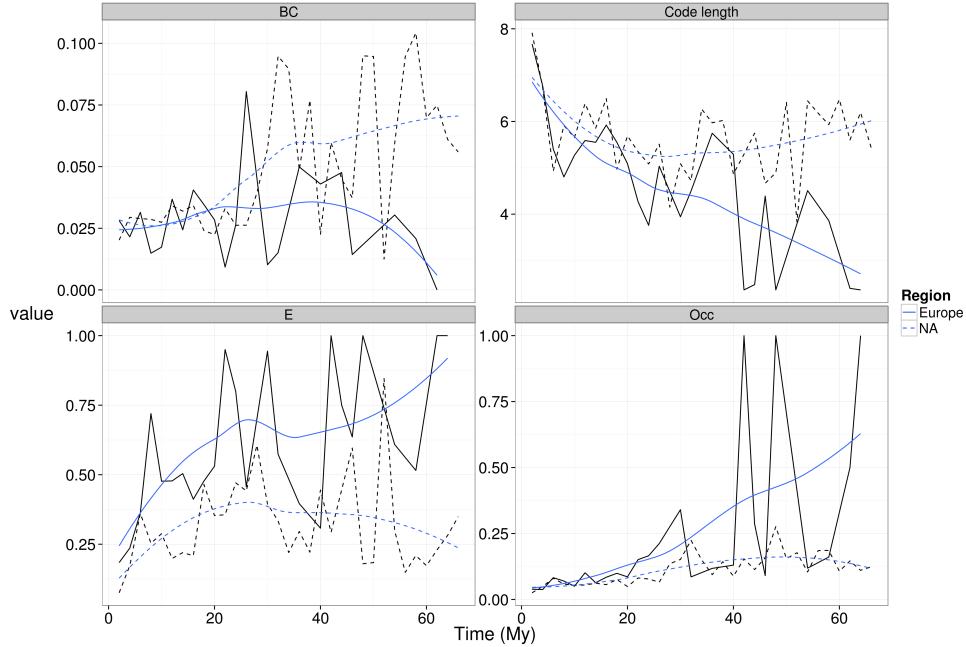


Figure 5: Biogeographic network summary statistics for mammalian communities in North America (dashed line) and Europe (solid line). The summary statistics are, clockwise from top left: biogeographic connectedness (BC), code length, average relative locality occupancy per taxon (Occ), and average relative number of endemic taxa per locality (E). Blue lines are generalized additive model smooths and are presented to illustrate the overall pattern for each region.

entire Cenozoic and almost always lower than that observed for Europe. This means that, on average, North American taxa are present in very few localities at any given point in time.

In Europe there is a qualitative rise in *BC* in the first few million years of the Cenozoic, but afterwards remains relatively stationary meaning that the average proportion of shared taxa remained qualitatively stationary. In comparison, North American *BC* remains stationary with a greater amount of shared taxa than Europe for the first half of the Cenozoic followed by a decrease and another plateau at the end of the Cenozoic.

In Europe, there is a over all qualitative decrease in *E* while in North America there is a qualitatively constant *E* over the Cenozoic with a slight decrease in the Neogene. As discussed above, *E* is a measure of relative uniqueness of a locality on average. Qualitatively, North America retained approximately the same amount of site uniqueness through out the Cenozoic. While the pattern of the European record shows a qualitatively nonmonotonic decrease in locality uniqueness.

The code length of European biogeographic networks increases qualitatively over the entire Cenozoic, while code length of North American networks remains relatively constant until the Neogene when there is a qualitative increase. Initial interpretation of these results indicates

790 that North America maintains a relatively stationary degree of provinciality while Europe has a qualitatively decreasing degree of provinciality.

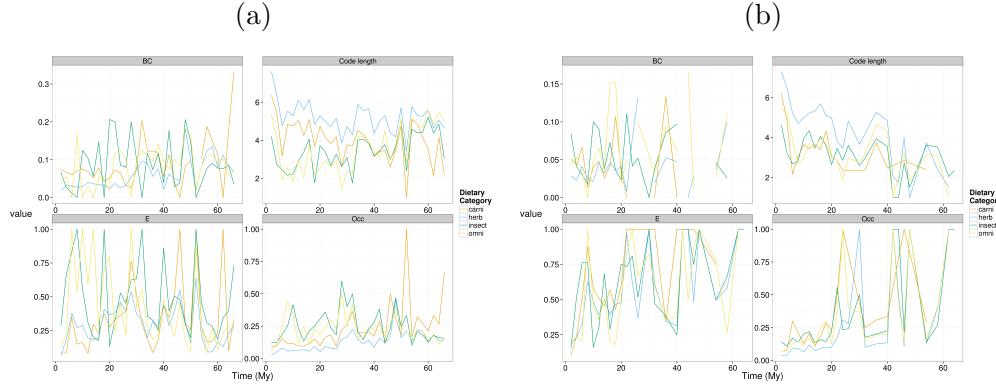


Figure 6: Time series of summary statistics for biogeographic networks determined by dietary category for North America (a) and Europe (b). The summary statistics are, clockwise from top left: biogeographic connectedness (BC), code length, average relative locality occupancy per taxon (Occ), and average relative number of endemic taxa per locality (E).

792 When taxa are separated by dietary categories, the amount of noise associated with each  
 793 statistic increases greatly (Fig. 6). In North America, *BC*, while variable, appears to qualita-  
 794 tively demonstrate no net change. Carnivores and herbivores to qualitatively become less  
 795 volatile during the Neogene compared to the Paleogene (Fig. 6a). *BC* for Europe is also very  
 796 volatile, though impossible to measure for dietary categories individually for much of the  
 797 Paleogene (Fig. 6b).

798 Code length for North American qualitatively shows a stationary pattern with an up-tick in  
 799 the Recent and a major drop at approximately 50-55 My (Fig. 6a). Additionally, herbivore  
 800 and carnivore patterns appear qualitatively similar. In comparison, the European record for  
 801 code length shows a qualitatively slight increase over the entire Cenozoic (Fig. 6b). Also, the  
 802 patterns of herbivore and carnivores appear qualitatively less similar than for North America.  
 803 For both Europe and North America, herbivores have the over all highest code length. In  
 804 North America, carnivores arguably have the second highest code length. In all other cases,  
 805 the ranks are qualitatively ambiguous.

806 *E* for North American appears to qualitatively have two categories (Fig. 6a). Herbivore and  
 807 carnivore patterns are qualitatively stationary and low during the Neogene, while the omnivore  
 808 and insectivore patters are qualitatively more variable and higher during the Neogene. In  
 809 comparison, all four categories of European mammals demonstrate a slight decrease during  
 810 the Cenozoic (Fig. 6b).

811 For North America, *Occ* are qualitatively stationary throughout the Cenozoic with one spike  
 812 in carnivore *Occ* at approximately 50-55 My (Fig. 6a). In contrast, European values are  
 813 highly volatile throughout the Paleogene and then less volatile during the Neogene (Fig. 6b).

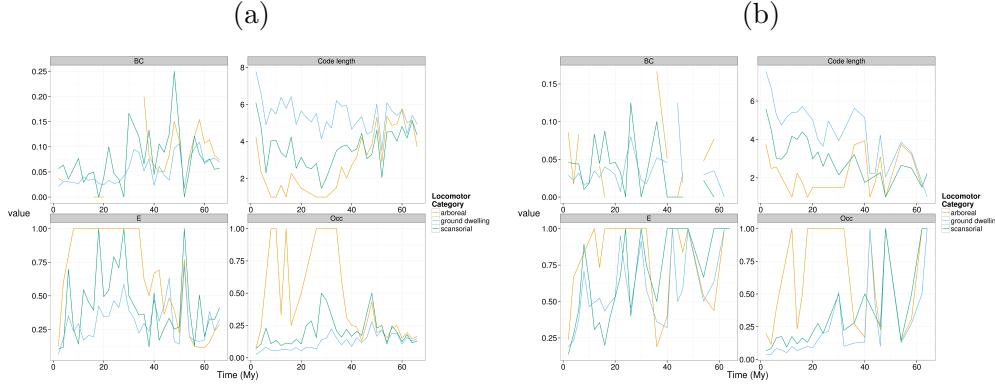


Figure 7: Time series of summary statistics for biogeographic networks determined by locomotor category for North America (a) and Europe (b). The summary statistics are, clockwise from top left: biogeographic connectedness (BC), code length, average relative locality occupancy per taxon (Occ), and average relative number of endemic taxa per locality (E).

When taxa are separated by locomotor category, there is qualitatively less noise than is the case for by dietary category (Fig. 7). *BC* for North America has qualitative differences between each of the three categories (Fig. 7a). Arboreal taxa can only be measured for *BC* predominately during the Paleogene where there is no qualitative pattern beyond high variance. Scansorial taxa have a qualitative decline in volatility and was stationary during the Neogene. European values of *BC* were generally more volatile and very difficult to measure during the Paleogene because of the paucity of geographically spaced localities (Fig. 7b). Qualitatively, values of *BC* for scansorial taxa are more volatile than for ground dwelling taxa.

For North American values of code length, there are a few clear qualitative patterns (Fig. 7a). Ground dwelling taxa have generally the highest code length values, followed by scansorial and arboreal taxa. Interestingly, all three of these categories have almost identical code length values until approximately 50 My. Following this, arboreal taxa have a qualitative decrease in code length, while scansorial taxa are qualitatively stationary with a slight decrease, and ground dwelling taxa have a slight increase though are mostly stationary. European code length values show a general increase during the entire Cenozoic, though this is mostly confined to scansorial and ground dwelling taxa (Fig. 7b).

The *E* series for North America demonstrates qualitatively distinct patterns for the three locomotor categories (Fig. 7a). *E* increases dramatically for arboreal taxa, has a moderate increase for scansorial taxa, and is qualitatively stationary for ground dwelling taxa during the Cenozoic. In comparison for Europe, values of *E* are generally high throughout the entire Cenozoic and vary with much greater volatility (Fig. 7b). Qualitatively there is a decrease in *E* for ground dwelling and scansorial taxa during the Neogene.

Values of *Occ* for both North America and Europe show respectively qualitatively similar patterns to patterns of *E*, though are less volatile. *Occ* increases in North American arboreal taxa at approximately 40 My years ago while both scansorial and ground dwelling taxa are qualitatively stationary (Fig. 7a). The pattern of *Occ* for scansorial taxa appears to qualitatively be a more exaggerated version of the pattern for ground dwelling taxa. All three appear correlated during the earliest Cenozoic. As with *E*, European patterns of *Occ* are volatile, particularly during the early Cenozoic (Fig. 7b). At approximately 40 My, patterns of *Occ* become less volatile and qualitatively decrease for ground dwelling and scansorial taxa. In comparison, *Occ* values for arboreal taxa become qualitatively much higher during the late Cenozoic with a massive decrease near the Recent.

These analyses will be greatly improved by varying locality “size”, comparison with South American patterns, comparison of major orders, and other ideas stated above (Section 3.3.3). Additionally, quantitatively analysis of these patterns and what correlations might exist, especially in a phylogenetic context, are necessary in order to better understand what processes might dominate and when.

852 **4 Synthesis of proposed research**

Underlying all of the above is a foundational question in paleobiology: why do certain taxa  
854 go extinct while others do not? In the context of evolutionary paleoecology, this question can  
be rephrased as “how do the set of all biotic–biotic and biotic–abiotic interactions a taxon  
856 experiences over time (i.e. adaptive zone 122) affect extinction risk?” Related to this is the  
Law of Constant Extinction which states that extinction risk for a given adaptive zone is  
858 taxon–age independent [132]. It is asserted that the Law of Constant Extinction only holds  
during periods of relatively constant environment, even though this was not the context for  
860 the initial observation [77, 132], which can be interpreted as the set of dominant non-organism  
mediated processes do not fluctuate or fluctuate in a known manner. By understanding which  
862 non-organism mediated processes may be shaping the environment (set of all possible biotic  
and abiotic interactors) and how they change over time and phrasing analysis of extinction  
864 in this context, it may be possible to “test” the Law of Constant Extinction.

The two studies proposed above (Sections 2.2 and 3.2) investigate how organismal traits  
866 potentially related to environmental preference affect extinction rate. In effect, these traits  
may determine the “bounds” of a taxon’s adaptive zone by limiting the total set of interactions  
868 to just those for which the taxon is adapted. The other two proposed studies (Sections 2.3 and  
3.3) aim to estimate what non-organism mediated processes (global, regional, and/or local)  
870 may be dominate in shaping the environment and the related set of adaptive zones. Between  
these studies, as well the use of two disparate groups, it should be possible to determine  
872 when, what, and if certain variables matter for survival and, potentially, how they matter.

## 5 Timeline

- <sup>874</sup> Spring/Summer 2014
- Evolution Meeting: preliminary brachiopod survival results
- <sup>876</sup> • South American fossil mammal data from American Museum of Natural History collections
- <sup>878</sup> Fall 2014/Winter 2015
- GSA: survivorship simulation for anagenesis and sampling
  - Doctoral Dissertation Improvement Grant
- Spring/Summer 2015
- Evolution Meeting: mammalian survivorship analysis for North America and Europe
  - write and submit survivorship simulation paper
  - possible South American fossil mammal data from American Museum of Natural History collections
- <sup>886</sup> Fall 2015/Winter 2016
- SVP: mammalian biogeographic connectedness
  - write and submit mammal connectedness paper
- Spring/Summer 2016
- Evolution Meeting: brachiopod survival analysis
  - write and submit brachiopod survival paper
- <sup>892</sup> Fall 2016/Winter 2017
- GSA: brachiopod community connectedness
  - write and submit mammal survival paper
- Spring/Summer 2017
- Evolution Meeting: survival and communities together
  - write and submit brachiopod community paper
  - write and review/philosophy paper
  - **Defend**

900    6    Bibliography

- 902 [1] R. R. Alexander. Generic longevity of articulate brachiopods in relation to the mode of  
stabilization on the substrate. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 21:  
209–226, 1977.
- 904 [2] W. D. Allmon. Taxic evolutionary paleoecology and the ecological context of macroevo-  
lutionary change. *Evolutionary Ecology*, 8(2):95–112, Mar. 1994. ISSN 0269-7653. doi:  
10.1007/BF01238244.
- 908 [3] J. Alroy. Constant extinction, constrained diversification, and uncoordinated stasis  
in North American mammals. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 127:  
285–311, 1996.
- 910 [4] J. Alroy. Cope’s rule and the dynamics of body mass evolution in North American  
fossil mammals. *Science*, 280:731–734, 1998. doi: 10.1126/science.280.5364.731.
- 912 [5] J. Alroy, P. L. Koch, and J. C. Zachos. Global climate change and North American  
mammalian evolution. *Paleobiology*, 26(1981):259–288, 2000.
- 914 [6] L. W. Alvarez. Experimental evidence that an asteroid impact led to the ex-  
tinction of many species 65 million years ago. *Proceedings of the National  
Academy of Sciences of the United States of America*, 80(2):627–42, Jan. 1983.  
ISSN 0027-8424. URL [http://www.ncbi.nlm.nih.gov/article/abstract.fcgi?  
artid=393431&tool=pmcentrez&rendertype=abstract](http://www.ncbi.nlm.nih.gov/article/abstract.fcgi?artid=393431&tool=pmcentrez&rendertype=abstract).
- 920 [7] B. M. Anderson, D. Pisani, A. I. Miller, and K. J. Peterson. The environmental affinities  
of marine higher taxa and possible biases in their first appearances in the fossil record.  
*Geology*, 39(10):971–974, Sept. 2011. ISSN 0091-7613. doi: 10.1130/G32413.1. URL  
<http://geology.gsapubs.org/cgi/doi/10.1130/G32413.1>.
- 924 [8] N. W. Archbold and G. R. Shi. Permian brachiopod faunas of Western Australia:  
Gondwanan-Asian relationships and Permian climate. *Journal of Southeast Asian Earth  
Sciences*, 11(3):207–215, 1995.
- 926 [9] C. Badgley and J. A. Finarelli. Diversity dynamics of mammals in relation to tec-  
tonic and climatic history: comparison of three Neogene records from North America.  
*Paleobiology*, 39(3):373–399, Apr. 2013. ISSN 0094-8373. doi: 10.1666/12024.
- 930 [10] A. D. Barnosky. Distinguishing the effects of the Red Queen Court Jester on Miocene  
Mammal Evolution in the Northern Rocky Mountains. *Journal of Vertebrate Paleontology*,  
21(1):172–185, 2001.
- 932 [11] T. K. Baumiller. Survivorship analysis of Paleozoic Crinoidea: effect of filter morphology  
on evolutionary rates. *Paleobiology*, 19(3):304–321, 1993.
- 934 [12] A. Birand, A. Vose, and S. Gavrilets. Patterns of species ranges, speciation, and

- extinction. *The American naturalist*, 179(1):1–21, Jan. 2012. ISSN 1537-5323. doi: 10.1086/663202. URL <http://www.ncbi.nlm.nih.gov/pubmed/22173457>.
- [13] L. P. Birgenheier, T. D. Frank, C. R. Fielding, and M. C. Rygel. Coupled carbon isotopic and sedimentological records from the Permian system of eastern Australia reveal the response of atmospheric carbon dioxide to glacial growth and decay during the late Palaeozoic Ice Age. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 286(3-4): 178–193, 2010. URL <http://dx.doi.org/10.1016/j.palaeo.2010.01.008>.
- [14] J. L. Blois and E. A. Hadly. Mammalian Response to Cenozoic Climatic Change. *Annual Review of Earth and Planetary Sciences*, 37(1):181–208, May 2009. ISSN 0084-6597. doi: 10.1146/annurev.earth.031208.100055.
- [15] D. J. Bottjer and D. Jablonski. Paleoenvironmental patterns in the evolution of Post-Paleozoic benthic marine invertebrates. *Palaios*, 3(6):540–560, 1988.
- [16] S. Brin and L. Page. The anatomy of a large-scale hypertextual Web search engine. *Computer Networks and ISDN Systems*, 30(1-7):107–117, Apr. 1998. ISSN 01697552. doi: 10.1016/S0169-7552(98)00110-X. URL <http://linkinghub.elsevier.com/retrieve/pii/S016975529800110X>.
- [17] J. H. Brown and B. A. Maurer. Evolution of species assemblages: effects of energetic constraints and species dynamics on the diversification of the North American avifauna. *The American Naturalist*, 130(1):1–17, 1987.
- [18] K. P. Burnham and D. R. Anderson. *Model selection and multi-model inference: a practical information-theoretic approach*. Springer, New York, 2nd edition, 2002.
- [19] M. E. Clapham and D. J. Bottjer. Permian marine paleoecology and its implications for large-scale decoupling of brachiopod and bivalve abundance and diversity during the Lopingian (Late Permian). *Palaeogeography, Palaeoclimatology, Palaeoecology*, 249 (3-4):283–301, June 2007. ISSN 00310182. doi: 10.1016/j.palaeo.2007.02.003. URL <http://linkinghub.elsevier.com/retrieve/pii/S0031018207000600>.
- [20] M. E. Clapham and D. J. Bottjer. Prolonged Permian Triassic ecological crisis recorded by molluscan dominance in Late Permian offshore assemblages. *Proceedings of the National Academy of Sciences of the United States of America*, 104(32):12971–5, Aug. 2007. ISSN 0027-8424. doi: 10.1073/pnas.0705280104. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1941817&tool=pmcentrez&rendertype=abstract>.
- [21] M. E. Clapham and N. P. James. Paleoenvironment Of Early-Middle Permian Marine Communities In Eastern Australia: Response To Global Climate Change In the Aftermath Of the Late Paleozoic Ice Age. *Palaios*, 23(11):738–750, Nov. 2008. ISSN 0883-1351. doi: 10.2110/palo.2008.p08-022r. URL <http://palaios.sepmonline.org/cgi/doi/10.2110/palo.2008.p08-022r>.

- [22] M. E. Clapham and N. P. James. Regional-scale marine faunal change in Eastern Australia during Permian climate fluctuations and its relationship to local community restructuring. *Palaios*, 27:627–635, Oct. 2012. ISSN 0883-1351. doi: 10.2110/palo.2012.p12-003r. URL <http://palaios.sepmonline.org/cgi/doi/10.2110/palo.2012.p12-003r>.
- [23] M. E. Clapham, D. J. Bottjer, C. M. Powers, N. Bonuso, M. L. Fraiser, P. J. Marenco, S. Q. Dornbos, and S. B. Pruss. Assessing the Ecological Dominance of Phanerozoic Marine Invertebrates. *Palaios*, 21(5):431–441, Oct. 2006. ISSN 0883-1351. doi: 10.2110/palo.2005.P05-017R. URL <http://palaios.sepmonline.org/cgi/doi/10.2110/palo.2005.P05-017R>.
- [24] A. Clauset and D. H. Erwin. The evolution and distribution of species body size. *Science*, 321:399–401, 2008.
- [25] J. Damuth. Home range, home range overlap, and species energy use among herbivorous mammals. *Biological Journal of the Linnean Society*, 15:185–193, 1979.
- [26] J. Damuth. Population density and body size in mammals. *Nature*, 290:699–700, 1981. URL <http://www.jstor.org/stable/10.2307/2461771> <http://www.nature.com/nature/journal/v290/n5808/abs/290699a0.html>.
- [27] A. D. Davidson, A. G. Boyer, H. Kim, S. Pompa-Mansilla, M. J. Hamilton, D. P. Costa, G. Ceballos, and J. H. Brown. Drivers and hotspots of extinction risk in marine mammals. *Proceedings of the National Academy of Sciences*, 109(9):3395–400, Feb. 2012. ISSN 1091-6490. doi: 10.1073/pnas.1121469109. URL <http://www.ncbi.nlm.nih.gov/articlerender.fcgi?artid=3295301&tool=pmcentrez&rendertype=abstract>.
- [28] R. Diestel. *Graph Theory*. Springer, Berlin, 3rd edition, 2005.
- [29] J. M. Drake. Tail probabilities of extinction time in a large number of experimental populations. *Ecology*, page 140206083444001, Feb. 2014. ISSN 0012-9658. doi: 10.1890/13-1107.1. URL <http://www.esajournals.org/doi/abs/10.1890/13-1107.1>.
- [30] C. R. Fielding, A. J. Falkner, and S. G. Scott. Fluvial response to foreland basin overfilling; the Late Permian Rangal Coal Measures in the Bowen Basin, Queensland, Australia. *Sedimentary Geology*, 85:475–497, 1993.
- [31] C. R. Fielding, K. L. Bann, J. A. Maceachern, S. C. Tye, and B. G. Jones. Cyclicality in the nearshore marine to coastal, Lower Permian, Pebbley Beach Formation, southern Sydney Basin, Australia: a record of relative sea-level fluctuations at the close of the Late Palaeozoic Gondwanan ice age. *Sedimentology*, 53(2):435–463, 2006. URL <http://doi.wiley.com/10.1111/j.1365-3091.2006.00770.x>.
- [32] C. R. Fielding, T. D. Frank, L. P. Birgenheier, M. C. Rygel, A. T. Jones, and J. Roberts. Stratigraphic record and facies associations of the late Paleozoic ice age in eastern

- 1010 Australia (New South Wales and Queensland). *Geological Society of America Special*  
1011 *Papers*, 441:41–57, 2008. doi: 10.1130/2008.2441(03).
- 1012 [33] C. R. Fielding, T. D. Frank, and J. L. Isbell. The late Paleozoic ice age – a review of  
1013 current understanding and synthesis of global climate patterns. *Geological Society of*  
1014 *America Special Papers*, 441:343–354, 2008. doi: 10.1130/2008.2441(24).
- 1016 [34] C. R. Fielding, T. D. Frank, J. L. Isbell, L. C. Henry, and E. W. Domack. Stratigraphic  
1017 signature of the late Palaeozoic Ice Age in the Parameener Supergroup of Tasmania,  
1018 SE Australia, and inter-regional comparisons. *Palaeogeography, Palaeoclimatology,*  
1019 *Palaeoecology*, 298(1-2):70–90, Dec. 2010. ISSN 00310182. doi: 10.1016/j.palaeo.2010.05.  
1020 023. URL <http://linkinghub.elsevier.com/retrieve/pii/S003101821000310X>.
- 1022 [35] B. Figueirido, C. M. Janis, J. A. Pérez-Claros, M. De Renzi, and P. Palmqvist. Cenozoic  
1023 climate change influences mammalian evolutionary dynamics. *Proceedings of the*  
1024 *National Academy of Sciences*, 109(3):722–727, Jan. 2012. ISSN 1091-6490. doi:  
1025 10.1073/pnas.1110246108.
- 1026 [36] S. Finnegan, J. L. Payne, and S. C. Wang. The Red Queen revisited: reevaluating the  
1027 age selectivity of Phanerozoic marine genus extinctions. *Paleobiology*, 34(3):318–341,  
1028 Sept. 2008. ISSN 0094-8373. doi: 10.1666/07008.1. URL <http://www.bioone.org/doi/abs/10.1666/07008.1>.
- 1029 [37] J. J. Flynn and A. R. Wyss. Recent advances in South American mammalian paleontology.  
1030 *Trends in ecology & evolution*, 13(11):449–54, Nov. 1998. ISSN 0169-5347. URL  
1031 <http://www.ncbi.nlm.nih.gov/pubmed/21238387>.
- 1032 [38] M. Foote. Survivorship analysis of Cambrian and Ordovician Trilobites. *Paleobiology*,  
1033 14(3):258–271, 1988.
- 1034 [39] M. Foote. Estimating taxonomic durations and preservation probability. *Paleobiology*,  
1035 23(3):278–300, 1997. URL <http://www.psjournals.org/paleoonline/?request=get-abstract&issn=0094-8373&volume=23&issue=3&page=278>.
- 1036 [40] M. Foote. Substrate affinity and diversity dynamics of Paleozoic marine animals.  
1037 *Paleobiology*, 32(3):345–366, Sept. 2006. ISSN 0094-8373. doi: 10.1666/05062.1. URL  
1038 <http://www.bioone.org/doi/abs/10.1666/05062.1>.
- 1040 [41] M. Foote and A. I. Miller. Determinants of early survival in marine animal genera.  
1041 *Paleobiology*, 39(2):171–192, Mar. 2013. ISSN 0094-8373. doi: 10.1666/12028. URL  
1042 <http://www.bioone.org/doi/abs/10.1666/12028>.
- 1044 [42] M. Foote and D. Raup. Fossil preservation and the stratigraphic ranges of taxa.  
1045 *Paleobiology*, 22(2):121–140, 1996. URL <http://www.psjournals.org/paleoonline/?request=get-abstract&issn=0094-8373&volume=022&issue=02&page=0121>.
- [43] M. Fortelius, J. Eronen, J. Jernvall, L. Liu, D. Pushkina, J. Rinne, A. Tesakov, I. Vis-

- 1046 lobokova, Z. Zhang, and L. Zhou. Fossil mammals resolve regional patterns of Eurasian  
climate change over 20 million years. *Evolutionary Ecology Research*, 4:1005–1016, 2002.
- 1048 [44] T. D. Frank, S. G. Thomas, and C. R. Fielding. On Using Carbon and Oxygen Isotope  
Data from Glendonites as Paleoenvironmental Proxies: A Case Study from the Permian  
1050 System of Eastern Australia. *Journal of Sedimentary Research*, 78(11):713–723, Nov.  
2008. ISSN 1527-1404. doi: 10.2110/jsr.2008.081. URL <http://jsedres.sepmonline.org/cgi/doi/10.2110/jsr.2008.081>.
- 1052 [45] K. J. Gaston. Geographic range limits: achieving synthesis. *Proceedings. Biological  
1054 sciences / The Royal Society*, 276(1661):1395–406, Apr. 2009. ISSN 0962-8452. doi:  
10.1098/rspb.2008.1480. URL <http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2677218/>&tool=pmcentrez&rendertype=abstract.
- 1056 [46] G. Gunnell, M. Morgan, M. C. Mass, and P. D. Gingerich. Comparative paleoecology  
1058 of Paleogene and Neogene mammalian faunas: trophic structure and composition.  
*Palaeogeography, Palaeoclimatology, Palaeoecology*, 115:265–286, 1995. URL <http://www.sciencedirect.com/science/article/pii/0031018294001150>.
- 1060 [47] E. A. Hadly and B. A. Maurer. Spatial and temporal patterns of species diversity  
1062 in montane mammal communities of western North America. *Evolutionary Ecology  
Research*, 3:477–486, 2001.
- 1064 [48] P. G. Harnik, C. Simpson, and J. L. Payne. Long-term differences in extinction  
risk among the seven forms of rarity. *Proceedings of the Royal Society B: Biological  
1066 Sciences*, (October), Oct. 2013. ISSN 0962-8452. doi: 10.1098/rspb.2012.1902. URL  
<http://rsb.rsa.org/doi/10.1098/rspb.2012.1902>.
- 1068 [49] S. P. Hawley, R. A. Glen, and C. J. Baker. Newcastle Coalfield Regional Geology 1:100  
000. Technical report, Geological Survey of New South Wales, Sydney, 1995.
- 1070 [50] N. A. Heim and S. E. Peters. Regional environmental breadth predicts geographic range  
and longevity in fossil marine genera. *PLoS one*, 6(5):e18946, Jan. 2011. ISSN 1932-  
1072 6203. doi: 10.1371/journal.pone.0018946. URL <http://www.ncbi.nlm.nih.gov/pmc/articles/PMC3137268/>&tool=pmcentrez&rendertype=abstract.
- 1074 [51] M. R. Helmus, T. J. Bland, C. K. Williams, and A. R. Ives. Phylogenetic Measures  
of Biodiversity. *The American naturalist*, 169(3), Jan. 2007. ISSN 1537-5323. doi:  
10.1086/511334. URL <http://www.ncbi.nlm.nih.gov/pubmed/17230400>.
- 1076 [52] D. Huffman. A Method for the Construction of Minimum-Redundancy Codes.  
*Proceedings of the IRE*, 40(9):1098–1101, Sept. 1952. ISSN 0096-8390. doi: 10.  
1109/JRPROC.1952.273898. URL <http://ieeexplore.ieee.org/lpdocs/epic03/wrapper.htm?arnumber=4051119>.
- 1078 [53] C. M. Hurvich and C.-L. Tsai. Regression and time series model se-  
lection in small samples. *Biometrika*, 76(2):297–307, 1989. URL  
1080 <http://biomet.oxfordjournals.org/content/76/2/297>.

<http://biomet.oxfordjournals.org/content/76/2/297.shortpapers2://publication/uuid/FC795396-A722-468D-AF8E-4E1F7C97998C>.

- [54] D. Jablonski. Background and mass extincitons: the alternation of macroevolutionary regimes. *Science*, 231(4734):129–133, 1986.
  - [55] D. Jablonski. Heritability at the species level: analysis of geographic ranges of cretaceous mollusks. *Science*, 238(4825):360–363, Oct. 1987. ISSN 0036-8075. doi: 10.1126/science.238.4825.360. URL <http://www.ncbi.nlm.nih.gov/pubmed/17837117>.
  - [56] D. Jablonski. Species Selection: Theory and Data. *Annual Review of Ecology, Evolution, and Systematics*, 39(1):501–524, Dec. 2008. ISSN 1543-592X. doi: 10.1146/annurev.ecolsys.39.110707.173510. URL <http://www.annualreviews.org/doi/abs/10.1146/annurev.ecolsys.39.110707.173510>.
  - [57] D. Jablonski and D. J. Bottjer. Environmental patterns in the origins of higher taxa: the post-paleozoic fossil record. *Science (New York, N.Y.)*, 252(5014):1831–3, June 1991. ISSN 0036-8075. doi: 10.1126/science.252.5014.1831. URL <http://www.ncbi.nlm.nih.gov/pubmed/17753259>.
  - [58] D. Jablonski and G. Hunt. Larval ecology, geographic range, and species survivorship in Cretaceous mollusks: organismic versus species-level explanations. *The American naturalist*, 168(4):556–64, Oct. 2006. ISSN 1537-5323. doi: 10.1086/507994. URL <http://www.ncbi.nlm.nih.gov/pubmed/17004227>.
  - [59] D. Jablonski and R. a. Lutz. Larval Ecology of Marine Benthic Invertebrates: Paleo-biological Implications. *Biological Reviews*, 58(1):21–89, Feb. 1983. ISSN 1464-7931. doi: 10.1111/j.1469-185X.1983.tb00380.x. URL <http://doi.wiley.com/10.1111/j.1469-185X.1983.tb00380.x>.
  - [60] D. Jablonski and K. Roy. Geographical range and speciation in fossil and living molluscs. *Proceedings. Biological sciences / The Royal Society*, 270(1513):401–6, Feb. 2003. ISSN 0962-8452. doi: 10.1098/rspb.2002.2243. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=1691247&tool=pmcentrez&rendertype=abstract>.
  - [61] D. Jablonski, J. J. Sepkoski, D. J. Bottjer, and P. M. Sheehan. Onshore-offshore patterns in the evolution of Phanerozoic shelf communities. *Science*, 222(4628):1123–1125, 1983.
  - [62] C. M. Janis. Tertiary mammal evolution in the context of changing climates, vegetation, and tectonic events. *Annual Review of Ecology and Systematics*, 24:467–500, 1993.
  - [63] C. M. Janis, J. Damuth, and J. M. Theodor. Miocene ungulates and terrestrial primary productivity: where have all the browsers gone? *Proceedings of the National Academy of Sciences*, 97(14):7899–904, July 2000. ISSN 0027-8424. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=16642&tool=pmcentrez&rendertype=abstract>.
  - [64] J. Jernvall and M. Fortelius. Common mammals drive the evolutionary increase of

- 1120        hypsodonty in the Neogene. *Nature*, 417(6888):538–40, May 2002. ISSN 0028-0836. doi:  
1122        10.1038/417538a.
- 1124 [65] J. Jernvall and M. Fortelius. Maintenance of trophic structure in fossil mammal  
communities: site occupancy and taxon resilience. *American Naturalist*, 164(5):614–624,  
Nov. 2004. ISSN 1537-5323. doi: 10.1086/424967.
- 1126 [66] M. B. Johansen. Adaptive radiation, survival and extinction of brachiopods in  
the northwest European upper cretaceous-lower paleocene chalk. *Palaeogeography,  
Palaeoclimatology, Palaeoecology*, 74(3-4):147–204, Nov. 1989. ISSN 00310182. doi:  
1128 10.1016/0031-0182(89)90060-6. URL [http://linkinghub.elsevier.com/retrieve/  
pii/0031018289900606](http://linkinghub.elsevier.com/retrieve/pii/0031018289900606).
- 1130 [67] C. N. Johnson. Determinants of loss of mammal species during the Late Quaternary  
'megafauna' extinctions: life history and ecology, but not body size. *Proceedings of the  
Royal Society B: Biological Sciences*, 269:2221–2227, 2002. doi: 10.1098/rspb.2002.2130.
- 1132 [68] A. T. Jones, T. D. Frank, and C. R. Fielding. Cold climate in the eastern  
1134 Australian mid to late Permian may reflect cold upwelling waters. *Palaeogeography,  
Palaeoclimatology, Palaeoecology*, 237(2-4):370–377, 2006. URL  
1136 <http://linkinghub.elsevier.com/retrieve/pii/S0031018205007327>papers2:  
//publication/doi/10.1016/j.palaeo.2005.12.009.
- 1138 [69] D. Kendall. On the generalized "birth-and-death" process. *The Annals of Mathe-  
1140 matical Statistics*, 19(1):1–15, 1948. URL [http://projecteuclid.org/euclid.aoms/  
1177730285](http://projecteuclid.org/euclid.aoms/1177730285).
- 1142 [70] W. Kiessling and M. Aberhan. Environmental determinants of marine benthic biodi-  
versity dynamics through Triassic Jurassic time. *Paleobiology*, 33(3):414–434, 2007.
- 1144 [71] W. Kiessling, M. Aberhan, B. Brenneis, and P. J. Wagner. Extinction trajec-  
1146 tories of benthic organisms across the TriassicJurassic boundary. *Palaeogeography,  
Palaeoclimatology, Palaeoecology*, 244(1-4):201–222, Feb. 2007. ISSN 00310182. doi:  
10.1016/j.palaeo.2006.06.029. URL [http://linkinghub.elsevier.com/retrieve/  
pii/S0031018206004494](http://linkinghub.elsevier.com/retrieve/pii/S0031018206004494).
- 1148 [72] J. Kitchell. Biological selectivity of extinction. In E. G. Kaufman and O. H. Walliser,  
editors, *Extinction Events in Earth History*, pages 31–43. Springer-Verlang, Berlin,  
1150 1990. URL <http://link.springer.com/chapter/10.1007/BFb0011132>.
- 1152 [73] J. A. Kitchell. Evolutionary paleoecology: recent contributions to evolutionary theory.  
*Paleobiology*, 11(1):91–104, 1985.
- 1154 [74] D. G. Kleinbaum and M. Klein. *Survival analysis: a self-learning text*. Springer, New  
York, NY, 2 edition, 2005.
- [75] M. LaBarbera. Brachiopod orientation to water movement: functional morphology.

- 1156        *Lethaia*, 11(1):67–79, Jan. 1978. ISSN 0024-1164. doi: 10.1111/j.1502-3931.1978.tb01219.x. URL <http://doi.wiley.com/10.1111/j.1502-3931.1978.tb01219.x>.
- 1158 [76] M. LaBarbera. Water flow patterns in and around three species of articulate brachiopods. *Journal of Experimental Marine Biology and Ecology*, 55:185–206, 1981.
- 1160 [77] L. H. Liow. Does versatility as measured by geographic range, bathymetric range and morphological variability contribute to taxon longevity? *Global Ecology and Biogeography*, 16(1):117–128, 2007. URL <http://doi.wiley.com/10.1111/j.1466-8238.2006.00269.x>.
- 1162 [78] L. H. Liow, M. Fortelius, E. Bingham, K. Lintulaakso, H. Mannila, L. Flynn, and N. C. Stenseth. Higher origination and extinction rates in larger mammals. *Proceedings of the National Academy of Sciences*, 105(16):6097, 2008. URL <http://papers2://publication/doi/10.1111/j.1466-8238.2006.00269.x>.
- 1164 [79] L. H. Liow, L. Van Valen, and N. C. Stenseth. Red Queen: from populations to taxa and communities. *Trends in ecology & evolution*, 26(7):349–58, July 2011. ISSN 0169-5347. doi: 10.1016/j.tree.2011.03.016. URL <http://www.ncbi.nlm.nih.gov/pubmed/21511358>.
- 1166 [80] S. K. Lyons. A quantitative model for assessing community dynamics of pleistocene mammals. *The American naturalist*, 165(6):E168–85, June 2005. ISSN 1537-5323. doi: 10.1086/429699. URL <http://www.ncbi.nlm.nih.gov/pubmed/15937741>.
- 1168 [81] S. K. Lyons, P. J. Wagner, and K. Dzikiewicz. Ecological correlates of range shifts of Late Pleistocene mammals. *Philosophical transactions of the Royal Society of London. Series B, Biological sciences*, 365(1558):3681–93, Nov. 2010. ISSN 1471-2970. doi: 10.1098/rstb.2010.0263. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=2981999&tool=pmcentrez&rendertype=abstract>.
- 1170 [82] B. J. Macfadden. Origin and evolution of the grazing guild in New World terrestrial mammals. *Trends in ecology & evolution*, 12(5):182–187, 1997.
- 1172 [83] B. J. Macfadden. Extinct mammalian biodiversity of the ancient New World tropics. *Trends in ecology & evolution*, 21(3):157–65, Mar. 2006. ISSN 0169-5347. doi: 10.1016/j.tree.2005.12.003. URL <http://www.ncbi.nlm.nih.gov/pubmed/16701492>.
- 1174 [84] G. A. Mark and K. W. Flessa. A test for evolutionary equilibria: Phanerozoic brachiopods and Cenozoic mammals. *Paleobiology*, 3(1):17–22, 1977.
- 1176 [85] L. G. Marshall, S. D. Webb, J. J. Sepkoski, and D. M. Raup. Mammalian evolution and the Great American interchange. *Science*, 215(4538):1351–1357, 1982.
- 1178 [86] A. P. Martin and S. R. Palumbi. Body size, metabolic rate, generation time, and the molecular clock. *Proceedings of the National Academy of Sciences*, 90(9):4087–91, May 1993. ISSN 0027-8424. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=46451&tool=pmcentrez&rendertype=abstract>.

- [87] A. I. Miller and S. R. Connolly. Substrate affinities of higher taxa and the Ordovician Radiation. *Paleobiology*, 27(4):768–778, Dec. 2001. ISSN 0094-8373. doi: 10.1666/0094-8373(2001)027<0768:SAOHTA>2.0.CO;2. URL <http://www.bioone.org/doi/abs/10.1666/0094-8373%282001%29027%3C0768%3ASAHTA%3E2.0.CO%3B2>.
- [88] S. Nee. Birth-Death Models in Macroevolution. *Annual Review of Ecology, Evolution, and Systematics*, 37(1):1–17, Dec. 2006. ISSN 1543-592X. doi: 10.1146/annurev.ecolsys.37.091305.110035. URL <http://www.annualreviews.org/doi/abs/10.1146/annurev.ecolsys.37.091305.110035>.
- [89] S. Nee, R. May, and P. Harvey. The reconstructed evolutionary process. *Philosophical Transactions of the Royal Society B: Biological Sciences*, 344:305–311, 1994. doi: 10.1098/rstb.1994.0068. URL <http://rstb.royalsocietypublishing.org/content/344/1309/305.short>.
- [90] S. Nürnberg and M. Aberhan. Habitat breadth and geographic range predict diversity dynamics in marine Mesozoic bivalves. *Paleobiology*, 39(3):360–372, Apr. 2013. ISSN 0094-8373. doi: 10.1666/12047. URL <http://www.bioone.org/doi/abs/10.1666/12047>.
- [91] T. Olszewski and D. Erwin. Dynamic response of Permian brachiopod communities to long-term environmental change. *Nature*, 428(April):2–5, 2004. doi: 10.1038/nature02471.1. URL <http://www.nature.com/nature/journal/v428/n6984/abs/nature02464.html>.
- [92] R. Othman. *Petroleum geology of the Gunnedah-Bowen-Surat Basins, Northern New South Wales: stratigraphy, organic petrology and organic geochemistry*. PhD thesis, University of New South Wales, 2003.
- [93] R. Othman and C. R. Ward. Thermal maturation pattern in the southern Bowen, northern Gunnedah and Surat Basins, northern New South Wales, Australia. *International Journal of Coal Geology*, 51(3):145–167, Aug. 2002. ISSN 01665162. doi: 10.1016/S0166-5162(02)00082-4. URL <http://linkinghub.elsevier.com/retrieve/pii/S0166516202000824>.
- [94] B. Patterson and R. Pascual. The fossil mammal fauna of South America. *The Quarterly review of biology*, 43(4):409–451, 1968.
- [95] I. G. Percival, N. S. Meakin, L. Sherwin, T. A. Vanderlaan, and P. A. Flitcroft. Permian fossils and palaeoenvironments of the northern Sydney Basin, New South Wales. *Quarterly Notes Geological Survey of New South Wales*, 138:1–24, 2012.
- [96] R. H. Peters. *The ecological implications of body size*. Cambridge University Press, Cambridge, 1983.
- [97] S. E. Peters. Environmental determinants of extinction selectivity in the fossil record.

- 1230 *Nature*, 454(7204):626–9, July 2008. ISSN 1476-4687. doi: 10.1038/nature07032. URL <http://www.ncbi.nlm.nih.gov/pubmed/18552839>.
- 1232 [98] M. G. Powell. Latitudinal diversity gradients for brachiopod genera during late Palaeozoic time: links between climate, biogeography and evolutionary rates. *Global Ecology and Biogeography*, 16(4):519–528, July 2007. ISSN 1466-822X. doi: 10.1111/j.1466-8238.2007.00300.x. URL <http://doi.wiley.com/10.1111/j.1466-8238.2007.00300.x>.
- 1234 [99] S. A. Price, S. S. B. Hopkins, K. K. Smith, and V. L. Roth. Tempo of trophic evolution and its impact on mammalian diversification. *Proceedings of the National Academy of Sciences of the United States of America*, 109(18):7008–12, May 2012. ISSN 1091-6490. doi: 10.1073/pnas.1117133109. URL <http://www.ncbi.nlm.nih.gov/pubmed/22509033>.
- 1240 [100] D. L. Rabosky. Extinction rates should not be estimated from molecular phylogenies. *Evolution*, 64(6):1816–24, June 2010. ISSN 1558-5646. doi: 10.1111/j.1558-5646.2009.00926.x. URL <http://www.ncbi.nlm.nih.gov/pubmed/20030708>.
- 1242 [101] P. Raia, P. Piras, and T. Kotsakis. Turnover pulse or Red Queen? Evidence from the large mammal communities during the Plio-Pleistocene of Italy. *Palaeogeography, Palaeoclimatology, Palaeoecology*, 221(3-4):293–312, June 2005. ISSN 00310182. doi: 10.1016/j.palaeo.2005.02.014. URL <http://linkinghub.elsevier.com/retrieve/pii/S0031018205001033>.
- 1244 [102] P. Raia, C. Meloro, A. Loy, and C. Barbera. Species occupancy and its course in the past: macroecological patterns in extinct communities. *Evolutionary Ecology Research*, 8:181–194, 2006.
- 1248 [103] P. Raia, F. Carotenuto, J. T. Eronen, and M. Fortelius. Longer in the tooth, shorter in the record? The evolutionary correlates of hypsodonty in Neogene ruminants. *Proceedings. Biological sciences / The Royal Society*, 278(1724):3474–81, Dec. 2011. ISSN 1471-2954. doi: 10.1098/rspb.2011.0273. URL <http://www.pubmedcentral.nih.gov/articlerender.fcgi?artid=3189366&tool=pmcentrez&rendertype=abstract>.
- 1252 [104] D. M. Raup. Taxonomic survivorship curves and Van Valen’s Law. *Paleobiology*, 1(1):82–96, Jan. 1975. ISSN 0036-8075. doi: 10.1126/science.49.1254.50. URL <http://www.ncbi.nlm.nih.gov/pubmed/17777225>.
- 1256 [105] D. M. Raup. A kill curve for Phanerozoic marine species. *Paleobiology*, 17(1):37–48, 1991.
- 1260 [106] D. M. Raup and G. E. Boyajian. Patterns of generic extinction in the fossil record. *Paleobiology*, 14(2):109–125, 1988.
- 1262 [107] D. M. Raup and S. J. Gould. Stochastic simulation and evolution of morphology – towards a nomothetic paleontology. *Systematic Zoology*, 23(3):305–322, 1974.
- 1264 [108] J. R. Richardson. Ecology of articulated brachiopods. In A. Williams, C. H. C. Brunton,

- 1266 and S. J. Carlson, editors, *Treatise on Invertebrate Paleontology, Part H, Brachiopoda*  
1, pages 441–462. The Geological Society of America, Boulder, Colorado, 1997.
- 1268 [109] J. R. Richardson. Biogeography of articulated brachiopods. In A. Williams, C. H. C.  
1270 Brunton, and S. J. Carlson, editors, *Treatise on Invertebrate Paleontology, Part H,*  
*Brachiopoda 1*, pages 463–472. The Geological Society of America, Boulder, Colorado,  
1997.
- 1272 [110] K. D. Rose. *The beginning of the age of mammals*. Johns Hopkins University Press,  
Baltimore, Md, 2006.
- 1274 [111] M. Rosvall and C. T. Bergstrom. Maps of random walks on complex net-  
1276 works reveal community structure. *Proceedings of the National Academy of*  
*Sciences*, 105(4):1118–23, Jan. 2008. ISSN 1091-6490. doi: 10.1073/pnas.  
0706851105. URL <http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2234100/>&tool=pmcentrez&rendertype=abstract.
- 1280 [112] M. Rosvall, D. Axelsson, and C. Bergstrom. The map equation. *The European Physical*  
*Journal Special Topics*, 178(14):13–24, 2009. URL <http://www.springerlink.com/index/H8193132U6432363.pdf>.
- 1282 [113] K. Roy, G. Hunt, D. Jablonski, A. Z. Krug, and J. W. Valentine. A macroevo-  
1284 lutionary perspective on species range limits. *Proceedings. Biological sciences /*  
*The Royal Society*, 276(1661):1485–93, Apr. 2009. ISSN 0962-8452. doi: 10.1098/  
rspb.2008.1232. URL <http://www.ncbi.nlm.nih.gov/pmc/articles/PMC2677224/>&tool=pmcentrez&rendertype=abstract.
- 1286 [114] M. J. S. Rudwick. *Living and fossil brachiopods*. Hutchinson and Co, London, 1970.
- 1288 [115] J. J. Sepkoski. Stratigraphic biases in the analysis of taxonomic survivorship. *Paleobi-  
ology*, 1(4):343–355, 1975.
- 1290 [116] J. J. Sepkoski. A model of onshore-offshore change in faunal diversity. *Paleobiology*, 17  
(1):58–77, 1991.
- 1292 [117] C. E. Shannon. A mathematical theory of communication. *The Bell System Technical*  
*Journal*, 27:379–423,623–656, 1948. URL <http://dx.doi.org/10.1002/j.1747-2524.1948.tb00930.x>
- 1294 [118] C. A. Sidor, D. A. Vilhena, K. D. Angielczyk, A. K. Huttenlocker, S. J. Nesbitt, B. R.  
1296 Peecook, J. S. Steyer, R. M. H. Smith, and L. A. Tsuji. Provincialization of terrestrial  
faunas following the end-Permian mass extinction. *Proceedings of the National Academy*  
*of Sciences*, 110(20):8129–33, May 2013. ISSN 1091-6490. doi: 10.1073/pnas.1302323110.
- 1298 [119] M. Silva, J. H. Brown, and J. a. Downing. Differences in Population Density and  
1300 Energy Use between Birds and Mammals: A Macroecological Perspective. *The Journal*  
*of Animal Ecology*, 66(3):327, May 1997. ISSN 00218790. doi: 10.2307/5979. URL  
<http://www.jstor.org/stable/5979?origin=crossref>.

- [120] C. Simpson. *Levels of selection and large-scale morphological trends*. PhD thesis, University of Chicago, 2006.
- [121] C. Simpson and P. G. Harnik. Assessing the role of abundance in marine bivalve extinction over the post-Paleozoic. *Paleobiology*, 35(4):631–647, Dec. 2009. ISSN 0094-8373. doi: 10.1666/0094-8373-35.4.631. URL <http://www.bioone.org/doi/abs/10.1666/0094-8373-35.4.631>.
- [122] G. G. Simpson. *Tempo and Mode in Evolution*. Columbia University Press, New York, 1944.
- [123] G. G. Simpson. *The Major Features of Evolution*. Columbia University Press, New York, 1953.
- [124] F. A. Smith, J. Brown, J. Haskell, and S. Lyons. Similarity of mammalian body size across the taxonomic hierarchy and across space and .... *The American Naturalist*, 2004. URL <http://www.journals.uchicago.edu/doi/abs/10.1086/382898papers2://publication/uuid/D5606802-FD91-49EB-BE2F-E2D314A5E71D>.
- [125] F. A. Smith, S. K. Lyons, S. Morgan Ernest, and J. H. Brown. Macroecology: more than the division of food and space among species on continents. *Progress in Physical Geography*, 32(2):115–138, Apr. 2008. ISSN 0309-1333. doi: 10.1177/0309133308094425. URL <http://ppg.sagepub.com/cgi/doi/10.1177/0309133308094425>.
- [126] A. Solow and W. Smith. On fossil preservation and the stratigraphic ranges of taxa. *Paleobiology*, 23(3):271–277, 1997. URL <http://www.psjournals.org/doi/abs/10.1666/0094-8373-23.3.271>.
- [127] D. Strauss and P. M. Sadler. Classical confidence intervals and Bayesian probability estimates for ends of local taxon ranges. *Mathematical Geology*, 21(4):411–427, May 1989. ISSN 0882-8121. doi: 10.1007/BF00897326. URL <http://link.springer.com/10.1007/BF00897326>.
- [128] C. A. E. Strömberg. Decoupled taxonomic radiation and ecological expansion of open-habitat grasses in the Cenozoic of North America. *Proceedings of the National Academy of Sciences of the United States of America*, 102(34):11980–4, Aug. 2005. ISSN 0027-8424. doi: 10.1073/pnas.0505700102. URL <http://www.ncbi.nlm.nih.gov/articlerender.fcgi?artid=1189350&tool=pmcentrez&rendertype=abstract>.
- [129] C. A. E. Strömberg, R. E. Dunn, R. H. Madden, M. J. Kohn, and A. A. Carlini. Decoupling the spread of grasslands from the evolution of grazer-type herbivores in South America. *Nature communications*, 4:1478, Jan. 2013. ISSN 2041-1723. doi: 10.1038/ncomms2508. URL <http://www.ncbi.nlm.nih.gov/pubmed/23403579>.
- [130] S. G. Thomas, C. R. Fielding, and T. D. Frank. Lithostratigraphy of the late Early Permian (Kungurian) Wandrawandian Siltstone, New South Wales: record of glaciation? *Australian Journal of Earth Sciences*, 54(8):1057–1071, Dec. 2007. ISSN 0812-0099.

- 1340 doi: 10.1080/08120090701615717. URL <http://www.tandfonline.com/doi/abs/10.1080/08120090701615717>.
- 1342 [131] S. Tomiya. Body Size and Extinction Risk in Terrestrial Mammals Above the Species Level. *The American Naturalist*, pages E000–E000, Sept. 2013. ISSN 00030147. doi: 10.1086/673489. URL <http://www.jstor.org/stable/info/10.1086/673489>.
- 1344 [132] L. Van Valen. A new evolutionary law. *Evolutionary Theory*, 1:1–30, 1973. URL <http://ci.nii.ac.jp/naid/10011264287/>.
- 1346 [133] L. Van Valen. Taxonomic survivorship curves. *Evolutionary Theory*, 4:129–142, 1979.
- 1348 [134] L. Van Valen. Three paradigms of evolution. *Evolutionary Theory*, 9:1–17, 1989.
- 1350 [135] J. Waterhouse and G. Shi. Climatic implications from the sequential changes in diversity and biogeographic affinities for brachiopods and bivalves in the Permian of eastern Australia and New Zealand. *Gondwana Research*, 24(1):139–147, July 2013. ISSN 1342937X. doi: 10.1016/j.gr.2012.06.008. URL <http://linkinghub.elsevier.com/retrieve/pii/S1342937X12002171>.
- 1354 [136] J. B. Waterhouse. Late Palaeozoic Mollusca and correlations from the south-east Bowen Basin, east Australia. *Palaeontographica Abteilung A*, 198:129–233, 1987.
- 1356 [137] C. O. Webb, D. D. Ackerly, M. a. McPeek, and M. J. Donoghue. Phylogenies and Community Ecology. *Annual Review of Ecology and Systematics*, 33(1):475–505, Nov. 2002. ISSN 0066-4162. doi: 10.1146/annurev.ecolsys.33.010802.150448. URL <http://arjournals.annualreviews.org/doi/abs/10.1146%2Fannurev.ecolsys.33.010802.150448>.
- 1360 [138] E. P. White, S. K. M. Ernest, A. J. Kerkhoff, and B. J. Enquist. Relationships between body size and abundance in ecology. *TRENDS in Ecology and Evolution*, 22(6):323–30, June 2007. ISSN 0169-5347. doi: 10.1016/j.tree.2007.03.007. URL <http://www.ncbi.nlm.nih.gov/pubmed/17399851>.
- 1364 [139] J. C. Zachos, M. Pagani, L. Sloan, E. Thomas, and K. Billups. Trends, rhythms, and aberrations in global climate 65 Ma to present. *Science*, 292:686–693, 2001. doi: 10.1126/science.1059412. URL <http://www.sciencemag.org/content/292/5517/686.short>.
- 1366 [140] J. C. Zachos, G. R. Dickens, and R. E. Zeebe. An early Cenozoic perspective on greenhouse warming and carbon-cycle dynamics. *Nature*, 451(7176):279–283, Jan. 2008. ISSN 1476-4687. doi: 10.1038/nature06588.

1370 **A Permian lithology and paleoenvironment**

Lithological and paleoenvironmental assignments available in the PBDB can be poorly  
1370 resolved or missing in the case of environment. Because these assignments are critical in the  
proposed study of brachiopod survival and distribution (Section 2), it is necessary to improve  
1372 these values with more precise information available in the paleontological and geological  
literature. Currently, none of these improved assignments have been included in any of the  
1374 above analyses (Section 2).

The geological unit reference guide on from Geosciences Australia (<http://www.ga.gov.au/>),  
1378 the lithological information for many of the brachiopod occurrences can be improved and  
made more precise (Table 3). The lithological assignments below are based on the order  
1380 with which rock types are named in the lithological description of a geological unit. These  
were extracted automatically using a very simple algorithm. While more than two rock types  
1382 may be listed for a geological unit, only the first two are reported below. These formations  
represent 3533 of 5737 (61%) total occurrences across all of Australia.

Table 3: Australian formations included in the study of brachiopod survival and distribution (Section 2). Improvements are based on lithological descriptions from Geosciences Australia (<http://www.ga.gov.au/>).

geological unit	PDBD lithology 1	PBDB lithology 2	my lithology 1	my lithology 2
Aldebaran Sandstone	sandstone		conglomerate	siltstone
Allandale Formation	siliciclastic		conglomerate	sandstone
Alum Rock Conglomerate	siliciclastic		tuff	limestone
Bon-donga/Pikedale/Silver Spur and beds				
Baker Formation	siltstone		siltstone	quartz
Bakers Blue Granite	siltstone		granodiorite	
Bakers Creek Diorite	siltstone		diorite	quartzbiotite
Bakers Creek Suite	siltstone		gabbros	diorites
Bakerville Granodiorite	siltstone		granodiorite	
Barfield Formation	sandstone		tuff	conglomerate
Baywulla Formation	siltstone		limestone	conglomerate
Beekeeper Formation	not reported		carbonatesiliciclastic	carbonatesiliciclastic
Berserker Group	siliciclastic		conglomerates	breccia
Billidee Formation	sandstone		siltstone	shale
Bingleburra Formation	siltstone	limestone	mudstone	siltstone
Black Alley Shale	shale		shale	siltstone
Black Jack Granodiorite	siliciclastic		granite	granodiorite
Black Jack Group	siliciclastic		sandstone	
Blenheim Formation	sandstone		sandstone	coquinite
Branch Creek Rhyolite	sandstone		rhyolite	breccia
Broughton River Granodiorite	sandstone		granodiorite	granite
Broughton River Suite	sandstone		granodiorite	
Buffel Formation	siliciclastic		limestone	limestone
Bulgadoo Shale	shale		shale	siltstone
Burnett Formation	siltstone		arenite	siltstone
Callytharra Formation			calcarenite	conglomerate
Carmila beds	siliciclastic		siltstone	basalt
Carolyn Formation	sandstone	claystone	sandstone	sandstone
Carrandibby Formation	siliciclastic		claystone	siltstone
Catherine Sandstone	sandstone		siltstone	mudstone
Cattle Creek Formation	siliciclastic		mudstone	quartzose
Condamine beds	mudstone		conglomerate	tuff
Cookilya Sandstone	sandstone		quartz	siltstone
Coyrie Formation	siliciclastic		shale	siltstone
Crocker Well Suite	siliciclastic		granodiorite	
Cundlego Formation			siltstone	shale
Darlington Limestone	limestone		limestones	calcirudites
Eight Mile Creek beds	siliciclastic		conglomerate	sandstone
Eight Mile Creek Granite	siliciclastic		granite	
Eight Mile Creek Granodiorite	siliciclastic		granite	
Flat Top Diorite	sandstone		diorite	diorite
Flat Top Formation	sandstone		tuff	sandy
Freitag Formation	sandstone		sandstone	sandstone
Gilgurra Mudstone	mudstone		mudstone	sandstone
Glencoe Gabbro	mudstone		gabbro	gabbro
Glencoe Limestone Member	mudstone		limestone	
Glenmore Creek Granite	siliciclastic		monzogranite	
Gray Creek Complex	siltstone		metagabbro	
Gudman Formation	grainstone		feldsarenitesiltstone	feldsarenitesiltstone
Hardman Formation	sandstone		sandstone	limestone
Hickman Creek Granite	siliciclastic		monzogranite	
High Cliff Sandstone	sandstone		siltstone	shale
Holmwood Shale	siliciclastic		limestone	shale
Ingelara Formation	siliciclastic		sandy	siltstone
Ingliston Granite	siliciclastic		granite	
Kockatea Shale	shale		shale	
Lakes Creek Formation	not reported		volcanics	siltstone
Lizzie Creek Volcanic Group/Mount Wickham Rhyolite	sandstone		andesite	sandstones
Lochinvar Formation				rhyolite
Manning Group	limestone		basalt	siltstone
Maria Formation	siliciclastic		mudstone	conglomerate
Maria Island Granite	siliciclastic		mudstone	shale
Marra Creek Formation	siliciclastic		granite	
Marra Formation	siliciclastic		sandy	carbonate
Marrangaroo Conglomerate	siliciclastic		sandstone	siltstone
Marrar Dyke	siliciclastic		monzogabbro	conglomerate
Merlewood Formation			sandstone	
Mistletoe Granite	siliciclastic		granite	conglomerate
Moonlight Valley Tillite	sandstone		conglomerate	sandstone
Mooraback beds	siliciclastic		sandstone	siltstone
Mount Poole Monzogranite	siliciclastic		monzogranite	
Muggleton Formation	siliciclastic		shale	quartzose
Mulbring Siltstone	sandstone		claystone	sandstone
Muree Sandstone	siltstone		sandstone	conglomerate
Namoi Formation	siliciclastic		siltstone	litharenite
Narayan beds			conglomerate	siltstone
Nowra Sandstone	sandstone		conglomerate	quartzose

Oxtrack Formation	siltstone	chert	siltstone
Peawaddy Formation	siliciclastic	siltstone	siltstone
Poole Sandstone	siliciclastic	conglomerate	quartzose
Porcupine Creek Granodiorite	siliciclastic	granodiorite	
Porcupine Creek rhyolite	siliciclastic	ignimbrite	
Porcupine Formation	siliciclastic	conglomerate	sandstone
Quinnanie Shale	shale	shale	siltstone
Rammutt Formation	siliciclastic	mudstone	basaltic
Rhyolite Range beds	siliciclastic	sandstone	siltstone
Risdon Stud Formation	sandstone	tuff	arenite
Rutherford Formation	siliciclastic	marl	sandstone
Silver Spur beds	siliciclastic	conglomerate	mudstone
Snapper Point Formation/Wandrawandian Siltstone	siltstone	sandstone	siltstone
South Curra Limestone	limestone	grainstone	calcilutite
Tamby Creek Formation	siliciclastic	andesite	breccia
Tomago Coal Measures	siliciclastic	tuff	siltstone
Towgon Grange Tonalite	mudstone	granodiorite	diorite
Tulcumbe Sandstone	sandstone	sandstone	sandstone
Wandagee Formation		siltstone	quartz
Wandrawandian Siltstone	siltstone	siltstone	quartzlithic
Watermark Formation	siliciclastic	siltstone	claystone
Waverley Formation		sandstone	conglomerate
Waverly Formation/Dangerfield Formation		sandstone	conglomerate
Werri Basalt	siliciclastic	basaltic	tuffs
Woolooma Formation		tuff	limestone
Yagon Siltstone	siltstone	mudstone	sandstone
Yagon Siltstone/Booti Booti Sandstone	siltstone	siltstone	coalrich
Yessabah Limestone	limestone	limestone	mudstone

<sup>1384</sup> Below is a set of representative PBDB assignments and my improvements, focused primarily  
<sup>1386</sup> on paleoenvironmental reconstruction (Table 4). These assignments are very preliminary and  
<sup>1388</sup> based only on a few key papers and associated maps [30, 31, 49, 92, 93, 95]. There are a total  
<sup>1390</sup> of 5737 Permian Australian brachiopod occurrences in the PBDB, from both eastern and  
Western Australia. Within is, there are 4711 occurrences that are not from Western Australia.  
The geological units listed in Table 4, which are from Eastern Australia, account for 1897  
of these occurrences which is about 33% of the total samples or 40% of the east Australian  
samples.

formation	PBDB paleoenvironment	my paleoenvironment 1	my paleoenvironment 2
Aldebaran	offshore	deltaic/coastal plain	nearshore marine
Allandale	coastal indet	sublittoral strand	marine shelf
Barfield	coastal indet	prograding shelf	deep shelf
Berry	marine indet	offshore marine	
Black Alley	offshore	alluvial plain	delta
Black Jack	marine indet	alluvial	delta
Branxton	coastal indet	fan delta	delta plain
Buffel	coastal indet	delta	shallow shelf/coastal
Camboon	coastal indet	alluvial	lacustrine
Catherine	coastal indet	prograding shelf	nearshore marine
Cattle Creek	coastal indet	delta	nearshore marine
Farley	marine indet	delta plain	delta front
Flat Top	coastal indet	prograding shelf	deep shelf
Freitag	marine indet	coastal plain	offshore marine
Ingelara	marine indet	prograding shelf	offshore marine
Lizzie Creek	coastal indet	alluvial	lacustrine
Lochinvar	marine indet	sublittoral strand	marine shelf
Mulbring	marine indet	marine shelf	
Muree	coastal indet	alluvial fan	fan delta
Nowra	coastal indet	nearshore marine	coastal
Oxtrack	coastal indet	deltaic/coastal plain	shallow shelf/coastal
Peawaddy	marine indet	prograding shelf	nearshore marine
Porcupine	marine indet	marine shelf	
Rutherford	coastal indet	delta front	marine shelf
Snapper Point	shoreface	fluvial coastal	nearshore marine
Wandrawandian	offshore	offshore marine	
Wasp Head	shoreface	alluvial valley fill	nearshore marine
Watermark	marine indet	delta	marine shelf

Table 4: Australian geological units included in the study of brachiopod survival and distribution (Section 2). Both PBDB assignments and those sourced from the literature are included. Improvements are obvious, especially in regards to paleoenvironmental reconstruction.