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**Introduction**

Snake venom is perhaps best known for its ability to incapacitate biological systems, a property which has made it both a source of novel biomedical compounds ([1](#_ENREF_1)) and also a major health concern due to the estimated millions of human envenomation cases each year ([2](#_ENREF_2)). However, snake venom is also an ideal system of predatory trait evolution ([1](#_ENREF_1)), particularly as its predatory functionality, the ability to subjugate prey, can be quantified by measured venom potency and the quantity a species possess. Despite significant developments in our understanding of the evolutionary origin ([3-5](#_ENREF_3)) and the subsequent evolution of its complex compounds ([1](#_ENREF_1), [6](#_ENREF_6)), surprisingly little is known about the evolutionary pressures that drive this ability to incapacitate prey. For example, many species, such as the Gold-ringed cat snake (*Boiga dendrophila*), are incapable of subduing laboratory test models (ref) while other species, such as Russel’s viper (*Daboia russelii*), possess enough potent venom to incapacitate hundreds of thousands of potential prey items (ref). This lack of a clear pattern relating to the ability to subdue prey using venom suggests drivers assumed to be important in driving predator trait evolution, such as trophic factors, are less important than expected (ref). One potential explanation for this lack of a pattern is that weak selective pressures are caused by an “overkill” effect, were due the capacity of many species venoms to kill prey numbers far in excess of any biological requirement, such as in the case of Russel’s viper, neutral evolutionary process becoming the main driver of trait variation ([7](#_ENREF_7), [8](#_ENREF_8)). Such a scenario predicts that traits relating to venom, such as potency and quantity, would be idiosyncratic in nature and also raises the question of the general importance of ecological factors on the evolution of predatory traits. Here, we use a comparative approach that accounts for both snake phylogenetic history and the evolutionary naivety of species which venom potency is measured on to test whether snake venom traits follow idiosyncratic patterns expected from neutral selection or patterns predicted from macroevolutionary and trophic theory.

As predation plays a key role in both ecosystems dynamics ([9](#_ENREF_9)) and the evolution of the species within these ecosystems ([10](#_ENREF_10)) much attention has been given to understanding the general patterns related to these trophic interactions and the drivers behind them. These include macroecological patterns derived from body mass and habitat complexity which predict allometric scaling relationships for traits underlining trophic interactions such as search and encounter rates ([11-13](#_ENREF_11)); the ability to spot and track prey ([14](#_ENREF_14), [15](#_ENREF_15)); and ingestion rates ([13](#_ENREF_13), [16](#_ENREF_16)). Other general predictions for predatory traits stem from the co-evolution between both predators and their prey, were arms race dynamics may drive the evolution of predatory traits that increase the ability to capture prey and conversely the evolution of traits in prey species that increases their ability to escape ([17](#_ENREF_17), [18](#_ENREF_18)). While these two general drivers are likely to have important influences on trophic trait evolution they can be difficult to decouple as predatory traits are often multifaceted, for example claws can be used for predation, climbing, digging, defence etc., and difficult to compare across the large number of species required for macroecological studies (ref). Snake venom however, offers a primarily predatory system which can be quantitatively compared across a large group of diverse species allowing the importance of both trophic and macroecological factors on its evolution to be tested.

Predatory foraging traits are expected to be adapted towards their main target prey species. For example, jaw morphology in cichlid fish is strongly selected by factors such as prey type ([19](#_ENREF_19)). Similar selection pressures from prey type is also seen in snake venom. For example, a switch in diet from fish to eggs has resulted in the almost complete atrophy of the venom apparatus in the egg eating marble sea snake *Aipysurus eydouxii* (refs), demonstrating the importance of predation in venom evolution (ref). Further such trophic selection is also seen in cases of prey-specific venoms were potency is higher when tested on species commonly found within the snakes diet, such as observed in Malayan pitvipers ([20](#_ENREF_20)), coral snakes ([21](#_ENREF_21)), the viper genus *Echis* ([22](#_ENREF_22)), saw-scaled vipers ([23](#_ENREF_23)) and insect eating *Pelias* vipers ([24](#_ENREF_24)). However, such co-evolution is not apparent in all snake species, such as in tiger snakes where variation in venom across populations is independent of dietary differences ([25](#_ENREF_25)). Furthermore, prey species can also respond in such co-evolutionary arms races through the development of immunity. Evidence for this is seen in venoms that show weaker potencies in potential prey items such as in the cases of Opossums and Neotropical pitvipers ([26](#_ENREF_26)); eels and *Laticauda colubrine* ([27](#_ENREF_27)); and between ground squirrels and rattlesnakes ([28](#_ENREF_28), [29](#_ENREF_29)) . This idiosyncratic pattern of some species displaying prey-specific venom while others do not may mirror a pattern predicted from the Overkill hypotheses were neutral processes are the main driver of venom potency ([8](#_ENREF_8)). However, many measures of venom potency are not typically tested on natural prey species leading to the question of whether these examples represent an idiosyncratic pattern expected under weak selection or whether a general pattern will emerge under appropriate analysis (ref). While measures of potency using natural prey species are becoming more common, these are taxonomically restricted making large comparative analysis difficult ([21](#_ENREF_21), [22](#_ENREF_22)). Here we account for the species a venom is tested on and measure the phylogenetic distance of this species to the snake’s species natural diet which, in turn, allows us to test whether venom is generally adapted towards common target prey species. However, while venom potency is an important aspect of these species ability to incapacitate prey, the volume of venom and the role of other macro-ecological drivers are also likely to be important when considering venom evolution.

The ability to subdue prey is not only dependent on the potency of a species venom but also the amount of venom available. As such, venom volume is also expected to be under positive selection from potential trophic and macroecological factors. As venom production incurs a metabolic cost ([30](#_ENREF_30)) (although the level of this cost is debated ([31](#_ENREF_31))) and requires storage, the volume of venom a snake species can produce is likely to be heavily linked to one of the strongest determinants of trophic ecology; body size. In general, larger predators eat larger prey ([16](#_ENREF_16)). If venom is under selection for its ability to kill prey it would be expected that larger snake species would need to produce larger quantities of venom to keep pace with the subsequent increased in prey size. Such an increase of venom volume to compensate for larger prey is supported by the metering of venom in response to prey size seen in several species ([32](#_ENREF_32), [33](#_ENREF_33)). However, while bigger snakes are known to have larger amounts of venom in general ([34](#_ENREF_34)) little is known about the interspecific scaling of this trait. One prediction is that venom volume increases with snake body size with an exponent relating to their predator-prey body size scaling as described in equation 1;

Were the scaling exponent *a* is approximately 0.88 ([16](#_ENREF_16)). However, venom volume would not be expected to scale according to this exponent as the effects of toxicological agents also follows an allometric relationship ([35](#_ENREF_35)) were the amount of venom required (*V*) to induce the same incapacitating effect on a prey of mass (*Mprey*) would follow (equation 2);

were *b* is the scaling coefficient of venoms toxicological effects ([35](#_ENREF_35)). Hence to calculate the expected allometry of venom volume with snake body mass in a case were venom volume increases at a rate to match predator-prey size scaling after accounting for scaling of toxicological effects we substitute from equation 1 into equation 2 to get (equation 3);

If we take the commonly used value of 0.75 for the interspecific scaling of drug dosages ([35](#_ENREF_35)) for *b* and the value of 0.68 for *a* from the scaling predator-prey mass relationship for snakes ([16](#_ENREF_16)) we would expect a scaling exponent of approximately 0.51 between snake venom volume and snake mass. Other predictions relating to venom volume scaling include the overkill hypothesis which predicts no relationship between venom volume and prey size, or alternatively venom volume may scale according to constraints such as metabolic costs, were a scaling of 0.75 would be expected ([36](#_ENREF_36)). At the other extreme superlinear allometeries would suggests patterns associated with drivers such as sexual selection, such as proposed by the weapons hypothesis ([37](#_ENREF_37)), or defenses requiring increased effectiveness with size, such as seen in the allometry of horn growth in horned lizards ([38](#_ENREF_38)) (Figure 1).

Finally, an overlooked feature that may also drive both venom volume and toxicity evolution is habitat structure ([39](#_ENREF_39)). The structural complexity of a habitat, such as whether it's a 2-dimensional terrestrial surface or a complex 3-dimensional forest canopy, can influence both encounter rates ([12](#_ENREF_12), [16](#_ENREF_16)) and the escape rates of prey, with higher dimensional spaces increasing both ([40](#_ENREF_40), [41](#_ENREF_41)). Hence predators in high dimensional habitats with associated increased escape rates may compensate through larger volumes of more potent venom in order to increase capture rates. For example, strike and release behaviors may be less successful in either 3-dimensional arboreal or aquatic environments requiring higher toxicities to incapacitate prey quickly. Conversely there may be less of a requirement for high potencies and venom volumes due to increases in encounter rates, and hence feeding opportunities, in high dimensional habitats which may compensate for possible increases in escape rates.

Here we test the importance of these multiple potential drivers of both venom quantity and potency in a phylogenetically corrected comparative analysis of ninety-nine species of venomous snakes. We test the importance of snake body mass; habitat dimensionality; prey type; and prey size on the variation of both venom toxicity, as measured using the median lethal dose (LD50), and venom volume. We achieve this by using a novel metric of the evolutionary distance between the model animal on LD50 is measured and the typical species found in each snakes’ diet in order to test the general pattern of snake venom prey-specificity which predicts higher potencies when tested on species phylogenetically close to natural target prey. This approach also allows us to control for the variance associated with the LD50 model used and hence test the general influence of macroecological factors on factors such as venom volume.

Using these corrections and a series of models accounting for phylogenetic similarity between snake species we test; (1) the overkill hypothesis: that there is no relationship between venom volume or toxicity with prey size; (2) the importance of trophic drivers on venom evolution including; that venom potency is higher (lower LD50) when tested on model species phylogenetically closer to species found in the diet; and that snake species which include eggs in their diets have lower venom potencies (higher LD50) or volumes; (3) the importance of macorecological drivers on venom evolution including; that venom volume is higher in species with larger prey, with a predicted scaling of approximately 0.51 or that (4) venom volume scales according to 0.75 as expected if metabolic cost is the main driver; (5) the importance of habitat dimensionality on venom evolution, in particular that; species in high dimensional habitats show either higher or lower potencies or venom volumes depending on encounter and escape rates; and finally (6) the importance of other potential drivers such as superlinear scaling of venom volume or potency as expected with the weapons hypothesis or sexual selection drivers. We show that both trophic and macroecological factors are important in driving venom evolution with patterns supporting prey-specific venom in general and venom volume scaling as predicted by metabolic cost constraints.

**Results**

Our final compiled dataset of venom traits and corresponding trophic and macroecological data consisted of 275 observations over 99 species which corresponds to the data used in the main analysis. We also conducted supplementary analysis were we included whether species to are known to using constricting behaviors; the inclusion of habitat type and finally an analysis including prey body size which was conducted using a reduced dataset of 177 observations across 68 species. We report the results across all models relating to the importance of each driving factor in order below.

**Predator-prey coevolution**

Despite the presence of only eight egg eating species in our dataset LD50 was found to be significantly higher in species with eggs in their diet in both the main and constriction models, while venom volume had a negative, but non-significant, association with ovivorous behavior in all models (Table 2; Tables A2-4).

Of the species included within the analysis only 14 species had a diet completely matching that of the LD50 model their venom was tested on, i.e. the LD50 of a species with a diet including 100% mammals tested using a mouse model. Most species in the dataset had a diet with a least some component not matching the LD50 model as reflected by a median evolutionary distance of 211.3 million years between the common ancestors of the dietary taxa and the LD50 species. In all models species LD50 increased with mean phylogenetic distance between the diet and the LD50 model with snake species with diets phylogentically close to the LD50 model species hence having higher potencies (table 2; tableA2-3; Figure 1B). From the main model, after back transforming the mean centred log10 value, LD50 was found to increase by 1.44 for every 100 million years between the species in the diet and that of the LD50 model. This was after correcting for LD50 injection route were intravenous and Intraperitoneal routes were found to have lower LD50 values in comparison to a subcutaneous route (table 2; tableA2-3).

**Macroecological drivers; body size and habitat dimensionality**

The mean volume of venom, as measured using dried weight, ranged from 0.15 mg in the egg-eating sea snake (*Emydocephalus annulatus*) to 571 mg in the forest cobra (*Naja melanoleuca*). Body size, prey size and habitat dimensionality all correlated with venom volume. The main correlate with venom volume was snake body size with a log-log scaling of between 0.74 and 0.76 across all models after back transforming (Table 2, A2-4; Figure 1). Venom volume also showed a positive increase with prey body mass, with a log10-log10 increase of 0.139, however only 90% of the posterior samples are above the zero threshold (Table A2). In the model corresponding to equation 2, which only included venom volume and prey mass, we found a significant exponent of 0.37 (Table A5) while in the model corresponding to equation 1 of predator prey body mass scaling we found an exponent of 0.52 (Table A6). Whether using the scaling exponent of 0.51 as calculated in equation 2 or using an exponent of 0.19 as calculated using a value of 0.37 found here between venom volume and prey size for *b* in equation 2 and the value of 0.52 found here between prey and predator scaling for *a* in equation 1, both values are far lower than the exponent of 0.75 between venom volume and snake body mass found across all models (Table 2, A2-4; Figure 1). Snake body size was also found to have a significantly positively correlation with LD50, meaning larger snakes showed decreased potency. However, this relationship was only significant in the main analysis (Table 2).

The next most significant driver of venom volume was the dimensionality of the habitat with the 27 species in high dimensional environments (arboreal = 9, aquatic = 18) showing lower venom volumes in comparison to species in lower dimensional habitats (Table 2). A sensitivity analysis were habitat was include as terrestrial, arboreal and aquatic also showed similar significant reductions in both arboreal and aquatic habitats (Table A4).

**Phylogeny, constriction behaviour and covariance between venom volume and LD50**

In all models there is was a high phylogenetic signal for both LD50 and venom volume with lambda values of x for LD50 and x for venom volume (Table 2, A2-4). For all models LD50 showed a higher phylogenetic signal (Table 2, A2-4). The presence of constricting behaviour was found to have no effect on either venom volume or LD50(Table A3). Finally, there is no significant covariance between either the residuals of phylogenetic terms of both venom volume and LD50 across all models (Table 2, A2-4).

**Discussion**

Predator traits are thought to be heavily shaped by both predator-prey evolutionary dynamics and macroecological forces such as imposed by the limitations of body size and habitat structure. Here we show that traits associated with snake venom follow patterns predicted from a scenario were these factors are important evolutionary drivers of predator trait. This includes that venom potency is prey specific, with higher potencies found when venom is testes on species more phylogenetically similar to a snakes natural diet, and that venom volume scales with snakes body size with an allometry expected if venom volume is determined by the energetic costs of its production. We also find that snake venom is influenced, as expected by the presence of ovivorous feeding behaviours and, in a less expected direction, by the dimensionality of the environment. Hence, our results suggest that while aspects of neutral selection, such as genetic drift, may play an important role in generated the observed variance in predator traits, positive selection forces associated with trophic and macroevolutionary drivers still play a key role in the evolution of these traits.

Trophic factors play an important role in the maintenance of venom. This is supported by cases of its evolutionary loss with certain dietary changes (ref), the absence of an increased lifespan associated with venomous as generally seen in species with toxic defence systems (ref), and also by our findings here that ovivorous feeding is associated with lower potencies and venom volumes. However, the role of trophic drivers in shaping the subsequent evolution of venom has been much more hotly debated, in particularly relating to the generality of prey-specific venoms ([1](#_ENREF_1), [7](#_ENREF_7), [8](#_ENREF_8)). By flipping the inconvenience usually associated with the use of none-natural test models for venom potency ([21](#_ENREF_21)) we show that venom is generally prey-specific with higher potencies associated with species that more closely resemble a species diet. This prey-specific effect found in our analysis is also likely to underestimate the prey specific nature of venom in general due to our use of LD50. While LD50 measures the lethality of a venom, it is likely that venom is selected to simply subdue prey, as opposed to cause mortality, in order to reduce the chances of prey escaping or retaliating ([22](#_ENREF_22)). Even though prey mortality achieves this, other measures of venoms ability to subdue prey, such as the speed at which a venom affects prey, may give clearer reflection of venoms prey-specificity ([22](#_ENREF_22)). Despite such limitations, we find a clear pattern of prey-specificity suggesting that cases of non-prey specific venom ([25-29](#_ENREF_25)) are more likely to be the exception as opposed to the general rule.

While the predator-prey dynamics of snake venom has received relatively significant attention the importance of macroecological factors, namely body size and habitat complexity, have received surprisingly little despite their known importance in driving trophic interaction in general (refs). Our results support the importance of these factors with venom volume in particular showing strong relationships to both snake body size and habitat dimensionality. Unsurprisingly, we found that larger snakes have larger quantities of venom, more surprisingly however, these increases did not follow the expected scaling allometery predicted from a predator-prey scaling perspective ([16](#_ENREF_16)) with venom volume increasing faster with snake body size than expected. Even when considering potential variation in the allometric scaling of toxicological effects (equation 2) ([35](#_ENREF_35)), an exponent far in excess of 1 is required for *b* in order for the observed scaling of 0.75 between venom volume and snake mass to match. Furthermore, our analysis shows that the predatory-prey scaling of venomous snakes is much lower in comparison to snakes as a whole ([16](#_ENREF_16)), with larger venomous snakes feeding on much smaller prey items than expected, a trend that may also explain the reduced venom potency with snake size seen in our main analysis. A more likely explanation for the venom volume scaling found here is that it relates to limitations relating to metabolic rate ([67](#_ENREF_67)), which increases with a scaling coefficient of 0.75 with respect to body mass ([36](#_ENREF_36), [67](#_ENREF_67)). In particular, it might be expected that the production of a potentially metabolically costly material such as venom ([30](#_ENREF_30)), would follow such a metabolic scaling, particularly if the costs of venom were maintained at a constant proportion of overall energy budgets across snake species.

Another potential macroecological factor shaping the available volume of venom to a species is habitat dimensionality. While we expected that species in high dimensional habitats may have higher venom volumes to compensate for higher escape rates of prey ([39](#_ENREF_39)) we found that, counter to our expectation, these species had lower venom volumes in comparison species in low dimensional habitats (terrestrial and fossorial). This may be associated with differences in prey capturing behaviour as it might be expected that high dimensional environments require more holding behaviours during attacks in order to prevent the loss of prey, hence allowing a more accurate delivery of venom. However, the presence of constriction in venomous snakes ([43](#_ENREF_43)), the most extreme form of prey holding behaviours, is present in both arboreal and terrestrial species and was found to have no effect when included within the analysis. Furthermore bite and release behaviours are known in arboreal species such as the black mamba (*Dendroaspis polylepis*) suggesting this behaviour is not fully restricted to low dimensional environments ([51](#_ENREF_51)). Another potential explanation is that higher encounter rates in high dimensional environments ([12](#_ENREF_12)) may reduce the missed opportunity of feeding cost associated with replenishing venom. Rates of replenishing venom can be substantial with estimates of replenishment rate ranging from 3-7 days ([68](#_ENREF_68)) to 30-50 days ([33](#_ENREF_33), [69-71](#_ENREF_69)). These long periods of replenishment may hence select for larger reserves in species where encounter rates with prey are lower in order to minimise potential missed opportunity costs. While further research on the role of habitat dimensionality will allow more detailed understanding of the mechanisms driving this trend our results highlight that venom may not only be selected according to metabolic constraints but that factors relating to encounter rate may also have important influences.

Our analysis shows that both predator-prey dynamics and macroecological factors shape the evolution of venom in snakes. Through establishing these patterns we reject the strongest form of the overkill hypothesise which posits that positive selection plays a minor role in shaping venom evolution. However, while the above factors play an important role in the evolution of venom, much of the variation relating to the ability to incapacitate prey is still unexplained. Outside selection pressures related to ecology and physiology other evolutionary mechanisms, such as gene duplication events, play an important role in venom evolution ([66](#_ENREF_66)). In fact, such mechanisms are likely to be reflected in the high phylogenetic signal found for both venom volume and potency, were rare duplication and mutation events result in closely related species showing similar trait values ([72](#_ENREF_72)). While future research will continue to unravel the huge variation associated with snakes ability to kill prey, our analysis shows that both predatory-prey dynamics and macroecological factors are important drivers in the evolution of venom. These results show that by testing general drivers of predatory traits we can further understand the evolution of not only unusual physiologies, such as the venomous system, but also through understanding the evolution of predator traits in general, trophic ecology as a whole.

**Methods**

**Data**

We collected data on venom volume and toxicity from the literature, along with our predicted drivers. We used mean dry weight (mg) as a measure of venom volume as it represents the amount of active ingredients available and is the most available reported measure. As a measure of venom lethality we used median lethal dose (LD50). We only included intravenous (IV), subcutaneous (SC), Intraperitoneal(IP) or intramuscular routes (IM) of injecting the venom as other routes were too uncommon to include within the analysis. We include LD50 values measured on all animal models as we were interested in including variation relating to the potential prey specific nature of venom. While other measures of venom toxicities are available, only measures of LD50 were numerous enough to carry out a large scale comparative analysis.

To test whether venom is prey specific we calculated the phylogenetic distance between the model animal species used to measure LD50 for each snake species and the species naturally present in its diet. We calculated this as the sum of the phylogenetic distance, using mean estimates from TimeTree ([42](#_ENREF_42)), between each prey group and the LD50 model multiplied by the proportion of each prey group reported in each snake species diet. For example, a species with a diet comprising of 20% mammals, 50% fish and 30% reptiles with a LD50 measured using mice would have a diet with an average phylogenetic distance of 0.2(0) + 0.5(400.1) + 0.3(296) = 288.85 million years from the common ancestor of the LD50 model.

Diet data was collated from the literature using studies with quantitative estimates of prey proportions, mainly from studies of stomach contents. As prey items were rarely identified to lower taxonomic levels diet was categorized into six prey categories; invertebrates, fish, amphibians, lizards, birds and mammals.

Species habitat was categorized as either terrestrial, fossorial, aquatic or arboreal based on accounts in the literature. In order to directly test the expected effect of the dimensionality of habitat environment each environment was scored, as in Pawar et al (2012)need ref, with terrestrial and fossorial environments scored as two-dimensional and arboreal and aquatic scored as three-dimensional. As some venomous species also engage in constriction behavior we collected data on any observation of constriction behavior in capturing prey from the literature ([43](#_ENREF_43)).

For snake body size we used total length values from the literature ([44-50](#_ENREF_44)) and field guides ([46-48](#_ENREF_46), [51-56](#_ENREF_51)) as these were the most common measures available. All lengths were then converted to mass using family-level allometric scaling ([57](#_ENREF_57)). Prey size data was included from dietary studies when available. When prey size was not reported in the dietary studies and were prey species were identified to the species level, we used mean prey species body mass from available databases ([44](#_ENREF_44), [57](#_ENREF_57), [58](#_ENREF_58)). In cases were only body lengths were available for prey species allometric scaling were used to convert to mass ([44](#_ENREF_44), [59](#_ENREF_59)). For species that were only identified to the genus level the genus mean body mass was used if possible. The estimate mean prey size for each snake species was then calculated using a weighted mean based on the proportion each prey species/genus or group within the diet.

Snake mass, prey mass, LD50, venom volume and phylogenetic distance between diet and model were all log10 transformed, mean centered and expressed in units of standard deviation prior to analysis. Significance was determined for the fixed effects when 95% of the data is greater or less than 0. The phylogeny from Pyron RA & Burbrink ([60](#_ENREF_60)) was included in all analyses to account for similarities in traits due to common descent.

**Analysis**

To test our hypotheses we fit Bayesian multivariate phylogenetic mixed models using the MCMCglmm package ([61](#_ENREF_61)) in R v 3.2.4 ([62](#_ENREF_62)). As venom volume and LD50 are likely to have co-evolved, both were included as correlated (??) multivariate normal response variables in all models. Phylogeny was controlled by including it using the animal term in the MCMCglmm model while variation due to multiple measures on individual species was included using a separate random term. For the main model snake body mass; LD50 inoculation method (SC, IM, IV, IP); habitat dimensionality (2D, 3D); the presence of eggs in the diet (absent, present); and the phylogenetic distance of diet species to LD50 model were included as explanatory variables to give the analysis;

1. Volume + LD50 = *f*(Snake mass + LD50 method + Presence of eggs in diet + Phylogenetic distance between diet and model species + Habitat dimensionality) (275 observations over 99 species).

A similar model was also fit including prey size as an explanatory model which resulted in a smaller dataset of 177 observations across 68 species. To estimate the direct scaling exponents relating to prey mass, venom volume and predator mass as referred to in equations (1 and 3) we also ran the following model;

1. Volume = *f*(Prey mass)
2. Prey mass = *f*(Predator mass)

Finally, we also fitted a final set of sensitivity analysis including the main model with constriction behavior included as a categorical factor (absent, present) and a model with habitat type included instead of habitat dimension with the levels of terrestrial; aquatic and arboreal.

All models were fitted with parameter expanded priors (Hedfield 2010) with standard non-informative priors also tested separately to ensure that choice of prior had no affect on model results. A burn-in, thinning and number of iterations was determined for each model separately to ensure effective sample sizes exceeded 1000 for all parameter estimates. We tested for convergence using the Gelman-Rubin statistic over three separate chains ([63](#_ENREF_63)).

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