**Snake venom potency and volume are driven by metabolism, dimensionality and prey characteristics**

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**Significance**

Snake venom is best known for its ability to incapacitate prey, a property that makes it of both biomedical interest and public health concern. However, there is considerable variation in both the volume and potency of venom among snake species, which offers an opportunity to understand a novel aspect of predator-prey co-evolution. We investigate how this variation in snake venom is related to key characteristics of their prey, snake size, metabolic rates and complexity (dimensionality) of their habitat. Here, using comparative analysis, we show that venom has higher potency towards species closely resembling their natural diet, suggesting a co-evolution of predator to prey types, while macroecological drivers, such as habitat dimensionality and metabolic constraints shape the quantity of venom available.

**Abstract (250 words – 250 max)**

Snake venom is best known for its ability to incapacitate and kill prey. However, venom potency and volume vary greatly across species, with some species possessing venoms which are seemingly harmless, while others carry enough venom to kill vast numbers of potential prey. This highlights the need for a multi-species comparison to identify possible environmental and ecological drivers of snake venom evolution. However, studies commonly use non-native prey species as models to assess venom potency, an approach which may confound comparative analyses if the toxic effects of each snake species venom is adapted to its specific prey. Here we compare 99 species of snakes to test a range of hypotheses relating to the drivers of venom evolution. We assess potency (LD50) by accounting for the phylogenetic distance between the often non-natural model species used to measure potency and the species naturally occurring in their diets. We also examine variation in yield among species in relation to biological and habitat characteristics. We show that snake venom potency is prey-specific, with higher potencies when venoms were tested on species phylogenetically similar to common prey species. We also show that venom yield scales positively with snake body mass supporting a link with metabolic rate, but is lower for species found in high dimensionality habitats. These results underline the importance of ecological, physiological and environmental factors in the evolution of novel predatory traits and highlight the wider potential of using venom as a system to understand the evolution of predator-prey coevolution more generally.

**Keywords:** Venom, Body size, Comparative analysis, Scaling, trophic ecology, Macroecology, LD50, phylogenetic analysis

**Introduction**

Snake venom is perhaps best known for its ability to incapacitate and disrupt the physiological systems of animals. This is particularly well demonstrated by extremely lethal species, such as Russel’s viper (*Daboia russelii*), which possess enough potent venom to incapacitate tens of thousands of potential prey items (Figure 1). From a human perspective this property of venom has made it not only a source of novel biomedical compounds ([1](#_ENREF_1)) but also a major health concern, with snakebites estimated to cause up to 94,000 deaths annually ([2](#_ENREF_2)). Although the most lethal venomous snake species often gain the most attention ([3](#_ENREF_3)), the prey incapacitating ability of different species venoms ranges widely, from those unable to subjugate prey larger than a few grams to those capable of subduing vast numbers of laboratory animals ([4](#_ENREF_4)) (Figure 1). While understanding this variation is important from both a medical ([2](#_ENREF_2)) and evolutionary viewpoint ([1](#_ENREF_1)), little is known about what drives it. One reason for this is the lack of multi-species comparisons across taxonomically diverse groups. For example, while several studies have explored whether venoms have evolved to specifically target particular prey species these studies are typically focused at the genus level ([5-8](#_ENREF_5)), making general inferences regarding the evolution of fundamental aspects of venom characteristics difficult ([9-11](#_ENREF_9)). Here we conduct a comparative analysis across a taxonomically and ecologically diverse range of venomous snakes allowing us to test fundamental aspects of both the evolution of venom and predator traits in general.

Variation in predatory traits, are typically associated with differences in trophic ecology. For example, selection on jaw and beak morphology in cichlid fish and birds is strongly associated with trophic factors such as prey type ([12](#_ENREF_12), [13](#_ENREF_13)). Apart from prey type, other components of trophic interactions such as search and encounter rates ([14-16](#_ENREF_14)); the ability to spot, track and capture prey ([17](#_ENREF_17), [18](#_ENREF_18)); and the rate of ingestion of such captured prey ([16](#_ENREF_16), [19](#_ENREF_19" \o "Carbone, 2014 #93)) are also likely to influence predatory traits. However, while morphological measures of trophic traits can be linked to such potential drivers ([13](#_ENREF_13)) it is difficult to accurately quantify how changes in morphology map to trophic functional ability. Snake venom however offers a system were foraging capabilities can be quantified directly by measuring both venom potency, such as by measuring the median lethal dose (LD50), and the quantity available. This direct measure of the predatory ability of venom allows for the fundamental evolutionary drivers of venom and predator traits in general to be tested. However, to compare such measures of venom potency across snakes the route of administration (Figure 1) and the species used to test toxicity must be standardized in some way so as not to confound comparisons ([5](#_ENREF_5)). If venom characteristics are adapted to incapacitate prey species commonly found in their diets, we might expect the use model species not typical of a snake species diet would result in the underestimation of its potency ([1](#_ENREF_1)). However, the confounding effects of using measures of potency on standard laboratory model species can be minimized by controlling for the phylogentic distance between the model species and those prey species naturally found in the snakes diet. Furthermore, this approach not only provides for the corrections to allow for comparisons across species with diverse diets but also acts as a test of whether venom evolution results in prey-specific characteristics in general.

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Figure 1. Histogram and density plot of the animal mass (kg) that snakes species can impart a 50% mortality rate on for 282 observations of 104 species). This was calculated as the average volume of dried venom divided by its as LD50 (mg/kg). Measurement routes of LD50 administration are shown for intraperitoneal (light blue), intramuscular IM (dark blue), intravenous (red) and subcutaneous (orange). Species form left to right are *Thamnophis elegans, Emydocephalus annulatus, Causus rhombeatus, Atractaspis bibronii, Hydrophis elegans, Agkistrodon piscivorus, Ophiophagus hannah, Daboia russelii, Bungarus multicinctus, Oxyuranus scutellatus*.

Whether snake venom potency is prey-specific as a general rule has been the subject of much debate ([9-11](#_ENREF_9)). While prey-specificity has been demonstrated in several groups ([5-8](#_ENREF_5), [20](#_ENREF_20)), other examples have shown either no relationship between prey and venom lethality ([21](#_ENREF_21)), or cases were the prey species have evolved tolerance towards their predators venoms ([22](#_ENREF_22), [23](#_ENREF_23)). Of these outcomes predator-prey arms race dynamics ([24](#_ENREF_24)) predicts cases of prey-specific venoms and the evolution of prey tolerances depending on the level of selection on both predator and prey ([24](#_ENREF_24)). In contrast no relationship between potency and prey identity is predicted by the overkill hypothesis ([9-11](#_ENREF_9)) (Figure 2). The overkill hypothesis is based on the observation that venoms often display levels of lethality far in excess of the requirements needed to kill prey (Figure 1), resulting in predator-prey dynamics playing a minor role in the evolution of venom potency due to weak selection. As these hypotheses predict different outcomes when non-native species are used to test potency, the inclusion of the phylogenetic distance between the model species and natural prey not only allows comparison across species but also provides a test between these competing hypothesis (Figure 2). In particular, no relationship between venom potency phylogentic distance, would support the overkill mechanism, while if venom was evolved to be prey-specific, we would be expected to see higher potencies in model species more closely related to their typical prey species (Figure 2).

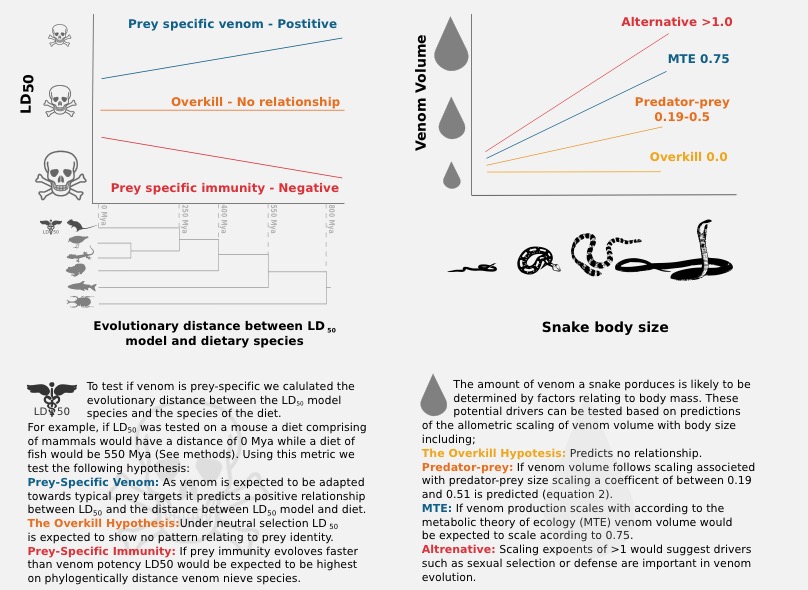


Figure 2. Summary of hypothesis relating to potential drivers of snake venom evolution.

As the ability to incapacitate prey is also determined by the amount of venom available it would also be predicted that venom volume would be under similar selective pressures. In particular, as venom production incurs an energetic cost ([25](#_ENREF_25)) (although the level of this cost is debated ([26](#_ENREF_26))) and requires storage, the volume of venom a snake species can produce is likely to be linked to prey size and or metabolic rate. Both prey size and metabolic rate are strongly determined by body size ([9](#_ENREF_9), [19](#_ENREF_19)). In terms of prey size, in general larger terrestrial predators eat larger prey ([19](#_ENREF_19)). It would hence be expected that larger snake species would need to produce larger quantities of venom to keep pace with subsequent increases in prey size. Metering of venom in response to prey size seen in several species ([27](#_ENREF_27), [28](#_ENREF_28)) supports the presence of such selection on venom quantity. However, while bigger snakes are known to have larger amounts of venom in general ([4](#_ENREF_4)) it is not known whether venom yield scales interspecifically according to any general pattern.

The overkill hypothesis would predict no relationship between venom yield and prey size, or a scaling exponent with snake mass of 0. Alternatively, another prediction is that venom yield increases with snake body size with an exponent relating to their predator-prey body size scaling such as described by (equation 1);

Where for snakes the scaling exponent *a* is approximately 0.68 ([19](#_ENREF_19)). However, venom yield would not be expected to scale according to this exponent as the effects of toxicological agents also follows an allometric relationship ([29](#_ENREF_29)) where the amount of venom required (*V*) to induce the same incapacitating effect on a prey of mass (*Mprey*) would be expected to follow (equation 2);

where *b* is the scaling coefficient, commonly estimated as 0.75, of venoms’ toxicological effects ([29](#_ENREF_29)). Hence to calculate the expected allometry of venom yield with snake body mass in a case where yield increases at a rate to match predator-prey size scaling after accounting for scaling of toxicological effects we substitute from equation 1 for to get (equation 3);

If we take the commonly used value of 0.75 for the interspecific scaling of drug dosages ([29](#_ENREF_29)) for *b* and the value of 0.68 for *a* from the scaling predator-prey mass relationship for snakes ([19](#_ENREF_19)) we would expect a scaling exponent of approximately 0.51 between snake venom yield and snake mass. Otherwise venom yield may scale according to constraints such as metabolic costs matched with metabolic rate, where a scaling of 0.75 would be expected ([30](#_ENREF_30)). At the other extreme super-linear allometries (exponents >1) would suggest patterns associated with drivers such as sexual selection, such as proposed by the weapons hypothesis ([31](#_ENREF_31)), or defenses requiring increased effectiveness with size, such as seen in the allometry of horn growth in horned lizards ([32](#_ENREF_32)) (Figure 1).

Finally, an overlooked feature that may also drive the evolution of both venom quantity and toxicity is habitat structure ([33](#_ENREF_33)). 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 ([15](#_ENREF_15), [19](#_ENREF_19)) and the escape rates of prey, with higher dimensional spaces increasing both ([34](#_ENREF_34), [35](#_ENREF_35)). Hence predators in high dimensional habitats with associated increased escape rates may compensate through larger yields 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 large reservoirs of venom 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, as measured using median lethal dose (LD50), in a phylogenetically corrected comparative analysis of ninety-nine species of venomous snakes. Using the phylogenetic distance between species used to measure LD50 and dietary species, we test;

1. the overkill hypothesis: that there is no relationship between venom potency and the species on which it was measured or between venom yield and 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 yields.
3. the importance of macorecological drivers on venom evolution with scaling of venom yield with snake body mass predicted from predator-prey size scaling to be approximately 0.51; from metabolic constraints to be 0.75; and from other potential drivers such as sexual selection and the weapons hypothesis to be superlinear.
4. the importance of habitat dimensionality on venom evolution, in particular that; species in high dimensional habitats show either higher or lower potencies.

**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 where we included whether species to are known to using constricting behaviors in place of or augmenting venom delivery; 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 (Table A1).

**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 yield had a negative, but non-significant, association with ovivorous behavior in all models (Figure 2; Tables A1-5).

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 such that snake species with diets phylogenetically close to the LD50 model species having higher potencies (Figure 2; table A3-4; Figure 1B). From the main model, after back-transforming the mean centered 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. The route venom was administered was also found to affect values of LD50 where intravenous and Intraperitoneal routes were found to have lower LD50 values in comparison to a subcutaneous route (Figure 2; Table A1-3).

Figure 3. Posterior distributions of LD50 and mean venom volume estimates (represented by dots) and higher and lower 95% credibility intervals (represented by dotted horizontal bar). Fixed factors include mass; LD50 method (subcutaneous (SC), intravenous (IV), intrapulmonary (IP) and intramuscular (IM)); habitat dimensionality (Dim- 2D and 3D); Presence of eggs in diet (Eggs in Diet) and the mean phylogenetic distance between diet species and the LD50 model (Diet-LD50 Dist). The random terms and the co-variance (CV) between LD50 and yield are also presented. Signifiance is determined when 95% of the data is above or below zero. The model was run with 12,000,000 iterations with a 2,000,000 burn-in and a thinning of 5000.

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

The mean yield 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 yield. The main correlate with average yield was snake body size with a log-log scaling of between 0.74 and 0.76 across all models, after back-transforming (Figure 3, Tables A2-6), an exponent far higher than the predicted scaling of 0.51 for predator-prey body size scaling (equation 3). Venom yield also showed a positive increase with prey body mass, with a log10-log10 slope of 0.139 (equating to approximately a 10% increase in venom volume with a doubling of body mass), however only 90% of the posterior samples are above the zero threshold (Table A3). Snake body size was also found to have a significantly positively correlation with LD50 in the main analysis of 99 species, meaning larger snakes showed decreased potency (Figure 3). However, this relationship was not significant in any of the sub analysis (Table A3-4).

The next most significant driver of venom yield was the dimensionality of the habitat with the 27 species in high dimensional environments (arboreal = 9, aquatic = 18) showing lower venom yields in comparison to species in lower dimensional habitats (Figure 2). A sensitivity analysis where habitat was included as terrestrial, arboreal and aquatic also showed similar significant reductions in both arboreal and aquatic habitats (Table A5).

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Figure 4. (A) Mean phylogenetic distance between diet species and LD50 model (Myr) against log10 LD50 (intercept = -0.58, slope = 0.002). Hollow points represent silhouette species which are from left to right; *Bungarus multicinctus*; *Oxyuranus microlepidotus*; *Echis carinatus*; *Causus rhombeatus*.

(B) Relationship between log10 mass (g) against log10 venom yield (mg). Red points and fitted line (intercept = -0.58, slope = 0.75) represent species in 2D habitats and the blue points and fitted line (intercept = -1.14, slope = 0.75) represent species in 3D habitats. Hollow points represent silhouette species which are from left to right *Atractaspis bibronii*; *Emydocephalus annulatus*; *Naja\_melanoleuca*; *Agkistrodon piscivorus*; *Ophiophagus hannah*. All intercepts and slopes are back transformed from the values in Figure 3 and Table A1.

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

In all models there is was an intermediate to high phylogenetic signal for both LD50 and venom yield with phylogeny explaining 0.69 of the residual variation for LD50 and 0.49 for venom yield, with LD50 showing a higher phylogenetic signal in all models (Figure 2, A2-5). The presence of constricting behaviour was found to have no effect on either yield or LD50(Table A4). Finally, there is no significant covariance between either the residuals or phylogenetic terms of both yield and LD50 across all models (Figure 2, A2-5).

**Discussion**

Predator traits are predicted to be strongly shaped by both predator-prey co-evolution and macroecological forces such as body size and habitat structure. Traits such as jaw morphology are inextricably linked to diet (ref), while the size, encounter rate and escape rate of potential prey is limited by predators size and the habitat in which it forages (refs). Here we find that snake venom follows such expected patterns finding with venom haining an increased potency when tested on species more closely resembling target prey while body size and habitat dimensionality influenced the quantity of venom available to a snake species. These results demonstrate that the trophic traits of an ecological and phylogenetical diverse taxon follows general predictable patterns. By using venom as a system where the trophic ability of a trait can be quantified and by accounting for factors affect its comparability across species, our findings show that a trait targeting diverse prey species, ranging from insects to other snakes,

patterns consistent with the notion that it has evolved under predator-prey selection pressures. Venom potency was found to be prey-specific and in general the scaling of venom volume, paralleled that snake metabolism. In addition venon volume decreased with increasing habitat dimensionality. Overall, these results show that by controlling for the model species used to measure potency in comparative analysis across large taxonomic groups, hypothesis regarding the evolutionary drivers of venom can be robustly tested.

The importance of trophic factors in the evolutionary maintenance of venom is well demonstrated by cases of dietary switches to immobile or unprotected prey, such as seen by the almost complete atrophy of the venom apparatus in the marbled sea snake (*Aipysurus eydouxii*) due switiching to an egg based diet ([36](#_ENREF_36)). This maintenance mechanism is also supported by our findings that ovivorous feeding is associated with lower potencies and venom yield. However, beyond the role of maintaining the possession of venom the role of trophic drivers in shaping venoms subsequent evolution has been much more hotly debated, in particularly relating to the generality of prey-specific venoms ([1](#_ENREF_1), [9](#_ENREF_9), [10](#_ENREF_10)). By accounting for the phylogenetic relatedness between the model species and natural prey when estimating venom potency ([6](#_ENREF_6)) we show that venom is generally prey-specific. Moreover, our analysis is likely to underestimate the generality of the prey-specific nature of venom 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, in order to reduce the chances of prey escaping or retaliating ([5](#_ENREF_5)). Even though prey mortality achieves this, other measures of venoms ability to subdue prey, such as the speed at which a venom affects prey or measures of sub-lethal incapacitating effects, may further our understanding of how predator-prey dynamics shape venom potency ([5](#_ENREF_5)). However, despite such limitations, we find a clear pattern of prey-specificity suggesting that cases of non-prey specific venom ([21-23](#_ENREF_21), [37](#_ENREF_37), [38](#_ENREF_38)) are more likely to be exceptions to the general rule.

In terms of macroecological patterns, unsurprisingly we found that larger snakes had larger quantities of venom. However, these increases did not follow the scaling of 0.51 predited from the scaling of prey size with respect to predator size ([19](#_ENREF_19)). Venom yield increased more steeply than this. Even when considering potential variation in the allometric scaling of toxicological effects (equation 3) ([29](#_ENREF_29)), a value far in excess of 1 is required for the scaling of toxicological effects (*b*), in order for our results to match the predator-prey scaling prediction: a situation that seems unlikely. Furthermore, our analysis revealed that the predatory-prey body mass scaling of venomous snakes, (*a* in equation 3) is much lower than expected, based on comparisons to previous body mass scaling conducted on all snakes ([19](#_ENREF_19)), meaning that larger venomous snakes feed on much smaller prey items than predicted for their size. This lower scaling of prey size may explain the reduced venom potency with snake size seen in our main analysis.

While venom yield did not follow predictions arising from predator-prey body mass scaling it did match an allometry of 0.75 expected from metabolic theory assuming snakes invest a constant proportion of their metabolism to produce venom ([30](#_ENREF_30), [39](#_ENREF_39)). However, other studies have also showed that the amount administered in a single bite is closely related to prey size ([28](#_ENREF_28)). As yield scales with body size according to a higher exponent than prey size, larger species would be expected to have the capacity to envenomate more prey items before depleting their reservoir in comparison to smaller species which may be constrained to something closer to a one shot strategy.

Habitat dimensionality was another factor found to influence the volume of venom. We expected that species in high dimensional habitats may have higher venom yields to compensate for higher escape rates of prey ([33](#_ENREF_33)), however, we found that these species had lower yields 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 ([40](#_ENREF_40)), 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 ([41](#_ENREF_41)). Another potential explanation is that higher encounter rates in high dimensional environments ([15](#_ENREF_15)) may reduce the missed opportunity of feeding cost associated with replenishing venom. Rates of replenishing venom can be substantial with estimates ranging from 3-7 days ([42](#_ENREF_42)) to 30-50 days ([28](#_ENREF_28), [43](#_ENREF_43)). These long periods of replenishment may hence select for larger reserves in species where prey encounter rates are low 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 behind this difference our results highlight that prey encounter rates may be an important factor in venom evolution.

While our analysis demonstrates the importance of trophic and macroecological drivers in snake venom evolution these drivers are also expected to influence the evolution of venom in other taxa ([1](#_ENREF_1)). For example, prey-specific venom is seen in cone snails and spiders ([1](#_ENREF_1)), while the energetic costs of producing venom is also suggested by venom metering in scorpions ([44](#_ENREF_44)). Future analysis that include other venomous taxa in a comparative approach such as used here, will further test whether venom fundamentally follows such similar patterns. Certain elements of prey-specify and macroecological constraints are also likely to generally apply across other non-venomous predatory traits. For example, possible predator-prey arms dynamics relating to bite force and prey size ([45](#_ENREF_45)), or macroecological constraints relating to pursuit speed ([14](#_ENREF_14)). By using venom as a system of predator trait evolution the importance of multiple evolutionary drivers can be robustly tested and hence offer a window not only into the evolution of venomous systems, but of predatory traits and trophic ecology as a whole.

**Methods**

**Data**

We collected data on venom yield and toxicity from the literature, along with our predicted drivers. We used mean dry weight (mg) extracted as a measure of venom yield 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) due to its wide availability. We only included intravenous (IV), subcutaneous (SC), Intraperitoneal(IP) or intramuscular routes (IM) of administering 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.

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 ([46](#_ENREF_46)), between each prey taxa 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 ([15](#_ENREF_15)), 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 ([40](#_ENREF_40)).

For snake body size we used total length values from the literature and field guides as these were the most common measures available (See Appendix A). All lengths were then converted to mass using family-level allometric scaling ([47](#_ENREF_47)). 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 ([47-49](#_ENREF_47)). In cases were only body lengths were available for prey species allometric scaling were used to convert to mass ([48](#_ENREF_48), [50](#_ENREF_50)). 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 yield 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 ([51](#_ENREF_51)) was included in all analyses to account for non-independence in traits due to common descent.

**Analysis**

To test our hypotheses we fitted Bayesian multivariate phylogenetic mixed models using the MCMCglmm package ([52](#_ENREF_52)) in R v 3.2.4 ([53](#_ENREF_53)). As venom yield and LD50 are likely to have co-evolved, both were included as response variables in a series of multivariate analysis. 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. The relative variance attributable to the phylogenetic random effect component (*H* 2) was calculated as the ratio of variance explained by phylogeny to the sum of phylogenetic variance, species variance and residual variance. 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. Yield + 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 fitted 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 yield and predator mass as referred to in equations (1 and 3) we also ran the following model;

1. Yield = *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 effect 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 ([54](#_ENREF_54)).

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**References**

Uncategorized References

1. Casewell NR, Wüster W, Vonk FJ, Harrison RA, & Fry BG (2013) Complex cocktails: the evolutionary novelty of venoms. *Trends in ecology & evolution* 28(4):219-229.

2. Kasturiratne A*, et al.* (2008) The global burden of snakebite: a literature analysis and modelling based on regional estimates of envenoming and deaths. *PLoS Med* 5(11):e218.

3. Weinstein SA, Warrell DA, White J, & Keyler D (2011) Venomous" bites from non-venomous snakes. *A Critical Analysis of Risk and Management of “Colubrid” Snake Bites*, (Elsevier London).

4. Chippaux J-P, Williams V, & White J (1991) Snake venom variability: methods of study, results and interpretation. *Toxicon* 29(11):1279-1303.

5. Barlow A, Pook CE, Harrison RA, & Wüster W (2009) Coevolution of diet and prey-specific venom activity supports the role of selection in snake venom evolution. *Proceedings of the Royal Society of London B: Biological Sciences* 276(1666):2443-2449.

6. da Silva NJ & Aird SD (2001) Prey specificity, comparative lethality and compositional differences of coral snake venoms. *Comparative Biochemistry and Physiology Part C: Toxicology & Pharmacology* 128(3):425-456.

7. Daltry JC, Wuester W, & Thorpe RS (1996) Diet and snake venom evolution. *Nature* 379(6565):537-540.

8. Starkov VG, Osipov AV, & Utkin YN (2007) Toxicity of venoms from vipers of Pelias group to crickets Gryllus assimilis and its relation to snake entomophagy. *Toxicon* 49(7):995-1001.

9. Sasa M (1999) Diet and snake venom evolution: can local selection alone explain intraspecific venom variation? *TOXICON-OXFORD-* 37:249-252.

10. Mebs D (2001) Toxicity in animals. Trends in evolution? *Toxicon* 39(1):87-96.

11. WuÈster W, Daltry JC, & Thorpe RS (1999) Can diet explain intraspecific venom variation? Reply to Sasa. *TOXICON-OXFORD-* 37:253-258.

12. Albertson RC, Markert J, Danley P, & Kocher T (1999) Phylogeny of a rapidly evolving clade: the cichlid fishes of Lake Malawi, East Africa. *Proceedings of the National Academy of Sciences* 96(9):5107-5110.

13. Cooney CR*, et al.* (2017) Mega-evolutionary dynamics of the adaptive radiation of birds. *Nature*.

14. Domenici P (2001) The scaling of locomotor performance in predator–prey encounters: from fish to killer whales. *Comparative Biochemistry and Physiology Part A: Molecular & Integrative Physiology* 131(1):169-182.

15. Pawar S, Dell AI, & Savage VM (2012) Dimensionality of consumer search space drives trophic interaction strengths. *Nature* 486(7404):485-489.

16. Kane A*, et al.* (2016) Body Size as a Driver of Scavenging in Theropod Dinosaurs. *The American Naturalist* 187(6):706-716.

17. Healy K, McNally L, Ruxton GD, Cooper N, & Jackson AL (2013) Metabolic rate and body size are linked with perception of temporal information. *Animal behaviour* 86(4):685-696.

18. Kiltie R (2000) Scaling of visual acuity with body size in mammals and birds. *Functional Ecology* 14(2):226-234.

19. Carbone C, Codron D, Scofield C, Clauss M, & Bielby J (2014) Geometric factors influencing the diet of vertebrate predators in marine and terrestrial environments. *Ecology letters* 17(12):1553-1559.

20. Richards D, Barlow A, & Wüster W (2012) Venom lethality and diet: differential responses of natural prey and model organisms to the venom of the saw-scaled vipers (Echis). *Toxicon* 59(1):110-116.

21. Williams V, White J, Schwaner T, & Sparrow A (1988) Variation in venom proteins from isolated populations of tiger snakes (Notechis ater niger, N. scutatus) in South Australia. *Toxicon* 26(11):1067-1075.

22. Heatwole H & Poran NS (1995) Resistances of sympatric and allopatric eels to sea snake venoms. *Copeia*:136-147.

23. Voss RS (2013) Opossums (Mammalia: Didelphidae) in the diets of Neotropical pitvipers (Serpentes: Crotalinae): Evidence for alternative coevolutionary outcomes? *Toxicon* 66:1-6.

24. Van Valen L (1973) A new evolutionary law. *Evolutionary theory* 1:1-30.

25. McCue MD & Mason R (2006) Cost of producing venom in three North American pitviper species. *Copeia* 2006(4):818-825.

26. Pintor AF, Krockenberger AK, & Seymour JE (2010) Costs of venom production in the common death adder (Acanthophis antarcticus). *Toxicon* 56(6):1035-1042.

27. Hayes WK (1995) Venom metering by juvenile prairie rattlesnakes, Crotalus v. viridis: effects of prey size and experience. *Animal Behaviour* 50(1):33-40.

28. Hayes WK, Herbert SS, Rehling GC, & Gennaro JF (2002) Factors that influence venom expenditure in viperids and other snake species during predatory and defensive contexts. *Biology of the Vipers*:207-233.

29. Nestorov I (2003) Whole body pharmacokinetic models. *Clinical pharmacokinetics* 42(10):883-908.

30. Isaac NJ & Carbone C (2010) Why are metabolic scaling exponents so controversial? Quantifying variance and testing hypotheses. *Ecology Letters* 13(6):728-735.

31. Kodric-Brown A, Sibly RM, & Brown JH (2006) The allometry of ornaments and weapons. *Proceedings of the National Academy of Sciences* 103(23):8733-8738.

32. Bergmann PJ & Berk CP (2012) The evolution of positive allometry of weaponry in horned lizards (Phrynosoma). *Evolutionary Biology* 39(3):311-323.

33. Arbuckle K (2015) Evolutionary Context of Venom in Animals.

34. Heithaus MR, Wirsing AJ, Burkholder D, Thomson J, & Dill LM (2009) Towards a predictive framework for predator risk effects: the interaction of landscape features and prey escape tactics. *Journal of Animal Ecology* 78(3):556-562.

35. Møller A (2010) Up, up, and away: relative importance of horizontal and vertical escape from predators for survival and senescence. *Journal of evolutionary biology* 23(8):1689-1698.

36. Li M, Fry B, & Kini RM (2005) Eggs-only diet: its implications for the toxin profile changes and ecology of the marbled sea snake (Aipysurus eydouxii). *Journal of Molecular Evolution* 60(1):81-89.

37. Biardi JE & Coss RG (2011) Rock squirrel (Spermophilus variegatus) blood sera affects proteolytic and hemolytic activities of rattlesnake venoms. *Toxicon* 57(2):323-331.

38. Poran NS, Coss RG, & Benjamini E (1987) Resistance of California ground squirrels (Spermophilus beecheyi) to the venom of the northern Pacific rattlesnake (Crotalus viridis oreganus): a study of adaptive variation. *Toxicon* 25(7):767-777.

39. Brown JH, Gillooly JF, Allen AP, Savage VM, & West GB (2004) Toward a metabolic theory of ecology. *Ecology* 85(7):1771-1789.

40. Shine R & Schwaner T (1985) Prey constriction by venomous snakes: a review, and new data on Australian species. *Copeia* 1985(4):1067-1071.

41. Branch WR (1998) *Field guide to snakes and other reptiles of southern Africa* (Struik).

42. Currier RB*, et al.* (2012) Unusual stability of messenger RNA in snake venom reveals gene expression dynamics of venom replenishment. *PloS one* 7(8):e41888.

43. Hayes W (2008) The snake venom-metering controversy: levels of analysis, assumptions, and evidence. *The biology of rattlesnakes*:191-220.

44. Nisani Z, Dunbar SG, & Hayes WK (2007) Cost of venom regeneration in Parabuthus transvaalicus (Arachnida: Buthidae). *Comparative Biochemistry and Physiology Part A: Molecular & Integrative Physiology* 147(2):509-513.

45. Wroe S, McHenry C, & Thomason J (2005) Bite club: comparative bite force in big biting mammals and the prediction of predatory behaviour in fossil taxa. *Proceedings of the Royal Society of London B: Biological Sciences* 272(1563):619-625.

46. Hedges SB, Dudley J, & Kumar S (2006) TimeTree: a public knowledge-base of divergence times among organisms. *Bioinformatics* 22(23):2971-2972.

47. Meiri S (2010) Length–weight allometries in lizards. *Journal of Zoology* 281(3):218-226.

48. Feldman A & Meiri S (2013) Length–mass allometry in snakes. *Biological Journal of the Linnean Society* 108(1):161-172.

49. Myhrvold NP*, et al.* (2015) An amniote life‐history database to perform comparative analyses with birds, mammals, and reptiles. *Ecology* 96(11):3109-3109.

50. Pough FH (1980) The advantages of ectothermy for tetrapods. *American Naturalist*:92-112.

51. Pyron RA & Burbrink FT (2014) Early origin of viviparity and multiple reversions to oviparity in squamate reptiles. *Ecology Letters* 17(1):13-21.

52. Hadfield JD (2010) MCMC methods for multi-response generalized linear mixed models: the MCMCglmm R package. *Journal of Statistical Software* 33(2):1-22.

53. Team RC (2016) R: A Language and Environment for Statistical Computing. R Foundation for Statistical Computing.

54. Brooks SP & Gelman A (1998) General methods for monitoring convergence of iterative simulations. *Journal of computational and graphical statistics* 7(4):434-455.