

RH: Skink Anomaly Zone

Detecting the anomaly zone in species trees and evidence for a misleading signal in higher-level skink phylogeny (Squamata: Scincidae).

CHARLES W. LINKEM¹, VLADIMIR N. MININ^{1,2}, AND ADAM D. LEACHÉ^{1,3}

¹*Department of Biology, University of Washington, Seattle, Washington, 98195, USA.*

²*Department of Statistics, University of Washington, Seattle, Washington, 98195, USA.*

³*Burke Museum of Natural History and Culture, University of Washington, Seattle, Washington, 98195, USA.*

Corresponding author: Charles W. Linkem, Department of Biology, University of Washington, Box 351800, Seattle, Washington, 98195-1800, USA; E-mail: cwlinkem@gmail.com.

Abstract.—The anomaly zone, defined by the presence of gene tree topologies that are more probable than the true species tree, presents a major challenge to the accurate resolution of many parts of the Tree of Life. This discrepancy can result from consecutive rapid speciation events in the species tree. Similar to the problem of long-branch attraction, including more data via loci concatenation will only reinforce the support for the incorrect species tree. Empirical phylogenetic studies often employ coalescent-based species tree methods to avoid the anomaly zone, but to this point these studies have not had a method

for providing any direct evidence that the species tree is actually in the anomaly zone. In this study, we use 16 species of lizards in the family Scincidae to investigate whether nodes that are difficult to resolve place the species tree within the anomaly zone. We analyze new phylogenomic data (429 loci), using both concatenation and coalescent-based species tree estimation, to locate conflicting topological signal. We then use the unifying principle of the anomaly zone, together with estimates of ancestral population sizes and species persistence times, to determine whether the observed phylogenetic conflict is a result of the anomaly zone. We identify at least three regions of the Scincidae phylogeny that provide demographic signatures consistent with the anomaly zone, and this new information helps reconcile the phylogenetic conflict in previously published studies on these lizards. The anomaly zone presents a real problem in phylogenetics, and our new framework for identifying anomalous relationships will help empiricists leverage their resources appropriately for investigating and overcoming this challenge. (Keywords: anomalous gene trees, IDBA, incomplete lineage sorting, phylogenetics, probes, sequence capture, ultraconserved elements, UCE)

The field of phylogenetics is poised to benefit tremendously from genomics, since resolving evolutionary relationships often requires massive amounts of data. Empirical phylogenetic researchers foresaw genomic data as a holy grail for resolving difficult relationships, such as rapid speciation events (Rokas et al. 2003; Dunn et al. 2008; Edwards 2009). However, phylogenetic conflict persists with genomic data, often with greater support, generating more questions than answers for many studies (Song et al. 2012;

Gatesy and Springer 2014; Springer and Gatesy 2014; Pyron et al. 2014; Zhang et al. 2014). In addition, reanalyses of previously published phylogenomic datasets often produce conflicting results (Dunn et al. 2008; Philippe et al. 2009, 2011), suggesting that analytical approach and model assumptions are critical to phylogenomic studies.

Processes such as horizontal gene transfer, gene duplication, and incomplete lineage sorting can lead to differences between species trees and gene trees (Maddison 1997). Species histories containing rapid diversification will have a high prevalence of incomplete lineage sorting due to few generations between speciation events. Rapid diversification in combination with a large effective population size can result in higher probabilities for gene trees that do not match the species tree than for gene trees that do match. These non-matching gene trees with high probability from incomplete lineage sorting are referred to as anomalous gene trees (AGT), and the species tree branches that produce them are considered to be in the anomaly zone (Degnan and Rosenberg 2006). When the demographic history of the species tree is in the anomaly zone, sampling independent loci from the genome will result in AGT topologies being recovered at higher frequency than gene trees that match the species tree. Concatenation of these independent loci will result in strong support for the AGT topology whereas analyses using a species tree approach may recover the correct species tree (Kubatko and Degnan 2007; Liu and Edwards 2009).

Coalescent theory characterizes the anomaly zone for a four-taxon tree (Degnan and Rosenberg 2006) by showing that short internal branch lengths for an asymmetric topology will result in high probability for a symmetric AGT (Figure 1). The boundary of the anomaly zone $a(x)$ in the four-taxon case is defined by the following equation:

$$a(x) = \log \left[\frac{2}{3} + \frac{3e^{2x} - 2}{18(e^{3x} - e^{2x})} \right], \quad (1)$$

where x is the length of the branch in the species tree that has a descendant internal

branch. If the length of the descendant internal branch, y , is less than $a(x)$, then the species tree is in the anomaly zone. As values of x get small, $a(x)$ goes to infinity and therefore the value of the descendant branch y can be very long and still produce AGTs. In the four-taxon case, values of x greater than 0.27 coalescent units are outside the anomaly zone, when $a(x)$ approaches zero, but this value increases with the addition of more taxa. In a four-taxon tree there is only one set of x and y internodes to consider and only three possible AGTs. With the addition of a single taxon, the five-taxon species tree has multiple sets of x and y and z internode branches that can have as many as 45 AGTs (Rosenberg and Tao 2008). The calculation of the multidimensional anomaly zone in trees larger than five taxa is impractical, but a conservative simplification of the theory can be used for any species tree (Rosenberg 2013).

Species tree topologies of five taxa or more can be broken up into sets of four-taxon trees, which can individually be used in the anomaly zone calculation of equation (1). Rosenberg (2013) showed that focusing on sets of consecutive internal branches consistent with a four-taxon topology is a conservative estimate of the presence of the anomaly zone in any species tree. If the set of internodes fits the anomaly zone for the four-taxon case, at least one AGT exists, though more AGTs may occur due to nearby branches not considered in the isolated calculation (Rosenberg 2013). The unifying principle of the anomaly zone, that the anomaly zone condition can be checked for any four-taxon case within a larger phylogeny independently of other subtrees, allows for the estimation of the anomaly zone in trees of any size.

The theoretical predictions of the anomaly zone are well characterized (Rosenberg and Tao 2008; Degnan and Rosenberg 2009; Rosenberg and Degnan 2010; Degnan et al. 2012a,b; Degnan 2013), and simulations have identified situations where certain phylogenetic methods succeed or fail under the anomaly zone (Steel and Rodrigo 2008; Huang and Knowles 2009; Liu and Edwards 2009; Liu et al. 2010b; Liu and Yu 2011; Liu

et al. 2015). Empirical studies of genomic data have shown that incomplete lineage sorting can be pervasive throughout the genome in rapidly speciating groups (Suh et al. 2015). However, an empirical example of the anomaly zone has yet to be demonstrated. The lack of empirical evidence for the anomaly zone has led to doubt about the influence of the anomaly zone on real data (Huang and Knowles 2009; Tonini et al. 2015), and the utility of coalescent methods for phylogenetic inference has been questioned (Gatesy and Springer 2014; Springer and Gatesy 2014). An investigation of the anomaly zone in an empirical setting requires an accurate species tree topology and estimates for ancestral branch lengths and population sizes, parameters that can be inferred accurately with hundreds of loci. Until recently, obtaining loci at this magnitude was not feasible for most non-model organisms, but new methods for obtaining large phylogenomic datasets are quickly changing the scale and scope of phylogenetic studies (Faircloth et al. 2012; McCormack et al. 2012; Song et al. 2012).

Here we present an empirical example of the anomaly zone in a diverse radiation of lizards, the Scincidae. Using theoretical expectations, we define the set of species tree branches expected to generate anomalous gene trees based on the multispecies coalescent, and we apply these predictions to an empirical species tree. We use a new phylogenomic dataset collected using sequence capture of ultraconserved elements (Faircloth et al. 2012) and protein-coding genes (Wiens et al. 2012) to estimate the species tree, branch lengths, and population sizes required for identifying the anomaly zone in an empirical phylogeny.

Scincidae

Scincidae is the largest family of lizards comprising >25% of all lizard species. This globally distributed family of 1,583 species (Uetz 2015) can be found from deserts to rain forests, across continents and on tiny Pacific islands (Greer 1970). This relatively ancient group (80–110 ma; Mulcahy et al. (2012)) exhibits extreme variation in body-form

evolution with repeated limb loss (Greer 1991; Skinner et al. 2008) and variation in reproductive biology through multiple transitions to live birth (Blackburn 1982). The first major division of higher-level skink taxonomy was largely based on skull morphology, dividing the family into four subfamilies: Acontinae, Felylinae, Lygosominae, and Scincinae (Greer 1970). Synapomorphies were defined for all but Scincinae, which was described as possessing the “primitive” form from which the other forms evolved, indicating Scincinae was not a natural group (Greer 1970). Molecular phylogenies have largely supported the morphological hypotheses of Lygosominae and Acontinae monophyly but have found Lygosominae and Feylinae nested within Scincinae (Whiting et al. 2003; Brandley et al. 2005, 2012; Wiens et al. 2012; Pyron et al. 2013; Lambert et al. 2015). A few molecular studies found paraphyly in Lygosominae (Whiting et al. 2003; Siler et al. 2011), though the majority of molecular phylogenies (Wiens et al. 2012; Pyron et al. 2013; Reeder 2003; Pyron et al. 2013; Lambert et al. 2015) and a series of morphological synapomorphies (Greer 1970, 1986) support the monophyly of this subfamily. Feylinae is no longer considered a valid subfamily and the species within it are part of the broader paraphyletic Scincinae (Whiting et al. 2003; Brandley et al. 2005). The relationships among the major groups within Scincinae have remained difficult to decipher (Brandley et al. 2005, 2012; Wiens et al. 2012; Pyron et al. 2013). Most studies have found Scincinae to be paraphyletic with the genus *Brachymeles* sister to Lygosominae, but two recent papers (Pyron et al. 2013; Lambert et al. 2015) found a monophyletic Scincinae.

Molecular phylogenies of skinks have sampled up to 44 genes for a few taxa (Wiens et al. 2012; Lambert et al. 2015) or a few genes for up to 683 taxa (Pyron et al. 2013) but still lack resolution or present strong conflict for many relationships. The deep branches across this group are often found to be short when estimated from individual genes and concatenated genes, indicating diversification may have been rapid in this group. Strong support for different relationships from concatenated gene trees and coalescent species trees

(Lambert et al. 2015) in combination with a history of rapid diversification indicate that speciation history may fit the demographic parameters consistent with the anomaly zone.

MATERIALS AND METHODS

Identification of the causes of gene tree—species tree conflict in empirical phylogenies requires a large number of sampled genes on taxa that span difficult nodes (Liu and Edwards 2009). To accomplish this, we use a sequence capture next-generation sequencing approach to obtain 429 independent loci for all taxa of interest.

Taxon sampling

Sampling was focused on fifteen species of skinks that span the deep nodes in the Scincidae tree where topological conflict is high. From Scincinae we include *Brachymeles bonitae*, *Chalcides ocellatus*, *Eurylepis taeniolatus*, *Mesoscincus manguae*, *Ophiomorus raithmai*, *Plestiodon fasciatus*, and *Scincus scincus*, which represent the broad diversity in this difficult-to-resolve subfamily. These species have been used in prior studies of skink relationships based on Sanger sequencing (Brandley et al. 2005, 2012; Pyron et al. 2013) and have shown very short internode lengths and multiple alternative topologies suggesting anomalous gene trees may be present. A single Acontinae sample, *Typhlosaurus sp.*, is used to represent this well-supported subfamily (Lamb et al. 2010). We include at least one sample of four of the five groups in Lygosominae (missing *Egernia* group): *Mabuya unimarginata* for the *Mabuya* group; *Lygosoma brevicaudis* for the *Lygosoma* group; *Emoia caeruleocauda* for the *Eugongylus* group; and *Lobulia elegans*, *Sphenomorphus tridigitus*, *Sphenomorphus variegatus*, and *Tytthoscincus parvus* for the *Sphenomorphus* group. Previous studies (Honda et al. 2003; Reeder 2003; Skinner 2007; Skinner et al. 2011) have shown variation in the relationships between the *Lygosoma*, *Mabuya*, and *Eugongylus*

groups, some with short internode lengths. A single outgroup taxon, *Xantusia vigils*, was chosen to root phylogenetic analyses.

Probe design, library preparation, sequencing, and data processing

We designed a unique set of sequence-capture probes targeting a combination of UCE loci and loci used in squamate phylogenetics. The subset of UCE probes used are derived from the *Tetrapods-UCE-5kv1* probe set (Faircloth et al. 2012) that most closely match the *Anolis* genome (Alföldi et al. 2011) and the *Sceloporus occidentalis* genome (Genomic Resources Development Consortium et al. 2014). Additionally, we designed 120 bp probes with 2x tiling (60 bp overlap) for the 44 genes used in the Squamate Tree of Life project (Wiens et al. 2012). This resulted in a sequence capture probe set targeting 585 single copy loci that can be used across squamates. Details of probe design and synthesis are available in the Supplemental Materials.

Whole genomic DNA was broken through sonication using a Bioruptor Pico (Diagenode Inc.) and prepared for sequencing using an Illumina Truseq Nano library preparation kit. Libraries were pooled into sets of eight and hybridized to the RNA probes using a MYBaits target enrichment kit (MYcroarray). Post-hybridized libraries were quantified using qPCR (Applied Biosystems Inc.) and quality was assessed using an Agilent TapeStation 2200 (Agilent Tech.). Samples were sequenced at the QB3 facility at UC Berkeley on an Illumina HiSeq2500 on a single lane 150 bp, paired-end rapid run. Details of library preparation, sequence-capture hybridization, and sequencing are available in the Supplemental Materials.

Sequence data were processed using *Illuminaprocessor* v.2.0 (Faircloth 2013) and *Trimmomatic* (Bolger et al. 2014) to remove low-quality reads, trim low-quality ends, and remove adapter sequence. De novo assembly for each species was conducted with the iterative de Bruijn graph short-read assembler IDBA (Peng et al. 2010) over a set of k-mer

values from 50 to 90 with a step length of 10. We used **Phyluce** (Faircloth et al. 2012; Faircloth 2014) to align species-specific contigs to the set of probes and create an **SQL** database of contig-to-probe matches for each taxon. We queried this database to create the datasets for the 429 loci that were sequenced across all taxa. Expanded details of data processing are available in the Supplemental Materials. Datasets and analysis results are available on Dryad (doi:10.5061/dryad.sf6s9). Sequences are available on Genbank (KU343288–KU350151).

Model testing, gene trees, and concatenation analyses

We identified the models of substitution in the 95% credible set of the BIC for each locus using **jModelTest v2.1.5** (Guindon and Gascuel 2003; Darriba et al. 2012). The model with the lowest BIC score was chosen as the preferred model (Supplemental Materials). Each locus was evaluated for the number of parsimony-informative sites, number of constant sites, and number of variable sites using **PAUP* v.40b10** (Swofford 2003)(Supplemental Materials). A maximum likelihood phylogenetic analysis was conducted on each locus using **RAxML v7.2.8** (Stamatakis 2006) with 1000 rapid-bootstrap replicates with the **GTRGAMMA** model.

All loci were concatenated into a single alignment for maximum-likelihood (**RAxML**) and Bayesian analysis (**ExaBayes**) (Aberer et al. 2014). The concatenated data set was partitioned by locus (429 partitions) for both analyses. The ML analysis was conducted with the **GTRGAMMA** model with 1000 rapid-bootstrap replicates. **ExaBayes** analyses were run with the **GTRGAMMA** model with branch lengths linked across partitions and a parsimony starting tree with heated chains using different starting trees than the cold chain. Four independent runs were conducted, each with four chains, sampling every 500 generations. **ExaBayes** runs continued until the termination condition of mean topological difference less than 5% with at least 500,000 generations was met. Posterior distributions

of trees were summarized with the `consense` script and posterior sample of parameters were assessed with `Tracer v1.5` (Rambaut and Drummond 2007) and combined with the `postProcParam` script.

Species tree estimation

Due to the large number of genes sampled we limit our species tree estimation to a summary statistic approach. Species tree accuracy in summary statistic approaches is dependent on gene tree accuracy (Huang and Knowles 2009; Mirarab et al. 2014) since the methods rely solely on the structure of the fully resolved gene trees. Loci with few informative sites, often seen in NGS datasets, may not give strong support for all splits in the gene trees. This can potentially bias the species tree estimate and analyses relying on the species tree topology, such as the identification of the anomaly zone. We use the maximum pseudo-likelihood estimation of species trees `MP-EST v1.4` (Liu et al. 2010a) because it can accurately estimate the species tree topology despite the anomaly zone and in the presence of gene tree estimation error (Liu et al. 2010a). We accounted for gene tree and species tree uncertainty by running `MP-EST` using each iteration of the 1000 maximum-likelihood bootstrap replicates from `RAxML`. We created 1000 new tree files consisting of 429 trees, one bootstrap replicate of each locus. `MP-EST` was run on each of these files and an extended majority-rule consensus (eMRC) tree of the resulting species trees was calculated using `sumtrees` in `Dendropy v4.0.3` (Sukumaran and Holder 2010). The support values from this approach are a summary of bootstrap replicates from the gene trees to account for error in gene tree estimates but should not be considered a true bootstrap replicate of the species tree.

Identifying anomalous nodes

We use the unifying principle of the anomaly zone (Rosenberg 2013) to determine which, if any, parts of the Scincidae eMRC species tree topology should produce AGTs. This procedure requires an estimate of internal branch lengths in coalescent units. We estimated branch lengths using two different methods. First, we used the branch lengths estimated by MP-EST, which jointly estimates internal branch lengths in coalescent units (based on $\lambda = 2\tau/\theta$) while maximizing the pseudo-likelihood of the species topology given the set of triplet topologies for each gene tree (Liu et al. 2010a, Equations: 6–8). Internal branches estimated by MP-EST may be shorter than expected when gene tree error is high (Mirarab et al. 2014), which may give an overestimation of internode pairs in the anomaly zone. Second, we estimated branch lengths using BP&P v2.1b (Yang and Rannala 2010) with the original sequence data for 429 loci and the eMRC topology from MP-EST. BP&P uses a fixed species tree topology and the multispecies coalescent along with gene trees estimated using the Jukes-Cantor model to estimate branch lengths (τ) and population sizes (θ) (Rannala and Yang 2003; Burgess and Yang 2008). A gamma prior on θ ($\alpha = 2.0$, $\beta = 200$) with a mean of 0.01 was used for population size estimates on nodes. A gamma prior on τ ($\alpha = 4.0$, $\beta = 10.0$) was used for the root node height with other times generated from the Dirichlet distribution (Yang and Rannala 2010). Rate variation between loci was accommodated with the random-rates model (Burgess and Yang 2008), in which the average rate for all loci is fixed at 1 and the rates among loci are generated from a Dirichlet distribution. We used an α of 2.0 for moderate variation among loci. The MCMC chain was run for 100,000 samples, sampling every 10 generations for a total of 1,000,000 sampled states with a burnin of 150,000 states. Three independent analyses were conducted to verify convergence on a stable posterior. BP&P results were converted to coalescent units ($\lambda = 2\tau/\theta$) consistent with those calculated by MP-EST.

Each pair of parent–child internodes were compared to the anomaly zone based on values of λ calculated from BP&P and MP-EST. The value for the parent nodes (x -nodes)

were put into equation (1) for the limit of the anomaly zone in a four-taxon asymmetric tree to determine if they are inside the zone and would therefore produce anomalous gene trees. If the value of the child (y -node) is less than $a(x)$, the pair of internodes are in the anomaly zone and AGTs are expected. This calculation was first conducted on the median values of branch lengths from the eMRC tree of the MP-EST species tree replicates and the median values of τ and θ from the eMRC tree of the BP&P posterior distribution.

Additionally, for the MP-EST bootstrap replicates, anomaly zone calculations were done for each internode pair for each species tree bootstrap replicate, accounting for topological error in estimates of branch lengths. For BP&P, 1000 random draws of joint values of τ and θ for the internode pairs were made from the posterior distribution and compared to the anomaly zone. The BP&P analysis is only performed on the eMRC topology. Scripts (https://github.com/cwlinkem/anomaly_zone) to perform these functions relied on the Dendropy package (Sukumaran and Holder 2010). We report the proportion of bootstrap replicates that match the eMRC tree and are in the anomaly zone with MP-EST and the proportion of the 1000 draws from BP&P posterior distribution that are in the anomaly zone.

RESULTS

Genomic data and assemblies

All samples were successfully sequenced at sufficient levels to result in high coverage of target loci. Some samples represented a larger portion of the sequencing (Table 1) potentially due to unequal pooling prior to hybridization. Raw reads averaged 9.8 million (range 5.4–21.6) reads per species with most reads being of high quality, resulting in a low rate of trimming and removal. Sequencing resulted in higher coverage than needed for the sequence capture approach due to using 24 samples on the sequencing lane instead of the

full potential of 96 samples. The added sequence coverage resulted in a higher proportion of off-target sequencing than would be expected, including complete mitochondrial genomes for many taxa. Results from other experiments have shown that off-target sequencing is reduced when more samples are included in the sequencing lane without a loss of target sequence coverage (unpublished data).

The number of contigs found for IDBA for each individual is large, averaging over 49,000 (range 18,850–155,978) contigs across the 16 species. The IDBA assemblies match most of the 585 loci targeted. Datasets were assembled for complete taxon sampling for all loci. IDBA assemblies resulted in 429 loci across all taxa. Of the 44 loci used in previous squamate systematics studies, only two loci were present in IDBA assemblies for all species.

Loci informativeness and model choice

The 429 loci from the IDBA assembly totals 276,480 nucleotide positions with 5.28 % missing data and an average length of 644 base pairs (range 338–1070). Individual loci vary in character variability with an average of 6% parsimony-informative sites (range 0–15%) and 18% variable sites (range 1–48%). Most loci have a best-fit model matching HKY or K80 (400 out of 429) suggesting a prevalence of transition/transversion bias in these genomic loci (Table 2). Model testing shows a preference for either a gamma or invariant-sites model (376 out 429) but the combination is rarely preferred (38 out of 429). Only 18 loci have a preferred model that does not accommodate among-site rate heterogeneity (Table 2).

Gene trees

Maximum Likelihood searches for individual loci resulted in 429 unique topologies, one for each locus. These topologies also differ from the concatenation ML tree and the

species tree. Bootstrap replicates average 925 (range 163–1000) unique topologies out of the 1000 replicates indicating that gene tree resolution is low for most individual loci.

Concatenated gene trees are largely congruent between the ML and Bayesian runs (Figure 2). Acontinae is sister to all other taxa with strong support (100 bootstrap and 100 posterior probability). Lygosominae is monophyletic and the *Sphenomorphus* group is sister to the clade of the *Mabuya*, *Lygosoma*, and *Eugongylus* groups. The *Eugongylus* group is sister to the *Lygosoma* group. All of these relationships have strong support in both analyses. There is also strong support for *Brachymeles* to be sister to Lygosominae, making Scincinae paraphyletic. The only topological difference between the two analyses is the placement of *Ophiomorus*. In the ML analysis *Ophiomorus* is sister to the other genera in Scincinae (minus *Brachymeles*) with a bootstrap score of 51, whereas in the Bayesian analysis *Ophiomorus* is sister to all Scincinae and Lygosominae with a posterior probability of 71. In both cases the support for the placement of *Ophiomorus* is low despite the large amount of data used in these analyses.

Species trees

There were 110 unique species tree topologies found across the replicate species trees. The MP-EST eMRC species tree is 100% consistent for many relationships (Figure 2c), but some key relationships vary among species tree replicates and differ from the concatenation tree. Most significantly, the placement of *Brachymeles* in the species trees recovers a monophyletic Scincinae, but with moderate (77% of replicates) frequency. The alternative placement of *Brachymeles* sister to Lygosominae (as seen in the concatenation analyses) has a frequency of 21%. The sister relationships of *Scincus* and *Mesoscincus* has the lowest frequency (48% of replicates). *Ophiomorus* is sister to *Plestiodon* with low frequency (53% of replicated). The Lygosominae portion of the species tree is identical to

the concatenation topologies. The concatenation topology from RAxML, which had >99% support for all but one node, only matches 3% of the species tree replicates.

Nodes in the anomaly zone

MP-EST.—Median internode lengths calculated with **sumtrees** (Sukumaran and Holder 2010) were used to calculate the anomaly zone for each pair of internodes in the eMRC (equation (1)). A large region of the phylogeny is in the anomaly zone based on these median values (Figure 3a). The majority of relationships in Scincinae have internode lengths that are expected to produce AGTs. Examining the anomaly zone across the species tree replicates (Figure 3b) shows that most pairs of parent–child internodes found to be in the anomaly zone with median branch lengths remain in the anomaly zone across replicates. One region of the phylogeny leading to Lygosominae (yellow branches) is only in the anomaly zone for 6% of the replicates that match the eMRC topology. The species tree replicates include relationships not found in the eMRC that represent the alternate resolutions of the poorly resolved nodes. Many of these alternative species tree relationships in the replicate set of trees are also in the anomaly zone (results not shown).

BP&P.—The posterior distribution from BP&P was summarized using **sumtrees** to obtain the median branch length τ and population size θ for internodes. BP&P calculates τ and θ individually, which were used to calculate λ ($\lambda = 2\tau/\theta$) (Table 3). Internodes B, C, and N (Figure 2c) are particularly short and internode G is the longest. Calculations of $a(x)$ (equation (1)) show that six internodes (B, C, J, K, L, N) have $a(x)$ values above zero indicating they may produce AGTs depending on the length of the descendant internode. Of these, B, C, J, and L have descendant internodes. The internode pairs B/C, B/I, and L/N are in the anomaly zone because the lengths of internodes C and I are shorter than $a(x)$ for B and internode N is shorter than $a(x)$ for L (Figure 3c). To account for the range

in branch length estimates across the posterior distribution, 1000 random draws of τ and θ were made. Each draw was calculated for occurrence of the anomaly zone for each pair of internodes. Internodes inferred to be in the anomaly zone using median branch lengths were all inferred at high frequency (Figure 3d). Additional pairs of internodes are in the anomaly zone when taking into account the range of τ and θ values in the posterior distribution, showing a similar pattern of anomaly zone nodes as found with estimates from MP-EST.

DISCUSSION

Species trees estimated using coalescent methods often differ from concatenated gene trees, and the anomaly zone may be contributing to these disagreements. The anomaly zone could also explain why some phylogenomic studies find low support for relationships despite the inclusion of hundreds to thousands of loci. In the empirical example of skinks presented here, we find strong conflict between species trees and concatenated gene trees, as well as conflict between individual gene trees. This type of conflict is typical for phylogenomic studies of rapid diversification events (Zou et al. 2008; McCormack et al. 2012; Jarvis et al. 2014). Our examination of the anomaly zone in skinks shows that the parts of the tree in conflict correspond with areas of the tree that are also estimated to be in the anomaly zone. The anomaly zone is a potential explanation for why conflicting relationships persist in some phylogenomic studies.

The anomaly zone in empirical phylogenies

We find that the anomaly zone likely occurs in empirical studies and that it may be more pervasive than previously assumed (but see Huang and Knowles (2009)). Although there might be insufficient variation to resolve gene trees that are derived from a species tree in

the anomaly zone, most studies are not specifically interested in whether or not we can estimate individual gene trees. The species tree is the target of analysis, not the individual gene trees (Edwards 2009). While it is true that estimation of any particular gene tree is hindered by low genetic variation, this does not change whether or not speciation events occurred quickly enough to place them into the anomaly zone. Phylogenetic analysis of concatenated loci, as a proxy for the species tree, can result in strong support as more loci are combined, even if there is low genetic variation in individual loci. If the speciation history is in the anomaly zone, then the resulting phylogeny will be erroneous (Kubatko and Degnan 2007). Our study shows that there is low support for relationships in most individual gene trees, but after concatenating the 276,480 characters they provide strong support for most relationships (Figure 2a–b). Huang and Knowles (2009) showed that when the species history is in the anomaly zone, gene tree estimation error will be high and that the lack of variation will make estimation of individual gene trees difficult. The observation of significant gene tree discordance across hundreds of loci may be a good sign that the species history is in the anomaly zone, and the framework that we provide here offers one way to test this hypothesis.

Summary statistic methods for species tree inference have become a necessity, since the more statistically rigorous full-Bayesian approaches (*BEAST and BEST) cannot handle hundreds of loci (Bayzid and Warnow 2013). Summary methods use gene trees to estimate species trees, and as previously discussed, gene tree estimation error is typically high in cases where an anomaly zone is suspected. Gene tree estimation error reduces the accuracy of species tree estimation (Bayzid and Warnow 2013; Mirarab et al. 2014), which makes species tree inference in the anomaly zone more difficult (Liu and Edwards 2009). We accounted for gene tree error by repeating the species tree estimation procedure using the gene trees constructed from the bootstrap replicates, which can provide a measure of accuracy that is not available when only using the ML gene trees (Mirarab et al. 2014).

Despite this, it is still possible that gene tree estimation error is biasing the species tree. Ideally, these inaccuracies should be reflected by the low frequency of occurrence for inaccurate nodes in the species tree. By estimating the anomaly zone across all topologies in the replicate species trees from MP-EST we find that the anomaly zone can be inferred even when species trees contain low supported nodes. Species trees in the anomaly zone are likely to have nodes with low frequency due to the high frequency of AGTs and the occurrence of gene tree estimation error. In these situations, including more loci may not increase node frequency due to the addition of more gene tree estimation error. Increasing the accuracy of gene trees through sampling longer loci (McCormack et al. 2009), or by combining loci with a shared history into larger gene fragments (Bayzid and Warnow 2013; Betancur-R et al. 2013), may improve inference of species trees in the anomaly zone. Inferring relationships in recently diverged species may also benefit from increased sampling of individuals within each species (Rokas and Carroll 2005; Maddison and Knowles 2006), though this benefit would be lost in deep divergences (Kubatko and Degnan 2007).

Frequency of anomalous gene trees

While we cannot be certain that the skink species tree that we estimated is correct, it is clear that the set of species tree topologies estimated from these phylogenomic data show signs of the presence of the anomaly zone over multiple pairs of internodes (Figure 3). The extent and frequency of the anomaly zone in this empirical example indicates that many AGTs may exist across the genomes of these taxa. It is important to keep in mind that each inference of the anomaly zone is limited to pairs of internal branches, without consideration of neighboring relationships. Only considering neighboring internal branches is a simplification; accounting for other branches would increase the size of the anomaly zone (Rosenberg and Tao 2008). Focusing on pairs of internal branches makes our investigation of the anomaly zone a conservative approximation of the extent of AGTs.

When considered together, the extent of the anomaly zone in our empirical example has the potential to produce many AGTs.

Rosenberg and Tao (2008) showed that the number of AGTs increases rapidly as the number of short internodes increases and that an anomaly zone that includes three internodes (five-taxon trees or larger) can produce as many as 45 AGTs. Estimates for trees larger than five taxa or an anomaly zone spanning more than three nodes have not been estimated, but are expected to increase exponentially. In the example of skinks, the 429 loci sampled are all estimated to have unique ML topologies. Our sample of loci does not show a trend for a particular topology to be more frequent, but this is not unexpected given the large number of possible topologies (4.22×10^{13}) for a 16-taxon rooted tree. Across the 429 alternate topologies sampled, an extended majority-rule consensus of these topologies shows that most splits occur in low frequency. Splits associated with internode pairs estimated to be in the anomaly zone occur at lower frequency among the gene trees than splits not estimated in the anomaly zone. It is unclear how species tree methods perform when the anomaly zone is larger than a four-taxon case as we see in skinks.

Simulation studies testing the ability of different species tree methods to overcome the four-taxon anomaly zone (Kubatko and Degnan 2007; Huang and Knowles 2009; Liu and Edwards 2009) have shown that species tree methods can accurately estimate the true species tree when provided with enough data. For concatenated gene trees, we know that combining loci with conflicting topologies can result in strong support for a single topology (Rokas et al. 2003). In some cases, relationships within the concatenated gene tree may not occur within the individual gene trees, a result referred to as hidden likelihood support (Gatesy and Baker 2005). This hidden support has been argued to be an advantage of concatenation methods over species tree methods (Springer and Gatesy 2014; Gatesy and Springer 2014). In cases where the speciation history fits the demographic scenario of the anomaly zone, the hidden likelihood support in the concatenated gene tree will be for an

incorrect topology, resulting in strong inference of an incorrect species history (Kubatko and Degnan 2007). The method for inferring the anomaly zone presented here allows researchers to identify potential conflicts and decide on appropriate analytical solutions.

Edwards (2009) predicted that phylogenomic studies would find lower support for relationships using species tree methods than would be obtained from concatenation, especially in older clades. This prediction is based on the idea that missing data would have a larger effect on species trees, and that species trees use more complex models compared to concatenation. We find these predictions to be accurate, though their cause may be different than originally proposed. We propose that low support in species tree analyses is due to a combination of gene tree estimation error and the presence of AGTs that bias phylogenetic signal. Simulation studies have shown that many species tree methods can overcome the anomaly zone in simple four-taxon scenarios (Liu and Edwards 2009), but no study has looked at the effect of larger anomaly zone problems on trees with more taxa. We predict that when the anomaly zone occurs across more than two internodes, the greater number of AGTs will provide support for multiple species trees, reducing the support for some species tree nodes. This will likely occur even when gene trees are estimated with certainty (i.e., using simulated data), a luxury not available with empirical studies. The increased number of AGTs may also result in low support in concatenation analyses even when analyzing hundreds of genes if the AGTs are in direct topological conflict. In this study, we find two nodes with **RAxML** and a different node in **EXaBayes** that have low support despite having over 276,000 (50,666 variable; 15,524 parsimony-informative) characters in the analyses. With multiple AGTs there may be alternative topologies with high probability in the set of candidate genes. These alternative topologies should lower branch support in concatenation analyses. This low support for nodes near short branches may be an indication of an anomaly zone problem in phylogenomic datasets.

Higher-level skink relationships

Resolving the relationships within Scincidae is an ongoing challenge (Brandley et al. 2012; Wiens et al. 2012; Pyron et al. 2013; Lambert et al. 2015) which our current study addresses with a slightly different approach compared to previous studies. Pyron et al. (2013) used hundreds of taxa in a sparse matrix, while Wiens et al. (2012); Lambert et al. (2015) used many loci with few taxa. Our taxon sampling is most similar to Brandley et al. (2012) but utilizes 429 loci and species tree analyses to estimate relationships. Our preferred estimate of species relationships (Figure 2c) shows subfamily relationships concordant with results in Pyron et al. (2013) and Lambert et al. (2015). Acontinae is sister to all other skinks and the subfamilies Scincinae and Lygosominae are monophyletic. The monophyly of Scincinae conflicts with most studies and our own results from analyses of concatenated data (Whiting et al. 2003; Brandley et al. 2005, 2012; Wiens et al. 2012) but is likely the more accurate relationship based on our inference of the anomaly zone in relation to the nodes preceding and in this subfamily. Similar to Lambert et al. (2015) we find *Brachymeles* to be related to other Scincinae as opposed to sister to all Lygosominae (Brandley et al. 2012). The relationships among Scincinae genera differs from those presented in Brandley et al. (2012) though we have far fewer genera sampled and low support so many comparisons should be made with caution. We consider the alternative placement of *Brachymeles* sister to Lygosominae, seen in concatenated gene trees, to be an artifact of rapid diversification resulting in a series of AGTs. A detailed examination of the relationships among the genera in Scincinae with broad taxon sampling is clearly warranted. Our inference of the anomaly zone among some Scincinae genera suggests that hundreds of loci and species tree analyses will be necessary to accurately estimate the phylogenetic relationships within this group.

Relationships within Lygosominae are largely concordant with previous studies in finding the *Sphenomorphus* group sister to all other groups (Honda et al. 2003; Reeder

2003; Skinner et al. 2011). We find the *Lygosoma* group to be sister to the *Eugongylus* group and that this pair is sister to the *Mabuya* group consistent with the results of Reeder (2003) and Skinner et al. (2011). Relationships among the sampled genera in the *Sphenomorphus* group are consistent with previous results (Linkem et al. 2011). Over three-quarters of all skink species are in Lygosominae and it appears based on our limited sampling that the broad groupings of genera can be consistently resolved and the anomaly zone is not an issue at this level. Within the *Sphenomorphus* group, previous studies have reported short branches separating major groups (Linkem et al. 2011; Skinner et al. 2011). We suspect that there will be anomaly zone issues within the *Sphenomorphus* radiation, especially in Australia. The *Eugongylus* group will likely also present anomaly zone issues given the large number of species in the group and relatively recent origin. Studies at a more fine scale will likely find similar rapid diversification patterns that are consistent with the anomaly zone. Further work on these diverse groups is needed to better understand their systematic relationships.

A monophyletic Scincinae necessitates an investigation into morphological features to define this group. Greer (1970) considered Scincinae to be the primitive morphological form from which all other skinks were derived, but molecular studies find Scincinae taxa to be nested and sister to Lygosominae, conflicting with the prevailing morphological interpretation. A detailed and rigorous analysis of the morphological evolution and biogeography of skinks will require much more rigorous taxon sampling.

Overcoming the anomaly zone

Sequencing hundreds of unlinked single copy loci provides an opportunity to explore the conflicts between loci and analytical approaches, as well as address what may be the source of conflict. Our work shows that researchers conducting empirical studies should closely consider the potential impact the anomaly zone has on their phylogenetic analyses. A

common trend with phylogenomic studies is to analyze the data with concatenation, an approach that has the advantage of faster computation times and simplicity, but that provides overwhelmingly and likely erroneous strong support across most of the tree. Species tree analyses often result in lower support for difficult parts of the tree than concatenation. Instead of marginalizing the species tree results, we should acknowledge that they are a likely consequence of the speciation history for the group. The lower support provided by coalescent-based species tree inference is potentially a more accurate reflection of the support for the tree given the data.

As we show here, the anomaly zone is likely more pervasive than previously suggested and should be accounted for when studying taxa that may have diverged rapidly, even if that rapid event was in the distant past. Combining hundreds to thousands of independent loci together with coalescent-based species tree inference is the most effective way of getting an accurate result. Targeting longer loci will help reduce gene tree estimation error, resulting in a better estimate of the species tree. In the most extreme cases, resolving the nodes of a species tree with strong support may not be possible even when sampling the entire genome.

FUNDING

C.W.L. was supported by a National Science Foundation Postdoctoral Research Fellowship in Biology (Award 1202754). Data collection was funded by National Science Foundation grant DBI-1144630 awarded to A.D.L. This work used the Vincent J. Coates Genomics Sequencing Laboratory at UC Berkeley, supported by National Institutes of Health S10 Instrumentation Grants S10RR029668 and S10RR027303.

ACKNOWLEDGEMENTS

We would like to thank R. Brown of the University of Kansas, C. Spencer and J. McGuire of UC Berkeley, A. Resetar of the Field Museum, and K. Imada and A. Allison of the Bishop Museum for loans of tissues necessary for this work. B. Faircloth provided detailed help in troubleshooting early attempts at sequence capture, and we thank him for quickly responding to our queries. R. Harris helped with designing this new probe set. R. Bryson provided useful feedback and discussion. Members of the Leache lab contributed to early discussions of this work.

*

References

- Aberer, A. J., K. Kobert, and A. Stamatakis. 2014. Exabayes: massively parallel bayesian tree inference for the whole-genome era. *Molecular Biology and Evolution* 31:2553–2556.
- Alföldi, J., F. Di Palma, M. Grabherr, C. Williams, L. Kong, E. Mauceli, P. Russell, C. B. Lowe, R. E. Glor, J. D. Jaffe, et al. 2011. The genome of the green anole lizard and a comparative analysis with birds and mammals. *Nature* 477:587–591.
- Bayzid, M. S. and T. Warnow. 2013. Naive binning improves phylogenomic analyses. *Bioinformatics* 29:2277–2284.
- Betancur-R, R., G. J. Naylor, and G. Ortí. 2013. Conserved genes, sampling error, and phylogenomic inference. *Systematic Biology* 63:257–262.
- Blackburn, D. G. 1982. Evolutionary origins of viviparity in the Reptilia. i. Sauria. *Amphibia-Reptilia* 3:185–205.
- Bolger, A. M., M. Lohse, and B. Usadel. 2014. Trimmomatic: A flexible trimmer for illumina sequence data. *Bioinformatics* 30:2114–2120.
- Brandley, M. C., H. OTA, T. Hikida, A. Nieto Montes De Oca, M. Fería-ortíz, X. Guo, and Y. Wang. 2012. The phylogenetic systematics of blue-tailed skinks (Plestiodon) and the family Scincidae. *Zoological Journal of the Linnean Society* 165:163–189.
- Brandley, M. C., A. Schmitz, and T. W. Reeder. 2005. Partitioned Bayesian analyses, partition choice, and the phylogenetic relationships of scincid lizards. *Systematic Biology* 54:373–390.

- Burgess, R. and Z. Yang. 2008. Estimation of hominoid ancestral population sizes under Bayesian coalescent models incorporating mutation rate variation and sequencing errors. *Molecular Biology and Evolution* 25:1979–1994.
- Darriba, D., G. L. Taboada, R. Doallo, and D. Posada. 2012. jModelTest 2: more models, new heuristics and parallel computing. *Nature Methods* 9:772–772.
- Degnan, J. H. 2013. Anomalous unrooted gene trees. *Systematic Biology* 62:574–590.
- Degnan, J. H. and N. A. Rosenberg. 2006. Discordance of species trees with their most likely gene trees. *PLoS Genetics* 2:762–768.
- Degnan, J. H. and N. A. Rosenberg. 2009. Gene tree discordance, phylogenetic inference and the multispecies coalescent. *Trends in Ecology & Evolution* 24:332–340.
- Degnan, J. H., N. A. Rosenberg, and T. Stadler. 2012a. A characterization of the set of species trees that produce anomalous ranked gene trees. *IEEE/ACM Transactions on Computational Biology and Bioinformatics (TCBB)* 9:1558–1568.
- Degnan, J. H., N. A. Rosenberg, and T. Stadler. 2012b. The probability distribution of ranked gene trees on a species tree. *Mathematical Biosciences* 235:45–55.
- Dunn, C. W., A. Hejnol, D. Q. Matus, K. Pang, W. E. Browne, S. A. Smith, E. Seaver, G. W. Rouse, M. Obst, G. D. Edgecombe, et al. 2008. Broad phylogenomic sampling improves resolution of the animal tree of life. *Nature* 452:745–749.
- Edwards, S. V. 2009. Is a new and general theory of molecular systematics emerging? *Evolution* 63:1–19.
- Faircloth, B. C. 2013. illumiprocessor: a trimmomatic wrapper for parallel adapter and quality trimming. <http://dx.doi.org/10.6079/j9ill> .

- Faircloth, B. C. 2014. phyluce: phylogenetic estimation from ultraconserved elements. doi:10.6079/j9phyl .
- Faircloth, B. C., J. E. McCormack, N. G. Crawford, M. G. Harvey, R. T. Brumfield, and T. C. Glenn. 2012. Ultraconserved elements anchor thousands of genetic markers spanning multiple evolutionary timescales. *Systematic Biology* 61:717–726.
- Gatesy, J. and R. H. Baker. 2005. Hidden likelihood support in genomic data: can forty-five wrongs make a right? *Systematic Biology* 54:483–492.
- Gatesy, J. and M. S. Springer. 2014. Phylogenetic analysis at deep timescales: Unreliable gene trees, bypassed hidden support, and the coalescence/concatalescence conundrum. *Molecular Phylogenetics and Evolution* 80:231–266.
- Genomic Resources Development Consortium, W. Arthofer, B. L. Banbury, M. Carneiro, F. Cicconardi, T. F. Duda, R. B. Harris, D. S. Kang, A. D. Leaché, V. Nolte, C. Nourisson, N. Palmieri, B. C. Schlick-Steiner, C. Schlatterer, F. Sequeira, C. Sim, F. M. Steiner, M. Vallinoto, and D. A. Weese. 2014. Genomic resources notes accepted 1 august 2014–30 september 2014. *Molecular Ecology Resources* 15:228–229.
- Greer, A. E. 1970. A subfamily classification of scincid lizards. *Bulletin of the Museum of Comparative Zoology* 139:151–184.
- Greer, A. E. 1986. Lygosomine (Scincidae) monophyly: a third, corroborating character and a reply to critics. *Journal of Herpetology* 20:123–126.
- Greer, A. E. 1991. Limb reduction in squamates: identification of the lineages and discussion of the trends. *Journal of Herpetology* 25:166–173.
- Guindon, S. and O. Gascuel. 2003. A simple, fast, and accurate algorithm to estimate large phylogenies by maximum likelihood. *Systematic Biology* 52:696–704.

- Honda, M., H. Ota, G. Köhler, I. Ineich, L. Chirio, S.-L. Chen, and T. Hikida. 2003. Phylogeny of the lizard subfamily Lygosominae (Reptilia: Scincidae), with special reference to the origin of the new world taxa. *Genes & Genetic Systems* 78:71–80.
- Huang, H. and L. L. Knowles. 2009. What is the danger of the anomaly zone for empirical phylogenetics? *Systematic Biology* 58:527–536.
- Jarvis, E. D., S. Mirarab, A. J. Aberer, B. Li, P. Houde, C. Li, S. Y. Ho, B. C. Faircloth, B. Nabholz, J. T. Howard, et al. 2014. Whole-genome analyses resolve early branches in the tree of life of modern birds. *Science* 346:1320–1331.
- Kubatko, L. S. and J. H. Degnan. 2007. Inconsistency of phylogenetic estimates from concatenated data under coalescence. *Systematic Biology* 56:17–24.
- Lamb, T., S. Biswas, and A. M. Bauer. 2010. A phylogenetic reassessment of African fossorial skinks in the subfamily Acontinae (Squamata: Scincidae): evidence for parallelism and polyphyly. *Zootaxa* 2657:33–46.
- Lambert, S. M., T. W. Reeder, and J. J. Wiens. 2015. When do species-tree and concatenated estimates disagree? An empirical analysis with higher-level scincid lizard phylogeny. *Molecular Phylogenetics and Evolution* 82:146–155.
- Linkem, C. W., A. C. Diesmos, and R. M. Brown. 2011. Molecular systematics of the Philippine forest skinks (Squamata: Scincidae: Sphenomorphus): testing morphological hypotheses of interspecific relationships. *Zoological Journal of the Linnean Society* 163:1217–1243.
- Liu, L. and S. V. Edwards. 2009. Phylogenetic analysis in the anomaly zone. *Systematic Biology* 58:452–460.

- Liu, L., S. Wu, and L. Yu. 2015. Coalescent methods for estimating species trees from phylogenomic data. *Journal of Systematics and Evolution* 53:380–390.
- Liu, L. and L. Yu. 2011. Estimating species trees from unrooted gene trees. *Systematic Biology* 60:661–667.
- Liu, L., L. Yu, and S. V. Edwards. 2010a. A maximum pseudo-likelihood approach for estimating species trees under the coalescent model. *BMC Evolutionary Biology* 10:302.
- Liu, L., L. Yu, and D. K. Pearl. 2010b. Maximum tree: a consistent estimator of the species tree. *Journal of Mathematical Biology* 60:95–106.
- Maddison, W. P. 1997. Gene trees in species trees. *Systematic Biology* 46:523–536.
- Maddison, W. P. and L. L. Knowles. 2006. Inferring phylogeny despite incomplete lineage sorting. *Systematic Biology* 55:21–30.
- McCormack, J. E., B. C. Faircloth, N. G. Crawford, P. A. Gowaty, R. T. Brumfield, and T. C. Glenn. 2012. Ultraconserved elements are novel phylogenomic markers that resolve placental mammal phylogeny when combined with species-tree analysis. *Genome Research* 22:746–754.
- McCormack, J. E., H. Huang, and L. L. Knowles. 2009. Maximum likelihood estimates of species trees: how accuracy of phylogenetic inference depends upon the divergence history and sampling design. *Systematic Biology* 58:501–508.
- Mirarab, S., M. S. Bayzid, and T. Warnow. 2014. Evaluating summary methods for multi-locus species tree estimation in the presence of incomplete lineage sorting. *Systematic Biology* doi:10.1093/sysbio/syu063.
- Mulcahy, D. G., B. P. Noonan, T. Moss, T. M. Townsend, T. W. Reeder, J. W. Sites, and J. J. Wiens. 2012. Estimating divergence dates and evaluating dating methods using

phylogenomic and mitochondrial data in squamate reptiles. *Molecular Phylogenetics and Evolution* 65:974–991.

Peng, Y., H. Leung, S. Yiu, and F. Chin. 2010. IDBA a practical iterative de bruijn graph de novo assembler. Pages 426–440 *in* *Research in Computational Molecular Biology* (B. Berger, ed.) vol. 6044 of *Lecture Notes in Computer Science*. Springer Berlin Heidelberg.

Philippe, H., H. Brinkmann, D. V. Lavrov, D. T. J. Littlewood, M. Manuel, G. Wörheide, and D. Baurain. 2011. Resolving difficult phylogenetic questions: why more sequences are not enough. *PLoS Biology* 9:e1000602.

Philippe, H., R. Derelle, P. Lopez, K. Pick, C. Borchellini, N. Boury-Esnault, J. Vacelet, E. Renard, E. Houliston, E. Quéinnec, et al. 2009. Phylogenomics revives traditional views on deep animal relationships. *Current Biology* 19:706–712.

Pyron, R. A., F. T. Burbrink, and J. J. Wiens. 2013. A phylogeny and revised classification of Squamata, including 4161 species of lizards and snakes. *BMC Evolutionary Biology* 13:93.

Pyron, R. A., C. R. Hendry, V. M. Chou, E. M. Lemmon, A. R. Lemmon, and F. T. Burbrink. 2014. Effectiveness of phylogenomic data and coalescent species-tree methods for resolving difficult nodes in the phylogeny of advanced snakes (Serpentes: Caenophidia). *Molecular Phylogenetics and Evolution* 81:221–231.

Rambaut, A. and A. Drummond. 2007. Tracer v1.5, available from: <http://beast.bio.ed.ac.uk/tracer> .

Rannala, B. and Z. Yang. 2003. Bayes estimation of species divergence times and ancestral population sizes using DNA sequences from multiple loci. *Genetics* 164:1645–1656.

- Reeder, T. W. 2003. A phylogeny of the Australian *Sphenomorphus* group (Scincidae: Squamata) and the phylogenetic placement of the crocodile skinks (*Tribolonotus*): Bayesian approaches to assessing congruence and obtaining confidence in maximum likelihood inferred relationships. *Molecular Phylogenetics and Evolution* 27:384–397.
- Rokas, A. and S. B. Carroll. 2005. More genes or more taxa? The relative contribution of gene number and taxon number to phylogenetic accuracy. *Molecular Biology and Evolution* 22:1337–1344.
- Rokas, A., B. L. Williams, N. King, and S. B. Carroll. 2003. Genome-scale approaches to resolving incongruence in molecular phylogenies. *Nature* 425:798–804.
- Rosenberg, N. A. 2013. Discordance of species trees with their most likely gene trees: A unifying principle. *Molecular Biology and Evolution* 30:2709–2713.
- Rosenberg, N. A. and J. H. Degnan. 2010. Coalescent histories for discordant gene trees and species trees. *Theoretical Population Biology* 77:145–151.
- Rosenberg, N. A. and R. Tao. 2008. Discordance of species trees with their most likely gene trees: the case of five taxa. *Systematic Biology* 57:131–140.
- Siler, C. D., A. C. Diesmos, A. C. Alcala, and R. M. Brown. 2011. Phylogeny of Philippine slender skinks (Scincidae: *Brachymeles*) reveals underestimated species diversity, complex biogeographical relationships, and cryptic patterns of lineage diversification. *Molecular Phylogenetics and Evolution* 59:53–65.
- Skinner, A. 2007. Phylogenetic relationships and rate of early diversification of Australian *Sphenomorphus* group scincids (Scincoidea, Squamata). *Biological Journal of the Linnean Society* 92:347–366.

- Skinner, A., A. F. Hugall, and M. N. Hutchinson. 2011. Lygosomine phylogeny and the origins of Australian scincid lizards. *Journal of Biogeography* 38:1044–1058.
- Skinner, A., M. S. Lee, and M. N. Hutchinson. 2008. Rapid and repeated limb loss in a clade of scincid lizards. *BMC Evolutionary Biology* 8:310.
- Song, S., L. Liu, S. V. Edwards, and S. Wu. 2012. Resolving conflict in eutherian mammal phylogeny using phylogenomics and the multispecies coalescent model. *Proceedings of the National Academy of Sciences* 109:14942–14947.
- Springer, M. S. and J. Gatesy. 2014. Land plant origins and coalescence confusion. *Trends in Plant Science* 19:267–269.
- Stamatakis, A. 2006. RAxML-VI-HPC: maximum likelihood-based phylogenetic analyses with thousands of taxa and mixed models. *Bioinformatics* 22:2688–2690.
- Steel, M. and A. Rodrigo. 2008. Maximum likelihood supertrees. *Systematic Biology* 57:243–250.
- Suh, A., L. Smeds, and H. Ellegren. 2015. The dynamics of incomplete lineage sorting across the ancient adaptive radiation of neoavian birds. *PLoS Biology* 13:e1002224.
- Sukumaran, J. and M. T. Holder. 2010. Dendropy: a Python library for phylogenetic computing. *Bioinformatics* 26:1569–1571.
- Swofford, D. 2003. PAUP*. Phylogenetic Analysis Using Parsimony (*and other methods). Version 4. Sinauer Associates, Sunderland, Massachusetts.
- Tonini, J., A. Moore, D. Stern, M. Shcheglovitova, and G. Ortí. 2015. Concatenation and species tree methods exhibit statistically indistinguishable accuracy under a range of simulated conditions. *PLoS Currents* 7.

- Uetz, P. 2015. The reptile database. [Http://reptile-database.reptarium.cz](http://reptile-database.reptarium.cz).
- Whiting, A. S., A. M. Bauer, and J. W. Sites Jr. 2003. Phylogenetic relationships and limb loss in sub-Saharan African scincine lizards (Squamata: Scincidae). *Molecular Phylogenetics and Evolution* 29:582–598.
- Wiens, J. J., C. R. Hutter, D. G. Mulcahy, B. P. Noonan, T. M. Townsend, J. W. Sites, and T. W. Reeder. 2012. Resolving the phylogeny of lizards and snakes (Squamata) with extensive sampling of genes and species. *Biology Letters* 8:1043–1046.
- Yang, Z. and B. Rannala. 2010. Bayesian species delimitation using multilocus sequence data. *Proceedings of the National Academy of Sciences* 107:9264–9269.
- Zhang, G., C. Li, Q. Li, B. Li, D. M. Larkin, C. Lee, J. F. Storz, A. Antunes, M. J. Greenwold, R. W. Meredith, et al. 2014. Comparative genomics reveals insights into avian genome evolution and adaptation. *Science* 346:1311–1320.
- Zou, X.-H., F.-M. Zhang, J.-G. Zhang, L.-L. Zang, L. Tang, J. Wang, T. Sang, and S. Ge. 2008. Analysis of 142 genes resolves the rapid diversification of the rice genus. *Genome Biology* 9:R49.

Table 1: Genomic data collected and assembly results for 15 species of skinks and the outgroup.

Species	Voucher	Total reads (Million)	Clean reads (Million)	IDBA ^a	
				Contigs	Loci
<i>Brachymeles bonitae</i>	AJB077 ^c	8.71	7.71	29,942	550
<i>Chalcides ocellatus</i>	MVZ242790 ^b	6.59	5.93	26,745	569
<i>Emoia caeruleocauda</i>	KU307155 ^c	6.08	5.64	21,414	563
<i>Eurylepis taeniolatus</i>	MVZ246017 ^b	7.42	6.72	30,163	572
<i>Lobulia elegans</i>	BPBM18690 ^d	10.2	9.44	49,859	556
<i>Lygosoma brevicaudis</i>	MVZ249721 ^b	8.40	7.70	47,774	553
<i>Mabuya unimarginata</i>	CWL615 ^e	19.5	17.9	134,666	533
<i>Mesoscincus manguae</i>	CWL614 ^e	11.4	10.5	38,828	550
<i>Ophiomorus raithmai</i>	MVZ248453 ^b	7.78	7.00	35,506	568
<i>Plestiodon fasciatus</i>	KU289464 ^c	12.6	11.7	67,908	542
<i>Scincus scincus</i>	MVZ234538 ^b	6.54	5.94	20,231	566
<i>Sphenomorphus tridigitus</i>	FMNH258830 ^f	8.39	7.82	39,350	560
<i>Sphenomorphus variegatus</i>	KU315087 ^c	8.99	8.31	43,927	550
<i>Typhlosaurus sp</i>	MVZ164850 ^b	8.35	7.60	31,261	549
<i>Tytthoscincus parvus</i>	JAM6275 ^b	5.41	5.00	18,850	568
<i>Xantusia vigilis</i>	KU220092 ^c	21.5	19.9	155,978	550

^a Iterative de Bruijn graph short-read assembler.

^b Museum of Vertebrate Zoology, UC Berkeley, CA.

^c University of Kansas, Lawrence KS.

^d Bernice Pauahi Bishop Museum, Honolulu, HI.

^e No voucher specimen. Tissue deposited at KU.

^f Field Museum of Natural History, Chicago IL.

Table 2: Summary of model testing results ranked by model complexity. The model chosen for each locus can be found in the supplemental information.

Model	# free parameters	# of Loci	Avg % informative	Avg % variable
K80	31	1	0.8	4.5
K80+I	32	13	5.1	14.5
K80+G	32	22	6.9	21.3
K80+I+G	33	9	8.0	22.3
F81	33	3	0.2	1.6
F81+G	34	3	2.3	8.6
HKY	34	11	1.2	7.8
HKY+I	35	111	3.9	13.2
HKY+G	35	209	6.3	21.1
HKY+I+G	36	26	6.5	20.0
SYM+I	36	1	6.6	19.1
SYM+G	36	3	8.6	23.7
GTR+G	39	13	7.5	22.0
GTR+I	39	1	6.1	16.5
GTR+I+G	40	3	8.4	23.6

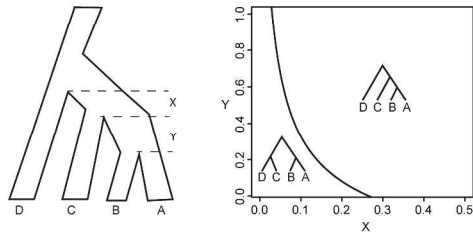
Table 3: Median branch lengths (τ) and population sizes (θ) used to calculate the branch length in coalescent units (λ) for the internodes labelled on the MP-EST species tree (Figure 2c). $a(x)$ indicates the limit of the anomaly zone for that internode length. If the length of the descendant internode is smaller than $a(x)$, then the internode pair is in the anomaly zone. If $a(x)$ is < 0 the internode pair will not produce anomalous trees. Descendant internodes in bold indicate anomaly zone pair.

Internode	τ	θ	λ	$a(x)$	Descendant internode(s)
A	0.006391	0.016352	0.781678	< 0	B
B	0.000036	0.012459	0.005778	1.020	C, I
C	0.000117	0.008112	0.028846	0.443	D, E
D	0.001574	0.00686	0.45889	< 0	G
E	0.002552	0.007271	0.701966	< 0	F
F	0.001553	0.01005	0.309054	< 0	—
G	0.007702	0.007601	2.02657	< 0	H
H	0.001286	0.00518	0.49652	< 0	—
I	0.005576	0.014087	0.79165	< 0	J
J	0.001754	0.024723	0.14189	0.0832	L, K
K	0.000902	0.013989	0.128958	0.0982	—
L	0.000588	0.010911	0.10778	0.1284	M, N
M	0.001272	0.00959	0.26527	0	—
N	0.000154	0.012777	0.024105	0.4994	—

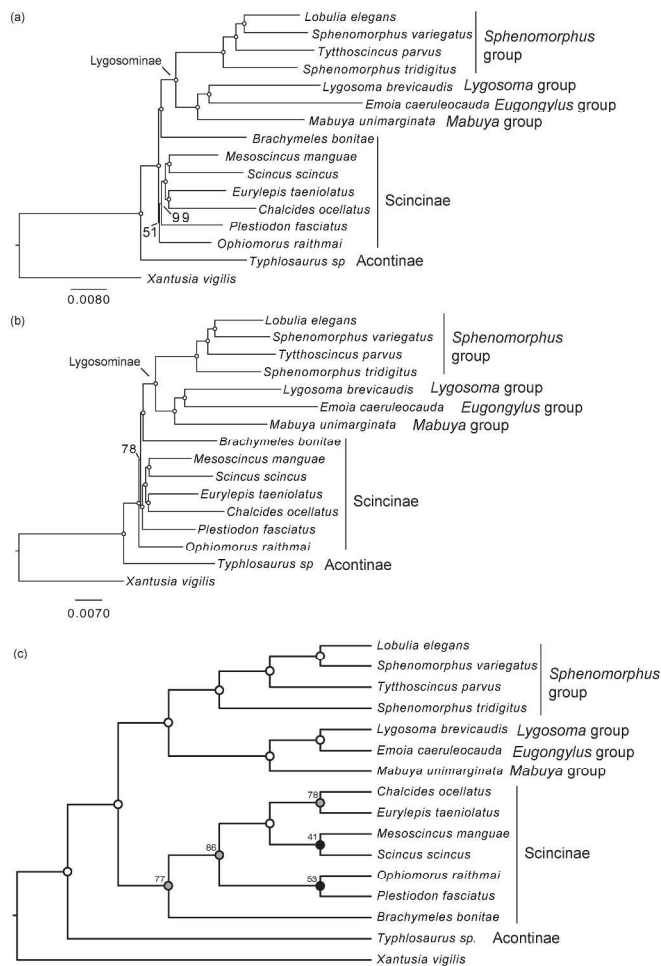
Figure 1: The length of branches X and Y in coalescent units in the species tree determine the probability of the gene tree topology. For branches under the anomaly zone curve the symmetric anomalous gene tree will have a higher probability than the asymmetric gene tree that matches the species tree.

Figure 2: Majority-rule consensus trees from the concatenated loci run in **RAxML** (a), **ExaBayes**, (b) and the species tree from **MP-EST** (c). Open circles indicate 100% bootstrap support or a Bayesian posterior probability of 1.0. Nodes with lower support are labelled. Both concatenation analyses have similar topologies with differences in the placement of *Ophiomorus* and the **RAxML** tree is slightly longer. The species tree (c) is shown as a cladogram. Relationships within Lygosominae are the same across analyses. The relationships within Scincinae differ both between the concatenation analyses and in comparison to the species tree. Letters on species tree nodes are used for Table 3 and discussion of internode pairs.

Figure 3: Majority-rule consensus topology shown as a cladogram. Pairs of internodes that are under the anomaly zone curve are highlighted in bold on each tree. Tree a) is based on median values of branch lengths calculated from MP-EST. Tree b) shows the frequency of the internodes in the anomaly zone across species tree replicates that match the eMRC topology. Tree c) shows the occurrence based on the median values of branch lengths from the posterior distribution of BP&P. Tree d) shows the frequency of the internode in the anomaly zone based on 1000 draws of joint internode values from the posterior distribution.



The length of branches X and Y in coalescent units in the species tree determine the probability of the gene tree topology. For branches under the anomaly zone curve the symmetric anomalous gene tree will have a higher probability than the asymmetric gene tree that matches the species tree.
 279x361mm (300 x 300 DPI)



Majority-rule consensus trees from the concatenated loci run in RAXML (a), ExaBayes, (b) and the species tree from MP-EST (c). Open circles indicate 100% bootstrap support or a Bayesian posterior probability of 1.0. Nodes with lower support are labelled. Both concatenation analyses have similar topologies with differences in the placement of *Ophiomorus* and the RAXML tree is slightly longer. The species tree (c) is shown as a cladogram. Relationships within the Lygosominae are the same across analyses. The relationships within Scincinae differ both between the concatenation analyses and in comparison to the species tree. Letters on species tree nodes are used for Table 3 and discussion of internode pairs.

279x361mm (300 x 300 DPI)

