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The origin(s) of modern amphibians: a commentary

By

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Anderson (2008) recently reviewed the controversial topic of extant amphibian origins, on which three (groups of) hypotheses exist at the moment. Anderson favors the “polyphyly hypothesis” (PH), which considers the extant amphibians to be polyphyletic with respect to many Paleozoic limbed vertebrates and was most recently supported by the analysis of Anderson *et al.* (2008). Another is the “temnospondyl hypothesis” (TH – lissamphibians nested within temnospondyls), most recently supported by Ruta and Coates (2007). We prefer the “lepospondyl hypothesis” (LH – lissamphibians nested within “lepospondyls”; most recently supported by Vallin and Laurin, 2004 and Marjanović and Laurin, 2008a). We would like to clarify important points that were not discussed in Anderson’s review, or for which crucial arguments were left out.

Anderson (2008) argues that most molecular dates favor the PH because they suggest a Devonian or Early Carboniferous diversification of Lissamphibia. This is inaccurate, since the confidence intervals of the dates obtained by Hugall *et al.* (2007) range from Early Carboniferous to Middle Permian, and our own molecular dating suggests a Permian origin. Indeed, three methods (molecular dating, a paleontological supertree and a confidence interval on the stratigraphic range of Lissamphibia) all hint at a Permian or (less likely) a Late Carboniferous origin of Lissamphibia (Marjanović and Laurin, 2007, 2008b).

Citing Schoch and Milner (2004), Anderson (2008: 234) argues that the LH is mainly supported by loss characters, and that this is problematic “given the relative ease that these losses can arise via pedomorphosis, which appears to evolve repeatedly.” This is especially surprising because we count (Supplementary Table 1) about fifty loss characters in the matrix by Anderson *et al.* (2008) – more than one out of five characters –, including several that describe the loss of bones that ossified late in the ontogeny of branchiosaurids (Schoch, 1992) and/or the aïstopod *Phlegethontia* (Anderson, 2002) and are absent in lissamphibians.

Furthermore, Anderson's remark amounts to criticizing the use of loss characters simply because they *could* be homoplastic. Yet, Anderson (2008) emphasizes development characters such as digit development and skull ossification order, which are *known* to be homoplastic. For instance, under Anderson's version of the PH, the similarity between the digit development orders of the branchiosaurid temnospondyl *Apateon* and the urodeles is either convergent, or homologous between these two taxa but reversed in anurans; indeed, Johanson *et al.* (2007) suggest that the digits of tetrapods are homologous to the "radials" of other sarcopterygians and find the "radials" of the Australian lungfish to develop independently of the rest of the forelimb (pectoral fin), like in urodeles and *Apateon* (and unlike in anurans and amniotes, where the limb chondrifies in a strict proximal-to-distal sequence), strongly suggesting that the urodele-*Apateon* pattern is plesiomorphic – regardless of whether the PH, the TH, or the LH is (closest to) correct. Anderson (2008: 242) furthermore mentions that "the pattern of cranial ossification [...] has compared very closely with the sequence of cranial ossification seen in salamanders" (making explicit on the next page that these similarities are plesiomorphic); more recently, however, Germain and Laurin (2009) have shown that the ancestral urodele sequence, even though poorly constrained, differed in several respects from that of *Apateon*.

Nonetheless, the suggestion that loss characters are likely to be strongly correlated to each other and homoplastic is serious; and indeed the matrix by Vallin and Laurin (2004) contains a greater proportion of loss characters (Supplementary Table 2) than that by Anderson *et al.* (2008). Unfortunately, no rigorous test of this possibility exists. The closest workable approximation we are aware of is the skewness test proposed by Huelsenbeck (1991), which measures how well a set of characters is compatible with different numbers of phylogenetic hypotheses: the smaller the g_1 statistic, the fewer trees are compatible with the character set, and the stronger is therefore the coherent phylogenetic signal in that character

set. The g_1 of the loss characters is much lower (and thus their phylogenetic signal higher) than that of the others in the matrix of Vallin and Laurin (2004), but higher in that of Anderson *et al.* (2008), and highly significant in all four partitions (Supplementary Table 3), showing that loss characters are not necessarily strongly correlated to each other, and that they appear to contain a phylogenetic signal.

Matrices of various sizes support the various hypotheses on lissamphibian origins. Anderson (2008: 234), citing Schoch and Milner (2004), criticizes the “limited number of taxa analyzed” by LH proponents. There is a trade-off between quantity and quality in a data matrix for a given time investment, and quality is as important as quantity for reaching accurate phylogenetic results. Even small matrices are sometimes not coded carefully enough. We recently showed that 35% of the cells of the first matrix that supported the PH, that by McGowan (2002), were problematic (Marjanović and Laurin, 2008a); with only the original taxa included, our modified version supports the TH, while adding *Gerobatrachus* and the “lepospondyl” *Brachydectes* results in the LH.

Until recently, few studies supported the LH, but two recent dissertations which contain large phylogenetic analyses bolster it (Pawley, 2006: 239 and appendix 16; Germain, 2008a). Pawley (2006) built upon the data matrix by Ruta *et al.* (2003), which originally supported the TH; Germain (2008a) merely took the matrix by Ruta and Coates (2007), which likewise originally supported the TH, did not change its taxon or character sampling (except for fusing four correlated characters), and improved the accuracy of the scoring. Although Germain (2008a) found the LH to be only one step more parsimonious than the TH, continuation of his work by D. M. (supplemented by the fusion of further correlated characters and the addition of information published in or after 2007) has so far increased the difference to eight steps. Incidentally, the LH is no longer supported only by M. L. and his students; neither of us knew of Pawley’s dissertation till two years after the defense.

As with that by McGowan (2002), we find much to disagree with in the matrix of Anderson *et al.* (2008). Our rescored and/or recoded of 39 characters and all but five taxa (and ordering of 38 of the multistate characters for reasons explained by Marjanović and Laurin, 2008a: 163, partially overlapping with the set of rescored characters) supports the LH, with the lysorophian *Brachydectes* as the sister-group of Lissamphibia; the bootstrap and Bremer values for a “lepospondyl”-lissamphibian clade (Amphibia) which excludes all temnospondyls are high. Again as with the matrix by McGowan (2002), our changes range from the correction of probable typographic errors (like the alleged lack of a cleithrum in *Triadobatrachus* and “frogs” in character 193, or the alleged absence of a caudolateral flange or corner on the pterygoid that constricts the subtemporal fenestra in *Seymouria* and *Limnoscelis* in character 120) to disagreements of interpretation; for example, having compared all described temnospondyl tarsi, we are not convinced that an *os basale commune* (character 207) is present in *Gerobatrachus* (Marjanović and Laurin, 2008a: 168–169), and although strong cases for tooth pedicely (character 99) in closely related temnospondyls have been made, we do not think the published evidence establishes whether *Gerobatrachus* shares this feature (Marjanović and Laurin, 2008a: 179), so we had to score these two characters as unknown in *Gerobatrachus*. See the Electronic Supplementary Material for more information about the cells that were rescored, the supporting references, the methods, the resulting topology, the support values, and the modified matrix.

Furthermore, possibly pedicellate teeth have now been described in an aïstopod (Germain, 2008b) and may be present in another (Carroll, 1998: fig. 4B). In addition, Milner (1980: 392) calls the teeth of the nectridean *Scincosaurus* “pedicellate [...] without a line of abscission”; to the best of our knowledge, the teeth of *Scincosaurus* have never been mentioned in the literature before or since (most notably not by Bossy and Milner, 1998).

Further research will be necessary to determine if pedicely is much more widespread than previously thought.

For all these reasons, we conclude that the polyphyly hypothesis is less likely than suggested by Anderson (2008), and that the lepospondyl hypothesis seems to be at least as well supported as the temnospondyl hypothesis.

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References

- Anderson JS. 2002. Revision of the aïstopod genus *Phlegethontia* (Tetrapoda: Lepospondyli). *J Paleont* 76:1029–1046.
- Anderson JS. 2008. Focal review: The origin(s) of modern amphibians. *Evol Biol* 35:231–247.
- Anderson JS, Reisz RR, Scott D, Fröbisch NB, Sumida SS. 2008. A stem batrachian from the Early Permian of Texas and the origin of frogs and salamanders. *Nature* 453:515–518.
- Bolt JR. 1980. New tetrapods with bicuspid teeth from the Fort Sill Locality (Lower Permian, Oklahoma). *N Jb Geol Paläont Mh* 1980:449–459.
- Bossy KA, Milner AC. 1998. Order Nectridea MIALL 1875. In: Carroll RL, Bossy KA, Milner AC, Andrews SM, Wellstead CF. *Lepospondyli*. Part 1 of Wellnhofer P, editor. *Encyclopedia of Paleoherpétology*. Munich: Dr. Friedrich Pfeil. p 73–131.
- Carroll RL. 1998. Cranial anatomy of ophiderpetontid aïstopods: Palaeozoic limbless amphibians. *Zool J Linn Soc* 122:143–166.

- Germain D. 2008a. Anatomie des Lépospondyles et origine des Lissamphibiens [PhD thesis]. Paris: Muséum National d'Histoire Naturelle. 351 p.
- Germain D. 2008b. A new phlegethontiid specimen (Lepospondyli, Aistopoda) from the Late Carboniferous of Montceau-les-Mines (Saône-et-Loire, France). *Geodiversitas* 30:669–680.
- Germain D, Laurin M. 2009. Evolution of ossification sequences in salamanders and urodele origins assessed through event-pairing and new methods. *Evol Dev.* 11:170–190.
- Huelsenbeck JP. 1991. Tree-length distribution skewness: an indicator of phylogenetic information. *Syst Zool* 40:257–270.
- Hugall AF, Foster R, Lee MSY. 2007. Calibration choice, rate smoothing, and the pattern of tetrapod diversification according to the long nuclear gene RAG-1. *Syst Biol* 56:543–563.
- Johanson Z, Joss J, Boisvert CA, Ericsson R, Sutija M, Ahlberg PE. 2007. Fish fingers: digit homologues in sarcopterygian fish fins. *J Exp Zool B (Mol Dev Evol)* 308:757–768.
- Marjanović D, Laurin M. 2007. Fossils, molecules, divergence times, and the origin of lissamphibians. *Syst Biol* 56:369–388.
- Marjanović D, Laurin M. 2008a. A reevaluation of the evidence supporting an unorthodox hypothesis on the origin of extant amphibians. *Contrib Zool* 77:149–199.
- Marjanović D, Laurin M. 2008b. Assessing confidence intervals for stratigraphic ranges of higher taxa: the case of Lissamphibia. *Acta Palaeont Pol* 53:413–432.
- McGowan GJ. 2002. Albanerpetontid amphibians from the Lower Cretaceous of Spain and Italy: a description and reconsideration of their systematics. *Zool J Linn Soc* 135:1–32.
- Milner AC. 1980. A review of the Nectridea (Amphibia). In: Panchen AL, editor. *The Terrestrial Environment and the Origin of Land Vertebrates*. London and New York:

Academic Press. p 377–405.

- Pawley K. 2006. The postcranial skeleton of temnospondyls (Tetrapoda: Temnospondyli) [PhD thesis]. Melbourne: La Trobe University. 442 p. Downloadable (free access) at <http://www.lib.latrobe.edu.au/thesis/public/adt-LTU20061124.124055/index.html>
- Ruta M, Coates MI. 2007. Dates, nodes and character conflict: addressing the lissamphibian origin problem. *J Syst Paleontol* 5:69–122.
- Schoch RR. 1992. Comparative ontogeny of Early Permian branchiosaurid amphibians from southwestern Germany. *Palaeontogr A* 222:43–83.
- Schoch RR, Milner AR. 2004. Structure and implications of theories on the origin of lissamphibians. In: Arratia G, Wilson MVH, Cloutier R, editors. *Recent Advances in the Origin and Early Radiations of Vertebrates*. Munich: Dr. Friedrich Pfeil. p 345–377.
- Vallin G, Laurin M. 2004. Cranial morphology and affinities of *Microbrachis*, and a reappraisal of the phylogeny and lifestyle of the first amphibians. *J Vertebr Paleontol* 24:56–72.

Electronic Supplementary Material 1 to

The origin(s) of modern amphibians: a commentary

by

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This supplement contains the methods, materials and results of our reanalysis of the data matrix by Anderson *et al.* (2008) and of our application of the skewness test by Huelsenbeck (1991) to the matrices by Anderson *et al.* (2008) and Vallin and Laurin (2004), including supplementary references and the Supplementary Tables. The Supplementary Figure (ESM2), the data matrix (ESM3), and the trees used to find the ancestral states of the “frogs” (ESM4) and “salamanders” OTUs (ESM5) are separate files.

Legend to the Supplementary Figure (ESM2): Strict consensus of the four most parsimonious trees (see Results section). Numbers below internodes are bootstrap percentages (in bold if 50 or higher; “–” indicates clades contradicted by the bootstrap tree, always by clades with bootstrap percentages of 40 or less), numbers above internodes are Bremer values. Some or all of the Bremer values shown as “ ≥ 5 ” might actually be 5, because we were unable to find all trees that were up to 5 steps longer than the most parsimonious trees, although the fact that an earlier iteration of this analysis, with a dataset that differed only in two cells, found the same results makes this possibility unlikely.

All amphibians that are not lissamphibians are “lepospondyls” (alternatively, a case could be made that Lepospondyli is – under the present phylogenetic hypothesis – a junior synonym of Amphibia). Strictly speaking, Amphibia is defined with respect to Amniota and therefore cannot be applied to this tree, which lacks amniotes, but the close relationship of *Limnoscelis* and Amniota has never been doubted. The temnospondyl *Gerobatrachus* is marked with an arrow.

Methods

The matrix was manipulated in Mesquite 2.6+ (Maddison and Maddison, 2009) on an Intel Macintosh; the analyses were conducted in PAUP* 4.0b10 (Swofford, 2003) on a G5 Macintosh.

We treated polymorphism differently from uncertainty (PAUP* command: “pset mstaxa=variable”). For the reasons explained below, characters 103, 146, 163 and 217 were excluded from all analyses (but kept in the matrix to retain the original numbering for all characters); furthermore, characters 137, 161, 188, and (now) 207 are parsimony-uninformative.

The main analysis was a heuristic search with 10,000 addition-sequence replicates (random addition sequence, 10 trees held at each step, TBR swapping, no limit on rearrangements). Five additional analyses were performed to find all trees that are up to one to five steps longer (the search for those up to five steps longer than the minimum length had to be terminated after 370,000 trees due to lack of memory), in order to find Bremer support values; these analyses had otherwise identical settings to the main analysis, except for consisting of only 200 addition-sequence replicates. We also conducted a bootstrap analysis with 200 bootstrap replicates (100 addition-sequence replicates within each, 10 trees held at each step, TBR swapping).

The term “loss character” seems never to have been defined. We opt for a strict definition: the wholesale loss (on the most parsimonious trees of the publication in question) of entire bones or structures between bones (like fontanelles or the parietal foramen) or on them (canals for the lateral-line system, dermal sculpturing). We have not counted the loss of processes of bones, even conspicuous ones like the tabular “horns”, or other potentially continuous characters; we have also not counted meristic characters (like the number of vertebrae in certain parts of the column). The lists of loss characters in the matrices of Anderson *et al.* (2008) and Vallin and Laurin (2004) are presented in Supplementary Tables 1 and 2, respectively.

We calculated the g_1 statistic for the loss characters and the remaining characters of both matrices (using PAUP*; Swofford, 2003). Unfortunately, Huelsenbeck (1991: table 1; reproduced in Supplementary Table 3, left side) only lists the 5% and 1% significance thresholds for 6, 7 and 8 taxa; the matrices by Vallin and Laurin (2004) and Anderson *et al.* (2008) are both much larger. However, from 6 to 8 taxa, the thresholds decrease with increasing taxon number. We conclude that all values (Supplementary Table 3, right side) are most likely highly significant, so that all four partitions contain a phylogenetic signal of comparable and high strength.

Modified scores, character definitions and state delimitations

All characters were left unordered by Anderson *et al.* (2008). However, potentially continuous multistate characters should always be ordered, because the assumption that any character state can more easily change into a similar state than into a very different one was already used for subdividing the potential continuum into discrete states (Wiens, 2001; see Marjanović and Laurin, 2008, for a previous application of this principle). Altogether, we ordered characters 1, 2, 16, 29, 32, 34, 39, 74, 75, 82, 87, 103, 115, 126, 128, 130, 134, 136, 138, 144, 145, 149, 155, 159, 170, 172, 179, 181, 183, 195, 197, 198, 200, 201, 204, 208, 209, and 211. In four cases, the state numbers were not in the appropriate order for this. We exchanged states 0 and 1 of characters 16 and 115. The former state 3 of character 34 was put at the other end, so that the new states 0, 1, 2 and 3 correspond to the old states 3, 0, 1 and 2. The states of character 145 had to be rearranged more extensively: the new states 0, 1, 2 and 3 correspond to the old states 2, 0, 1 and 3, respectively.

More trivially, Anderson *et al.* (2008) used “?” for missing data and “-” to indicate inapplicable codings, a distinction that no currently existing phylogenetics program can deal with. The hyphen is interpreted as a gap (in a DNA or protein sequence, implying a mixed dataset) by PAUP*; fortunately, the default setting is to interpret gaps as missing data. To avoid problems with the other setting (which is to interpret gaps as a 5th base/21st amino acid), we have replaced every hyphen by a question mark in the NEXUS file (ESM3).

Yet more trivially, we have corrected the spelling “*Hapsidoparion*” to *Hapsidopareion*.

Because of the size of this matrix and various time constraints, we have not checked the accuracy of every cell; we have revised mostly those characters whose distribution of states across taxa seemed anomalous and those where we redelimited the states. Indeed, we only noticed the errors in character 193 (see below) after the first submission and had to repeat all analyses; it is therefore possible that the matrix still contains errors.

Despite this, only changes are listed below (as in Marjanović and Laurin, 2008); listing the cells we found to be accurate would require too much space and time.

Character 16: lacrimal possesses both dorsal (prefrontal/frontal) and ventral (jugal/maxillary) processes (0), ventral process only (1), or neither (2) (ordered). In *Acanthostega*, as correctly coded by character 15, the lacrimal does not participate in the orbit

margin (being excluded by prefrontal-jugal contact), so we have scored it as inapplicable (= unknown).

At least some specimens of *Micropholis* possess the caudal process (Schoch and Rubidge, 2005: figs. 3B, E), giving it states 1 and 2 (a polymorphism).

We have also changed Branchiosauridae from state 2 to state 1, because the morphologically most mature known specimens possess the ventral process (Schoch and Fröbisch, 2006: figs. 1C, D).

The ventral process is likewise present in Albanerpetontidae (McGowan, 2002: fig. 5B).

This also seems to be the ancestral state for those salamanders (that is, the hynobiids) which possess a lacrimal which participates in the orbit margin, for example *Hynobius* (Carroll and Holmes, 1980: fig. 4A, C).

Limnoscelis appears to lack both processes (Fracasso, 1983: fig. 3A), although this could be a matter of definition.

Character 22: Prefrontal-postfrontal suture (0); frontal participates in margin of orbit (1). Anderson *et al.* (2008) defined state 0 only as the opposite of state 1, without mentioning the fact that it contains several different states (for example, *Brachydectes* lacks postfrontals and has a prefrontal-parietal contact which excludes the frontal from the orbit margin). Furthermore, redefining state 0 as we have done makes explicit that many animals that, at face value, show state 1 should actually be scored as inapplicable.

This includes all lissamphibians in the present matrix except *Eocaecilia* (which was correctly scored as possessing state 0), because they all lack postfrontals and therefore cannot help lacking a prefrontal-postfrontal contact.

Character 26: Dorsal process of premaxilla: broad, low, indistinct (0); alary process (broad, vaguely triangular) (1); moderately high, vaguely rectangular, or acutely triangular linked directly to base (2); narrow and long, along the sagittal plane or parasagittal (3) (unordered). This coding differs from that by Anderson *et al.* (2008) so as to better fit the morphological diversity seen especially in extant amphibians. It is congruent with the findings of Good and Wake (1992), but it recognizes two states within the condition that Good and Wake (1992) considered primitive.

Greererpeton shows state 1 (Smithson, 1982).

Seymouria baylorensis is borderline between states 0 and 2 (see Laurin, 1996); we have decided on state 2.

Limnoscelis possesses state 2 (Fracasso, 1983).

All states except 0 occur in salamanders: *Karaurus* has state 1 (Ivachnenko, 1978: fig. 1); Hynobiidae shows states 3 (*Hynobius tsuensis*), 1 (*Batrachuperus sinensis*), and indeterminate (*Hynobius naevius*) (Carroll and Holmes, 1980: fig. 4). According to figure 5 of the same paper, state 2 occurs in Cryptobranchidae. *Ambystoma* (*ibid.*, fig. 6) and plethodontids (*ibid.*, fig. 7) possess state 3. Salamandrids can have state 3 (*ibid.*, fig. 8A) or be indeterminate (*ibid.*, fig. 8B). Proteidae exhibits state 3 (*ibid.*, fig. 9), as do Amphiumidae (*ibid.*, fig. 10) and Sirenidae (*ibid.*, fig. 11). To code the single OTU “salamanders”, we have used the same approach as in Marjanović and Laurin (2008): optimizing this diversity onto the phylogenetic hypothesis shown in fig. 8 of Wiens *et al.* (2005) when *Karaurus* is added as the sister-group of Urodela, states 1 and 3 emerge as most parsimoniously plesiomorphic for Caudata as a whole, so we have assigned it state 1 or 3 (partial uncertainty).

Most anurans have state 3 (Duellman and Trueb, 1986: figs. 13-17 and 13-18).

Most “microsaurs” also had to be rescored according to Carroll and Gaskill (1978).

Character 34: Caudal margin of the skull roof: undulating (0); concave (1); straight (2); convex (3) (ordered). The place of state 0 in this sequence is certainly debatable; an alternative would have been to consider the median caudal projection of the skull roof a separate character, as Ruta and Coates (2007: 96) did (twice: characters POSPAR 4 and POSPAR 8), defensible by the fact that most but not all taxa with an undulating margin would otherwise count as concave.

The margin is straight in *Brachydectes* (Wellstead, 1991: figs. 2, 3, 8) and undulating in *Sauropseura* (Bossy and Milner, 1998: fig. 53B), arguably *Ptyonius* (ibid., fig. 53G), and *Eocaecilia* (Jenkins *et al.*, 2007: figs. 1, 2).

Character 39: Large otic notch approaching orbit: absent (0); intermediate (1); close (2) (ordered). Because the albanerpetontids lack an otic or other temporal notch or other embayment, we consider this character (which appears to describe the distance between the rostral margin of the “otic notch” and the caudal margin of the orbit) inapplicable to them.

Character 51: Parietal-squamosal contact: absent (0), present (1). This character is only applicable when the supratemporal is absent. (When present, the supratemporal extends from the tabular and/or the caudal margin of the skull roof to the postorbital, unless a temporal fenestra intervenes as it does in many amniotes and aistopods; in the latter the fenestra separates the parietal from the squamosal, making the present character likewise inapplicable.) The presence of the supratemporal is already coded as state 0 of character 5; keeping state 51(0) for taxa with 5(0) would therefore correlate these two characters.

Both states occur in *Microbrachis* (Carroll and Gaskill 1978: figs. 77, 78).

Character 59: Tabular: present (0); absent (1). Having examined the only known (and very confusing) specimen of *Triadobatrachus* (see Marjanović and Laurin, 2008: appendix-table 1), we provisionally disagree with J. Anderson’s otherwise unpublished reinterpretation of its skull roof and agree with the literature (*e.g.*, Rage and Roček, 1989) that tabulars are absent in this animal.

Character 74: Number of premaxillary teeth: ≥ 10 (0); 5–9 (1); < 5 (2) (ordered). All three states occur in salamanders: *Karaurus* has about 25 premaxillary teeth (Ivachnenko, 1978: 364, fig. 1a); *Batrachuperus sinensis* (Hynobiidae) possesses 9; *Cryptobranchus* and *Ambystoma* both possess more than 10; *Phaeognathus hubrichti* (Plethodontidae) shows about 8; *Salamandra atra* exhibits only about 4; more than 10 are present in *Notophthalmus viridescens* (Salamandridae) and *Necturus* (Proteidae); only about 5 occur in *Amphiuma* and in *Habrosaurus* (Sirenidae). Using the same approach and the same references as for character 26, states 0 and 1 emerge as candidates for the plesiomorphic state, so we have coded the salamander OTU as possessing state 0 or 1.

Using the same approach, only state 0 (rather than 0 or 1) emerges as plesiomorphic for frogs: *Yizhoubatrachus* (10 on the right, 11 on the left premaxilla: Gao and Chen, 2004), *Notobatrachus* (reconstructed with 14 to 15: Sanchíz, 1998: fig. 20B), *Mesophryne* (16 teeth are preserved on an incomplete premaxilla: Gao and Wang, 2001: 461), *Eodiscoglossus* (at least 15 tooth positions: Evans *et al.*, 1990: 302), and *Ascaphus* (Carroll and Holmes, 1980: fig. 3A) all show well over 10 teeth; the variation in *Leiopelma* even includes 15 to 25 teeth (Sanchíz, 1998: 16, fig. 19G).

Both *Albanerpeton* (McGowan, 2000: 367; Venczel and Gardner, 2005: 1282) and *Celtdens* (McGowan, 2002: 5) possess both state 0 and state 1. The only known premaxilla of *Anoualerpeton* with a complete tooththrow preserves 10 teeth (Gardner *et al.*, 2003: 308). Taken at face value, this would make state 0 more parsimonious as the plesiomorphic

condition of Albanerpetontidae, but because of the limited sample size we have preferred to score Albanerpetontidae as polymorphic.

Character 75: Number of maxillary teeth: ≥ 30 (0); 20–29 (1); < 20 (2) (ordered). *Albanerpeton* possesses states 1 and 2 (Gardner, 1999: 536; 2000: 367; Venczel and Gardner, 2005: 1282), while the composite maximum estimate for *Anoualerpeton* (Gardner *et al.*, 2003: 308) is 25 teeth, thus staying in state 1. In the absence of evidence to the contrary from *Celtedens*, and due to the small sample size of *Anoualerpeton*, we have extrapolated the polymorphism of *Albanerpeton* to Albanerpetontidae as a whole.

Character 84: Occipital condyle: concave (0); convex (1). State 1 was ascribed to *Gerobatrachus*, but the occipital condyle is not preserved (Anderson *et al.*, 2008: figs. 1, 2). We have therefore scored *Gerobatrachus* as unknown.

Character 85: Occipital condyle: single (0); double (1). We have changed the score of *Gerobatrachus* from state 1 to unknown for the reasons mentioned under character 84.

Ecolsonia, on the other hand, possesses a “distinctly double occipital condyle” (Berman *et al.*, 1985: 16).

State 1 is shared by *Asaphstera* (Carroll and Gaskill, 1978: fig. 6E), *Hapsidopareion* (*ibid.*, fig. 13B), *Pelodosotis* (*ibid.*, fig. 48A), *Micraroter* (*ibid.*, fig. 53), *Pantylus* (*ibid.*, fig. 25), *Cardiocephalus sternbergi* (*ibid.*, figs. 30B, C), *Cardiocephalus peabodyi* (judging from the articular surfaces on the atlas: *ibid.*, fig. 31), *Euryodus primus* (for the same reason: *ibid.*, figs. 37, 38, 41), *Euryodus dalyae* (*ibid.*, figs. 42–44), *Rhynchonkos* (*ibid.*, figs. 63–65), *Microbrachis* (*ibid.*, figs. 77–79; Vallin and Laurin, 2004: 62), *Stegotretus* (Berman *et al.*, 1988: 310), *Sauropleuria scalaris* (Bossy, 1976: 96 and fig. 25b), *Ptyonius* (Bossy, 1976: 145), and *Brachydectes* (Wellstead, 1991: figs. 2, 15C).

“The surfaces that formed part of the occipital condyle cannot be seen” in *Tuditonus* (Carroll and Baird, 1968: 12), suggesting that the reconstruction in ventral view by Carroll and Gaskill (1978: fig. 4) is just that – a reconstruction. We have therefore changed its score to unknown.

Much the same holds for *Batropetes* (Carroll and Gaskill, 1971: 454) and *Saxonerpeton* (Carroll and Gaskill, 1978: 34).

Bossy (1976: 209) describes the occipital condyle of *Urocordylus* as “semidouble”, suggesting an intermediate morphology which we have coded as unknown for the sake of simplicity.

Character 86: Jugular foramen: between opisthotic and exoccipital (0); through exoccipital (1). This character has not been described in *Doleserpeton*, so we have changed its score to unknown.

In frogs, the exoccipital is almost always fused to other braincase bones (Duellman and Trueb, 1986: table 13-1), making it impossible to evaluate the position of the jugular foramen relative to the opisthotic and forcing us to score the frogs as unknown.

Character 87: Jaw articulation: caudal to occiput (0); even with occiput (1); rostral to occiput (2); far rostral to occiput (over 20% of basal skull length) (3) (ordered). The frogs were scored as 0 (a state that really occurs in the separate OTU *Triadobatrachus*, which was scored correctly; Rage and Roček, 1989), but *Notobatrachus* (Sanchíz, 1998), *Vieraella* (Carroll and Holmes, 1980: fig. 3), *Ascapus* (same figure, and Noble, 1931: fig. 81A), and others like *Pelobates* possess state 2. *Mesophryne* appears to show state 1, but is incompletely preserved and somewhat disarticulated (Gao and Wang, 2001: fig. 4). We have scored it as unknown (though we should have restricted this somewhat).

Character 89: Palatine fangs: present (0); absent (1). This character originally referred to “palatal teeth” in general, but this would correlate to several others like 94 (presence and arrangement of vomerine teeth) and especially 90, which treated fangs and tooththrows as states of the same character, even though both can occur together (like in *Acanthostega*: Clack, 1994: fig. 11C).

Asaphestera is restored by Carroll and Gaskill (1978: fig. 7) as possessing a row of very large teeth on the palatine and ectopterygoid. We count this as state 0 (see Marjanović and Laurin, 2008: 180), even though we also count it as state 1 of the next character (as originally scored).

The palate of *Diceratosaurus* is illustrated in Jaekel (1902: plate III); it shows state 1.

The palatine of *Adelogyrinus* is unknown (Andrews and Carroll, 1991).

Character 90: Palatine teeth (marginal-sized): absent (0); multiple in rows (1); multiple random (2) (unordered). See character 89.

Greererpeton possesses a row of palatine teeth (Smithson, 1982).

Limnoscelis lacks palatine teeth (Fracasso, 1983: fig. 2).

Micropholis possesses a palatine tooththrow (Schoch and Rubidge, 2005: fig. 1D).

The only known adult specimen of *Amphibamus* possesses fangs instead of a tooththrow (Daly, 1994).

The palatine teeth of *Gerobatrachus* are small enough to count as denticles, and are not arranged in rows (Anderson *et al.*, 2008: fig. 2a).

All teeth on the palatine of *Microbrachis* are small enough to count as denticles (Carroll and Gaskill, 1978); this means state 0.

The palate of *Diceratosaurus* is illustrated in Jaekel (1902: plate III); it shows state 1.

The palatine of *Adelogyrinus* is unknown (Andrews and Carroll, 1991).

The palatine of *Brachydectes* is toothless (Wellstead, 1991).

Triadobatrachus has a preserved and apparently toothless palatine, so we have given it state 1 (Rage and Roček, 1989).

Palatine teeth are likewise always absent in the “frogs” OTU (Duellman and Trueb, 1986: 318).

Character 94: Vomerine teeth: absent (0); forming a single row (1); forming a field (2) (unordered). This definition is new. “Teeth” are about the size of the marginal teeth, markedly larger than denticles and smaller than fangs; state 2 is what is exhibited by the sirenids.

Doleserpeton possesses states 1 and 2; there is both a row (near the choana) and a field (Bolt 1969). This might, however, be an artefact of the fact that the marginal teeth are tiny – no teeth of *Doleserpeton* are sufficiently smaller to count as denticles.

Amphibamus, *Gerobatrachus*, and *Limnoscelis* show state 0.

Character 99: Tooth pedicely: absent (0); present (1). We have scored *Gerobatrachus* as unknown for the reasons explained in Marjanović and Laurin (2008: 179) and done the same with *Oestocephalus*, *Phlegethontia*, and *Scincosaurus* for the reasons explained in the main text of the present paper.

Character 100: Denticles on vomers: present (0); absent (1). *Gerobatrachus* was scored 0, but, judging from the illustrations rather than the text of Anderson *et al.* (2008), its vomerine teeth are unambiguously small enough to count as denticles, giving it state 0.

Character 102: Denticles on parasphenoid: present (0), absent (1). *Triadobatrachus*, frogs, salamanders and albanerpetontids (*Celtdens*: McGowan, 2002: fig. 13; denticles are

also not mentioned for the incomplete parasphenoid part of the os basale of *Albanerpeton inexpectatum* by Estes and Hoffstetter, 1976: 314) possess state 1, while *Eocaecilia* shows state 0 (Jenkins *et al.*, 2007).

Character 103: Palatal teeth: larger than marginals (0), equal to marginals (1). This character is correlated to several others (such as 89, 90, 93 and 94); we have therefore excluded it from our analyses.

Character 105: Parasphenoid basal plate: roughly quadrangular, basipterygoid articulations narrowly spaced (0); about as broad as long, articulations moderately distant (1); rectangular laterally, anteroposteriorly narrow, basipterygoid articulations distant (2) (ordered). State 1 was added because the condition in the following OTUs fits neither state 0 nor state 2 (the former state 1):

Tuditanus (Carroll and Gaskill, 1978: fig. 4); *Hapsidopareion* (*ibid.*, fig. 13); *Saxonerpeton* (*ibid.*, fig. 22); *Cardiocephalus sternbergi* (*ibid.*, fig. 30); *Euryodus primus* (*ibid.*, figs. 36–38); *Euryodus dalyae* (*ibid.*, figs. 42, 43); *Pelodosotis* (*ibid.*, fig. 48); *Micraroter* (*ibid.*, fig. 53); *Rhynchonkos* (*ibid.*, figs. 63–65); *Microbrachis* (*ibid.*, fig. 77; Vallin and Laurin, 2004: fig. 5B); *Stegotretus* (Berman *et al.*, 1988: fig. 8); *Ptyonius* (Bossy, 1976: fig. 49; Bossy and Milner, 1998: fig. 55B); *Diploceraspis* (Beerbower, 1963: fig. 4); *Scincosaurus* (Bossy and Milner, 1998: fig. 57D); *Oestocephalus* (Carroll, 1998: fig. 3); *Brachydectes* (Wellstead, 1991: fig. 2); *Eocaecilia* (Jenkins *et al.*, 2007); salamanders (Carroll and Holmes, 1980).

Furthermore, the condition is unknown in the following OTUs: *Gerobatrachus* (where the caudal part of the palate is very fragmentary and somewhat disarticulated, so that the statement in the text about the shape of the parasphenoid may not be warranted; Anderson *et al.*, 2008); *Urocordylus* (Bossy and Milner, 1998: fig. 55A); *Batrachiderpeton* (where the parasphenoid has a unique shape and is incompletely preserved; Bossy and Milner, 1998: fig. 57B); *Adelogyrinus* (Andrews and Carroll, 1991: fig. 6).

The condition in *Phlegethontia* counts as state 0 (Anderson, 2002: fig. 4:2).

Character 109: Stapes orientation: lateral, towards quadrate (0); dorsal, towards squamosal embayment, elongate columella (1). Salamanders possess state 0.

Character 120: Pterygoid extending into subtemporal fenestra, constricting it: no (0); yes (1). The pterygoid sometimes extends a corner (especially when a transverse flange is present on the pterygoid, like in *Seymouria* [Laurin, 1996] or *Limnoscelis* [Fracasso, 1983]; incidentally, both were erroneously scored as 0, which we can only explain as a typographic error; but also in the absence of a flange, such corners occur, like in the correctly scored *Eryops*: Sawin, 1941: plate 2) or a rounded flange (like in the correctly scored *Balanerpeton*: Milner and Sequeira, 1994) into the subtemporal fenestra, which was erroneously called “posttemp fen” in the supplementary information of Anderson *et al.* (2008), the posttemporal fenestra being on the occiput.

State 1 is further present in *Dendrerpeton* (Milner, 1996: fig. 6), *Asaphestera*, *Hapsidopareion*, *Saxonerpeton* (all Carroll and Gaskill, 1978: fig. 108), *Euryodus primus* and *E. dalyae* (both Carroll and Gaskill, 1978: fig. 109), *Microbrachis* (Vallin and Laurin, 2004), and *Eocaecilia* (Jenkins *et al.*, 2007).

Branchiosauridae is polymorphic (Boy, 1978, 1986, 1987).

The condition in *Tuditanus* is unknown because the lower jaws are in place (Carroll and Baird, 1968).

In *Cardiocephalus sternbergi*, the condition is so intermediate (Carroll and Gaskill, 1978: fig. 30) that we score it as unknown.

The palate of *Batropetes* is too incomplete to tell (Carroll, 1991: fig. 5B).

The only illustration of the palate of *Sauropleura scalaris* that we know about is fig. 55C of Bossy and Milner (1998). It shows a juvenile. This may be interesting because the adult of *S. pectinata*, illustrated in fig. 55D, exhibits state 1. Therefore we do not take fig. 55C at face value and score *S. scalaris* as unknown rather than 0.

The only illustration of the palate of *Ptyonius* known to us is fig. 55C of Bossy and Milner (1998). It shows state 0 on the left side and state 1 on the right. Taking into account that this line drawing is a somewhat schematic reconstruction, we have scored *Ptyonius* as unknown.

The palate is entirely unknown in *Adelogyrinus*. The only adelogyrinid palate illustrated by Andrews and Carroll (1991: fig. 13C) is that of *Adelospondylus* (which, incidentally, appears to possess state 1, even though the reconstruction in fig. 13D omits it almost entirely); *Adelospondylus* is not *Adelogyrinus*.

State 0 is present in all salamanders (except *Hynobius*, *Ambystoma*, and the salamandrid *Tylototriton*: Ivachnenko, 1978; Carroll and Holmes, 1980), as well as in *Triadobatrachus* (Rage and Roček, 1989: fig. 2B).

In crown-group frogs, state 0 is present (Duellman and Trueb, 1986: figs. 13-3, 13-4), except, under a generous interpretation, in *Ascaphus* (Carroll and Holmes, 1980: fig. 3A). State 0 is also found in *Vieraella* (Carroll and Holmes, 1980: fig. 3B) and *Yizhoubatrachus* (Gao and Chen, 2004), while *Notobatrachus* (Sanchíz, 1998: fig. 20B) and *Mesophryne* (Gao and Wang, 2001) possess state 1. In the continued absence of data from animals like *Prosalirus* (Shubin and Jenkins, 1995), whether the optimization for the “frogs” OTU is state 0 or ambiguous depends on how generous we are with *Ascaphus*; because there is no flange or other process in *Ascaphus* – instead, the rod-shaped pterygoid is bent as a whole to make the medial margin of the subtemporal fenestra concave instead of straight or convex –, we have decided to count *Ascaphus* and thus the “frogs” as a whole as having state 0 (which happens to be shared by *Triadobatrachus* and the salamanders, see above).

Character 121: Ectopterygoid: present, with fang-pit pair (0); present, without fang-pit pair (1); absent (2) (unordered). Although this character could be ordered on the grounds that the present sequence of states could be interpreted as a gradual shrinking of the ectopterygoid, we have not ordered it because we have not tested the assumption that there is a strong correlation between the size of the ectopterygoid and the presence of fangs.

State 2 is new (Anderson *et al.* did not distinguish it from state 1). It occurs in *Doleserpeton* (Bolt, 1969), both pantylids (*Pantylus*: Carroll and Gaskill, 1978: fig. 25; *Stegotretus*: Berman *et al.*, 1988: 308), *Brachydectes* (Wellstead, 1991: fig. 2), salamanders (Carroll and Holmes, 1980), *Triadobatrachus* (Rage and Roček, 1989) and the other frogs (Duellman and Trueb, 1986: figs. 13-17 and 13-18).

While the presence of a very small ectopterygoid (too small to bear fangs; known, if indeed homologous to the ectopterygoid, in several extant teresomatan caecilians – see Marjanović and Laurin, 2008: 181, for a list and some discussion) cannot be excluded in *Eocaecilia* (Jenkins *et al.*, 2007: 304), it is at least as probable that this bone was entirely absent (as reconstructed in Jenkins *et al.*, 2007: fig. 1). We have therefore changed its score to state 1 or 2.

Character 146: Vertebral development: arches, then centra (0); arches and centra simultaneously (1). In the absence of a very detailed growth series, it is impossible to distinguish state 1 from a simple lack of temporal resolution. There is widespread agreement about this among development biologists (*e.g.*, Bininda-Emonds *et al.*, 2003: 341). Such detailed growth series that show the stages of the ossification of the vertebral column are probably not available in any fossil limbed vertebrates other than the “branchiosaurs”, where

the development of most endochondral bones was delayed (Schoch and Fröbisch, 2006). We have therefore excluded this character, which was already coded for only nine taxa (including two extant ones, one of which is polymorphic), from our analyses.

Perhaps not surprisingly, however, state 0 is known to occur in *Utaherpeton* (Carroll and Chorn, 1995: fig. 4b).

Character 163: Neural arches of the trunk: paired (0), fused (1). Except (presumably) in *Acanthostega* (which is polymorphic), this character is correlated to ontogenetic age and paedomorphosis. Therefore, all occurrences of state 0 except (presumably) *Acanthostega* should be scored as unknown (Wiens et al., 2005); this would make the character parsimony-uninformative. We have excluded this character from all analyses.

Character 168: Cranial surface of atlas centrum: same size as caudal surface (0); laterally expanded (1). The cranial surface is slightly expanded in *Sauroplorea scalaris* (Bossy, 1976: 102 and fig. 25b), giving it state 1.

The same holds for *Ptyonius* (*ibid.*, fig. 51a) and *Urocordylus* (*ibid.*, 215).

The only preserved atlas centrum of *Keraterpeton* we know of (Huxley and Wright, 1867; Jaekel, 1902) is in dorsal view, hiding the cranial and the caudal surface and forcing us to score *Keraterpeton* as unknown.

The condition seems to be likewise unknown in *Diceratosaurus* and *Scincosaurus*.

Brachydectes possesses state 1 (Wellstead, 1991: figs. 11C, 18).

Character 179: Ribs: at least as long as three successive articulated vertebrae and curved in adults (0); like state 0, but straight (1); shorter than three successive articulated vertebrae in adults (2) (unordered). Anderson *et al.* (2008) called the states “elongated and sometimes curved”, “straight”, and “short, simple rod”, but rib length and curvature are continua, so quantification was necessary to make reproducible coding possible. We have imported the quantification we used in Marjanović and Laurin (2008: 177); all the cautionary notes mentioned there apply here.

While close to the limit (like, *e.g.*, those of *Micraroter*: Carroll and Gaskill, 1978: fig. 57), the longest ribs of *Greererpeton* are longer than three successive vertebrae and are curved ventrally and (slightly) caudally (Godfrey, 1989: figs. 1, 3, 4); they are not outside the range of other OTUs with state 0, unlike the case of *Diplocaulus* with its long, very straight ribs.

Both the cervical (Berman *et al.*, 1985: fig. 13) and the proximal caudal (*ibid.*, fig. 10C, D) ribs of *Ecolsonia* – others are not preserved – are longer than three successive vertebrae and “moderately curved” (*ibid.*: 20).

The ribs of *Eocaecilia* are slightly curved, but curvature is not mentioned in either the original or our wording of state 2, and the ribs are only about as long as one vertebra (Jenkins *et al.*, 2007); we have accordingly scored *Eocaecilia* as possessing state 2 (like, incidentally, all other lissamphibians).

As explained in Marjanović and Laurin (2008: 177–178), we have also scored *Balanerpeton*, *Dendrerpeton*, *Acheloma*, *Cardiocephalus peabodyi*, and *Rhynchonkos* as having state 1.

In *Euryodus primus*, no complete ribs seem to be known (Carroll and Gaskill, 1978: 64), and the rib fragments illustrated by Carroll and Gaskill (1978: fig. 39) are much shorter than three vertebrae. We have therefore scored *E. primus* as unknown.

We are not aware of a description or illustration of the ribs of *Stegotretus*; Berman *et al.* (1988: 312) state that “[a]lthough ribs are ubiquitous, none are sufficiently preserved or exposed to warrant description.” Accordingly, we have scored *Stegotretus* as unknown.

The ribs of *Adelogyrinus* (Andrews and Carroll, 1991: fig. 6) are very similar to those of *Greererpeton*, but shorter (and somewhat straighter), fitting the definition of state 2.

Even though *Lethiscus* just reaches state 0 (Wellstead, 1982: figs. 1, 9A), the two aïstopods in the present matrix, *Ophiderpeton* and *Phlegethontia*, have clear cases of state 2 (Baird, 1964: fig. 1; Anderson, 2002, figs. 9, 10).

With both *Greererpeton* and *Ecolsonia* corrected to state 0, state 1 turns out to be unique to *Diplocaulus*. A case could be made for merging it into state 0 (ribs in state 2 are not necessarily straight), but we deem such a change unnecessary.

Character 181: Number of sacrals: one (0), two (1), three (2) (ordered). Several “microsaurs” were scored as possessing two sacral vertebrae. We cannot replicate this:

Carroll and Baird (1968: 14) argue that *Tuditonus* has two sacral vertebrae: “[...] two vertebrae with atypical ribs. The rib of the first, best seen in the type specimen, is thickened at the base and is evidently a sacral [rib]; the rib of the second vertebra is obscure. From the narrowness of the iliac blade it is doubtful that more than a single sacral rib could have articulated directly. Aside from its specialized rib, the principal [= first; Carroll and Gaskill, 1978: 16] sacral vertebra does not appear to differ significantly from its neighbors. The second appears to have a considerably larger transverse process.” There are only two specimens. In the type (Carroll and Baird, 1968: figs. 4, 5B) the sacral region (with the pelvis, the proximal part of the left femur, and the entire right hindlimb and the tail) is missing – except for the mentioned rib, which is clearly disarticulated and may therefore belong to a vertebra that is not preserved. The referred specimen (*ibid.*, figs. 6, 7) is preserved in ventral view, so that the pelvis obscures the sacral region more or less completely. We find ourselves unable to judge how many sacrals *Tuditonus* has (even the number 3 cannot be excluded) and therefore have to score it as unknown.

Carroll and Gaskill (1978: 57; see also fig. 31) state that, in the only articulated specimen of *Cardiocephalus peabodyi*, “[t]here are thirty-seven presacral vertebrae and two sacrals. The block is truncated in the middle of the second sacral, and there is not a trace of caudals.” The second sentence means that state 2 (three sacrals) cannot be excluded.

Two sacral ribs are known from *Pelodosotis*, but it is apparently not known if they are from the same vertebra, and then the specimen ends (Carroll and Gaskill, 1978: 85; see also fig. 58A). It must therefore be scored as unknown.

The sacral region of *Rhynchonkos* is apparently known only in the specimen FM-UR 2414, which is stated by Carroll and Gaskill (1978: 109) to possess two sacrals, but illustrated in a way that shows it is much too fragmentary to tell (*ibid.*, fig. 70). We conclude that *Rhynchonkos* must be scored as unknown.

In *Utaherpeton*, “[t]wo vertebrae lie adjacent to the central portion of the pelvic girdle. The more anterior is associated with a structure that can be identified as a sacral rib. The second is crushed below the level of the rest of the column and partially covered by the head of the femur. It is uncertain whether it was a second sacral or a first caudal. The ilium is so narrow that it is unlikely to have accommodated more than a single sacral rib.” (Carroll *et al.*, 1991: 318) Evidently, *Utaherpeton* must be scored as possessing one or two sacrals (and, of course, the latter possibility is clearly less probable than the former).

Character 183: Number of caudal rib pairs: 5 or more (0), 4 (1), 3 (2), 2 or fewer (3) (ordered). *Triadobatrachus* was given states 0 or 3; we have changed this to unknown because we cannot see a reason to exclude all other states (Rage and Roček, 1989: 11).

Character 191: Cleithrum head: aligned along cranial rim of scapula (0); caudodorsally enlarged head wrapping around dorsal scapula (1). *Sauroplorea scalaris* (Bossy, 1976: fig. 33) and *Brachydectes* (Wellstead, 1991: fig. 19D) show state 1.

Character 192: Cleithrum head: dorsally greatly expanded, much wider than shaft (0); simple rod without or with slight dorsal expansion (1). *Sauropleura scalaris* and *Brachydectes* again possess state 1 (same references as above). So does *Ptyonius* (Bossy, 1976: fig. 57a).

Character 193: Cleithrum: ossified (0); unossified (1). *Triadobatrachus* and frogs possess state 0 (Rage and Roček, 1989; Duellman and Trueb, 1986), leaving state 1 only to Albanerpetontidae and salamanders.

Character 201: Deltopectoral crest: weak (0), intermediate (1), prominent (2) (ordered). State 2 occurs in *Acanthostega* (Coates, 1996: fig. 16), *Proterogyrinus* (Holmes, 1984: fig. 26) and *Greererpeton* (Godfrey, 1989: fig. 18), while *Eocaecilia* possesses state 0 (Jenkins *et al.*, 2007: fig. 41).

The plesiomorphic condition for salamanders appears to be state 0, judging from *Chunerpeton* (assigned to Cryptobranchidae; Gao and Shubin, 2003: figs. 1, 2), *Ranodon* (Hynobiidae; Averianov, 1995: fig. 4), *Palaeoamphiuma* (Amphiumidae; Rieppel and Grande, 1998: fig. 5), and *Necturus* (Proteidae; Wischnitzer, 1979: fig. 2-8), even though *Parahynobius betfianus* (Hynobiidae; Venczel, 1999: fig. 2E).

Character 203: Humerus length: long, > 4 trunk centra (0); short (1). Where known (*Celtesdens*: McGowan, 2002: fig. 3), albanerpetontids possess state 1.

Character 206: Carpals: fully or partly ossified (0); unossified (1). Where known (*Celtesdens*: McGowan, 2002: 11), albanerpetontids possess state 0.

Character 207: Basale commune: absent (0); present (1). For the reasons explained in Marjanović and Laurin (2008) – such as comparisons to all other known temnospondyl tarsi and the fact that the tarsus of *Gerobatrachus* is highly incomplete and disarticulated from the lower leg on both sides –, we have changed the score of *Gerobatrachus* to unknown. This makes the character parsimony-uninformative.

Character 208: Number of digits in manus: 5 or more (0), 4 (1), 3 (2) (ordered). Apparently like Anderson *et al.* (2008), we have not counted the prepollex as a digit, and we have counted the metacarpal as part of the digit (so that *Scincosaurus*, which possesses what “may represent a very reduced fifth” metacarpal, retains state 0; Bossy and Milner, 1998: 99, fig. 66G).

We have furthermore accepted the decision by Anderson *et al.* (2008) to score *Eocaecilia* as having three digits per hand (and foot: character 219), something that Jenkins *et al.* (2007) consider highly likely but not certain.

Greererpeton was long interpreted as having four fingers per hand, and was scored accordingly. However, Coates (1996: 415) mentions and illustrates a fairly well preserved specimen with five fingers and mentions another that preserves four, one of which is the (distinctively small) fifth. Consequently, there is no evidence for four fingers (state 1) in any specimen of *G.* in the literature, and we code *G.* as pentadactyl (state 0).

Dendrerpeton preserves four distal carpals (Holmes *et al.*, 1998), which means that four or more fingers were present (state 0 or 1).

We infer that *Acheloma* had four or more fingers (state 0 or 1) because it appears to have had five distal carpals (Olson, 1941), of which the tiny cranialmost one could belong to a prepollex. As in *Dendrerpeton*, no metacarpals or fingers are preserved.

Micropholis possessed four fingers per hand (Schoch and Rubidge, 2005: 512), giving it state 1.

Tambachia was scored as unknown, but state 2 can be excluded (Sumida *et al.*, 1998), so we have scored it as showing state 0 or 1.

Saxonerpeton, on the other hand, is so incompletely preserved that none of the states can be ruled out (Carroll and Gaskill, 1978: 38).

Rhynchonkos was scored as 1, but while four metacarpals are preserved, a higher number cannot be excluded based on the preservation (Carroll and Gaskill, 1978: 111).

Urocordylus has state 0 (Bossy and Milner, 1998: 98–99, fig. 66A).

Bossy and Milner (1998) say *Keraterpeton* has state 1 as originally scored. However, Jaekel (1902) provides a line drawing of *Keraterpeton* (fig. 2) that shows articulated but incomplete hands which may well have had five fingers. Huxley and Wright (1867) claim five metacarpals and fingers to be present, but their plate XIX (or at least our bad photocopy of it) appears to show only four; Bossy (1976:304), contradicting Bossy and Milner (1998), writes “My own investigations corroborate the existence of the five-digit manus in this genus.” Awaiting a reply from Milner, we have scored *Keraterpeton* as possessing state 0 or 1.

Jaekel (1902) says three times explicitly that *Diceratosaurus* had pentadactyl hands, and illustrates a convincing pentadactyl hand in the specimen drawing (plate IV-6), even though Bossy & Milner (1998) explicitly disagree, stating the animal to be tetradactyl as originally coded. This is all the more surprising because Bossy earlier (1976: 304) wrote “*Diceratosaurus* definitely has five forefingers (pers. obs.)”. Because we have not been able to find the reason for this discrepancy, we have scored *Diceratosaurus* as possessing state 0 or 1.

Character 217: Number of distal tarsals: 6 (0), 5 or fewer (1). Three OTUs were scored as possessing six distal tarsals: *Acanthostega*, *Tuditanus*, and part of the salamanders (the “salamanders” OTU is scored as having state 0 or 1, but polymorphism was clearly intended). Regardless of which condition is plesiomorphic for salamanders, we think the extra distal tarsal is not homologous between any of these taxa. In *Acanthostega*, the caudalmost “distal tarsal”, to which three “metatarsals” attach directly, is better interpreted as the fibulare (*e.g.*, Coates, 1996: fig. 24; Johanson *et al.*, 2007: fig. 1); in the fully articulated referred specimen of *Tuditanus*, the distal tarsal that does not correspond to a metatarsal lies on the caudal side (Carroll and Baird, 1968: figs. 6, 7) and could therefore represent a postminimus (or, of course, something entirely different like a sesamoid), but neither the fibulare nor a prehallux; in the salamanders with an extra distal tarsal, this is the prehallux (as the first digital ray in the foot of *Acanthostega*, including the “first” distal tarsal rather than the “sixth”, might be) or the centrale 1 (the so-called “prehallux” in Duellman and Trueb, 1986: fig. 13-33B; the “centrale” in that figure is the centrale 4). We have therefore excluded this character from our analyses.

Results and conclusion

Four most parsimonious trees were found (length = 1264 steps, consistency index excluding parsimony-uninformative characters = 0.2659, retention index = 0.6003, rescaled consistency index = 0.1610). Their only differences lie in whether *Ecolsonia* (in two trees) or *Tambachia* (in the two others) is the sister-group of *Acheloma*, and in whether Pantylidae (= *Pantylus* + *Stegotretus*) is the sister-group of Ostodolepididae (= *Pelodosotis* + *Micraroter*) (in two trees) or of *Batropetes* + (*Rhynchonkos* + Gymnarthridae [= *Cardiocephalus* + *Euryodus*]) (in the two others). The strict consensus is shown in the Supplementary Figure.

All most parsimonious trees support the lepospondyl hypothesis, with a monophyletic Lissamphibia as the sister-group of *Brachydectes* and nested inside the “microsaurs”, even though the positions of the paraphyletic “nectrideans” and the aïstopods (*Oestocephalus*, *Phlegethontia*) as well as the topology of the “microsaurs” differ from those found by Vallin and Laurin (2004).

The bootstrap values (see the Supplementary Figure) are generally low – indeed, several clades are contradicted by the bootstrap tree, which for example finds Nectridea to be monophyletic (if only in 39% of the replicates) and arranges the “microsaurs” in a rather different way. The monophyly of Lissamphibia is found in only 23% of the bootstrap replicates, and the *Brachydectes*-Lissamphibia clade in only 22.

Nonetheless, Amphibia (= everything more closely related to Lissamphibia than to Amniota, here including Lissamphibia and the “lepospondyls”) has a bootstrap percentage of 68. A clade composed of all temnospondyls and all lissamphibians but no “lepospondyls” does not occur in the list put out by PAUP* that shows all clades found in 5% or more of the bootstrap replicates (reproduced below as Supplementary Table 4), arguing strongly against the temnospondyl hypothesis.

No clade that includes *Gerobatrachus* and some or all lissamphibians but no “lepospondyls” has a bootstrap percentage greater than 14 (Supplementary Tables 4 and 5), and no clade that includes all temnospondyls and some lissamphibians, but no “lepospondyls”, has a bootstrap value greater than 7. We think that the support for a special relationship between *Gerobatrachus* and some lissamphibians is not significant, because it is not only small in absolute terms and much smaller than the support for the clade composed of all “lepospondyls” and lissamphibians (68, see above), but also smaller than the support for clades which contain frogs, salamanders and lepospondyls but neither any temnospondyls nor *Eocaecilia*, contradicting both the monophyly of extant amphibians and all existing versions of the polyphyly hypothesis (18 and smaller); see Supplementary Table 5 for a more detailed comparison. At 22, the highest bootstrap percentage for any clade compatible with the polyphyly and not the lepospondyl hypothesis is not significantly higher either (the clade in question consists of *Eocaecilia* and the “microsaur” *Rhynchonkos*, and is the only clade with a bootstrap value above 14% that is compatible with the polyphyly hypothesis).

127 trees have the minimum length or are one step longer. Their strict consensus is poorly resolved (all nodes marked “1” in the supplementary figure collapse), e.g., *Eocaecilia* is no longer guaranteed to be the sister-group of the clade composed by the other “lissamphibians”, but it still contains a “nectridean”-aïstopod-“lissamphibian”-*Brachydectes* clade nested inside two more clades of amphibians, from which all of the “temnospondyl” clades (Temnospondyli having collapsed) are excluded. The majority-rule consensus is practically identical in topology to the strict consensus of the most parsimonious trees.

1635 trees are up to two steps longer than the minimum length. In the strict consensus, Amphibia survives and still excludes all “temnospondyls” while still including the fully resolved Albanerpetontidae + (salamanders + (*Triadobatrachus* + frogs)) clade. The topology of the majority-rule consensus has not changed.

13,396 trees are up to three steps longer than the minimum length. The only noteworthy change in the strict consensus is that Albanerpetontidae is no longer guaranteed to be the sister-group of the salamanders + (*Triadobatrachus* + frogs); Amphibia survives. The topology of the majority-rule consensus has changed imperceptibly.

The same holds for the 81,754 trees that are up to four steps longer than the minimum.

Because the computer had run out of memory, the search for trees up to five steps longer than the minimum length was terminated after 367,200 trees had been found. Their strict consensus does not contain Amphibia anymore (Amphibia thus has a Bremer value of 5); even salamanders and (*Triadobatrachus* + frogs) take part in the basal polytomy. The majority-rule consensus has changed imperceptibly in topology and is still almost completely resolved; Amphibia, for example, still occurs in all but four of the 367,200 trees.

The lepospondyl hypothesis is thus strongly supported by the present matrix, even though no characters or taxa were added to the matrix by Anderson *et al.* (2008), and even though the taxon sampling outside of Amphibamidae and Amphibia is poor. For further conclusions regarding errors in data matrices see Marjanović and Laurin (2008).

References

- Anderson JS. 2002. Revision of the aïstopod genus *Phlegethontia* (Tetrapoda: Lepospondyli). *J Paleontol* 76:1029–1046.
- Anderson JS, Reisz RR, Scott D, Fröbisch NB, Sumida SS. 2008. A stem batrachian from the Early Permian of Texas and the origin of frogs and salamanders. *Nature* 453:515–518.
- Andrews SM, Carroll RL. 1991. The order Adelospondyli: Carboniferous lepospondyl amphibians. *Trans R Soc Edinburgh* 82:239–275.
- Averianov AO. 1995. *Ranodon* cf. *sibiricus* (Amphibia, Caudata) from the upper Pliocene of southern Kazakhstan: the first fossil record of the family Hynobiidae. *Paläontol Z* 69:257–264.
- Baird D. The aïstopod amphibians surveyed. *Breviora Mus Comp Zool* 206:1–17.
- Berman DS, Reisz RR, Eberth DA. 1985. *Ecolsonia cutlerensis*, an Early Permian dissorophid amphibian from the Cutler Formation of north-central New Mexico. *Circ 191 N Mex Bureau Mines & Mineral Resources*:1–31.
- Berman DS, Eberth DA, Brinkman DB. 1988. *Stegotretus agyrus*[,] a new genus and species of microsaur (amphibian) from the Permo-Pennsylvanian of New Mexico. *Ann Carnegie Mus* 57:293–323.
- Bininda-Emonds ORP, Jeffrey JE, Richardson MK. 2003. Is sequence heterochrony an important evolutionary mechanism in mammals? *J Mammal Evol* 10:335–361.
- Bolt JR. 1969. Lissamphibian origins: possible protolissamphibian from the Lower Permian of Oklahoma. *Science* 166:888–891.
- Bossy KVH. 1976. Morphology, paleoecology, and evolutionary relationships of the Pennsylvanian urocordylid nectrideans (subclass Lepospondyli, class Amphibia [PhD thesis]. New Haven: Yale University. 370 + 108 p.
- Bossy KA, Milner AC. Order Nectridea MIALL 1875. In: Carroll RL, Bossy KA, Milner AC, Andrews SM, Wellstead CF. Lepospondyli. Part 1 of Wellnhofer P, editor. *Encyclopedia of Paleoherpetology*. Munich: Dr. Friedrich Pfeil. p 73–131.
- Carroll RL. 1991. *Batropetes* from the Lower Permian of Europe – a microsaur, not a reptile. *J Vertebr Paleontol* 11:229–242.
- Carroll RL, Baird D. 1968. The Carboniferous amphibian *Tuditanus* [*Eosauravus*] and the distinction between microsaur and reptiles. *Am Mus Nov* 2337:1–50. [Brackets in the original.]
- Carroll RL, Chorn J. 1995. Vertebral development in the oldest microsaur and the problem of “lepospondyl” relationships. *J Vertebr Paleontol* 15:37–56.
- Carroll RL, Gaskill P. 1971. A captorhinomorph reptile from the Lower Permian of Europe. *J Paleontol* 45:450–463.
- Carroll RL, Gaskill P. 1978. The order Microsauria. Philadelphia: American Philosophical Society.
- Carroll RL, Holmes R. 1980. The skull and jaw musculature as guides to the ancestry of salamanders. *Zool J Linn Soc* 68:1–40.
- Carroll RL, Bybee P, Tidwell WD. 1991. The oldest microsaur (Amphibia). *J Paleontol* 65:314–322.
- Clack JA. 1994. *Acanthostega gunnari*, a Devonian tetrapod from Greenland; the snout, palate and ventral parts of the braincase, with a discussion of their significance. *Medd Grøn Geosci* 31:1–24.
- Coates MI. 1996. The Devonian tetrapod *Acanthostega gunnari* Jarvik: postcranial anatomy, basal tetrapod interrelationships and patterns of skeletal evolution. *Trans R Soc Edinburgh* 87:363–421.
- Daly E. 1994. The Amphibamidae (Amphibia: Temnospondyli), with a description of a new genus from the upper Pennsylvanian of Kansas. *Univ Kans Mus Nat Hist Misc Publ*

85:1–40.

- Duellman WE, Trueb L. 1986. *Biology of Amphibians*. New York: McGraw-Hill.
- Evans SE, Milner AR, Mussett F. 1990. A discoglossid frog from the Middle Jurassic of England. *Palaeontology* 33:299–311.
- Estes R, Hoffstetter R. 1976. Les urodèles du Miocène de la Grive-Saint-Alban (Isère, France). *Bulletin du Muséum national d'Histoire naturelle de Paris* 3^{ème} série 398:297–343.
- Fracasso MA. 1983. Cranial osteology, functional morphology, systematics and paleoenvironment of *Limnoscelis paludis* Williston [PhD thesis]. New Haven: Yale University. 624 p.
- Gao K, Chen S. 2004. A new frog (Amphibia: Anura) from the Lower Cretaceous of western Liaoning, China. *Cret Res* 25:761–769.
- Gao K, Shubin NH. 2003. Earliest known crown-group salamanders. *Nature* 422:424–428.
- Gao K, Wang Y. 2001. Mesozoic anurans from Liaoning Province, China, and phylogenetic relationships of archaeobatrachian anuran clades. *J Vertebr Paleontol* 21:460–476.
- Gardner JD. 1999. The amphibian *Albanerpeton arthridion* and the Aptian–Albian biogeography of albanerpetontids. *Palaeontology* 42:529–544.
- Gardner JD. 2000. Albanerpetontid amphibians from the Upper Cretaceous (Campanian and Maastrichtian) of North America. *Geodiversitas* 22:349–388.
- Gardner JD, Evans SE, Sigogneau-Russell D. 2003. New albanerpetontid amphibians from the Early Cretaceous of Morocco and Middle Jurassic of England. *Acta Palaeont Pol* 48:301–319.
- Godfrey SJ. 1989. The postcranial skeletal anatomy of the Carboniferous tetrapod *Greererpeton burkemorani* Romer, 1969. *Phil Trans R Soc Lond B* 323:75–133.
- Good DA, Wake DB. 1992. Geographic variation and speciation in the torrent salamanders of the genus *Rhyacotriton* (Caudata: Rhyacotritonidae). *Univ Cal Publ Zool* 126:1–91.
- Holmes R. 1984. The Carboniferous amphibian *Proterogyrinus scheelei* Romer, and the early evolution of tetrapods. *Phil Trans R Soc Lond B* 306:431–527.
- Holmes RB, Carroll RL, Reisz RR. 1998. The first articulated skeleton of *Dendrerpeton acadianum* (Temnospondyli, Dendrerpetontidae) from the lower Pennsylvanian locality of Joggins, Nova Scotia, and a review of its relationships. *J Vertebr Paleontol* 18:64–79.
- Huelsenbeck JP. 1991. Tree-length distribution skewness: an indicator of phylogenetic information. *Syst Zool* 40:257–270.
- Huxley TH, Wright EP. 1867. On a collection of fossil Vertebrata, from the Jarrow Colliery, county of Kilkenny, Ireland. *Trans R Ir Acad* XXIV:351–368.
- Ivachnenko MF. 1978 [English translation 1979]. Urodélans from the Triassic and Jurassic of Soviet Central Asia. *Paleont Jour* 1978:362–368.
- Jaekel O. 1902. Ueber Ceraterpeton [sic], Diceratosaurus und Diplocaulus. *Z dt geol Ges* 54 109–134.
- Jenkins FA Jr, Walsh DM, Carroll RL. 2007. Anatomy of *Eocaecilia micropodia*, a limbed caecilian of the Early Jurassic. *Bull Mus Comp Zool* 158:285–365.
- Johanson Z, Joss J, Boisvert CA, Ericsson R, Sutija M, Ahlberg PE. 2007. Fish fingers: digit homologues in sarcopterygian fish fins. *J Exp Zool B (Mol Dev Evol)* 308:757–768.
- Laurin M. 1996. A redescription of the cranial anatomy of *Seymouria baylorensis*, the best[-] known seymouriamorph (Vertebrata: Seymouriamorpha). *PaleoBios* 17:1–16.
- Laurin M. 1998. The importance of global parsimony and historical bias in understanding tetrapod evolution. Part I. Systematics, middle ear evolution and jaw suspension. *Ann Sci Nat* 1:1–42.
- Maddison WP, Maddison DR. 2009. Mesquite: a modular system for evolutionary analysis. Version 2.6+ (build 490). <http://mesquiteproject.org>.

- Marjanović D, Laurin M. 2008. A reevaluation of the evidence supporting an unorthodox hypothesis on the origin of extant amphibians. *Contrib Zool* 77:149–199.
- McGowan GJ. 2002. Albanerpetontid amphibians from the Lower Cretaceous of Spain and Italy: a description and reconsideration of their systematics. *Zool J Linn Soc* 135:1–32.
- Milner AR. 1996. A revision of the temnospondyl amphibians from the Upper Carboniferous of Joggins, Nova Scotia. *Spec Pap Palaeontology* 52:81–103.
- Milner AR, Sequeira SEK. 1994. The temnospondyl amphibians from the Viséan of East Kirkton, West Lothian, Scotland. *Trans R Soc Edinburgh* 84:331–361.
- Olson EC. 1941. The family Trematopsidae. *J Geol* 49:149–176.
- Rage J-C, Roček Z. 1989. Redescription of *Triadobatrachus massinoti* (Piveteau, 1936)[,] an anuran amphibian from the Early Triassic. *Palaeontographica A* 206:1–16.
- Rieppel O, Grande LA. 1998. A well-preserved fossil amphiumid (Lissamphibia: Caudata) from the Eocene Green River Formation of Wyoming. *J Vertebr Paleontol* 18:700–708.
- Ruta M, Coates MI. 2007. Dates, nodes and character conflict: addressing the lissamphibian origin problem. *J Syst Palaeontol* 5:69–122.
- Sanch[í]z B. 1998. Salientia. Part 4 of Wellnhofer P, editor. *Encyclopedia of Paleoherpitology*. Munich: Dr. Friedrich Pfeil.
- Sawin HJ. 1941. The cranial anatomy of *Eryops megacephalus*. *Bull Mus Comp Zool* 88:407–463.
- Schoch RR, Fröbisch NB. 2006. Metamorphosis and neoteny: alternative pathways in an extinct amphibian clade. *Evolution* 60:1467–1475.
- Schoch RR, Rubidge BS. 2005. The amphibamid *Micropholis* from the *Lystrosaurus* Assemblage Zone of South Africa. *J Vertebr Paleontol* 25:502–522.
- Shubin NH, Jenkins FA Jr. 1995. An Early Jurassic jumping frog. *Nature* 377:49–52.
- Smithson TR. 1982. The cranial morphology of *Greererpeton burkemorani* Romer (Amphibia: Temnospondyli). *Zool J Linn Soc* 76:29–90.
- Sumida SS, Berman DS, Martens T. 1998. A new trematopid amphibian from the Lower Permian of central Germany. *Palaeontology* 41:605–629.
- Swofford DL. 2003. PAUP*: Phylogenetic Analysis Using Parsimony (*and other methods). Version 4.0b10. Sunderland, Massachusetts: Sinauer Associates.
- Vallin G, Laurin M. 2004. Cranial morphology and affinities of *Microbrachis*, and a reappraisal of the phylogeny and lifestyle of the first amphibians. *J Vertebr Paleontol* 24:56–72.
- Venczel M. 1999. Land salamanders of the family Hynobiidae from the Neogene and Quaternary of Europe. *Amphibia-Reptilia* 20:401–412.
- Venczel M, Gardner JD. 2005. The geologically youngest albanerpetontid amphibian, from the Lower Pliocene of Hungary. *Palaeontology* 48:1273–1300.
- Wellstead CF. 1982. A Lower Carboniferous aïstopod amphibian from Scotland. *Palaeontology* 25:193–208.
- Wellstead CF. 1991. Taxonomic revision of the Lysorophia, Permo-Carboniferous lepospondyl amphibians. *Bull Am Mus Nat Hist* 209:1–90.
- Wiens JJ. 2001. Character analysis in morphological phylogenetics: problems and solutions. *Syst Biol* 50:689–699.
- Wiens JJ, Bonett RM, Chippindale PT. 2005. Ontogeny discombobulates phylogeny: paedomorphosis and higher-level salamander relationships. *Syst Biol* 54:91–110.
- Wischnitzer S. 1979. Atlas and dissection guide for comparative anatomy. Vol. 1. New York: W. H. Freeman and company.

Supplementary Table 1: List of loss characters in the matrix by Anderson *et al.* (2008). All of them are retained in our version of the matrix (ESM3), although 137 is parsimony-uninformative.

- 4 Intertemporal: present (0), absent (1)
- 5 Supratemporal: present (0), absent (1)
- 10 Squamosal-tabular [contact]: absent (0), present (1), fused (2)¹
- 13 Lacrimal: present (0), absent (1)
- 18 Quadratojugal: present (0), absent (1)
- 24 Nasals: present (0), absent (1)
- 27 Internarial fontanelle: absent (0), present (1)
- 28 Septomaxilla: ossified (0), unossified (1)²
- 38 Otic notch: present (0), absent (1)
- 45 Postorbital: present (0), absent (1)
- 53 Postparietals: paired (0), fused (1), absent (2)
- 54 Parietal foramen: present (0), absent (1)
- 59 Tabular: present (0), absent (1)
- 64 Lateral line canal grooves: present (0), absent (1)
- 65 Dermal sculpturing: circular pits (0), shallow ridges and grooves (1), little to none (2)
- 81 Labyrinthine in-folding [sic]: present (0), absent (1)
- 83 Supraoccipital: absent (0), present (1)
- 89 Palatal teeth: present (0), absent (1)
- 92 Anterior [part of] palatine: short anteromedial process articulating with vomer at choana (0), long anteromedial process, more medial than lateral (1), palatine absent (2)
- 93 Vomerine teeth: present (0), absent (1)
- 96 Intervomerine rostral fenestration: absent (0), present (1)
- 97 Denticles on pterygoid: present (0), absent (1)
- 98 Teeth on pterygoid: absent (0), present (1)
- 100 Denticles on vomers: present (0), absent (1)
- 101 Denticles on palatines: present (0), absent (1)
- 102 Denticles on parasphenoid: present (0), absent (1)
- 107 Basicranial articulation: loose (0), sutured or fused (1)
- 108 Stapes: perforated stem (0), imperforate stem (1), no stem (2)³
- 114 Sphenethmoid: ossified (0), unossified (1)
- 115 Interpterygoid vacuities [sic]: narrow (“closed”) (0), wide (1), fused at midline (2)⁴
- 121 Ectopterygoid: present with fang-pit pair (0), present[,] lacking fang-pit pair (1), absent (2)⁵
- 126 Surangular: normal (0), reduced (1), absent (2)
- 128 Number of splenials: 2 (0), 1 (1), 0 (2)⁶
- 130 Meckelian fossae [sic – actually fenestrae, lacking a bony floor]: 2 or more (0), 1 (1), 0 (2)
- 136 Number of coronoids: 3 (0), 2 (1), 1 (2), 0 (3)⁷
- 137 Coronoid teeth: present (0), absent (1)
- 140 Jaw sculpture: present (0), absent (1)
- 141 Ossified hyoids: present (0), absent (1)
- 148 Trunk intercentra: present (0), absent (1)
- 156 Haemal arches: present (0), absent (1)
- 167 Atlas-axis intercentra: present (0), absent (1)
- 169 Atlas centrum: multipartite (0), single notochordal (1), single odontoid (2)⁸
- 174 Proatlantes: present (0), absent (1)
- 176 Atlas ribs: one pair (0), two pairs (1), absent (2)⁹

- 190 Interclavicle sculpture: present (0), absent (1)
 193 Cleithrum: ossified (0), unossified (2)²
 196 Supraglenoid foramen: present (0), absent (1)
 208 Number digits – manus: 5 or more (0), 4 (1), 3 (2)¹⁰
 219 Number of digits, pes: 5 or more (0), 4 or less (1)¹¹

¹ State 2, the presence of a single bone in the area normally occupied by the squamosal and the tabular, can at least as easily be interpreted as the loss of the tabular with replacement by the squamosal (or imaginably the other way around).

² These are dermal bones, so “unossified” equals “absent”.

³ State 1 refers to the loss of the stapedial foramen.

⁴ This obviously refers to the pterygoids touching each other and/or the parasphenoid throughout the length of their medial margins, in other words, the loss of the interpterygoid vacuities as seen for example in *Seymouria* and *Batrachiderpeton*.

⁵ This character refers to the loss of the fangs and the loss of the bone that carries them and therefore counts as two loss characters.

⁶ This character refers to the sequential loss of two bones and therefore counts as two loss characters. We have opted not to treat character 53 the same way, because there is evidence for the fusion of the two postparietals (left and right), but none for the fusion of the two splenials (“presplenic” and “postsplenic” in each hemimandible).

⁷ Referring to the sequential loss of three bones, this character counts as three loss characters.

⁸ States 1 and 2 both refer to the absence of the atlas intercentrum, so this character counts as a single loss character.

⁹ State 1 refers to the fusion of atlas and axis, so this character counts as a single loss character.

¹⁰ Counted as two loss characters.

¹¹ Counted as one loss character.

Supplementary Table 2: List of loss characters in the matrix by Vallin and Laurin (2004). Except for character 3, taken from the NEXUS file of Vallin and Laurin (2004), the spellings of the names of the characters and their states are taken from Laurin (1998); the six characters added later are not loss characters.

1 Lateral-line location in adults: in channels below the surface of the dermal bones (0), in grooves at the surface of the dermal bones (1), in soft tissues or absent (2)

2 Lateral-line grooves in ontogeny: present at least in the adults (may also be present in larvae) (0), present only in the larvae (1), never present (2)

3 Dermal sculpturing: ‘honeycomb’ pattern of ridges and pits (0), cosmine (1), shallow pits widely spaced from each other on an otherwise smooth surface (2), narrowly separated protuberances (3), smooth (4), broadly separated low tubera (5)¹

4 Rostrum: absent, mouth terminal (0), short, high rostrum protruding anterior to the premaxillary teeth (1), long, low (2)

5 Orbit shape: ovoid (0), confluent with a large antorbital fenestra (1), open posteriorly (2)²

10 Anterior tectal: present (0), absent (1)

15 Pineal foramen: present (0), absent (1)

16 Intertemporal: present (0), absent (1)

17 Supratemporal: present (0), absent (1)

18 Tabular: present (0), absent (1)

22 Postparietal number: two (paired) (0), single, median (1), none (2)

25 Prefrontal: present (0) absent (1)

27 Postfrontal: present (0), absent (1)

- 28 Postorbital: reaches orbit (0), excluded from orbit (1), absent (2)
- 29 Lacrimal: reaches orbit (0), excluded from orbit (1), absent (2)
- 30 Jugal: borders orbit (0), excluded from orbit (1), absent (2)
- 31 Temporal emargination: absent, area covered by opercular bones (0), present, bordered by squamosal, tabular, and (sometimes) supratemporal (1), absent, area covered by squamosal, supratemporal, and tabular (2), present, bordered by quadrate (3), present, bordered by squamosal (4)³
- 35 Quadratojugal: present (0), absent (1)
- 37 Lateral palatal tooth row: present, complete (0), incomplete (1), absent (2)
- 39 Palatal recess: median (0), divided medially (1), absent (2)
- 40 Vomerine fangs: present (0), absent (1)
- 41 Vomerine shagreen of denticles: absent (0), present (1)
- 42 Palatine: discrete (0), fused or absent (1)
- 44 Palatine fangs: present (0), absent (1)
- 45 Palatine shagreen of denticles: absent (0), present (1)
- 46 Ectopterygoid: present (0), absent (1)
- 47 Ectopterygoid fangs: present (0), absent (1)
- 48 Ectopterygoid shagreen of denticles: absent (0), present (1)
- 51 Pterygoid shagreen of denticles: present (0), absent (1)
- 54 Interpterygoid vacuity: narrow (0), broad (1), absent (2)
- 58 Parasphenoid denticles: present (0), absent (1)
- 60 Ventral cranial fissure: present (0), absent (1)
- 65 Braincase endochondral roof: ossified as a unit (0), unossified (1), composed of exoccipital (2), composed of a discrete supraoccipital (3)⁴
- 67 Basioccipital and exoccipital: indistinguishably fused in adults (0), suturally distinct throughout ontogeny (1), basioccipital never distinct (2)⁵
- 70 Preopercular: present (0), absent (1)
- 71 Subopercular: present (0), absent (1)
- 72 Epipterygoid ossification: present (0), absent (1)
- 74 Basicranial articulation: not fused, potentially mobile (0); sutured, immobile (1)
- 78 Stapedial foramen: present (0), absent (1)
- 79 Mandibular fenestrae: absent (0), small fenestrae present in splenial, postsplenial, and angular (1), large fenestra(e) present between angular, postsplenial, splenial, and prearticular (2)
- 80 Anterior coronoid: present (0), fused or absent (1)
- 81 Middle coronoid: present (0), fused or absent (1)
- 82 Posterior coronoid: present (0), fused or absent (1)
- 83 Anterior splenial: present (0), fused or absent (1)
- 84 Postsplenial: present (0), fused or absent (1)
- 85 Angular: present (0), fused or absent (1)
- 86 Surangular: present (0), fused or absent (1)
- 88 Coronoid fangs: present (0), absent (1)
- 89 Coronoid denticles: absent (0), present (1)
- 90 Dentary: dentigerous (0), edentulous
- 92 Medial mandibular tooth row: on coronoids (0), absent (1), on pseudodentary (2)
- 96 Labyrinthine infolding: present (0), absent (1)
- 98 Presacral centra (excluding atlas-axis complex): large, crescentic intercentra and small, paired, dorsal pleurocentra (0); cylindrical intercentra only (1); crescentic intercentra only (2); crescentic intercentra and cylindrical pleurocentra (3); circular intercentra and pleurocentra (4); cylindrical pleurocentra only (5)⁶
- 109 Atlantal intercentrum: present (0), absent (1)

- 115 Number of sacral vertebrae: none (0), one (1), two or more (2)
- 117 Discrete dorsal fin: present (0), absent (1)
- 118 Radials in caudal fin: present (0), absent (1)
- 120 Interclavicle: without a parasternal process (0), with a parasternal process (1), absent (2)
- 121 Clavicle: present (0), absent (1)
- 123 Cleithrum: with a dorsal expansion (0), slender, without a discrete dorsal expansion (1), with a ventral expansion (2), absent (3)
- 125 Anocleithrum: present (0), absent (1)
- 126 Lateral extrascapular: present (0), absent (1)⁷
- 127 Median extrascapular: present (0), absent (1)⁷
- 130 Humerus: present (0), absent (1)
- 141 Number of digits in manus: none (0), eight (1), five (2), four (3)⁸
- 142 Number of pelvic ossifications: one (0), three (1), two (2)⁹
- 150 Number of pedal digits: none (0), seven (1), five (2), four (3)¹⁰
- 151 Number of phalanges in second pedal digit: two (0), three (1)
- 152 Number of phalanges in third pedal digit: three (0), four (1)
- 153 Number of phalanges in fourth pedal digit: four (0), five (1), three (2)
- 154 Number of phalanges in fifth pedal digit: four (0), five (1), three (2), two (3)
- 155 Lepidotrichia in caudal fin: present (0), absent (1)

¹ State 4 refers to the loss of sculpturing.

² State 2 refers to the loss of bones such as the postorbital.

³ State 2 refers to loss of the emargination.

⁴ Under the lepospondyl hypothesis, loss of the suproccipital (from state 3 to state 1) is a lissamphibian autapomorphy; this disappears under the temnospondyl hypothesis, however. This character could thus be said to be a loss character that supports the temnospondyl hypothesis.

⁵ State 2 refers to the loss of the basioccipital: "The basioccipital of lissamphibians is never distinct and there does not appear to be a discrete ossification centre for this element (2)." (Laurin, 1998)

⁶ States 1 and 2 refer to the loss of pleurocentra, and state 5 to the loss of intercentra. Therefore this character is counted as two loss characters.

⁷ Counted as a single loss character because a functional correlation is hypothesized and no taxon is known to possess either lateral extrascapulars or the median one but not both.

⁸ Counted as three loss characters: no taxon with seven manual digits is known, but one with six (*Tulerpeton*) is, even though it is not present in the matrix.

⁹ State 2 refers to loss of ossification of the pubis.

¹⁰ Counted as three loss characters (even though the only known taxon with six pedal digits, *Tulerpeton*, is not present in the matrix).

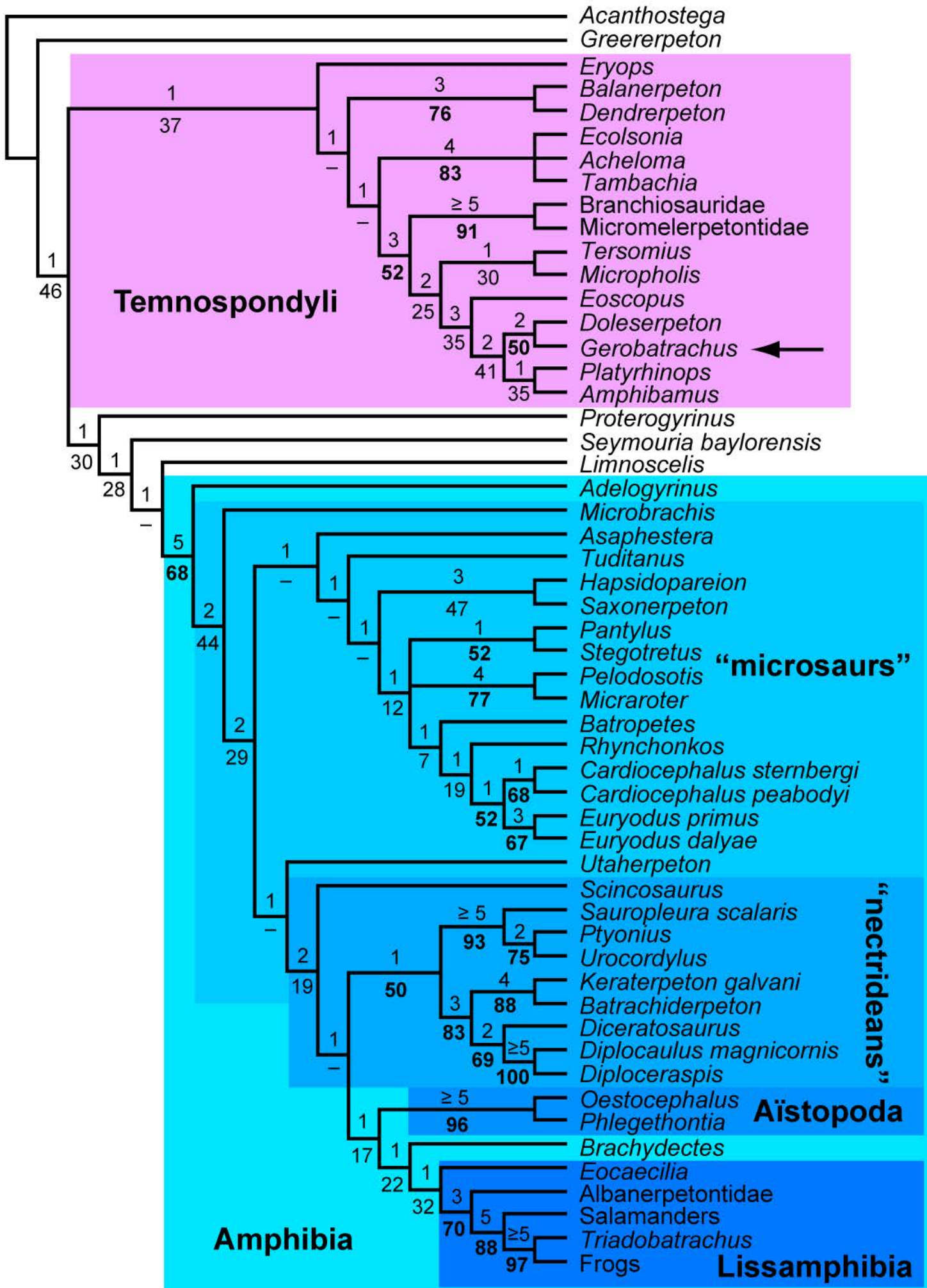
.....*****.....	78.56	39.3%
..**.....	77.15	38.6%
.....*****.....	73.01	36.5%
.....*****.....	71.65	35.8%
.....**.....	70.02	35.0%
.....**.....	69.70	34.9%
.....*.....	67.41	33.7%
.....*****.....	60.55	30.3%
.....*.....	60.19	30.1%
.....*****.....*****.....*****.....	57.17	28.6%
.....*****.....*****.....*****.....	55.51	27.8%
.....*****.....	55.29	27.6%
.....**.....**.....**.....**.....	51.79	25.9%
.....*.....**.....**.....	50.69	25.3%
.....*.....**.....	48.15	24.1%
.....*****.....	48.04	24.0%
.....*.....**.....*	47.39	23.7%
.....*****.....	46.21	23.1%
.....*****.....	45.39	22.7%
.....*****.....*****.....*****.....*****.....*****.....	44.91	22.5%
.....*****.....*.....*	44.81	22.4%
.....*****.....	44.24	22.1%
.....*.....**.....	40.50	20.3%
.....*.....*.....**.....	40.20	20.1%
.....**.....	39.81	19.9%
.....*****.....*****.....	38.59	19.3%
.....*****.....*	38.55	19.3%
.....*****.....*	38.37	19.2%
.....**.....	37.35	18.7%
.....*****.....*****.....*****.....*****.....*****.....*****.....	36.94	18.5%
.....*****.....**.....*****.....	36.29	18.1%
.....**.....*****.....	36.23	18.1%
.....*****.....*****.....	34.55	17.3%
.....*****.....**.....*****.....	33.98	17.0%
.....**.....**.....*****.....*****.....*****.....*****.....*****.....	33.13	16.6%
.....*.....*	33.05	16.5%
.....*****.....**.....*****.....	31.88	15.9%
.....**.....*.....**.....	30.42	15.2%
.....*****.....**.....	30.37	15.2%
.....*.....*	29.52	14.8%
.....*****.....*****.....*****.....*****.....*****.....*****.....	28.82	14.4%
.....*****.....*.....*****.....*****.....	28.78	14.4%
.....**.....	28.24	14.1%
.....*****.....*.....*****.....*****.....*****.....	28.16	14.1%
.....*.....*****.....	27.88	13.9%
.....*****.....*****.....*****.....*****.....*****.....*****.....	27.79	13.9%
.....*****.....	27.64	13.8%
.....*****.....*****.....	24.39	12.2%
.....**.....	24.39	12.2%
.....*****.....*****.....*****.....*****.....	24.39	12.2%
.....*****.....*.....*	24.28	12.1%
.....**.....*	24.15	12.1%
.....*.....**.....*	23.86	11.9%
.....*****.....*****.....*****.....*****.....*****.....	22.20	11.1%
.....*****.....	22.08	11.0%
.....**.....	21.73	10.9%
.....**.....*****.....	21.69	10.8%
.....**.....*.....**.....	20.51	10.3%
.....*.....**.....*	20.32	10.2%
.....*****.....	20.03	10.0%
.....*.....*.....*****.....*****.....*****.....*****.....	19.14	9.6%
.....*.....*****.....*****.....*****.....*****.....	18.89	9.4%
.....*.....*.....*	18.80	9.4%
.....*****.....*.....*	18.70	9.3%
.....**.....*****.....*****.....*****.....*****.....*****.....	18.69	9.3%
.....*****.....**.....*****.....	18.49	9.2%
.....**.....*	18.24	9.1%
.....*****.....*****.....*****.....*****.....*****.....*****.....	18.04	9.0%
.....**.....**.....**.....**.....*****.....*****.....*****.....*****.....	18.04	9.0%
.....**.....**.....	17.94	9.0%

Abbreviations: LH, lepospondyl hypothesis; PH, polyphyly hypothesis; TH, temnospondyl hypothesis; Salientia: *Triadobatrachus* and the “frogs” OTU.

Contents of the clade	Clade contradicts:			Bootstrap percentage
	LH	PH	TH	
all “lepospondyls”, Lissamphibia	no	yes	yes	67.5
all “lepospondyls” except <i>Adelogyrinus</i> , Lissamphibia	no	yes	yes	43.6
Temnospondyli	no	yes	yes	36.5
all “lepospondyls”, Lissamphibia, <i>Limnoscelis</i>	no	yes	yes	35.8
all “lepospondyls”, Lissamphibia, <i>Limnoscelis</i> , <i>Seymouria</i> , <i>Proterogyrinus</i>	no	yes	yes	30.3
all “lepospondyls” except <i>Adelogyrinus</i> and <i>Microbrachis</i> , Lissamphibia	no	yes	yes	28.6
all “lepospondyls”, Lissamphibia, <i>Limnoscelis</i> , <i>Seymouria</i>	no	yes	yes	27.8
all temnospondyls except <i>Eryops</i> , <i>Ecolsonia</i> , <i>Acheloma</i> and <i>Tambachia</i>	no	yes	yes	25.9
<i>Tersomius</i> , <i>Doleserpeton</i> , <i>Micropholis</i> , <i>Eoscopus</i> , <i>Platyrrhinops</i> , <i>Amphibamus</i> , <i>Gerobatrachus</i> (thus same as above except <i>Balanerpeton</i> , <i>Dendrerpeton</i> , Branchiosauridae and Micromelerpetontidae)	no	rather yes	rather yes	25.3
<i>Doleserpeton</i> , <i>Amphibamus</i> , <i>Gerobatrachus</i>	no	rather no	rather no	24.1
all temnospondyls except <i>Balanerpeton</i> , <i>Dendrerpeton</i> and <i>Eryops</i>	no	yes	yes	23.1
Lissamphibia	no	yes	no	22.7
<i>Rhynchonkos</i>, <i>Eocaecilia</i>	yes	no	yes	22.4
<i>Brachydectes</i> , Lissamphibia	no	yes	yes	22.1
<i>Tersomius</i> , <i>Doleserpeton</i> , <i>Eoscopus</i> , <i>Platyrrhinops</i> , <i>Amphibamus</i> , <i>Gerobatrachus</i>	no	rather yes	rather yes	20.1
all “nectrideans”, Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	19.3
all “nectrideans”, Aïstopoda, Albanerpetontidae, salamanders, Salientia	yes	yes	yes	18.1
Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	17.3
all “nectrideans”, Aïstopoda, <i>Brachydectes</i> , Albanerpetontidae, salamanders, Salientia	yes	yes	yes	17.0
all temnospondyls except <i>Balanerpeton</i> , <i>Dendrerpeton</i> , <i>Eryops</i> , <i>Ecolsonia</i> , <i>Acheloma</i> and <i>Tambachia</i> , all “lepospondyls”, Lissamphibia	no	no ¹	rather yes	16.6
Aïstopoda, Albanerpetontidae, salamanders, Salientia	yes	yes	yes	15.9
all “lepospondyls” except <i>Microbrachis</i> , <i>Utaherpeton</i> and <i>Adelogyrinus</i> , Lissamphibia	no	yes	yes	14.4
same as above except <i>Tuditanus</i> , <i>Asaphestera</i> , <i>Hapsidopareion</i> and <i>Saxonerpeton</i>	no	yes	yes	14.1
<i>Gerobatrachus</i>, salamanders, Salientia	yes	no	no	13.9
all temnospondyls except <i>Balanerpeton</i> , <i>Dendrerpeton</i> and <i>Eryops</i> , all “lepospondyls”, Lissamphibia	no	no ¹	rather yes	13.9
all “nectrideans” except <i>Scincosaurus</i> , Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	12.2
all “lepospondyls” except <i>Utaherpeton</i> and	no	yes	yes	12.2

<i>Adelogyrinus</i> , Lissamphibia				
<i>Doleserpeton</i> , <i>Platyrhinops</i> , <i>Gerobatrachus</i>	no	rather	rather	11.9
		no	no	
all “lepospondyls” except <i>Utaherpeton</i> , Lissamphibia	no	yes	yes	11.1
<i>Greererpeton</i> , Temnospondyli	no	yes	yes	10.8
<i>Rhynchonkos</i> , <i>Batropetes</i> , all “nectrideans”, Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	9.6
same as above except <i>Rhynchonkos</i>	no	yes	yes	9.4
<i>Hapsidopareion</i> , <i>Brachydectes</i>	no	no	no	9.4
Gymnarthridae, <i>Rhynchonkos</i>, <i>Eocaecilia</i>	yes	no	yes	9.3
all temnospondyls except <i>Eryops</i> , all “lepospondyls”, Lissamphibia	no	no ¹	rather	9.3
			no	
Aïstopoda, <i>Brachydectes</i> , Lissamphibia except <i>Eocaecilia</i>	yes	yes	yes	9.2
all “lepospondyls”, <i>Eocaecilia</i>, Albanerpetontidae	yes	no	yes	9.0
all temnospondyls except <i>Eryops</i> , <i>Ecolsonia</i> , <i>Acheloma</i> and <i>Tambachia</i> , all “lepospondyls”, Lissamphibia	no	no ¹	rather	9.0
			no	
<i>Platyrhinops</i> , <i>Amphibamus</i> , <i>Gerobatrachus</i>	no	rather	rather	9.0
		no	no	
Lissamphibia without Albanerpetontidae	no	yes	no	8.7
<i>Utaherpeton</i> , all “nectrideans”, Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	8.4
Pantylidae, Gymnarthridae, Ostodolepididae, <i>Rhynchonkos</i>, <i>Batropetes</i>, <i>Eocaecilia</i>	yes	no	yes	8.3
all “lepospondyls” except <i>Tuditonus</i> , <i>Asaphestera</i> , <i>Microbrachis</i> and <i>Adelogyrinus</i> , Lissamphibia	no	yes	yes	8.1
Albanerpetontidae, salamanders	no	no	no	7.9
<i>Batropetes</i> , <i>Utaherpeton</i> , all “nectrideans”, Aïstopoda, <i>Brachydectes</i> , Lissamphibia	no	yes	yes	7.4
<i>Doleserpeton</i> , <i>Platyrhinops</i> , <i>Amphibamus</i> , <i>Gerobatrachus</i> , salamanders, Salientia	yes	no	yes	7.4
<i>Brachydectes</i> , <i>Eocaecilia</i>	yes	no	yes	7.3
Temnospondyli, salamanders, Salientia	yes	no	yes	7.3
all “lepospondyls” except <i>Adelogyrinus</i> , <i>Eocaecilia</i> , Albanerpetontidae	yes	no	yes	7.1
<i>Doleserpeton</i> , <i>Gerobatrachus</i> , salamanders, Salientia	yes	no	yes	7.0
<i>Batropetes</i> , <i>Eocaecilia</i>	yes	no	yes	6.4
<i>Gerobatrachus</i> , Lissamphibia except <i>Eocaecilia</i>	yes	no	yes	6.3
<i>Brachydectes</i> , Lissamphibia except <i>Eocaecilia</i>	yes	yes	yes	6.1
<i>Rhynchonkos</i> , <i>Batropetes</i> , <i>Eocaecilia</i>	yes	no	yes	6.1
Pantylidae, Gymnarthridae, <i>Rhynchonkos</i> , <i>Batropetes</i> , <i>Eocaecilia</i>	yes	no	yes	5.9
all “lepospondyls” except <i>Utaherpeton</i> and <i>Adelogyrinus</i> , <i>Eocaecilia</i> , Albanerpetontidae	yes	no	yes	5.2
same as above except Albanerpetontidae and all “nectrideans” other than <i>Ptyonius</i> (!)	yes	no	yes	5.1

¹ This only holds if highly unconventional arrangements like those found by McGowan (2002), where “lepospondyls” and lissamphibians are found inside Temnospondyli, are counted as acceptable within the polyphyly hypothesis.



#NEXUS

[written Sat Jun 13 21:47:32 CEST 2009 by Mesquite version 2.6+ (build 494) at hdt203ml1.mnhn.fr/10.8.64.235]

BEGIN TAXA;

TITLE Untitled_Taxa_Block;

DIMENSIONS NTAX=54;

TAXLABELS

Acanthostega Proterogyrinus Greererpeton Seymouria_baylorensis Limnoscelis Balanerpeton
Dendrerpeton Eryops Branchiosauridae Micromelerpetontidae Tersomius Ecolsonia Acheloma Doleserpeton
Micropholis Eoscopus Tambachia Platyrhinops Amphibamus Gerobatrachus Tuditanus Asaphestera Hapsidopareion
Saxonerpeton Pantylus Cardiocephalus_sternbergi Cardiocephalus_peabodyi Euryodus_primus Euryodus_dalyae
Pelodosotis Micraroter Rhynchonkos Microbrachis Batropetes Utaherpeton Stegotretus Sauropleura_scalaris
Ptyonius Urocordylus Keraterpeton_galvani Batrachiderpeton Diceratosaurus Diplocaulus_magnicornis
Diploceraspis Scincosaurus Adelogyrinus Oestocephalus Phlegethontia Brachydictes Eocaecilia Albanerpetontidae
Salamanders Triadobatrachus Frogs

;

END;

BEGIN CHARACTERS;

TITLE morphology;

DIMENSIONS NCHAR=219;

FORMAT DATATYPE = STANDARD GAP = - MISSING = ? SYMBOLS = " 0 1 2 3";

CHARSTATELABELS

1 Basal_Skull_Length / '>70mm' '50-70mm' '30-50mm' '<30mm', 2 'Skull:trunk' / '>=0.45' '0.30-0.45'
'0.20-0.29' '<0.20', 3 Skull_proportions / longer_than_wide wider_than_long, 4 Intertemporal / present absent, 5
Supratemporal / present absent, 6 ST_exposure_on_occiput / absent present, 7 'Tabular-postorbital' / absent
Present, 8 'T-PF' / absent present, 9 Postfrontal_shape / broadly_quadrangular falciform, 10 'Squamosal-Tabular' /
absent present fused, 11 'Squamosal-Temporal area' / weakly_overlapping sutural, 12 'Lacrimal-prefrontal suture' /
simple_butt_joint interdigitating prefrontal_broadly_underplates_lacrimal, 13 Lacrimal / present absent, 14
L_to_naris / present absent, 15 L_to_orbit / absent present, 16 lacrimal_orbital_processes /
dorsal_and_ventral_present only_ventral_present neither_present, 17 'lacrimal-jugal contact' / present absent, 18
Quadratojugal / present absent, 19 'Quadratojugal-Jugal contact' / present absent, 20 'Quadratojugal-Maxillary
contact' / present absent, 21 Frontals / paired_along_midline fused, 22 Prefrontal_and_postfrontal / 'meet,
excluding frontal from orbit' fail_to_meet, 23 Anterior_laterally_flaring_frontals / absent present, 24 Nasals /
present absent, 25 Narial_flange / absent present, 26 Dorsal_processes_of_premax / 'broad, low, indistinct dorsal
process' 'alary process (broad, vaguely triangular process)' 'moderately high, vaguely rectangular process, or acutely
triangular linked directly to base' 'narrow, long dorsal process, along the midline or parasagittal', 27
Internarial_fontanelle / absent present, 28 Septomaxilla / ossified unossified, 29 Prefrontal_into_external_naris /
distant_from near present, 30 external_naris_in_dorsal_view / exposed not_exposed, 31 External_naris_shape /
circular 'posteriorly_extended,_along_L-PF_suture' 'posteriorly_extended,_excavation_of_L_only', 32
dorsal_exposure_of_premax / broad narrow none, 33 Dorsal_shape_of_skull / triangular diamond rounded, 34
Posterior_skull_margin / undulating. concave straight convex, 35 snout / blunt pointed, 36 snout / short long, 37 'Q
-internal flange of Sq' / absent present, 38 otic_notch / present absent, 39 Large_otic_notch_approaching_orbit /
absent intermediate close, 40 otic_notch / open_posteriorly closed_posteriorly, 41
semilunar_flange_of_supratemporal / absent present, 42 supratympanic_flange / absent 'present_' 'trematopid''
'present_'dissorophid'', 43 supratympanic_shelf / absent present, 44 raised_orbital_rim / absent present, 45
Postorbital / present absent, 46 'J-PO interfingered processes' / absent present, 47 PO_in_orbital_margin / present
absent, 48 shape_of_postorbital / irregular_trapezoid 'triangular,_apex_caudal', 49 Palpebral_ossifications / absent
mosaic_of_bone_plates_present_in_orbit, 50 'Parietal-postorbital contact' / absent present, 51 'parietal-squamosal
contact' / absent present, 52 'parietal-tabular contact' / absent present, 53 Postparietals / paired fused absent, 54
parietal_foramen / present absent, 55 postparietals / moderate large, 56 'postparietal-squamosal contact' / absent
present, 57 Postparietal_length / 'large,_quadrangular' 'abbreviated_anteroposteriorly,_elongate_lateral_rectangle',
58 'squamosal-jugal contact' / present absent, 59 Tabular / present absent, 60
Posterolateral_projection_from_lateral_margin_of_tabular_above_squamosal_embayment / absent present, 61
'Tabular horns' / absent present, 62 Tabular_horns / parallel_or_slightly_divergent widely_divergent, 63

Sq_forms_base_of_tabular_horn / absent present, 64 Lateral_line_canal_grooves / present absent, 65 Dermal_sculpturing / circular_pits shallow_ridges_and_grooves little_to_none, 66 premaxilla_anterior_margin / vertical overturned, 67 Maxilla_into_orbit / no yes, 68 Maxilla_into_external_naris / present absent, 69 Maxilla_entire_ventral_naris / absent present, 70 maxilla / longer_than_palatine shorter_than_palatine, 71 marginal_teeth / vertical turned_medially, 72 marginal_teeth_largest_anterior / absent present, 73 marginal_teeth / pointed_pegs blunt_pegs large_cones, 74 Number_of_premax_teeth / '>=10' '5-9' '<5', 75 Number_of_max_teeth / '>=30' '20-29' '<20', 76 teeth_laterally_compressed / no yes, 77 'Enlarged teeth mid-toothrow' / absent present, 78 teeth / simple_points multiple_cusps, 79 Multiple_Cusp_Orientation / 'Labio-lingual' 'antero-posterior', 80 Enamel_fluting / absent present, 81 'labyrinthine in-folding' / present absent, 82 occipital_profile / low_and_wide high_and_wide high_and_narrow, 83 supraoccipital / absent present, 84 Occipital_condyle / concave convex, 85 occipital_condyle / 'single,_with_basioccipital' double, 86 jugular_foramen / between_opisthotic_and_exoccipit through_exoccipital, 87 jaw_articulation / posterior_to_occiput even_with_occiput anterior_to_occiput 'far_anterior_(>20%_BSL)', 88 Internal_nares / widely_separated narrowly_separated, 89 Palatine_fangs / present absent, 90 Palatine_teeth / absent multiple_in_rows multiple_random, 91 LEP / absent present, 92 Anterior_palatine / short_anteromedial_process_articulating_with_vomer_at_choana 'long_anteromedial_process,_more_medial_than_lateral' palatine_absent, 93 Vomerine_fangs / present absent, 94 Vomer_teeth / absent 'present, in a single row' 'present, as a field', 95 Intervomerine_depression / absent present, 96 Intervomerine_rostral_fenestration / absent present, 97 denticles_on_pterygoid / present absent, 98 teeth_on_pterygoid / absent present, 99 tooth_pedicely / absent present, 100 denticles_on_vomers / present absent, 101 denticles_on_palatines / present absent, 102 denticles_on_parasphenoid / present absent, 103 DELETED_was_Palatal_teeth / larger_than_marginals equal_to_marginals smaller_than_marginals, 104 parasphenoid / medial_of_stapes under_footplate_of_stapes, 105 Parasphenoid_basal_plate / 'roughly_quadrangular,_basipterygoid_articulations_narrowly_spaced' 'about as broad as long, articulations moderately distant' 'rectangular_laterally,_anteroposteriorly_narrow,_basipterygoid_articulations_distant', 106 cultriform_process_contact_with_vomer / narrow broad, 107 basicranial_articulation / loose sutured_or_fused, 108 stapes / perforated_stem imperforate_stem no_stem, 109 Stapes_orientation / 'lateral,_towards_quadrate' 'dorsal,_towards_squamosal_embayment,_elongate_columella', 110 footplate_of_stapes / oval round palmate, 111 dorsal_process_of_stapes / absent present, 112 "'accessory ossicle'" / absent present, 113 pleurosphenoid / unossified ossified, 114 sphenethmoid / ossified unossified, 115 interpterygoid_vacuities / wide 'narrow_("closed")' fused_at_midline, 116 pterygoids_contact_anteriorly / present absent, 117 'pterygoid-exoccipital contact' / absent present, 118 'Pterygoid-palatine suture' / present absent, 119 'Pterygoid-vomer contact' / present absent, 120 'lat process of pt into subtemp (transverse flange)' / absent present, 121 ectopterygoid / 'present_with_fang-pit_pair' 'present_lacking_fang-pit_pair' absent, 122 'Ectopterygoid-palatine width' / wider_than_maxilla narrower_than_maxilla, 123 "'pharyngeobranchial pouches'" / absent present, 124 dentary / long short, 125 dentary_forms_coronoid_process / absent present, 126 surangular / normal reduced absent, 127 angular / narrow deep, 128 number_of_sphenals / 2 1 0, 129 splenal_exposed_laterally / present absent, 130 meckelian_fossae / 2_or_more 1 0, 131 'Ventral border of Meckel's foss' / splenal angular, 132 retroarticular_process / absent present, 133 retroarticular_process / straight hooked, 134 articulation_to_tooth_row / above equal below, 135 'angular extends to (lat view)' / posterior_tooth_row middle_of_tooth_row, 136 number_of_coronoids / 3 2 1 0, 137 coronoid_teeth / present absent, 138 coronoid_teeth / larger_than_marginals equal_to_marginals smaller_than_marginals, 139 symphysis / dentary_and_sphenal dentary_alone, 140 jaw_sculpture / present absent, 141 ossified_hyoids / present absent, 142 Gill_Osteoderms / absent 'present,_noninterdigitating' 'toothed,_interdigitating_rakers', 143 parahyoid / absent present, 144 number_of_accessory_articulation / 0 1 2_or_more, 145 number_of_presacrals / '>35' '25-35' '20-24' '<20', 146 DELETED_was_vertebral_development / 'arches,_then_centra' centra_and_arches_simultaneously, 147 caudal_processes_bt看_depression / absent present, 148 trunk_intercentra / present absent, 149 trunk_neural_arch_to_centrum / loosely_articulated sutured fused, 150 base_of_neural_spine / equal_to_or_wider_than_haemal smaller_than_haemal_spine, 151 height_of_neural_spines / even alternating, 152 dorsal_neural_spine / narrow_and_smooth laterally_broad_and_sculpted, 153 'neural spine shape (lat)' / 'ant-post_sides_parallel_(rect)' 'non-parallel_(triangular)', 154 Neural_spine_lateral_surface / smooth crenulated, 155 Pleurocentra / paired_rhachitomous closely_approaching_ventrally 'fused,_dominant_weight_bearing_element', 156 haemal_arches / present absent, 157 haemal_arches / 'loosely_articulated_(intercentra)' 'fused_to_mid-length_of_centrum', 158 haemal_arches / longer_than_or_equal_to_neurals shorter_than_neurals, 159 haemal_accessory_articulations / none one two, 160 haemal_arch_shape / 'non-parallel_(triangular)' 'parallel_(rectangular)', 161 tail / tapers deep_with_sudden_end, 162 Tail / 'elongate,_equal_to_or_exceeding_trunk_and_skull_length' 'forshortened,_markedly_shorter_than_trunk', 163 DELETED_was_trunk_arches / paired fused, 164 spinal_nerve_foramina / absent present, 165

Euryodus_primus 2?011?11011100100000000?2001102121001??0001000000010110000?0???
1000010222010?1100011201100110000000121100?0???0?1000010000111112?11203??1?1000???02?
00102???????1000?????????????0??1111?1?????1??0(0 1)2101?100?102010?0??
Euryodus_dalyae 2?011?110111001000010000?(0 2)001102021001??0001000000010010000?0??
121000010222010?010001020110?????00000021100000010?1000010000111112?0?203??111000???02?
00102???????10000122000100??0?0?????????????????????1????0?????????
Pelodosotis 13011?110111001000110000?2001000001001??00000000000010110010?0??12101?
010012010?01210102?1100????000??0211001021100110000?000{0 1}0111111020?021100000??01?
10102???????10001121?101100100?0?0111?11100011?10220021??00201000??
Micraroter 13011?110111001110110000?2000000000001??00010000100101(0 1)0010?0??
121100010112010?0101011201100120000000020100001010?100000000010101??1020??1100001?
00101010200001001000012200011000002?????????1000?0?1??0??????102110?????
Rhynchonkos 33011?110111001100010000?2000102131001??0001000010010010000?0??
121000010012000?010111131110011001001101110000001101100010000100001110101011110000??02?
00102???????10000122?201121020??3??????????0??122012100{0 1}1020100010?
Microbrachis 32011?100111011100010000?2010000020001??0000000011(0 1)10000100?0??
010000000(0 1)11000?010?01?201000??00000000?
110000000011100010000101001110000021001000101100010200000011010112121011000000020110?
00010000?102001100?21011110??00
Batropetes 32011?10111100110??01000?012102100001??00000000011120??00?0??12101??
10012001101111??3??00??100110?120010000001?0????001??11??101????1110003?
0120001020010000100011220?011000000001110?11??0102002201210?10020110?110
Utaherpeton 310????00??100101??0000??0?20000?0001??0000000?01??0??0?0??12010??001?
1000?01????{1 2}??0????000?0??0??????????????????00??10??0?21??110?01001200000200000??(0 1)0?
01????????0000{0 1}?00?000(0 1)??????102201011?11?1?101??0
Stegotretus 3?011?1101110010000{0 1}0000?2001101020001??000100000001001000000??
111001010222010?0101010210200120001000011100?????0010000020001?001110?00?0(0 1)0?10?0??11?
00102???????10001121?????????0??1111??????011012200210??2?00100?????
Sauroplesura_scalaris 210100000100001010001000?0012000001101??0000010101?10000000?0??1101(0
1)0001012000?011??1?000100????01011120000?????001?000?1100000100010003??1000011?01200011201010?
01010111222111210000010?0010011000??0100110000100211100?10
Ptyonius 110101000101001000010000?0012000001001??0000010101?10000000?0??
110000000012000?011??1?001100110010001120100?????0100000?0100001?1??1101??1000022?112100(0
1)120102100101011222211211000020?0000001000??10001001?12021111??1
Urocordylus 22010??0?000010000?????????0?00?1001?????00100?1??????0??0??1100??010??
000?01?????00110?????000?1?2??0?0??0010?0??1000?0?0??1100??100??22?112100112011(0
1)1010101112221?210000030?000?000000??010201111?01021011??0
Keraterpeton_galvani 32011?000111001100010000?00120000100?1??00000000?1110010000?
101010000?00022000?010?11?20110?????0000111??0??????0020??????0100010111111??0000013?
012000112011000010101?1222111210000030?000?010012??10001011?{0 1}2021111??0
Batrachiderpeton ??011?000111001100010000?0011000010011??0000000001110010000?
110010100100012000?011?11?20110011000000111??01?????0020000011{0 1}10001011111120(1 2)0000013?
012000(0 1)12011000010101112221122?0?00?10?000?010012?????????????20?????????
Diceratosaurus ?1011?000111001110010000?2012000010011??00000000011100100000100010100?
00????00?010?11?2?110??????0?????01?????00101?????000?11011102?2011000013?
012001012010000010101?1222111210000030?000?010012??210001011?{0 1}2021111??0
Diplocaulus_magnicornis 01011?000111000?1001110102011100010011??
00000010011100100000111010000100012000?010011{0 1}301100110010011110001?????
000110101010020010(0 1)0?202010000013?0120010120100000101111222112200100130000010???12??
200001101?120??111??0
Diploceraspis 0?111?00011?1??0001110102012000010001??
00000010011100100000111010000100012000?0100110301100??0010011110101?????
00011010001010110111020201000002??012001012010100?1011112221122??0?1?0?00010??
12????????????????????
Scincosaurus 33011?110111001100010000?20120000{0 2}0011??00000000000120??0000??
010100000012000?01??11?301?00??0010?101?0101?????00110000100?0??????????01?000012?
012100012011110010001?1222?11210000?20?000?000002??0022001000010201000?10

Adelogyrinus 2?011?10021?1????0000000?20110000000000?00000010?111000000{0 1}10??
0101???000?0000?01000012???0?????0?0001?0?00?????01?0?000000?00??100(0 1)???100000?010?
00102???????0001020220112??20???0?00000???0???2?????????????????????
Oestocephalus 230100?0010100111011000002012101030001?00001???0??0001001000??
120100100010000?01200000011?????10??101?1?0???000100000100000001110?003??1110010?012?
000021????00111010122200121?21??0?????????00?????????????????????????
Phlegethontia 23011???0?1?1???????110002012000031001???0001???1???21???01?0??120101?
00022000?012000?2?????????10???1?0??020000001?0??010000??2?2?0?0?3??11100101012?000021???
00111010{1 2}22200121?21??0?????????00?????????????????????????????
Brachydectes 33011?????1110010?1?0-000{1 2}012000?20001??00001???0?1101110?000??
120100000012000?012101031100011001001111110?10100101100002001100111110203??1100000?
011000102000010100101121200112100000311000101100010011001(0 1)11?12021110?00
Eocaecilia 33011?0011?1????0000000?2012101101001?00011???0?110111000?0??
1201000000000010010011?3011001100001110111000000?0110011{1 2}000???2???10103??1110000?002?00?
02?????0111010122?201121020013?????????????????01200121002?0?011001?1
Albanerpetontidae ??011?????01001011??1-00?20?2000120001??00001???1?0?21???01?0??110100?
0001(1 2)1011011011?2???0??????0????????????????????????????????002?2?1110213??01?000(1 2)?012?00002?????
011001122220122002001??????????1???01???1??00??0??10?0?0?
Salamanders ?2(0 1)11?????000010?0110-00?(1 3)1000002{2 3}0001??00001???0?1?21????1?0??
120100?000(0 1)00010010011?(2 3)01(1 2)021{1 2}011011?110110(0 1)0{0 1}00?1010?102?0002?2?2?0?10{2 3}
{0 1}1010(1 2)003(0 1)0120000020100100111011222201120020013??????????1??111110001001120211000{0
1}00
Triadobatrachus 31111???1?????????0?0-00?????????22???020?0101???0?0?20????100??12?
1????????????????0011?0?10?1???10???1?02?01?0001001???02?0?????????0?0????????00103?012?00?021???
011010?1???20?1?002001?????????00??0110001100?(1 2)12?1001?0?
Frogs ?1111?????0?1???1??{0 1}-00?310?0002200002000101???0?0?20????1?0??120100?
000000010010011??0100111011011?1102(0 1)011{0 1}00?001011{0 1}2?0002?{1 2}12?0?113??01001030012?
000021?????1101011122201120020013?????????001100111001000101211001100

;
END;
BEGIN TREES;
Title Results;
LINK Taxa = Untitled_Taxa_Block;
TRANSLATE
1 Acanthostega,
2 Proterogyrinus,
3 Greererpeton,
4 Seymouria_baylorensis,
5 Limnoscelis,
6 Balanerpeton,
7 Dendrerpeton,
8 Eryops,
9 Branchiosauridae,
10 Micromelerpetontidae,
11 Tersomius,
12 Ecolsonia,
13 Acheloma,
14 Doleserpeton,
15 Micropholis,
16 Eoscopus,
17 Tambachia,
18 Platyrrhinops,
19 Amphibamus,
20 Gerobatrachus,
21 Tuditanus,

22 *Asaphestera*,
 23 *Hapsidopareion*,
 24 *Saxonerpeton*,
 25 *Pantylus*,
 26 *Cardiocephalus_sternbergi*,
 27 *Cardiocephalus_peabodyi*,
 28 *Euryodus_primus*,
 29 *Euryodus_dalyae*,
 30 *Pelodosotis*,
 31 *Micraroter*,
 32 *Rhynchonkos*,
 33 *Microbrachis*,
 34 *Batropetes*,
 35 *Utaherpeton*,
 36 *Stegotretus*,
 37 *Sauropleura_scalaris*,
 38 *Ptyonius*,
 39 *Urocordylus*,
 40 *Keraterpeton_galvani*,
 41 *Batrachiderpeton*,
 42 *Diceratosaurus*,
 43 *Diplocaulus_magnicornis*,
 44 *Diploceraspis*,
 45 *Scincosaurus*,
 46 *Adelogyrinus*,
 47 *Oestocephalus*,
 48 *Phlegethontia*,
 49 *Brachydectes*,
 50 *Eocaecilia*,
 51 *Albanerpetontidae*,
 52 *Salamanders*,
 53 *Triadobatrachus*,
 54 *Frogs*;

TREE Strict_consensus_of_MPs = (1,(3,((8,((6,7),(12,13,17)Trematopidae,(9,10),(11,15),(16,((14,20),(18,19))))))Dissorophoidea)Temnospondyli,(2,(4,(5,(46,(33,((22,(21,((23,24)Hapsidopareiontidae,(25,36)Pantylidae,(30,31)Ostodolepididae,(34,(32,((26,27),(28,29))Gymnarthridae)))Recumbirostra))),35,(45,(((37,(38,39))Urocordylidae,((40,41),(42,(43,44)))Diplocaulidae,((47,48)Aïstopoda,(49,(50,(51,(52,(53,54)Salientia)Batrachia)Parotoïdia)Lissamphibia))))))Amphibia)Tetrapoda)))));

TREE MP_1_of_4 = (1,(3,((8,((6,7),(17,(12,13))Trematopidae,(9,10),(11,15),(16,((14,20),(18,19))))))),(2,(4,(5,(46,(33,((22,(21,((23,24),((25,36),(30,31)),(34,(32,((26,27),(28,29))))))Recumbirostra))),35,(45,(((37,(38,39)),((40,41),(42,(43,44))),((47,48),(49,(50,(51,(52,(53,54)))))))))))));

TREE MP_2_of_4 = (1,(3,((8,((6,7),(17,(12,13))Trematopidae,(9,10),(11,15),(16,((14,20),(18,19))))))),(2,(4,(5,(46,(33,((22,(21,((23,24),((30,31),((25,36),(34,(32,((26,27),(28,29))))))Recumbirostra))),35,(45,(((37,(38,39)),((40,41),(42,(43,44))),((47,48),(49,(50,(51,(52,(53,54)))))))))))));

TREE MP_3_of_4 = (1,(3,((8,((6,7),(12,(13,17))Trematopidae,(9,10),(11,15),(16,((14,20),(18,19))))))),(2,(4,(5,(46,(33,((22,(21,((23,24),((25,36),(30,31)),(34,(32,((26,27),(28,29))))))Recumbirostra))),35,(45,(((37,(38,39)),((40,41),(42,(43,44))),((47,48),(49,(50,(51,(52,(53,54)))))))))))));

TREE MP_4_of_4 = (1,(3,((8,((6,7),(12,(13,17))Trematopidae,(9,10),(11,15),(16,((14,20),(18,19))))))),(2,(4,(5,(46,(33,((22,(21,((23,24),((30,31),((25,36),(34,(32,((26,27),(28,29))))))Recumbirostra))),35,(45,(((37,(38,39)),((40,41),(42,(43,44))),((47,48),(49,(50,(51,(52,(53,54)))))))))))));

TREE Strict_consensus_of_trees_up_to_1_step_longer = (1,(2,3,4,5,8,(6,7),(12,13,17)Trematopidae,(9,10),(11,15,(16,(18,19,(14,20))))),(46,(33,(21,22,25,26,27,32,34,35,36,(23,24)Hapsidopareiontidae,(28,29),(30,31)Ostodolepididae,(45,49,50,(47,48)Aïstopoda,(37,(38,39))Urocordylidae,((40,41),(42,(43,44)))Diplocaulidae,(51,(52,(53,54)Salientia)Batrachia)Parotoïdia)))Amphibia)Tetrapoda);

TREE Strict_consensus_of_trees_up_to_2_steps_longer = (1,(2,3,4,5,(6,7),8,(12,13,17)Trematopidae,(9,10),11,15,(14,16,18,19,20)),(46,21,22,(23,24)Hapsidopareiontidae,25,26,27,(28,29),(30,31)Ostodolepididae,32,33,34,35,36,(37,38,39)Urocordylidae,((40,41),42,(43,44))Diplocaulidae,45,

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(47,48)Aistopoda,49,50,(51,(52,(53,54)Salientia)Batrachia)Parotoidia)Amphibia)Tetrapoda);
TREE Strict_consensus_of_trees_up_to_3_steps_longer = (1,(2,3,4,5,6,7,8,(9,10),11,
(12,13,17)Trematopidae,14,15,16,18,19,20,(46,21,22,23,24,25,26,27,28,29,(30,31)Ostodolepididae,32,33,34,35,36,
(37,38,39)Urocordylidae,(40,41),42,(43,44),45,(47,48)Aistopoda,49,50,51,(52,
(53,54)Salientia)Batrachia)Amphibia)Tetrapoda);
TREE Strict_consensus_of_trees_up_to_4_steps_longer = (1,(2,3,4,5,6,7,8,
(9,10),11,12,13,14,15,16,17,18,19,20,(46,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,
(37,38,39)Urocordylidae,40,41,42,(43,44),45,(47,48)Aistopoda,49,50,51,(52,
(53,54)Salientia)Batrachia)Amphibia)Tetrapoda);
TREE Strict_consensus_of_trees_up_to_5_steps_longer = (1,(2,3,4,5,6,7,8,
(9,10),11,12,13,14,15,16,17,18,19,20,21,22,23,24,25,26,27,28,29,30,31,32,33,34,35,36,
(37,38,39)Urocordylidae,40,41,42,(43,44),45,46,(47,48)Aistopoda,49,50,51,52,(53,54)Salientia)Tetrapoda);
TREE [!Percentage not marked if 100. Four of the 367200 trees do not contain Amphibia.] 'Majority-rule
consensus of trees up to 5 steps longer' = (1,(3,((8,((6,7)99,((17,(12,13)61)Trematopidae,((9,10),(11,15,(16,
((14,20)91,(18,19)68)98)97)89)99)69)59)94_Temnospondyli,(2,(4,(5,(46,(33,((21,22,((23,24)99,((25,36)96,(30,31),
(34,(32,((26,27)78,(28,29)97)89_Gymnarthridae)66)56)59_Recumbirostra)63)62,35,(45,(((37,
(38,39)98)Urocordylidae,((40,41),(42,(43,44))89)99_Diplocaulidae)54,((47,48)Aistopoda,(49,(50,(51,(52,
(53,54)Salientia)Batrachia)99_Parotoidia)66_Lissamphibia)71)66)66)93)91)91)Amphibia)60_Tetrapoda)77)83)76))
;
TREE Bootstrap_tree = (1:0.0,(3:100.0,(((8:50.0,(17:100.0,
(12:100.0,13:100.0)61:67.207458)83_Trematopidae:91.315681)24:50.0,((6:100.0,7:100.0)76:73.432388,
((9:100.0,10:100.0)91:87.416183,((11:100.0,15:100.0)30:29.816542,(16:100.0,((14:100.0,20:100.0)50:63.010063,
(18:100.0,19:100.0)35:34.675461)41:45.568012)35:39.394554)25:23.766048)52:42.3649065)26:42.3649065)37_Te
mnospondyli:62.2652499999999995,(2:100.0,((4:100.0,5:100.0)39:43.098595,(46:100.0,(33:100.0,(35:100.0,
((21:100.0,22:100.0)40_Tuditanidae:33.360523,((23:100.0,24:100.0)47_Hapsidopareiontidae:51.669201,
(((25:100.0,36:100.0)52_Pantylidae:56.898338,(30:100.0,31:100.0)77_Ostodolepididae:79.032425)10:7.018339,
(34:100.0,(32:100.0,((26:100.0,27:100.0)68:70.956215,
(28:100.0,29:100.0)67:64.295372)52_Gymnarthridae:52.899769)19:10.173872)7:10.173872)12_Recumbirostra:21.
328098,((45:100.0,((37:100.0,(38:100.0,39:100.0)75:78.576607)93_Urocordylidae:98.166664,
((40:100.0,41:100.0)88:88.51355,(42:100.0,
(43:100.0,44:100.0)100:100.0)69:72.758209)83_Diplocaulidae:79.589012)50:48.661064)39_Nectridea:34.552895,
((47:100.0,48:100.0)96_Aistopoda:97.81675,(49:100.0,(50:100.0,(51:100.0,(52:100.0,
(53:100.0,54:100.0)97_Salientia:99.193863)88_Batrachia:94.06057)70_Parotoidia:83.267189)23_Lissamphibia:30.
000544)22:25.682316)17:21.598787)19:29.115671)14:12.156971)9:10.932807)14:13.878294)29:35.719635)44:52.
283924)68_Amphibia:78.760185)28_Tetrapoda:35.696217)30:36.594379)46:40.75029):100.0);

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END;

BEGIN ASSUMPTIONS;

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OPTIONS DEFTYPE = unord PolyTcount = MINSTEPS;
TYPESET * UNTITLED = unord: 3 - 15 17 - 28 30 - 31 33 35 - 38 40 - 73 76 - 81 83 - 86 88 - 102 104
- 114 116 - 125 127 129 131 - 133 135 137 139 - 143 146 - 148 150 - 154 156 - 158 160 - 169 171 173 - 174
176 - 178 180 182 184 - 194 196 199 202 - 203 205 - 207 210 212 - 219, ord: 1 - 2 16 29 32 34 39 74 - 75 82
87 103 115 126 128 130 134 136 138 144 - 145 149 155 159 170 172 175 179 181 183 195 197 - 198 200 - 201
204 208 - 209 211;

```

WTSET * UNTITLED = 1: 1 - 219 ;

END;

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begin paup;
pset mstaxa = variable;
exclude 103 146 163 217;

```

END;

BEGIN NOTES;

TEXT CHARACTER = 12 TEXT = recode_to_remove_lower_orbital_process;

TEXT TAXON = 1 CHARACTER = 16 TEXT = 'Neither process present, but lacrimal is excluded from orbit. See Clack, 2002: fig. 6A. Lacrimal excluded from orbital margin, apparently as in Gerobatrachus. Thus, inapplicable.';

TEXT TAXON = 4 CHARACTER = 16 TEXT = Fine.;

TEXT TAXON = 5 CHARACTER = 16 TEXT = 'Borderline, but OK.';

TEXT TAXON = 9 CHARACTER = 16 TEXT = 'Fröbisch & Schoch, 2009, fig. c clearly show that the absence of the ventral process reflects the immature status of the other specimens.';

TEXT TAXON = 11 CHARACTER = 16 TEXT = Fine.;

TEXT TAXON = 12 CHARACTER = 16 TEXT = LEP_covers_ventral;

TEXT TAXON = 15 CHARACTER = 16 TEXT = 'See Schoch & Rudibge 2005, fig. 3 B, E. At least some specimens have the ventral process.';

TEXT TAXON = 25 CHARACTER = 16 TEXT = Fine.;

TEXT TAXON = 37 CHARACTER = 16 TEXT = Probably_correct.;

TEXT TAXON = 51 CHARACTER = 16 TEXT = 'Both processes present. See McGowan, 2002: fig. 5B.';

TEXT TAXON = 52 CHARACTER = 16 TEXT = 'In Karaurids, the lacrimal is excluded from the orbit. See Ivachnenko, 1978: fig. 1a. But in Hynobius, the lacrimal has both processes ; see Carroll & Holmes, 1980, fig. 4A.';

TEXT TAXON = 22 CHARACTER = 22 TEXT = Fine.;

TEXT TAXON = 23 CHARACTER = 22 TEXT = 'Borderline. See Carroll & Gaskill, 1978: 27 and fig. 13.';

TEXT TAXON = 24 CHARACTER = 22 TEXT = 'Fine. Borderline. See Carroll & Gaskill, 1978: fig. 22.';

TEXT TAXON = 25 CHARACTER = 22 TEXT = 'Fine. See Carroll & Gaskill, 1978: fig. 25.';

TEXT TAXON = 26 CHARACTER = 22 TEXT = 'Fine. See Carroll & Gaskill, 1978: fig. 30.';

TEXT TAXON = 27 CHARACTER = 22 TEXT = 'Fine. See Carroll & Gaskill, 1978: fig. 31?32.';

TEXT TAXON = 28 CHARACTER = 22 TEXT = 'Fine. See Carroll & Gaskill, 1978: fig. 37.';

TEXT TAXON = 29 CHARACTER = 22 TEXT = 'Fine. See Carroll & Gaskill, 1978: fig. 43.';

TEXT TAXON = 3 CHARACTER = 26 TEXT = 'Greererpeton changed from 0 to 1. See Smithson (1982).^n';

TEXT TAXON = 5 CHARACTER = 26 TEXT = 'Limnoscelis changed from 0 to 2. See Fracasso 1983.^n';

TEXT TAXON = 21 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 4.';

TEXT TAXON = 22 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 6, 7.';

TEXT TAXON = 23 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 4.';

TEXT TAXON = 24 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 17, 22.';

TEXT TAXON = 25 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 25.';

TEXT TAXON = 26 CHARACTER = 26 TEXT = Premaxilla_apparently_not_well_preserved.;

TEXT TAXON = 27 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 32.';

TEXT TAXON = 28 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 36, 37.';

TEXT TAXON = 29 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 43, 43.';

TEXT TAXON = 30 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 48.';

TEXT TAXON = 31 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 52-54.';

TEXT TAXON = 32 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: figs 63, 64.';

TEXT TAXON = 33 CHARACTER = 26 TEXT = 'See Carroll & Gaskill, 1978: fig. 76.';

TEXT TAXON = 34 CHARACTER = 26 TEXT = 'Not well preserved. See Carroll & Gaskill, 1971.';

TEXT TAXON = 35 CHARACTER = 26 TEXT = 'Not well preserved. See: Carroll R. L., P. Bybee, & W. D. Tidwell. 1991. The oldest microsauro (amphibia). Journal of Paleontology 65: 314-322.^n^n';

TEXT TAXON = 36 CHARACTER = 26 TEXT = 'See figs. 6, 10, in: Berman D. S., D. A. Eberth, & D. B. Brinkman. 1988. Stegotretus agyrus a new genus and species of microsauro (amphibian) from the Permo-Pennsylvanian of New Mexico. Annals of Carnegie Museum 57: 293-323.^n^n';

TEXT TAXON = 37 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 53A, B.';

TEXT TAXON = 38 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 53G.';

TEXT TAXON = 39 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 53F.';

TEXT TAXON = 40 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56, 57.';

TEXT TAXON = 41 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56B.';

TEXT TAXON = 42 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56C.';

TEXT TAXON = 43 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56E.';

TEXT TAXON = 44 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56D.';

TEXT TAXON = 45 CHARACTER = 26 TEXT = 'See Bossy & Milner, 1998: fig. 56C.';

TEXT TAXON = 46 CHARACTER = 26 TEXT = 'Andrews & Carroll, 1991: fig. 7.';

TEXT TAXON = 47 CHARACTER = 26 TEXT = 'Carroll, 1998: fig. 7.';

TEXT TAXON = 48 CHARACTER = 26 TEXT = 'Figs 2, 3 in: Anderson J. S. 2002. Revision of the Aïstopod genus *Phlegethontia* (Tetrapoda-Lepospondyli). *Journal of Paleontology* 76: 1029-1046.';

TEXT TAXON = 49 CHARACTER = 26 TEXT = 'Wellstead, 1991, figs. 2, 3 (reconstructions), suggest state 2. Figs 7, 8 (specimen drawings) suggest states 1 or 2.';

TEXT TAXON = 50 CHARACTER = 26 TEXT = 'See Jenkins et al., figs 1, 2, 4, 7.';

TEXT TAXON = 51 CHARACTER = 26 TEXT = 'See Estes, 1969: fig. 2; Fox & Naylor, 1982: figs 1, 2; McGowan, 2002: fig. 5.';

TEXT TAXON = 52 CHARACTER = 26 TEXT = 'Karaurus has state 1 (Ivachnenko, 1978: fig. 1). Hynobiidae has states 3 (*Hynobius tsuensis*) or 1 (*Batrachuperus sinensis*), or indeterminate (*Hynobius naevius*), according to Carroll & Holmes, 1980: fig. 4. Cryptobranchidae has state 2, according to Carroll & Holmes, 1980: fig. 4. Ambystomatidae has state 2 (fig. 6). Plethodontids have state 3 (fig. 7). Salamandrids have state 3 (fig. 8A) or indeterminate (fig. 8B). Proteidae has state 3 (fig. 9). Amphiumidae has state 3 (fig. 10). Sirenidae has state 3 (fig. 11). If this is optimized (unordered) on the phylogeny in fig. 8 of this paper: ^nWiens J. J., R. M. Bonett, & P. T. Chippindale. 2005. Ontogeny discombobulates phylogeny: paedomorphosis and higher-level salamander relationships. *Systematic Biology* 54: 91-110.^nStates 1 or 3 may be primitive for the whole clade (including *Karaurus*). Coded as such.';

TEXT TAXON = 54 CHARACTER = 26 TEXT = 'According to Duellman & Trueb, 1986, we find the following conditions (figs. 13-17 and 13-18): *Barbourula busuquanensis* (Discoglossidae), state 3. *Rhinophryus dorsalis* (Rhinophrynidae), state 3, etc. Most have state 3, although a few may have state 0.';

TEXT TAXON = 49 CHARACTER = 34 TEXT = 'Changed *Brachydectes* from 2 (convex) to 1 (straight). See Wellstead, 1991: figs 2, 3, and especially, 8.^n';

TEXT TAXON = 50 CHARACTER = 34 TEXT = 'Changed *Eocaecilia* from 2 (convex) to 3 (undulating). See Jenkins et al, 2007: figs 1, 2.^n';

TEXT TAXON = 51 CHARACTER = 39 TEXT = 'Changed from 2 to ? (there is no otic notch).';

TEXT CHARACTER = 43 TEXT = not_as_margin_of_otic_notch;

TEXT CHARACTER = 51 TEXT = 'Inapplicable when the supratemporal is present (5(0)).';

TEXT TAXON = 33 CHARACTER = 51 TEXT = 'Polymorphic. See Carroll & Gaskill, 1978; figs 77, 78.';

TEXT TAXON = 53 CHARACTER = 59 TEXT = Triadobatrachus_changed_from_0_to_1.;

TEXT TAXON = 50 CHARACTER = 74 TEXT = '10-12 positions (Jenkins et al., 2007: 298).';

TEXT TAXON = 51 CHARACTER = 74 TEXT = 'Changed *Albanerpetontidae* from ? to 1. McGowan (2000: 367) wrote, about *Albanerpeton nexuosus*, that '?Premaxillae with a complete tooth row have eight (n = 4) or nine (n = 5) loci.? McGowan 2002: 5, ?there are 8?10 teeth in *Celtdens*?, so the average is presumably about 9 teeth. In *Albanerpeton pannonicus*, Vencel & Gardner, 2005: 1282 wrote: '?The number of tooth positions ranges from 7 to 10 (x = 8; n = 25) on the premaxilla, 15?21 (x = 18; n = 16) on the maxilla and 22?29 (x = 25; n = 27) on the dentary.? ^n';

TEXT TAXON = 52 CHARACTER = 74 TEXT = 'Salamanders were dealt with as for the dorsal process of the premaxilla, character 26 (same phylogeny and references). Distribution: *Karaurus*, about 25 teeth, according to the text (Ivachnenko, 1978: 364); 9 teeth in *Batrachuperus sinensis* (Hynobiidae); more than 10 in *Cryptobranchus*, *Ambystoma*; about 8 in *Phaeognathus hubrichti* (Plethodontidae), about 4 in *Salamandra atra* (Salamandridae), but more than 10 in *Notophthalmus viridescens* (Salamandridae), more than 10 in *Necturus* (Proteidae). About 5 in *Amphiuma* (Amphiumidae) and in *Habrosaurus* (Sirenidae). The ancestral condition is ambiguous. Coded as polymorphic.^n';

TEXT TAXON = 54 CHARACTER = 74 TEXT = 'Frogs were dealt in the same way. *Viaerella* seems to have numerous teeth, although this is based only on the reconstruction of Shubin & Jenkins (1995: fig. 3a). *Eodiscoglossus* has at least 15 tooth positions (Evans et al., 1990: 302). *Ascaphus* has well over 10 teeth (Carroll & Holmes, 1980: fig. 3).^nEvans S. E., A. R. Milner, & F. Mussett. 1990. A discoglossid frog from the Middle Jurassic of England. *Palaeontology* 33: 299-311.^nIt seems fair to code them as 0 (at least 10 tooth positions). Changed from

0/1 to 0.^n';

TEXT TAXON = 51 CHARACTER = 75 TEXT = 'Changed Albanerpetontidae from ? to 1&2. McGowan (2000: 367) wrote, about Albanerpeton nexuosus, that ?No maxilla available to me has an intact tooth row: UALVP 39973 (Fig. 3C) and 39977 (Fig. 3E) preserve the anterior 17 and 18 tooth positions, respectively, and I estimate that each bone probably held about 25 loci when complete.? In Albanerpeton pannonicus, Vencel & Gardner, 2005: 1282 wrote: The number of tooth positions ranges from 7 to 10 (x = 8; n = 25) on the premaxilla, 15?21 (x = 18; n = 16) on the maxilla and 22?29 (x = 25; n = 27) on the dentary. I believe that x represents the average. McGowan (1999: 536) wrote, about Albanerpeton arthridion: ?Neither maxilla has an intact tooth row: OMNH 33284 preserves the posteriormost 14 loci; OMNH 34072 preserves 12 loci along the anterior part of the tooth row, but the anteriormost end of the row may be absent. Judging by the overlap between OMNH 33284 and 34072, I estimate that the maxilla held about 23 loci when complete.?^n';

TEXT TAXON = 52 CHARACTER = 75 TEXT = 'Urodeles were dealt with as for the dorsal process of the premaxilla (ch. 26), except that the states were ordered. Karaurus, up to 40 positions (Ivachnenko, 1979: 364). Batrachuperus sisnesis (Hynobiidae) has 16?20 teeth (Carroll & Holmes, 1980: fig. 4B). Cryptobranchus and Ambystoma, Salamandra, and Plethodon glutinosus (Noble, 1931: fig. 81) have more than 30. Notophthalmus (Salamandridae) has about 22 teeth, and Tylostrotion (Salamandridae), barely 20. Amphiuma has 17-18 teeth and Habrosaurus (Sirenidae) has about 9 teeth. The ancestral condition is unambiguously 0.^n';

TEXT TAXON = 20 CHARACTER = 84 TEXT = 'In Gerobatrachus, the occipital condyle is apparently not preserved! Anderson et al. (2008) wrote: ?The parabasisphenoid complex is fragmentary, preserving only portions of the basicranial articulation, and a portion of the cultriform process; however, the overall shape of the parasphenoid plate can be determined to have been much wider than long, as is common for amphibamids, branchiosaurids, frogs and salamanders.? The figures show no condyle. Coding changed from 1 to ?^n';

TEXT TAXON = 12 CHARACTER = 85 TEXT = 'Ecolsonia : Changed from 0 to 1. See Berman et al., 1985: p. 16 "...the fused basioccipital-exoccipital complex has been somewhat rotated anteriorly so that the articular surfaces of the distinctly double occipital condyle face mainly ventrally rather than posteriorly.'";

TEXT TAXON = 14 CHARACTER = 85 TEXT = Fine.;

TEXT TAXON = 20 CHARACTER = 85 TEXT = 'In Gerobatrachus, the occipital condyle is apparently not preserved! Anderson et al. (2008) wrote: "The parabasisphenoid complex is fragmentary, preserving only portions of the basicranial articulation, and a portion of the cultriform process; however, the overall shape of the parasphenoid plate can be determined to have been much wider than long, as is common for amphibamids, branchiosaurids, frogs and salamanders." The figures show no condyle. Coding changed from 1 to ?^n';

TEXT TAXON = 21 CHARACTER = 85 TEXT = 'Tuditonus. Changed from 0 to ?. See Carroll & Baird, 1968: 12 : « The surfaces that formed part of the occipital condyle cannot be seen. This is confirmed by the illustrations, although a non-diagnostic reconstruction of this area in ventral is given in Carroll & Gaskill, 1978 : fig. 4.^n';

TEXT TAXON = 22 CHARACTER = 85 TEXT = 'Asaphestera. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 6E.^n';

TEXT TAXON = 23 CHARACTER = 85 TEXT = 'Hapsidopareion. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 13B.^n';

TEXT TAXON = 24 CHARACTER = 85 TEXT = 'Saxonerpeton. Changed from 0 to ?. See Carroll & Gaskill, 1978: 34. The condyle was probably double, but this area is poorly preserved.^n';

TEXT TAXON = 25 CHARACTER = 85 TEXT = 'Pantylus. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 25.^n';

TEXT TAXON = 26 CHARACTER = 85 TEXT = 'Cardiocephalus sternbergi. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 30B, C.^n';

TEXT TAXON = 27 CHARACTER = 85 TEXT = 'Cardiocephalus peabodyi. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 31. Although this area is poorly exposed, the shape of the atlantal articular surfaces indicate a typical microsaurian cotyle.^n';

TEXT TAXON = 28 CHARACTER = 85 TEXT = 'Euryodus primus. Changed from 0 to 1. See Carroll & Gaskill, 1978: figs 37, 38, 41. Although this area is poorly exposed, the shape of the atlantal articular surfaces indicate a typical microsaurian cotyle.^n';

TEXT TAXON = 29 CHARACTER = 85 TEXT = 'Euryodus dalyae. Changed from 0 to 1. See Carroll & Gaskill, 1978: figs 42?44. ^n';

TEXT TAXON = 30 CHARACTER = 85 TEXT = 'Pelodosotis. Changed from 0 to 1. See Carroll & Gaskill,

1978 : fig. 48A.^n';

TEXT TAXON = 31 CHARACTER = 85 TEXT = 'Micraroter. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 53.^n';

TEXT TAXON = 32 CHARACTER = 85 TEXT = 'Rhynchonkos. Changed from 0 to 1. See Carroll & Gaskill, 1978 : figs 63?65 (under the name Goniorhynchus).^n';

TEXT TAXON = 33 CHARACTER = 85 TEXT = 'Microbrachis. Changed from 0 to 1. See Carroll & Gaskill, 1978: fig. 77?79; Laurin & Vallin, 2004: 62.^n';

TEXT TAXON = 34 CHARACTER = 85 TEXT = 'Batropetes. Changed from 0 to ?. See Carroll & Gaskill, 1971 :454. This area is not well exposed.^n';

TEXT TAXON = 36 CHARACTER = 85 TEXT = 'Stegotretus. Changed from 0 to 1. See Berman et al., 1988: 310.';

TEXT TAXON = 37 CHARACTER = 85 TEXT = 'Sauropleura scalaris. Changed from 0 to 1. See Bossy, 1976: 96 and fig. 25b.^n';

TEXT TAXON = 39 CHARACTER = 85 TEXT = 'Urocordylus. Changed from 0 to ?. See Bossy, 1976: 209. She wrote that it must be ?semidouble?, which suggests an intermediate morphology.^n';

TEXT TAXON = 14 CHARACTER = 86 TEXT = 'In Doleserpeton, it was not described, so we changed the coding from 1 to ?';

TEXT TAXON = 54 CHARACTER = 86 TEXT = 'In anurans, there is no prootic, and the exoccipital is fused in most taxa to form a compound bone, so the position of this foramen cannot be assessed relative to the opisthotic. There is no place for it to exit but through the exoccipital. Anurans were thus changed from 1 to ?';

TEXT TAXON = 54 CHARACTER = 87 TEXT = 'Highly variable in frogs. Unknown in Vieraella. Posterior in Barbourula (Discoglossidae) (Duellman & Trueb, 1986: fig. 13-17), Caudiverbera (Leptodactylidae). Even with occiput in Leptodactylus. Anterior in Ascaphus (Noble, 1932: fig. 81A; Carroll & Holmes, 1980: fig. 3), Pelobates (Pelobatidae), Rhamphophyrne (Bufonidae). Far anterior in Rhinophyus (check spelling) (Rhinophyrnidae), Notaden (Myobatrachidae), Brachycephalus (Brachycephalidae). Highly variable in hylids (Duellman & Trueb, 1986: fig. 13-18). Can conservatively be scored ? (initially 0).^n';

TEXT CHARACTER = 89 TEXT = 'Characters 89 and 90 remade, see text.';

TEXT TAXON = 53 CHARACTER = 89 TEXT = 'Triadobatrachus has a preserved palatine, illustrated and described by Rage & Rocek (1989). It appears to be edentulous, so it was coded as such (changed from ?).^n';

TEXT TAXON = 54 CHARACTER = 89 TEXT = 'Frogs never have palatine teeth (Duellman & Trueb, 1986: 318).';

TEXT CHARACTER = 90 TEXT = 'Characters 89 and 90 remade, see text.';

TEXT TAXON = 3 CHARACTER = 90 TEXT = 'Smithson_1982.';

TEXT TAXON = 5 CHARACTER = 90 TEXT = 'Fracasso_1983_fig_2.';

TEXT TAXON = 15 CHARACTER = 90 TEXT = 'Schoch_&_Rubidge_2005_fig_1D.';

TEXT TAXON = 33 CHARACTER = 90 TEXT = 'Carroll_&_Gaskill_1978.';

TEXT TAXON = 49 CHARACTER = 90 TEXT = 'Brachydectes changed from 1 to 0.^n';

TEXT TAXON = 53 CHARACTER = 90 TEXT = 'Triadobatrachus has a preserved palatine, illustrated and described by Rage & Rocek (1989). It appears to be edentulous, so it was coded as such (changed from ?).';

TEXT TAXON = 54 CHARACTER = 90 TEXT = 'Frogs changed from 1&2 to 0, since there are no teeth on the palatine.^n';

TEXT CHARACTER = 91 TEXT = 'if_ex_fused_in_os_basiale,_if_pt_contacts_the_ex_area';

TEXT TAXON = 5 CHARACTER = 94 TEXT = 'Limnoscelis changed from 2 to 0.^n';

TEXT TAXON = 14 CHARACTER = 94 TEXT = 'Doleserpeton, changed from 1/2 to 1&2.';

TEXT TAXON = 19 CHARACTER = 94 TEXT = 'Amphibamus changed from 1 to 0.^n';

TEXT TAXON = 20 CHARACTER = 94 TEXT = 'Gerobatrachus_changed_from_1_to_0.';

TEXT TAXON = 54 CHARACTER = 94 TEXT = 'Correct. See Duellman & Trueb, 1986: 317. Also see Carroll & Holmes, 1980, fig 3B (Vieraella), Duellman & Trueb, 1986: fig. 13-17 A (Barbourula busuquanensis, Discoglossidae), C (Pelobates fuscus, Pelobatidae), E (Leptodactylus bolivianus, Leptodactylidae), Noble, 1931, fig. 80b (Rana adspersa), 81A (Ascaphus truei).^n';

TEXT TAXON = 37 CHARACTER = 97 TEXT = 'Fine. See Bossy, 1976: fig 24.';

TEXT TAXON = 39 CHARACTER = 97 TEXT = 'Fine. See Bossy, 1976: fig 69.';

TEXT TAXON = 41 CHARACTER = 97 TEXT = 'Eocaecilia: Changed from 1 to 0. See Jenkins et al., 2007: figs 1, 3, 7.^n';

TEXT TAXON = 20 CHARACTER = 99 TEXT = 'Changed Gerobatrachus from 0 to ?. See Marjanovic & Laurin, 2008: 179.^n';

TEXT TAXON = 45 CHARACTER = 99 TEXT = 'Milner 1980: 392 says they're pedicellate. No answer yet. The teeth of Scincosaurus have never been mentioned before or since.';

TEXT TAXON = 47 CHARACTER = 99 TEXT = 'Interpretation of Carroll, 1998: fig. 4B.';

TEXT TAXON = 48 CHARACTER = 99 TEXT = 'Germain, 2008.';

TEXT TAXON = 20 CHARACTER = 100 TEXT = 'Changed from 1 to 0. Gerobatrachus has denticles, not teeth, according to the illustrations (contra the text) provided by Anderson et al. (2008). They are much smaller than the marginal teeth (about a third).^n';

TEXT TAXON = 54 CHARACTER = 100 TEXT = 'Frogs also sometimes have teeth, but not denticles.';

TEXT TAXON = 52 CHARACTER = 102 TEXT = 'Changed frogs and salamanders from 0 to 1. Carroll & Holmes (1980) show no denticles on the parasphenoid of any urodele, although salamandrids are shown as having a row of teeth. Noble (1931: fig. 81b) shows that plethodontids can have patches of teeth (I would not call these denticles, given that they are as large as marginal teeth). Given the deeply nested position of salamandrids and plethodontids, this is surely not the primitive condition for urodeles.^n';

TEXT TAXON = 53 CHARACTER = 102 TEXT = 'Triadobatrachus also lacked denticles (Rage & Rocek, 1989).^n';

TEXT TAXON = 54 CHARACTER = 102 TEXT = 'Noble (1931: fig. 81a) shows that Ascaphus lacks denticles on the parasphenoid. Duellman & Trueb (1986: figs 13-17 and 13-18) show no parasphenoid denticles on any anuran.^n';

TEXT CHARACTER = 103 TEXT = 'Correlated to lots of other stuff, see text.';

TEXT TAXON = 20 CHARACTER = 105 TEXT = 'The parasphenoid of Gerobatrachus is extremely fragmentary; it seems fair to score it as ? See Anderson et al., 2008 (the illustration; the text contradicts it).';

TEXT TAXON = 21 CHARACTER = 105 TEXT = 'Tuditanus. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 4.^n';

TEXT TAXON = 23 CHARACTER = 105 TEXT = 'Hapsidoparierion. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 13.^n';

TEXT TAXON = 24 CHARACTER = 105 TEXT = 'Saxonerpeton. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 22.^n';

TEXT TAXON = 26 CHARACTER = 105 TEXT = 'Cardiocephalus sternbergi. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 30.^n';

TEXT TAXON = 28 CHARACTER = 105 TEXT = 'Euryodus primus. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 36?38.^n';

TEXT TAXON = 29 CHARACTER = 105 TEXT = 'Euryodus delyae. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 42?43.^n';

TEXT TAXON = 30 CHARACTER = 105 TEXT = 'Pelodosotis. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 48.^n';

TEXT TAXON = 31 CHARACTER = 105 TEXT = 'Microrator. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 53.^n';

TEXT TAXON = 32 CHARACTER = 105 TEXT = 'Rhynchonkos. Changed from 0 to 1. See Carroll & Gaskill, 1978 : figs 63?65.^n';

TEXT TAXON = 33 CHARACTER = 105 TEXT = 'Microbrachis. Changed from 0 to 1. See Carroll & Gaskill, 1978 : fig. 77.^n';

TEXT TAXON = 36 CHARACTER = 105 TEXT = 'Stegotretus. Changed from 0 to 1. See Berman et al., 1988: fig. 8.^n';

TEXT TAXON = 37 CHARACTER = 105 TEXT = Borderline.;

TEXT TAXON = 38 CHARACTER = 105 TEXT = 'Ptyonius. Changed from 0 to 1. See Bossy, 1976: fig. 49 and Bossy & Milner, 1998: fig. 55B.^n';

TEXT TAXON = 39 CHARACTER = 105 TEXT = 'Urocordylus. Changed from 0 to ?. Bossy & Milner, 1998: fig. 55A.^n';

TEXT TAXON = 41 CHARACTER = 105 TEXT = 'Batachiderpeton. Changed from 0 to ?. See Bossy & Milner, 1998: fig. 57B. The parasphenoid is shown, but has a strange morphology unlike any other state, and seem poorly preserved, if at all, posteriorly.^n';

TEXT TAXON = 44 CHARACTER = 105 TEXT = 'Diploceraspis. Changed from 0 to 1. See Beerbower, 1963: fig. 4. A bit unlike others, but closer to 2 than to 0.^n';

TEXT TAXON = 45 CHARACTER = 105 TEXT = 'Scincosaurus. Changed from 0 to 1. See Bossy & Milner, 1998: fig. 57D.^n';

TEXT TAXON = 46 CHARACTER = 105 TEXT = 'Adelogyrinus. Changed from 0 to ?. See Andrews & Carroll, 1991: fig. 6. If it was like Palaeomolgophis, it probably had state 1 (fig. 2).^n';

TEXT TAXON = 47 CHARACTER = 105 TEXT = 'Oestocephalus. Changed from 0 to 1. See Carroll, 1998: fig. 3.^n';

TEXT TAXON = 49 CHARACTER = 105 TEXT = 'Brachydectes changed from 1 to 1. See Wellstead, 1991: fig. 2.^n';

TEXT TAXON = 50 CHARACTER = 105 TEXT = 'Eocaecilia changed from 1 to 1. See Jenkins et al., 2007.^n';

TEXT TAXON = 52 CHARACTER = 105 TEXT = 'Salamanders changed from 1 to 1. See Carroll & Holmes, 1980.^n';

TEXT TAXON = 4 CHARACTER = 113 TEXT = '=orbitosphenoid_of_White_1938_(Romer,_1956)';

TEXT TAXON = 4 CHARACTER = 120 TEXT = Changed_from_0_to_1.;

TEXT TAXON = 5 CHARACTER = 120 TEXT = Changed_from_0_to_1.;

TEXT TAXON = 7 CHARACTER = 120 TEXT = Milner_1996_fig._6.;

TEXT TAXON = 9 CHARACTER = 120 TEXT = 'Boy 1978, 1986, 1987.';

TEXT TAXON = 21 CHARACTER = 120 TEXT = 'Unknown (C&G fig. 108, Carroll & Baird 1968).';

TEXT TAXON = 22 CHARACTER = 120 TEXT = C&G_fig._108;

TEXT TAXON = 23 CHARACTER = 120 TEXT = C&G_fig._108;

TEXT TAXON = 24 CHARACTER = 120 TEXT = C&G_fig._108;

TEXT TAXON = 26 CHARACTER = 120 TEXT = 'Precisely intermediate (C&G fig. 30).';

TEXT TAXON = 28 CHARACTER = 120 TEXT = C&G_fig._109;

TEXT TAXON = 29 CHARACTER = 120 TEXT = C&G_fig._109;

TEXT TAXON = 32 CHARACTER = 120 TEXT = 'C&G fig. 64 is somewhat borderline, but can still be counted. Fig. 63 is unambiguously state 1.';

TEXT TAXON = 33 CHARACTER = 120 TEXT = Vallin_&_Laurin_2004.;

TEXT TAXON = 34 CHARACTER = 120 TEXT = 'Too incomplete to tell (Carroll 1991 fig. 5B).';

TEXT TAXON = 37 CHARACTER = 120 TEXT = Bossy_&_Milner_1998_fig._55C_is_juvenile._This_is_interesting_because_S._pectinata_has_state_1.;

TEXT TAXON = 38 CHARACTER = 120 TEXT = 'Bossy & Milner 1998 fig. 55B: absent on the left side, present on the right...';

TEXT TAXON = 46 CHARACTER = 120 TEXT = 'Palate unknown (Andrews & Carroll 1991).';

TEXT TAXON = 50 CHARACTER = 120 TEXT = Jenkins_et_al._2007.;

TEXT TAXON = 52 CHARACTER = 120 TEXT = 'Changed from 0&1 to 0. See Carroll & Holmes, 1980. Hynobiids seem to be polymorphic, but the flange is absent in Cryptobranchus and all other urodeles figured in that paper, except for very weak flanges in Ambystoma maculatum and Tylostrotion, but even in these two genera, the structure is so slight that it can be considered absent.';

TEXT TAXON = 53 CHARACTER = 120 TEXT = 'Changed from 0/1 to 0. See Rage & Rocek, 1989: fig. 2B.';

TEXT TAXON = 54 CHARACTER = 120 TEXT = 'Changed from 0&1 to 0/1. See text.';

TEXT CHARACTER = 121 TEXT = 'triangular=non_parallel-sided__rect=parallel-sided';

TEXT TAXON = 14 CHARACTER = 121 TEXT = 'Changed Doleserpeton from 1 to 2. See Bolt, 1969.^n';

TEXT TAXON = 25 CHARACTER = 121 TEXT = 'Changed Pantylus from 1 to 2. See Carroll & Gaskill, 1978: fig. 25.^n';

TEXT TAXON = 36 CHARACTER = 121 TEXT = 'Changed from 1 to 2: Berman et al. 1988: 308.';

TEXT TAXON = 49 CHARACTER = 121 TEXT = 'Changed Brachydectes from 1 to 2. See Wellstead, 1991:';

fig. 2.^n';

TEXT TAXON = 50 CHARACTER = 121 TEXT = 'Changed Eocaecilia from 1 to 2. See Jenkins et al, 2007, fig. 1, and text (that region is supposedly hidden). That is true, to a small extent, but the drawings and pictures show plainly that there could not be one. The quote of Nussbaum and Wilkinson (1989) cannot be substantiated; personal observations by one of us (ML) in Marvalee Wake's collection have failed to reveal a single gymnophionan ectopterygoid, and no more recent investigation has substantiated the presence of this bone. -- Changed to 1 or 2 because a very small one (as found in several extant caecilians) could have been present; Jenkins et al. 2007: 304.';

TEXT TAXON = 52 CHARACTER = 121 TEXT = 'Changed salamanders from 1 to 2. See Carroll & Holmes, 1980.^n';

TEXT TAXON = 53 CHARACTER = 121 TEXT = 'Changed Triadobatrachus from 1 to 2. See Rage & Rocek, 1989.^n';

TEXT TAXON = 54 CHARACTER = 121 TEXT = 'Changed frogs from 1 to 2. See Duellman & Trueb, 1986: figs 13-17 and 13-18.^n';

TEXT CHARACTER = 131 TEXT = 'at_least_one_zygo"s_width_again_from_centrum';

TEXT TAXON = 50 CHARACTER = 138 TEXT = 'is_second_row_of_equal_teeth,_but_pseudentary,_no_coronoid';

TEXT CHARACTER = 146 TEXT = 'When this is coded as simultaneous, this probably just reflects a lack of temporal resolution. There is widespread agreement about this among developmental biologists (e.g., Bininda-Emonds et al., 2003: 341). This character should be deleted; it is scored for very few taxa anyway.^nBininda-Emonds O. R. P., J. E. Jeffrey, & M. K. Richardson. 2003. Is sequence heterochrony an important evolutionary mechanism in mammals? Journal of Mammalian Evolution 10: 335-361.^n';

TEXT TAXON = 35 CHARACTER = 146 TEXT = 'Changed Utaherpeton from ? to 0. See Carroll & Chorn, 1995: fig. 4b. ^n';

TEXT CHARACTER = 153 TEXT = 'triangular=non_parallel-sided__rect=parallel-sided';

TEXT CHARACTER = 165 TEXT = 'at_least_one_zygo"s_width_again_from_centrum';

TEXT TAXON = 37 CHARACTER = 168 TEXT = 'Perhaps initially based on Bossy & Milner, 1998: fig. 59C, that is a bit ambiguous. Bossy, 1976: fig. 25B suggests an expanded anterior surface. Confirmed by text, p. 102: "the lateral facets for the exoccipitals are only slightly expanded, so that the anterior face of the atlas is not much broader than the more posterior centra." Thus, there is some expansion. Changed to 1.';

TEXT TAXON = 38 CHARACTER = 168 TEXT = 'Ptyonius. Changed from 0 to 1. See Bossy, 1976: fig. 51a. Again, the expansion is very modest, but it is there.^n';

TEXT TAXON = 39 CHARACTER = 168 TEXT = 'Urocordylus. Changed from 0 to 1. See Bossy, 1976: 215: ?Its anterior face is slightly concave and broadened laterally, probably for articulation with the exoccipital?.';

TEXT TAXON = 40 CHARACTER = 168 TEXT = 'Keraterpeton. Changed from 0 to ?. Does not seem to be known, although a specimen certainly has a preserved atlas (but in dorsal view, which hides the centrum). See Huxley & Wright, 1867: plate XIX, and Jaekel, 1903. ^nJaekel O. 1903. Ueber Ceraterpeton, Diceratosaurus und Diplocaulus. Neues Jahrbuch für Mineralogie, Geologie und Paläontologie 109-134.^n^n^n';

TEXT TAXON = 42 CHARACTER = 168 TEXT = Changed_from_0_to_?._Seems_unknown.;

TEXT TAXON = 45 CHARACTER = 168 TEXT = Changed_from_0_to_?._Seems_unknown.;

TEXT TAXON = 46 CHARACTER = 168 TEXT = 'Correct. See Andrews & Carroll, 1991: fig. 6.';

TEXT TAXON = 49 CHARACTER = 168 TEXT = 'Brachydictes changed from 0 to 1. See Wellstead, 1991: figs 11C, 18.';

TEXT TAXON = 6 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008).';

TEXT TAXON = 7 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008).';

TEXT TAXON = 12 CHARACTER = 179 TEXT = 'All preserved ribs are long and "moderately curved" (Berman et al. 1985).';

TEXT TAXON = 13 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008).';

TEXT TAXON = 27 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008).';

TEXT TAXON = 28 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008) and the text.';

TEXT TAXON = 32 CHARACTER = 179 TEXT = 'See our Cont. Zool. paper (2008).';

TEXT TAXON = 36 CHARACTER = 179 TEXT = 'Unknown and more or less unknowable (Berman et al. 1988: 312).';

TEXT TAXON = 46 CHARACTER = 179 TEXT = Andrews_&_Carroll_1991_fig._6.;

TEXT TAXON = 47 CHARACTER = 179 TEXT = Baird_1964_fig._1.;

TEXT TAXON = 48 CHARACTER = 179 TEXT = Baird_1964_fig._1.;

TEXT TAXON = 50 CHARACTER = 179 TEXT = 'The ribs are tiny, tiny, tiny! (Jenkins et al., 2007)';

TEXT TAXON = 21 CHARACTER = 181 TEXT = 'Completely impossible to tell (Carroll & Baird 1968, figures 4 to 7).';

TEXT TAXON = 27 CHARACTER = 181 TEXT = 'The fossil ends in the middle of the second sacral (Carroll & Gaskill 1986: 57, fig. 31).';

TEXT TAXON = 30 CHARACTER = 181 TEXT = 'Completely unknown (Carroll & Gaskill 1978: 85 and fig. 58A).';

TEXT TAXON = 32 CHARACTER = 181 TEXT = 'Impossible to tell (Carroll & Gaskill 1978 fig. 70).';

TEXT TAXON = 35 CHARACTER = 181 TEXT = 'Carroll et al. 1991:318 actually _say_ "It is uncertain whether it was a second sacral or a first caudal" (and then go on to point out that "The ilium is so narrow that it is unlikely to have accommodated more than a single sacral rib", but let's be generous).';

TEXT TAXON = 53 CHARACTER = 183 TEXT = 'Can't see a reason to exclude states 1 and 2 (Rage & Rocek 1989: 11).';

TEXT TAXON = 37 CHARACTER = 191 TEXT = 'Changed Sauropleura scalaris from 0 to 1. See Bossy, 1976: fig. 33.');

TEXT TAXON = 40 CHARACTER = 191 TEXT = 'Correct. See Bossy & Milner, 1998: fig. 65B.^n';

TEXT TAXON = 49 CHARACTER = 191 TEXT = 'See Wellstead, 1991: fig. 19D.');

TEXT TAXON = 15 CHARACTER = 192 TEXT = 'May be correct. See plate II from Broili & Schrödre 1937. Better, see Schoch & Rubidge, 2005: fig. 6D.^nBroili F. & J. Schröder. 1937. Beobachtungen an Wirbeltieren der Karrooformation. XXV. Über Micropholis Huxley. Sitzungsberichte der Bayerischen Akademie der Wissenschaften 19-38.^nSchoch R. R. & B. S. Rubidge. 2005. The amphibamid Micropholis from the Lystrosaurus Assemblage Zone of South Africa. Journal of Vertebrate Paleontology 25: 502-522.');

TEXT TAXON = 37 CHARACTER = 192 TEXT = 'Changed Sauropleura scalaris from 0 to 1. See Bossy, 1976: fig. 33.');

TEXT TAXON = 38 CHARACTER = 192 TEXT = 'Changed Ptyonius from 0 to 1. See Bossy, 1976: fig. 57a.^n';

TEXT TAXON = 39 CHARACTER = 192 TEXT = 'Borderline, but correct. See Bossy, 1976: fig. 77a.');

TEXT TAXON = 49 CHARACTER = 192 TEXT = Changed_from_0_to_1.;

TEXT TAXON = 1 CHARACTER = 201 TEXT = 'Changed from 0 to 2. See Coates 1996: fig. 16.');

TEXT TAXON = 2 CHARACTER = 201 TEXT = 'Changed Proterogyrinus from 0 to 2. See Holmes, 1984: fig. 26.');

TEXT TAXON = 3 CHARACTER = 201 TEXT = 'Changed Greererpeton from 1 to 2. See Godfrey, 1989: fig. 18.');

TEXT TAXON = 49 CHARACTER = 201 TEXT = 'Correct. See Wellstead, 1991: fig. 21A-D.');

TEXT TAXON = 50 CHARACTER = 201 TEXT = 'Changed Eocaecilia from 2 to 0. See Jenkins et al., 2007: fig. 41.');

TEXT TAXON = 52 CHARACTER = 201 TEXT = 'Salamanders: Weak in Chunerpeton (Cryptobranchidae; Gao & Shubin, 2003: figs 1, 2), Ranodon (Hynobiidae; Averianov, 1995: fig. 4), Palaeoamphiuma (Amphiumidae; Rieppel & Grande, 1998: fig. 5), Necturus (Proteidae; Wischnitzer, 1979: fig. 2-8). Moderate in Parahynobius betfianus (Veanzel, fig. 2E). The primitive condition is clearly for a weak crest. Changed from 2 to 0.^n';

TEXT TAXON = 54 CHARACTER = 201 TEXT = 'Frogs: Prominent in Latonia (Discoglossidae; Rocek, 1994: fig. 17); intermediate in Beleaphryne (Discoglossidae; Sanchiz & Alcover, 1982: figs 1-3); weak in Eoxenopoides reuningi (Pipidae; Estes, 1977: fig. 8), Thoraciliacus (Trueb, 1999: fig. 2), Pipa pipa (Trueb et al., 2000: fig. 7), Xenopus arabiensis (Henrici & Báez, 2001: figs 2-4). Intermediate seems a reasonable guess about the primitive condition.^n';

TEXT TAXON = 51 CHARACTER = 203 TEXT = 'Changed Albanerpetontidae from 0 to 1. See McGowan

2002: fig. 3 for Celtedens.');

TEXT TAXON = 38 CHARACTER = 206 TEXT = 'Correct. See Bossy 1976: 168.';
TEXT TAXON = 39 CHARACTER = 206 TEXT = 'Probably correct. See Bossy 1976: 226.';
TEXT TAXON = 49 CHARACTER = 206 TEXT = 'Correct. See Wellstead, 1991: 32';

TEXT TAXON = 20 CHARACTER = 207 TEXT = 'Changed Gerobatrachus from 1 to ? See Marjanovic & Laurin 2008.^n^n';

TEXT TAXON = 3 CHARACTER = 208 TEXT = 'Changed (Coates 1996:415). See text.';
TEXT TAXON = 7 CHARACTER = 208 TEXT = 'Changed (Holmes et al. 1998); see text.';
TEXT TAXON = 13 CHARACTER = 208 TEXT = 'State 2 can be excluded (Olson 1941).';
TEXT TAXON = 15 CHARACTER = 208 TEXT = 'Changed (Schoch & Rubidge 2005: 512).';
TEXT TAXON = 17 CHARACTER = 208 TEXT = 'State 2 can be excluded (Sumida et al. 1998).';
TEXT TAXON = 24 CHARACTER = 208 TEXT = 'Nothing can be ruled out (C&G 38).';
TEXT TAXON = 32 CHARACTER = 208 TEXT = '4 are preserved, but a higher number cannot be excluded based on the preservation (C&G: 111)';
TEXT TAXON = 39 CHARACTER = 208 TEXT = Bossy_&_Milner_1998.;
TEXT TAXON = 40 CHARACTER = 208 TEXT = Contradictions_in_the_literature._See_text.;
TEXT TAXON = 42 CHARACTER = 208 TEXT = Contradictions_in_the_literature._See_text.;

TEXT CHARACTER = 217 TEXT = "The three cases of 6 are obviously not homologous: in Acanthostega it's the fibulare, in Tuditanus the postminimus or a sesamoid or who knows what, and in salamanders the prehallux. Therefore excluded from all analyses.';

TEXT TAXON = 49 CHARACTER = 219 TEXT = 'Correct. See Wellstead, 1991: 25.';

Version 4.0 87;
LastModified -1017388859;
FileSettings editor '0' '0' '92' '1';
Singles 010;
Editor 0001100111111100100010010 '0' '24' Geneva '9' '100' '1' all;
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TreeWindowPosition '46' '6' '719' '974';
ListWindow Characters closed Geneva '9' '50' '10' '273' '644' 000;
ListWindow Taxa open Geneva '9' '479' '360' '273' '295' 100000000;
ListWindow Trees closed Geneva '9' '564' '39' '129' '393';
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OtherSymbols & / 00 ? -;
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Salmo 00000001;
EditorFile '2';
ExportHTML _ MOSS '100' 110000;
PrettyPrint 10;
EditorToolsPosition '492' '136' '115' '165';
TreeToolsPosition '583' '25' '126' '138';
TreeWindowProgram 10;

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Continuous '0' '3'          1;
Calculations 0000001;
SummaryMode '0' '0'          0;
Charts Geneva '9' (          normal )          0010;
NexusOptions '0' '0' '50'          001011011;
TipLabel '1';
TreeFont Geneva '9' (          normal );
TreeShape 1.0 1.0 1110;
TraceLabels 0101;
ChartColors '0' '0' '65535' '9' '0'          1;
ChartBiggestSpot 1;
ChartPercent 10;
ChartBarWidth '10'          1;
ChartVerticalAxis 10101;
ChartMinMax '0';
TraceAllChangesDisplay '1'          1;
BarsOnBranchesDisplay '0' '0' '60000' '10000' '10000' '10000' '10000' '60000' '65000' '65000' '65000' '6' '1'
0000101;
ContinuousBranchLabels 0;
AllStatesBranchLabels 1;
IndexNotation '2'          1;
PrintTree 10.00 '2' '2' '2' '2' '2' '2' '2' '2' '2' '2' '2'          Geneva '9' (          normal )
Geneva '10' (          normal )          Geneva '9' (          normal )          Geneva '9' (
normal )          Geneva '9' (          bold )          Geneva '9' (          normal )
Geneva '9' (          normal ) '83' '115' '13' '45' '116' '148' '13' '45' '149' '181' '13' '45' '182' '214' '13' '45' '226' '75'
'0' '0' '-15492' '456' '636' '35' '678' '35' '678' '288' '24' '35' '1' '1' '1' '1' '29440' '-17792' '-39' '4' '-40' '0' '1' '2' '4' '8' '8' '8'
'0' '2'          0100110000000000010100000111000;
MatchChar 00 .;
EntryInterpretation 00;
ColorOptions 00;
TreeTools '0' '5' '4' '0' '10' '4' '0'          00100111111101110;
EditorTools '0' '0' '0' '1000' '0' '0' '6' '3' '1'          100010101110011;
PairAlign '2' '2' '3' '2' '1' '1' '2' '1' '3'          1010;
BothTools '1';

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END;

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ProbModelSet * UNTITLED = 'Mk1 (est.): 1 - 219;

END;

Begin MESQUITE;

MESQUITESCRIPTVERSION 2;

TITLE AUTO;

tell ProjectCoordinator;

getEmployee #mesquite.minimal.ManageTaxa.ManageTaxa;

tell It;

setID 0 5128558849605471043;

tell It;

setDefaultOrder 0 1 34 35 36 2 3 42 37 38 39 40 41 43 47 48 49 52 53 51 4 5 6 7 8 9 10 11 12

13 14 15 17 18 19 20 21 22 23 24 25 26 27 28 29 31 32 33 30 16 46 44 50 45;

attachments ;

endTell;

endTell;

getEmployee #mesquite.charMatrices.ManageCharacters.ManageCharacters;

tell It;

```
setID 0 8479452511181763361;
tell It;
  setDefaultOrder 0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15 16 17 18 19 20 21 22 23 24 25 26 27 28
29 30 31 32 33 34 35 36 37 38 39 40 41 42 43 44 45 46 47 48 49 50 51 52 53 54 55 56 57 58 59 60 61 62 63 64 65
66 67 68 69 70 71 72 73 74 75 76 77 78 79 80 81 82 83 84 85 86 87 88 89 90 91 221 93 94 95 96 97 98 99 100 101
102 103 104 105 106 107 108 109 110 111 112 113 114 115 116 117 118 119 120 121 122 123 124 125 126 127
128 129 130 131 132 133 134 135 136 137 138 139 140 141 142 143 144 145 146 147 148 149 150 151 152 153
154 155 156 157 158 159 160 161 162 163 164 165 166 167 168 169 170 171 172 173 174 175 176 177 178 179
180 181 182 183 184 185 186 187 188 189 190 191 192 193 194 195 196 197 198 199 200 201 202 203 204 205
206 207 208 209 210 211 212 213 214 215 216 217 218;
  attachments ;
endTell;
checksumv 0 2 262862292 null;
endTell;
getWindow;
tell It;
  suppress;
  setResourcesState false false 100;
  setPopoutState 400;
  setExplanationSize 0;
  setAnnotationSize 0;
  setFontIncAnnot 0;
  setFontIncExp 0;
  setSize 1920 1079;
  setLocation 0 22;
  setFont SanSerif;
  setFontSize 10;
  getToolPalette;
  tell It;
  endTell;
  desuppress;
endTell;
getEmployee #mesquite.trees.ManageTrees.ManageTrees;
tell It;
  showTrees 0 #mesquite.lists.TreesList.TreesList;
  tell It;
    setTreeBlock 1;
    getWindow;
    tell It;
      newAssistant #mesquite.lists.NumForTreeList.NumForTreeList;
    tell It;
      suppress;
      setValueTask #mesquite.trees.NumberOfTaxa.NumberOfTaxa;
      desuppress;
    endTell;
    newAssistant #mesquite.lists.TreeListRooted.TreeListRooted;
    newAssistant #mesquite.lists.TreeListPolys.TreeListPolys;
    newAssistant #mesquite.lists.TreeListPolyAssumption.TreeListPolyAssumption;
    getTable;
    tell It;
      rowNamesWidth 327;
    endTell;
    setExplanationSize 30;
    setAnnotationSize 20;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 1820 1012;
```

```

        setLocation 0 22;
        setFont SanSerif;
        setFontSize 10;
        getToolPalette;
        tell It;
            setTool mesquite.lists.TreesList.TreesListWindow.ibeam;
        endTell;
    endTell;
    showWindow;
endTell;
showTreeBlocks #mesquite.lists.TreeblockList.TreeblockList;
tell It;
    getWindow;
    tell It;
        newAssistant #mesquite.lists.TreeblocksListNumber.TreeblocksListNumber;
        newAssistant #mesquite.lists.TreeblocksListTaxa.TreeblocksListTaxa;
        setExplanationSize 30;
        setAnnotationSize 20;
        setFontIncAnnot 0;
        setFontIncExp 0;
        setSize 1820 1012;
        setLocation 0 22;
        setFont SanSerif;
        setFontSize 10;
        getToolPalette;
        tell It;
            setTool mesquite.lists.lib.ListableVectorWindow.ibeam;
        endTell;
        setActive;
    endTell;
    showWindow;
endTell;
endTell;
getEmployee #mesquite.trees.BasicTreeWindowCoord.BasicTreeWindowCoord;
tell It;
    makeTreeWindow #5128558849605471043
#mesquite.trees.BasicTreeWindowMaker.BasicTreeWindowMaker;
    tell It;
        suppressEPCResponse;
        setTreeSource #mesquite.trees.StoredTrees.StoredTrees;
        tell It;
            setTreeBlock 1;
            toggleUseWeights off;
        endTell;
        setAssignedID 1293.1239345682121.8984554425152804462;
        getTreeWindow;
        tell It;
            popAsTile false;
            popOut;
            setExplanationSize 30;
            setAnnotationSize 20;
            setFontIncAnnot 0;
            setFontIncExp 0;
            setSize 1121 708;
            setLocation 3 34;
            setFont SanSerif;
            setFontSize 10;

```



```

getToolPalette;
tell It;
  setTool mesquite.trees.NodeNamer.NodeNamerExtra.nodeNamer;
endTell;
getTreeDrawCoordinator
#mesquite.trees.BasicTreeDrawCoordinator.BasicTreeDrawCoordinator;
tell It;
  suppress;
  setTreeDrawer #mesquite.trees.SquareTree.SquareTree;
  tell It;
    setNodeLocs #mesquite.trees.NodeLocsStandard.NodeLocsStandard;
    tell It;
      branchLengthsToggle off;
      toggleScale on;
      toggleBroadScale off;
      toggleCenter on;
      toggleEven on;
    endTell;
    setEdgeWidth 6;
    orientUp;
    toggleCorners off;
  endTell;
  setBackground White;
  setBranchColor Black;
  showNodeNumbers off;
  showBranchColors on;
  labelBranchLengths off;
  centerBrLenLabels on;
  showBrLensUnspecified on;
  showBrLenLabelsOnTerminals on;
  setBrLenLabelColor 0 0 255;
  setNumBrLenDecimals 6;
  desuppress;
  getEmployee #mesquite.trees.BasicDrawTaxonNames.BasicDrawTaxonNames;
  tell It;
    setColor Black;
    toggleColorPartition on;
    toggleShadePartition off;
    toggleShowFootnotes on;
    toggleNodeLabels on;
    toggleCenterNodeNames off;
    toggleShowNames on;
    namesAngle ?;
  endTell;
endTell;
  setTreeNumber 1;
  setDrawingSizeMode 0;
  toggleLegendFloat on;
  scale 0;
  toggleTextOnTree off;
  showWindow;
  newAssistant #mesquite.trees.TreeLegendMaker.TreeLegendMaker;
tell It;
  setOffsetsX 4;
  setOffsetsY 4;
  getLegendsVector;
  tell It;

```

```

    distributeCommands;
        setBounds 4 4 217 39;
        setOffsetX 4;
        setOffsetY 4;
    endDistributeCommands;
endTell;
newLegendItemNoCalc #mesquite.trees.TreeValueUsingMatrix.TreeValueUsingMatrix;
tell It;
    getEmployee #mesquite.parsimony.TreelengthForMatrix.TreelengthForMatrix;
    tell It;
        getEmployee #mesquite.parsimony.ParsCharSteps.ParsCharSteps;
        tell It;
            setModelSource
#mesquite.parsimony.CurrentParsModels.CurrentParsModels;
            endTell;
        endTell;
    getEmployee #mesquite.charMatrices.CharMatrixCoordIndep.CharMatrixCoordIndep;
    tell It;
        setCharacterSource #mesquite.charMatrices.StoredMatrices.StoredMatrices;
        tell It;
            setDataSet #8479452511181763361;
            endTell;
        endTell;
    endTell;
    calculate;
endTell;
endTell;
desuppressEPCResponse;
getEmployee #mesquite.trees.ColorBranches.ColorBranches;
tell It;
    setColor Red;
    removeColor off;
endTell;
getEmployee #mesquite.ornamental.BranchNotes.BranchNotes;
tell It;
    setAlwaysOn off;
endTell;
getEmployee #mesquite.ornamental.ColorTreeByPartition.ColorTreeByPartition;
tell It;
    colorByPartition off;
endTell;
getEmployee #mesquite.ornamental.DrawTreeAssocDoubles.DrawTreeAssocDoubles;
tell It;
    setOn on;
    toggleShow consensusFrequency;
    setDigits 4;
    writeAsPercentage off;
    toggleCentred on;
    toggleHorizontal on;
    setFontSize 10;
    setOffset 0 0;
endTell;
getEmployee #mesquite.trees.TreeInfoValues.TreeInfoValues;
tell It;
    panelOpen false;
endTell;
endTell;

```

```

endTell;
getEmployee #mesquite.charMatrices.BasicDataWindowCoord.BasicDataWindowCoord;
tell It;
  showDataWindow #8479452511181763361
#mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindowMaker;
tell It;
  getWindow;
  tell It;
    getTable;
    tell It;
      rowNamesWidth 110;
    endTell;
    setExplanationSize 30;
    setAnnotationSize 120;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 1820 912;
    setLocation 0 22;
    setFont SanSerif;
    setFontSize 10;
    getToolPalette;
    tell It;
      setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
    endTell;
    setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
    colorCells #mesquite.charMatrices.NoColor.NoColor;
    colorRowNames #mesquite.charMatrices.TaxonGroupColor.TaxonGroupColor;
    colorColumnNames #mesquite.charMatrices.CharGroupColor.CharGroupColor;
    colorText #mesquite.charMatrices.NoColor.NoColor;
    setBackground White;
    toggleShowNames on;
    toggleShowTaxonNames on;
    toggleTight off;
    toggleThinRows off;
    toggleShowChanges on;
    toggleSeparateLines off;
    toggleShowStates on;
    toggleAutoWCharNames on;
    toggleShowDefaultCharNames off;
    toggleConstrainCW on;
    toggleBirdsEye off;
    toggleAllowAutosize on;
    toggleColorsPanel off;
    toggleDiagonal on;
    setDiagonalHeight 80;
    toggleLinkedScrolling on;
    toggleScrollLinkedTables off;
  endTell;
  showWindow;
  getWindow;
  tell It;
    forceAutosize;
  endTell;
  getEmployee #mesquite.charMatrices.ColorCells.ColorCells;
  tell It;
    setColor Red;
    removeColor off;

```

```

endTell;
getEmployee #mesquite.categ.StateNamesStrip.StateNamesStrip;
tell It;
    showStrip off;
endTell;
getEmployee #mesquite.charMatrices.AnnotPanel.AnnotPanel;
tell It;
    togglePanel off;
endTell;
getEmployee #mesquite.charMatrices.CharReferenceStrip.CharReferenceStrip;
tell It;
    showStrip off;
endTell;
getEmployee #mesquite.charMatrices.QuickKeySelector.QuickKeySelector;
tell It;
    autotabOff;
endTell;
getEmployee #mesquite.stratigraphictools.ColorScaleCells.ColorScaleCells;
tell It;
    setColor Red;
    removeColor off;
endTell;
getEmployee #mesquite.categ.SmallStateNamesEditor.SmallStateNamesEditor;
tell It;
    panelOpen true;
endTell;
endTell;
endTell;
getEmployee #mesquite.charMatrices.ManageCharacters.ManageCharacters;
tell It;
    showCharacters #8479452511181763361 #mesquite.lists.CharacterList.CharacterList;
tell It;
    setData 0;
    getWindow;
tell It;
    newAssistant #mesquite.lists.DefaultCharOrder.DefaultCharOrder;
    newAssistant #mesquite.lists.CharListInclusion.CharListInclusion;
    newAssistant #mesquite.lists.CharListPartition.CharListPartition;
    newAssistant #mesquite.lists.CharacterStats.CharacterStats;
tell It;
    toggleSelectedOnly off;
endTell;
    newAssistant #mesquite.parsimony.CharListParsModels.CharListParsModels;
    newAssistant #mesquite.lists.CharListWeights.CharListWeights;
    setExplanationSize 30;
    setAnnotationSize 20;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 1820 1012;
    setLocation 0 22;
    setFont SanSerif;
    setFontSize 10;
    getToolPalette;
    tell It;
        setTool mesquite.lists.CharacterList.CharacterListWindow.ibeam;
    endTell;
endTell;
endTell;

```

```
        showWindow;  
        getEmployee #mesquite.lists.CharListAnnotPanel.CharListAnnotPanel;  
        tell It;  
            togglePanel off;  
        endTell;  
    endTell;  
endTell;  
endTell;  
end;
```

#NEXUS

[written Tue Apr 21 15:12:16 CEST 2009 by Mesquite version 2.6+ (build 487) at G5-de-David-Marjanovic.local/10.8.64.236]

BEGIN TAXA;

TITLE Taxa;

DIMENSIONS NTAX=6;

TAXLABELS

Notobatrachus Yizhoubatrachus Vieraella Mesophryne Ascaphus rest

;

END;

BEGIN CHARACTERS;

TITLE Character_Matrix;

DIMENSIONS NCHAR=2;

FORMAT DATATYPE = STANDARD GAP = - MISSING = ? SYMBOLS = " 0 1";

CHARSTATELABELS

1 'Pterygoid constricts subtemporal fenestra; generous interpretation of Ascaphus', 2 'Pterygoid constricts subtemporal fenestra; strict interpretation of Ascaphus' ;

MATRIX

Notobatrachus 11

Yizhoubatrachus 00

Vieraella 00

Mesophryne 11

Ascaphus 10

rest 00

;

END;

BEGIN TREES;

Title Trees;

LINK Taxa = Taxa;

TRANSLATE

1 Notobatrachus,

2 Yizhoubatrachus,

3 Vieraella,

4 Mesophryne,

5 Ascaphus,

6 rest;

TREE tree = ((1,2),(3,(4,(5,6))));

END;

BEGIN ASSUMPTIONS;

TYPESET * UNTITLED = unord: 1 - 2;

END;

BEGIN MESQUITECHARMODELS;

ProbModelSet * UNTITLED = 'Mk1 (est.): 1 - 2;

END;

Begin MESQUITE;

```

MESQUITESCRIPTVERSION 2;
TITLE AUTO;
tell ProjectCoordinator;
getEmployee #mesquite.minimal.ManageTaxa.ManageTaxa;
tell It;
    setID 0 3366549656872469409;
endTell;
getEmployee #mesquite.charMatrices.ManageCharacters.ManageCharacters;
tell It;
    setID 0 9176248464225086950;
    checksumv 0 2 888337455 null;
endTell;
getWindow;
tell It;
    suppress;
    setResourcesState false false 100;
    setPopoutState 400;
    setExplanationSize 0;
    setAnnotationSize 0;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 761 521;
    setLocation 8 22;
    setFont SanSerif;
    setFontSize 10;
    getToolPalette;
    tell It;
    endTell;
    desuppress;
endTell;
getEmployee #mesquite.trees.ManageTrees.ManageTrees;
tell It;
    showTreeBlocks #mesquite.lists.TreeblockList.TreeblockList;
    tell It;
        getWindow;
        tell It;
            newAssistant #mesquite.lists.TreeblocksListNumber.TreeblocksListNumber;
            newAssistant #mesquite.lists.TreeblocksListTaxa.TreeblocksListTaxa;
            setExplanationSize 30;
            setAnnotationSize 20;
            setFontIncAnnot 0;
            setFontIncExp 0;
            setSize 661 454;
            setLocation 8 22;
            setFont SanSerif;
            setFontSize 10;
            getToolPalette;
            tell It;
            endTell;
        endTell;
        showWindow;
    endTell;
endTell;
getEmployee #mesquite.trees.BasicTreeWindowCoord.BasicTreeWindowCoord;
tell It;
    makeTreeWindow #3366549656872469409
#mesquite.trees.BasicTreeWindowMaker.BasicTreeWindowMaker;

```

```

tell It;
  suppressEPCResponse;
  setTreeSource #mesquite.trees.StoredTrees.StoredTrees;
  tell It;
    setTreeBlock 1;
    toggleUseWeights off;
  endTell;
  setAssignedID 984.1239909960369.4011266357751987828;
  getTreeWindow;
  tell It;
    setExplanationSize 30;
    setAnnotationSize 20;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 661 454;
    setLocation 8 22;
    setFont SanSerif;
    setFontSize 10;
    getToolPalette;
    tell It;
      endTell;
      setActive;
      getTreeDrawCoordinator
#mesquite.trees.BasicTreeDrawCoordinator.BasicTreeDrawCoordinator;
    tell It;
      suppress;
      setTreeDrawer #mesquite.trees.SquareTree.SquareTree;
      tell It;
        setNodeLocs #mesquite.trees.NodeLocsStandard.NodeLocsStandard;
        tell It;
          branchLengthsToggle off;
          toggleScale on;
          toggleBroadScale off;
          toggleCenter on;
          toggleEven on;
        endTell;
        setEdgeWidth 6;
        orientUp;
        toggleCorners off;
      endTell;
      setBackground White;
      setBranchColor Black;
      showNodeNumbers off;
      showBranchColors on;
      labelBranchLengths off;
      centerBrLenLabels on;
      showBrLenUnspecified on;
      showBrLenLabelsOnTerminals on;
      setBrLenLabelColor 0 0 255;
      setNumBrLenDecimals 6;
      desuppress;
      getEmployee #mesquite.trees.BasicDrawTaxonNames.BasicDrawTaxonNames;
      tell It;
        setColor Black;
        toggleColorPartition on;
        toggleShadePartition off;
        toggleNodeLabels on;

```



```

toggleCenterNodeNames off;
toggleShowNames on;
namesAngle ?;
endTell;
endTell;
setTreeNumber 1;
setDrawingSizeMode 0;
toggleLegendFloat on;
scale 0;
toggleTextOnTree off;
showWindow;
newAssistant #mesquite.ancstates.TraceCharacterHistory.TraceCharacterHistory;
tell It;
suspend ;
setDisplayMode #mesquite.ancstates.ShadeStatesOnTree.ShadeStatesOnTree;
tell It;
toggleLabels off;
toggleGray off;
endTell;
setHistorySource #mesquite.ancstates.RecAncestralStates.RecAncestralStates;
tell It;
getCharacterSource #mesquite.charMatrices.CharSrcCoordObed.CharSrcCoordObed;
tell It;
setCharacterSource #mesquite.charMatrices.StoredCharacters.StoredCharacters;
tell It;
setDataSet #9176248464225086950;
endTell;
endTell;
setMethod #mesquite.parsimony.ParsAncestralStates.ParsAncestralStates;
tell It;
setModelSource #mesquite.parsimony.CurrentParsModels.CurrentParsModels;
toggleMPRsMode off;
endTell;
endTell;
setCharacter 2;
setMapping 1;
toggleShowLegend on;
toggleGray off;
toggleWeights on;
setInitialOffsetX -162;
setInitialOffsetY -183;
setLegendWidth 142;
setLegendHeight 170;
resume ;
endTell;
endTell;
desuppressEPCResponse;
getEmployee #mesquite.trees.ColorBranches.ColorBranches;
tell It;
setColor Red;
removeColor off;
endTell;
getEmployee #mesquite.ornamental.BranchNotes.BranchNotes;
tell It;
setAlwaysOn off;
endTell;
getEmployee #mesquite.ornamental.ColorTreeByPartition.ColorTreeByPartition;

```

```
tell It;
    colorByPartition off;
endTell;
getEmployee #mesquite.ornamental.DrawTreeAssocDoubles.DrawTreeAssocDoubles;
tell It;
    setOn on;
    toggleShow consensusFrequency;
    setDigits 4;
    writeAsPercentage off;
    toggleCentred on;
    toggleHorizontal on;
    setFontSize 10;
    setOffset 0 0;
endTell;
getEmployee #mesquite.trees.TreeInfoValues.TreeInfoValues;
tell It;
    panelOpen false;
endTell;
endTell;
endTell;
getEmployee #mesquite.charMatrices.BasicDataWindowCoord.BasicDataWindowCoord;
tell It;
    showDataWindow #9176248464225086950
#mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindowMaker;
tell It;
    getWindow;
    tell It;
        setExplanationSize 30;
        setAnnotationSize 20;
        setFontIncAnnot 0;
        setFontIncExp 0;
        setSize 661 454;
        setLocation 8 22;
        setFont SanSerif;
        setFontSize 10;
        getToolPalette;
        tell It;
            setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
        endTell;
        setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
        colorCells #mesquite.charMatrices.NoColor.NoColor;
        colorRowNames #mesquite.charMatrices.TaxonGroupColor.TaxonGroupColor;
        colorColumnNames #mesquite.charMatrices.CharGroupColor.CharGroupColor;
        colorText #mesquite.charMatrices.NoColor.NoColor;
        setBackground White;
        toggleShowNames on;
        toggleShowTaxonNames on;
        toggleTight off;
        toggleThinRows off;
        toggleShowChanges on;
        toggleSeparateLines off;
        toggleShowStates on;
        toggleAutoWCharNames on;
        toggleShowDefaultCharNames off;
        toggleConstrainCW on;
        toggleBirdsEye off;
        toggleAllowAutosize on;
```

```
toggleColorsPanel off;
toggleDiagonal on;
setDiagonalHeight 80;
toggleLinkedScrolling on;
toggleScrollLinkedTables off;
endTell;
showWindow;
getWindow;
tell It;
    forceAutosize;
endTell;
getEmployee #mesquite.charMatrices.ColorCells.ColorCells;
tell It;
    setColor Red;
    removeColor off;
endTell;
getEmployee #mesquite.categ.StateNamesStrip.StateNamesStrip;
tell It;
    showStrip off;
endTell;
getEmployee #mesquite.charMatrices.AnnotPanel.AnnotPanel;
tell It;
    togglePanel off;
endTell;
getEmployee #mesquite.charMatrices.CharReferenceStrip.CharReferenceStrip;
tell It;
    showStrip off;
endTell;
getEmployee #mesquite.charMatrices.QuickKeySelector.QuickKeySelector;
tell It;
    autotabOff;
endTell;
getEmployee #mesquite.stratigraphictools.ColorScaleCells.ColorScaleCells;
tell It;
    setColor Red;
    removeColor off;
endTell;
getEmployee #mesquite.categ.SmallStateNamesEditor.SmallStateNamesEditor;
tell It;
    panelOpen true;
endTell;
endTell;
endTell;
end;
```

#NEXUS

[written Tue Apr 21 15:34:46 CEST 2009 by Mesquite version 2.6+ (build 487) at G5-de-David-Marjanovic.local/10.8.64.236]

BEGIN TAXA;

TITLE Urodeles;

DIMENSIONS NTAX=9;

TAXLABELS

Karaurus Hynobiidae Cryptobranchidae Ambystomatidae Plethodontidae Salamandridae Proteidae
Amphiumidae Sirenidae

;

END;

BEGIN CHARACTERS;

TITLE Character_Matrix;

DIMENSIONS NCHAR=3;

FORMAT DATATYPE = STANDARD GAP = - MISSING = ? SYMBOLS = " 0 1 2 3";

CHARSTATELABELS

1 Dorsal_processes_of_premax / 'broad, low, indistinct dorsal process' 'alary process (broad, vaguely triangular process)' 'moderately high, vaguely rectangular process, or acutely triangular linked directly to base' 'narrow, long dorsal process, along the midline or parasagittal', 2 Number_of_premax_teeth / '>=10' '5-9' '<5', 3 Number_of_max_teeth / '>=30' '20-29' '<20' ;

MATRIX

Karaurus 100

Hynobiidae {1 3}1{1 2}

Cryptobranchidae 200

Ambystomatidae 200

Plethodontidae 310

Salamandridae 3(0 1)(0 1)

Proteidae 30?

Amphiumidae 312

Sirenidae 312

;

END;

BEGIN TREES;

Title Trees;

LINK Taxa = Urodeles;

TRANSLATE

1 Karaurus,

2 Hynobiidae,

3 Cryptobranchidae,

4 Ambystomatidae,

5 Plethodontidae,

6 Salamandridae,

7 Proteidae,

8 Amphiumidae,

9 Sirenidae;

TREE Alary_process_optimization = (1,((2,3),(9,((5,8),(7,(4,6))))));

END;

BEGIN ASSUMPTIONS;

TYPESET * UNTITLED = unord: 1 - 2, ord: 3;

END;

BEGIN MESQUITECHARMODELS;

ProbModelSet * UNTITLED = 'Mk1 (est.)': 1 - 3;

END;

Begin MESQUITE;

MESQUITESCRIPTVERSION 2;

TITLE AUTO;

tell ProjectCoordinator;

getEmployee #mesquite.minimal.ManageTaxa.ManageTaxa;

tell It;

setID 0 1289524600427350746;

endTell;

getEmployee #mesquite.charMatrices.ManageCharacters.ManageCharacters;

tell It;

setID 0 437647273697402900;

checksumv 0 2 2499153713 null;

endTell;

getWindow;

tell It;

suppress;

setResourcesState false false 100;

setPopoutState 400;

setExplanationSize 0;

setAnnotationSize 0;

setFontIncAnnot 0;

setFontIncExp 0;

setSize 1168 464;

setLocation 207 87;

setFont SanSerif;

setFontSize 10;

getToolPalette;

tell It;

endTell;

desuppress;

endTell;

getEmployee #mesquite.trees.BasicTreeWindowCoord.BasicTreeWindowCoord;

tell It;

makeTreeWindow #1289524600427350746

#mesquite.trees.BasicTreeWindowMaker.BasicTreeWindowMaker;

tell It;

suppressEPCResponse;

setTreeSource #mesquite.trees.StoredTrees.StoredTrees;

tell It;

setTreeBlock 1;

toggleUseWeights off;

endTell;

setAssignedID 1841.1239624217592.7879340418447371515;

getTreeWindow;

tell It;

setExplanationSize 30;

setAnnotationSize 20;

setFontIncAnnot 0;

setFontIncExp 0;

```

setSize 1068 397;
setLocation 207 87;
setFont SanSerif;
setFontSize 10;
getToolPalette;
tell It;
    setTool mesquite.trees.BasicTreeWindowMaker.BasicTreeWindow.arrow;
endTell;
setActive;
getTreeDrawCoordinator
#mesquite.trees.BasicTreeDrawCoordinator.BasicTreeDrawCoordinator;
tell It;
    suppress;
    setTreeDrawer #mesquite.trees.SquareTree.SquareTree;
    tell It;
        setNodeLocs #mesquite.trees.NodeLocsStandard.NodeLocsStandard;
        tell It;
            branchLengthsToggle off;
            toggleScale on;
            toggleBroadScale off;
            toggleCenter on;
            toggleEven on;
        endTell;
        setEdgeWidth 6;
        orientUp;
        toggleCorners off;
    endTell;
    setBackground White;
    setBranchColor Black;
    showNodeNumbers off;
    showBranchColors on;
    labelBranchLengths off;
    centerBrLenLabels on;
    showBrLensUnspecified on;
    showBrLenLabelsOnTerminals on;
    setBrLenLabelColor 0 0 255;
    setNumBrLenDecimals 6;
    desuppress;
    getEmployee #mesquite.trees.BasicDrawTaxonNames.BasicDrawTaxonNames;
    tell It;
        setColor Black;
        toggleColorPartition on;
        toggleShadePartition off;
        toggleNodeLabels on;
        toggleCenterNodeNames off;
        toggleShowNames on;
        namesAngle ?;
    endTell;
endTell;
setTreeNumber 1;
setDrawingSizeMode 0;
toggleLegendFloat on;
scale 0;
toggleTextOnTree off;
showWindow;
newAssistant #mesquite.ancstates.TraceCharacterHistory.TraceCharacterHistory;
tell It;

```

```

suspend ;
setDisplayMode #mesquite.ancstates.ShadeStatesOnTree.ShadeStatesOnTree;
tell It;
    toggleLabels off;
    toggleGray off;
endTell;
setHistorySource #mesquite.ancstates.RecAncestralStates.RecAncestralStates;
tell It;
    getCharacterSource #mesquite.charMatrices.CharSrcCoordObed.CharSrcCoordObed;
    tell It;
        setCharacterSource #mesquite.charMatrices.StoredCharacters.StoredCharacters;
        tell It;
            setDataSet #437647273697402900;
        endTell;
    endTell;
    setMethod #mesquite.parsimony.ParsAncestralStates.ParsAncestralStates;
    tell It;
        setModelSource #mesquite.parsimony.CurrentParsModels.CurrentParsModels;
        toggleMPRsMode off;
    endTell;
endTell;
setCharacter 1;
setMapping 1;
toggleShowLegend on;
toggleGray off;
toggleWeights on;
setInitialOffsetX -142;
setInitialOffsetY -199;
setLegendWidth 142;
setLegendHeight 186;
resume ;
endTell;
endTell;
desuppressEPCResponse;
getEmployee #mesquite.trees.ColorBranches.ColorBranches;
tell It;
    setColor Red;
    removeColor off;
endTell;
getEmployee #mesquite.ornamental.BranchNotes.BranchNotes;
tell It;
    setAlwaysOn off;
endTell;
getEmployee #mesquite.ornamental.ColorTreeByPartition.ColorTreeByPartition;
tell It;
    colorByPartition off;
endTell;
getEmployee #mesquite.ornamental.DrawTreeAssocDoubles.DrawTreeAssocDoubles;
tell It;
    setOn on;
    toggleShow consensusFrequency;
    setDigits 4;
    writeAsPercentage off;
    toggleCentred on;
    toggleHorizontal on;
    setFontSize 10;
    setOffset 0 0;

```

```

    endTell;
    getEmployee #mesquite.trees.TreeInfoValues.TreeInfoValues;
    tell It;
        panelOpen false;
    endTell;
endTell;
endTell;
getEmployee #mesquite.charMatrices.BasicDataWindowCoord.BasicDataWindowCoord;
tell It;
    showDataWindow #437647273697402900
#mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindowMaker;
tell It;
    getWindow;
    tell It;
        setExplanationSize 30;
        setAnnotationSize 20;
        setFontIncAnnot 0;
        setFontIncExp 0;
        setSize 1068 397;
        setLocation 207 87;
        setFont SanSerif;
        setFontSize 10;
        getToolPalette;
        tell It;
            setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
        endTell;
        setTool mesquite.charMatrices.BasicDataWindowMaker.BasicDataWindow.ibeam;
        colorCells #mesquite.charMatrices.NoColor.NoColor;
        colorRowNames #mesquite.charMatrices.TaxonGroupColor.TaxonGroupColor;
        colorColumnNames #mesquite.charMatrices.CharGroupColor.CharGroupColor;
        colorText #mesquite.charMatrices.NoColor.NoColor;
        setBackground White;
        toggleShowNames on;
        toggleShowTaxonNames on;
        toggleTight off;
        toggleThinRows off;
        toggleShowChanges on;
        toggleSeparateLines off;
        toggleShowStates on;
        toggleAutoWCharNames on;
        toggleShowDefaultCharNames off;
        toggleConstrainCW on;
        toggleBirdsEye off;
        toggleAllowAutosize on;
        toggleColorsPanel off;
        toggleDiagonal on;
        setDiagonalHeight 80;
        toggleLinkedScrolling on;
        toggleScrollLinkedTables off;
    endTell;
    showWindow;
    getWindow;
    tell It;
        forceAutosize;
    endTell;
    getEmployee #mesquite.charMatrices.ColorCells.ColorCells;
    tell It;

```



```

        setColor Red;
        removeColor off;
    endTell;
    getEmployee #mesquite.categ.StateNamesEditor.StateNamesEditor;
    tell It;
        makeWindow;
        tell It;
            setExplanationSize 30;
            setAnnotationSize 20;
            setFontIncAnnot 0;
            setFontIncExp 0;
            setSize 1068 397;
            setLocation 207 87;
            setFont SanSerif;
            setFontSize 10;
            getToolPalette;
            tell It;
                setTool mesquite.categ.StateNamesEditor.StateNamesWindow.ibeam;
            endTell;
            rowsAreCharacters on;
            toggleConstrainChar on;
            toggleConstrainCharNum 3;
            togglePanel off;
            toggleSummaryPanel off;
        endTell;
        showWindow;
    endTell;
    getEmployee #mesquite.categ.StateNamesStrip.StateNamesStrip;
    tell It;
        showStrip off;
    endTell;
    getEmployee #mesquite.charMatrices.AnnotPanel.AnnotPanel;
    tell It;
        togglePanel off;
    endTell;
    getEmployee #mesquite.charMatrices.CharReferenceStrip.CharReferenceStrip;
    tell It;
        showStrip off;
    endTell;
    getEmployee #mesquite.charMatrices.QuickKeySelector.QuickKeySelector;
    tell It;
        autotabOff;
    endTell;
    getEmployee #mesquite.stratigraphictools.ColorScaleCells.ColorScaleCells;
    tell It;
        setColor Red;
        removeColor off;
    endTell;
    getEmployee #mesquite.categ.SmallStateNamesEditor.SmallStateNamesEditor;
    tell It;
        panelOpen true;
    endTell;
endTell;
endTell;
getEmployee #mesquite.charMatrices.ManageCharacters.ManageCharacters;
tell It;
    showCharacters #437647273697402900 #mesquite.lists.CharacterList.CharacterList;

```

```
tell It;
  setData 0;
  getWindow;
  tell It;
    newAssistant #mesquite.lists.DefaultCharOrder.DefaultCharOrder;
    newAssistant #mesquite.lists.CharListInclusion.CharListInclusion;
    newAssistant #mesquite.lists.CharListPartition.CharListPartition;
    newAssistant #mesquite.stochchar.CharListProbModels.CharListProbModels;
    newAssistant #mesquite.parsimony.CharListParsModels.CharListParsModels;
    setExplanationSize 30;
    setAnnotationSize 20;
    setFontIncAnnot 0;
    setFontIncExp 0;
    setSize 1068 397;
    setLocation 207 87;
    setFont SanSerif;
    setFontSize 10;
    getToolPalette;
    tell It;
      endTell;
    endTell;
  showWindow;
  getEmployee #mesquite.lists.CharListAnnotPanel.CharListAnnotPanel;
  tell It;
    togglePanel off;
  endTell;
endTell;
endTell;
end;
```